

A photograph showing two hands, palms up, holding a small, green, leafy tree against a light background. The hands are positioned on either side of the tree, with fingers slightly curled as if supporting it. The background is a soft, out-of-focus light green and yellow.

**BIODIVERSITY CONSERVATION OF  
SOUTHERN EASTERN GHATS,  
ANDHRA PRADESH.**

**Dr. S. K. M.Basha  
P. Siva Kumar Reddy  
M. John Paul**



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# BIODIVERSITY CONSERVATION OF PULICATLAKE

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## BIOPLASTICS AND THE DAWN OF NEW MATERIALS- A REVIEW

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**“Bioplastics are the workhorse material of the modern economy”**

### ABSTRACT

Plastics that have been universally used in our daily lives are now causing serious environmental problems. Petroleum derived plastics dominate the packaging industry even today. It accounts for the largest usage of plastics worldwide and is used in numerous applications. Millions of tons of these non-degradable plastics accumulate in the environment each year. Bioplastics are a suitable alternative to petroleum-based plastics. They are highly complex and sophisticated materials that can help make plastic products more sustainable and eco-friendly. Bioplastics today are a large family of materials with differing properties and applications. The review is an attempt to investigate the facts about different types of bioplastics, their current status, major advantages and other related issues.

Key words: Bio-plastics, Petro-plastics, Sustainability

### INTRODUCTION:

Plastics have become a vital part of our life. The environmental crisis arising from the use of petroleum based non-degradable plastics led the globe to find alternatives. Biomaterial based plastic is one such alternative. They comprise bio-based materials featuring identical properties compared to their fossil-based versions as well as new materials featuring additional properties. These add-on qualities like biodegradability and bio-based property can reduce the impact on environment significantly. Moreover, the emission of greenhouse gases during production and degradation of bioplastics is very low when compared to conventional plastics.

### BIOPLASTICS: THE NOVEL MATERIALS

Bioplastics are novel materials of 21st century and would be of great importance to the material world (Mohanty et al., 2002). The keen interest in bioplastics started way back in early 20th century when Henry Ford used corn and soybean oils to manufacture his automotive parts (Stevens E.S., 2003). Bioplastics are made from a number of renewable resources such as plant oils, cellulose, starches, sugars, carbohydrates, bacteria and algae. Bioplastics are not single substances; they comprise of a whole family of materials with differing properties and applications. According to European Bioplastics, a plastic material is defined as a bioplastic, if it is either bio-based or biodegradable or features both the properties (Eu.Bio., 2016). Bioplastics are eco-friendly alternative to traditional plastics and are extracted from renewable resources like corn, sugarcane, cellulose, potato or starch. These are 100% degradable, equally resistant and versatile, already used in agriculture, textile industry, medicine and for packaging.

### CLASSIFICATION OF BIOPLASTICS:

The world of bioplastics has exploded, and there is an amazing breadth of materials that can be classified as bioplastics. The term bioplastic refers to several groups of plastics namely bio-based plastics (from renewable resources) and bio-degradable including compostable plastics.

#### Biobased plastics

Biobased plastics are those that are derived from plant-based polymers, such as corn starch, sugarcane or cellulose, potatoes, rice, soy, wheat and vegetable oil and are not derived from petroleum resources. The Business-NGO (non-government organization) Working

Group for Safer Chemicals and Sustainable Materials defines bio-based bioplastics as “plastics in which 100% of the carbon is derived from renewable agricultural and forestry resources such as corn starch, soybean protein and cellulose” (Alvarezetal.,2011)

### **Biodegradable bioplastics**

Biodegradable bioplastics are fully degraded by microorganisms without leaving any visible toxic remainders. The term “biodegradable” refers to materials that can disintegrate naturally into biogas and biomass (mostly carbon dioxide and water) as a result of being exposed to a microbial environment and humidity (Alvarezetal.,2011)

### **Compostable bioplastics**

Compostable bioplastics will biodegrade in a compost site. Microorganisms break it down into carbon dioxide, water, inorganic compounds and biomass leaving no toxic residues. The most commonly used raw material is corn starch, which is converted into a polymer with similar properties to traditional polyethylene plastic products. Other compostable plastics available are made from potato starch, soya bean protein, cellulose as well as from petroleum and its by products. This shows that compostable plastics may be derived from both plantbased and petroleumderived polymers.

### **TYPES OF BIOPLASTICS:**

Bioplastics are currently used in disposable items like packaging, containers, straws, bags and bottles, and in non-disposable carpets, plastic piping, phone casings, 3-D printing, car insulation and medical implants.

#### **A. Starch based plastics**

Starch is the key raw material for bioplastics which is found in seeds and in tubers or roots of the plants. Most of the starch produced worldwide is derived from corn (Asaf,2008). Today, thermoplastic starch, accounting for about 50% to 80% of the global bioplastics market, is the most significant and widely used bioplastic. Applications of thermoplastic starch are bags, yogurt tubs, cups, plant pots, cutlery, diaper foil, coated paper and cardboard. **Poly(lactic acid) (PLA):** Poly(lactic acid) is the polymer with the highest potential for a commercial production as bioplastic. PLA is a 100% bio-based plastic that is currently being used in packaging applications. Poly(lactic acid), the secondmost important bioplastic of the world in regard to consumption in volume (Garima et al). It is derived from renewable resources like corn starch or sugar cane. PLA is biodegradable and suitable for the manufacture of compostable packaging products.

**ii. PSM (Plastarch Material):** PSM is a biodegradable, thermoplastic resin. It is composed of starch combined with several other biodegradable materials. PSM is currently used for a wide variety of applications in the plastic market, such as food packaging and utensils, personal care items, plastic bags, temporary construction tubing, industrial foam packaging, industrial and agricultural film, window insulations, construction stakes, and in horticulture (GGP,2009-10)

**iii. MATER-BI :** MATER-BI is biodegradable and compostable bioplastic synthesized mainly by starches, cellulose, vegetable oils and their combinations. BIO BIG is the world’s largest brand of certified compostable bags and films made from Mater-BI (NOVAMONT)

**B. Cellulose based Bioplastics:** Produced using cellulose esters and cellulose derivatives.

**i. Cellophane:** Cellophane is made from cellulose which is biodegradable. Cellophane films can be coloured and are well known as candy wrappings. The biodegradable films are available in a wide range of grades, and they can be used to pack products ranging from cheese to coffee and chocolates (O.J.P.C.,2018).

**ii. Cellulose acetate:** Cellulose acetate is thermoplastic which is rather expensive and rarely used in packaging applications.

**iii. Polybutylene succinate (PBS):** Biodegradable, particularly used in packaging industry. Other applications include disposable products such as tableware or medical articles.

In agriculture, PBS finds interest in the fabrication of mulching films or delayed release of materials for pesticides and fertilizers.

### C. Microbial synthesis:

**i. Poly hydroxybutyrate (PHB):** Poly hydroxybutyrate (PHB) is biodegradable; used in a wide variety of fields including packaging, ropes, bank notes and car parts. PHB has potential applications in a wide variety of fields such as industrial, biomedical, agricultural, domestic, and automobile.

**ii. Polyhydroxyalkanoate (PHA):** PHAs are polyesters produced in nature by numerous microorganisms through bacterial fermentation of sugar or lipids (Jingnanetal,2009). They are biodegradable and are used in the production of bioplastics. PHA is often used for medical applications such as sutures, slings, bone plates and skin substitutes; it is also used for single-use food packaging.

### D. Synthesis from oils:

**Polyamides 11: PA11** a biopolymer derived from natural oil known by the trade name Rilson B commercialized by Arkoma. It is prized for its thermal resistance that makes it valued for use in car fuel lines, pneumatic air brake tubing, electrical anti-termite cable sheathing and oil and gas flexible pipes and control fluid umbilicals(Nifisaetal.,2015).It is non biodegradable.

### E. Other Biopolymers :

**Polyethylene(PE):** Polyethylene (PE) is a bio-based polymer generally extracted from sugar used in carry bags, films and bottles.

**Polyethylene terephthalate(PET):** Polyethylene terephthalate (PET) is one of the major polymers produced worldwide representing about 18 % of world polymer production and comes in third after Polyethylene and Polypropylene. It is known for its use in beverage bottles.Coca Cola introduced Plant Bottle Technology where the PET had been made from bio-based mono-ethylene glycol from sugarcane and terephthalic acid from petrochemicals.

**Polyglycolic acid (PGA):** This is used in medicine and specialized applications.

**Polyhydroxyvalerate (PHBV):** Used for films and paper coatings, with possible markets including biomedical applications and veterinary science.

**Polyvinyl alcohol (PHV):** Used in packaging designed to dissolve in water to release products such as laundry detergent, pesticides, and hospital washables.

**Polycaprolactone(PCL):** Biodegradable polyester, with low melting temperature and easily biodegradable. Widely used for mulch, seeding containers and biomedical applications.

**Poly butyrate adipate terephthalate (PBAT):** PBAT is a biodegradable and compostable biopolymer and is known for its flexibility and toughness which makes it ideal for combination with other biodegradable polymers that are brittle with high modulus and strength. Widely used in garbage bags, wrapping films, disposable plastic products.

**Polyethylene furanoate(PEF)** PEF, a new polymer that is expected to enter the market in 2020. PEF is made entirely from vegetable raw materials and is also recyclable. It is considered to be the packaging material of the future, particularly for food and beverages. PEF finds applications in the packaging industry for fruit juices, milk, soft drinks, fresh tea or water.

**Bioplastics packaging:** Bioplastic packaging is being used extensively in a wide array of industries including textiles, consumer goods, automotive and transport sectors. In 2018, global production capacities of bioplastics amounted to about 2.11 million tons with almost 65 percent of the volume destined for the packaging market, the biggest market segment within the bioplastics industry (Euro.Bio). There is a high demand for packaging made from bioplastics to be used for wrapping organic foods as well as for premium and branded products with a particular requirement. All types of bioplastics used in packaging offer one

Advantage over fossil-based products is the usage of renewable resources to overcome the environmental impact. Bio-PET and PLA are the major types of bioplastics for packaging around the world.

**Advantages of Bioplastics:** Bioplastic is a much younger industry than conventional oil plastics. Bio-based plastics often shows good performance and a better environmental footprint when compared to conventional plastics. They have a less carbon footprint helps to save resources and reduce crude oil dependency. Bioplastics can help the plastic industry and plastics consuming brands grow more sustainably. Bioplastics and bioeconomy overall have the potential to unite the agricultural, chemical and industrial sectors. Bioplastics and bio-economy also help to create new jobs elsewhere both within industry as well as in the entire value chain.

**Disadvantages of Bioplastics:** Although bioplastics show promising potential as alternative to conventional plastics, it is equally important to consider their drawbacks. Many people believe that bioplastics are biodegradable. This is only partially true, as some bioplastics are biodegradable while others are not (Grabianowski, 2018). Surprisingly, many bioplastics won't degrade at all, if placed in landfill with other garbage. Although bioplastics don't result in fossil-fuel emissions, they do require large amounts of fertilizers and pesticides (Grabianowski, 2018). More over Bioplastics are basically designed to be composted, not recycled. Another drawback of bioplastics is their indistinguishability. To overcome this users must be educated about bioplastics and proper recycling techniques to discern which bioplastics are biodegradable and which are compostable (Creative Mechanisms Staff, 2017). Further, most bioplastics require high temperatures to degrade, and most cities lack the infrastructure for proper bioplastic composting. As a result, many bioplastics end up in the trash, are deprived of oxygen, and release harmful methane into the environment (Creative Mechanisms Staff, 2017). Waste collection and resource management needs to be standardized along with the packaging industry to create some synchronized systems for waste to resource management. Bioplastics are often produced from genetically modified food crops such as corn, potatoes and soybeans, a practice that carries a high risk of contaminating our food supply. Also, corn and soybean producers typically apply large amounts of chemical pesticides and fertilizers that pollute our air and water.

#### **Innovations in the field of Bioplastics continues –**

- Researchers around the world continue to develop greener varieties of Bioplastics that can be more effectively reduce plastic pollution and the carbon footprint.
- More durable bioplastics are being designed for automotive, electronics and consumer goods such as Electrolux's new refrigerator made from corn and sugarcane bioplastic material.
- Newer bioplastics are being made in laboratories from straw, wood chips and food waste.
- Bioplastics can be genetically engineered from *Pseudomonas* by the mutation of some of the genes involved in the  $\beta$ -oxidation pathway.

**Conclusion:** Even though the production of bioplastics is costly, their applications are of interest owing to the drawbacks of conventional plastics. Industrial progress in packaging technology in future appears to be moving towards newer breed of bio-materials. The trend is to maximize the production of bioplastics by enhancing the properties of bioplastics producing microorganisms, plants and algae. It is clear that bioplastics can reduce many of the environmental problems posed by the conventional plastics.

**REFERENCES:**

- Alvarez-Chavez, C. R., Edwards, S., Moure-Eraso, R. L., and Geiser, K. 2011. "Sustainability of Bio-based Plastics: General Comparative Analysis and Recommendations for Improvement." *Journal of Cleaner Production* 23 (1): 46-7.
- Asaf Kleopas, S. (2008) *Synthesis and Properties of Starch Based Bio-Materials*. University of Groningen,
- Cho, Renee. "The Truth About Bioplastics" *State of the Planet*, 15 Dec. 2017. [blogs.ei.columbia.edu/2017/12/13/the-truth-about-bioplastics/](http://blogs.ei.columbia.edu/2017/12/13/the-truth-about-bioplastics/)
- Dr. Garima Goswami, Sudhanshu Joshi, Ujjawal Sharma: *International Journal of Engineering Research & Technology (IJERT)*; ISSN: 2278-0181, [www.ijert.org](http://www.ijert.org); ETRASCT' 14 Conference Proceedings
- European Bioplastics (a), 2016. <http://en.european-bioplastics.org/bioplastics/> (Date of Consult: 10.01.2016).
- GGP-Library, 2009-10 : Bioplastics
- Grabianowski, Ed. "What is the future of Bioplastics?" *How Stuff Works. Science, How Stuff Works*, 8 Mar. 2018. [Science.howstuffworks.com/environmental/green-science/future-of-bioplastics.html](http://Science.howstuffworks.com/environmental/green-science/future-of-bioplastics.html)
- Lu, Jingnan; Tappel, Ryan C.; Nomura, Christopher T. (2009-08-05). "Mini-Review: Biosynthesis of Poly(hydroxyalkanoates)". *Polymer Reviews*. 49 (3): 226-248. doi:10.1080/15583720903048243. ISSN 1558-3724.
- Mohanty, A. K., Misra, M., and Drzal, L.T (2002): "Sustainable Bio-composites from Renewable Resources: Opportunities and Challenges in the Green Materials World." *Journal of Polymers and the Environment* 10 (1-2): 19-26.
- Nafisa Jabeen, Ishrat Majid & Gulzar Ahmad Nayik | Fatih Yildiz (2015): REVIEW FOR POLYAMIDES: Bioplastics and food packaging: A review, Accepted paper, 21 Oct 2015, Article: 1117749
- Novamont.com/northamerica/page=74&idfirst=74
- Open Journal of Polymer Chemistry, 2018, 8, 21-33 [http:// www.scirp.org/journal/ojchem](http://www.scirp.org/journal/ojchem)  
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- Stevens, E.S. (2003): 'What makes green plastics green', *Bi oCycle*, 44(3), 24,4

**BIODEGRADABLE PLASTICS –AN OVERVIEW**

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**ABSTRACT**


Plastics are synthetic or semi-synthetic materials which are typically polymers of high molecular mass obtained from petroleum and natural gas. Bioplastic is the universal term for polymers made of renewable biomass sources. Bioplastics are an alternative to traditional plastics. As a consequence, bioplastic is biodegradable in about 180 days in opposition to traditional plastics, which remain in the environment for 500 to 1000 years. Bioplastics are made of different sources of biomass, such as corn or potato starch and cellulose. These substances are naturally produced by plants. Biodegradable plastics are the type of plastics that undergo decomposition over a period of time under composting conditions. The global biodegradable plastics market accounts for less than 1% of the overall plastics market, however, it is expected to grow at a fast pace over the next 5 years. Organic



**DEPARTMENT OF ZOOLOGY**

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# **MEDICINAL PLANTS RESEARCH**

**INDIRA PRIYADARSINI A  
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# Impact of Electronic Resources on Academics

(Festschrift in Honour of Prof. V. Pulla Reddy)

Editors:

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**Use and effectiveness of E-ShodhSindhu in promotion of higher education among the research scholar faculty members of Dravidian University: A Study**

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**Abstract** - The present study aims to analyse the use and effectiveness of E-ShodhSindhu consortium e-journals in promotion of higher education among the faculty members and research scholars of Dravidian University. The study also intends to determine the frequency of use of E-ShodhSindhu, the types of e-journals used in E-ShodhSindhu, access place for E-ShodhSindhu, types of e-portals used, effectiveness of E-ShodhSindhu on academic activities, factors affecting the use of E-ShodhSindhu and overall satisfaction with E-ShodhSindhu. Questionnaire was adopted to elicit data from faculty and research scholars of Dravidian University. A total 110 respondents participated in the survey from which 60 are faculty members and 48 are research scholars. The study reveals that faculty members are using E-ShodhSindhu online databases subscribed by Dravidian University. Majority of faculty and research scholars are using E-ShodhSindhu for research purpose and the department is the most preferred location for accessing e-journals.

**Keywords:** E-ShodhSindhu, INFLIBNET, E-journals, e-resources, databases, Dravidian University

**Introduction**

The recent developments in Information Technology change the scenario. Libraries and Information Centres have embraced information technology more profoundly than many other institutions. Most of them are currently using electronic products and services which offers tremendous opportunities to provide solutions to some of the challenges now libraries are facing. Electronic Publishing or

### Conclusion

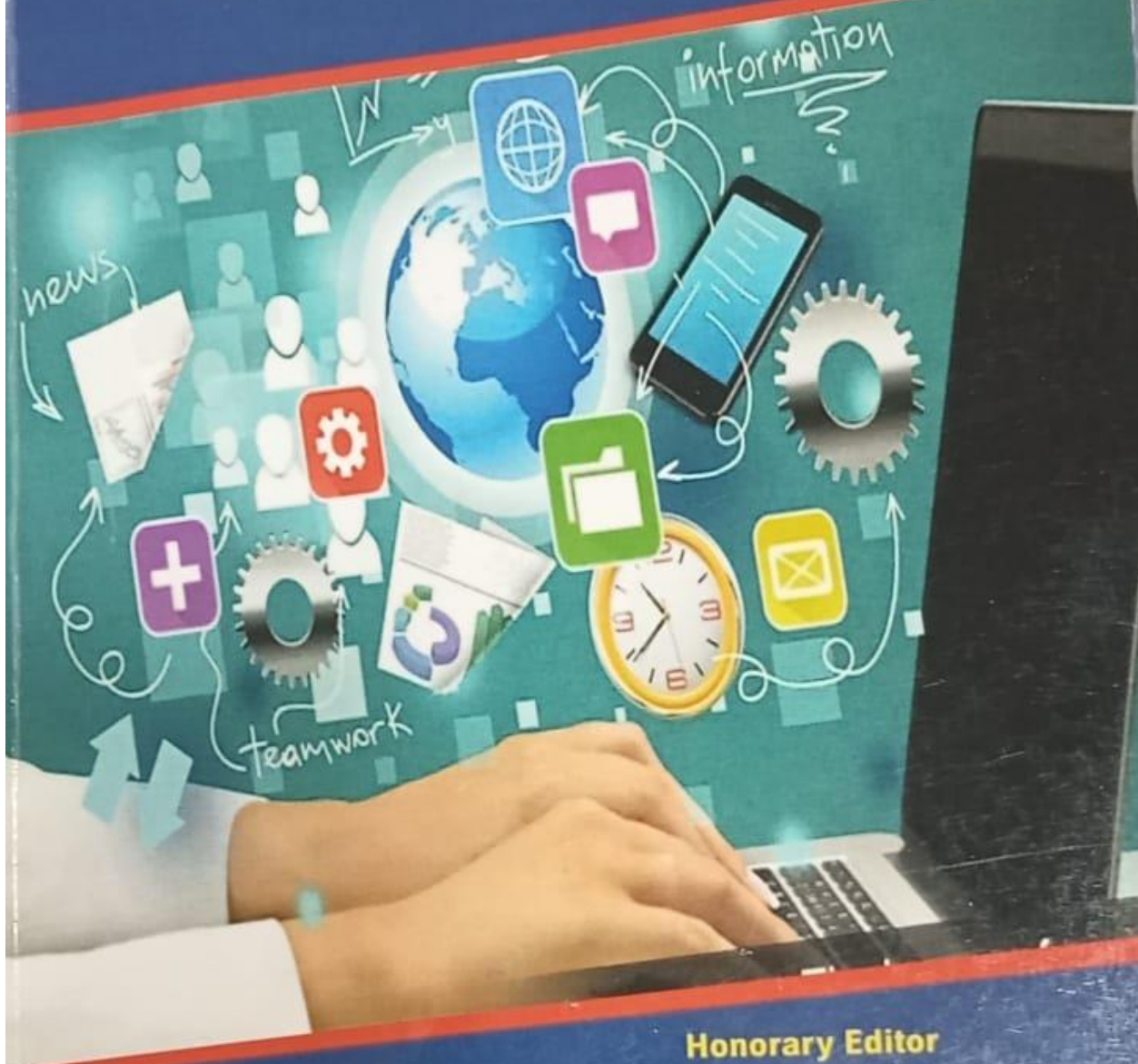
The study reveals that the faculty members and research scholars of Dravidian University mostly rely on E-ShodhSindhu e-journals for research purpose and publishing articles. They are using the available e-journals satisfactorily. The library professionals tried their best for promotion, assistance and guidance in accessing e-journals. The increase in e-journals subscription indicates that the library has made drastic changes in E-ShodhSindhu e-journals collection and services. The library has to employ different strategies for effective use of available online resources through E-ShodhSindhu. Central Library has to conduct the orientation programme for the best use of E-ShodhSindhu e-resources.

### References

1. Archita Nanda (2017). Use and Awareness of E-Journals by the Faculty and Research Scholars of Veer Surendra Sai University of Technology. *DESIDOC Journal of Library & Information Technology*, 37(4),274-280 .
2. Chauhan, Vasantray A (2017). E-Shodhsindhu Consortia: A Boon to User of Indian Academic Libraries. *Indian Journal of Information Sources and Services*. 7(2),34-36.
3. Karan Singh. (2017). E-Shodh Sindhu: Consortium For Higher Education Electronic Resources For Scholarly Content: An Overview. *International Journal of Advanced Multidisciplinary Research*, 4(3), 43-50.
4. Lingaiah, Vuppala (2016). A Study on use of E-Shodhsindhu Resources by the Scholars and Faculty Members in JNTU. *Journal of Advances in Library and Information Science*, 5(3), 268-272.
5. Praveenkumar Kumbargoudar (2017). Knowledge and Use of E-Shodhsindhu Consortium in Karnataka University, Dharwad. *E-Library Science Research Journal*, 5 (3), 1-6.
6. <http://www.dravidianuniversity.ac.in/central-library.php>
7. <http://ess.inflibnet.ac.in/about.php>



# User Perceptions and Expectations from Academic Libraries in Digital Information Society



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## Use of Social Media in Library Services

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*Abstract - Traditional Libraries gives emphasis on storage and preservation of physical items, particularly books and periodicals those in which librarian were a custodian of the library. In the past years, libraries have increasingly developed into a provider of information resources and services that do not even require a building. It is about new demands of competence within education, trade and industry, the public's demand for new services, new media and information resources, the range of possibilities offered by the Internet, digital services etc. The main aim users ,it will help in discover user's need for effective and efficient library services. This paper is to focus on creatively engage in marketing library services through social media. Now a days for excellent library services social media has become an integral part. Libraries should design a workable plan with total emphasis on the delivery.*

**Keywords:** Social Media, Library Services.

### 1. Introduction

Social Media deliver several opportunities to reach and interact with the society. In recent years social media and social network have grown amazingly. Social Media is being used all over world for manifold purposes in Libraries and Knowledge centre. Currently libraries and information centres are facing many problems and challenges. It is very difficult for library professionals to manage and share their resources with others manually. For solving these problems, many technological development has been discovered and creating new forms of information, new sources of information and new ways of providing information bypassing traditional institutional like libraries.

Now, the actual challenge for information professionals is not to handle the collection, staff and technology, but to turn these resources into services which are user oriented. Web 2.0 applications in libraries have acquired growing reputation all over world. It appears that the library must think about marketing its services more regularly through the internet, taking advantage of Social networks to get better access to its users and to promote information services. Social media is a powerful new form of communication.

## 2. Objectives of the Study

The study seeks to achieve the following objectives:

1. To investigate the problems in utilizing social media for marketing purposes in libraries.
  2. To promote the library services
  3. Review actual study on the Utilization of Social media platforms in library and information centres.
3. Social Media

Social media is the collective of online communications channels dedicated to community-based input, interaction, content-sharing and collaboration. Social Media can be described as a group of web-based and mobile applications that allow users to share and create knowledge in a real time social interaction. It is user-centric, multi-purpose and it is not time and location bound. Social Media consists of various users driven marketing channels, e.g. Face book, Twitter, Blogs, YouTube, Flickr, Pinterest. Social Media provides more opportunities to reach the user community target specific audiences and give users a chance to interact with Library. Libraries can market their services and products using different social media platforms, for Example publicize their different upcoming events and newly acquired information materials through the Face book. Different programs such as conference and workshops can be marketed by uploading videos on the YouTube. The pictures of different library events and services can be shared using Flickr. Blogs can be used to market library services among distance learners. Twitter and Instance messaging can be used to market a Library's Reference. Quora can be used for A Questions and Answers. Library services the most widely used social media platforms follow.

### 3.1 Face book

Face book is a social networking website intended to connect friends, family, and business associates. It is the largest of the networking sites,

with the runner up being My Space. Face book the largest number of active users and is easy to get started with. When looking for a business or information Face book is one of the first places many people look. It is a place for community members to connect online. Therefore every library in the Peace Library System should have a Face book page. Through the face book page / Accounts librarians can draw user's attention to useful hidden treasures of the library that library users have forgotten or are not aware of such a grey literature.

### 3.2 Twitter

Twitter is second most popular social media platform. It is micro blogging network of real posts and all post is limit to 140 characters or less. Twitter can be used to keep library staff and patrons updated on library's daily activities for exam frequently updated library collections. User can utilize this platform to type in short messages or status updates. Often library users prefer twitter to interact with librarians because this is more influential than other social media platforms and what happens on Twitter does not stay in twitter. Twitter is a marketing tool for libraries including tips, best practices, evaluation and assessment of a twitter account.

### 3.3 Blog

A blog is a discussion or informational website published on the World Wide Web consisting of discrete, often informal diary-style text entries ("posts"). Posts are typically displayed in reverse chronological order, so that the most recent post appears first, at the top of the web page. Blogs are popularly used in library to broadcast library news and market other library resources. Librarians can also develop subject specific blogs and play a leading role in advocating the use of blogs for scholarly communication and commenting on research findings.

### 3.4 Flickr

It is an image- and video-hosting website and web services suite that was created by Ludicorp in 2004 and acquired by Yahoo on 20th March 2005. In addition to being a popular website for users to share and embed personal photographs, and effectively an online community, the service is widely used by photo researchers and by bloggers to host images that they embed in blogs and media. Librarians can use it to market general library services to their users. Most students / users

are not aware of the different services offered in the library such as reservation of Books, Reference services and strategic dissemination of information. Flickr is good to sensitize users with library services.

### 3.5 Pinterest

Pinterest is a web and mobile application company that operates a software system designed to discover information on the World Wide Web, mainly using images and on a shorter scale, GIFs and videos. It provides great venues to market library resources. A library can make its own profile and create boards, pinning photos and video showcasing the library.

## 4. Advantages of using Social Media

Use of Social media by the library has now become Mainstream. As per the survey 80% librarians now feel that use of Social Media is important.

1. Financial costs of using Social media are perceived to be low. It requires little training.
2. It connects users with the library activities and engages them in library activities. It increases interactions with library users.
3. It helps gather feedback of libraries for redesign the library services.
4. To marketing the product and services of library.
5. To promote information on new arrivals on library resources and process to use it.
6. It also creates awareness and innovate the way users live.

## 5. Challenges of using Social Media

1. Social Media can require considerable time from Library Professional.
2. The Users can create a false account and do anything without being traced.
3. There are limited funds to support more advanced social media usage / features and the training that would be required to enable this.
4. Library professionals need to work hard to maintain engagement with library users.
5. Personal Data and privacy can easily be hacked and shared on the internet.

6. Factors such as internet connectivity, technological infrastructure and government restrictions on the use of Social Media may restrict access.

## 6. Conclusion

His research contributes about various techniques used to improve library services and resources through social media. It has practical implications for how to effectively promote future services and resources and contributes to future researchers wanting to explore library promotional techniques. By using social media libraries can fulfill users need. Study gathers that according to the changing needs of library users, libraries need to be changed correctly. Libraries should be facilitated with internet service and develop their web page and social media page. Social media is used by libraries to supply a merging of user services, news, content etc.

## References

1. Burkhardt, "Four reasons libraries should be on social media", 2009.
2. Chand S.N, Suman.D &.Nirmalendu.P, "Application of Web 2.0 in library and information science: With special Reference to RSS", 2008.
3. Kamdani,S & Kumbar. B.D, "Web-based services expected from libraries: A case study of Management Institutes in Mumbai City "2006.
4. Kaba, "Marketing information resources and services on the web: current status of academic libraries in the United Arab Emirates" Information Development,2011 27(58), 58-65.
5. Fang.F, "Using Google analytics for improving library Website content and design: a case study" , Library Philosophy and Practice, 9(2), 2007
6. Khan. S.A. & Bhatti.R, "Application of social media in marketing of library and informationservices: A case study from Pakistan" Webology, 9(1), 1, 2012



**USAGE OF E-SHODHSINDHU RESOURCES BY USERS OF SRI  
PADMAVATHI MAHILA UNIVERSITY: A STUDY**

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**Abstract**

Based on the recommendation of an Expert Committee Ministry of Education has formed e-Shodh Sindhu merging three consortia initiatives, namely UGC-INFONET Digital Library Consortium, NLIST and INDEST-AICTE Consortium. The e-Shodh Sindhu will continue to provide current as well as archival access to more than 10,000 core and peer-reviewed journals and a number of bibliographic, citation and factual databases in different disciplines from a large number of publishers and aggregators to its member institutions including centrally-funded technical institutions, universities and colleges that are covered under 12(B) and 2(f) Sections of the UGC Act.

**Keywords:** E-Shodh Sindhu, UGC Infonet, Infflibnet, E-Journals, E-Resources.

**Introduction**

Modern libraries provides literature and information through electronic sources and online services to support the academic activity and satisfy the user's information needs, so it is considered as the backbone of any research organization. The development of technology is changing the pattern of library resources and also changing the process of storage and retrieval in the digital environment from the traditional physical environment. Now the modern library is acquiring e-resources through consortia approach. Consortia approach is one of the ways to maintain cooperation, coordination and collaboration between libraries for Resources sharing and work as a technology for collection development and minimize the cost.

**Sri Padmavati Mahila Visvavidyalayam** (University for Women) was founded in the year 1983 by Sri N.T.Rama Rao, the then Chief Minister of Andhra Pradesh, with the fervent desire to train women students as better builders of the Nation and to inculcate skills of leadership in all aspects of life. The University was established under the Sri Padmavati Mahila Visvavidyalayam Act of 1983, which has come into force on the 14 of April 1983. It was started with ten faculties and 300 students and twenty staff members. Today the University has a student population nearly 4000 and an academic staff of 105 and 131 academic consultants.

**Objectives of study**

1. To look into the information search habits of Faculty Members and Research Scholars and PG Students using ICT facilities.
2. To study the knowledge of Faculty Members, Research Scholars and PG Students



in use of E- Shodh Sindhu Consortium

3. To know whether there is essential cooperation in terms of user orientation is available to the faculty members, PG Students and research scholars to use E-Shodh Sindhu Consortium.
4. To know regarding frequency of usage of E-Shodh Sindhu
4. To know the purpose of usage of E-Shodh Sindhu
5. To know about level of satisfaction in using E-Shodh Sindhu
6. To identify the problems in facing E-Shodh Sindhu

#### Review of Literature

- 1) Liao<sup>1</sup> conducted a study on information seeking behavior of agricultural researchers as teachers in Taiwan. He investigated that the critical ways for the teachers to get needed Literature and the use pattern of information sources is primarily the primary sources of information
- 2) Majid<sup>2</sup> and Tan investigated the information needs and information seeking behavior of Computer engineering undergraduate students at Nanyang Technological University (NTU), Singapore. The purpose was to investigate the types of information sources used by the students, the information formats they preferred, the importance of and reasons for using certain Information sources and the use of various electronic information sources. A questionnaire was Distributed to 200 randomly selected students and 102 completed questionnaires were returned. The study found that printed materials were the most preferred information format among the Students. The top five most preferred information sources, in the order of importance, were Books, lecturers, the Internet, friends and manuals. Unexpectedly, the use of databases and Electronic journals were quite low among the computer engineering students. The study Recommends a promotional campaign for introducing electronic information
- 3) Sbeba<sup>4</sup> discusses agricultural information seeking behavior and use patterns among the African farmers and extension workers. Study suggests for establishment of agricultural Advisory board comprising both librarians and extension workers to make extensive use of Non book material as a means to overcome the handicap of illiteracy among farmers.
- 4) UNESCO (1998) observed that the rapid breakthrough in new information and Communication technologies would further change the way knowledge was developed, acquired and delivered. It was also important to note that the new technologies have offered opportunities to innovate on course content and teaching methods and to widen access to higher learning.

#### Methodology and Limitations

As discussed above, the present study was confined to Sri Padmavathi Mahila University. 2301 post- graduate students, 483 research scholars are studying in the University. Further there are more than 231 Faculty members working in the university. Due to the limited time frame, a sample survey random



Technique used questionnaire method was adopted. Total 50 questionnaires were distributed to the faculty members and of which 40 faculty members were responded to the survey. Further, of the 100 research scholars to whom questionnaires were distributed, only 55 were responded to the present study, further 125 PG Students were distributed and 76 responded. The collected primary data is analyzed and discussed under.

#### Data Analysis and Interpretation

The data collected by the questionnaires were analyzed and Interpreted and present in Tables & Figures

**Table 1 Distribution of Respondents**

S.No	Questionnaires distributed	Questionnaires received	Percentage
Faculty	75	40	53.33%
Research Scholars	100	55	55.00%
PG Students	145	76	52.00%

The study of data in Table 1 shows that 75 Questionnaires distributed among faculty members and 40 Questionnaires were received that amount to percentage of 53.33 %. 100 Questionnaires were distributed among research scholars which include both full time and part time and 55 Questionnaires were received that amount to 55 % and 145 Questionnaires were distributed and 76 received which amount to 52.00%.

**Table 2 Internet Skills**

S.No	Very Good	Fair	Uncertain	Total
Faculty	37(92.5%)	2(.05%)	1(.025%)	40
Research Scholars	40(72.0%)	12(21%)	3(.05%)	55
PG Students	45(59%)	20(26%)	11(14%)	76

The study of data in Table 2 describe 92.5 % of faculty had very good Internet Skills .05% are fair and .02 % are uncertain. Regarding Internet skills for scholars 72 % had very good Internet Skills 21 % were fair and .05 % were uncertain with their internet skills. Regarding PG Students.

**Table 3 Awareness on E-ShodhSindhu**

S.No	Faculty	Research Scholars	PG Students
Yes	37 (92.5%)	52(94%)	30(39%)
No	03(0.075%)	03(0.05%)	46(60%)
Total	40	55	76

Table 3 tells regarding awareness on E-Shodh Sindhu. 92.5 % faculty had awareness on E-Shodh Sindhu, 94% Scholars had awareness on E-Shodh Sindhu. 39 % of PG Students have awareness on E-Shodh Sindhu.



**Table 4 Frequency of use of E-Shodh Sindhu**

Library Visit	Faculty	Research Scholars	PG Students
Daily	3 (7.5%)	25 (45%)	06(7%)
Twice a Week	09(22.5%)	07(12.7%)	25(32%)
once a Week	15(37.5%)	08(14.5%)	15(19.7%)
Once in fortnight	13(32.5%)	15(27%)	30(39%)
Total	40	55	76

Table 4 shows that 45% of Research Scholars visit library daily, 32 % of PG Students visit library twice a week 37.5% of faculty visits library once a week, 39 % of PG Students visit library once in a fortnight.

**Table 5 Search preference of respondents**

Search Preference of Respondents	Faculty	Research Scholars	PG Students
Author	15(37%)	5(9%)	14(18%)
Title	19(47%)	15(27%)	28(36%)
Subject	8(20%)	20(36%)	15(19%)
Keyword	7(17%)	10(18%)	13(17%)
Publisher	1(.025)	5(9%)	06(7%)
Total	40	55	76

Table 5 explains that 37% of faculty search by author, 47% of faculty search by Title, 36% of research scholars search by subject, 18% of research scholars search by key word, 9% of research scholars search by publisher. Here we took only majority of respondents among different variables.

**Table 6 Purpose of using E-Shodh Sindhu**

Purpose	Faculty	Research Scholars	PG Students
To keep abreast with the latest developments	5(12.5%)	7(12.7%)	5(6.5%)
For research work	11(27.5%)	19(34.5%)	15(19.7%)
For prepare research article	18(45%)	18(32.7%)	21(27.6%)
For seminar, workshop, presentation	6(15%)	13(23.6%)	35(46%)
Total	40	55	76

Table 6 explain us 45% of faculty use E-Shodh Sindhu for writing research article, 34.7% of Research scholars use E-Shodh Sindhu for research work, 46% of PG Students use for seminar, workshop and presentation.



**Table 9 Opinion towards E-Shodh Sindhu Consortia Resources**

Opinion	Faculty	Research Scholars	PG Students
Excellent	6(15%)	13(23.6%)	15(19.7%)
Very Good	11(27.5%)	7(12.7%)	5(6.5%)
Good	5(12.5%)	19(34.5%)	21(27.6%)
Poor	18(45%)	18(32.7%)	35(46%)
Total	40	55	76

Table 8 explain us respondents view while using E-Shodh Sindhu 23.6% of Research Scholars view as Excellent, 27.5% faculty view it as very good while 32.7% Research scholars view it as good while 46% PG Students express their view as Poor.

#### Findings

- 1) While responding to present study research scholars have shown more interest 55%
- 2) It is found that faculty had excellent Internet skills and majority of students are uncertain while using Internet.
- 3) Research scholars are more aware regarding awareness on E-Shodh Sindhu
- 4) Majority of research scholars use E-Shodh Sindhu on daily basis while majority off aculty use it as twice aweek
- 5) Majority of faculty search by title while scholars search by subject and PG Students by Title only.
- 6) Faculty use E-Shodh Sindhu for preparation of Article, Scholars uses it for research work while students use it for seminars and presentation.
- 7) Regarding popularity faculty, scholars and PG Students use Springer link,
- 8) Problems while accessing E-Shodh Sindhu faculty responds to Lack of sufficient Internet nodes in University Library, scholars respond to Slow Internet bandwidth, PG Students to Lack of knowledge to use.
- 9) With regard to E-Shodh Sindhu Consortia Resources faculty have poor opinion, scholars & PG Students had good in opinion.

#### Conclusion

E-Shodh Sindhu not only reduces budgetary expenses of various university libraries but also increases quality of producing articles and research production. It is highly recommended for sufficient training programmes on continuous basis for every academic year. Respondents recommended inclusion of some publishers in databases.

#### References

- 1) Gulati A, 2004. Use of information and communication technology in libraries and information centres: an Indian scenario, The Electronic Library, Vol. 22, No. 4, pp. 335- 350.
- 2) Inflibnet. <https://ess.inflibnet.ac.in/>



- 3) Komuravelli P, 2014. E-Resources in UGC-Infonet Digital Library Consortium: A Profile, International Journal of Digital Library Services, Vol. 4, No. 3, pp.263-275.
- 4) Lingaiah V, 2016. A Study on Use of E-ShodhSindhu Resources by the Scholars and Faculty Members in Jawaharlal Nehru Technological University, Hyderabad, Journal of Advances in Library and Information Science, Vol. 5, No. 3, pp.268-272.
- 5) Vishala BK and Bhandi MK, 2008. Use of UGC-Infonet Digital Library Consortium Resources: Its Impact on University Libraries of Karnataka, 6th International CALIBER- 2008, University of Allahabad, Allahabad, February 28- 29 & March 1, 2008



## Usage of e-Shodh Sindhu in Promotion of Research and Academic Knowledge by Scholars and PG Students in Shri Padmavati Mahila Vishvavidyalayam: A Study

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### Abstract

*The aim of this paper is to know about usage of E-shodhsindhu by research scholars and PG Students their awareness, area of specialization, availability, level of satisfaction, and future requirements for their study. It also explains their experience in usage of E-shodhsindhu and productivity.*

**Keywords:** *E-ShodhSindhu, E-Books, E-Journals, E-Databases, INFLIBNET, Resource Sharing Sri Padmavati Mahila Vishvavidyalayam.*

### Introduction

Libraries play an important role in vast changing e environment. E resources change the role of libraries, its scope and users. It also changed its nature of job what it has done earlier. Modern information and communication technologies had profound influence on its access to various types of online e resources like e journals, databases, e books method of resource sharing and upgrade its resources to attract its users.

### UGC INFONET- Digital Library consortium

The University Grants Commission (UGC) has initiated a programme to provide e resource over Internet to scholarly literature in all areas of learning to the university sector in India. The programme is wholly funded by the UGC. All universities which come under UGC are the members of this programme and it also be extended to degree colleges also.

### E-ShodhSindhu

Based on the recommendations of an expert committee, the MHRD has formed e-shodhsindhu merging three consortia Initiatives, namely UGC INFONET Digital Library Consortium, NLIST and INDEST-AICTE Consortium. The e-ShodhSindhu will continue to provide current as well as archival access to more than 15,000 core and peer-reviewed journals and a number of bibliographic, citation and factual databases in different disciplines from a large number of publishers and aggregators to its member institutions including centrally-funded technical institutions, universities and colleges that are covered under 12 (B) and 2(f) Sections of the UGC Act.

### Objectives of Study

- The major objectives of the study follow
- To find out the awareness and utility of e shodhsindhu among research scholars and PG Students
- To identify area of interest on databases on e shodhsindhu
- To find out the purpose and utilization on e shodhsindhu
- To estimate level of satisfaction regarding usage of e shodhsindhu
- To find out the suggestions and improvement on usage of e shodhsindhu

### Methodology

This present study used survey method to conduct research. A well structured questionnaire covering entire components was distributed to collect data from respondents.

### Scope and Limitations

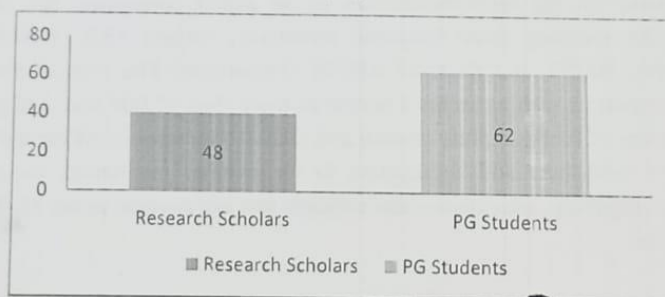
The present study confined was carried out by distributing 50 Questionnaire to Research Scholars and 75 Questionnaire to PG Students of Shri Padmavathi Mahila Vishvavidyalayamu in the year 2019. Only 40 Scholars and 62 PG Students responded.

### Literature Review

At this stage it is important to view research gap need and importance of present study. It is revealed that majority of users are aware of e shodhsindhu online resources available in library. YCH Venkateswarlu ( 2015) " Problems In accessing UGC INFONET E-Journals consortium among Research scholars : A Survey of Sri Venkateswara University Library , Tirupathi, AP" have indicated Increase acceptance of electronic resources by research scholars in present environment in which the UGC INFONET has played a meaningful role to assist academic and research community.

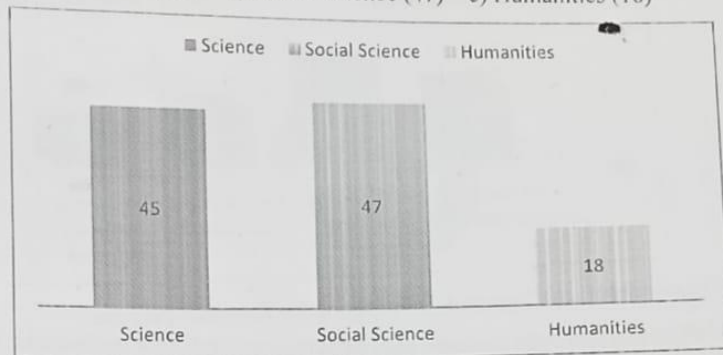
### Data Analysis and Interpretations

1) Designation Research Scholars (48), PG Students (62)



The above graph explains usage E-ShodhSindhu by PG students more than Research scholars.

2) Departments (a) Science (45) b) Social Science (47) c) Humanities (18)

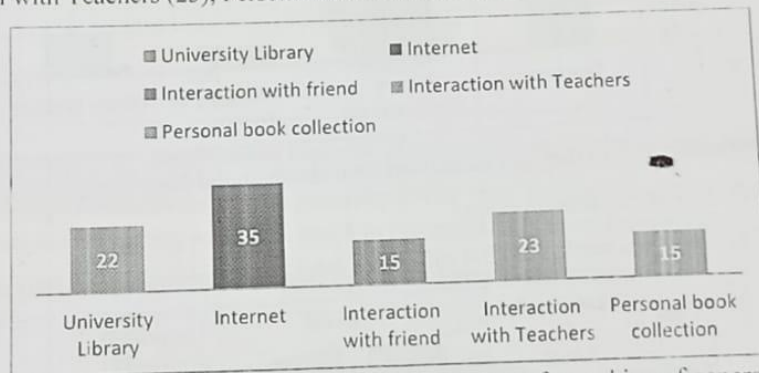


The above graph explains usage of e-shodhsindhu by social science department more when compare to sciences and humanities

3) When you are seeking information, which source of information do you generally consult? Tick all the relevant Colum?

University Library (32), Internet (35), Interaction with friend and colleague (15)

Interaction with Teachers (23), Personal book collection (15)

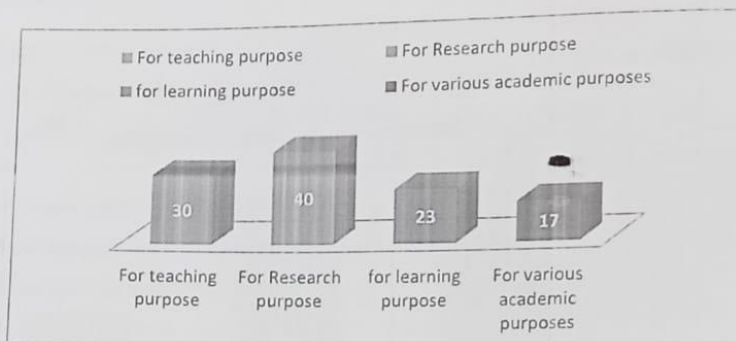


The above graph explains usage of Internet more for seeking of general information.

4) For what purpose do you seek information (Tick all the relevant columns)?

(30) For teaching purpose, (40) For Research purpose, (23) for learning purpose,

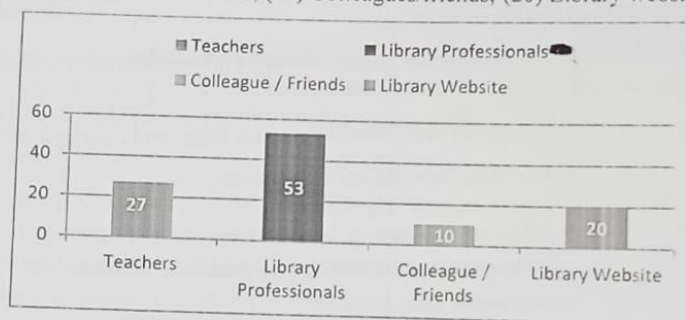
(17) For various academic purposes.



Majority of respondents says they seek information for Research purpose.

5) From which source you came to know about E-ShodhSindhu?

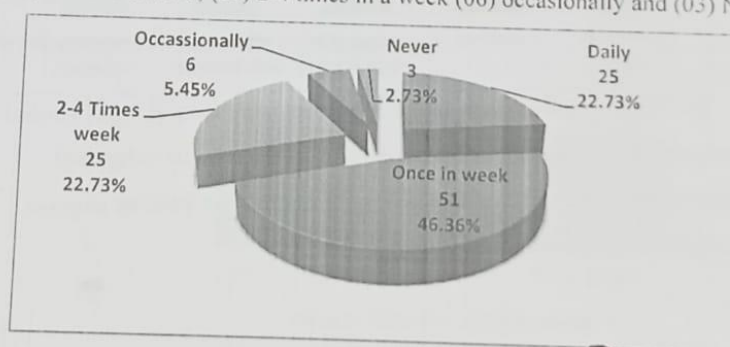
(27) Teachers, (53) Library Professionals, (10) Colleagues/friends, (20) Library website



Majority of respondents says they get awareness on e shodhsindhu from Library Professionals.

6) How often do you use E-ShodhSindhu?

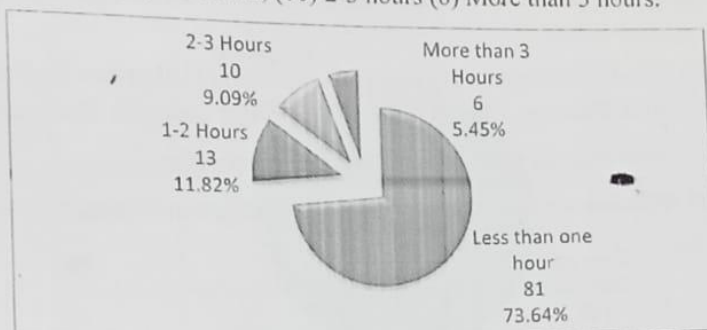
(25) Daily, (51) Once in a week, (25) 2-4 times in a week (06) occasionally and (03) Never.



Majority of respondents said they use e-shodhsindhu once in a week.

7) How much time do you spend in E-ShodhSindhu in a visit?

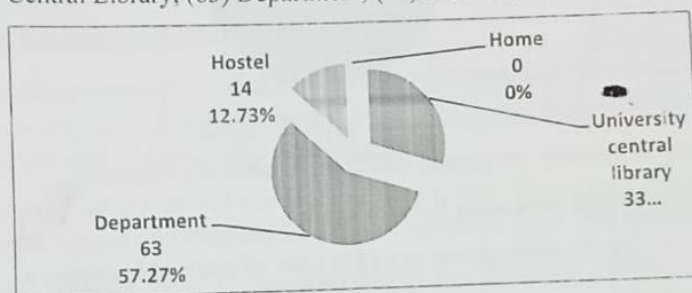
(81) Less than one hour, (13) 1-2 hours, (10) 2-3 hours (6) More than 3 hours.



Majority of respondents says that they spend on e shodhsindhu by less than one hour

8) Where do you access E-ShodhSindhu?

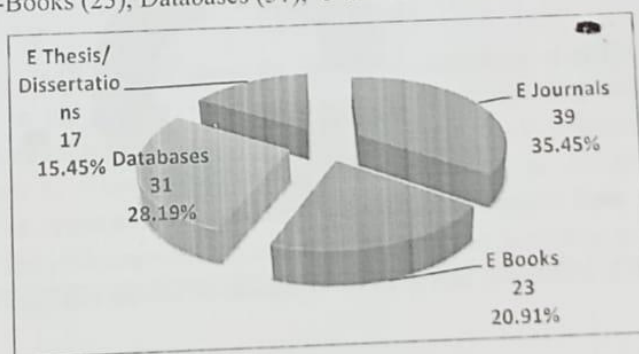
(33) University Central Library, (63) Department, (14) Hostel, (Nil) Home



Regarding access to e – shodhsindhu majority of respondent access from their departments.

9) Which types of e-resources do you generally prefer from E-ShodhSindhu? (Please indicate your preferences by putting as 1, 2, 3, and 4 in the boxes/ 4 is more important)

E- Journals (39), E-Books (23), Databases (31), e-theses/dissertations (17).



Majority of respondents prefer to use E- Journals from E-ShodhSindhu.

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Dr. R. Sarangapani, is working as University Librarian, Bharathiar University, Coimbatore. He finished his PG & Doctorate Degree in LIS at Annamalai University. He has presented 47 papers in various conference and published 15 articles in journals. He also published a book "Information Literacy Expertise". He got Best National Librarian Awards-2017 from MALA, Chennai. He is Resources Person, Invited Speaker, Key Note Speaker, Inaugural Address in various programme. He also Selection Committee Member and External Examiner in various universities in Tamil Nadu. He is a life member of ILA, TLA, ISTE, MALA & SALIS.



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# ಜುಂತ್ರುಶಾಸ್ತ್ರಂ

ಬಿ.ಯಸ್ಸಿ-IV ವ ಸೆಮಿಸ್ಟರ್



e-content

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ಡಾ.ಯನ್. ಶ್ರೀನಿವಾಸ

ಜುಂತ್ರುಶಾಸ್ತ್ರ ಅಧ್ಯಾಪಕುಲು

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ಡಾ.ನಿ. ಅನಿಲ್ ಕುಮಾರ್

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ಅಕಡಮಿಕ್ ಆಫಿಸರ್

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# Biogenic Nanoparticles: A Comprehensive Review to Explore Multidrug Resistance Mechanisms among Microbes

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## ABSTRACT

Microbial resistance has been an intense hindrance in preventing diseases for many decades. The rapid emergence of resistance towards several antibiotics could help the bacteria to become stronger to the existing antibiotics and becoming fatal to the mankind. This multidrug resistance is paving a challenging threat to human beings globally due to indiscriminate use of synthetic antibiotics and several other chemical compounds. Several factors have been influencing in developing resistance either by vertical or horizontal gene transfer among microbial species through different ecosystems. Microorganisms are altering genetic makeup to tackle the existing antibiotics very rapidly. Therefore, there is an urgent need to overcome this resistance and to develop new forms of antibiotics that are cost-effective, biocompatible, showing fewer side effects and a single-step fabricated approach that helps for large scale production. Biogenic mediated metallic nanoparticles became a

hopeful alternative to fight against the microbial multidrug resistance and also used to change the microbial ecology to keep up non-pathogenic microbial flora. These are being safely and effectively used clinically for a broad range of gene therapy applications and also to treat multidrug resistance among pathogenic microbes. The present review summarizes the synthesis of biogenic mediated nanoparticles and their interactions with the biological environment for treating infectious diseases.

**Keywords:** Antibiotic resistance, Conventional antibiotics, Multidrug-resistant bacteria, Metallic nanoparticles, Biogenic nanoparticles.

## INTRODUCTION

The term 'Antibiotic' arises from the word Antibiosis (Against life). Antibiotics are composed of chemical compounds and biological elements, which can either inhibit or kill the growth of microorganisms. Antibiotics can be classified as Antibacterial, Antiviral, and Antifungal, depending on their target group. The term Antibiotic is generally used to explain antibacterial compounds (Etebu and Arkekepar, 2016). Since the invention of Penicillin, the first antibiotic, released and used without clinical trials, scientists started working on invention of new antibiotics and are being used to treat infectious diseases. Besides, they support in various procedures like chemotherapy and organ transplants and also in major surgeries. For the last few decades' multiple classes of antibiotics have been produced synthetically as well biologically to overcome the drug-resistance. The drug-resistance of bacteria will be developed through changes in DNA, RNA, protein, enzyme inhibition, and membrane structure disruption (Kohanski and Dwyer *et al.*, 2010). Therefore, to fight against resistant microbes, antibiotics are essential and without which we can't imagine the world (Padiyara and Inoue *et al.*, 2018; Gonzalez-Candelas *et al.*, 2017). Antibiotic resistance was a major issue for human beings, which causes infectious diseases (Gonzalez-Candelas and Comas *et al.*, 2017). Usage of high doses of drugs, high toxicity, and improper hygienic conditions in the hospital environment leads to development of resistance in microbes (Pelgrift and Friedman, 2013). Some factors which develop antibiotic resistance in microorganisms are misuse and overuse of antibiotics, usage in agriculture, and availability of new antibiotics (Ventola, 2015). The present era acts as a medium of fast-spreading multidrug-resistant microorganisms globally. An antibiotic-resistant microorganism poses a severe threat of epidemic infections, and it spreads an unconstructive social and economic effect on society (Baluaand Nabi, 2018). The World Health Organization (WHO) has confirmed antibiotic-resistant microorganisms as one of the significant threats to global health (Davis *et al.*, 2018).

The mechanisms of antibiotic resistance in microbes were displayed through antimicrobial drug reduction, increased drug efflux, alteration

of drug targets, and modification of enzymes. Bacteria play a significant role in the degrading of drugs through the formation of the biofilm layer (Bhattacharand Nasir *et al.*, 2011). These results in less accumulation of drugs in bacterial cells, and the drug cannot be easily bound to the active target site (Huh and Kwon, 2011). The higher amount and repeated administration of drugs usage lead to developing side effects on human beings and animals. Many existing antibiotics that we are using were miserably failed to respond to the infectious diseases caused by the pathogenic microorganism by developing resistance against the antibiotics (Teixeira and Sanchez-Lopez, 2018). In the past few decades, there have been no reports on the development of new antibiotic classes. Innovation and commercialization of antibiotics and the discovery of new antibiotics is a long process, high-priced and licensing (Bartlett and Gilbert, 2013). This condition revealed the fact that multidrug-resistant bacteria can come out quickly to any new antibiotics, which results in a reduction of antibiotic usage (Adeniji, 2018). To tackle these problems, there is an imperative need to discover new drugs against infectious diseases caused by antibiotic-resistant bacteria. To solve these problems scientists have done many experiments to achieve targeted therapy and rapid diagnostics by modifying antibiotics or avoiding the use of conventional antibiotics. To overcome this obstacle, investigating the metals like platinum, gold, silver, copper, and zinc displayed a better alternative for antimicrobial agents. From past few decades, metals are used as antimicrobial agents. Metals act against microbes like antibiotics through several mechanisms such as cellular components disruption, damage of cell membranes and generation of reactive oxygen species (Ahmand Singh, 2018). The unique properties of these metals can be understood by exploring with the help of nanotechnology. The nanoparticles have been establishing by an improvement of all the features of metals at the nanoscale (Singh and Singh *et al.* 2017). The transformation of metals to nanoparticles depends on the size and shape of nanoparticles. They may occur in the form of flower, rod, spherical, triangular and octahedral shapes (Singh and Kim *et al.*, 2015; 2016). (Fig. 1)



FIG. 1: Different shapes of nanoparticles

The antimicrobial action of nanoparticles is directly proportional to the area of surface which interacts with biological compounds. Consequently, the metal nanoparticles became the most confident choice. Consequently, the resistant microbes and fight against the multiplying to overcome the resistant microorganisms (Huh and Kwon, 2011). The production of resistant nanoparticles consists of various conventional methods being metallic nanoparticles which include physical methods such as used for the past few decades, which include physical methods being laser ablation, physical vapour deposition, melt mixing, sputtering and chemical methods comprises photoreduction, thermolysis, sol-gel and microemulsion. These physical and chemical synthetic methods give microemulsion. These nanoparticles, and it will become toxic in the course of instability to the nanoparticles, and it will become toxic in the course of treatment. The nanoparticles can be stabilized by capping agents like polyethylene glycol or starch etc but this stabilization causes the production of toxic substances. Contrary to these problems, green methodologies have substituted the conventional synthesis of nanoparticles. (Singh and Pandit *et al.*, 2018). In green methods, the resources of microorganisms and plants are being used to produce metallic nanoparticles (Singh and Kim *et al.*, 2016). Bio reduction is the best method of the accumulation of metal ions and reduces their toxicity with the help of various reducing species present inside or outside of the cell wall of bacteria, whereas plants contain multiple flavonoids, proteins, and water-soluble biomolecules as reducing agents. These reducing agents help to detoxify the metals and help in containing antibiotic resistance. Different methods of nanoparticle synthesis were given in the following picture: (Fig 2)

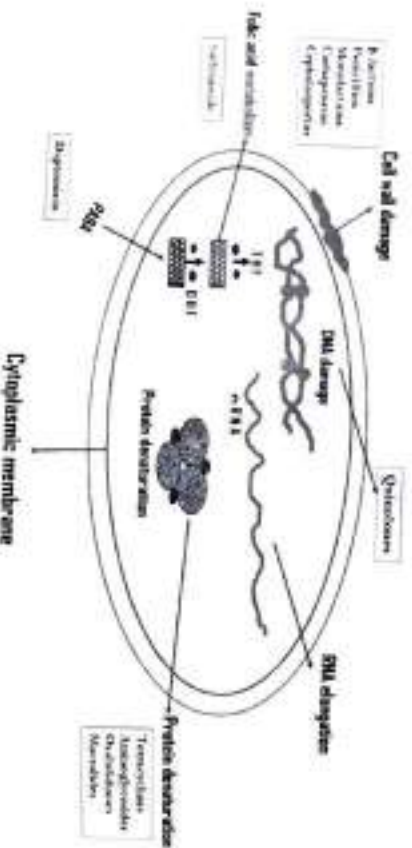
The green nanotechnology methods contain many advantages, which include stable biomolecule production, cell surface coated with nanoparticles that possess new active cell surface for the interaction of surrounding biological environment and this leads to the need for additional stabilizing agents and reducing compounds (Singh *et al.*, 2016; 2017). These characteristics make the nanoparticles to contain a high economic value in the global arena. Nanoparticles that are arising from microbes through the capping layer and nucleation possess high stability in biogenic nanoparticles (Abhai *et al.*, 2016). Capping layer generally forms during biogenic nanoparticle synthesis and arise from the biological extracts used for the synthesis of correlated nanoparticles. It helps the biological nanoparticles to increase their long-term stability. Green nanoparticles production by microorganisms requires heavy instruments for the creation, maintenance, and purification of biogenic nanoparticles. All these sophisticated instruments used for the production and purification of biogenic nanoparticles are expensive (Singh *et al.*, 2016 and Jo *et al.*, 2016), wherein the case of plants, they do not require costly instruments for production of biogenic nanoparticles (Singh *et al.*, 2018; Hsu *et al.*, 2018). The advantages of biogenic metallic nanoparticles over the physicochemically obtained nanoparticles, however, the benefits of biogenic metallic nanoparticles over physico chemically



In this review, we focused on the limitations of conventional antibiotics, biogenic nanoparticle development, multidrug resistance implications, biogenicity of metallic nanoparticles, therapeutic application of biogenic metal nanoparticles, and their future perspectives.

## CONVENTIONAL ANTIBIOTICS AND THEIR TARGETS ON THE CELL

The first antibiotic penicillin was discovered in 1928, and it led to the beginning of the modern era in antibiotics (Ventola, 2015). The mid-20<sup>th</sup> century was the era that the majority of the antibiotics were discovered and called "Age of Antibiotics." With the discovery of antibiotics and their innovation, resistance remains a significant and growing challenge to infectious diseases (Fischbach and Walsh, 2009). They can be classified based on their action of the spectrum, mode of action, or their chemical structure. Antibiotics can act on microbes, either bactericidal or bacteriostatic. The narrow-spectrum antibiotics work on either Gram-positive or Gram-negative bacteria, while the broad-spectrum antibiotics target both the Gram-negative and Gram-positive bacteria (Adzity, 2015). Based on their molecular structure, antibiotics can be classified as sulphonamides, oxazolidinones, and macrolides (Elebu and Arikekar, 2016).



**FIG 4:** Mechanism of microbial cell death by conventional antibiotics.

The beta-lactam antibiotics interfere with cell wall synthesis and destroy the bacteria by cell lysis and other categories of antibiotics target on different areas of the cell (Fig-4). These antibiotics are further divided

into penicillins, monolactam, carbapenems, and cephalosporins. In the early 1960s, penicillin-resistant bacteria were observed. These bacteria were able to produce an enzyme called beta-lactamase; it degrades the beta-lactam antibiotics. To overcome these problems, carbapenems were developed. These are resistant to beta-lactamases and contain broad-spectrum activity (Etebu and Arikekpar, 2016).

The tetracyclines, aminoglycosides, oxazolidinones, and macrolides suppress the growth of bacteria by targeting protein synthesis in cell walls. Macrolides act on protein synthesis during translation (Etebu and Arikekpar, 2016). Oxazolidinones also act on protein synthesis like macrolides but bind to 50S ribosomal subunit. Instead of inhibition of protein synthesis, it enhances the translation initiation complex (Pandit *et al.*, 2012). Aminoglycosides and tetracyclines are 30S inhibitor groups, which binds to 30S ribosomal subunit and turn down aminocycl 1-RNA from the ribosome and inhibit the translation (Fig-4). Depending upon the mode of action, tetracyclines and macrolides are bacteriostatic. Aminoglycosides are broad-spectrum antibiotics (Etebu and Arikekpar, 2016). Quinolones act on DNA helicases and inhibit the growth of bacteria and also RNA synthesis through interfering with topoisomerases, which leads to the effect on RNA polymerase (Etebu and Arikekpar, 2016). Sulphonamide structure is like para-aminobenzoic acid (PABA), it blocks folic acid synthesis by competing with para-aminobenzoic acid (PABA), and it is a substrate for the synthesis of folic acid in microbial cells. Daptomycin breaks down of the cytoplasmic membrane structure or function through depolarizes calcium-dependent membrane leads to disruption of the cytoplasmic membrane in bacteria (Alborn *et al.*, 1991). Previous records have reported conventional antibiotics, and these are used for treating infectious diseases from the past few decades. At present, these antibiotics are not responding to treat the infections caused by microorganisms. Hence these organisms develop resistance against conventional antibiotics. Some microbes like *Enterococcus* species like *E. faecalis*, *E. faecium* and *Staphylococcus aureus* (Ceitkaya *et al.*, 2000) *Streptococcus pneumoniae*, *Mycobacterium tuberculosis*, *Salmonella enterica*, *Vibrio cholerae*, *Enterobacteriaceae* and *Acinetobacter baumannii* shows resistance against antibiotics by developing resistant genes via horizontal gene transfer by: transformation, transduction and conjugation (Hajipour *et al.*, 2012), some spontaneous mutations in the existing genes makes the bacteria as multidrug-resistant bacteria. Finally, microorganisms contain resistance genes towards the antibiotics. This could happen when bacteria prone to prolong exposure of antibiotics.

To understand the regulation of genes responsible for antibiotic resistance, need to be studied at genetic and molecular levels (Frye *et al.* 2013). The changes in the cellular genome lead to the development of resistance or by the addition of extrachromosomal genes in the genome, helps to enhance resistance, termed as acquired resistance. Likewise, the

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various mechanisms make the bacteria to adapt resistance to two or more antibiotics and referred to as multidrug-resistant bacteria or superbugs.

## BIOGENIC METALLIC NANOPARTICLES FOR ANTIBACTERIAL APPLICATIONS

The nanoparticle is a particulate matter which comes under the nanometres size. Nowadays, the use of nanotechnology for the production of particulate matter, which has a high surface area, has become a trend. Nanotechnology helps in many ways to achieve good targets for drug delivery systems. Nanomaterials also being used in biological applications. Nanomaterials include carbon nanotubes, polymeric nanomaterials, and metallic nanoparticles (Gloria Medugno *et al.*, 2014). As the chemical synthesis of nanoparticles has many disadvantages, the use of the biological synthesis of nanoparticles has taken advantage. The biogenic synthesis and application of metallic nanoparticles have proven a good record as antimicrobials (Dalia and Salem *et al.*, 2019). At present the use of metals like platinum, gold, silver, zinc, copper, titanium, magnesium, and silica for green synthesis of nanoparticles has been accelerating since the dawn of the 21<sup>st</sup> century. The wide range of applications of biogenic nanoparticles in various fields used as follows like catalytic (Jiang *et al.*, 2005), bactericidal (Sekar *et al.*, 2011), optical devices (Anderson and Moskovits, 2008), anti-biofouling (Agrawal *et al.*, 2014), dye degradation (Priyadarini *et al.*, 2014), electronic (Kao *et al.*, 2000), sensor technology (Sharpe and Andreescu, 2015), some cancers treatments (Pratetorius and Mandal, 2007) and malaria treatment (Kathik *et al.*, 2013) and so on.

Metallic nanoparticles may vary in their size and shape and depends on viable properties. Metallic nanoparticles such as platinum, gold, silver, copper, and zinc can be used against antibiotic-resistant bacteria (Jin and Kwon, 2011; Fernandez-Moure and Evangelopoulos, 2017). Due to their long-term stability and biocompatibility, biogenic nanoparticles are used for antimicrobial applications against many infections caused by pathogenic microorganisms. The mechanism of the antimicrobial effect of biogenic nanoparticles involves the release of a metal ion through oxidative stress, non-oxidative stress. The biogenic silver nanoparticles synthesized from the exudates of indigenous plants *Azadirachta indica* were reported effective against pathogenic organisms (Jannu and Bhatt, 2018). Copper nanoparticles synthesized from *Sida acuta* displayed enhanced antimicrobial activity against *Proteus mirabilis*, *Staphylococcus aureus* and *Escherichia coli* (Sathiyavimal *et al.*, 2018). It also increases the efficacy of conventional antibiotics like rifampicin, penicillin G, neomycin, vancomycin, oleandomycin, and linezolidin. Microbial metabolites also used for the synthesis of biogenic nanoparticles. The nanoparticles obtained from microorganisms like *Brevibacterium Frigidolerans* DC2 (Singh and Kim, 2015), *Sporosarcina karwasii* DC4 (Singh and Singh

2016) and *Rhizoglyphus tricola* DC1 (Singh and Kim *et al.*, 2015) showed antibacterial activity against some species like *Escherichia coli*, *Salmonella cholerae*, *Bacillus subtilis*, *Bacillus cereus*, *Canalida albicans* and *Vibrio parahaemolyticus*. Metallic zinc nanoparticles also showed significant antimicrobial activity against *E. coli*, *P. aeruginosa*, and *S. aureus* (Pasquet and Chevalier, 2014). *Salmonella typhimurium* ATCC 14028, *B. subtilis* ATCC 6633, and *Micrococcus luteus* ATCC 9341 are most susceptible to biological nanoparticles when compared with chemically synthesized zinc nanoparticles (Singh and Kim, 2016).

Biogenic nanoparticles are most effective against drug-resistant bacteria when compared to chemically synthesized nanoparticles. Biological nickel nanoparticle synthesized from *Desmodium gangeticum* is having biocompatibility and antibacterial activity against Gram-positive (*Staphylococcus aureus*) and Gram-negative (*Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Vibrio cholerae*, and *Proteus vulgaris*) organisms. These biogenic nanoparticles were more efficient and less toxic than chemically synthesized nickel nanoparticles (Sudhasree *et al.*, 2014). For the past few decades, nanoparticles, especially metal like silver, are being used in dental implants, surgical instruments, and bone prostheses as an antimicrobial preventive measure and also used to combat the microbes in wound dressing (Correa *et al.*, 2015; Burdusel *et al.* 2018). The activity of various metal nanoparticles like silver, platinum, gold, zinc, and copper targets the primary function of cell membranes such as membrane permeability and membrane respiration (Dakal *et al.*, 2016; Slavin *et al.*, 2017). These metal nanoparticles react with intracellular components such as nucleic acids and proteins leads to the reduction of gene transfer mechanisms and cell division (Dakal *et al.* 2016; Slavin *et al.* 2017). The mechanism of activity of nanoparticles and antibiotics is similar in interaction and membrane disruption (Kohanski *et al.*, 2010; Dakal *et al.* 2016). Metallic nanoparticles mostly exhibit antimicrobial activity through multiple mechanisms, which decrease the possibility of the development of resistance against them in microorganisms. In microbes, gene mutations rarely occur and develop resistance against nanoparticles. The biogenic nanoparticles activity can be increased by synthesizing them with compounds like proteins, polysaccharides, and small bioactive compounds. These kinds of molecules have been proved to be effective and enhance the antimicrobial activity towards drug-resistant microorganisms. In this context, the overall effects and operators of metallic nanoparticles and their synthetic biogenic methods on the different multidrug-resistant microorganisms were discussed.

### Platinum Nanoparticles

Platinum is a scarce and vital metal available on the earth. It can be reduced to synthesize nanoparticles by applying various physical, chemical, and biological methods. Depending on the methods of synthesis and

production, platinum nanoparticles vary in their shape and size. Biogenic platinum nanoparticles are enormously studied for their antimicrobial properties (Kim *et al.*, 2008). Platinum nanoparticles contain antimicrobial activity against multidrug-resistant bacteria, and some applications are used to develop hydrogen storage material and fuel cells (Wen *et al.*, 2008; Li *et al.*, 2007). Mostly platinum nanoparticles can be used as coating materials and development of polymer membranes and nanofibers. Some platinum-based chemotherapeutic drugs like oxaliplatin, cisplatin and polyphenol (TPP) extracts were used as surface modifying reducing agent to synthesize platinum biogenic nanoparticles (Ali *et al.*, 2016). Some of the combinations of platinum nanoparticles with irradiation showed fast ion efficiency and can be used in hadron therapy. These nanoparticles increase damage in DNA, thus confirmed that the sensitization in nanoparticles is increased due to auto-amplified electronic cascades inside the nanoparticles, which supports the energy deposition in the close vicinity of the metal. The combination of fast ion radiation (hadron therapy) with platinum nanoparticles also improves cancer therapy protocols (Erika Porcel *et al.*, 2010).

### Gold Nanoparticles

Gold nanoparticles (AuNPs) are widely used in various applications. The shape of the synthesized gold nanoparticles can be triangular (Suman *et al.*, 2014), hexagonal (Shenoy *et al.*, 2012), spherical (Aromal *et al.*, 2012) and rod-shaped. Biogenic gold nanoparticles are synthesized by using plant extract, microbial exudates *etc.*, (Ahal Bharat *et al.*, 2007), which acts as reducing agents in stabilizing the nanoparticles (Cecilia Fernandez-Ponce *et al.*, 2018). The type of extracts, like bio reductant, defines the shape and size of synthesized nanoparticles. Gold nanoparticles synthesized from *Galaxaura elongata*, are in a range of 4-77nm size and with shapes like a rod, spherical, triangular, hexagonal (Abdel-Raouf *et al.*, 2017). Some studies reported that pH affects the size of gold nanoparticles. Yang *et al.*, (2014) reported that mango peel extract mediated nanoparticles were in the range of 6nm and 18 nm at pH 9 and pH 2 respectively. Gold nanoparticles showed biocompatibility to bacterial cells without bactericidal and bacteriostatic activity. However, when gold nanoparticles were conjugated with antibiotics develops a strong bactericidal effect against antibiotic-resistant bacteria. Gold nanoparticles conjugated with ampicillin have been involved in the disruption of ampicillin-resistant microbes and enhanced the susceptibility of multidrug-resistant bacteria like *E. coli* K-12 sub-strain DH5-alpha, MRSA, *Enterobacter aerogenus* and *P. aeruginosa* (Brown *et al.*, 2012).

### Silver Nanoparticles

Silver nanoparticles have been utilized in many pharmaceutical industries.

food and packaging industries, paints, and ointments industries (Suresh *et al.*, 2012). Plants, bacteria, fungi, and yeast are extensively used for biogenic production of silver nanoparticles (Singh *et al.*, 2016). Silver nanoparticles have remarkable bactericidal and fungicidal properties. The antimicrobial mechanism involved in the silver nanoparticles is either disruption of energy metabolism or due to release of silver ions (Hosainogad *et al.* 2018; Kim *et al.*, 2016), and majorly involvement of disruption of energy metabolism, cell membrane damage, oxidative stress generation due to formation of ROS and transcription inhibition. Silver nanoparticles release silver ions to interact with sulphur and phosphorous, containing groups in the cell wall and plasma membrane of microbes (Hindi *et al.*, 2009). Interaction between silver and microbial cell membranes is due to the binding of cationic silver with negatively charged bacterial cells and produces multiple pores in the cell membrane, which leads to the efflux of intracellular components outside. This causes an electrochemical imbalance in the bacterial cell and allows the silver ions to balance the intracellular components and leads to complete cell damage (Dakai *et al.*, 2016). Silver ions inhibit the protein and enzyme activity that are essential for the production of ATP. ROS production leads to inhibition of respiratory enzymes, damages the DNA and RNA and content disrupt and destabilize the outer membrane. Due to their size and large surface area, nanoparticles have a high capability to cross the cell membranes and peptidoglycan layer in bacteria (Blecher *et al.*, 2011; Lara *et al.*, 2010). Silver nanoparticles affect mostly the gram-negative bacteria when compared with gram-positive bacteria which are having thick peptidoglycan layer (Singh *et al.*, 2018). This property makes the gram-positive bacteria to contain more resistance against the nanoparticles. Many studies revealed that antimicrobial activity of silver nanoparticles and their toxicity depends upon the size and shape of nanoparticles (Raza *et al.*, 2016). The nanoparticles contain more surface area and have been releasing a higher rate of silver ion, and they contain high antimicrobial activity (Tang *et al.* 2018). Antimicrobial activity of silver nanoparticles has been studied against some multidrug-resistant microbes like *E. coli*, *S. aureus*, *P. aeruginosa*, *Enterococcus SP.*, *Streptococcus pyogenes*, *Salmonella SP* and *Klebsiella pneumonia* (Jinu *et al.*, 2017; Gopinath *et al.*, 2015). The bactericidal effect of silver nanoparticles against resistant bacteria is through lysis of cell wall synthesis and disruption of protein synthesis. Silver nanoparticles have the capability of enhancing the antibacterial activity of antibiotics like vancomycin, amoxicillin, clindamycin, penicillin G and erythromycin against *E. coli* and *S. aureus* (Shahverdi *et al.*, 2007). Additionally, the silver nanoparticles encapsulated with antibiotics act against multidrug-resistant bacteria, which include MRSA, *P. aeruginosa*, *A. baumannii*, *K. pneumonia* and *Burkholderia cepacia* (Leid *et al.*, 2012). Recent studies involving pre-treatment of microbial cells with a sub-lethal concentration of silver nanoparticles showed a low level of membrane damage and intracellular ROS and production of a higher amount of

intracellular ATP'. This reveals that pre-treatment of microbial cells with sublethal concentrations of silver nanoparticles provides long-lasting responses, which enhance the antibiotic stress resistance in bacteria at various levels (Kawceetceerawat *et al.*, 2017).

### Zinc Oxide Nanoparticles

Zinc oxide nanoparticles are synthesized from different parts of plants such as roots, leaves, rhizomes, flowers, fruits, bark (Ahmed *et al.*, 2017) and synthesized by various biological resources as reducing agents (Madhumitha *et al.*, 2016). Zinc oxide nanoparticles act as catalyzing agent nontoxic and show a potential antimicrobial activity (Bhuyan *et al.*, 2015). These are also used in applications of drug delivery systems (Ali *et al.*, 2016) and anticancer therapy due to their photocatalysis activity (Vimala *et al.*, 2014). Zinc oxide nanoparticles have a wide range of both Gram-positive and Gram-negative bacteria. The microorganisms like *E. coli*, *S. aureus*, *Listeria monocytogenes*, and *Salmonella* have exhibited sensitivity towards Zinc oxide nanoparticles (Jones *et al.*, 2008; Liu *et al.*, 2009). Zinc oxide nanoparticles act on microbial cells and lead to leakage of the membrane, reducing sugars, proteins, DNA content and generation of Reactive Oxygen Species (ROS), lipid peroxidation (Tiwari *et al.*, 2018). The production of ROS, such as superoxide anion and hydrogen peroxide in cells leads to membrane leakage of nucleic acids and proteins by increasing lipid peroxidation on the membrane (Kumar *et al.*, 2011; Horie *et al.*, 2012). Some zinc nanoparticles release Zn<sup>2+</sup> ions and also interact with intracellular compounds and damage the cell membrane (Mc Devitt *et al.*, 2011; Li *et al.*, 2011). Studies of zinc nanoparticles on *Carbapenem-resistant A. baumannii* displayed inhibition of bacterial growth by the production of ROS that leads to cell damage. This suggests that zinc oxide nanoparticles play an important role as an alternative to beta-lactams (Tiwari *et al.*, 2018).

### Copper Nanoparticles

Biogenic cupric oxide nanoparticles are synthesized from biogenic polysaccharides such as chitosan, pectin, alginate, bacteria, and leaf extract. Unlike silver, gold, and other nanoparticles, cupric oxide nanoparticles also act as antimicrobial agents. Cupric oxide nanoparticles are easy to synthesize but contain some disadvantages like low potency, environmentally unfriendliness, high toxicity, and high cost. Cupric oxide is majorly inert from copper salt, and in some conditions, these are non-inert (El-Batal *et al.*, 2018). Compared to other nanoparticles the production of biogenic cupric oxide nanoparticles is a relatively new topic, and it is being made accessible and eco-friendly. The mechanism involved in antimicrobial activity of cupric oxide is an electrostatic attraction between Cu<sup>2+</sup> and plasma membrane leading to damage of cell membrane and cell death (Hoshino *et al.*, 1999; Bogdanovic *et al.* 2014).

The  $\text{Cu}^{2+}$  ions are small and easy to move across a lipid bilayer, and it forms ROS, protein oxidation, and lipid peroxidation (Bogdanovic *et al.*, 2014). Cupric oxide nanoparticles had an intense antimicrobial activity against the Gram-positive and Gram-negative bacteria (Bogdanović *et al.*, 2014; DeAlba-Montero *et al.*, 2017). As the copper nanoparticles showing strong microbial effects, these are used to treat wound healing in bacterial plasters and bandages due to its illegible sensibility of human tissue and strong microbial effect of copper compounds (Borkow *et al.*, 2004; Hostynek *et al.*, 2003).

TABLE 1: Properties & Applications of Metallic Nanoparticles

Nanoparticles	Properties	Applications
Platinum	Antioxidant (Kim <i>et al.</i> , 2008)	To develop hydrogen storage material and fuel cells (Wen <i>et al.</i> , 2006; Li <i>et al.</i> , 2007)
Gold	Act as a reducing agent to stabilize the nanoparticle (Cecilia Fernandez-Ponce <i>et al.</i> , 2018).	Biocompatibility and enhance the susceptibility of multidrug-resistant bacteria (Brown <i>et al.</i> , 2012).
Silver	Antimicrobial activity (Tang <i>et al.</i> , 2018).	Utilized in many pharmaceutical industries, food and packaging industries, paints, and ointments industries (Suresh <i>et al.</i> , 2012).
Zinc oxide	Act as catalyzing agent nontoxic and show a potential antimicrobial activity (Bhuyan <i>et al.</i> , 2015).	Used in drug delivery systems (Ali <i>et al.</i> , 2016) and anticancer therapy (Vimala <i>et al.</i> , 2014).
Copper	Antimicrobial activity against the Gram-positive and Gram-negative bacteria (Bogdanovic <i>et al.</i> , 2014; DeAlba-Montero <i>et al.</i> , 2017).	Used to treat wound healing in bacterial plasters and bandages (Borkow <i>et al.</i> , 2004; Hostynek <i>et al.</i> , 2003).

## Future Perspectives

Multidrug-resistant bacterial infections constitute a significant threat to global healthcare. Microorganisms commonly develop resistance against drugs due to overdoses, incapability of medicines, and inferior diagnostic methods. To overcome these problems, the biogenic metallic nanoparticles may be an alternative. These nanoparticles, either individually or in conjugation with antibiotics or with biomolecules produce intense antimicrobial activity against multidrug-resistant bacteria. Depending upon nanoparticle distribution, their bioavailability, and active targeting, excretion is highly concentrated to overcome the classical antibacterial resistant mechanisms. Nowadays, nanotechnology is helping majorly to overcome pathogenic infections (Singh *et al.*, 2018). Biogenic metallic nanoparticles and their applications are not only limited to biomedical

areas and also extended to textiles, food packaging, cosmetics, agriculture in the form of nano pesticides and nano fertilizers, wastewater treatment, washing machines, computer keyboards and self-cleaning coating on mobile phones. Biogenic nanoparticles are nontoxic and have excellent biocompatibility when compared to chemically synthesized nanoparticles. Nanoparticles also have great potential as a strategy for gene therapy and can be used to treat genetic defects *in vitro* and *in vivo*. The use of nanoparticles as carriers for the delivery of therapeutic materials to target tissues has become popular in recent years and has demonstrated high potentials for the treatment of a wide range of diseases like ocular therapy that have been successfully tested in the eye, lung, and brain. These particles resulted in higher transfection efficiency and longer duration of expression than other non-viral vectors without any toxicity or other side effects. These are being safely used clinically and are efficient for a broad range of gene therapy applications (Konstan *et al.*, 2004). Majority of the studies were evidenced for antibacterial activity, but scientists have to extend and explore the research on anti-viral activities of these nanoparticles to take containment measures to prevent epidemics, pandemics and also syndemics. The high production and accessibility of the biogenic metallic nanoparticles for biomedical applications is a unique challenge at the global market. Better research is to be engaged to increase the output with minimum cost-effective to fulfil the requirements. This would be a great help from the scientific community to the global arena in treating multidrug resistance and future therapeutic applications.

## REFERENCES

- Abbai R, Mathiyalagan R, Markus J, Kim Y J, Wang C, Singh P, Ahn S, Farh Mel A, Yang D C. Green synthesis of multifunctional silver and gold nanoparticles from the oriental herbal adaptogen: Siberian ginseng. *Int. J. Nanomed.* 2016;11: 3131-3143.
- Abdel-Raouf N, Al-Enazi N M, Ibraheem I B M. Green biosynthesis of gold nanoparticles using *Galaxaura elongata* and characterization of their antibacterial activity. *Arabian J. Chem.* 2017;10: S3029-S3039.
- Adeniji F. Global analysis of strategies to tackle antimicrobial resistance. *Int. J. Pharm. Pract.* 2018; 26: 85-89.
- Adzitey F. Antibiotic Classes and Antibiotic Susceptibility of Bacterial Isolates from Selected Poultry. A Mini Review. *World's Vet. J.* 2015;5: 36-41.
- Agarwal A, Mehra A, Karthik L, Kumar G, Rao K V B. Anti-bio fouling property of marine actinobacteria and its mediated nanoparticle. *Int. J. Nanoparticles.* 2014;7(3-4): 294-306.
- Ahmed S, Annu Chaudhry S A, Ikram S. A review on biogenic synthesis of ZnO nanoparticles using plant extracts and microbes: A prospect towards green chemistry. *J. Photochem. Photobiol. B Biol.* 2017;166: 272-284.
- Ahn S, Singh P, Jang M, Kim Y J, Castro-Aceituno V, Simu S Y, Kim Y J, Yang D C. Gold nano flowers synthesized using *Acanthopanax cortex* extract inhibit inflammatory mediators in LPS-induced RAW264.7 macrophages

- via NF-kappaB and AP-1 pathways. *Colloids Surf. B Biointerfaces*, 2018, 162: 398–404.
- Alborn W E, Allen NE, Preston A. Daptoraycin Disrupts Membrane Potential in Growing *Staphylococcus aureus*. *Antimicrobial agents and chemotherapy*. 1991, 35(11): 2282–2287.
- Ali A, Alshatwi, Jegan, Athinarayanan, Periasamy, Vaiyapuri Subbarayan. Green synthesis of platinum nanoparticles that induce cell death and G2/M-phase cell cycle arrest in human cervical cancer cells. 2015. *J Mater Sci: Mater Med*. 26:7.
- Ali K, Dwivedi S, Azam A, Saquib Q, Al-Said M S, Alkhedhairy A A, Musarrat J. *Aloe vera* extract functionalized zinc oxide nanoparticles as nanoantibiotics against multi-drug resistant clinical bacterial isolates. *J. Colloid Interface Sci*. 2016, 472:145–156.
- Anderson D J, Moskovits M. "A SERS-active system based on silver nanoparticles tethered to a deposited silver film". *J. Phys. Chem. B*, 2006, 110(28): 13722–13727.
- Aromal S A, Vidhu V K, Philip D. Green synthesis of well-dispersed gold nanoparticles using *Macrotylomauniflorum*. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc*. 2012, 85: 99–104.
- Atul Bharde, Aarohi Kulkarni, Mala Rao, Asmita Prabhune, Murali Sastry. "Bacterial Enzyme Mediated Biosynthesis of Gold Nanoparticles. *Journal of Nanoscience and Nanotechnology* 2007, 7: 4369–4377.
- Baluja Z, Nabi N, Ray A. Challenges in Antimicrobial Resistance: An Update. *EC Pharmacol. Toxicol*. 2018, 6: 865–877.
- Bartlett J G, Gilbert D N, Spellberg B. 2013. Seven ways to preserve the miracle of antibiotics. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am*, 56: 1445–1450.
- Bhuyan T, Mishra K, Khanuja M, Prasad R, Varma A. Biosynthesis of zinc oxide nanoparticles from *Azadirachta indica* for antibacterial and photocatalytic applications. *Mater. Sci. Semicond. Process*. 2015, 32: 55–61.
- Blecher K, Nasir A, Friedman A. The growing role of nanotechnology in combating infectious disease. *Virulence*. 2011, 2: 395–401.
- Bogdanovic U, Lazic V, Vodnik V, Budimir M, Markovic Z, Dimitrijevic S. Copper nanoparticles with high antimicrobial activity. *Mater. Lett*. 2014, 128: 75–78.
- Borkow G, Gabbay J. Putting copper into action: Copper-impregnated products with potent biocidal activities. *FASEB J. Official Publ. Fed. Am. Soc. Exp. Biol*. 2004, 18: 1728–1730.
- Brown AN, Smith K, Samuels T A, Lu J, Obare S O, Scott M E. Nanoparticles functionalized with ampicillin destroy multiple-antibiotic-resistant isolates of *Pseudomonas aeruginosa* and *Enterobacter aerogenes* and methicillin-resistant *Staphylococcus aureus*. *Appl. Environ. Microbiol*. 2012, 78:2768–2774.
- Burdusel A C, Gherasim O, Crumezescu A M, Mogoanta L, Fici A, Andronescu E. Biomedical Applications of Silver Nanoparticles: An Up-to-Date Overview. *Nanomaterials*. 2018, 8: 681.
- Cecilia Fernandez-Ponce, Juan P. Muñoz-Miranda, Desire M. de los Santos, Enrique Aguado, Francisco Garcia-Cozar, Rocio Litran. 2018. Influence of size and surface capping on photoluminescence and cytotoxicity of gold

- nanoparticles. *J Nanopart Res*, 20: 305.
- Cetinkaya Y, Falk P, Mayhall C G. Vancomycin-resistant enterococci. *Clin. Microbiol. Rev.*13: 2000.686–707.
- Correa J M, Mori M, Sanches H L, da Cruz A D, Poiate E, Jr Poiate I A. Silver nanoparticles in dental biomaterials. *Int. J. Biomater.* 2015.485275.
- Dakal T C, Kumar A, Majumdar R S, Yadav V. Mechanistic Basis of Antimicrobial Actions of Silver Nanoparticles. *Front. Microbiol.* 2016. 7: 1831.
- Dalia M S A, Salem Mona M, Ismail, Mohamed A. Aly-Eldeen. Biogenic synthesis and antimicrobial potency of iron oxide (Fe<sub>3</sub>O<sub>4</sub>) nanoparticles using algae harvested from the Mediterranean Sea. *Egyptian Journal of Aquatic Research.* 2019.45: 197–204.
- Davis M, Whittaker A, Lindgren M, Djerf-Pierre M, Manderson L, Flowers P. Understanding media publics and the antimicrobial resistance crisis. *Glob. Public Health.* 2018.13: 1158–1168.
- De Alba-Montero I, Guajardo-Pacheco J, Morales-Sanchez E, Araujo-Martinez R, Loredó-Becerra G M, Martínez-Castanón G A, Ruiz F, Compeán Jasso M E. Antimicrobial Properties of Copper Nanoparticles and Amino Acid Chelated Copper Nanoparticles Produced by Using a Soya Extract. *Bioinorg. Chem. Appl.* 2017.1064918.
- El-Batal A I, Al-Hazmi N E, Mosallam F M, El-Sayyad G S. Biogenic synthesis of copper nanoparticles by natural polysaccharides and *Pleurotus ostreatus* fermented fenugreek using gamma rays with antioxidant and antimicrobial potential towards some wound pathogens. *Microb. Pathog.* 2018. 118: 159–169.
- Erika Porcel, Samuel Liehn, Hynd Remita, Noriko Usami, Katsumi Kobayashi, Yoshiya Furusawa, Claude Le Sechand Sandrine Lacombe. Platinum nanoparticles: a promising material for future cancer therapy? 2010. *Nanotechnology.*21: 085103.
- Etebu E, Arikekpar I. Antibiotics: Classification and mechanisms of action with emphasis on molecular perspectives. *Int. J. Appl. Microbiol. Biotechnol.* 2016.4:90–101.
- Fernandez-Moure J S, Evangelopoulos M, Colvill K, Van Eps J L, Tasciotti E. Nanoantibiotics: A new paradigm for the treatment of surgical infection. *Nanomed. (Lond).* 2017.12: 1319–1334.
- Fischbach Michael, Walsh Christopher T. Antibiotics for Emerging Pathogens. *Science.* 2009.28: 325(5944): 1089–1093.
- Frye Jonathan G, Charlene R. Jackson. Genetic mechanisms of antimicrobial resistance identified in *Salmonella enterica*, *Escherichia coli*, and *Enterococcus* spp. isolated from U.S. food animals. *Front. Microbiol.* 2013.
- Gloria Modugno, Cécilia Ménard-Moyon, Maurizio Prato, Alberto Bianco. Carbon nanomaterials combined with metal nanoparticles for theranostic applications. *British Journal of Pharmacology.* 2014.172:975–991.
- González-Candelas F, Comas I, Martínez J I, Galán J C, Baquero F. 12–The Evolution of Antibiotic Resistance. In *Genetics and Evolution of Infectious Diseases*, 2nd ed. Tibayrenc, Elsevier: London, UK. 2017.257–284.
- Gopinath P M, Narchonai G, Dhanasekaran D, Ranjani A, Thajuddin N. Mycosynthesis, characterization and antibacterial properties of AgNPs

- against multidrug resistant (MDR) bacterial pathogens of female infertility cases. *Asian J. Pharm. Sci.* 2015.10: 138–145.
- Hajipour M J, Fromm K M, Akbar Ashkarran A, Jimenez de Aberasturi D, Larramendi I R D, Rojo, T, Serpooshan V, Parak W J, Mahmoudi M. Antibacterial properties of nanoparticles. *Trends Biotechnol.* 2012.30: 499–511.
- Hindi K M, Ditto A J, Panzner M J, Medvetz D A, Han D S, Hovis C E, Hilliard J K, Taylor J B, Yun Y H, Cannon C L *et al.*, The antimicrobial efficacy of sustained release silver-carbene complex-loaded L-tyrosine polyphosphate nanoparticles: Characterization, in vitro and in vivo studies. *Biomaterials.* 2009.30: 3771–3779.
- Hoseinnejad M, Jafari S M, Katouzian I. Inorganic and metal nanoparticles and their antimicrobial activity in food packaging applications. *Critical Rev. Microbiol.* 2018. 44:161–181.
- Hoshino N, Kimura T, Yamaji A, Ando T. "Damage to the cytoplasmic membrane of *Escherichia coli* by catechin-copper (II) complexes". *Free Radic. Biol. Med.* 1999.27: 1245–1250.
- Hostynek J J, Maibach H I. Copper hypersensitivity: Dermatologic aspects - An overview. *Rev. Environ. Health.* 2003.18: 153–183.
- Huh A J, Kwon YJ. Nanoantibiotics. A new paradigm for treating infectious diseases using nanomaterials in the antibiotic's resistant era. *J. Control. Release: Off. J. Controll. Release Soc.* 2011. 156:128–145.
- Huo Y, Singh P, Kim YJ, Soshnikova V, Kang J, Markus J, Ahn S, Castro-Aceituno V, Mathiyalagan R, Chokkalingam M. Biological synthesis of gold and silver chloride nanoparticles by *Glycyrrhiza uralensis* and in vitro applications". *Artif. Cells Nanomed. Biotechnol.* 2018.46: 303–312.
- Jamiu A T, Bello S A. Biosynthesis of silver nanoparticles using *Azadirachta indica* leaf extract and assessment of its antibacterial activity on some pathogenic enteric bacteria. *International Journal of Novel Research in Life Sciences.* 2018. 5 (2): 25-31.
- Jiang Z J, Liu C Y, Sun L W. Catalytic properties of silver nanoparticles supported on silica spheres. *J. Phys. Chem. B.*, 2005.109(5): 1730-17305.
- Jinu U, Jayalakshmi N, Sujima Anbu A, Mahendran D, Sahi S, Venkatachalam P. Biofabrication of Cubic Phase Silver Nanoparticles Loaded with Phytochemicals from *Solanum nigrum* Leaf Extracts for Potential Antibacterial, Antibiofilm and Antioxidant Activities against MDR Human Pathogens. *J. Clust. Sci.* 2017. 28: 489–505.
- Jo J H, Singh P, Kim Y J, Wang C, Mathiyalagan R, Jin C G, Yang D C. *Pseudomonas deceptionensis* DC5-mediated synthesis of extracellular silver nanoparticles. *Artif. Cells Nanomed. Biotechnol.* 2016.44: 1576–1581.
- Jones N, Ray B, Ranjit K T, Manna A C. Antibacterial activity of ZnO nanoparticle suspensions on a broad spectrum of microorganisms". *FEMS Microbiol. Lett.* 2008. 279: 71–76.
- Karthik L, Kumar G, Keswani T, Bhattacharyya A, Reddy BP, Rao KVB. Marine Actinobacterial mediated Gold nanoparticles synthesis and their antimalarial activity. *Nanome.* 2013.9(7): 951-960.
- Kaweeteerawat C, Na Übol P, Sangmuang S, Aueviriyavit S, Maniratanachote R. Mechanisms of antibiotic resistance in bacteria mediated by silver

- nanoparticles. *J. Toxicol. Environ. Health*. 2017. 80:1276–1289.
- Kim J, Takahashi M, Shimizu T, Shirasawa T, Kajita M, Kanayama A, Miyamoto Y. Effects of a potent antioxidant, platinum nanoparticle, on the lifespan of *Caenorhabditis elegans*. *Mech. Ageing Dev.* 2008.129(6): 322–331.
- Kim T, Braun G B, She Z G, Hussain S, Ruoslahti E, Sailor M J. "Composite Porous Silicon-Silver Nanoparticles as Theranostic Antibacterial Agents". *ACS Appl. Mater. Interfaces*. 2016. 8: 30449–30457.
- Kohanski M A, Dwyer D J, Collins J J. 2010. How antibiotics kill bacteria: From targets to networks. *Nat. Rev. Microbiol.* 8: 423–435.
- Konstan M W, Davis P B, Wagener J S, Hilliard K A, Stern R C, Milgram I. J. Compacted DNA nanoparticles administered to the nasal mucosa of cystic fibrosis subjects are safe and demonstrate partial to complete cystic fibrosis transmembrane regulator reconstitution. *Human Gene Therapy*. 2004. 5, 1255–1269.
- Lara H H, Ayala-Núñez N V, IxtepanTurrent L D C, Rodríguez Padilla C. Bactericidal effect of silver nanoparticles against multidrug-resistant bacteria. *World J. Microbiol. Biotechnol.* 2010.26: 615–621.
- Leid J G, Ditto A J, Knapp A, Shah P N, Wright B D, Blust R, Christensen L, Clemons C B, Wilber J P, Young G W. In vitro antimicrobial studies of silver carbene complexes: Activity of free and nanoparticle carbene formulations against clinical isolates of pathogenic bacteria. *J. Antimicrob. Chemother.* 2012.67: 138–148.
- Li Y, Yang R T, Liu C J, Wang Z. Hydrogen storage on carbon doped with platinum nanoparticles using plasma reduction. *Ind. Eng. Chem. Res.* 2007. 46: 8277–8281.
- Liu Y, He L, Mustapha A, Li H, Hu Z Q, Lin M. Antibacterial activities of zinc oxide nanoparticles against *Escherichia coli* O157:H7. *J. Appl. Microbiol.* 2009.107: 1193–1201.
- Madhumitha G, Elango G, Roopan S M. Biotechnological aspects of ZnO nanoparticles: Overview on synthesis and its applications. *Appl. Microbiol. Biotechnol.* 2016.100:571–581.
- Padiyara P, Inoue H, Sprenger M. Global Governance Mechanisms to Address Antimicrobial Resistance. *Infect. Dis. Res. Treat.* 2018.11.
- Pandit N, Singla R K, Shrivastava B. Current Updates on Oxazolidinone and Its Significance. *International Journal of Medicinal Chemistry*. 2012:159285. 2012.
- Pasquet J, Chevalier Y, Pelletier J, Couval E, Bouvier D, Bolzinger M.-A. The contribution of zinc ions to the antimicrobial activity of zinc oxide. *Colloids Surf. A Physicochem. Eng. Asp.* 2014. (457): 263–274.
- Pelgrift R Y, Friedman A J. Nanotechnology as a therapeutic tool to combat microbial resistance. *Adv. Drug Deliv. Rev.* 2013. 65:1803–1815.
- Praetorius N P, Mandal T K. Engineered nanoparticles in cancer therapy. *Recent Pat. Drug Deliv. Formula.* 2007. 1(1): 37–51.
- Priyragini S, Veena S, Swetha D, Karthik L, Kumar G, Rao K VB. Evaluating the effectiveness of marine actinobacterial extract and its mediated titanium dioxide nanoparticle in the degradation of azo dyes. *J. Environ. Sci.* 2014. 26(4): 775–782.
- Rao C N R, Kulkarni G U, Thomas P J, Edwards P P. Metal nanoparticles and

- their assemblies. *Chem. Soc. Rev.* 2000. 29: 27-35.
- Raza M A, Kanwal Z, Rauf A, Sabri A N, Riaz S, Naseem S. Size- and Shape-Dependent Antibacterial Studies of Silver Nanoparticles Synthesized by Wet Chemical Routes. *Nanomaterials.* 2016.6: 74.
- Sathiyavimal S, Vasantharaj S, Bharathi D, Saravanan M, Manikandan E, Kumar S S, Pugazhendhi A. Biogenesis of copper oxide nanoparticles (CuONPs) using *Sidaacuta* and their incorporation over cotton fabrics to prevent the pathogenicity of Gram negative and Gram positive bacteria. *J. Photochem. Photobiol. B Biol.* 2018.188: 126-134.
- Sekar D K, Karthik L, Kumar G, Rao K V B. Biosynthesis of silver nanoparticles from marine yeast and their antimicrobial activity against multidrug resistant pathogens. *Pharmacol. Online Newslett.* 2011.3: 1100-1111.
- Shahverdi A R, Fakhimi A, Shahverdi H R, Minaian S. Synthesis and effect of silver nanoparticles on the antibacterial activity of different antibiotics against *Staphylococcus aureus* and *Escherichia coli*. *Nanomed. Nanotechnol. Biol. Med.* 2007. 3: 168-171.
- Sharpe E, Andreescu S. Portable nanoparticle based sensors for antioxidant analysis. *Methods Mol. Biol.* 2015.1208: 221-231.
- Sheny D S, Mathew J, Philip D. Synthesis characterization and catalytic action of hexagonal gold nanoparticles using essential oils extracted from *Anacardium occidentale*. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* 2012. 97: 306-310.
- Singh P, Kim Y J, Wang C, Mathiyalagan R, Yang D C. Microbial synthesis of Flower-shaped gold nanoparticles. *Artif. Cells Nanomed. Biotechnol.* 2016.44:1469-1474.
- Singh P, Kim Y J, Wang C, Mathiyalagan R, El-Agamy Farh M, Yang D C. Biogenic silver and gold nanoparticles synthesized using red ginseng root extract, and their applications. *Artif. Cells Nanomed. Biotechnol.* 2016. 44:811-816.
- Singh H, Du J, Singh P, Yi T H. Ecofriendly synthesis of silver and gold nanoparticles by *Euphrasiaofficinalis* leaf extract and its biomedical applications. *Artif. Cells Nanomed. Biotechnol.* 2018. 46:1163-1170.
- Singh P, Ahn S, Kang J P, Veronika S, Huo Y, Singh H, Chokkaligam M, El-AgamyFarh M, Aceituno V C, Kim Y J. In vitro anti-inflammatory activity of spherical silver nanoparticles and monodisperse hexagonal gold nanoparticles by fruit extract of *Prunus serrulata*: A green synthetic approach. *Artif. Cells Nanomed. Biotechnol.* 2017. 46: 2022-2032.
- Singh P, Kim Y J, Singh H, Mathiyalagan R, Wang C, Yang D C. Biosynthesis of Anisotropic Silver Nanoparticles by *Bhargavaea indica* and Their Synergistic Effect with Antibiotics against Pathogenic Microorganisms. *J. Nanomater.* 2015.10.
- Singh P, Kim Y J, Singh H, Wang C, Hwang K H, Farh Mel A, Yang D C. Biosynthesis, characterization, and antimicrobial applications of silver nanoparticles. *Int. J. Nanomed.* 2015.10: 2567-2577.
- Singh P, Kim Y J, Wang C, Mathiyalagan R, Yang D C. The development of a green approach for the biosynthesis of silver and gold nanoparticles by using *Panax ginseng* root extract, and their biological applications. *Artif. Cells Nanomed. Biotechnol.* 2016.44:1150-1157.

- Singh P, Kim Y J, Wang C, Mathiyalagan R, Yang D C. Weissellaoryzae DC6-facilitated green synthesis of silver nanoparticles and their antimicrobial potential. *Artif. Cells Nanomed. Biotechnol.* 2016. 44: 1569–1575.
- Singh P, Kim Y J, Yang D C. A strategic approach for rapid synthesis of gold and silver nanoparticles by *Panax ginseng* leaves. *Artif. Cells Nanomed. Biotechnol.* 2016. 44:1949–1957.
- Singh P, Kim Y J, Zhang D, Yang D C. Biological Synthesis of Nanoparticles from Plants and Microorganisms. *Trends Biotechnol.* 2016. 34: 588–599.
- Singh P, Pandit S, Garnaes J, Tunjic S, Mokkaapati V R, Sultan A, Thygesen A, Mackevica A, Mateiu R V, Daugaard A E. Green synthesis of gold and silver nanoparticles from *Cannabis sativa* (industrial hemp) and their capacity for biofilm inhibition". *Int. J. Nanomed.* 2018. (13): 3571–3591.
- Singh P, Singh H, Kim Y J, Mathiyalagan R, Wang C, Yang D C. Extracellular synthesis of silver and gold nanoparticles by *Sporosarcina koreensis* DC4 and their biological applications". *Enzyme Microb. Technol.* 2016. 86: 75–83.
- Singh, P, Singh H, Ahn S, Castro-Aceituno V, Jimenez Z, Simu S Y, Kim Y J, Yang D C. Pharmacological importance, characterization and applications of gold and silver nanoparticles synthesized by *Panax ginseng* fresh leaves. *Artif. Cells Nanomed. Biotechnol.* 2017.45:1415–1424.
- Slavin Y N, Asnis J, Hafeli U O, Bach H. Metal nanoparticles: Understanding the mechanisms behind antibacterial activity. *J. Nanobiotechnol.* 2017.15: 65.
- Sudhasree S, ShakilaBanua A, Brindhab P, Gino A, Kuriana. Synthesis of nickel nanoparticles by chemical and green route and their comparison in respect to biological effect and toxicity. *Toxicological and Environmental Chemistry.* 2014. 96 (5):743-754.
- Suman T Y, Rajasree S R, Ramkumar R, Rajthilak C, Perumal P. The Green synthesis of gold nanoparticles using an aqueous root extract of *Morinda citrifolia* L. *Spectrochim. Acta Part A Mol. Biomol. Spectrosc.* 2014. (118): 11–16.
- Suresh A K, Pelletier D A, Wang W, Morrell-Falvey J L, Gu B, Doktycz M J. Cytotoxicity induced by engineered silver nanocrystallites is dependent on surface coatings and cell types. *Langmuir ACS J. Surf. Colloids.* 2012. 28: 2727–2735.
- Tang S, Zheng J. Antibacterial Activity of Silver Nanoparticles: Structural Effects. *Adv. Healthc. Mater.* 2018. 7: 701503.
- Teixeira M C, Sanchez-Lopez E, Espina M, Calpena A C, Silva A M, Veiga F J, Garcia M L, Souto E B. Advances in antibiotic nanotherapy: Overcoming antimicrobial resistance. In: *Emerging Nanotechnologies in Immunology*; Shegokar, R., Souto, E.B., Eds.; Elsevier: Boston, MA, USA. 2018. pp. 233–259.
- Tiwari V, Mishra N, Gadani K, Solanki P S, Shah N A, Tiwari M. Mechanism of Anti-bacterial Activity of Zinc Oxide Nanoparticle against Carbapenem-Resistant *Acinetobacter baumannii*. *Front. Microbiol.* 2018. 9: 1218.
- Ventola C L. The antibiotic resistance crisis: Part 1: Causes and threats. *P T Peer-Rev. J. Formul. Manag.* 2015.40: 277–283.
- Vimala K, Sundarraj S, Paulpandi M, Vengatesan S, Kannan S. Green

synthesized doxorubicin loaded zinc oxide nanoparticles regulates the Bax and Bcl-2 expression in breast and colon carcinoma. *Process. Biochem.* 2014. 4:60–172.

Wen Z, Liu j, Li J. Core/shell Pt/C nanoparticles embedded in mesoporous carbon as a methanol-tolerant cathode catalyst in direct methanol fuel cells. *Adv. Materials*, 2008. 20(4): 743-747.

Yang N, WeiHong L, HaoL. Biosynthesis of Au nanoparticles using agricultural waste mango peel extract and its in vitro cytotoxic effect on two normal cells. *Mater. Lett.* 2014. 134: 67–70.

# A Critical Review on Potent Prospects of Biogenic Metallic Nanoparticles in Cancer Therapy

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## ABSTRACT

Cancer is one of the deadliest diseases all over the globe, causing social and economic inequalities among people. The incidence is being increased each year and diagnosed with different types of cancers with drug resistance followed by metastasis. Many cancer cells have type specific cancer protein all over their surface, which is lacking in normal cells. The nanoparticles like metal, Biogenic metal and other types conjugate or bind to the cancer cells and helps to increase the specific antibodies to unravel the treatment mechanisms. Many works indicate that the nanoparticles help in drug delivery systems as a result unique cancer treatment will be enabled. Therefore, nanotherapeutics is rapidly developing in cancer treatment, diagnostics, biomarker identification, detection of multiple genes and matrix RNA. The conjugated nanoparticles allow to detect cancer related proteins, resulting in to new method of analyzing proteome of each tumor

type. Magnetic nanoparticles are also giving exciting results for cancer detection and treatment *in vivo*. These Nano formulations are being used with chemotherapeutic agents to reduce the toxic side effects of conventional formulations. Therefore, nanoparticles play crucial role in arresting the cell proliferation mechanisms. Apoptotic pathways, Cell cycle pathways, phagocytic mechanism's and epigenetic pathways. In this regard, it has been opined that biogenic metal nanoparticles would overcome the obstacles of cancer treatment. Biogenic nanoparticles have displayed potential access towards the tumor cells with high specificity, smaller size, and bioavailability. Tumor cells promising response to the biogenic nanoparticles is opening a new avenue in the cancer therapy. In this review, an overview of the use of biogenic metallic nanoparticles in drug delivery and treatment of cancer has been discussed.

**Keywords:** Cancer, Green nanotechnology, Biogenic nanoparticles, Cancer therapy.

## INTRODUCTION

Cancer has been identified as one of the lethal diseases in the world (Bray *et al.*, 2018). Cancer has been accepted as a disease of humans in which a tumor can feed, expand, and invade. Structurally tumor is a complex tissue that consists of several cells that multiply and lose control of their proliferation and evade an immune response which leads to metastasis. A tumor cell undergoes several alterations that include undefined proliferation, inhibition of apoptosis, induction of angiogenesis, evasion, and invasion of the immune response. These alterations, in turn, create instability that leads to modification in energy metabolism, genome instability, genetic diversity, and immune evasion. These manifestations considered as hallmarks of cancer. The causes of cancer can be classified broadly into external and internal factors. Exposure to the chemicals and microorganisms like viruses can trigger an internal cancer initiation (Manzoor *et al.*, 2016), while external factors include mutations in the genes, hormonal imbalances, immune system abnormalities could strategically promote the origin of carcinogenesis (Anand *et al.*, 2008). Traditionally cancer treatment involves complex methodologies. Despite the treatment practices the metastasis has been increasing day by day. Definitely the drugs which have been using in the cancer treatment should act as one of the hallmark targets for the inhibition of cancer metastasis.

For the past two decades, there has been a paradigm shift in cancer treatment and therapeutics from traditional cytotoxic drugs to the usage of novel nano drugs. Functionally targeting cancer is a multifactorial concept and has constantly been evolving from pre genomic chemotherapy to the emergence of combinational therapy (Al-Lazikani *et al.* 2012). As the existing oncotherapeutics are evicting less response, there has been an emergence of novel technologies for the enhancement of drug response.

The usage of nanotechnology is one of the recent advancements for drug therapeutics in the present century. Nanotechnology involves the engineering of the matter at the nanoscale ( $10^{-9}$  m). Nanotechnology acts as one of the best alternatives to detect even a single cancer cell in vivo and interferes with enhancing the drug delivery to the target site. Several severe side effects are involved in the treatment of cancers like chemotherapy, radiation therapy, immunotherapy, vaccination, stem cell transformation, photodynamic therapy, or the combination of the therapies. The side effects conventional cancer therapeutic agent treatment includes toxicity, limited bioavailability and fast clearance of the drug from the site of target. For instance, the commonly used chemotherapeutic drug 5- fluorouracil induces constriction of blood vessels, cardiotoxicity, and myelotoxicity (Macdonald, 1999). Similarly, doxorubicin using in the cancer treatment evicts cardiotoxicity, renal toxicity, and myelotoxicity (Avilés *et al.* 1993). Keeping in view the instances of conventional cancer therapy we are exploring the biogenic metal nanotherapy, a newer approach involving the biogenic nanoparticles for cancer treatment

Nanomedicine is the newest concept of nanotechnology in which the bioengineering of materials at nanoscale for efficient disease diagnosis, drug delivery, and therapeutics. Nanomedicine employs the usage of nanomaterials like carbon nanotubes, nanoshells, nanoparticles, nanowires, and quantum dots (Jaishree and Gupta, 2012). Among the nanomaterials, metal nanoparticles stabilized with biological metabolites from plants, animals, and microbes have been occupying a good stage in the present world. In this context, we are aimed to give comprehensive glimpses of biological nanoparticles and their applications in cancer therapeutics.

## ASSOCIATION OF CANCER, NANOTECHNOLOGY, PLANTS AND MICROORGANISMS

There is an emergence of the need for cancer treatment with minimum side effects in contrast to the existing conventional cancer therapies. Phytochemicals from the plants would mediate a cheaper means by employing novel strategies in interactions with the tumor cell environment (Singh *et al.* 2016). Medicinal plants showed a ray of hope for cancer treatment in terms of providing compounds that are highly specific towards the cancer cell. The green compounds, when given in small quantity similar to the nano level, were dynamic in the pharmacokinetics. The plant compounds, when capped onto the surface, noble metals, provides a higher target specificity and long run of bioavailability in the tumor environment. New approaches in nanotechnology research include drug delivery diagnosis and treatment has opened a new avenue in the cancer treatment (Klefenz, 2004). The remarkable properties of nanoparticles like size, surface area to volume made them ideal for

several biological applications (Salata, 2004). Though the synthesis of nanoparticles carried out in different ways of physical and chemical methods, biological method of synthesis has paved an alternate way in the synthesis of metal nanoparticles (Rai *et al.* 2014). Living organisms like fungi and bacteria are also used to synthesize nanoparticles, but the plant-mediated synthesis provides an eco-friendly platform devoid of using toxic and harmful chemicals. Rapid and one-time synthesis can be exploited through plant-derived synthesis. Besides different size and shape of nanoparticles are produced using plants products contrary to microorganisms (Iravani, 2011).

## Nanoparticles in Cancer Therapy

Inorganic and organic nanoparticles have been using in therapeutics over a decade. Inorganic nanoparticles include iron, copper, silver, zinc, titanium. While organic nanoparticles are synthesized by conjugating with carrier molecules. They include chitosan, polylysine, liposome, *etc.* Liposome, a lipid nanoparticle is a form of vesicle-mediated carrier nanoparticle for the drug delivery at the target site.

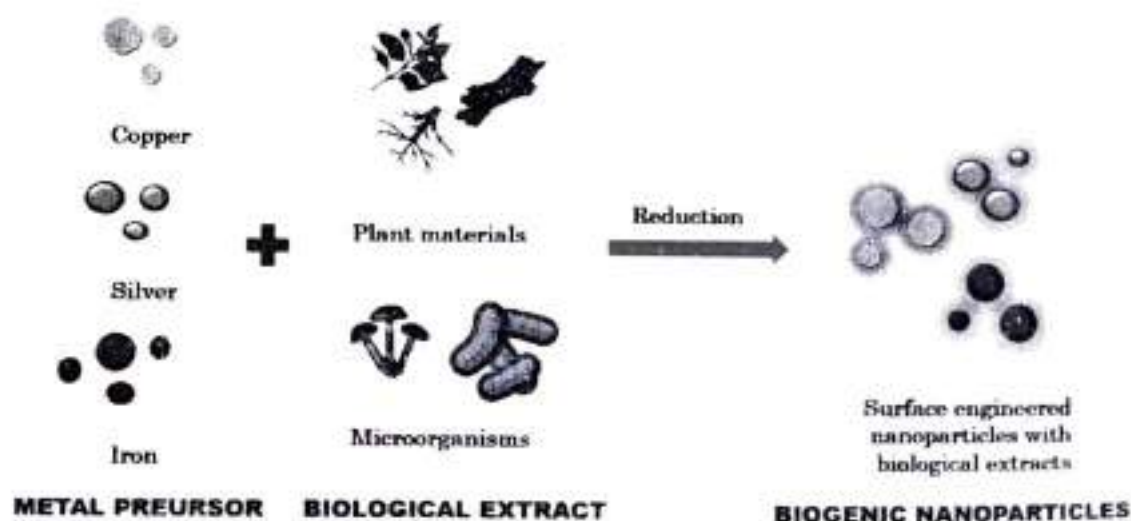


FIG. 1: Synthesis of biogenic metal nanoparticles

The commercial usage of the liposome nanoparticles has not achieved due to their instability, leakage, oxidation (Petersen *et al.* 2016). Organic nanoparticles are generally less stable due to their instability at high temperatures (Maier-Hauff *et al.* 2011). On the other hand, metallic nanoparticles showing a better alternative for cancer treatment. Herbal medicine has been in practice from the ancient days to the present. Numerous plant products were discovered to cure diseases. Phytochemical drugs act as good anti-cancer over chemical drugs and are less toxic. But the lack of targeting methods the drugs have not been achieved target specificity. The metal nanoparticles conjugated with metabolites of plant

and microbes displaying a significant result in the in-vitro treatments. These biogenic nanoparticles have a good advantage over conventional chemotherapies. Figure -1 describes the synthesis of Biogenic metal nanoparticles

### Dynamics of Interactions of Nanoparticles with Cancer Cells

The nanoparticles target delivery to the cancerous cells is accomplished in two ways, either by active targeting or by passive targeting. The basic principle involved in the active targeting is the ligand-receptor efficacy. The active targeting is performed by the recognition of tumor-specific surface receptors by the nanoparticle surface ligands. The ligands that are present on the surface of nanoparticle are actively involved in binding includes peptides, sugars, antibodies, phytochemicals (Bazak *et al.* 2015; Saha *et al.* 2010). Similarly, the receptor moieties include the sugars, proteins that are exposed on the cancer cell surface (des Rieux *et al.* 2013; Yu *et al.* 2010).

Challenges are also taking place in the active targeting of nanoparticles. The biogenic nanoparticles that are targeted should be available for a longer duration towards active antigen or receptor. This prospect is being affected due to a lack of proper blood supply to the cancerous cell (Alhaddad *et al.* 2012; Klein, 2018). Attempts having been made to escape this drawback and to enhance the bioavailability and internalization of nanoparticles. The tailoring of the surface with biogenic phytochemical and microbial metabolites paves as one of the alternatives to overcome this prospect in active targeting. Encapsulation of nanoparticles with biogenic materials have increased the therapeutic efficacy and also helps in escaping the lysosomal and endosomal environment.

The lack of tumor specificity and selectivity in active targeting searched for the passive targeting of nanoparticles. Enhanced permeability and retention effect (EPR) play an essential role in passive targeting. The large-sized nanoparticles are internalized in the tumor tissues. These nanoparticle does not penetrate the tight junctions of endothelial cells and increases their plasma half-life. The abnormal vascularization of tumor tissue is exploited in selective targeting to solid tumor tissue. The nanoparticles that are accumulated enhances their concentration over time and extravasation within the tumor tissues due to a lack of efficient lymphatic drainage. This EPR based nanotherapy proves superior to oncotherapy in a selective localizing high concentration of nanoparticles in tumor tissues with minimal side effects (Greish, 2010; Swartz *et al.* 2007). Passive targeting of nanoparticles in tumor tissue was described pictorially in Figure -2 and the role of different biogenic metallic nanoparticles in cancer studies was given in Table-1.

TABLE 1: Role of biogenic metallic nanoparticles in cancer studies

Sl. No.	Metal precursor	Source of Plant/ Microorganism	Type of biological extract	Cancer cell/ Cell lines	Activity on cancer cell	Reference
1	Gold	<i>Curcuma longa</i> (Plant)	Root extract	MCF-7, MDA-MB 231	Enhanced inhibition	(Vemuri et al. 2019)
2	Gold	<i>Paracoccus haemidaensis</i> BC74717 (Bacteria)	Extracellular extract	HEK293, HaCaT, A549	Antiproliferative	(Patil et al. 2019)
3	Gold	<i>Halymenia dilatata</i> (Seaweed)	Extracellular extract	HT-29	Cytotoxic	(Vinosha et al. 2015)
4	Gold	<i>Acacia Senegal</i> (Gum Arabic)	Bark exudate	HepG2	Inhibitory effect	(Gamal-Eldeen et al. 2016)
5	Gold	<i>Trapa natans</i> (Plant)	Leaf extract	A431	Cytotoxic	(Saber et al. 2018)
6	Silver	<i>Spinacia oleracea</i> (Plant)	Leaf extract	C2C12	Cytotoxic	(Ramachandran et al. 2017)
7	Silver	<i>Commelina nudiflora</i> (Plant)	Leaf extract	HCT116	Reduced cell viability, increased cytotoxicity	(Kuppusamy et al. 2016)
8	Silver	<i>Acalypha indica</i> (Plant)	Leaf extract	MDA-MB-231	Cytotoxic	(Krishnaraj et al. 2014)
9	Silver	<i>Humicola</i> (Fungus)	Extracellular	MDA-MB-231	Cytotoxic	(Syed et al. 2013)
10	Silver	<i>Vitex negundo</i> L (Plant)	Leaf extract	HCT15	Cytotoxic	(Prabhu et al. 2013)
11	Silver	<i>Citrullus colocynthis</i> (Plant)	Callus extract	HEP-2	Reduced cell viability	(KS et al. 2011)

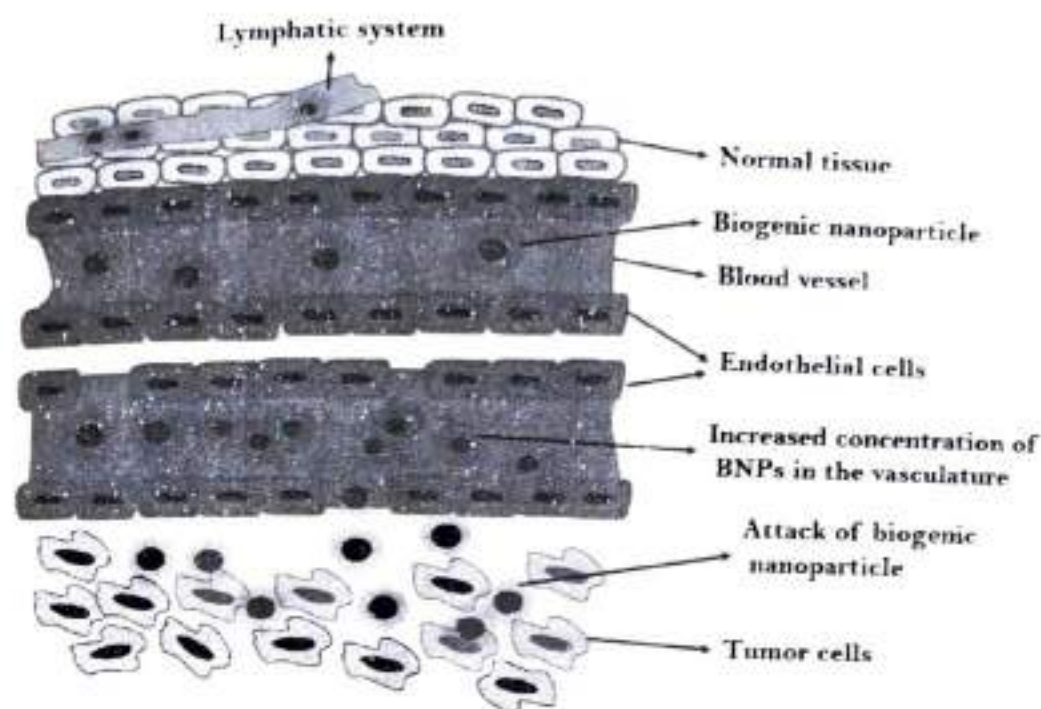


FIG. 2: Passive targeting of nanoparticles in tumor tissue

## Biogenic Gold Nanoparticles in Cancer Therapeutics

Gold nanoparticles are excellent nanoparticles with remarkable absorption capacity and scattering of light. The gold nanoparticle can convert optical energy into heat energy through non-radiative electron relaxation dynamics. The surface of the gold nanoparticle can be easily tailored and have remarkable stability and biocompatibility. They can be easily functionalized with high drug loads and less toxicity. AuNPs can be synthesized in the shape of a solid sphere, shell, cage, rods, and chains (Karathanasis and Ghaghada, 2016). As the nanoparticles are water-soluble and biocompatible drugs can be quickly delivered to the target site. From the favorable surface characteristics and fewer toxicity mechanisms, gold nanoparticles can be efficiently targeted to cancer cells, ligands, and malignant cells only be in disturbed.

The primary reason for considering nanoparticles in cancer therapies is selectively functionalizing the surface to deliver the drug to the tumor. To accomplish the functionalizing of the AuNP, the functionalizing agent should be attached to the surface layer. However, the feasible and maximum surface should be provided to advantage the surface volume ratio. Smaller, spherical nanoparticles have more surface area than the larger nanoparticles. As the radius decreases surface volume increases. Gold nanoparticles comprise numerous applications in medicine like medicals devices (Hashimoto *et al.* 2013), cancer stem cell therapy (Shen *et*

al. 2016), and biosensors (Sattarahmady *et al.* 2016).

The cancer cell inhibition was exhibited by chitosan biogenic nanoparticles. There was a lower level of cell viability of MCF-7 breast cancer cell lines when treated with biogenic chitosan conjugated gold nanoparticles (Bilal *et al.* 2019). *Mangifera indica* seed extract mediated synthesized gold nanoparticle study observed the suppression of gastric tumors (Vimalraj *et al.* 2018). Pollen extracts of *Phoenix dactylifera* (Date Palm) acted as good conjugates of gold nanoparticle (DPP-GNPs) for the treatment of human breast adenocarcinoma cells. MCF-7 breast cell lines displayed an apoptotic cell death invitro when incubated with the date palm gold nanoparticles. The apoptotic death was due to the modulation of pro-apoptotic protein p53 and anti-apoptotic protein Bcl-2 (Banu *et al.* 2018). The Au NPs synthesized using tea leaves have effectively internalized into prostate and breast cancer cell lines (Nune *et al.* 2009). The mechanism of action of Biogenic nanoparticles was described in Figure-3.

The studies on the possible pathway elucidation for MCF-7 breast cancer cell line inhibition by *Ferulago angulata* leaves hexane (extract FAHE) demonstrated that flow cytometric analysis of annexin-V induced the apoptosis of MCF-7 (Karimian *et al.* 2014). It also further revealed that MCF-7 cells were arrested in the G1 phase of the cell cycle and suppressed by an intrinsic apoptotic pathway.

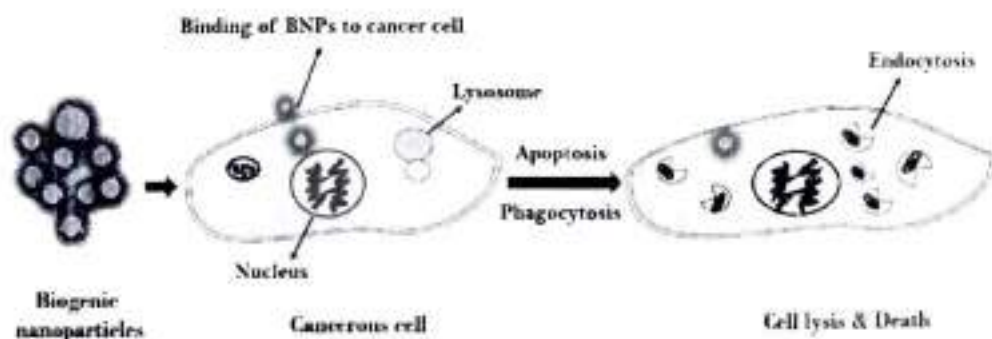


FIG. 3: Attack of biogenic nanoparticles on cancer cell

### Biogenic Silver Nanoparticles in Cancer Therapeutics

Biogenic silver nanoparticles are fabricated by using eco-friendly, nontoxic chemicals, and biodegradable polymers. The phytoconstituents include polyphenols (Nadagouda and Varma, 2008), polysaccharides (Huang and Yang, 2004), polymetalates (Troupis *et al.* 2002), etc., These acts as both reducing agents and capping agents and stabilize the silver ion into a colloidal silver nanoparticle (Nguyen *et al.* 2019). Silver has been using by the humankind since from the ancient days as in the

form of utensils and in the Ayurveda for treating ailments. The silver nanoparticles were acted as effective antimicrobials in ancient times (Wu *et al.* 2016). In the recent century usage of silver has been increasing in broad range of applications like dietary supplements, immune-boosting, sanitizing sprays (Rogers *et al.* 2018; Wasukan *et al.* 2015). Among metal nanoparticles, silver nanoparticles harbored with many attractive features that help in the usage of biology and medicine (Ullah Khan *et al.* 2018). The recent approaches of eco-friendly synthesis of silver nanoparticles not only add green-biogenic flavor but also helps as a new platform for cancer cell therapies.

The polysaccharide, Galactomannan isolated from the *Punica granatum* fruit, was conjugated with silver in preparing silver nanoparticles. The silver nanoparticles were stabilized by the capping agents as colloidal particles. This polysaccharide mediated silver nanoparticles displayed enhanced and selective cytotoxicity on human adenocarcinoma, colorectal carcinoma. The possible mechanism for the cytotoxicity was by the involvement of caspase-mediated cell death (Padinjarathil *et al.* 2018). Bilal *et al.* (2019) investigations on MCF-7 breast cancer cells revealed that biogenic silver nanoparticles synthesized from *Convolvulus arvensis* extract were inhibiting the MCF-7 cells. It also showed that C-AgNPs also inhibited the log phases of *S. aureus* and *E. coli*. The work of Rolim *et al.* (2019) exhibited the green tea extract mediated silver nanoparticles were effectively inhibiting cancer cell lines. Further, the size of the biogenic nanoparticle also a vital role in cancer therapy. Small size (10-30 nm) and spherical biogenic nanoparticles from the tea extract stabilization displayed the synergetic cytotoxic effect against the human leukemia cell lines (MoLT-4) and breast cancer cell lines (MCF-7) (Yadav and Mendhulkar, 2018). The biogenic potentiality of silver nanoparticles on lung cancer was also highly efficient. Apoptotic pathways were displayed when *Artemisia oliveriana* extract tailored silver nanoparticles were given to lung cancer cell lines (A549). The A549 cellular uptake showed substantial expression of apoptotic genes such as Bax, Bcl-2, caspase-3, caspase-9, and miR-192 (Fard *et al.* 2018). The observation of biogenic silver nanoparticles from the extracts of *Leptolyngbya* strain JSC-1 on HeLa cell line displayed the inhibition of cancerous cells due to the induction of apoptosis. (Zada *et al.* 2018). The biogenic nanoparticles were multifunctional besides inhibiting the cancer cell lines. The biogenic nanoparticles capped with the root extract of *Phoenix dactylifera* were significantly reduced the cell viability of MCF-7 breast cancer cell lines. They also exhibited the multifunction of biogenic silver nanoparticles by inhibiting the bacterial growth also (Oves *et al.* 2018). Phytochemicals capped on the biogenic nanoparticles from *Oxalis nana* Wall. ex Benth were also multifunctional. The stabilized O-AgNPs demonstrated theranostic properties towards red blood cells and macrophages (Oves *et al.* 2018). Eco-friendly stabilized silver nanoparticles from the extracts of *Rhynchosia suaveolens* demonstrated dose-dependent inhibition of variety of cancer

cell lines like human prostate carcinoma cell line (PC-3), human ovarian carcinoma cells (SKOV3), human lung adenocarcinoma cell line (A549) by the elevation of caspase-3 and caspase-7 activity (Bethu *et al.* 2018). Attempting the *Calotropis gigantea* capped silver nanoparticles against Ehrlich's ascites carcinoma (EAC) exhibited the inhibition of EAC cell line by upregulating Bax and caspase-3 genes along with the downregulation of Bcl-2 gene (Maity *et al.* 2018).

Further fungal extracts also act as good stabilizing agents for the synthesis of biogenic silver nanoparticles. Novel fungus *Piriformospora indica* extracts showed excellent capping activity towards the formation of P-AgNPs. The studies demonstrated high cytotoxicity against MCF-7, HeLa, HepG2 cell lines (Aziz *et al.* 2019). *Aspergillus niger* extract stabilized silver nanoparticles against the colon cancer cell line, HT-29 revealed that the spherical nanoparticles were potentially cytotoxic by induction of caspase-3 apoptotic pathway (Chengzheng *et al.* 2018).

## LIMITATIONS OF BIOGENIC NANOPARTICLES IN CANCER NANOTHERAPY

### 1. Toxicity of biogenic nanoparticles *in vivo*

Metallic nanoparticles as synthesized from the green chemistry; their interaction varies *in vitro* and *in vivo*. Pure metals like silver, gold, and copper were used in traditional medicinal practices since ancient times, the history of ayurvedic medicine revealed the use of silver, gold as potent therapeutic agents (Nowack *et al.* 2011). However, toxicity, side effects, bioavailability, membrane susceptibility should be thoroughly assessed before using on human beings. Most of the nanoparticles, including biogenic nanoparticles showing minimum to moderate toxicity when treated on animal models. The toxicity of metal nanoparticles *in vivo* would mainly be tracked to size, concentration, and morphology (Minai *et al.* 2013; Stensberg *et al.* 2011). Moreover, in nanotechnology, the nano-bio interfaces play a crucial role in cancer therapeutics. Nanoparticles should be primarily checked with single protein coronas like bovine serum albumin, tubulin, human serum albumin and as well as with complex protein coronas like yeast extract proteins, fetal bovine serum albumin (Durán *et al.* 2015).

### 2. Biological clearance and bioavailability

The metallic nanoparticles, when introduced *in vivo*, their kinetics have a significant concern in the removal. Micelles and polymeric nanoparticles are easily driven away from the organs in a short time, while the metal nanoparticles are looking more time for the clearance from the body (Zhou *et al.* 2011). The studies of biogenic nanoparticles clearance were highly effective in the renal cells. Glutathione coated biogenic nanoparticles

were cleared through urination. The bioavailability of the biogenic metal nanoparticles contrasts with the typical metallic nanoparticles, and their degree of bioavailability is attributed to high and low respectively. However, the detailed mechanism of clearance and biodegradable pathways is still unclear and needs further investigations before going to clinical studies (Rengan *et al.* 2015).

## CONCLUSION

The biocompatibility of nanoparticles has been exploring in the decade. Studies on the biogenic nanoparticles reported both positive and negative ways. Most of the studies *in-vitro* and reported the nontoxic levels in cell lines and biogenic nanoparticles acted as reducing agents on the growth of cancer cell lines through a mechanistic approach of apoptosis. The stimulation of the apoptotic pathway exerted the apoptotic protease caspase-3, upregulated the p53 pathway, and also elevated the sub-G1 arrest (Mukherjee *et al.* 2014). The acidic environment around the tumor tissues might have attracted the biogenic nanoparticles towards the cancerous tissue and involvement of releasing of phytoconstituents from the green colloidal nanoparticles (Gurunathan *et al.* 2013; Mukherjee *et al.* 2014). On the other hand, *in vivo* studies of biogenic metal nanoparticles revealed toxic levels to the treated individuals. The bioavailability, clearance of the metals would play a crucial role either in elevating or diminishing the toxicity. More studies to be conducted in reducing the harmfulness *in vivo* before going to the clinical trials.

For the past few years, the use of nanoparticles in cancer therapeutics is being increased. A report from the article "Nanotechnology Market Outlook 2020" forecasted the global nanotechnology market would reach high by 2020 (Anon, 2020). If the barriers of the biogenic nanoparticles are reduced, there will be a definite increase in utilizing the biogenic nanoparticles in the tumor treatments. It would certainly create a new era in cancer therapeutics.

## REFERENCES

- Al-Lazikani B, Banerji U, Workman P. (2012). Combinatorial drug therapy for cancer in the post-genomic era. *Nature Biotechnology* 30: 679–692. DOI: 10.1038/nbt.2284
- Alhaddad A, Durieu C, Dantelle G, Le Cam E, Malvy C, Treussart F, Bertrand J-R. (2012). Influence of the Internalization Pathway on the Efficacy of siRNA Delivery by Cationic Fluorescent Nanodiamonds in the Ewing Sarcoma Cell Model. Tajmir-Riahi H-A (ed). *PLoS ONE* 7: e52207. DOI: 10.1371/journal.pone.0052207
- Anand P, Kunnumakkara AB, Kunnumakara AB, Sundaram C, Harikumar KB, Tharakan ST, Lai OS, Sung B, Aggarwal BB. (2008). Cancer is a preventable disease that requires major lifestyle changes. *Pharmaceutical research* 25: 2097–116. DOI: 10.1007/s11095-008-9661-9

- Anon. ND. Nanotechnology Market Outlook (2020). [online] Available from: <https://www.reportlinker.com/p02162665/Nanotechnology-Market-Outlook.html> (Accessed 22 April 2019)
- Avilés A, Arévila N, Díaz Maqueo JC, Nambo MJ. (1993). Late Cardiac Toxicity of Doxorubicin, Epirubicin, and Mitoxantrone Therapy for Hodgkin's Disease in Adults. *Leukemia & Lymphoma* 11: 275–279. DOI: 10.3109/10428199309087004
- Aziz N, Faraz M, Sherwani MA, Fatma T, Prasad R. (2019). Illuminating the Anticancerous Efficacy of a New Fungal Chassis for Silver Nanoparticle Synthesis. *Frontiers in Chemistry* 7: 65. DOI: 10.3389/fchem.2019.00065
- Banu H *et al.* (2018). Gold and Silver Nanoparticles Biomimetically Synthesized Using Date Palm Pollen Extract-Induce Apoptosis and Regulate p53 and Bcl-2 Expression in Human Breast Adenocarcinoma Cells. *Biological Trace Element Research* 186: 122–134. DOI: 10.1007/s12011-018-1287-0
- Bazak R, Hourri M, El Achy S, Kamel S, Refaat T. (2015). Cancer active targeting by nanoparticles: a comprehensive review of literature. *Journal of Cancer Research and Clinical Oncology* 141: 769–784. DOI: 10.1007/s00432-014-1767-3
- Bethu MS, Netala VR, Domdi L, Tartte V, Janapala VR. (2018). Potential anticancer activity of biogenic silver nanoparticles using leaf extract of *Rhynchosia suaveolens*: an insight into the mechanism. *Artificial cells, nanomedicine, and biotechnology* 46: 1–11. DOI: 10.1080/21691401.2017.1414824
- Bilal M, Zhao Y, Rasheed T, Ahmed I, Hassan STS, Nawaz MZ, Iqbal HMN. (2019). Biogenic Nanoparticle–Chitosan Conjugates with Antimicrobial, Antibiofilm, and Anticancer Potentialities: Development and Characterization. *International journal of environmental research and public health* 16: 598. DOI: 10.3390/ijerph16040598
- Bray F, Ferlay J, Soerjomataram I, Siegel RL, Torre LA, Jemal A. (2018). Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA: A Cancer Journal for Clinicians* 68: 394–424. DOI: 10.3322/caac.21492
- Chengzheng W, Jiazhi W, Shuangjiang C, Swamy MK, Sinniah UR, Akhtar MS, Umar A. (2018). Biogenic Synthesis, Characterization and Evaluation of Silver Nanoparticles from *Aspergillus niger* JX556221 Against Human Colon Cancer Cell Line HT-29. *Journal of nanoscience and nanotechnology* 18: 3673–3681. DOI: 10.1166/jnn.2018.15364
- Durán N, Silveira CP, Durán M, Martínez DST. (2015). Silver nanoparticle protein corona and toxicity: a mini-review. *Journal of Nanobiotechnology* 13: 55. DOI: 10.1186/s12951-015-0114-4
- Fard NN, Noorbazargan H, Mirzaie A, Hedayati Ch M, Moghimian Z, Rahimi A. (2018). Biogenic synthesis of AgNPs using *Artemisia oliveriana* extract and their biological activities for an effective treatment of lung cancer. *Artificial cells, nanomedicine, and biotechnology* 46: S1047–S1058. DOI: 10.1080/21691401.2018.1528983
- Gamal-Eldeen AM. (2016). Photothermal therapy mediated by gum Arabic-conjugated gold nanoparticles suppresses liver preneoplastic lesions in mice. *Journal of Photochemistry and Photobiology B: Biology* 163: 47–56. DOI: 10.1016/j.jphotobiol.2016.08.009

- Greish K. (2010). Enhanced Permeability and Retention (EPR) Effect for Anticancer Nanomedicine Drug Targeting. In: 25–37. DOI: 10.1007/978-1-60761-609-2-3
- Gurunathan S, Han JW, Eppakayala V, Jeyaraj M, Kim J-H. (2013). Cytotoxicity of biologically synthesized silver nanoparticles in MDA-MB-231 human breast cancer cells. *BioMed research international* 2013: 535796. DOI: 10.1155/2013/535796
- Hashimoto M, Tong R, Kohane DS. (2013). Microdevices for Nanomedicine. *Molecular Pharmaceutics* 10: 2127–2144. DOI: 10.1021/mp300652m
- Huang H, Yang X. (2004). Synthesis of polysaccharide-stabilized gold and silver nanoparticles: a green method. *Carbohydrate Research* 339: 2627–2631. DOI: 10.1016/j.carres.2004.08.005
- Iravani S. (2011). Green synthesis of metal nanoparticles using plants. *Green Chemistry* 13: 2638. DOI: 10.1039/c1gc15386b
- Jaishree V, Gupta PD. (2012). Nanotechnology: A Revolution in Cancer Diagnosis. *Indian journal of clinical biochemistry: IJCB* 27: 214–20. DOI: 10.1007/s12291-012-0221-z
- KS, SG, TR, TtB. (2011). Biomedical potential of silver nanoparticles synthesized from calli cells of *Citrullus colocynthis* (L.) Schrad. *Journal of Nanobiotechnology* 9: 43. DOI: 10.1186/1477-3155-9-43
- Karathanasis E, Ghaghada KB. (2016). Crossing the barrier: treatment of brain tumors using nanochain particles. *Wiley interdisciplinary reviews. Nanomedicine and nanobiotechnology* 8: 678–95. DOI: 10.1002/wnan.1387
- Karimian H *et al.* (2014). *Ferulago angulata* activates intrinsic pathway of & nbsp; apoptosis in MCF-7& nbsp; cells associated with G1 & nbsp; cell cycle arrest via involvement of p21/p27. *Drug Design, Development and Therapy* 8: 1481. DOI: 10.2147/DDDT.S68818
- Klevenz H. (2004). Nanobiotechnology: From Molecules to Systems. *Engineering in Life Sciences* 4: 211–218. DOI: 10.1002/elsc.200402090
- Klein D. (2018). The Tumor Vascular Endothelium as Decision Maker in Cancer Therapy. *Frontiers in Oncology* 8: 367. DOI: 10.3389/fonc.2018.00367
- Krishnaraj C, Muthukumar P, Ramachandran R, Balakumaran MD, Kalaichelvan PT. (2014). *Acalypha indica* Linn: Biogenic synthesis of silver and gold nanoparticles and their cytotoxic effects against MDA-MB-231, human breast cancer cells. *Biotechnology Reports* 4: 42–49. DOI: 10.1016/j.btre.2014.08.002
- Kuppusamy P, Ichwan SJA, Al-Zikri PNH, Suriyah WH, Soundharrajan L, Govindan N, Maniam GP, Yusoff MM. (2016). In Vitro Anticancer Activity of Au, Ag Nanoparticles Synthesized Using *Commelina nudiflora* L. Aqueous Extract Against HCT-116 Colon Cancer Cells. *Biological Trace Element Research* 173: 297–305. DOI: 10.1007/s12011-016-0666-7
- Macdonald JS. (1999). Toxicity of 5-fluorouracil. *Oncology (Williston Park, N.Y.)* 13: 33–4.
- Maier-Hauff K, Ulrich F, Nestler D, Niehoff H, Wust P, Thiesen B, Orawa H, Budach V, Jordan A. (2011). Efficacy and safety of intratumoral thermotherapy using magnetic iron-oxide nanoparticles combined with external beam radiotherapy on patients with recurrent glioblastoma multiforme. *Journal of Neuro-Oncology* 103: 317–324. DOI: 10.1007/

- s11060-010-0389-0
- Maity P, Bepari M, Pradhan A, Baral R, Roy S, Maiti Choudhury S. (2018). Synthesis and characterization of biogenic metal nanoparticles and its cytotoxicity and anti-neoplasticity through the induction of oxidative stress, mitochondrial dysfunction and apoptosis. *Colloids and surfaces, B, Biointerfaces* 161: 111–120. DOI: 10.1016/j.colsurfb.2017.10.040
- Manzoor M, Khan AHA, Ullah R, Khan MZ, Ahmad I. (2016). Environmental Epidemiology of Cancer in South Asian Population: Risk Assessment Against Exposure to Polycyclic Aromatic Hydrocarbons and Volatile Organic Compounds. *Arabian Journal for Science and Engineering* 41: 2031–2043. DOI: 10.1007/s13369-016-2139-x
- Minai L, Yeheskely-Hayon D, Yelin D. (2013). High levels of reactive oxygen species in gold nanoparticle-targeted cancer cells following femtosecond pulse irradiation. *Scientific Reports* 3: 2146. DOI: 10.1038/srep02146
- Mukherjee S, Chowdhury D, Kotcherlakota R, Patra S, B V, Bhadra MP, Sreedhar B, Patra CR. (2014). Potential Theranostics Application of Bio-Synthesized Silver Nanoparticles (4-in-1 System). *Theranostics* 4: 316–335. DOI: 10.7150/thno.7819
- Nadagouda MN, Varma RS. (2008). Green synthesis of silver and palladium nanoparticles at room temperature using coffee and tea extract DOI: 10.1039/b804703k
- Nguyen NT-P, Nguyen LV-H, Thanh NT, Toi V Van, Ngoc Quyen T, Tran PA, Nguyen T-H. (2019). Stabilization of Silver Nanoparticles in chitosan and gelatin hydrogel and its applications. *Materials Letters* DOI: 10.1016/J.MATLET.2019.03.103
- Nowack B, Krug HF, Height M. (2011). 120 Years of Nanosilver History: Implications for Policy Makers. *Environmental Science & Technology* 45: 1177–1183. DOI: 10.1021/es103316q
- Nune SK, Chanda N, Shukla R, Katti K, Kulkarni RR, Thilakavathy S, Mekapothula S, Kannan R, Katti K V. (2009). Green nanotechnology from tea: phytochemicals in tea as building blocks for production of biocompatible gold nanoparticles. *Journal of Materials Chemistry* 19: 2912. DOI: 10.1039/b822015h
- Oves M, Aslam M, Rauf MA, Qayyum S, Qari HA, Khan MS, Alam MZ, Tabrez S, Pugazhendhi A, Ismail IML. (2018). Antimicrobial and anticancer activities of silver nanoparticles synthesized from the root hair extract of *Phoenix dactylifera*. *Materials science & engineering. C, Materials for biological applications* 89: 429–443. DOI: 10.1016/j.msec.2018.03.035
- Padinjarathil H, Joseph MM, Unnikrishnan BS, Preethi GU, Shiji R, Archana MG, Maya S, Syama HP, Sreelekha TT. (2018). Galactomannan endowed biogenic silver nanoparticles exposed enhanced cancer cytotoxicity with excellent biocompatibility. *International Journal of Biological Macromolecules* 118: 1174–1182. DOI: 10.1016/J.IJBIOMAC.2018.06.194
- Patil MP, Kang M, Niyonizigiye I, Singh A, Kim J-O, Seo YB, Kim G-D. (2019). Extracellular synthesis of gold nanoparticles using the marine bacterium *Paracoccus haendaensis* BC74171T and evaluation of their antioxidant activity and antiproliferative effect on normal and cancer cell lines. *Colloids and Surfaces B: Biointerfaces* 183: 110455. DOI: 10.1016/j.

COLSURFB.2019.110455

- Petersen GH, Alzghari SK, Chee W, Sankari SS, La-Beck NM. (2016). Meta-analysis of clinical and preclinical studies comparing the anticancer efficacy of liposomal versus conventional non-liposomal doxorubicin. *Journal of Controlled Release* 232: 255–264. DOI: 10.1016/j.jconrel.2016.04.028
- Prabhu D, Arulvasu C, Babu G, Manikandan R, Srinivasan P. (2013). Biologically synthesized green silver nanoparticles from leaf extract of *Vitex negundo* L. induce growth-inhibitory effect on human colon cancer cell line HCT15. *Process Biochemistry* 48: 317–324. DOI: 10.1016/j.PROCBIO.2012.12.013
- Rai M, Kon K, Ingle A, Duran N, Galdiero S, Galdiero M. (2014). Broad-spectrum bioactivities of silver nanoparticles: the emerging trends and future prospects. *Applied Microbiology and Biotechnology* 98: 1951–1961. DOI: 10.1007/s00253-013-5473-x
- Ramachandran R, Krishnaraj C, Sivakumar AS, Prasannakumar P, Abhay Kumar VK, Shim KS, Song C-G, Yun S-I. (2017). Anticancer activity of biologically synthesized silver and gold nanoparticles on mouse myoblast cancer cells and their toxicity against embryonic zebrafish. *Materials Science and Engineering: C* 73: 674–683. DOI: 10.1016/j.MSEC.2016.12.110
- Rengan AK, Bukhari AB, Pradhan A, Malhotra R, Banerjee R, Srivastava R, De A. (2015). In Vivo Analysis of Biodegradable Liposome Gold Nanoparticles as Efficient Agents for Photothermal Therapy of Cancer. *Nano Letters* 15: 842–848. DOI: 10.1021/nl5045378
- des Rieux A, Pourcelle V, Cani PD, Marchand-Brynaert J, Pr at V. (2013). Targeted nanoparticles with novel non-peptidic ligands for oral delivery. *Advanced drug delivery reviews* 65: 833–44. DOI: 10.1016/j.addr.2013.01.002
- Rogers KR *et al.* (2018). Characterization of engineered nanoparticles in commercially available spray disinfectant products advertised to contain colloidal silver. *Science of The Total Environment* 619–620: 1375–1384. DOI: 10.1016/j.scitotenv.2017.11.195
- Rolim WR, Pieretti JC, Ren  DLS, Lima BA, Nascimento MHM, Ambrosio FN, Lombello CB, Brocchi M, de Souza ACS, Seabra AB. (2019). Antimicrobial Activity and Cytotoxicity to Tumor Cells of Nitric Oxide Donor and Silver Nanoparticles Containing PVA/PEG Films for Topical Applications. *ACS Applied Materials & Interfaces* 11: 6589–6604. DOI: 10.1021/acsami.8b19021
- Saber MM, Mirtajani SB, Karimzadeh K. (2018). Green synthesis of silver nanoparticles using *Trapa natans* extract and their anticancer activity against A431 human skin cancer cells. *Journal of Drug Delivery Science and Technology* 47: 375–379. DOI: 10.1016/j.JDDST.2018.08.004
- Saha RN, Vasanthakumar S, Bende G, Snehalatha M. (2010). Nanoparticulate drug delivery systems for cancer chemotherapy. *Molecular membrane biology* 27: 215–31. DOI: 10.3109/09687688.2010.510804
- Salata O. (2004). Applications of nanoparticles in biology and medicine. *Journal of Nanobiotechnology* 2: 3. DOI: 10.1186/1477-3155-2-3
- Sattarahmady N, Movahedpour A, Heli H, Hatam GR. (2016). Gold nanoparticles-based biosensing of *Leishmania major* kDNA genome:

- Visual and spectrophotometric detections. *Sensors and Actuators B: Chemical* 235: 723–731. DOI: 10.1016/j.snb.2016.05.023
- Shen S, Xia J-X, Wang J. (2016). Nanomedicine-mediated cancer stem cell therapy. *Biomaterials* 74: 1–18. DOI: 10.1016/j.biomaterials.2015.09.037
- Singh P, Kim Y-J, Zhang D, Yang D-C. (2016). Biological Synthesis of Nanoparticles from Plants and Microorganisms. *Trends in Biotechnology* 34: 588–599. DOI: 10.1016/j.tibtech.2016.02.006
- Stensberg MC, Wei Q, McLamore ES, Porterfield DM, Wei A, Sepulveda MS. (2011). Toxicological studies on silver nanoparticles: challenges and opportunities in assessment, monitoring and imaging. *Nanomedicine* 6: 879–898. DOI: 10.2217/nnm.11.78
- Swartz HM *et al.* (2007). In vivo EPR for dosimetry. *Radiation Measurements* 42: 1075–1084. DOI: 10.1016/j.radmeas.2007.05.023
- Syed A, Saraswati S, Kundu GC, Ahmad A. (2013). Biological synthesis of silver nanoparticles using the fungus *Humicola* sp. and evaluation of their cytotoxicity using normal and cancer cell lines. *Spectrochimica Acta Part A: Molecular and Biomolecular Spectroscopy* 114: 144–147. DOI: 10.1016/j.SAA.2013.05.030
- Troupis A, Hiskia A, Papaconstantinou E. (2002). Synthesis of Metal Nanoparticles by Using Polyoxometalates as Photocatalysts and Stabilizers We thank the Ministry of Development, General Secretariat of Research and Technology of Greece, for supporting part of this work. We also thank Dr. A. Travlos for help with the TEM images. *Angewandte Chemie International Edition* 41: 1911. DOI: 10.1002/1521-3773(20020603)41:11<1911::AID-ANIE1911>3.0.CO;2-0
- Ullah Khan S, Saleh TA, Wahab A, Ullah Khan MH, Khan D, Ullah Khan W, Rahim A, Kamal S, Ullah Khan F, Fahad S. (2018). Nanosilver: new ageless and versatile biomedical therapeutic scaffold. *International Journal of Nanomedicine* Volume 13: 733–762. DOI: 10.2147/IJNS153167
- Vemuri SK, Banala RR, Mukherjee S, Uppala P, GPV S, A.V. GR, T. M. (2019). Novel biosynthesized gold nanoparticles as anti-cancer agents against breast cancer: Synthesis, biological evaluation, molecular modelling studies. *Materials Science and Engineering: C* 99: 417–429. DOI: 10.1016/j.MSEC.2019.01.123
- Vimalraj S, Ashokkumar T, Saravanan S. (2018). Biogenic gold nanoparticles synthesis mediated by *Mangifera indica* seed aqueous extracts exhibits antibacterial, anticancer and anti-angiogenic properties. *Biomedicine & Pharmacotherapy* 105: 440–448. DOI: 10.1016/j.biopha.2018.05.151
- Vinoshia M, Palanisamy S, Muthukrishnan R, Selvam S, Kannapiran E, You S, Prabhu NM. (2019). Biogenic synthesis of gold nanoparticles from *Halymenia dilatata* for pharmaceutical applications: Antioxidant, anti-cancer and antibacterial activities. *Process Biochemistry* 85: 219–229. DOI: 10.1016/j.PROCBIO.2019.07.013
- Wasukan N, Srisung S, Kuno M, Kulthong K, Maniratanachote R. (2015). Interaction evaluation of silver and dithizone complexes using DFT calculations and NMR analysis. *Spectrochimica acta. Part A. Molecular and biomolecular spectroscopy* 149: 830–8. DOI: 10.1016/j.saa.2015.04.064
- Wu M, Ma B, Pan T, Chen S, Sun J. (2016). Silver-Nanoparticle-Colored Cotton

Fabrics with Tunable Colors and Durable Antibacterial and Self-Healing Superhydrophobic Properties. *Advanced Functional Materials* 26: 569–576. DOI: 10.1002/adfm.201504197

Yadav A, Mendhulkar VD. (2018). Antiproliferative activity of *Camellia sinensis* mediated silver nanoparticles on three different human cancer cell lines. *Journal of cancer research and therapeutics* 14: 1316–1324. DOI: 10.4103/jcrt.JCRT\_575\_16

Yu B, Tai HC, Xue W, Lee LJ, Lee RJ. (2010). Receptor-targeted nanocarriers for therapeutic delivery to cancer. *Molecular Membrane Biology* 27: 286–298. DOI: 10.3109/09687688.2010.521200

Zada S *et al.* (2018). Biogenic synthesis of silver nanoparticles using extracts of *Leptolyngbya* JSC-1 that induce apoptosis in HeLa cell line and exterminate pathogenic bacteria. *Artificial Cells, Nanomedicine, and Biotechnology* 46: S471–S480. DOI: 10.1080/21691401.2018.1499663

Zhou C, Long M, Qin Y, Sun X, Zheng J. (2011). Luminescent Gold Nanoparticles with Efficient Renal Clearance. *Angewandte Chemie International Edition* 50: 3168–3172. DOI: 10.1002/anie.201007321.

# INDIA VIETNAM

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DEFENCE, STRATEGIC AND  
ECONOMIC COOPERATION

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JAYACHANDRA REDDY G  
RAMESH BABU V

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## Trends in India's Trade with Vietnam: An Empirical Analysis

*Venkata Prasad A and Krishnamoorthy D*

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### Abstract

India and Vietnam share close to half a century of relations with the establishment of formal diplomatic ties that began in 1972 between India and the then-North Vietnam. The relations were derived from common historical linkages as colonies of Western powers striving to achieve independence. Bilateral ties between India and Vietnam have got strengthened in recent years with a focus on regional security issues and trade. The adoption of the 'Look East Policy', imbued with the economic element, marked an important turning point in India-Vietnam economic relations. Vietnam has played an important role in India's "Act East" policy, which was enunciated as an upgrading of the earlier 'Look East Policy' and also she has been a crucial partner in sub-regional, regional and multilateral forums. The need for economic development increased the importance of bilateral economic and commercial linkages between both the countries. These efforts received fillip from regional arrangements, as for example, ASEAN-India cooperation and Mekong - Ganga Cooperation. India is Vietnam's fifth largest trading partner in ASEAN; Vietnam holds the 19<sup>th</sup> position worldwide for India, both the countries made conscious efforts to expand the areas of economic cooperation and to increase the volume of trade and investment. India's total trade comprising exports and imports skyrocketed from the abysmal level of US \$0.24 billion to the magnificent level of US \$12.27 billion between the years 2001 and 2017. India and Vietnam will explore "substantive and practical measures" to achieve the bilateral trade target of \$15 billion by 2020. Currently, India has 182 investment projects worth US \$816 million in Vietnam, while Vietnamese businesses have a mere seven projects worth some US \$6 million in India. The present paper reviews the performance of India's merchandise trade with Vietnam during 2000-01 to 2015-16 on the basis of the data released by the Ministry of Commerce and Industry, Government of India.

**Key Words:** ASEAN, Trade, Growth Rates, Commodity and Act East Policy



# Emerging Trends in Nanotechnology

## Innovations, Health and Risks

**S. Janardana Reddy**

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# A Critical Review on Potent Prospects of Biogenic Metallic Nanoparticles in Cancer Therapy

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## ABSTRACT

Cancer is one of the deadliest diseases all over the globe, causing social and economic inequalities among people. The incidence is being increased each year and diagnosed with different types of cancers with drug resistance followed by metastasis. Many cancer cells have type specific cancer protein all over their surface, which is lacking in normal cells. The nanoparticles like metal, Biogenic metal and other types conjugate or bind to the cancer cells and helps to increase the specific antibodies to unravel the treatment mechanisms. Many works indicate that the nanoparticles help in drug delivery systems as a result unique cancer treatment will be enabled. Therefore, nanotherapeutics is rapidly developing in cancer treatment, diagnostics, biomarker identification, detection of multiple genes and matrix RNA. The conjugated nanoparticles allow to detect cancer related proteins, resulting in to new method of analyzing proteome of each tumor.

# Biogenic Nanoparticles: A Comprehensive Review to Explore Multidrug Resistance Mechanisms among Microbes

N. VASAVI<sup>1</sup>, N. RAVI KUMAR<sup>1</sup>, A. T. VENKATRAMANA  
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## ABSTRACT

Microbial resistance has been an intense hindrance in preventing diseases for many decades. The rapid emergence of resistance towards several antibiotics could help the bacteria to become stronger to the existing antibiotics and becoming fatal to the mankind. This multidrug resistance is posing a challenging threat to human beings globally due to indiscriminate use of synthetic antibiotics and several other chemical compounds. Several factors have been influencing in developing resistance either by vertical or horizontal gene transfer among microbial species through different ecosystems. Microorganisms are altering genetic makeup to tackle the existing antibiotics very rapidly. Therefore, there is an urgent need to overcome this resistance and to develop new forms of antibiotics that are cost-effective, biocompatible, showing fewer side effects and a single-step fabricated approach that helps for large scale production. Biogenic mediated metallic nanoparticles became a

## ABOUT THE BOOK

Nanotechnology is an emerging interdisciplinary technology that has been flourishing in many areas during the recent decade, including material science, mechanics, electronics, optics, medicine, physics, energy electronics, biology, agriculture, nanosensors and nanorobots.

The applications of engineered nanomaterials are developing exponentially, along with the controversies at government, industry and public groups of nanotechnology issues. There is also growing public concern caused by negative perceptions among some high profile groups that nano-enabled products are being marketed indiscriminately and being released without adequate testing of their safety.

Many of the new particles presently contemplated harmless are likely to acquire unique properties when contained to a nanoscale size and could manifest toxic biological effects. It is well known that the toxicity of engineered nanoparticles (ENPs) may depend on the physical and chemical characteristics of the particles.

Nanotechnology is growing rapidly in industrial applications, medical imaging, cancer diagnosis, drug delivery, cancer treatment, and gene therapy, and also to aid in visual imaging. Nanotechnology is at the cutting edge of rapid healthcare product development as it has many potential human health benefits, but it is essential with some apprehensions for its potential human health risks.



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## ABSTRACT

Cancer is one of the deadliest diseases all over the globe, causing social and economic inequalities among people. The incidence is being increased each year and diagnosed with different types of cancers with drug resistance followed by metastasis. Many cancer cells have type specific cancer protein all over their surface, which is lacking in normal cells. The nanoparticles like metal, Biogenic metal and other types conjugate or bind to the cancer cells and helps to increase the specific antibodies to unravel the treatment mechanisms. Many works indicate that the nanoparticles help in drug delivery systems as a result unique cancer treatment will be enabled. Therefore, nanotherapeutics is rapidly developing in cancer treatment, diagnostics, biomarker identification, detection of multiple genes and matrix RNA. The conjugated nanoparticles allow to detect cancer related proteins, resulting in to new method of analyzing proteome of each tumor

## ABOUT THE BOOK

Nanotechnology is an emerging interdisciplinary technology that has been flowering in many areas during the recent decade, including material science, mechanics, electronics, optics, medicine, physics, energy, electronics, biology, agriculture, biotechnology and aerospace.

The approaches for engineered nanomaterials are developing exponentially, along with the controversies in government, industry and public groups of nanosafety issues. There is also growing public concern caused by negative perceptions among some high profile groups that nano-enabled products are proliferating uncontrollably and being released without adequate testing of their safety.

Many of the free particles presently contemplated harmless are likely to acquire unique properties when condensed to a nanomolecular size and could manifest toxic biological effects. It is well known that the toxicity of engineered nanoparticles (NPs) may depend on the physical and chemical characteristics of the particles.

Nanotechnology is evolving rapidly in industrial applications, medical imaging, disease diagnosis, drug delivery, cancer treatment, and gene therapy, and also to aid in visual imaging. Nanotechnology is at the cutting edge of rapid healthcare product development as it has many potential human health benefits, but it is perceived with some apprehension for its potential human health risks.



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# CHANGING INFORMATION LANDSCAPE AND ITS TRANSFORMATION IN LIS EDUCATION

**Editors**

**Prof. S. Thanuskodi**

**Dr. S. Kishore Kumar**

**Dr. S. Raja**

**Dr. A. Alagu**



**Department of Library and Information Science**  
**ALAGAPPA UNIVERSITY**

(A State University Accredited with A+ Grade by NAAC (CGPA: 3.64)  
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## USE OF INFORMATION TECHNOLOGY AMONG POSTGRADUATE PROFESSIONAL COURSE STUDENTS IN RAYALASEEMA AREA, ANDHRA PRADESH: A STUDY

Dr. Sudhakar, E. & Kiran Kumar, E.

### Abstract

*In this paper researchers studied Use of Information Technology among Postgraduate Professional Course Students in Rayalaseema Area, Andhra Pradesh: A Study, distributed 2000 questionnaires in colleges and collected 1655 questionnaires. It is found in this study the Use of information Technology library information resources, services (electronic & print), and library expressed their opinion about library working hours, physical facility, library information resources and services excellent and good.*

**Keywords:** Use of Information Technology; Professional colleges; Library and Information resources; ICT sources and services; Rayalaseema Area.

### Introduction

Education is the passport to accelerated economic growth. It is the key to build human capital. Human capital is the vital ingredient in building a nation. Education is a crucial building block, that recognized as pre-requisite for the general development programme. Since long time immemorial education formed a continuous basis for the development of society, "Education in whatever form has to respond expeditiously and effectively to the changing needs in an ever-changing society" as change is the only unchanging law. Education is a process of learning with the aim to develop the capabilities in the people at all levels. In this process, schools, colleges and universities play a vital role. The present educational system in India comprises both academic and professional studies by offering courses leading to bachelors and master's degree.

### Objectives of the Study

The following are the specific objectives of the study:

- To know the purpose of Information Technology services using by Postgraduate Professional course students in the library.
- To assess the suitable place to access the Internet services to the students.
- To understand the Utilization of different Internet services by the students.
- To identify the problems faced by Postgraduate students in accessing Information Technology services.
- To examine the overall satisfaction of postgraduate students with IT based Information services.

### Methodology

The present study focuses on the availability of Information technology resources, services and their utilization in professional college libraries in Rayalaseema area.

### Selection of Sample

Selection of sample is carried out at two levels namely college level and user level. The libraries of postgraduate professional colleges which are conducting M.Tech, MBA and MCA courses are known as postgraduate professional college libraries. There are 32 postgraduate professional college libraries in Rayalaseema area of Andhra Pradesh established on or before the year 2000. Rayalaseema area covers the districts of Anantapur, Kurumool, Y.S.R (Kadapa) and Chittoor. All 32 postgraduate colleges were taken for the study. The libraries which are attached to colleges are established in the year on 2000 before 2000 as sample for the present study. The population of this study consists of postgraduate students of these college libraries. There are a total number of 4204 users registered in the college libraries. As the population is large in terms of cost, time and labour involved, the investigator selected a sample of 1655 users out of 4204 users as using proportionate random sampling method.

### Data analysis and Interpretation

Table 1: Distribution of users according to their gender

S.No.	Gender Wise	Frequency	Percentage
1	Male	962	58.12
2	Female	693	41.87
	<b>Total</b>	<b>1655</b>	<b>100</b>

It is evident from the table-8.1 that out of total respondents gender-wise distribution of the M.Tech, MCA and MBA students 1655 respondents surveyed, majority of the users (58.12%) are males and remaining (41.87) of there are females. Hence it can be concluded that majority of the users (58.12%) are males and remaining (41.87) of there are females.

### Education qualifications

The success of IT application in libraries totally depends on the extent of users assistance and their satisfaction. User community should always be kept well informed about their IT application and its developments at regular intervals so that they can understand and make optimum use of IT based library services.

Table 2: Distribution of respondents according to their education qualifications

S.No.	Education	Frequency	Percentage
1	M.Tech	626	37.82
2	MCA	532	32.14
3	MBA	497	30.03
	<b>Total</b>	<b>1655</b>	<b>100</b>

From the above table it can be observed that 37.82% are having M. Tech qualification, 32.14% are having MCA qualification and the remaining 30.03% are having MBA qualification. Hence it can be concluded that a high percentage of respondents are studying M.Tech qualification, the remaining respondents are studying MCA and MBA.

### Frequency of visit to the library

Professional College Libraries is a place for research and education. Users visit the library to get their required/need information. When users get needed information, they get full satisfaction and visit the library again and again. Library may provide electronic information resources and services.

**Table 3. Frequency of visit to the library**

S.No.	Frequently	M.Tech	MCA	MBA	Total
1	Daily	156(9.42%)	122(7.37%)	102(6.16%)	380(22.96%)
2	Once in a week	198(11.96%)	185(11.17%)	153(9.24%)	536(32.38%)
3	Twice in week	188(11.36%)	149(9.0%)	159(9.60%)	496(29.96%)
4	Once in month	84(5.07%)	76(4.59%)	83(5.02%)	243(14.68%)
<b>Total</b>		<b>626(37.82%)</b>	<b>532(32.14%)</b>	<b>497(30.03%)</b>	<b>1655(100%)</b>

Table 3. shows that frequency of visiting library majority of 536(32.38%) respondents visit library once in a week, followed by 496(29.96%) respondents visit library twice in a week, 380(22.96%) respondents visit library daily. The remaining respondents visit library 243(14.68%) once in a month. Hence it can be concluded that majority of the respondents visit library 536 (32.38%) once in a week, followed by 496(29.96%) visit library twice in a week

### Average time spent in the library per day

Professional College Libraries Serious readers normally spend more time in libraries. Libraries are by their very nature, the centers for the spread of knowledge and information.

**Table 4: Time Spent in the Library Per Day**

S.No.	Time spent	M.Tech	MCA	MBA	Total
1	Less than one hour	86(5.19%)	78(4.71%)	74(4.47%)	238(14.38%)
2	1to 2 hours	327(19.75%)	183(11.05%)	166(10.03%)	676(40.85%)
3	2to 3 hours	124(7.49%)	135(8.15%)	143(8.64%)	402(24.29%)
4	More than 3 hours	89(5.38%)	136(8.21%)	114(6.88%)	339(20.48%)
<b>Total</b>		<b>626(37.82%)</b>	<b>532(32.14%)</b>	<b>497(30.03%)</b>	<b>1655(100%)</b>

The duration of visit is taken as less than one hour, 1 to 2 hours, 2 to 3 hours, more than 3 hours. From the above table 4 shows that 676 (40.85%) of respondents spent time 1 to 2 hours per day in the library, followed by 402 (24.29%) spent time 2 to 3 hours, 339(20.48%) spent more than 3 hours and the remaining 238(14.38%) spent time less than 1 hour in the library. Hence it can be concluded that majority of the respondents 40.85% spent time 1 to 2 hours per day in the library, followed by 24.29% spent time 2 to 3 hours.

### Most frequently used Location of Internet

Electronic resources can be accessed from anywhere in the world through Internet. Generally College library users access Internet and e-resources from library, department, or in campus at different locations. The following table presents the location, from where the users generally access internet resources. All the libraries that are taken for the study have Internet facility. Internet can be accessed via different service providers.

**Table 5: Most frequently used Location of Internet access by the users**

S.No.	Location of Internet access	Frequency	Percentage
1	Home	213	12.87
2	Library	374	23.80
3	College	292	17.64
4	Internet cafe	653	39.45
5	College hostel	123	7.43
<b>Total</b>		<b>1655</b>	<b>100</b>

It is evident from the table 5 that majority of the respondents 653(39.45%) prefer to access Internet in Internet café, 374(23.80%) of the respondents prefer using Internet access from College Library, 292(17.64%) of the respondents using Internet in college, 213(12.87%) of the respondents using Internet in Home and remaining 123(7.43%) of the respondents using Internet in College hostels. Hence it can be concluded that majority of the students 39.45% prefer to access Internet in Internet café, followed by 23.80% of the respondents prefer using Internet access from College Library.

### Facilities and services of Internet used

Internet is a dynamic source of information. Users' opinion regarding adequacy of internet accessing were sought and presented here. The users were asked about frequency of access to internet for different purpose.

**Table 6. Most frequently using services on Internet by the respondents**

S.No	Purpose	Frequency	Percentage
1	Web search engines	252	15.23
2	E-mail	218	13.17
3	Online reference sources	116	7
4	Online indexing, abstracting and full text databases	126	7.16
5	Online library catalogs (OPAC/Web OPACs)	79	4.77
6	Open access electronic journals and magazines	165	9.97
7	Online newspapers	95	5.74
8	Virtual libraries	85	5.13
9	Other libraries' websites	41	2.48
10	Online books	96	5.8
11	Web directories / Subject directories	35	2.11

12	Online bookstores / publishers	38	2.29
13	Online audio, video and multi-media collections	128	7.73
14	Blogs / Weblogs	78	4.71
15	Chatting	28	1.69
16	Others	75	4.53
<b>Total</b>		<b>1655</b>	<b>100</b>

Table 6 depicts that the majority of the respondents 252(15.23%) used web search engines. followed by 218(13.23%) of the respondents using e-mail, 165(9.97%) of the respondents using open access electronic journals and magazines, 128(7.73%) of the respondents using Online audio, video and multi-media collections, 126(7.16%) of the respondents using Online indexing, abstracting and full text databases, 116(7%) of the respondents using Online reference sources, 96(5.74%) of the respondents using online books, 95(5.74%) of the respondents using Online newspapers, 85(5.13%) of the respondents using Virtual libraries, 79(4.77%) of the respondents using Online library catalogs (OPAC/Web OPACs), 78(4.7%) of the respondents using Blogs / Weblogs, 75 (4.77%) of the respondents using other resources, 41 (2.48%) of the respondents using other libraries' websites, 38 (2.29%) of the respondents using Online bookstores / publishers, 35 (2.11%) of the respondents using Web directories / Subject directories and remaining 28(1.69%) of the respondents using chatting. Hence it can be concluded that majority of the respondents 252(15.23%) used web search engines. followed by 218(13.23%) of the respondents using e-mail.

#### Sources used academic Information needs

Table 7. Distribution of respondents according to their sources used

S. No	Academic information	Frequency	Percentage
1.	Google	818	49.42
2.	Classmates/Friends through Social Networking	433	26.16
3.	Professional Societies	152	9.18
4.	Databases	138	8.34
5.	Other websites	114	6.89
<b>Total</b>		<b>1655</b>	<b>100</b>

It is evident from the table 8 that majority of the respondents use for academic information needs 811 (49.42%) Google, followed by 433(26.16%) of respondents information needs classmates/friends through social networking, 152(9.18%) respondents information needs in professional societies, 138(8.34%) of the respondents using information needs for databases and remaining 114(6.89%) of the respondents using academic information needs for other websites. Hence it can be concluded that majority of the respondents use for academic information needs 811 (49.42%) Google, followed by 433(26.16%) of respondents information needs classmates/friends through social networking.

### **Suggestions**

A few suggestions are put forward by the researcher based on the observations gathered from the study.

- Need to Improve IT facilities, Provide needful info on usage of internet, WI-FI should be provided college campus.
- Develop IT communication in library, Increase computers with high band width and more computers.
- Better to have a digital library, Give training classes to students, Improve facilities of study oriented programs, Must improve IT in library and speed internet.

### **Conclusion**

All most all professional college libraries provide user training programs and computer literacy skills programs to their users. Majority of the respondents state that the library staff has helped you to utilize the Information Technology products in your library. The respondents (Students) stated that the user training and computer literacy programme is very useful for using of e-resources provided by their libraries

### **References**

- Akobundu, D.U.(2008). Availability and accessibility of information resources and use of library resources at Michael Okpara University of Agriculture. *Library Philosophy and Practice*, 1-8.
- Anjaiah, M. and Nageshwara Rao, P. (2015) Use of scholarly electronic information resources by faculty members of NBA accredited engineering college libraries: a survey, *International Journal of Innovation Sciences and Research* , 4 (11), 524-531.
- Chikkanmanju., and Kumbar, M. (2015). Use of Information Resources and Services by the Students of First Grade Colleges Affiliated to Tumkur University, Tumkur: A Comparative Study. *International Journal of Academic Library and Information Science*, 3(2), 53-64.
- Jaspal Kaur (2012). Use of electronic resources by teachers of degree colleges in Chandigarh. *Kelpro Bulletin*, 3(1), 23-27.
- Kaur, Baljinder, and Rama Verma (2009). Use of Electronic Information Resources: A Case Study of Thapar University. *DESIDOC Journal of Library and Information Technology*, 29(2), 67-73.

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# Transformation of Learning Resource Centres in the Digital Era

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Dr.A.M.Venkatachalam  
Dr.K.Karunai Raghavan  
Dr.M.Mandhirasalam  
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2022

## **Transformation of Learning Resource Centres in the Digital Era**

Department of Library, K.S.Rangasamy College of Technology (KSRCT)  
&  
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# Awareness and Usage of E-Shodhsindhu Consortium by Faculty and Research Scholars of Nagarjuna University: A Study

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## Abstract

The term "virtual library" refers to a library without walls where customers can access any sort of knowledge in electronic format without restriction. A digital library is a great resource for academic communities since it allows them to access electronic journals, databases, and e-books. The e-ShodhSindhu is that the Consortium for educational e-resources merging three consortia, i.e. UGC-INFONET Digital Library Consortium, INDEST-AICTE Consortium and N-LIST in December 2013. The e-ShodhSindhu is supposed to provide current and depository access to E-Resources i.e. e-books, e-journals, e-journal archives, on-line databases to its member establishments at a negotiated rate of subscription. The main purpose of this paper is to find out user awareness and usage of e-ShodhSindhu Consortium by Faculty Members and Research Scholars of AcharyaNagarjuna University (AU), in Andhra Pradesh.

**Keywords:** Virtual Library, e-ShodhSindhu, Digital Library, e-Resources

## Introduction

The significance of the name Acharya Nagarjuna University comes from the fact that it was established by the great Buddhist preceptor and philosopher Acharya Nagarjuna on the banks of the Krishna River many years ago. He turned it into a great centre of learning that drew many teachers and students from all over the world. The University was established by Act 43 of 1976 of the A.P. State Legislature and Governed by Act 4 of 1991 covering 6 Universities in the State.

The present Acharya Nagarjuna University is not very far from the haloed spot, as it derives its moral and intellectual sustenance from Acharya Nagarjuna in which context the University has been renamed Acharya Nagarjuna University through the A.P. Universities (Amendment) Ordinance, 2004, promulgated by the Governor of Andhra Pradesh. The aspirations of the southern coastal districts of Andhra Pradesh to pursue Post-Graduate education nearer their homes resulted in the establishment of Acharya Nagarjuna University.

## Dr.B.R.Ambedkar Memorial Library

The library building's plinth area is roughly 3,000 square feet in total. The Circulation, Periodicals, Reference, Acquisition, Binding, and Technical Sections, in addition to the Administrative and Computers Sections, perform the majority of the library's functions. The Dr. B.R.A.M. Library is open from 8 a.m. to 8 p.m. on all working days and from 8 a.m. to 1 p.m. on the first Saturday and Sunday of each month. For seating and reading purposes, the library mostly uses wooden chairs and tables. Steel book racks that meet specific criteria are utilized for shelving. For the year 2016, a total of Rs. 57,72,079 was spent on 270 publications, including daily newspapers, general magazines, and foreign and Indian periodicals.

### Objectives of the Library

- To inspire and encourage all who would seek knowledge through higher education and research.
- To provide quality instruction and research for the advancement of science and technology.
- To promote teaching and research studies in disciplines of societal relevance.
- To bridge the gap between theory and practice of the principles of higher education.
- To develop the human talent necessary for the industry.

**Table 1: Dr.B.R.Ambedkar Memorial Library Collection**

SLNo.	ResourceName	Description of Resource/URL	Total
<b>1. Print and Non-Print Resources</b>			
1.1	Books	TextBooks	1,35,796
1.2	ForeignJournals(Print)	AllSubjects	49
1.3	IndianJournals(Print)	AllSubjects	162
1.4	GeneralMagazines	English,Telugu, etc	28
1.5	Newspapers	English,Telugu, etc	15
1.6	BackVolumes	AllSubjects	2766
1.7	CD-ROMs	MagazinesandSubjectRelated	471
<b>2. University subscribed e-Resources</b>			
2.1	E-ShodhSindhu	<a href="http://www.inflibnet.ac.in/ess/eres.php?memID=93">http://www.inflibnet.ac.in/ess/eres.php?memID=93</a>	-
2.2	DELNET	<a href="http://www.delnet.nic.in/">http://www.delnet.nic.in/</a>	-
<b>3. e-Journals subscribed for the year 2021 through E-ShodhSindhu</b>			
3.1	AmericanChemical Society	<a href="https://pubs.acs.org/">https://pubs.acs.org/</a>	51
3.2	AnnualReviews	<a href="https://www.annualreviews.org/">https://www.annualreviews.org/</a>	43
3.3	EconomicandPolitical Weekly	<a href="https://www.epw.in/">https://www.epw.in/</a>	1
3.4	JSTOR	<a href="https://www.jstor.org/">https://www.jstor.org/</a>	3165
3.5	SpringerLink	<a href="https://link.springer.com/">https://link.springer.com/</a>	1725
3.6	Taylor&Francis	<a href="http://www.tandfonline.com/">http://www.tandfonline.com/</a>	1078
<b>4. Databases subscribed for the year 2021 through E-ShodhSindhu</b>			
4.1	WebofScience	<a href="http://webofknowledge.com/">http://webofknowledge.com/</a>	-
4.2	InstituteofStudiesin Industrialdevelopment	<a href="http://isid.org.in/">http://isid.org.in/</a>	-
4.3	J-gatePlus	<a href="https://jgateplus.com/search/">https://jgateplus.com/search/</a>	-

The variables used for this study are broadly divided into two, namely classificatory variables and study variables. The variables are selected by the literature reviewed for the purpose.

- University
- Gender of the respondent
- Age of the respondent
- Academic position of the respondent
- The academic area of the respondent

### Objectives of the Study

The proposed work considered Science and Social Science Research's magnitude in Nagarjuna universities for E-ShodhSindhu effects. Before deciding the research priorities for this study, previous research studies were utilized. An analysis of prior results has been undertaken, and some differences in the past research were identified. Based on the identified variations, the following research objectives are formulated by the researcher:

- To know the users' awareness about E-ShodhSindhu resources available in Nagarjuna university
- To know the usage pattern of the E-ShodhSindhu
- To identify the intent of accessing E-ShodhSindhu electronic resources.
- To evaluate the impact of E-ShodhSindhu electronic resources on the academic community of higher education.
- To find out the strategies adopted by user store cooperate data through E-ShodhSindhu electronic resources.
- To determine the user's challenges and problems while using the E-ShodhSindhu electronic resources.
- To find out the required training programs to improve the usage of E-ShodhSindhu electronic resources.
- To analyse the satisfactory level the users while using E-ShodhSindhu electronic resources
- To study the infrastructural facilities provided by the University Libraries associated with the E-ShodhSindhu electronic resources.

### Population of the Study

The research excludes PG students and administrative employees. This population is not having much awareness and usage of E-Shodh Sindhu. To obtain samples from both classes, the investigator divided the users into two categories based on their academic status and picked preventative sample from each category to perform the analysis. As a result, the following groups have been formed.

**Faculty** – Comprised faculty members from the relevant university departments.

**Research Scholars** – Includes those pursuing Ph.D. and M.Phil. Degrees in university departments

**Table 2: Total Population in the Study Area**

Variables	Status of the Respondents	ANU	% of Total Population
<b>Position of Respondents</b>	Professors	28	4.32
	Associate Professors	41	2.34
	Assistant Professors	34	10.65
	Research Scholars	1238	82.69
	<b>Total</b>	<b>1341</b>	<b>100</b>
<b>Gender</b>	Male	863	51.52
	Female	478	48.48
	<b>Total</b>	<b>1341</b>	<b>100</b>
<b>Academic area</b>	Science	686	47.18
	Social Sciences	655	52.82
	<b>Total</b>	<b>1341</b>	<b>100</b>
<b>Grand Total of Population</b>		<b>1341</b>	

From the above table we understood that the total population of ANU is 1341 and faculty are 103 and scholars are 1238.

**Table 3: Distribution of Respondents based on Subject of Study and Universities**

Subject		Acharya Nagarjuna University (ANU)
Science	N	104
	%	45.81
Social Sciences	N	123
	%	54.19
Total	N	227
	%	100.00

**Table 4: Source of Information the Respondents Generally Consult**

Source of Information		Academic Status		Universities
		Faculty	Research Scholars	ANU
University Library	N	38	153	42
	%	21.71	21.34	18.50
Internet	N	71	308	107
	%	40.57	42.96	47.14
Interaction with Friends / Colleagues	N	31	86	21
	%	17.71	11.99	9.25
Interaction with Teachers	N	25	118	44
	%	14.29	16.46	19.38
Family Members	N	4	36	5
	%	2.29	5.02	2.20
Personal Book Collection	N	6	16	8
	%	3.43	2.23	3.52
Total	N	175	717	227
	%	100	100	100

**Table 5: Availability of Library Consortium Facility based on Academic Status, Subject of Study and University**

Available Library consortium		Academic Status		Subject of Study		University
		Faculty	Research Scholars	Science	Social Sciences	ANU
E-Shodh Sindhu	N	68	322	181	209	158
	%	38.86	44.91	41.14	46.24	69.60
DELNET	N	37	161	96	102	34
	%	21.14	22.45	21.82	22.57	14.98
Any Other	N	70	234	163	141	35
	%	40.00	32.64	37.05	31.19	15.42

**Table 6: Frequency of Visiting E-ShodhSindhu Resources**

Frequency of Visit		Academic Status		Subject of Study		ANU
		Faculty	Research Scholars	Science	Social Sciences	
Daily	N	23	54	53	24	17
	%	13.14	7.53	12.05	5.31	7.05
Once in a week	N	46	27	27	46	20
	%	26.29	3.77	6.14	10.18	8.81
2-3 times in a week	N	43	241	152	132	114
	%	24.57	33.61	34.55	29.20	50.22
2-3 times in a month	N	39	223	106	156	52
	%	22.29	31.10	24.09	34.51	22.91
Occasionally	N	24	172	102	94	24
	%	13.17	23.99	23.18	20.80	10.54

**Table 7: Respondents spending time on E-ShodhSindhu Resources**

Time spent in E-Shodh Sindhu		Academic Status		Subject of Study		SPMU
		Faculty	Research Scholars	Science	Social Sciences	
Less than One hour	N	47	296	188	155	47
	%	26.86	41.28	42.73	34.29	35.61
1-2 hours	N	79	246	142	183	64
	%	45.14	34.31	32.27	40.49	48.48
2-3 hours	N	27	142	85	84	16
	%	15.43	19.80	19.32	18.58	12.12
More than 3 hours	N	22	33	25	30	5
	%	12.57	4.60	5.68	6.64	3.79
Total	N	175	717	440	452	132
	%	100.00	100.00	100.00	100.00	100.00

**Table 8: Usage of E-Journals based on Academic Status of Acharya Nagarjuna University**

Usage of E-Journals	Academic Status																
	Faculty (N=86)					WS	WAM Score	Rank	Research Scholars (N=192)					WS	WAM Score	Rank	
	1	2	3	4	5				1	2	3	4	5				
American Chemical Society	N	9	10	6	5	5	92	2.69	4	22	44	41	61	24	597	3.11	6
	%	25.71	28.57	17.14	14.29	14.29				11.46	22.92	21.33	31.77	12.50			
Annual Reviews	N	17	5	7	3	3	75	2.14	6	42	10	26	65	49	645	3.36	3
	%	48.57	14.29	20.00	8.57	8.57				21.88	5.21	13.54	33.85	25.52			
Economic and Political Weekly	N	10	12	4	3	6	88	2.51	6	21	16	69	30	56	660	3.44	1
	%	28.57	34.29	11.43	8.57	17.14				10.94	8.33	35.94	15.63	29.17			
ISTOR	N	7	6	6	7	9	110	3.14	1	35	24	22	61	50	643	3.35	4
	%	20.00	17.14	17.14	20.00	25.71				18.23	12.50	11.46	31.77	26.04			
Springer Link	N	7	6	11	4	7	103	2.94	2	48	18	47	43	36	577	3.01	6
	%	20.00	17.14	31.43	11.43	20.00				25.00	9.38	24.48	22.40	18.75			
Taylor and Francis	N	10	6	6	4	9	101	2.80	3	20	31	31	14	36	651	3.39	2
	%	28.57	17.14	17.14	11.43	25.71				10.42	16.15	16.15	38.54	18.75			

Table depicts the usage of E-Journals from E-ShodhSindhu by the Faculty and Research Scholars of Acharya Nagarjuna University. Based on the weighted arithmetic mean the purposes are ranked.

### **Major Findings**

It is evident from the table that all the faculty members have completely preferred and gave first rank for JSTOR followed by Springer Link (2<sup>nd</sup> rank), Taylor and Francis (3<sup>rd</sup> rank), American Chemical Society (4<sup>th</sup> rank), Economic and Political Weekly (5<sup>th</sup> rank) and Annual Reviews (6<sup>th</sup> rank). That means majority of the faculty preferred the usage of E-Journal 'JSTOR'.

The table analyses that all Research scholars have completely preferred and gave first rank to Economic and Political Weekly followed by Taylor and Francis (2<sup>nd</sup> rank), Annual Reviews (3<sup>rd</sup> rank), JSTOR (4<sup>th</sup> rank), American Chemical Society (5<sup>th</sup> rank) and Springer Link (6<sup>th</sup> rank). This means that majority of the research scholars from Acharya Nagarjuna University preferred the usage of E-Journal 'Economic and Political Weekly'.

### **References**

1. Allen B M., & Hirshon A. (1998). Hanging together to avoid hanging separately: opportunities for academic libraries and consortia. *Information Technology and Libraries*, 17(1), 36-44.
2. Anju, Saini. (2017). Library Consortia: An Overview. *International Journal of Digital Library Services*, 7(4), 119-123.
3. Arjun, Dinesh Kumar, Apurba Jyot Majumder, & Sharmila Bose (2010). Role of Library in Higher Education in India. *International Journal of Librarianship and Administration*, 1(1), 1-12.
4. Beth Ashmore., & Jill E. Grogg. (2009). The Art of the deal: The Power and pitfalls of Consortia Negotiation. *Searcher*, 17(3), 40-47.
5. Consortium (2013). In Merriam Webster Online Dictionary. Retrieved August 28 2013 from merriam - webster online. <http://www.merriam - webster.com/dictionary/consortium>
6. E-ShodhSindhu: Consortium for Higher Education Electronics ([www.inflibnet.ac.in](http://www.inflibnet.ac.in)).

## Recent Updates on the Bioactive Compounds of Ginger (*Zingiber officinale*) on Cancer: A Study with Special Emphasis of Gingerol and Its Anticancer Potential

### Effect of Ginger and Its Compounds in Cancer Subjects

Handbook of Oxidative Stress in Cancer: Therapeutic Aspects pp 1-18 | Cite as

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## Abstract

Medicinal plants have been used as therapeutic agents since the origin of mankind. Many medicinal plants like *Tinospora cordifolia*, *Andrographis paniculata*, *Curcuma longa*, *Withania somnifera*, *Zingiber officinale*, etc. are used to treat cancer. Ginger is reported to show anticancer effect in many cancer types like liver cancer, gastric cancer, oral cancer, prostate cancer, breast cancer, and ovarian cancers in animal models and cell lines. To date, over 400 bioactive compounds have been identified in ginger, they are gingerols, shogaols, and paradols. These compounds possess antioxidant, anti-inflammatory, antimicrobial, and anticancer properties. Gingerol especially shows anticancer effects in different cancer subjects. Gingerol

may act on the TNF- $\alpha$ , IL-6, NF- $\kappa$ B, cyclooxygenase-2 (COX-2), and caspase-3, and other tumor-metabolic pathway factors in the prevention of cancer. We hope that this chapter will attract more attention on ginger's therapeutic potential and impact on cancer subjects.

## Keywords

Ginger Bioactive compounds Gingerol Cancer

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## Introduction

The biological term cancer refers to a set of diseases in which the cells of an organ or tissue split uncontrollably and acquire the ability to attack other tissues. Cancer is a global health problem, and it is the top cause of deaths in the world. Cancer occurs through the mutations by stepwise process which results in malignancies. Metastasis is the process whereby cancer cells rupture from a malignant tumor and travel to and invade other tissues in the body (Butt and Sultan [2009](#)).

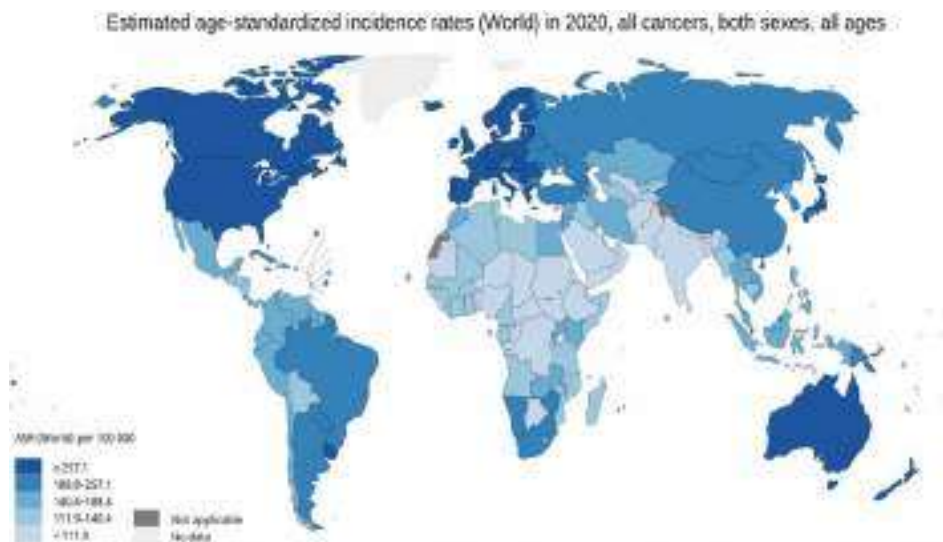
The most common types of cancer are liver cancer, esophageal cancer, breast cancer, oral cancer, lung cancer, prostate cancer, ovarian cancer, and stomach cancer. According to GLOBOCAN 2020 database in 2020, there were 19.1 million new cases of cancer and 10 million deaths from cancer worldwide, and the new cases will be upto 20 million globally by 2025 (Ferlay et al. [2019](#)). The principle malignant conditions of the cancer are breast cancer (2.26 million cases), lung cancer (2.20 million cases), stomach cancer (1.08 million cases), liver cancer (0.90 million cases), esophagus cancer (0.60 million cases), pancreatic cancer (0.49 million cases), and colorectum cancer (0.73 million cases) in 2020 (Table [1](#)).

### Table 1

New cancer cases and deaths as per global cancer statistics 2020

<b>S. no</b>	<b>Cancer type</b>	<b>No of cases (2020)</b>	<b>No of deaths</b>
1.	Breast cancer	2,261,419	684,996
2.	Lung cancer	2,206,771	1,796,144
3.	Prostate cancer	1,414,259	375,304
4.	Skin cancer	1,198,073	63,731
5.	Colon cancer	1,148,515	576,858
6.	Liver cancer	905,677	830,180
7.	Cervical cancer	604,127	341,831
8.	Esophageal cancer	604,100	544,076
9.	Thyroid cancer	586,202	43,646
10.	Pancreatic cancer	495,773	466,003
11.	Leukemia	474,519	311,594
12.	Kidney	431,288	179,368
13.	Oral cancer	377,713	177,757

India has 1.32 lakh new cancer cases in 2020, also the lowest rates of cancer in the world. Breast cancer, oral cancer, cervical cancer, and lung cancer are the top four cancers in India. Cancer deaths in India doubled from 1990 to 2016. It has been reported that genetic mutations, viruses, smoking, heavy metal ingestion, and dietary patterns are actively involved in cancer pathogenesis (Noonan et al. 2007). Despite huge progress and efforts have been made in the field of medicine for the prevention and treatment of cancer for years, cancer still remains the leading cause of deaths around the world (Figure 1).



**Figure 1**

Cancer patients in the world as per the Global Cancer report 2020 by WHO. Source: <https://www.uicc.org/news/globocan-2020-new-global-cancer-data#> (<https://www.uicc.org/news/globocan-2020-new-global-cancer-data#>)

## Carcinogenesis

Carcinogenesis is the process by which a normal cell is transformed into a tumor cell and its progression to a clinically observable tumor with high probability of metastasis. Cancer is considered a process with three steps: 1. initiation, 2. promotion, and 3. progression (Weston and Harris [2003](#)). This process can vary depending on the etiology of cancer. Normally, cancers are caused by chemical agents, viruses, and others by mutations of DNA, epigenetic changes of DNA (Baylin and Jones [2016](#)).

## Mechanism of Cancer

### Cancer: Reactive Oxygen Species (ROS)

Cancer is one of the stress-related disorders. During cancer condition, many free radicals or reactive oxygen species (ROS) are produced. The existence of oxidative stress resulting from increased free radicals has been postulated in cancer. Animal models and human studies and *in vitro* experiments suggest a role for oxidative stress, via an increased formation of free radicals in the pathophysiology of many complications, such as neurological, cardiovascular, renal, cancer, rheumatoid arthritis, cancer, and diabetes (Brownlee [2001](#)). According to Sies (Sies et al. [2017](#)), oxidative stress is defined as a shift in balance in cellular oxidation-reduction reactions in favor of oxidation, which leads to damage to the cell and formation of molecular products that are indicators of oxidative stress.

Reactive oxygen species are radicals or ions or molecules that have a one unpaired electron in their outermost shell of electrons. Due to this nature, ROS are highly reactive. Examples of ROS are superoxide ( $O_2^-$ ), hydrogen peroxide ( $H_2O_2$ ), hydroxyl radicals (OH), and nitric oxide ( $\bullet NO$ ). ROS and oxidative stress have been implicated in many diseases like cancer, diabetes, and hepatitis. ROS plays an important role in various cascading of the signals for the cancer cells to survival, proliferation, resistance to apoptosis, neovascularization, invasion, and extravacation. (Cullen et al. [2003](#)). In order to survive from oxidative stress condition, cancer cells adapt and acquire many mechanisms to counteract the potential toxic effects

of ROS stress in order to promote proximal protumorigenic signals. ROS also alters the DNA-binding sites of redox-sensitive transcription factors such as hypoxia-inducible factor-1 alpha (HIF-1 $\alpha$ ), NF $\kappa$ B, activator protein-1 (AP-1), and p53 (Trachootham et al. 2008). The antioxidant enzymes like SOD, CAT, GPx, GR, and GSH may act on these ROS, and hence these antioxidant enzymes activities are depleted in various cancer subjects.

Warburg (Liberti and Locasale 2016) described about cancer as metabolic alterations that represent a hallmark of cancer cells. Metabolic alterations and redox alterations which are important steps of cancer cell transformation make the mitochondria an attractive therapeutic target. Increased knowledge in the field of redox biology, its reactions, signaling networks, and interplay in disease and physiology has enabled not only a better realization of potential benefits but also grave dangers of ROS with respect to cancer phenotypes and drug resistance.

### Cancer Biomarkers

A thorough understanding of the roles of cancer biomarkers is essential for diagnostic purposes. A biomarker is “a characteristic that is objectively measured and evaluated as an indicator of normal biological processes, pathogenic processes, or pharmacologic responses to a therapeutic intervention.” Because early diagnosis of cancer will treat cancer patients early, we can save the patient’s life. Cancer biomarkers are produced by tumors or by the body in response to the presence of cancer. Cancer biomarkers can be used to determine the presence of malignancy and for the study of disease transmission. The important cancer biomarkers are Urokinase plasminogen activator (uPA), Carbonic anhydrase XII (CAXII), Cyclooxygenase (COX), Cytochrome P450 (CYP450), Telomerase, and Matrix metalloproteinases (MMPs).

The medical treatments for cancer are surgery, radiotherapy, and chemotherapy. These types of treatment are usually accompanied by a large number of side effects on patient health, like nausea, loss of appetite, weight loss, anemia, spinal cord injury, kidney damage, and mucositis. Hence, there is need for alternative therapy for cancer. So herbal medicine is practiced to treat cancer.

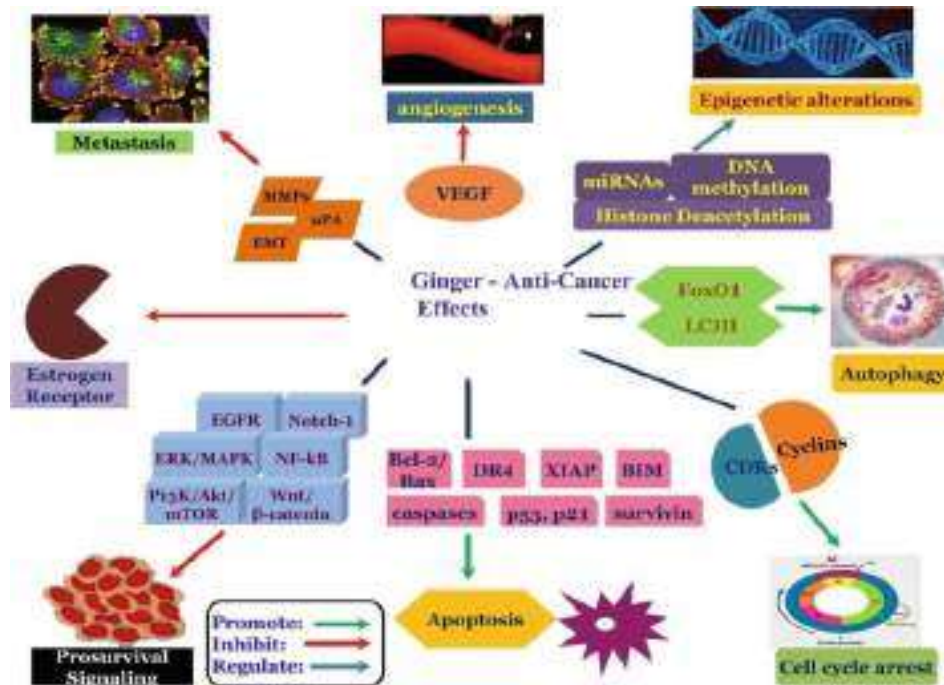
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## Medicinal Plants for the Treatment of Cancer

Medicinal plants are used to treat many diseases like diabetes, cancer, epilepsy, alzheimer’s, cough, fever, and other diseases. Drugs from medicinal plants focus on the most important active principles in terms of the quantity and the pharmacological actions. The information available on these medicinal plants has allowed the quantification of the active principle and the production of bioactive compounds. The bioactive compounds of medicinal plants have different chemical, pharmacological properties and actions. This has enabled us to evaluate the anticancer drugs from medicinal plants.

Many medicinal plants have been reported to possess anticancer properties. Due to the development of resistance against cancer effects, various plant-derived drugs have gained much attention in the recent years. Hence, it is of importance to focus on medicinal plants with anticancer properties that are easily available, culturally acceptable, and economically free. Herbal medicine has been used for thousands of years to cure cancer. Chemotherapeutic and chemopreventive effects of many herbal plants are attributed to phytochemicals, quinines, which are reported to induce antitumor effects *in vivo* and *in vitro* of cancer cells.

Recently, many well-known plant-derived compounds have been studied in animal models and cell lines. Among the above described plants and their formulations, *Terminalia chebula*, *Panax ginseng*, *Arachis hypogaea*, *Rauwolfia vomitoria*, *Azadirachta indica*, *Zingiber officinale*, *Commiphora mukul*, and *Rosa rugosa* are reported to be beneficial for cancer with less side effects as compared to conventional drugs. Hence, these medicinal plants could be a relatively safer and better therapeutic alternative for cancer treatment (Figure 2).



**Figure 2**  
Ginger and anticancer mechanism

## Ginger

Ginger is the rhizome of *Zingiber officinale*. Roscoe belongs to the family Zingiberaceae. Ginger has been used as a spice, food, supplement, and flavoring agent. Ginger is used to cure diseases and symptoms, such as headache, nausea, cold, rheumatism, diarrhea, and arthritis. It is also used as a carminative, digestant, and antifatulent (Ali et al. 2008). Ginger and its compounds are used as medicine in India, China, Burma, Germany, Japan, Indonesia, and the United States.

Ginger is reported to possess various medicinal properties like antibacterial, antiviral, antifungal, antiparasitic, anti-insecticidal, antianalgesic, antimutagen, anticarcinogenic, antispasmodic, and anti-inflammatory and antioxidant activities (Shanmugam et al. 2021). Ginger have radioprotective, hepatoprotective, gastroprotective, nephroprotective, and neuroprotective properties, and its mechanism of action at the cellular level has been studied by many scientists (Ali et al. 2008; Shanmugam et al. 2009, 2021) (Figure 2).

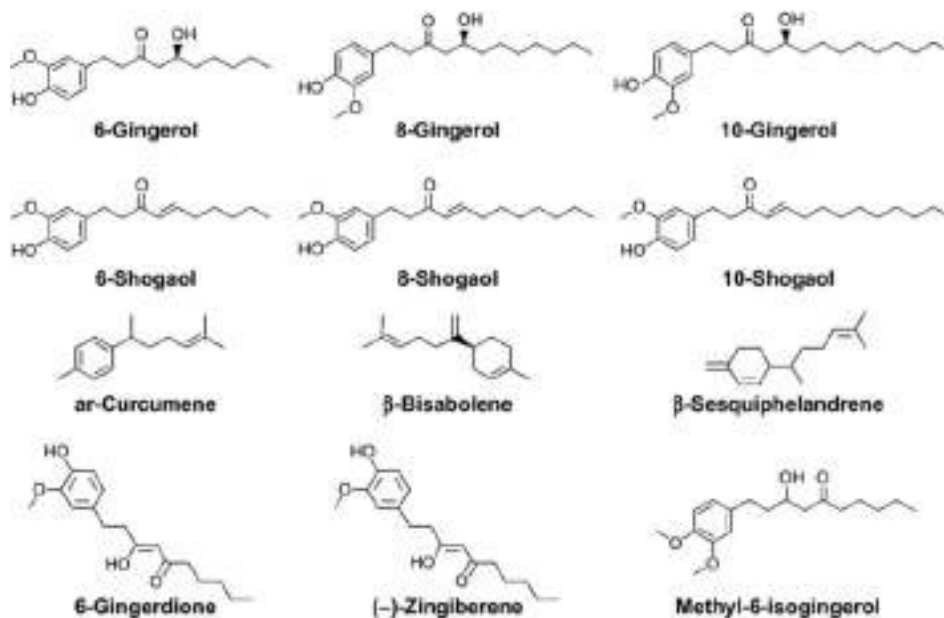


**Figure 3**  
Pharmacological and anticancer properties of Gingerol

#### Compounds of Ginger

The main constituents in ginger are terpenes, phenolic compounds, carbohydrates (50–70%), and lipids (3–8%). The terpenes (monoterpenes, sesquiterpenes, and sesquiterpene alcohols) are of 20–25%. The terpene compounds of ginger are zingiberene,  $\alpha$ -farnesene,  $\beta$ -sesquiphellandrene,  $\beta$ -bisabolene, and  $\alpha$ -curcumene. It has been identified that ginger has monoterpenes (such as  $\alpha$ -pinene, camphene, myrcene,  $\alpha$ -phellandrene, geranial, citronellal, neral, linalool, borneol, and alpha-terpineol). Phytosterols, amino acids, minerals, proteins, vitamins (vitamin A and nicotinic acid), and raw fiber, ash, are present in ginger (Shukla and Singh [2007](#)).

The phenolic compounds of ginger are gingerol, shogaol, and paradols. In ginger, gingerols are in higher concentration. Ginger-specific smell and odor are due to the presence of gingerol and shogaols. The other compounds of ginger are 6-paradol, 1-dehydrogingerdione, 8-gingerdiol, 10-gingerdiol, 4-gingerdiol, 6-gingerdiol, 6-gingerdione, and 10-gingerdione. Ginger also contains diterpenes, galanolactone, diterpenoid, and ginger glycolipids (Yeh et al. [2014](#)). Ginger has ascorbic acid, alkaloids, beta-carotene, and polyphenols. Ginger also has key volatile oils such as oleoresins, bisabolene, cineol, phellandrene, citral, borneol, and citronellol, vitamin B6, vitamin c, and linoleic acid. As per the available information, there are more than 400 bioactive compounds in ginger (Shukla and Singh [2007](#)) (Figure 4).



**Figure 4**  
Compounds of Ginger

#### Anticancer Effects of Ginger

Ginger has anticancer properties. In the methanolic, ethanolic, aqueous, n-hexane, and ethyl acetate, benzene extracts of ginger, many bioactive compounds are present which show antiproliferative, cytotoxic, and antiangiogenic activities. These pharmacological effects are due to bioactive compounds of ginger, which show antioxidant, anti-inflammatory, antioxidant, antihyperglycemic, antitumorigenic, and antilipidemic activities. These compounds may modulate the genetic expression and cause induction of apoptosis in cancer cells.

The anticancer effect of ginger may be due to the reduction of initiation, promotion, and progression. Hence, ginger can be used as anticancer agent (Zhang et al. 2017). Ginger has been reported to show its positive effects against many cancers, like liver cancer, gastric cancer, pancreatic cancer, breast cancer, prostate cancer, and other types of cancers. Ginger also has anticancer effects in different cancer cell lines in vitro, like lung, ovarian, liver, colon, cervical, and prostate cancer, through the stimulation of apoptosis and the reticence of cell proliferation (Zhang et al. 2017) (Table 2).

**Table 2**  
Effect of ginger and its compounds on different cancer subjects

<b>S. no.</b>	<b>Ginger extract/compounds of ginger</b>	<b>Study type</b>	<b>Subjects</b>	<b>Dose</b>	<b>Potential mechanisms</b>	<b>Referen</b>
1	Ginger ethanolic extract	<i>In vivo</i>	Wistar rats	50 mg/kg	NADH dehydrogenase activity elevated	Ali et al. ( <a href="#">2008</a> )
2	Ginger extract	<i>In vitro</i>	HepG2 human hepato-cellular carcinoma cells	1.11 mg/mL	Increasing the level of ROS levels increased and elevated p53 levels, thus promoting apoptosis	Li et al. ( <a href="#">2013</a> )
3	6-gingerol	<i>In vitro</i>	He La human adenocarcinoma cervical cells	60, 100, and 140µM	Cyclin A, cyclinD1, and cyclin E1 levels decreased, and caspase levels increased	Zhang et al. ( <a href="#">2017</a> )
4	10-gingerol	<i>In vitro</i>	Human and mouse breast carcinoma cells	50, 100, and 200µM	Cell growth and cell division are reduced	Bernard et al. ( <a href="#">2017</a> )
5	6-gingerol,10-gingerol,6-shogaol, and 10-shogaol	<i>In vitro</i>	PC-3 human prostate cancer cells	1,10, and 100µM	Inhibiting prostate cancer decreased, and the expression of MRP1 and GSTπ are lowered	Liu et al. ( <a href="#">2017</a> )

S. no.	Ginger extract/compounds of ginger	Study type	Subjects	Dose	Potential mechanisms	Referen
6	6-shogaol	<i>In vitro</i>	LNCaP, DU145, and PC-3 human prostate cancer cells	10, 20, and 40μM	Cyclin D1, surviving, c-Myc, and Bcl2 expression are decreased Inhibition of STAT3 and NF-κB signals	Saha et al. ( <a href="#">2014</a> )

Ginger has an antitumor activity by modulating of genetic pathways. It helps for the activation of suppressing gene of the tumor. Furthermore, ginger can inhibit the vascular endothelial growth factors and modulate apoptosis; thus, ginger supplementation can reduce cancer. Ginger extract prevents the initial stage of colon cancer. Supplementation of ginger to the mice with carcinogen 1,2-dimethylhydrazine (DMH) inhibited the levels of tissue cholesterol, HMG CoA reductase, free fatty acids, and triglycerides (Manju et al. [2006](#)). Hence, ginger treatment reduces the risk of cancer by antioxidative compounds.

For instance, it has been identified that the terpenoids, compound of ginger, induce apoptosis in uterus cancer cells via the activation of tumor protein p53. Water and organic solvent extracts of ginger reported to show anticancer activities in THP-1 AMoL cells in vitro. These reports show that ginger has anticancer activity (Prasad and Tyagi [2015](#)).

Ginger constituents ([6]-gingerol, [10]-gingerol, [6]-shogaol, and zerumbone) show anticancer effects. They are effective against many cancers (Habib et al. [2008](#)). Habib et al. ([2008](#)) reported that ginger extract inhibits liver carcinogenesis in wistar rat through the downregulation of elevated NF-κB and TNF-α. Hence, ginger may act as an anticancer agent, which may be helpful in treatment of cancer subjects. Ginger ingredients inhibit the development of diethylnitrosamine-(DEN-) induced premalignant phenotype in rat hepato-carcinogenesis. It has been reported that supplementation of ginger prevented decrease of the content of metallothionein and endostatin in the liver and elevated the growth factors induced by the carcinogen in wistar albino rats. Ginger also reverses the altered serum-hepatic tumor markers (Mansour et al. [2010](#)).

Wang et al. ([2008](#)) reported that beta-elemene compound of ginger induces caspase-3, -7, -9 activities, decreases Bcl-2 expression, which releases cytochrome c, and elevated the levels of cleaved caspase-9 and poly (ADP-ribose) polymerase in cells. Application of ginger extract to mouse skin afforded significant inhibition of TPA-caused epidermal edema (56%) and hyperplasia (44%) (Katiyar et al. [1996](#)). Kim et al. ([2005](#)) also reported that ginger induces programmed cell death in cell lines. Ethanol extract of ginger has antitumor-promoting effects in mouse skin tumorigenesis model, and it was concluded that animals pretreated with ginger showed depleted tumor compared with unginger-treated rats.

Zingerone with a dose of 100 mg mL suppressed LPS-induced NF-κB activities in cells. Dietary zingerone reduces proinflammatory cytokines. This study reports zingerone anti-inflammatory activity due to suppressing the activation of NF-κB, production of IL-1b, and the infiltration of inflammatory cells (Ganaie et al. [2019](#)). Ganaie et al. ([2019](#)) also reported the anticancer effect of zingerone due to activation of cytochrome P4502E1 and suppression of NF-κB-p65, iNOSCOX-2, and PCNA in cancer subjects.

Zerumbone, seen in ginger, shows antiproliferative and anti-inflammatory effects and mediates its activity through the modulation of NF-κB activation (Takada et al. [2005](#)). Zerumbone also inhibits the activation of NF-κB, and this inhibition may provide basis for the prevention and treatment of cancer. The antihpato-carcinogenic effect of zerumbone was due to suppression of PCNA and elevation of Bax and depletion of Bcl-2 protein expression (Gross et al. [1998](#)). Hence, zerumbone may act as anticancer agent in liver cancers

(Fuzer et al. 2017). Fuzere et al. (2017) reported that zerumbone induces the phase II detoxification enzymes depletion in rat liver epithelial cell line RL34. Hence, Zerumbone may act as a potential activator for the Nrf2-dependent detoxification pathway and provides a new insight into cancer management (Tsuboi et al. 2014) (Figure 5).



**Figure 5**

Suppression of tumors markers by ginger and its compounds

#### Ginger-Bioactive Compound: Gingerol and Its Anticancer Effects

Gingerols are part of the phenolic compounds and volatile organic compounds of ginger. 4-, 6, 8-, 10-, and 12-gingerol are types of gingerol. 6-gingerol is the main compound which is responsible for the strong aroma of ginger. The biological properties of gingerols are antimicrobial, anticancer, antioxidant, anti-inflammatory, and antiallergic (Akinyemi et al. 2015).

6-gingerol has been identified to show anticancerous effects. Gingerol helps in the suppression of the hyperproliferation, inflammatory processes, and transformation that engaged in various steps of angiogenesis and metastasis. For instance, through the activation of CD8+ T cells, it inhibited B16F10 melanoma cells of pulmonary metastasis in mice. Antitumoral activity showed by 6-gingerol through induction of reactive oxygen species (ROS) which, trigger p53 activation, apoptosis, and arrest the cell cycle (Lee et al. 2008). The study found that 6-G induced arrest of the cell cycle in both cell lines by reducing the expression of cyclin A and CDKA. In addition, Radhakrishnan et al. (2014) reported that 6-G induced cell death by apoptosis in the cell line with the mutated p53 gene. [6]-gingerol stimulated death receptor-mediated apoptosis in glioblastoma cells or p53-mediated apoptosis in skin tumor cells (Nakamura et al. 2004).

6-gingerol demonstrates an important potential to treat pancreatic cancer and has been shown to have a potential inhibitor of metastasis in *in vitro* and *in vivo* studies by different mechanisms, including a reduction in the expression of MMP and inhibiting angiogenesis (Kim et al. 2005). 6-gingerol targets many cellular molecules which promote cancer, for cell survival, cell proliferation, invasion, and angiogenesis. 6-Gingerol supplementation alters STAT3, NF- $\kappa$ B, Rb, MAPK, Akt, ERK, PI3K, and caspase-3/7, cIAP1. Thus, gingerol treatment may modulate molecular targets of cancer components, so gingerol may have the therapeutic potential for preventing and treating many cancers.

*In vitro* and *in vivo* analysis of [10]-gingerol has been reported against the metastatic triple negative breast cancer (TNBC) [97]. In addition, it was experimented that 10-gingerol inhibits cervical cancer (Zhang et al. 2017). A study carried out by Martin et al. (2017) showed the anticancer activity of 10-G *in vivo* and *in vitro* in triple negative breast cancer models through proapoptotic activity and the inhibition of metastasis. The activity of 10-G *in vitro* was dose dependent, finding that at the highest concentration (100 $\mu$ M) colony formation was completely inhibited and extensive cell death occurred. 10-gingerol caused a considerable upsurge in the initiation of caspase-3 and inhibited orthotopic tumor growth of spontaneous breast cancer metastasis. Zhang et al. (2017) reported that 10-gingerol inhibits metastasis to multiple organs like lung, bone, and brain.

### Therapeutical Potential of Gingerol and Its Effect on Metabolic Pathways in Cancer

Anticancer mechanisms of gingerol include free radical scavenging effect, antioxidant effect, modulation of various enzymes of inflammation, modulation of cell cycle proteins, induction of apoptosis, and arrest the cell cycle at G2/M phase in carcinoma which will provide basis for inhibition of tumor progression in experimental animals. TNF- $\alpha$ , IL-6, NF- $\kappa$ B, cyclooxygenase-2 COX-2, and caspase-3 are the most important cancer metabolic factors.

#### Free Radical Scavenging and Antioxidant Activity of Gingerol

Gingerol possesses antioxidant activity due to their free radical scavenging activity. It has been investigated that gingerol may modulate the antioxidant enzymes and suppress the lipid peroxidation products in cancer subjects. Hence, gingerol may prevent the pathogenesis of cancer-related disorders by its free radical scavenging activity. Alsahli et al. (2021) reported anticancer activity of gingerol in colon cancer, breast cancer, by elevating antioxidant enzymes and depleting lipid peroxidation in cancer condition.

#### Effect of Gingerol on Apoptotic Genes

Apoptosis is one of the prerequisites to maintain the normal and healthy internal milieu. Alteration in the normal process of apoptosis may raise cell survival and support the tumor growth and progression (Kim et al. 2005). Gingerol plays a vital role in the elevation of different proapoptotic genes and at the same time depletion of the antiapoptotic genes and by this way balances the apoptosis process. An interesting study showed that gingerol induces apoptosis in scleroderma lung fibroblasts without affecting normal lung fibroblasts. Furthermore, gingerol has shown an antitumor activity and was involved in the apoptosis induction and the modulation of key apoptotic proteins such as Bax and bcl-2 (Yu et al. 2011).

A study has reported that growth arrest and apoptosis of B cell lymphoma occur through the downregulation of c-myc, bcl-XL, and p53 with the treatment of gingerol. Another report in human breast cancer cell line showed that CD437 induces G0-G1 arrest and apoptosis via regulation of p21WAF1/CIPI, Bcl-2, and Bax in a p53-independent manner after treatment with gingerol. Another study on p53-null cells, as well as TR9-7 cells, reported that gingerol induces apoptosis in tumor cells via a p53-dependent pathway, and Bax acts as downstream effectors of p53. Gingerol induces apoptosis in a range of tumor cell lines through activation of caspase-3, cytochrome c release, and depletion of bcl-2 (de Lima et al. 2018).

Gingerol has shown an apoptotic effect by inhibiting various genes such as proteintyrosine kinase, protein kinase C, c-myc mRNA expression, and bcl-2 mRNA expression and also mitochondrial pathway. Earlier studies have shown that gingerol possesses an apoptotic activity in different types of cancer cells such as human colon cancer cells, stomach, and skin tumors, breast cancer cells, and prostate cancer cells (Nakamura et al. 2004). Gingerol may lower the incidence of various cancers and also induce apoptosis in MBT-2 cells and G2/M arrest of T24 cells (Park et al. 2014). Experimental studies showed that the downregulation of the expression of antiapoptotic protein occurs with gingerol treatment (Nakamura et al. 2004).

#### Effect of Gingerol on Tumor-Suppressor Genes

Tumor-suppressor genes play a vital and significant role in the inhibition of cancer formation and its progression. An alteration or mutations may occur in a gene, then tumor suppressor gene loses its ability to perform normal function and it transforms into tumor gene. p53 is one of the important suppressor genes, and it is the guardian of all genes and regulates the various cellular and molecular pathways and prevents the formation of cancer.

Numerous *in vivo* and *in vitro* reports showed that gingerol has a significant role in cancer prevention or inhibition. Another study showed that gingerol downregulates the expression of p53, as well as the survival genes *egr-1*, *c-myc* and *bcl-XL* in B cells. Another report also indicated that gingerol inhibits cell cycle progression of immortalized human umbilical vein-endothelial cells via upregulating the CDK inhibitors p21<sup>WAF1/CIP1</sup>, p27<sup>KIP1</sup>, and p53 (Park et al. [2014](#)).

Another tumor-suppressor gene, phosphatase and tensin homolog deleted on chromosome ten (PTEN), has a role in the progression of the cell cycle and apoptosis. The alteration or mutation of PTEN gene has been noticed in several types of cancers. A study of the gingerol has shown that PTEN increases the gingerol-induced apoptosis, whereas inactive PTEN decreases/inhibits the gingerol-induced apoptosis.

In mice, [6]-gingerol suppressed the promotion of skin cancer. Park et al. (Park et al. [2014](#)) reported that [6]-gingerol inhibited TPA skin tumor promotion in addition to the inhibition of epidermal ornithine decarboxylase activity in ICR mice. In a study, Surh et al. [[1999](#)] reported antitumor-promoting properties of both [6]-gingerol and [6]-paradol. [6]-gingerol treatment attenuated the skin papilloma genesis and inhibited the tumor-promoter genes, TNF-alpha production, and activation of epidermal ornithine decarboxylase in ICR mice (Nakamura et al. [2004](#)).

### **Effect of Gingerol on Cyclooxygenase Enzyme**

COX is an inducible enzyme in the conversion of arachidonic acid to prostaglandins (PGs). There are two types of cyclooxygenase COX-1 that play a vital role in physiological functions and COX-2, an enzyme responsible for inflammation and pain. COX-2 is upregulated or overexpressed in various types of cancers (Pournaderi et al. [2017](#)). It was previously stated that gingerol inhibits the critical stage of tumor initiation and promotion stages and COX inhibition. Gingerol also inhibits the COX2 expression on colon cancer cell lines (Nonn et al. [2007](#)). Previous reports states that gingerol plays an important role in the downregulation of the expression of COX-2 and finally suppresses the cancer progression (Kim et al. [2005](#)). Gingerol plays a significant role in the cancer prevention via controlling the activities of various genes in the initiation, promotion, and progression stage of tumor development and progression.

### **NF- $\kappa$ B and Gingerol in Cancer Prevention**

NF- $\kappa$ B family of transcription factors shows an important role in immune, inflammatory response and also stimulates the development and progression of cancer. In this regard, an important study demonstrated that gingerols have anticancer, antioxidant, and anti-inflammatory effects via the downregulation of the transcription factors NF- $\kappa$ B, AP-1, and Egr-1 (Han et al. [2002](#)) and repression of the genes for cell adhesion molecules (chemokines, TNF, Cox-2, and MMP-9). Another study showed that gingerol has been involved in the suppression of NF- $\kappa$ B activation and NF- $\kappa$ B gene products (Plummer et al. [1999](#)). An important study in pancreatic cancer cells reported that gingerol shows a vital role in the suppression of NF- $\kappa$ B activation by inhibiting I $\kappa$ B kinase, ultimately induces I $\kappa$ B $\alpha$  phosphorylation, and inhibits the NF- $\kappa$ B downstream gene expression. [6]-gingerol inhibited both the vascular endothelial growth factor (VEGF) and basic fibroblast growth factor(b-FGF)-induced proliferation of human endothelial cells and caused cell cycle arrest in the G<sub>1</sub> phase (Vijaya Padma et al. [2007](#)). Recently, anticancer and anti-inflammatory potential of 6-gingerol was reported by inactivating NF- $\kappa$ B through the suppression of the proinflammatory TNF-alpha (Kim et al. [2005](#); Habib et al. [2008](#)). Numerous studies have shown that gingerol is a potent inhibitor of NF- $\kappa$ B activation.

Kim et al. ([2005](#)) reported that 6-gingerol regulates tight junction-related proteins and suppresses invasion and metastasis of pancreatic cancer cells. 6-gingerol mediated through NF- $\kappa$ B inhibition via inhibition of the extracellular signal-regulated kinases (ERK) pathway. Thus, 6-gingerol suppresses the invasive activity of PANC-1 cells.

### **Impact of Gingerol on Angiogenesis**

Angiogenesis is a complex process involving widespread interaction between the cells, soluble factors, and ECM components. During cancer development, tumor growth is triggered by many signals in chain reaction manner. In cancer formation, there are stages and angiogenic factors such as vascular endothelial growth

factor (VEGF), basic fibroblast growth factor (bFGF), angiogenin, transforming growth factor (TGF- $\alpha$ , TGF- $\beta$ ), and epidermal growth factors, which play vital role in tumor angiogenesis through cancerous tumor cells by releasing molecules and sending signals to surrounding normal host tissue. VEGF is a crucial survival factor for endothelial cells in the process of physiological, tumor angiogenesis, and it induces the expression of antiapoptotic proteins in the endothelial cells (Shukla and Singh 2007). Many reports showed that many medicinal plants suppress the VEGF and other factors of cancer. We also reported that gingerol is best inhibitor of VEGF in different types of cancer (Weng et al. 2010).

*In vitro* studies on hepatic cancer cell lines conducted so far have demonstrated that ginger suppressed the growth of human hepatic cancer cell lines by inhibiting the phosphorylation of tyrosine-kinase receptor IGF-1R, inducing apoptosis by activating Caspase 9 and downregulating Bcl-2 and cyclooxygenase-2 (COX-2), modulating the levels of VEGF and its receptor (VEGFR-2), NF- $\kappa$ B, p53, and extracellular signal-regulated kinase 1/2 (ERK1/2), reducing the expression of lipogenic enzymes, certain types of RTKs, and their downstream pathways, and activating adenosine monophosphate-activated protein kinase (AMPK) and ROS-mediated lysosomal membrane permeabilization. Based on various liver carcinogenesis animal models with the intake of ginger, the inhibition of hepatoma growth, restriction of hepatic cancer cell line progression, and activation of apoptosis were observed; the probable mechanisms behind might be associated with suppression of hepatocyte progenitor cell/stem cell population, activation of AMPK protein in the liver, and modulation of self-renewal pathways and their related genes. *In vitro* and *in vivo* studies reported that gingerol suppresses the proliferation of human vascular endothelial cells and also abrogates the FGF-2-induced angiogenic response. Moreover, gingerol has the ability to inhibit both VEGF and its receptor in various cancer types; it might be useful as an antiangiogenic agent. Hence, gingerol acts as anticancer agent against many cancers through suppressing proliferation, angiogenesis, NF- $\kappa$ B, and NF- $\kappa$ B-regulated gene products.

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## Conclusion and Future Perspectives

Cancer is one of the deadliest diseases and a major health problem in the world. The present modes of treatments like chemotherapy and radiotherapy are very expensive and also exhibit many side effects in cancer patients. Ginger and its bioactive compounds are used to treat cancer. Gingerol a bioactive compound of ginger has been shown to target multiple signaling molecules in cancer metabolism and provides a basis for its therapeutic applications for cancer subjects. Moreover, most of the known activities of gingerol are based only on *in vitro* and *in vivo* studies, and some clinical studies in human subjects. Therefore, more extensive and well-controlled animal and human studies are required to demonstrate efficacy of gingerol and other compounds of ginger against cancer.

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## References

- Akinyemi AJ, Thome R, Morsch VM, Stefanello N, Goularte JF, Bello-Klein A, Oboh G, ChitolinaSchetinger MR (2015) Effect of dietary supplementation of ginger and turmeric rhizomes on angiotensin-1 converting enzyme (ACE) and arginase activities in L-NAME induced hypertensive rats. *J Funct Foods* 17:792–801  
[CrossRef](https://doi.org/10.1016/j.jff.2015.06.011) (https://doi.org/10.1016/j.jff.2015.06.011)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Effect%20of%20dietary%20supplementation%20of%20ginger%20and%20turmeric%20rhizomes%20on%20angiotensin-1%20converting%20enzyme%20%28ACE%29%20and%20arginase%20activities%20in%20L-NAME%20induced%20hypertensive%20rats&author=AJ.%20Akinyemi&author=R.%20Thome&author=VM.%20Morsch&author=N.%20Stefanello&author=JF.%20Goularte&author=A.%20Bello-Klein&author=G.%20Oboh&author=MR.%20ChitolinaSchetinger&journal=J%20Funct%20Foods&volume=17&pages=792-801&publication_year=2015) (http://scholar.google.com/scholar\_lookup?title=Effect%20of%20dietary%20supplementation%20of%20ginger%20and%20turmeric%20rhizomes%20on%20angiotensin-1%20converting%20enzyme%20%28ACE%29%20and%20arginase%20activities%20in%20L-NAME%20induced%20hypertensive%20rats&author=AJ.%20Akinyemi&author=R.%20Thome&author=VM.%20Morsch&author=N.%20Stefanello&author=JF.%20Goularte&author=A.%20Bello-Klein&author=G.%20Oboh&author=MR.%20ChitolinaSchetinger&journal=J%20Funct%20Foods&volume=17&pages=792-801&publication\_year=2015)
- Ali BH, Blunden G, Tanira MO, Nemmar A (2008) Some phytochemical, pharmacological and toxicological properties of ginger (*Zingiber officinale* roscoe): a review of recent research. *Food Chem Toxicol* 46:409–420  
[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=17950516) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=17950516)  
[CrossRef](https://doi.org/10.1016/j.jft.2007.09.085) (https://doi.org/10.1016/j.jft.2007.09.085)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Some%20phytochemical%20%20pharmacological%20and%20toxicological%20properties%20of%20ginger%20%28Zingiber%20officinale%20roscoe%29%3A%20a%20review%20of%20recent%20re) (http://scholar.google.com/scholar\_lookup?title=Some%20phytochemical%20%20pharmacological%20and%20toxicological%20properties%20of%20ginger%20%28Zingiber%20officinale%20roscoe%29%3A%20a%20review%20of%20recent%20re

search&author=BH.%20Ali&author=G.%20Blunden&author=MO.%20Tanira&author=A.%20Nemmar&journal=Food%20Chem%20Toxicol&volume=46&pages=409-420&publication\_year=2008)

Alsahli MA, Almatroodi SA, Almatroudi A, Khan AA, Anwar S, Almutary AG, Alrumaihi F, Rahmani AH (2021) 6-Gingerol, a major ingredient of ginger attenuates Diethylnitrosamine-induced liver injury in rats through the modulation of oxidative stress and anti-inflammatory activity. *Mediators Inflamm* 2021

[Google Scholar](https://scholar.google.com/scholar?) (<https://scholar.google.com/scholar?>

q=Alsahli%20MA%20Almatroodi%20SA%20Almatroudi%20AA%20Khan%20AA%20Anwar%20S%20Almutary%20AG%20Alrumaihi%20F%20Rahmani%20AH%20%282021%29%206-

Gingerol%20a%20major%20ingredient%20of%20ginger%20attenuates%20Diethylnitrosamine-induced%20liver%20injury%20in%20rats%20through%20the%20modulation%20of%20oxidative%20stress%20and%20anti-inflammatory%20activity.%20Mediators%20Inflamm%202021)

Baylin SB, Jones PA (2016) Epigenetic determinants of cancer. *Cold Spring Harb Perspect Biol* 8(9):019505

[CrossRef](https://doi.org/10.1101/cshperspect.a019505) (<https://doi.org/10.1101/cshperspect.a019505>)

[Google Scholar](http://scholar.google.com/scholar_lookup?) ([http://scholar.google.com/scholar\\_lookup?](http://scholar.google.com/scholar_lookup?)

title=Epigenetic%20determinants%20of%20cancer&author=SB.%20Baylin&author=PA.%20Jones&journal=Cold%20Spring%20Harb%20Perspect%20Biol&volume=8&issue=9&publication\_year=2016)

Bernard MM, McConnery JR, Hoskin DW (2017) [10]-Gingerol, a major phenolic constituent of ginger root, induces cell cycle arrest and apoptosis in triple-negative breast cancer cells. *Exp Mol Pathol* 102(2):370–376

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?) (<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?>

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=28315687)

[CrossRef](https://doi.org/10.1016/j.yexmp.2017.03.006) (<https://doi.org/10.1016/j.yexmp.2017.03.006>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=%5B10%5D-) ([Gingerol%20a%20major%20phenolic%20constituent%20of%20ginger%20root%20induces%20cell%20cycle%20arrest%20and%20apoptosis%20in%20triple-negative%20breast%20cancer%20cells&author=MM.%20Bernard&author=JR.%20McConnery&author=DW.%20Hoskin&journal=Exp%20Mol%20Pathol&volume=102&issue=2&pages=370-376&publication\\_year=2017\)](http://scholar.google.com/scholar_lookup?title=%5B10%5D-</a></p>
</div>
<div data-bbox=)

Brownlee M (2001) Biochemistry and molecular cell biology of diabetic complications. *Nature* 414:813–820

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?) (<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?>

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=11742414)

[CrossRef](https://doi.org/10.1038/414813a) (<https://doi.org/10.1038/414813a>)

[Google Scholar](http://scholar.google.com/scholar_lookup?) ([http://scholar.google.com/scholar\\_lookup?](http://scholar.google.com/scholar_lookup?)

title=Biochemistry%20and%20molecular%20cell%20biology%20of%20diabetic%20complications&author=M.%20Brownlee&journal=Nature&volume=414&pages=813-820&publication\_year=2001)

Butt MS, Sultan MT (2009) Green tea: nature's defense against malignancies. *Crit Rev Food Sci Nutr* 49(5):463–473

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?) (<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?>

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=19399671)

[CrossRef](https://doi.org/10.1080/10408390802145310) (<https://doi.org/10.1080/10408390802145310>)

[Google Scholar](http://scholar.google.com/scholar_lookup?) ([http://scholar.google.com/scholar\\_lookup?](http://scholar.google.com/scholar_lookup?)

title=Green%20tea%3A%20nature%27s%20defense%20against%20malignancies&author=MS.%20Butt&author=MT.%20Sultan&journal=Crit%20Rev%20Food%20Sci%20Nutr&volume=49&issue=5&pages=463-473&publication\_year=2009)

Cullen JJ, Weydert C, Hinkhouse MM, Ritchie J, Domann FE, Spitz D, Oberley LW (2003) The role of manganese superoxide dismutase in the growth of pancreatic adenocarcinoma. *Cancer Res* 15(63):1297–1303

[Google Scholar](http://scholar.google.com/scholar_lookup?) ([http://scholar.google.com/scholar\\_lookup?](http://scholar.google.com/scholar_lookup?)

title=The%20role%20of%20manganese%20superoxide%20dismutase%20in%20the%20growth%20of%20pancreatic%20adenocarcinoma&author=JJ.%20Cullen&author=C.%20Weydert&author=MM.%20Hinkhouse&author=J.%20Ritchie&author=FE.%20Domann&author=D.%20Spitz&author=LW.%20Oberley&journal=Cancer%20Res&volume=15&issue=63&pages=1297-1303&publication\_year=2003)

de Lima RMT, Dos Reis AC, de Menezes APM, Santos JVO, Filho JWGO, Ferreira JRO, de Alencar MVOB, da Mata AMOF, Khan IN, Islam A, Uddin SJ, Ali ES, Islam MT, Tripathi S, Mishra SK, Mubarak MS, Melo-Cavalcante AAC (2018) Protective and therapeutic potential of ginger (*Zingiber officinale*) extract and [6]-gingerol in cancer: a comprehensive review. *Phytother Res* 201832(10):1885–1907

[CrossRef](https://doi.org/10.1002/ptr.6134) (<https://doi.org/10.1002/ptr.6134>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Protective%20and%20therapeutic%20potential%20of%20ginger%20%28Zingiber%20officinale%29%20extract%20and%20%5B%5D-gingerol%20in%20cancer%3A%20a%20comprehensive%20review&author=RMT.%20Lima&author=AC.%20Dos%20Reis&author=APM.%20Menezes&author=JVO.%20Santos&author=JWGO.%20Filho&author=JRO.%20Ferreira&author=MVOB.%20Alencar&author=AMOF.%20Mata&author=IN.%20Khan&author=A.%20Islam&author=SJ.%20Uddin&author=ES.%20Ali&author=MT.%20Islam&author=S.%20Tripathi&author=SK.%20Mishra&author=MS.%20Mubarak&author=AAAC.%20Melo-Cavalcante&journal=Phytother%20Res&volume=201832&issue=10&pages=1885-1907&publication_year=2018) ([http://scholar.google.com/scholar\\_lookup?title=Protective%20and%20therapeutic%20potential%20of%20ginger%20%28Zingiber%20officinale%29%20extract%20and%20%5B%5D-gingerol%20in%20cancer%3A%20a%20comprehensive%20review&author=RMT.%20Lima&author=AC.%20Dos%20Reis&author=APM.%20Menezes&author=JVO.%20Santos&author=JWGO.%20Filho&author=JRO.%20Ferreira&author=MVOB.%20Alencar&author=AMOF.%20Mata&author=IN.%20Khan&author=A.%20Islam&author=SJ.%20Uddin&author=ES.%20Ali&author=MT.%20Islam&author=S.%20Tripathi&author=SK.%20Mishra&author=MS.%20Mubarak&author=AAAC.%20Melo-Cavalcante&journal=Phytother%20Res&volume=201832&issue=10&pages=1885-1907&publication\\_year=2018](http://scholar.google.com/scholar_lookup?title=Protective%20and%20therapeutic%20potential%20of%20ginger%20%28Zingiber%20officinale%29%20extract%20and%20%5B%5D-gingerol%20in%20cancer%3A%20a%20comprehensive%20review&author=RMT.%20Lima&author=AC.%20Dos%20Reis&author=APM.%20Menezes&author=JVO.%20Santos&author=JWGO.%20Filho&author=JRO.%20Ferreira&author=MVOB.%20Alencar&author=AMOF.%20Mata&author=IN.%20Khan&author=A.%20Islam&author=SJ.%20Uddin&author=ES.%20Ali&author=MT.%20Islam&author=S.%20Tripathi&author=SK.%20Mishra&author=MS.%20Mubarak&author=AAAC.%20Melo-Cavalcante&journal=Phytother%20Res&volume=201832&issue=10&pages=1885-1907&publication_year=2018))

Ferlay J, Colombet M, Soerjomataram I, Mathers C, Parkin DM, Piñeros M, Znaor A, Bray F (2019) Estimating the global cancer incidence and mortality in 2018: GLOBOCAN sources and methods. *Int J Cancer* 144(8):1941–1953

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=30350310) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=30350310](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=30350310))

[CrossRef](https://doi.org/10.1002/ijc.31937) (<https://doi.org/10.1002/ijc.31937>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Estimating%20the%20global%20cancer%20incidence%20and%20mortality%20in%202018%3A%20GLOBOCAN%20sources%20and%20methods&author=J.%20Ferlay&author=M.%20Colombet&author=I.%20Soerjomataram&author=C.%20Mathers&author=DM.%20Parkin&author=M.%20Pi%C3%B1eros&author=A.%20Znaor&author=F.%20Bray&journal=Int%20J%20Cancer&volume=144&issue=8&pages=1941-1953&publication_year=2019) ([http://scholar.google.com/scholar\\_lookup?title=Estimating%20the%20global%20cancer%20incidence%20and%20mortality%20in%202018%3A%20GLOBOCAN%20sources%20and%20methods&author=J.%20Ferlay&author=M.%20Colombet&author=I.%20Soerjomataram&author=C.%20Mathers&author=DM.%20Parkin&author=M.%20Pi%C3%B1eros&author=A.%20Znaor&author=F.%20Bray&journal=Int%20J%20Cancer&volume=144&issue=8&pages=1941-1953&publication\\_year=2019](http://scholar.google.com/scholar_lookup?title=Estimating%20the%20global%20cancer%20incidence%20and%20mortality%20in%202018%3A%20GLOBOCAN%20sources%20and%20methods&author=J.%20Ferlay&author=M.%20Colombet&author=I.%20Soerjomataram&author=C.%20Mathers&author=DM.%20Parkin&author=M.%20Pi%C3%B1eros&author=A.%20Znaor&author=F.%20Bray&journal=Int%20J%20Cancer&volume=144&issue=8&pages=1941-1953&publication_year=2019))

FuzerAM LSY, Mott JD, Cominetti MR (2017) [10]-Gingerol reverts malignant phenotype of breast cancer cells in 3D culture. *J Cell Biochem* 118(9):2693–2699

[CrossRef](https://doi.org/10.1002/jcb.25906) (<https://doi.org/10.1002/jcb.25906>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=%5B10%5D-Gingerol%20reverts%20malignant%20phenotype%20of%20breast%20cancer%20cells%20in%203D%20culture&author=LSY.%20FuzerAM&author=JD.%20Mott&author=MR.%20Cominetti&journal=J%20Cell%20Biochem&volume=118&issue=9&pages=2693-2699&publication_year=2017) ([http://scholar.google.com/scholar\\_lookup?title=%5B10%5D-Gingerol%20reverts%20malignant%20phenotype%20of%20breast%20cancer%20cells%20in%203D%20culture&author=LSY.%20FuzerAM&author=JD.%20Mott&author=MR.%20Cominetti&journal=J%20Cell%20Biochem&volume=118&issue=9&pages=2693-2699&publication\\_year=2017](http://scholar.google.com/scholar_lookup?title=%5B10%5D-Gingerol%20reverts%20malignant%20phenotype%20of%20breast%20cancer%20cells%20in%203D%20culture&author=LSY.%20FuzerAM&author=JD.%20Mott&author=MR.%20Cominetti&journal=J%20Cell%20Biochem&volume=118&issue=9&pages=2693-2699&publication_year=2017))

Ganaie MA, Al Saedan A, Madhkali H, Jan BL, Khatlani T, Sheikh IA, Rehman MU, Wani K (2019) Chemopreventive efficacy zingerone (4-[4-hydroxy-3-methylphenyl] butan-2-one) in experimental colon carcinogenesis in Wistar rats. *Environ Toxicol* 34(5):610–625

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=30720227) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=30720227](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=30720227))

[CrossRef](https://doi.org/10.1002/tox.22727) (<https://doi.org/10.1002/tox.22727>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Chemopreventive%20efficacy%20zingerone%20%284-%5B4-hydroxy-3-methylphenyl%5D%20butan-2-one%29%20in%20experimental%20colon%20carcinogenesis%20in%20Wistar%20rats&author=MA.%20Ganaie&author=A.%20Saedan&author=H.%20Madhkali&author=BL.%20Jan&author=T.%20Khatlani&author=IA.%20Sheikh&author=MU.%20Rehman&author=K.%20Wani&journal=Environ%20Toxicol&volume=34&issue=5&pages=610-625&publication_year=2019) ([http://scholar.google.com/scholar\\_lookup?title=Chemopreventive%20efficacy%20zingerone%20%284-%5B4-hydroxy-3-methylphenyl%5D%20butan-2-one%29%20in%20experimental%20colon%20carcinogenesis%20in%20Wistar%20rats&author=MA.%20Ganaie&author=A.%20Saedan&author=H.%20Madhkali&author=BL.%20Jan&author=T.%20Khatlani&author=IA.%20Sheikh&author=MU.%20Rehman&author=K.%20Wani&journal=Environ%20Toxicol&volume=34&issue=5&pages=610-625&publication\\_year=2019](http://scholar.google.com/scholar_lookup?title=Chemopreventive%20efficacy%20zingerone%20%284-%5B4-hydroxy-3-methylphenyl%5D%20butan-2-one%29%20in%20experimental%20colon%20carcinogenesis%20in%20Wistar%20rats&author=MA.%20Ganaie&author=A.%20Saedan&author=H.%20Madhkali&author=BL.%20Jan&author=T.%20Khatlani&author=IA.%20Sheikh&author=MU.%20Rehman&author=K.%20Wani&journal=Environ%20Toxicol&volume=34&issue=5&pages=610-625&publication_year=2019))

Gross A, Jockel J, Wei MC, Korsmeyer SJ (1998) Enforced dimerization of BAX results in its translocation, mitochondrial dysfunction and apoptosis. *EMBO J* 17(14):3878–3885

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=9670005) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=9670005](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=9670005))

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC1170723) (<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC1170723>)

[CrossRef](https://doi.org/10.1093/emboj/17.14.3878) (<https://doi.org/10.1093/emboj/17.14.3878>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Enforced%20dimerization%20of%20BAX%20results%20in%20its%20translocation%20C%20mitochondrial%20dysfunction%20and%20apoptosis&author=A.%20Gross&author=J.%20Jockel&author=MC.%20Wei&author=SJ.%20Korsmeyer&journal=EMBO%20J&volume=17&issue=14&pages=3878-3885&publication_year=1998) ([http://scholar.google.com/scholar\\_lookup?title=Enforced%20dimerization%20of%20BAX%20results%20in%20its%20translocation%20C%20mitochondrial%20dysfunction%20and%20apoptosis&author=A.%20Gross&author=J.%20Jockel&author=MC.%20Wei&author=SJ.%20Korsmeyer&journal=EMBO%20J&volume=17&issue=14&pages=3878-3885&publication\\_year=1998](http://scholar.google.com/scholar_lookup?title=Enforced%20dimerization%20of%20BAX%20results%20in%20its%20translocation%20C%20mitochondrial%20dysfunction%20and%20apoptosis&author=A.%20Gross&author=J.%20Jockel&author=MC.%20Wei&author=SJ.%20Korsmeyer&journal=EMBO%20J&volume=17&issue=14&pages=3878-3885&publication_year=1998))

Habib SH, Makpol S, Abdul Hamid NA, Das S, Ngah WZ, Yusof YA (2008) Ginger extract (*Zingiber officinale*) has anti-cancer and anti-inflammatory effects on ethionine-induced hepatoma rats. *Clinics (Sao Paulo)* 63(6):807–813

[CrossRef](https://doi.org/10.1590/S1807-59322008000600017) (<https://doi.org/10.1590/S1807-59322008000600017>)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Ginger%20extract%20%28Zingiber%20officinale%29%20has%20anti-cancer%20and%20anti-inflammatory%20effects%20on%20ethionine-induced%20hepatoma%20rats&author=SH.%20Habib&author=S.%20Makpol&author=NA.%20Abdul%20Hamid&author=S.%20Das&author=WZ.%20Ngah&author=YA.%20Yusof&journal=Clinics%20%28Sao%20Paulo%29&volume=63&issue=6&pages=807-813&publication_year=2008) ([http://scholar.google.com/scholar\\_lookup?title=Ginger%20extract%20%28Zingiber%20officinale%29%20has%20anti-cancer%20and%20anti-inflammatory%20effects%20on%20ethionine-induced%20hepatoma%20rats&author=SH.%20Habib&author=S.%20Makpol&author=NA.%20Abdul%20Hamid&author=S.%20Das&author=WZ.%20Ngah&author=YA.%20Yusof&journal=Clinics%20%28Sao%20Paulo%29&volume=63&issue=6&pages=807-813&publication\\_year=2008](http://scholar.google.com/scholar_lookup?title=Ginger%20extract%20%28Zingiber%20officinale%29%20has%20anti-cancer%20and%20anti-inflammatory%20effects%20on%20ethionine-induced%20hepatoma%20rats&author=SH.%20Habib&author=S.%20Makpol&author=NA.%20Abdul%20Hamid&author=S.%20Das&author=WZ.%20Ngah&author=YA.%20Yusof&journal=Clinics%20%28Sao%20Paulo%29&volume=63&issue=6&pages=807-813&publication_year=2008))

Han SS, Keum YS, Seo HJ, Surh Y-J (2002) Curcumin suppresses activation of NF- $\kappa$ B and AP-1 induced by phorbol ester in cultured human promyelocytic leukemia cells. *J Biochem Mol Biol* 35(3):337–342

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=12297018) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=12297018)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Curcumin%20suppresses%20activation%20of%20NF-%20Fo%9D%9C%85B%20and%20AP-1%20induced%20by%20phorbol%20ester%20in%20cultured%20human%20promyelocytic%20leukemia%20cells&author=SS.%20Han&author=YS.%20Keum&author=HJ.%20Seo&author=Y-J.%20Surh&journal=J%20Biochem%20Mol%20Biol&volume=35&issue=3&pages=337-342&publication_year=2002) (http://scholar.google.com/scholar\_lookup?

title=Curcumin%20suppresses%20activation%20of%20NF-%20Fo%9D%9C%85B%20and%20AP-1%20induced%20by%20phorbol%20ester%20in%20cultured%20human%20promyelocytic%20leukemia%20cells&author=SS.%20Han&author=YS.%20Keum&author=HJ.%20Seo&author=Y-J.%20Surh&journal=J%20Biochem%20Mol%20Biol&volume=35&issue=3&pages=337-342&publication\_year=2002)

Katiyar SK, Agarwa R, Mukhtar H (1996) Inhibition of tumor promotion in SENCAR mouse skin by ethanol extract of *Zingiber officinale* rhizome. *Cancer Res* 56(5):1023–1030

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=8640756) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=8640756)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Inhibition%20of%20tumor%20promotion%20in%20SENCAR%20mouse%20skin%20by%20ethanol%20extract%20of%20Zingiber%20officinale%20rhizome&author=SK.%20Katiyar&author=R.%20Agarwa&author=H.%20Mukhtar&journal=Cancer%20Res&volume=56&issue=5&pages=1023-1030&publication_year=1996) (http://scholar.google.com/scholar\_lookup?

title=Inhibition%20of%20tumor%20promotion%20in%20SENCAR%20mouse%20skin%20by%20ethanol%20extract%20of%20Zingiber%20officinale%20rhizome&author=SK.%20Katiyar&author=R.%20Agarwa&author=H.%20Mukhtar&journal=Cancer%20Res&volume=56&issue=5&pages=1023-1030&publication\_year=1996)

Kim EC, Min JK, Kim TY, Lee SJ, Yang HO, Han S, Kim YM, Kwon YG (2005) [6]-Gingerol, a pungent ingredient of ginger, inhibits angiogenesis *in vitro* and *in vivo*. *Biochem Biophys Res Commun* 335:300–308

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=16081047) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=16081047)

[CrossRef](https://doi.org/10.1016/j.bbrc.2005.07.076) (https://doi.org/10.1016/j.bbrc.2005.07.076)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=%5B6%5D-Gingerol%2C%20a%20pungent%20ingredient%20of%20ginger%2C%20inhibits%20angiogenesis%20in%20vitro%20and%20in%20vivo&author=EC.%20Kim&author=JK.%20Min&author=TY.%20Kim&author=SJ.%20Lee&author=HO.%20Yang&author=S.%20Han&author=YM.%20Kim&author=YG.%20Kwon&journal=Biochem%20Biophys%20Res%20Commun&volume=335&pages=300-308&publication_year=2005) (http://scholar.google.com/scholar\_lookup?title=%5B6%5D-

Gingerol%2C%20a%20pungent%20ingredient%20of%20ginger%2C%20inhibits%20angiogenesis%20in%20vitro%20and%20in%20vivo&author=EC.%20Kim&author=JK.%20Min&author=TY.%20Kim&author=SJ.%20Lee&author=HO.%20Yang&author=S.%20Han&author=YM.%20Kim&author=YG.%20Kwon&journal=Biochem%20Biophys%20Res%20Commun&volume=335&pages=300-308&publication\_year=2005)

Lee SH, Cekanova M, Baek SJ (2008) Multiple mechanisms are involved in 6-gingerol-induced cell growth arrest and apoptosis in human colorectal cancer cells. *Mol Carcinog* 47(3):197–208

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=18058799) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=18058799)

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2430145) (http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2430145)

[CrossRef](https://doi.org/10.1002/mc.20374) (https://doi.org/10.1002/mc.20374)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Multiple%20mechanisms%20are%20involved%20in%206-gingerol-induced%20cell%20growth%20arrest%20and%20apoptosis%20in%20human%20colorectal%20cancer%20cells&author=SH.%20Lee&author=M.%20Cekanova&author=SJ.%20Baek&journal=Mol%20Carcinog&volume=47&issue=3&pages=197-208&publication_year=2008) (http://scholar.google.com/scholar\_lookup?

title=Multiple%20mechanisms%20are%20involved%20in%206-gingerol-induced%20cell%20growth%20arrest%20and%20apoptosis%20in%20human%20colorectal%20cancer%20cells&author=SH.%20Lee&author=M.%20Cekanova&author=SJ.%20Baek&journal=Mol%20Carcinog&volume=47&issue=3&pages=197-208&publication\_year=2008)

Li F, Li S, Li HB, Deng GF, Ling WH, Xu XR (2013) Antiproliferative activities of tea and herbal infusions. *Food Funct* 4(4):530–538

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=23307138) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=23307138)

[CrossRef](https://doi.org/10.1039/c2fo30252g) (https://doi.org/10.1039/c2fo30252g)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Antiproliferative%20activities%20of%20tea%20and%20herbal%20infusions&author=F.%20Li&author=S.%20Li&author=HB.%20Li&author=GF.%20Deng&author=WH.%20Ling&author=XR.%20Xu&journal=Food%20Funct&volume=4&issue=4&pages=530-538&publication_year=2013) (http://scholar.google.com/scholar\_lookup?

title=Antiproliferative%20activities%20of%20tea%20and%20herbal%20infusions&author=F.%20Li&author=S.%20Li&author=HB.%20Li&author=GF.%20Deng&author=WH.%20Ling&author=XR.%20Xu&journal=Food%20Funct&volume=4&issue=4&pages=530-538&publication\_year=2013)

Liberti MV, Locasale JW (2016) The Warburg effect: how does it benefit cancer cells? *Trends Biochem Sci* 41(3):211–218

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=26778478) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=26778478)

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4783224) (http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4783224)

[CrossRef](https://doi.org/10.1016/j.tibs.2015.12.001) (https://doi.org/10.1016/j.tibs.2015.12.001)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=The%20Warburg%20effect%3A%20how%20does%20it%20benefit%20cancer%20cells%3F&author=MV.%20Liberti&author=JW.%20Locasale&journal=Trends%20Biochem%20Sci&volume=41&issue=3&pages=211-218&publication_year=2016) (http://scholar.google.com/scholar\_lookup?

title=The%20Warburg%20effect%3A%20how%20does%20it%20benefit%20cancer%20cells%3F&author=MV.%20Liberti&author=JW.%20Locasale&journal=Trends%20Biochem%20Sci&volume=41&issue=3&pages=211-218&publication\_year=2016)

- Liu CM, Kao CL, Tseng YT, Lo YC, Chen CY (2017) Ginger phytochemicals inhibit cell growth and modulate drug resistance factors in docetaxel resistant prostate cancer cell. *Molecules* 22(9):1477. <https://doi.org/10.3390/molecules22091477> (<https://doi.org/10.3390/molecules22091477>)  
[CrossRef](https://doi.org/10.3390/molecules22091477) (<https://doi.org/10.3390/molecules22091477>)  
[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC6151784) (<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC6151784>)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Ginger%20phytochemicals%20inhibit%20cell%20growth%20and%20modulate%20drug%20resistance%20factors%20in%20docetaxel%20resistant%20prostate%20cancer%20cell&author=CM.%20Liu&author=CL.%20Kao&author=YT.%20Tseng&author=YC.%20Lo&author=CY.%20Chen&journal=Molecules&volume=22&issue=9&pages=1477&publication_year=2017&doi=10.3390%2Fmolecules22091477) ([http://scholar.google.com/scholar\\_lookup?title=Ginger%20phytochemicals%20inhibit%20cell%20growth%20and%20modulate%20drug%20resistance%20factors%20in%20docetaxel%20resistant%20prostate%20cancer%20cell&author=CM.%20Liu&author=CL.%20Kao&author=YT.%20Tseng&author=YC.%20Lo&author=CY.%20Chen&journal=Molecules&volume=22&issue=9&pages=1477&publication\\_year=2017&doi=10.3390%2Fmolecules22091477](http://scholar.google.com/scholar_lookup?title=Ginger%20phytochemicals%20inhibit%20cell%20growth%20and%20modulate%20drug%20resistance%20factors%20in%20docetaxel%20resistant%20prostate%20cancer%20cell&author=CM.%20Liu&author=CL.%20Kao&author=YT.%20Tseng&author=YC.%20Lo&author=CY.%20Chen&journal=Molecules&volume=22&issue=9&pages=1477&publication_year=2017&doi=10.3390%2Fmolecules22091477))
- Manju V, Viswanathan P, Nalini N (2006) Hypolipidemic effect of ginger in 1,2-dimethyl hydrazine-induced experimental colon carcinogenesis. *Toxicol Mech Methods* 16(8):461–472  
[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20021021) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=20021021](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20021021))  
[CrossRef](https://doi.org/10.1080/15376520600728811) (<https://doi.org/10.1080/15376520600728811>)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Hypolipidemic%20effect%20of%20ginger%20in%201%2C2-dimethyl%20hydrazine-induced%20experimental%20colon%20carcinogenesis&author=V.%20Manju&author=P.%20Viswanathan&author=N.%20Nalini&journal=Toxicol%20Mech%20Methods&volume=16&issue=8&pages=461-472&publication_year=2006) ([http://scholar.google.com/scholar\\_lookup?title=Hypolipidemic%20effect%20of%20ginger%20in%201%2C2-dimethyl%20hydrazine-induced%20experimental%20colon%20carcinogenesis&author=V.%20Manju&author=P.%20Viswanathan&author=N.%20Nalini&journal=Toxicol%20Mech%20Methods&volume=16&issue=8&pages=461-472&publication\\_year=2006](http://scholar.google.com/scholar_lookup?title=Hypolipidemic%20effect%20of%20ginger%20in%201%2C2-dimethyl%20hydrazine-induced%20experimental%20colon%20carcinogenesis&author=V.%20Manju&author=P.%20Viswanathan&author=N.%20Nalini&journal=Toxicol%20Mech%20Methods&volume=16&issue=8&pages=461-472&publication_year=2006))
- Mansour MA, Bekheet SA, Al-Rejaie SS, Al-Shabanah OA, Al-Howiriny TA, Al-Rikabi AC, Abdo AA (2010) Ginger ingredients inhibit the development of diethylnitrosoamine induced premalignant phenotype in rat chemical hepato carcinogenesis model. *Biofactors* 36(6):483–490  
[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20872761) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=20872761](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20872761))  
[CrossRef](https://doi.org/10.1002/biof.122) (<https://doi.org/10.1002/biof.122>)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Ginger%20ingredients%20inhibit%20the%20development%20of%20diethylnitrosoamine%20induced%20pre-malignant%20phenotype%20in%20rat%20chemical%20hepato%20carcinogenesis%20model&author=MA.%20Mansour&author=SA.%20Bekheet&author=SS.%20Al-Rejaie&author=OA.%20Al-Shabanah&author=TA.%20Al-Howiriny&author=AC.%20Al-Rikabi&author=AA.%20Abdo&journal=Biofactors&volume=36&issue=6&pages=483-490&publication_year=2010) ([http://scholar.google.com/scholar\\_lookup?title=Ginger%20ingredients%20inhibit%20the%20development%20of%20diethylnitrosoamine%20induced%20pre-malignant%20phenotype%20in%20rat%20chemical%20hepato%20carcinogenesis%20model&author=MA.%20Mansour&author=SA.%20Bekheet&author=SS.%20Al-Rejaie&author=OA.%20Al-Shabanah&author=TA.%20Al-Howiriny&author=AC.%20Al-Rikabi&author=AA.%20Abdo&journal=Biofactors&volume=36&issue=6&pages=483-490&publication\\_year=2010](http://scholar.google.com/scholar_lookup?title=Ginger%20ingredients%20inhibit%20the%20development%20of%20diethylnitrosoamine%20induced%20pre-malignant%20phenotype%20in%20rat%20chemical%20hepato%20carcinogenesis%20model&author=MA.%20Mansour&author=SA.%20Bekheet&author=SS.%20Al-Rejaie&author=OA.%20Al-Shabanah&author=TA.%20Al-Howiriny&author=AC.%20Al-Rikabi&author=AA.%20Abdo&journal=Biofactors&volume=36&issue=6&pages=483-490&publication_year=2010))
- Martin ACBM, Fuzer AM, Becceneri AB, da Silva JA, Tomasin R, Denoyer D, Kim SH, McIntyre KA, Pearson HB, Yeo B, Nagpal A, Ling X, Selistre-de-Araújo HS, Vieira PC, Cominetti MR, Pouliot N (2017) [10]-gingerol induces apoptosis and inhibits metastatic dissemination of triple negative breast cancer in vivo. *Oncotarget* 8(42):72260–72271  
[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=29069785) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=29069785](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=29069785))  
[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC5641128) (<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC5641128>)  
[CrossRef](https://doi.org/10.18632/oncotarget.20139) (<https://doi.org/10.18632/oncotarget.20139>)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=%5B10%5D-10-gingerol%20induces%20apoptosis%20and%20inhibits%20metastatic%20dissemination%20of%20triple%20negative%20breast%20cancer%20in%20vivo&author=ACBM.%20Martin&author=AM.%20Fuzer&author=AB.%20Becceneri&author=JA.%20Silva&author=R.%20Tomasin&author=D.%20Denoyer&author=SH.%20Kim&author=KA.%20McIntyre&author=HB.%20Pearson&author=B.%20Yeo&author=A.%20Nagpal&author=X.%20Ling&author=HS.%20Selistre-de-Ara%20C3%20BAjo&author=PC.%20Vieira&author=MR.%20Cominetti&author=N.%20Pouliot&journal=Oncotarget&volume=8&issue=42&pages=72260-72271&publication_year=2017) ([http://scholar.google.com/scholar\\_lookup?title=%5B10%5D-10-gingerol%20induces%20apoptosis%20and%20inhibits%20metastatic%20dissemination%20of%20triple%20negative%20breast%20cancer%20in%20vivo&author=ACBM.%20Martin&author=AM.%20Fuzer&author=AB.%20Becceneri&author=JA.%20Silva&author=R.%20Tomasin&author=D.%20Denoyer&author=SH.%20Kim&author=KA.%20McIntyre&author=HB.%20Pearson&author=B.%20Yeo&author=A.%20Nagpal&author=X.%20Ling&author=HS.%20Selistre-de-Ara%20C3%20BAjo&author=PC.%20Vieira&author=MR.%20Cominetti&author=N.%20Pouliot&journal=Oncotarget&volume=8&issue=42&pages=72260-72271&publication\\_year=2017](http://scholar.google.com/scholar_lookup?title=%5B10%5D-10-gingerol%20induces%20apoptosis%20and%20inhibits%20metastatic%20dissemination%20of%20triple%20negative%20breast%20cancer%20in%20vivo&author=ACBM.%20Martin&author=AM.%20Fuzer&author=AB.%20Becceneri&author=JA.%20Silva&author=R.%20Tomasin&author=D.%20Denoyer&author=SH.%20Kim&author=KA.%20McIntyre&author=HB.%20Pearson&author=B.%20Yeo&author=A.%20Nagpal&author=X.%20Ling&author=HS.%20Selistre-de-Ara%20C3%20BAjo&author=PC.%20Vieira&author=MR.%20Cominetti&author=N.%20Pouliot&journal=Oncotarget&volume=8&issue=42&pages=72260-72271&publication_year=2017))
- Nakamura Y, Yoshida C, Murakami A, Ohigashi H, Osawa T, Uchida K (2004) Zerumbone, a tropical ginger sesquiterpene, activates phase II drug metabolizing enzymes. *FEBS Lett* 572(1–3):245–250  
[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=15304356) ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=15304356](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=15304356))  
[CrossRef](https://doi.org/10.1016/j.febslet.2004.07.042) (<https://doi.org/10.1016/j.febslet.2004.07.042>)  
[Google Scholar](http://scholar.google.com/scholar_lookup?title=Zerumbone%2C%20a%20tropical%20ginger%20sesquiterpene%2C%20activates%20phase%20I%20drug%20metabolizing%20enzymes&author=Y.%20Nakamura&author=C.%20Yoshida&author=A.%20Murakami&author=H.%20Ohigashi&author=T.%20Osawa&author=K.%20Uchida&journal=FEBS%20Lett&volume=572&issue=1&E2%80%93&pages=245-250&publication_year=2004) ([http://scholar.google.com/scholar\\_lookup?title=Zerumbone%2C%20a%20tropical%20ginger%20sesquiterpene%2C%20activates%20phase%20I%20drug%20metabolizing%20enzymes&author=Y.%20Nakamura&author=C.%20Yoshida&author=A.%20Murakami&author=H.%20Ohigashi&author=T.%20Osawa&author=K.%20Uchida&journal=FEBS%20Lett&volume=572&issue=1&E2%80%93&pages=245-250&publication\\_year=2004](http://scholar.google.com/scholar_lookup?title=Zerumbone%2C%20a%20tropical%20ginger%20sesquiterpene%2C%20activates%20phase%20I%20drug%20metabolizing%20enzymes&author=Y.%20Nakamura&author=C.%20Yoshida&author=A.%20Murakami&author=H.%20Ohigashi&author=T.%20Osawa&author=K.%20Uchida&journal=FEBS%20Lett&volume=572&issue=1&E2%80%93&pages=245-250&publication_year=2004))
- Nonn L, Duong D, Peehl DM (2007) 2006. Chemopreventive anti-inflammatory activities of curcumin and other phytochemicals mediated by MAP kinase phosphatase-5 in prostate cells. *Carcinogenesis* 28(6):1188–1196. <https://doi.org/10.1093/carcin/bgl241> (<https://doi.org/10.1093/carcin/bgl241>). *Epub* 2006 Dec 6

**CrossRef** (<https://doi.org/10.1093/carcin/bgl241>)

**PubMed** ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=17151092](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=17151092))

**Google Scholar** ([http://scholar.google.com/scholar\\_lookup?title=2006.%20Chemopreventive%20anti-inflammatory%20activities%20of%20curcumin%20and%20other%20phytochemicals%20mediated%20by%20MAP%20kinase%20phosphatase-5%20in%20prostate%20cells&author=L.%20Nonn&author=D.%20Duong&author=DM.%20Peehl&journal=Carcinogenesis&volume=28&issue=6&pages=1188-1196&publication\\_year=2007&doi=10.1093%2Fcarcin%2Fbgl241](http://scholar.google.com/scholar_lookup?title=2006.%20Chemopreventive%20anti-inflammatory%20activities%20of%20curcumin%20and%20other%20phytochemicals%20mediated%20by%20MAP%20kinase%20phosphatase-5%20in%20prostate%20cells&author=L.%20Nonn&author=D.%20Duong&author=DM.%20Peehl&journal=Carcinogenesis&volume=28&issue=6&pages=1188-1196&publication_year=2007&doi=10.1093%2Fcarcin%2Fbgl241))

Noonan DM, Benelli R, Albini A (2007) Angiogenesis and cancer prevention: a vision. *Recent Results Cancer Res* 174:219–224

**PubMed** ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=17302199](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=17302199))

**CrossRef** ([https://doi.org/10.1007/978-3-540-37696-5\\_19](https://doi.org/10.1007/978-3-540-37696-5_19))

**Google Scholar** ([http://scholar.google.com/scholar\\_lookup?title=Angiogenesis%20and%20cancer%20prevention%3A%20a%20vision&author=DM.%20Noonan&author=R.%20Benelli&author=A.%20Albini&journal=Recent%20Results%20Cancer%20Res&volume=174&pages=219-224&publication\\_year=2007](http://scholar.google.com/scholar_lookup?title=Angiogenesis%20and%20cancer%20prevention%3A%20a%20vision&author=DM.%20Noonan&author=R.%20Benelli&author=A.%20Albini&journal=Recent%20Results%20Cancer%20Res&volume=174&pages=219-224&publication_year=2007))

Park GH, Park JH, Song HM, Eo HJ, Kim MK, Lee JW, Lee MH, Cho KH, Lee JR, Cho HJ, Jeong JB (2014) Anti-cancer activity of ginger (*Zingiber officinale*) leaf through the expression of activating transcription factor 3 in human colorectal cancer cells. *BMC Complement Altern Med* 14:408

**PubMed** ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=25338635](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=25338635))

**PubMedCentral** (<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4210498>)

**CrossRef** (<https://doi.org/10.1186/1472-6882-14-408>)

**Google Scholar** ([http://scholar.google.com/scholar\\_lookup?title=Anti-cancer%20activity%20of%20ginger%20%28Zingiber%20officinale%29%20leaf%20through%20the%20expression%20of%20activating%20transcription%20factor%203%20in%20human%20colorectal%20cancer%20cells&author=GH.%20Park&author=JH.%20Park&author=HM.%20Song&author=HJ.%20Eo&author=MK.%20Kim&author=JW.%20Lee&author=MH.%20Lee&author=KH.%20Cho&author=JR.%20Lee&author=HJ.%20Cho&author=JB.%20Jeong&journal=BMC%20Complement%20Altern%20Med&volume=14&pages=408&publication\\_year=2014](http://scholar.google.com/scholar_lookup?title=Anti-cancer%20activity%20of%20ginger%20%28Zingiber%20officinale%29%20leaf%20through%20the%20expression%20of%20activating%20transcription%20factor%203%20in%20human%20colorectal%20cancer%20cells&author=GH.%20Park&author=JH.%20Park&author=HM.%20Song&author=HJ.%20Eo&author=MK.%20Kim&author=JW.%20Lee&author=MH.%20Lee&author=KH.%20Cho&author=JR.%20Lee&author=HJ.%20Cho&author=JB.%20Jeong&journal=BMC%20Complement%20Altern%20Med&volume=14&pages=408&publication_year=2014))

Plummer SMKA, Holloway MM, Manson et al (1999) Inhibition of cyclo-oxygenase 2 expression in colon cells by the chemopreventive agent curcumin involves inhibition of NF- $\kappa$ B activation via the NIK/IKK signalling complex. *Oncogene* 18(44):6013–6020

**Google Scholar** (<https://scholar.google.com/scholar?q=Plummer%20SMKA%20C%20Holloway%20MM%20C%20Manson%20et%20al%20%281999%29%20Inhibition%20of%20cyclo-oxygenase%202%20expression%20in%20colon%20cells%20by%20the%20chemopreventive%20agent%20curcumin%20involves%20inhibition%20of%20NF-%CE%BA%20activation%20via%20the%20NIK%20FIKK%20signalling%20complex.%20Oncogene%2018%2844%29%3A6013%E2%80%936020>)

Pournaderi PS, Yaghmaei P, Khodaei H, Noormohammadi Z, Hejazi SH (2017) 2017. The effects of 6-Gingerol on reproductive improvement, liver functioning and Cyclooxygenase-2 gene expression in estradiol valerate - induced polycystic ovary syndrome in Wistar rats. *Biochem Biophys Res Commun* 484(2):461–466. <https://doi.org/10.1016/j.bbrc.2017.01.057> (<https://doi.org/10.1016/j.bbrc.2017.01.057>). Epub 2017 Jan 16

**CrossRef** (<https://doi.org/10.1016/j.bbrc.2017.01.057>)

**PubMed** ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=28093231](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=28093231))

**Google Scholar** ([http://scholar.google.com/scholar\\_lookup?title=2017.%20The%20effects%20of%206-Gingerol%20on%20reproductive%20improvement%20C%20liver%20functioning%20and%20Cyclooxygenase-2%20gene%20expression%20in%20estradiol%20valerate%20-%20induced%20polycystic%20ovary%20syndrome%20in%20Wistar%20rats&author=PS.%20Pournaderi&author=P.%20Yaghmaei&author=H.%20Khodaei&author=Z.%20Noormohammadi&author=SH.%20Hejazi&journal=Biochem%20Biophys%20Res%20Commun&volume=484&issue=2&pages=461-466&publication\\_year=2017&doi=10.1016%2Fj.bbrc.2017.01.057](http://scholar.google.com/scholar_lookup?title=2017.%20The%20effects%20of%206-Gingerol%20on%20reproductive%20improvement%20C%20liver%20functioning%20and%20Cyclooxygenase-2%20gene%20expression%20in%20estradiol%20valerate%20-%20induced%20polycystic%20ovary%20syndrome%20in%20Wistar%20rats&author=PS.%20Pournaderi&author=P.%20Yaghmaei&author=H.%20Khodaei&author=Z.%20Noormohammadi&author=SH.%20Hejazi&journal=Biochem%20Biophys%20Res%20Commun&volume=484&issue=2&pages=461-466&publication_year=2017&doi=10.1016%2Fj.bbrc.2017.01.057))

Prasad S, Tyagi AK (2015) Ginger and its constituents: role in prevention and treatment of gastrointestinal cancer. *Gastroenterol Res Pract* 2015:142979

**PubMed** ([http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list\\_uids=25838819](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=25838819))

**PubMedCentral** (<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4369959>)

[CrossRef](https://doi.org/10.1155/2015/142979) (https://doi.org/10.1155/2015/142979)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Ginger%20and%20its%20constituents%3A%20role%20in%20prevention%20and%20treatment%20of%20gastrointestinal%20cancer&author=S.%20Prasad&author=AK.%20Tyagi&journal=Gastroenterol%20Res%20Pract&volume=2015&publication_year=2015) (http://scholar.google.com/scholar\_lookup?

title=Ginger%20and%20its%20constituents%3A%20role%20in%20prevention%20and%20treatment%20of%20gastrointestinal%20cancer&author=S.%20Prasad&author=AK.%20Tyagi&journal=Gastroenterol%20Res%20Pract&volume=2015&publication\_year=2015)

**Radhakrishnan EK, Bava SV, Narayanan SS, Nath LR, Thulasidasan AK, Soniya EV, Anto RJ (2014) [6]-Gingerol induces caspase-dependent apoptosis and prevents PMA-induced proliferation in colon cancer cells by inhibiting MAPK/AP-1 signaling. PLoS One 9(8):e104401**

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=25157570) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=25157570)

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4144808) (http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4144808)

[CrossRef](https://doi.org/10.1371/journal.pone.0104401) (https://doi.org/10.1371/journal.pone.0104401)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=%5B6%5D-Gingerol%20induces%20caspase-dependent%20apoptosis%20and%20prevents%20PMA-induced%20proliferation%20in%20colon%20cancer%20cells%20by%20inhibiting%20MAPK%20FAP-1%20signaling&author=EK.%20Radhakrishnan&author=SV.%20Bava&author=SS.%20Narayanan&author=LR.%20Nath&author=AK.%20Thulasidasan&author=EV.%20Soniya&author=RJ.%20Anto&journal=PLoS%20One&volume=9&issue=8&publication_year=2014) (http://scholar.google.com/scholar\_lookup?title=%5B6%5D-

Gingerol%20induces%20caspase-dependent%20apoptosis%20and%20prevents%20PMA-induced%20proliferation%20in%20colon%20cancer%20cells%20by%20inhibiting%20MAPK%20FAP-1%20signaling&author=EK.%20Radhakrishnan&author=SV.%20Bava&author=SS.%20Narayanan&author=LR.%20Nath&author=AK.%20Thulasidasan&author=EV.%20Soniya&author=RJ.%20Anto&journal=PLoS%20One&volume=9&issue=8&publication\_year=2014)

**Saha A, Blando J, Silver E, Beltran L, Sessler J, DiGiovanni J (2014) 6-Shogaol from dried ginger inhibits growth of prostate cancer cells both in vitro and in vivo through inhibition of STAT3 and NF-κB signaling. Cancer Prev Res (Phila) 7(6):627–638**

[CrossRef](https://doi.org/10.1158/1940-6207.CAPR-13-0420) (https://doi.org/10.1158/1940-6207.CAPR-13-0420)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=6-Shogaol%20from%20dried%20ginger%20inhibits%20growth%20of%20prostate%20cancer%20cells%20both%20in%20vitro%20and%20in%20vivo%20through%20inhibition%20of%20STAT3%20and%20NF-%CE%BA%20signaling&author=A.%20Saha&author=J.%20Blando&author=E.%20Silver&author=L.%20Beltran&author=J.%20Sessler&author=J.%20DiGiovanni&journal=Cancer%20Prev%20Res%20Phila%29&volume=7&issue=6&pages=627-638&publication_year=2014) (http://scholar.google.com/scholar\_lookup?title=6-

Shogaol%20from%20dried%20ginger%20inhibits%20growth%20of%20prostate%20cancer%20cells%20both%20in%20vitro%20and%20in%20vivo%20through%20inhibition%20of%20STAT3%20and%20NF-%CE%BA%20signaling&author=A.%20Saha&author=J.%20Blando&author=E.%20Silver&author=L.%20Beltran&author=J.%20Sessler&author=J.%20DiGiovanni&journal=Cancer%20Prev%20Res%20Phila%29&volume=7&issue=6&pages=627-638&publication\_year=2014)

**Shanmugam KR, Ramakrishna CH, Mallikarjuna K, Reddy KS (2009) Perturbation in kidney lipid metabolic profiles in diabetic rats with reference to alcoholic oxidative stress. Indian J Nephrol 19(3):101–106**

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20436729) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=20436729)

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2859474) (http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2859474)

[CrossRef](https://doi.org/10.4103/0971-4065.57106) (https://doi.org/10.4103/0971-4065.57106)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Perturbation%20in%20kidney%20lipid%20metabolic%20profiles%20in%20diabetic%20rats%20with%20reference%20to%20alcoholic%20oxidative%20stress&author=KR.%20Shanmugam&author=CH.%20Ramakrishna&author=K.%20Mallikarjuna&author=KS.%20Reddy&journal=Indian%20J%20Nephrol&volume=19&issue=3&pages=101-106&publication_year=2009) (http://scholar.google.com/scholar\_lookup?

title=Perturbation%20in%20kidney%20lipid%20metabolic%20profiles%20in%20diabetic%20rats%20with%20reference%20to%20alcoholic%20oxidative%20stress&author=KR.%20Shanmugam&author=CH.%20Ramakrishna&author=K.%20Mallikarjuna&author=KS.%20Reddy&journal=Indian%20J%20Nephrol&volume=19&issue=3&pages=101-106&publication\_year=2009)

**Shanmugam KR, Shanmugam B, Subbaiah GV, Ravi S, Reddy KS (2021) Medicinal plants and bioactive compounds for diabetes management: important advances in drug discovery. Curr Pharm Des 27(6):763–774**

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=32988345) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=32988345)

[CrossRef](https://doi.org/10.2174/1381612826666200928160357) (https://doi.org/10.2174/1381612826666200928160357)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Medicinal%20plants%20and%20bioactive%20compounds%20for%20diabetes%20management%3A%20important%20advances%20in%20drug%20discovery&author=KR.%20Shanmugam&author=B.%20Shanmugam&author=GV.%20Subbaiah&author=S.%20Ravi&author=KS.%20Reddy&journal=Curr%20Pharm%20Des&volume=27&issue=6&pages=763-774&publication_year=2021) (http://scholar.google.com/scholar\_lookup?

title=Medicinal%20plants%20and%20bioactive%20compounds%20for%20diabetes%20management%3A%20important%20advances%20in%20drug%20discovery&author=KR.%20Shanmugam&author=B.%20Shanmugam&author=GV.%20Subbaiah&author=S.%20Ravi&author=KS.%20Reddy&journal=Curr%20Pharm%20Des&volume=27&issue=6&pages=763-774&publication\_year=2021)

**Shukla Y, Singh M (2007) Cancer preventive properties of ginger: a brief review. Food Chem Toxicol 45(5):683–690**

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=17175086) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=17175086)

[CrossRef](https://doi.org/10.1016/j.fct.2006.11.002) (https://doi.org/10.1016/j.fct.2006.11.002)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Cancer%20preventive%20properties%20of%20ginger%3A%20a%20brief%20review&author=Y.%20Shukla&author=M.%20Singh&journal=Food%20Chem%20Toxicol&volume=45&issue=5&pages=683-690&publication_year=2007) (http://scholar.google.com/scholar\_lookup?

title=Cancer%20preventive%20properties%20of%20ginger%3A%20a%20brief%20review&author=Y.%20Shukla&author=M.%20Singh&journal=Food%20Chem%20Toxicol&volume=45&issue=5&pages=683-690&publication\_year=2007)

**Sies H, Berndt C, Jones DP (2017) Oxidative stress. Annu Rev Biochem 86(1):715–748.**

<https://doi.org/10.1146/annurev-biochem-061516-045037> (https://doi.org/10.1146/annurev-

biochem-061516-045037). Epub 2017 Apr 24

[CrossRef](https://doi.org/10.1146/annurev-biochem-061516-045037) (https://doi.org/10.1146/annurev-biochem-061516-045037)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Oxidative%20stress&author=H.%20Sies&author=C.%20Berndt&author=DP.%20Jones&journal=Annu%20Rev%20Biochem&volume=20&issue=86&pages=715-748&publication_year=2017&doi=10.1146%2Fannurev-biochem-061516-045037) (http://scholar.google.com/scholar\_lookup?

title=Oxidative%20stress&author=H.%20Sies&author=C.%20Berndt&author=DP.%20Jones&journal=Annu%20Rev%20Biochem&volume=20&issue=86&pages=715-748&publication\_year=2017&doi=10.1146%2Fannurev-biochem-061516-045037)

Surh YJ, Park KK, Chun KS, Lee LJ, Lee E, Lee SS (1999) Anti-tumor-promoting activities of selected pungent phenolic substances present in ginger. *J Environ Pathol Toxicol Oncol* 18(2):131–139

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=15281225) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=15281225)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Anti-tumor-promoting%20activities%20of%20selected%20pungent%20phenolic%20substances%20present%20in%20ginger&author=YJ.%20Surh&author=KK.%20Park&author=KS.%20Chun&author=LJ.%20Lee&author=E.%20Lee&author=SS.%20Lee&journal=J%20Environ%20Pathol%20Toxicol%20Oncol&volume=18&issue=2&pages=131-139&publication_year=1999) (http://scholar.google.com/scholar\_lookup?title=Anti-tumor-

promoting%20activities%20of%20selected%20pungent%20phenolic%20substances%20present%20in%20ginger&author=YJ.%20Surh&author=KK.%20Park&author=KS.%20Chun&author=LJ.%20Lee&author=E.%20Lee&author=SS.%20Lee&journal=J%20Environ%20Pathol%20Toxicol%20Oncol&volume=18&issue=2&pages=131-139&publication\_year=1999)

Takada Y, Murakami A, Aggarwal BB (2005) Zerumbone abolishes NF-kappa B and I kappa B kinase activation leading to suppression of antiapoptotic and metastatic gene expression, upregulation of apoptosis, and downregulation of invasion. *Oncogene* 24(46):6957–6969

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=16007145) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=16007145)

[CrossRef](https://doi.org/10.1038/sj.onc.1208845) (https://doi.org/10.1038/sj.onc.1208845)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Zerumbone%20abolishes%20NF-kappa%20B%20and%20I%20kappa%20B%20kinase%20activation%20leading%20to%20suppression%20of%20antiapoptotic%20and%20metastatic%20gene%20expression%20C%20upregulation%20of%20apoptosis%20C%20and%20downregulation%20of%20invasion&author=Y.%20Takada&author=A.%20Murakami&author=BB.%20Aggarwal&journal=Oncogene&volume=24&issue=46&pages=6957-6969&publication_year=2005) (http://scholar.google.com/scholar\_lookup?title=Zerumbone%20abolishes%20NF-kappa%20B%20and%20I%20kappa%20B%20kinase%20activation%20leading%20to%20suppression%20of%20antiapoptotic%20and%20metastatic%20gene%20expression%20C%20upregulation%20of%20apoptosis%20C%20and%20downregulation%20of%20invasion&author=Y.%20Takada&author=A.%20Murakami&author=BB.%20Aggarwal&journal=Oncogene&volume=24&issue=46&pages=6957-6969&publication\_year=2005)

Trachootham D, Lu W, Ogasawara MA, Nilsa RD, Huang P (2008) Redox regulation of cell survival.

*Antioxid Redox Signal* 10(8):1343–1374

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=18522489) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=18522489)

[PubMedCentral](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2932530) (http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2932530)

[CrossRef](https://doi.org/10.1089/ars.2007.1957) (https://doi.org/10.1089/ars.2007.1957)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Redox%20regulation%20of%20cell%20survival&author=D.%20Trachootham&author=W.%20Lu&author=MA.%20Ogasawara&author=RD.%20Nilsa&author=P.%20Huang&journal=Antioxid%20Redox%20Signal&volume=10&issue=8&pages=1343-1374&publication_year=2008) (http://scholar.google.com/scholar\_lookup?

title=Redox%20regulation%20of%20cell%20survival&author=D.%20Trachootham&author=W.%20Lu&author=MA.%20Ogasawara&author=RD.%20Nilsa&author=P.%20Huang&journal=Antioxid%20Redox%20Signal&volume=10&issue=8&pages=1343-1374&publication\_year=2008)

Tsuboi K, Matsuo Y, Shamoto T, Shibata T, Koide S, Morimoto M, Guha S, Sung B, Aggarwal BB,

Takahashi H, Takeyama H (2014) Zerumbone inhibits tumor angiogenesis via NF-κB in gastric cancer.

*Oncol Rep* 31(1):57–64

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=24220661) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=24220661)

[CrossRef](https://doi.org/10.3892/or.2013.2842) (https://doi.org/10.3892/or.2013.2842)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Zerumbone%20inhibits%20tumor%20angiogenesis%20via%20NF-%20CE%20B%20in%20gastric%20cancer&author=K.%20Tsuboi&author=Y.%20Matsuo&author=T.%20Shamoto&author=T.%20Shibata&author=S.%20Koide&author=M.%20Morimoto&author=S.%20Guha&author=B.%20Sung&author=BB.%20Aggarwal&author=H.%20Takahashi&author=H.%20Takeyama&journal=Oncol%20Rep&volume=31&issue=1&pages=57-64&publication_year=2014) (http://scholar.google.com/scholar\_lookup?

title=Zerumbone%20inhibits%20tumor%20angiogenesis%20via%20NF-%20CE%20B%20in%20gastric%20cancer&author=K.%20Tsuboi&author=Y.%20Matsuo&author=T.%20Shamoto&author=T.%20Shibata&author=S.%20Koide&author=M.%20Morimoto&author=S.%20Guha&author=B.%20Sung&author=BB.%20Aggarwal&author=H.%20Takahashi&author=H.%20Takeyama&journal=Oncol%20Rep&volume=31&issue=1&pages=57-64&publication\_year=2014)

Vijaya Padma V, Arul Diana Christie S, Ramkuma KM (2007) Induction of apoptosis by ginger in HEP-2 cell line is mediated by reactive oxygen species. *Basic Clin Pharmacol Toxicol* 100(5):302–307

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=17448115) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=17448115)

[CrossRef](https://doi.org/10.1111/j.1742-7843.2007.00046.x) (https://doi.org/10.1111/j.1742-7843.2007.00046.x)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Induction%20of%20apoptosis%20by%20ginger%20in%20HEP-2%20cell%20line%20is%20mediated%20by%20reactive%20oxygen%20species&author=V.%20Vijaya%20Padma&author=S.%20Arul%20Diana%20Christie&author=KM.%20Ramkuma&journal=Basic%20Clin%20Pharmacol%20Toxicol&volume=100&issue=5&pages=302-307&publication_year=2007) (http://scholar.google.com/scholar\_lookup?

title=Induction%20of%20apoptosis%20by%20ginger%20in%20HEP-2%20cell%20line%20is%20mediated%20by%20reactive%20oxygen%20species&author=V.%20Vijaya%20Padma&author=S.%20Arul%20Diana%20Christie&author=KM.%20Ramkuma&journal=Basic%20Clin%20Pharmacol%20Toxicol&volume=100&issue=5&pages=302-307&publication\_year=2007)

Wang CY, Staniforth V, Chiao MT, Hou CC, Wu HM, Yeh KC, Chen CH, Hwang PI, Wen TN, Shyur LF et al (2008) Genomics and proteomics of immune modulatory effects of a butanol fraction of

*echinacea purpurea* in human dendritic cells. *BMC Genomics* 2008(9):479

[CrossRef](https://doi.org/10.1186/1471-2164-9-479) (https://doi.org/10.1186/1471-2164-9-479)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Genomics%20and%20proteomics%20of%20immune%20modulatory%20effects%20of%20a%20butanol%20fraction%20of%20echinaceapurpurea%20in%20human%20dendritic%20cells&author=C.Y.%20Wang&author=V.%20Staniforth&author=MT.%20Chiao&author=CC.%20Hou&author=HM.%20Wu&author=KC.%20Yeh&author=CH.%20Chen&author=PI.%20Hwang&author=TN.%20Wen&author=LF.%20Shyur&journal=BMC%20Genomics&volume=2008&issue=9&pages=479&publication_year=2008) (http://scholar.google.com/scholar\_lookup?

title=Genomics%20and%20proteomics%20of%20immune%20modulatory%20effects%20of%20a%20butanol%20fraction%20of%20echinaceapurpurea%20in%20human%20dendritic%20cells&author=C.Y.%20Wang&author=V.%20Staniforth&author=MT.%20Chiao&author=CC.%20Hou&author=HM.%20Wu&author=KC.%20Yeh&author=CH.%20Chen&author=PI.%20Hwang&author=TN.%20Wen&author=LF.%20Shyur&journal=BMC%20Genomics&volume=2008&issue=9&pages=479&publication\_year=2008)

Weng CJ, Wu CF, Huang HW, Ho CT, Yen GC (2010) Anti-invasion effects of 6-shogaol and 6-gingerol, two active components in ginger, on human hepatocarcinoma cells. *Mol Nutr Food Res* 54(11):1618–1627

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=20521273) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=20521273)

[CrossRef](https://doi.org/10.1002/mnfr.201000108) (https://doi.org/10.1002/mnfr.201000108)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Anti-invasion%20effects%20of%206-shogaol%20and%206-gingerol%20in%20human%20hepatocarcinoma%20cells&author=CJ.%20Weng&author=CF.%20Wu&author=HW.%20Huang&author=CT.%20Ho&author=GC.%20Yen&journal=Mol%20Nutr%20Food%20Res&volume=54&issue=11&pages=1618-1627&publication_year=2010) (http://scholar.google.com/scholar\_lookup?title=Anti-

invasion%20effects%20of%206-shogaol%20and%206-gingerol%20in%20human%20hepatocarcinoma%20cells&author=CJ.%20Weng&author=CF.%20Wu&author=HW.%20Huang&author=CT.%20Ho&author=GC.%20Yen&journal=Mol%20Nutr%20Food%20Res&volume=54&issue=11&pages=1618-1627&publication\_year=2010)

Weston A, Harris CC (2003) Multistage Carcinogenesis. In: Kufe DW, Pollock RE, Weichselbaum RR et al (eds) *Holland-Frei cancer medicine*, 6th edn. BC Decker, Hamilton

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Multistage%20Carcinogenesis&author=A.%20Weston&author=CC.%20Harris&publication_year=2003) (http://scholar.google.com/scholar\_lookup?

title=Multistage%20Carcinogenesis&author=A.%20Weston&author=CC.%20Harris&publication\_year=2003)

Yeh H, Chuang C, Chen H, Wan C, Chen T, Lin L (2014) Bioactive components analysis of two various gingers (*Zingiber officinale* roscoe) and antioxidant effect of ginger extracts. *LWT-Food Sci Technol* 55:329–334

[CrossRef](https://doi.org/10.1016/j.lwt.2013.08.003) (https://doi.org/10.1016/j.lwt.2013.08.003)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Bioactive%20components%20analysis%20of%20two%20various%20gingers%20%28Zingiber%20officinale%20roscoe%29%20and%20antioxidant%20effect%20of%20ginger%20extracts&author=H.%20Yeh&author=C.%20Chuang&author=H.%20Chen&author=C.%20Wan&author=T.%20Chen&author=L.%20Lin&journal=LWT-Food%20Sci%20Technol&volume=55&pages=329-334&publication_year=2014) (http://scholar.google.com/scholar\_lookup?

title=Bioactive%20components%20analysis%20of%20two%20various%20gingers%20%28Zingiber%20officinale%20roscoe%29%20and%20antioxidant%20effect%20of%20ginger%20extracts&author=H.%20Yeh&author=C.%20Chuang&author=H.%20Chen&author=C.%20Wan&author=T.%20Chen&author=L.%20Lin&journal=LWT-Food%20Sci%20Technol&volume=55&pages=329-334&publication\_year=2014)

Yu JX, Zhou X, He M, Dai Q (2011) Zhang, Curcumin induces apoptosis involving Bax/Bcl-2 in human hepatoma SMMC-7721cells. *Asian Pac J Cancer Prev* 12:1925–1929

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=22292626) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=22292626)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Zhang%20Curcumin%20induces%20apoptosis%20involving%20Bax%20Bcl-2%20in%20human%20hepatoma%20SMMC-7721cells&author=JX.%20Yu&author=X.%20Zhou&author=M.%20He&author=Q.%20Dai&journal=Asian%20Pac%20J%20Cancer%20Prev&volume=12&pages=1925-1929&publication_year=2011) (http://scholar.google.com/scholar\_lookup?

title=Zhang%20Curcumin%20induces%20apoptosis%20involving%20Bax%20Bcl-2%20in%20human%20hepatoma%20SMMC-7721cells&author=JX.%20Yu&author=X.%20Zhou&author=M.%20He&author=Q.%20Dai&journal=Asian%20Pac%20J%20Cancer%20Prev&volume=12&pages=1925-1929&publication\_year=2011)

Zhang F, Thakur K, Hu F, Zhang JG, Wei ZJ (2017) 10-Gingerol, a phytochemical derivative from “Tongling white ginger”, inhibits cervical cancer: insights into the molecular mechanism and inhibitory targets. *J Agric Food Chem* 65:2089–2099

[PubMed](http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Abstract&list_uids=28230361) (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?

cmd=Retrieve&db=PubMed&dopt=Abstract&list\_uids=28230361)

[CrossRef](https://doi.org/10.1021/acs.jafc.7b00095) (https://doi.org/10.1021/acs.jafc.7b00095)

[Google Scholar](http://scholar.google.com/scholar_lookup?title=10-Gingerol%20a%20phytochemical%20derivative%20from%20E2%80%9CTongling%20white%20ginger%20inhibits%20cervical%20cancer%3A%20insights%20into%20the%20molecular%20mechanism%20and%20inhibitory%20targets&author=F.%20Zhang&author=K.%20Thakur&author=F.%20Hu&author=JG.%20Zhang&author=ZJ.%20Wei&journal=J%20Agric%20Food%20Chem&volume=65&pages=2089-2099&publication_year=2017) (http://scholar.google.com/scholar\_lookup?title=10-

Gingerol%20a%20phytochemical%20derivative%20from%20E2%80%9CTongling%20white%20ginger%20inhibits%20cervical%20cancer%3A%20insights%20into%20the%20molecular%20mechanism%20and%20inhibitory%20targets&author=F.%20Zhang&author=K.%20Thakur&author=F.%20Hu&author=JG.%20Zhang&author=ZJ.%20Wei&journal=J%20Agric%20Food%20Chem&volume=65&pages=2089-2099&publication\_year=2017)

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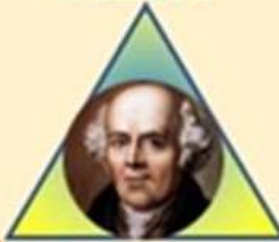
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# Lectures on Materia Medica of Selected Remedies



Master E.K.



Dr. Samuel Hahnemann



Dr. E.V.M. Acharya



Sri Ogirala Ramachandra Rao

**Lectures**  
**O. Ramachandra Rao**

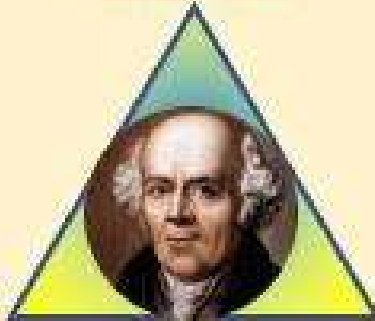
**Lecture Transcript**  
**Dr. I. S.Chakrapani**

# Homoeo Learning Classes

From August 11, 2021 to January 11, 2022



Master E.K.



Dr. Samuel Hahnemann



Dr. E.V.M. Acharya



Sri Ogirala Ramachandra Rao

Conducted By:

Sri Ogirala Ramachandra Rao

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## **FOREWORD**

*Those were the days of COVID-19 pandemic. Everywhere it was panic, turmoil, confusion, fear and doubt. Uncertainty, altered daily routine, social isolation & apprehensions ruled the society then. Restrictions & lock downs were the order of the day. Amidst this public psyche, we struggled a lot in discharging our duties in the dispensaries. As an ardent Homeopath, after a few days of observation, I could manage to serve the needy, by Master's grace, following the guidelines imposed by the government then.*

*While restricted at home, how to utilize this leisure was a big question. In a casual talk, Sri PSR Anjaneyulu, Nellore suggested me to conduct Homoeo classes online for the young generation. While I was hesitating, he encouraged me, saying that an online transaction of content was easier than conventional methods of teaching, and that would go a long way in the creation of learning resources. I hesitantly nodded.*

*I discussed the issue with a few enthusiastic youngsters like Prabhu, Aparna, Ravichandar, Padmaja and others. They started working as a team, and got it materialized. There was a brief discussion as to which platform must be selected, later on we decided to go online by youtube live streaming. We floated the idea to Sriman E.Anantkrishna, for which he was very happy and blessed us profusely. After a few initial hiccups during the trial run, it went on successfully from 11<sup>th</sup> August, 2021, on the birthday of Master EK.*

*These classes were received very well across the globe, an unexpected response! We were happy, at the same time it doubled our responsibility. I continued the classes, referring to the class notes of Master EK lectures, Kent, Dr.EVM Acharya's lectures, and of course the experiences provided by our Maser in the dispensary. More than 500 people, enthusiastic to learn Homeopathy followed the classes with utmost devotion, for which I wondered a lot. Many of those already working in our dispensaries followed and refreshed their knowledge. There were a few health issues, but the classes went on continuously till 11<sup>th</sup> January 2022 successfully. It was again a realization for all of us that when we start a work beneficial to all, resources and personnel come to us effortlessly.*

*While listening to the classes, Smt. Leela of Hyderabad made lecture notes in Telugu meticulously. Varchaswi & Tejaswi with the help of Sri Tarakeshwara Rao digitalized this notes diligently, and brought out a good volume. The same was provided to all by different social media. After some days, a few doctors have requested for English version of the lecture notes. This is taken as the Master's instruction, work is endorsed to Dr.I.S.Chakrapani, Nellore and the result is this compilation. Hope this will serve the purpose.*

***Vasudeva***

*Yours*



*Ogirala Ramachandra Rao*

*Guntur*

## *Before you start using this notes...*

*Let me say a few words...*

*After this mission of 'Online Homoeo Classes' was over, there was a period of brief silence. We took over the routine of our dispensaries.*

*On 23<sup>rd</sup> March 2022, Sriman Ogirala Ramachandra Rao called me and talked about the 'instruction'. A few members who followed the classes happened to be doctors, and they wanted the English version of this notes, as it was a bit difficult to follow in Telugu. As it is true with most of the young generation, an English version is always solicited. By His instruction, I took up the task.*

*I started working. I tried to prepare some notes based on the version in Telugu. While going through, I used to get a number of doubts, so I preferred watching the particular video lesson again, and then finishing the notes. This took lot of time, which practically became very difficult to cope up amidst the hectic daily work schedule. Sri Tarakeswara Rao came to my rescue, he started providing the draft notes of each remedy, one remedy a week. My 'bit rate' was slow, and the pace was not matching. Slowly I could gain the momentum to share the notes weekly. Lesson was about punctuality and keeping up the word.*

*Smt.Leela, from Hyderabad followed the classes ardently and made notes in Telugu. Her daughters, Chy.Tejaswi & Varchaswi carefully digitalized this notes. It helped me a lot, forming the basis for this notes in English. I would like to tell you that it is not exactly a translation. At times, I went beyond, at times I made a precise of the content. During this work, I consulted Dr.J.T.Kent, Master EK's lecture notes & Dr. EVM Acharya's homoeo classes, as per need. Preference was given for a concise expression, so that an enthusiastic learner easily finds the crux. Sri Subbarao of Tenali helped me improvise the presentation. Actually, when I accepted the assignment, I was afraid whether I could find time to complete. But the lesson was that, time is elastic & work gives us energy to work. The work is complete by 13.09.2022.*

*I thank Smt. Swarna & Chy. STG Sekhar for bringing out this compilation. They worked hard to re-align the content and made it into a book. The page is set so that wider margins will be useful to the young learners to make their own notes in the book itself.*

*I sincerely thank all those mentioned above. Without their dedicated contribution, this work would have not been shaped up as it is today.*

*I thank Sriman ORR for his benevolent guidance. Thanks to the invisible presence of Mater EK and Dr.EVM Acharya which guided me to be on the right path.*

*I humbly submit that merits of this work belong to all the people mentioned above. Errors if any, crept by chance are all mine. Please bring them to our notice for further correction.*

*Hope this work will serve the purpose of learning Homeopathy.*

*Submitted to the Lotus feet of Masters..*

**Vasudeva.**

A handwritten signature in black ink, appearing to read 'Chakrapani', with a horizontal line underneath it.

*Chakrapani.*

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# **Cell and Molecular Biology**

**Dr. A. Indira Priyadarsini**

**Dr. I. S. Chakrapani**

**AIIP**

# **CELL AND MOLECULAR BIOLOGY**

**FIRST EDITION**

## **Authors**

**Dr. A. Indira Priyadarsini**

**Dr. Inavolu Srinivasa Chakrapani**

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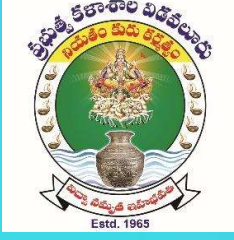
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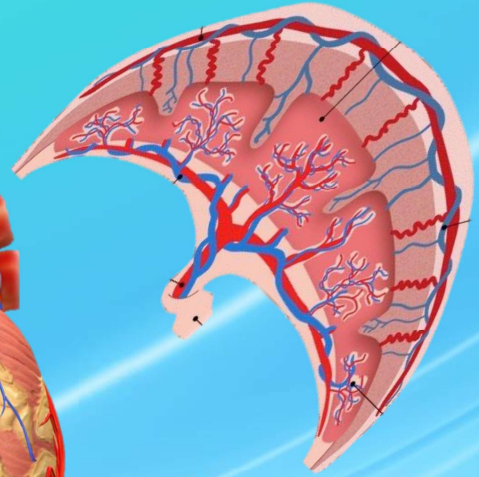
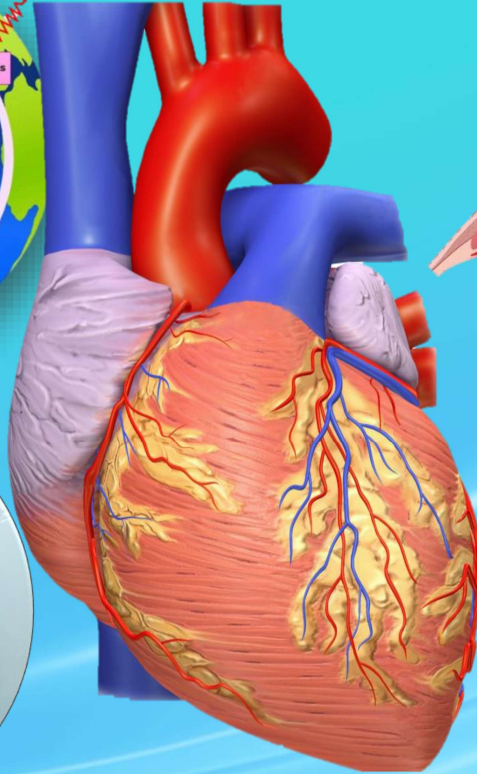
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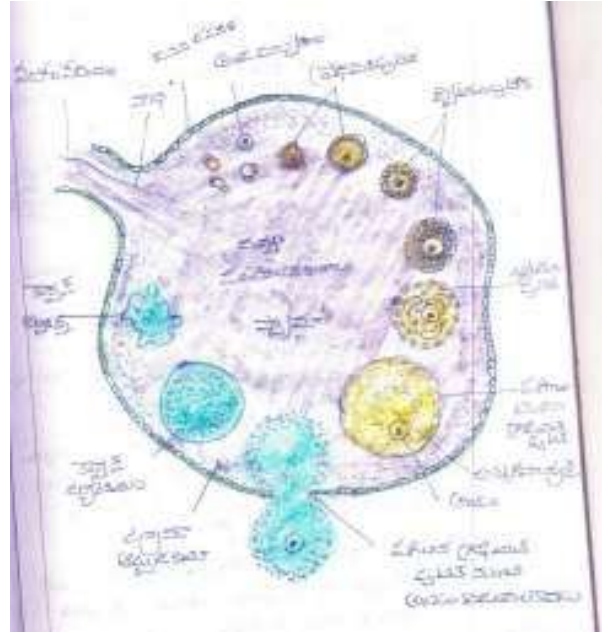
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# అండజననం (Oogenesis)

## స్త్రీబీజకోశం నిర్మాణం

అండజననం అండాశయంలో జరుగుతుంది. అండ మాతృకణాల నుంచి అండాలు ఏర్పడే విధానాన్ని అండ జననం అంటారు. సకశేరుకాలలో ఒక జత స్త్రీ బీజకోశాలు (అండాశయాలు) ఉదర కుహరంలో పృష్ఠ దేహ కుడ్య ప్రాంతంలో అతికి ఉంటాయి. పక్షులలో ఎడమ స్త్రీ బీజకోశం మాత్రం ఉంటుంది. మానవులలో స్త్రీ జీవి రెండు X క్రోమోజోమ్ ల కలయికవల్ల ఏర్పడుతుంది. X క్రోమోజోమ్ పై TDF కారకం లోపించడంవల్ల పిండాభివృద్ధి సమయంలో బీజకోశ వల్కలం (Gonadal cortex) స్త్రీ బీజకోశంగా విభేదనం చెందుతుంది. స్త్రీ బీజకోశం ఏర్పడిన సమయంలో ప్రాథమిక బీజ కణాలు (Primordial Germ cells) స్త్రీ బీజకోశం/అండాశయంలోకి వలస చెందుతాయి. పిండాభివృద్ధి సమయంలో స్త్రీ బీజకోశం అభివృద్ధి చెందుతుంది. స్త్రీ బీజకోశం వెలుపల ఉపకళా కణాలతో కూడిన ఒక స్తరం ఏర్పడుతుంది. ఈ స్తరంలోని కణాలను జనన ఉపకళాకణాలు (Germinal epithelial cells) అంటారు. అండాశయంలోని మిగిలిన లోపలి ప్రాంతం సంయోజక కణజాలంతో నింపబడుతుంది. దీన్ని స్ట్రోమా అంటారు. జనన ఉపకళాకణాలు

సమ విభజన చెందడం ద్వారా ఏర్పడే కొత్త కణాలు స్ట్రోమాలో చిన్న చిన్న గుంపులుగా ఏర్పడతాయి. ఒక్కొక్క గుంపును గ్రాఫియన్ పుటిక (Graffian follicle) గా పిలుస్తారు. ఒక గ్రాఫియన్ పుటికలో బాగా అభివృద్ధి చెందిన ఒక కణం అండ మాతృకణం (Oogonium) గా ఏర్పడుతుంది. మిగిలిన కణాలు పుటికాకణాలుగా ఉండి అభివృద్ధి చెందే అండానికి పోషక పదార్థాలను అందచేస్తాయి. అండ మాతృకణం విభజనలోకి ప్రవేశించడం ద్వారా ప్రాథమిక అండమాతృకణాలు (Primary oocytes)గా ఏర్పడతాయి.





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Pathology in Aquaculture for B.voc (CA) Core – XI , an e-book





## II SEMESTER



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## ANIMAL DIVERSITY-II

# BIOLOGY OF CHORDATES



**SREENIVAS-CHAKRAPANI-ANILKUMAR**

# Biology of Chordates

(Text book for UG II-Sem Zoology)

Dr.N.Srinivas, Dr.I.S.Chakrapani & Dr.P.Anilkumar

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**ANIMAL DIVERSITY-II**

# **BIOLOGY OF CHORDATES**

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- 1.1. General Characters and Classification of Chordates upto classes.
- 1.2 Salient features of Urochordata and Cephalochordata Salient features of Cyclostomes
- 1.3 General Characters and Classification of Fishes upto sub class level,
- 1.4. Structure and life-history of *Herdmania*, Significance of retrogressive Metamorphosis.

# **MODULE-I**

## **INTRODUCTION**

Animal kingdom is basically divided into two sub kingdoms:

(a) Nonchordata- including animals without notochord.

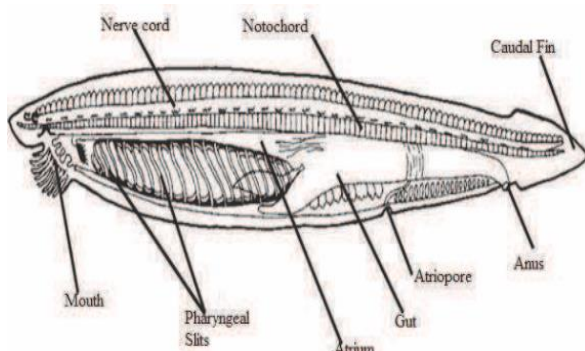
(b) Chordata- This comprising animal having notochord or chorda dorsalis. While the Chordata has a notochord at some stage during the life, it is not known to exist in the Nonchordata.

- The Chordata is the animal phylum with which everyone is most intimately familiar, since it includes humans and other vertebrates. However, not all chordates are vertebrates.

- All chordates have the following features at some stage in their life (in the case of humans and many other vertebrates, these features may only be present in the embryos).

- Pharyngeal slits – a series of openings that connect the inside of the throat to the outside of the

“neck”. These are often, but not always, used as gills.



- Dorsal tubular nerve cord – A bundle of nerve fibers which runs down the “back”. It connects the brain with the lateral muscles and other organs.

Pharyngeal slits dorsal nerve cord & Notochord

- Notochord – cartilaginous rod running underneath, and supporting, the nerve cord.

- Post-anal tail – an extension of the body posts the anal opening.

## **ORIGIN OF CHORDATES**

It is believed that chordates originated from invertebrates. However, it is difficult to determine from which invertebrate group of the chordate developed.

It is almost constant that chordate ancestors were soft bodied animals. Hence, they were not preserved as fossil.

Many theories have been put forward to explain the evolution of chordates, few of them are as follows:

### **(a) COELENTERATE THEORY:**

According to this theory chordates developed from coelenterates.

It is believed that radial symmetry coelenteron, cnidoblasts etc, disappeared and advanced characters developed to give rise the chordates.

This theory infers that chordate might have acquired higher characters independently. This theory is not acceptable.

**(b) ANNELID THEORY:** This theory suggests that the chordates have evolved from an annelid stock, like many chordates the annelids show bilateral symmetry, metasmerism, head, lateral coelome complete digestive tract, closed circulatory system, haemoglobin, etc.



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# **BIOLOGY OF CHORDATES**

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- 2.4. *Rana* : Morphology, respiratory system, structure of heart, Brain and reproductive systems only.

# **MODULE-II**

## **External Features of Dogfish (Scoliodon):**

### **Shape, Size and Colour:**

Dogfish (Scoliodon) has a long, laterally compressed spindle-shaped body tapering at both ends. The fully grown specimen measures from 30 to 60 cm in length. The colour of the body is dark grey above and pale white beneath, while the portions of the caudal fin are more or less dark. Body surface is rough due to backwardly directed spines of placoid scales embedded in the dermis.

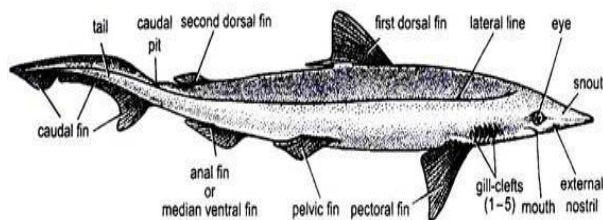


Fig. 14.1. Scoliodon. External features.

### **Division of Body:**

The body is divisible into head, trunk and tail, though there are no distinct boundaries between these regions.

#### **(i) Head:**

The head is strongly compressed dorso-ventrally and is produced in front into a wedge-shaped snout or rostrum.

#### **(ii) Trunk:**

The trunk is almost elliptical in transverse section. Its thickest part lying in front of the middle of the body. The trunk gradually tapers behind into the tail.

#### **(iii) Tail:**

The tail is laterally compressed and is bent upwards at a small angle and fringed with a caudal fin. Such a tail is known as heterocercal tail.

#### **Fins:**

Dogfish (Scoliodon) is provided with two sets of fins which are flattened expansions of the skin supported by cartilaginous rods and horny fin rays- these are unpaired or median fins and paired lateral fins.

#### **(i) Median Fins:**

The median fins are two dorsal fins, a ventral or anal fin and a caudal fin. The first dorsal fin is large and triangular in shape and is situated a little in front of the middle of the body. It has a basal lobe. The second dorsal fin is also triangular in outline but is very small and is situated midway between the first dorsal and the tip of the tail. The caudal fin extends along the dorsal and ventral



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# **BIOLOGY OF CHORDATES**

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# **MODULE-III**

## REPTILIA: CHARACTERS AND CLASSIFICATION

Reptiles are cold-blooded vertebrates, breath by lungs and having the body covered by scales or scutes.

General characters:

- ✚ They are inhabitants of terrestrial and aquatic (both marine and freshwaters) environments.
- ✚ Their skin is dry, cornified and usually covered by **epidermal** scales or scutes. There are a few integumentary scent glands secreting pheromones during breeding seasons.
- ✚ Single external nasal opening is present on the snout. Ear drums are slightly depressed.
- ✚ Two pairs of pentadactyle limbs are present. The limbs end in clawed digits.
- ✚ The cloacal opening is either transverse or longitudinal.
- ✚ A post-anal tail is present.
- ✚ The heart is composed of two auricles and a partially divided ventricle. There are right and left systemic arches.
- ✚ The kidney is metanephric type.
- ✚ Mullerian duct persists as oviduct in female and Wolffian duct is retained as vas deference in male. Males possess copulatory organs.
- ✚ Twelve pairs of cranial nerves are present.
- ✚ Vomero-nasal organ (Organ of Jacobson) is well-developed.
- ✚ Single occipital condyle in the skull is present for the attachment with atlas.
- ✚ Mandible consists usually six pieces of bones.
- ✚ Vertebrae are procoelous. Sternum is greatly developed with ribs.
- ✚ Cleidoic eggs are large. The calcareous shell serves for protection against desiccation and external injury. The shell is porous for gaseous exchange.

- ✚ Fertilisation is internal.
- ✚ Embryos are provided with extra-embryonic membranes, like amnion, chorion and allantois.



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- 4.4. Dentition in Mammals,

# **MODULE-IV**

# GENERAL CHARACTERS OF AVES

## Introduction:

Class Aves includes birds, bipedal vertebrates. Birds are unique in having feathers as their exoskeleton. Aves originated from theropod dinosaurs in Jurassic period and modernized in the cretaceous period. **T.H.Huxley** called birds

“**Glorified reptiles**”. **J.Z. Young** described them as the “**Masters of air**”. Ornithology is the study of birds.

## General characters of Aves

- Birds are cosmopolitan and found in all continents, seas and most islands.
- Their wide occurrence is due to their power of flight, which enables them to reach places unreachable to other animals.
- Most of them can fly and a few have lost the power of flight.

## Body temperature:

- They are **homeothermic (warm blooded)** and the body temperature provides high metabolic rate for quick energy supply.
- Birds are **endothermic**, and expend a lot of energy to keep warm.

## Body form and appendages:

- The body is boat shaped and streamlined.
- It is divisible into **head, neck, trunk** and **tail**.
- The forelimbs are modified into wings for flight which are supported by powerful flight muscles attached to the sternum.
- Each fore limb has 1 to 3 digits and each hind limb has 1 to 4 digits.

- The hind limbs are used for perching, walking, hopping, wading, swimming etc.
- Skin is dry and thin except for uropygial or oil gland on the tail.
- Body is covered by epidermal horny skeleton of feathers which conserve body heat, help in flight and provide colouration to the birds.

## Endoskeleton:

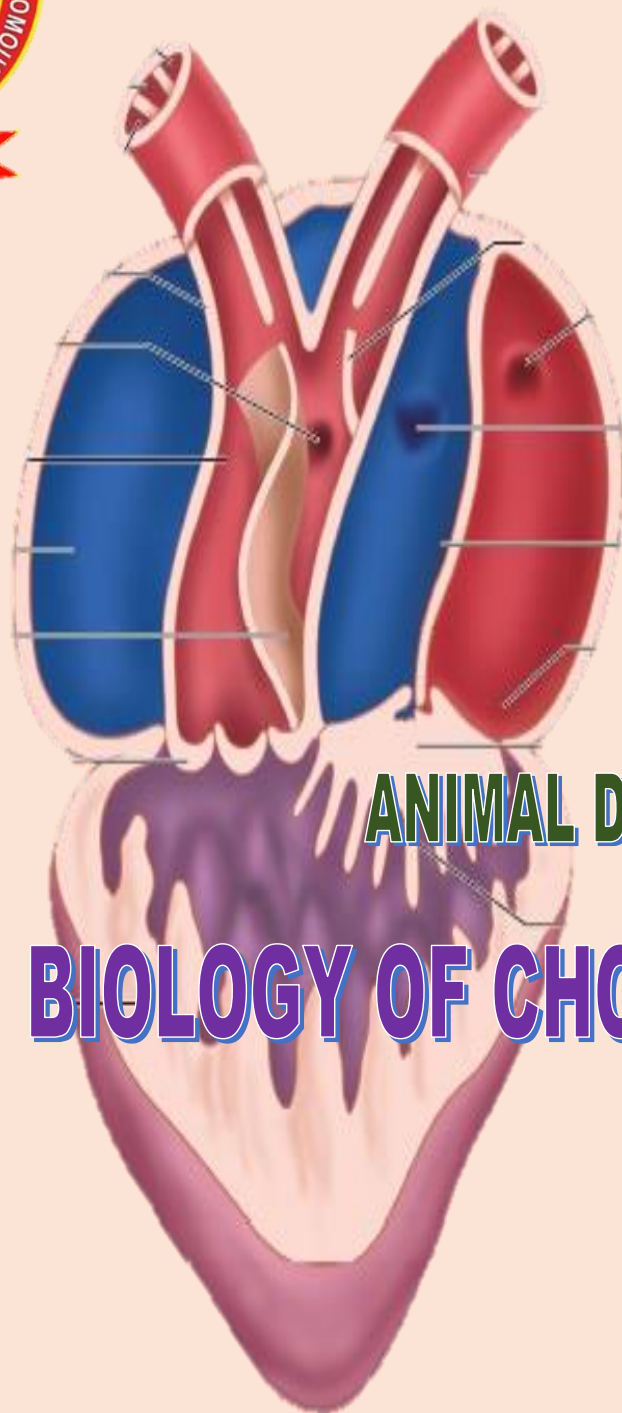
- The endoskeleton is bony, but delicate and light.
- Skull is **monocondylic**, i.e., with single occipital condyle.
- Sternum is usually large and with a median keel for the attachment of flight muscles.
- Bones are pneumatic, i.e., contain air cavities to reduce weight.

## Digestive system:

- Mouth has a wide gap and jaws are covered with horny sheaths to form strong beaks.
- Beaks are adapted to various modes of feeding: seed-crushing, fruit-scooping, fish-tearing, nectar-sipping, wood-chiselling, grain-pickling etc.
- Teeth are absent and food is swallowed un masticated.
- The **crop** stores and softens food.
- Alimentary canal often has additional chambers; **crop and gizzard**.
- Gizzard is muscular to crush and churn the softened food.
- Some birds keep stone in the gizzard to effectively crush seeds and grains. Alimentary canal leads to the cloaca.
- The cloaca is divided into three linear compartments: i.e., anterior **coprodaeum** middle **urodaeum**



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**ANIMAL DIVERSITY-II**

**BIOLOGY OF CHORDATES**

**SREENIVAS-CHAKRAPANI-ANILKUMAR**



# IoT Based Mobile App for Continuous Health Monitoring of the Person

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**Abstract**—In the sphere of medicine, IOT is meant to keep people safe and healthy plays a crucial part in communicating with doctors and patients through the use of health monitoring equipment and lowering healthcare costs in the future years. The internet of things (IoT) is making the world a smarter and more efficient village by allowing a variety of sensors and smart gadgets to gather and analyse data for a variety of reasons. As a result of these smart things, the healthcare system is growing wiser. When basic health facilities lack comprehensive medical care infrastructure, emerging countries gain. However, there is currently no specialized architecture for smart health units that can allow for this gathering and transferring patient health information to headquarters hospitals where live patient assistance is offered. Here, a smart IoT-based healthcare system is proposed, which includes a smart medical kit linked to sensors and a server for frequent health tracking. This smart medical kit is associated with sensors to measure the health parameters like body temperature, blood pressure, and heart rate for the effective function of the body. The proposed idea can alert the patient and their relatives in case of any abnormalities in their health parameters and also get suggestions from the doctor without physical contact with the doctor.

**Keywords**—Heart rate, Temperature, Blood Pressure, IoT, Blynk.

## I. INTRODUCTION

The Internet of Things (IoT) has become a globally acknowledged network technology and heavily researched area. Sensors are employed in practically every product today, from ordinary items to Sensor-based exhaustive medical systems, and industry surveillance systems are now booming. [1]. The Internet of Things (IoT) makes our lives smarter, efficient, and productive. The prototype device [2] uses a mobile phone as the data transferring platform to deliver user-friendly speech recognition and alert functionality. Because of insufficient and ineffective healthcare services to satisfy the expanding demands of a growing population with chronic conditions, health care is becoming ever more difficult to manage. Many smart or medical sensors have emerged as a result of technological advancements, which continually assess individual patient behavior and properly forecast a problem. Data collected by such sensors may be utilized for a variety of reasons by a variety of actors, including physicians, patients, family, and healthcare facilities. Reports that explain the patient's health state are created by combining analytics

and sensor data. This platform may also include data from a variety of sensors to provide mechanisms for monitoring, processing, visualizing, storing, and sending notifications about a patient's status and vital signs in real-time through Internet standards. Different types of sensing devices are employed in healthcare, depending on their qualities, usefulness, and efficiency. The growth of the IoT in the field of medicine is shown in Fig. 1 and the statistics taken from the grandviewresearch.com website.

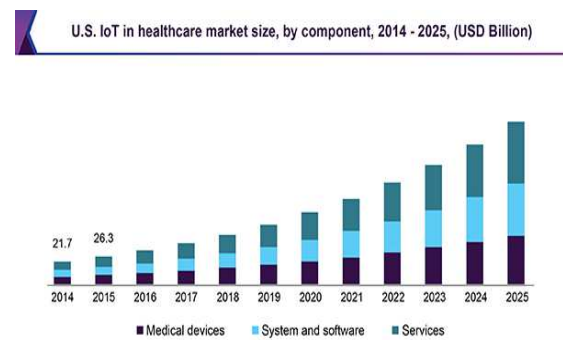


Fig. 1. IoT in health care

These networked IoT devices produce massive amounts of data that suppliers must handle well, which might be a big burden. The Internet of Things Analytics (Particle) process is used to address the difficulty of storing and analyzing large amounts of data. Using approaches such as information extraction and information analytics, the raw data is transformed into meaningful and therapeutically significant information. It is expected that by 2020, more than 50-55 % of raw data analysis approaches would be able to better handle the stream of data created by instrumented devices and applications. Mobile phone-based health assessing solutions are growing increasingly mostly as the information and technology revolution continues. These technologies can gather significant health data and deliver comments to doctors and patients [3]. Enabling everyone to assess their health to seek emergency treatment in the case of any unconditional emergency can save a person's life. In the long term, the adoption of such process moves may save the government money on medical expenditures [4]. Integrating cellular data with a healthcare system based on an open-source Android architecture has become fairly straightforward as a result of the high accessibility of mobile internet access [5].

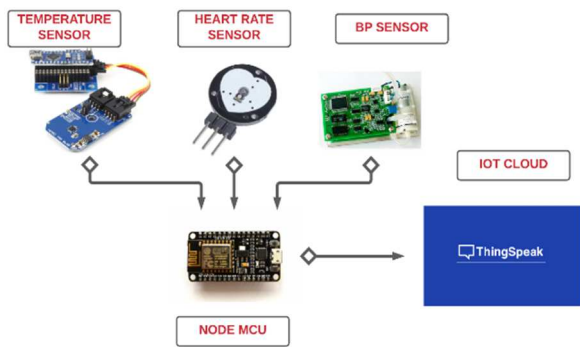


Fig. 2. Block diagram

The implementation of IoT in health monitoring systems has provided us with significant growth in the improvement of contemporary medical treatment [6]. Sensors have gotten smaller as a result of advancements in VLSI technology, allowing for the creation of wearable solutions. Devices are getting more effective and powerful as a result of persistent internet access. IoT-based health monitoring gadgets keep an eye on a patient 24 hours a day, seven days a week. The gadgets create essential signals by assessing statistical data at every critical point [7]. Patients may be remotely monitored and appropriate actions are taken in the event of an emergency since IoT-based gadgets are always linked to the internet. Health tracking systems and IoT-based health tracking systems have considerable distinctions. IoT integration in health monitoring systems is a difficult challenge [8]. The following are some of the difficulties: The majority of IoT efforts have yet to be deployed successfully. The Internet of Things produces a massive amount of data, which necessitates the use of specialized big data for proper management. For IoT systems, security is a major concern. In the case of flawed or obsolete security systems, hackers might readily get sensitive private data from users. Because it is not up to date with current security measures, obsolete infrastructure might cause difficulties. Fig. 2 shows the overall block diagram of the proposed idea.

## II. LITERATURE SURVEY

Pang and his team demonstrated a monitoring approach using an Android smart phone. The system may capture and process data on the server and send it to the smart phone terminal through a socket, allowing it to monitor the target location from anywhere within the range of a wireless network. Sensor data is collected from temperature, humidity, infrared, and CO2 sensors. A pervasive health system [9] using mobile phones was created to help chronic patients manage their diseases while out and about. This system includes patient health monitoring, diaries for various illnesses or symptoms, and social sharing of recorded information. Analyzes IoT-based patient monitoring. Chronically ill patients can benefit from an IoT-based patient monitoring system [10]. This method's goal is to improve patients' quality of life by giving them more control over their diet and exercise habits. Instructions to obtain measurements through sensors and recommendations and exercises are equally effective in improving patients' eating habits as the created model for the system [11]. Current human health monitoring systems have some flaws, such as a rising number of users and submitted data, no user assurance, poor real-time performance, and low data usage. This project develops an IoT-based health monitoring system [12]. The device can

continually monitor heart rate, blood pressure, pulse, body temperature, and other vital indications. In this project, wireless sensors save data for health monitoring. The data is integrated using the Internet of Things to enable real-time monitoring (IoT). The proposed approach improved the current health monitoring platform's accuracy and stability [13-14]. The author in suggests a smart monitoring system based on IoMT. This study describes a full health paradigm that includes a remote health-monitoring system. It gathers critical patient information and sends it to the physicians. This technique saves time for both the patients and the doctors. In the event of an emergency, the physicians will be able to assist the patient right away [15]. Many scholarly publications discuss IoT-enabled versus non-enabled health monitoring systems, both in terms of functionality and cost. R. Ali Khan et al. explain the system design of a wireless body area sensor network for health monitoring purposes [16-17]. Authors in proposed a mobile phone accessory-based wireless health monitoring system [18]. The patient's heart rate is communicated utilizing a smart phone with this method. It is made up of a single microcontroller chip. This type is the least but it has the benefit of extending the battery life.

## III. PROPOSED SYSTEM

A smart health assessment system checks body temperature, blood pressure, and heart rate using a variety of sensors, with the values being relayed using an ESP8266 with the support of an IoT platform to provide a comprehensive picture of one's health.

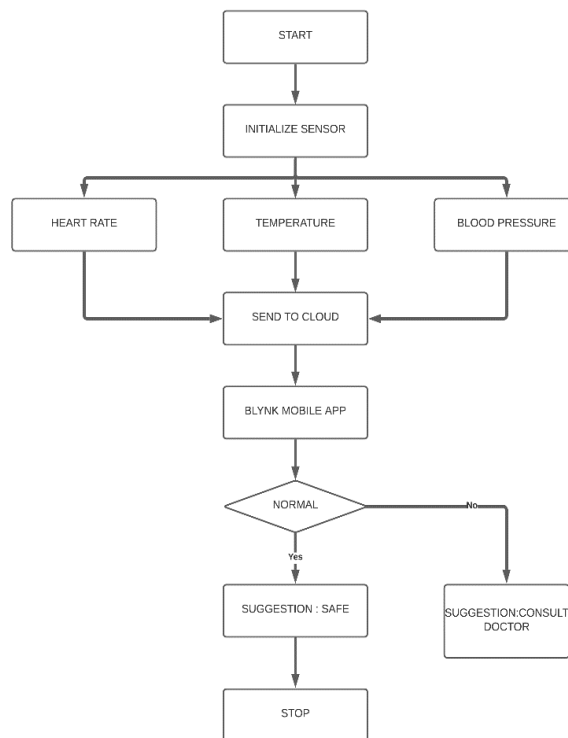


Fig. 3. Proposed Idea

The proposed concept is shown in Fig. 3. With a smart patient health monitoring system, the most fundamental physical parameters of the patient are measured and sent to the cloud for further analysis. A system for tracking the measured data in real time. Any disparities between the observed values and those established by the doctors as the standard of care will be communicated to the patients and their loved ones. A

direct effect of this is that the patient will be able to obtain prompt first aid without having to wait for or engage with the doctors personally.

The suggested system adds additional sensors to the physical layer, resulting in a significant shift in the healthcare paradigm, and analysis. Sensors are attached to the patient's body in various locations and wirelessly linked to the microcontroller. Heart Beat Sensor: Pulse waves, or changes in the volume of a blood artery caused by the heart pumping blood, are measured by an optical heart rate sensor. The range of heart rate for humans is shown in the table: 1. An optical sensor and a green LED are used to detect pulse waves by monitoring volume changes. To mitigate the influences of ambient light such as visible and infrared rays, the detector block utilizes an optic filter calibrated for pulse wave recognition. This facilitates the gathering of high-quality pulse impulses even in wide spaces. Furthermore, by leveraging optic sensor systems that had been developed over a long period, ROHM was capable of increasing the sensitivities of the sensing panel.

TABLE I. HEART BEAT SENSOR

Sl. No.	Age	Heart Rate
1	Less than 1 month	120-60
2	1-12 months	80-140
3	1-2 years	80-130
4	2-6 years	75-120
5	6-12 years	75-110
6	>12 years	60-100

#### A. Temperature Sensor

A temperature sensor is a device that employs an electrical signal to produce temperature measurement in a readable form, often a thermocouple or a resistance temperature detector. A thermometer is the most standard form of the temperature gauge, and it is used to detect exactly cold or hot something that is. Temperature meters are used in the geotechnical profession to monitor structural changes induced by seasonal oscillations in concrete, buildings, soil, water, bridges, and other structures. A temperature sensor is a gadget that senses an item's degree of warmth or coolness. How effectively a temperature meter works is determined by the voltage across the diode. The resistivity of the diodes varies in direct proportion to the temperature change. Whenever the temp drops, so does the resistance, and vice versa.

#### B. Pressure Sensor

The Blood Pressure Sensor is a non-invasive blood pressure monitor for people. Systolic, diastolic, and mean arterial pressure are all measured using the oscillometric approach. The pulse rate is also properly considered. Blood pressure is one of the most important essential indicators (BP). Moving blood exerts pressure on the vascular walls. Blood pressure is defined as the ratio of systolic to diastolic pressure. A mercury sphygmomanometer is used to check blood pressure. In this procedure, blood pressure is calculated using the height of the mercury column. The oscillometric method has been used for automated blood pressure monitoring since 1981. As technology progresses, devices for non-invasive oscillometric blood pressure monitoring are being developed. As technology progresses, devices for non-invasive oscillometric blood pressure monitoring are being developed. Such a gadget is the Blood Pressure Sensor: The diastolic and systolic pressure is monitored via oscillometric sensing and the disparity capacitive or variable piezo resistance concept,

respectively. Fig. 4 depicts a normal human Systolic vs. Diastolic blood pressure chart which is taken from [bloodpressureuk.org](http://bloodpressureuk.org).

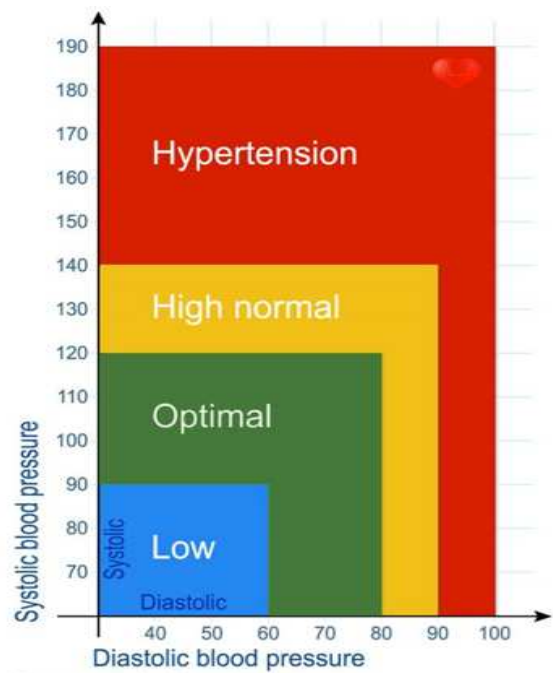


Fig. 4. Blood pressure range.

#### C. Node-MCU

Node MCU is an open-source IoT platform that includes both software and hardware. It performs the function of as like as the microcontrollers like Arduino and many. It is based on the ESP8266 microcontroller. It may be connected to the computer by USB connection, and programs can be installed onto it. A Wi-Fi module is included. The Node MCU is linked to a variety of health monitoring sensors, and data is gathered. These data are transferred to the cloud with the aid of the Wi-Fi module. It may be viewed from anywhere using e-mail from the cloud.

### IV. RESULTS AND DISCUSSION

Sensors in the proposed system measure and communicate all body variables in real time to the cloud, which is powered by an Internet of Things platform. It displays the heart rate at any given point in time during the day. Wearers of wearable devices can enjoy more comfort as a result of this feature. This method captures data that is more exact and accurate. If you compare it to the manual techniques, it saves you valuable time. Because data is collected on a constant basis, the burden on doctors is reduced as a result of these statistics. It is possible that the other specialist will also receive these numbers by email. The health status of a patient, for example, can be recognized and compared to established parameters defined by his or her family, and major health alarms can be triggered depending on the detected data. The temperature from the temperature sensor is monitored regularly, and the data in the Blynk app are updated regularly. The temperature measured graph is shown in Fig. 5, Fig. 6. The wearable blood pressure sensor measures both systolic and diastolic blood pressure. The systolic blood pressure is shown in Fig. 7, Fig. 8 and the diastolic blood pressure is shown in Fig. 9, Fig. 10. The heartbeat sensor is used to determine the heart rate. The heartbeat rate graph of the patient is shown in Fig. 11, Fig. 12.

Application software is designed to show the output status of the patient's body condition. Through this, uneducated people could also get to know their health condition with the help of a sensor is attached to the body. This particular sensor measures four important parameters. The detectable four parameters incorporated here are temperature, systolic, diastolic, and heart rate. As shown in Fig. 5 and Fig. 6. the temperature monitor is abnormal and normal respectively. Normally, a human's average temperature would be 98.6 degrees Fahrenheit. This tested output denotes us through the alert symbol accompanied with an alert beep. If the temperature is above the normal state, then the alert symbol and a beep sound will get initiated. From this Doctor, a visit has to be done by the patient for checking if any abnormality is noticed. The graph is depicted for legible understanding to uneducated people at the moment after testing. In Fig. 5, since the temperature exceeds the normal temperature, an alert beep is noticed. In 15 minutes, 1 hour, 6 hours, 1 day the temperature is noticed as 130 degrees, 120 degrees, 125 degrees, 124 degrees respectively. In Fig. 6, since the temperature doesn't exceed the normal temperature, an alert beep is not noticed. So that the patient could take self-care by him or herself. In the output, one of the patient's outputs is 140 degrees, so the alert is shown due to its abnormality. While another one is 101 degrees, so the alert is not shown due to its normality.

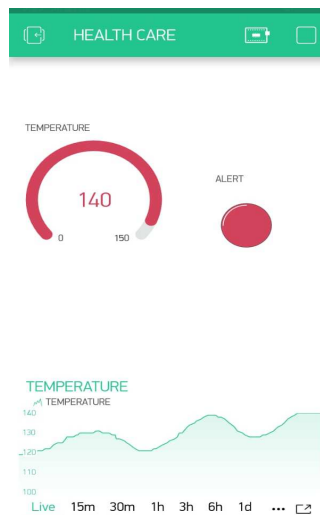


Fig. 5. Temperature monitor - abnormal

As shown in Fig. 7 and Fig. 8. the systolic monitor is abnormal and normal respectively. Normally, a human's average systolic pressure would be 80 to 120 mm Hg. This tested output denotes us through the alert symbol accompanied with an alert beep. If the systolic pressure is above the normal state, then the alert symbol and a beep sound will get initiated. From this Doctor, the visit has to be done by the patient for checking, if any abnormality is noticed. The graph is depicted for legible understanding to uneducated people at the moment after testing. In Fig. 7, since the systolic pressure exceeds the normal state, the alert beep is noticed. In 15 minutes, 1 hour, 6 hours, 1 day the systolic pressure is noticed as 120 mm Hg, 135 mm Hg, 140 mm Hg, 132 mm Hg respectively. In Fig. 8 since the systolic pressure doesn't exceed the normal state, an alert beep is not noticed. So that the patient could take self-care by him or herself without visiting the doctor. In the output, one of the patient's outputs is 134 mm Hg, so the alert is shown due to its abnormality.

While another one is 120 mm Hg, so the alert is not shown due to its normality.

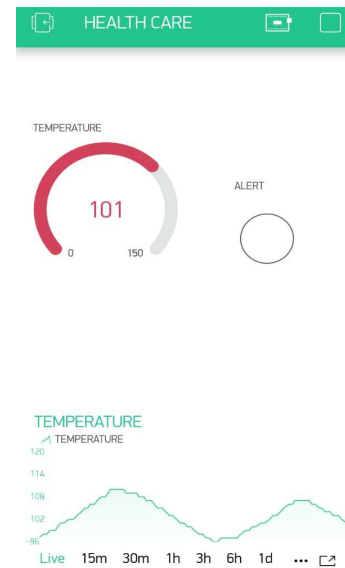


Fig. 6. Temperature monitor - normal



Fig. 7. Systolic monitor. – abnormal

As shown in Fig. 9 and Fig. 10. the diastolic monitor is abnormal and normal respectively. Normally, a human's average diastolic pressure would be 60 to 80 mm Hg. This tested output denotes us through the alert symbol accompanied with an alert beep. If the diastolic pressure is above the normal state, then the alert symbol and a beep sound will get initiated. From this Doctor, the visit has to be done by the patient for checking, if any abnormality is noticed. The graph is depicted for legible understanding to uneducated people at the moment after testing. In Fig. 9, since the diastolic pressure exceeds the normal state, an alert beep is noticed. In 15 minutes, 1 hour, 6 hours, 1 day the diastolic pressure is noticed as 82 mm Hg, 78 mm Hg, 90 mm Hg, 92 mm Hg respectively. In Fig. 10, since the diastolic pressure doesn't exceed the normal state, an alert beep is not noticed. So that the patient could take a self-care by him or herself without visiting the doctor. In the output, one of the patient's outputs is 99 mm Hg, so the alert is shown due to its abnormality. While another one is 73 mm Hg, so the alert is not shown due to its normality.



Fig. 8. Systolic monitor. - normal

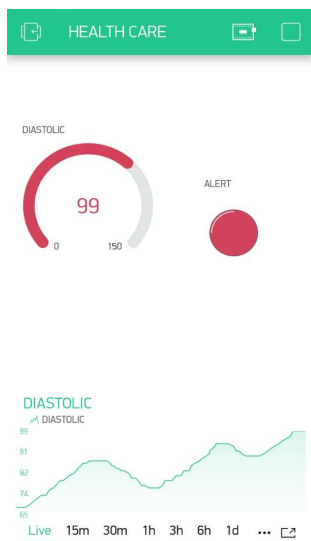


Fig. 9. Diastolic monitor. - abnormal



Fig. 10. Diastolic monitor. - normal

As shown in Fig. 11 and Fig. 12. the Heart rate monitor is abnormal and normal respectively. Normally, a human's average Heart rate would be 60 to 100 beats per minute. This tested output denotes us through the alert symbol

accompanied with an alert beep. If the diastolic pressure is above the normal state, then the alert symbol and a beep sound will get initiated. From this Doctor, the visit has to be done by the patient for checking, if any abnormality is noticed. The graph is depicted for legible understanding to uneducated people at the moment after testing. In Fig. 11, since the heart rate exceeds the normal state, an alert beep is noticed. In 15 minutes, 1 hour, 6 hours, 1 day the heart rate is noticed as 78 beats per minute, 82 beats per minute, 97 beats per minute, 95 beats per minute respectively. In Fig. 12, since the heart rate doesn't exceed the normal state, an alert beep is not noticed. So that the patient could take self-care by him or herself without visiting the doctor. In the output, one of the patient's output is 98 beats per minute, so the alert is shown due to its abnormality. While another one is 68 beats per minute, so the alert is not shown due to its normality.

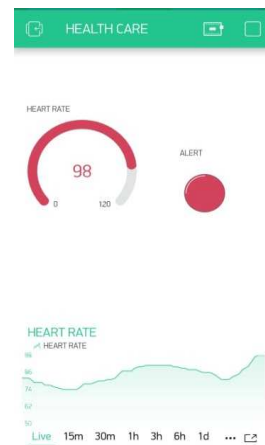


Fig. 11. Heart rate monitor abnormal

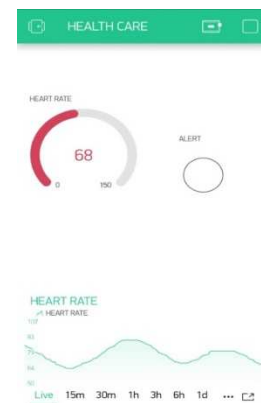


Fig. 12. Heart rate monitor - normal

The sensor's data is fed to the controller, which subsequently sends it to the IoT device. These values are uploaded in the Blynk app so that we can able to monitor them easily. The usual body temperature for a healthy person is between 97- and 99-degrees Fahrenheit. A healthy person's heart rate ranges from 60 to 100 beats per minute. The human body's typical systolic and diastolic rates are fewer than 120 and 80, respectively. If any of these numbers are abnormal, the controller will notify the patients, their families, and their physicians, who will administer emergency first aid.

## V. CONCLUSION

A low-cost, simple-to-implement health monitoring system is provided, which solves the shortcomings of the existing method of measuring the health parameters by

making it a more effective technique of monitoring patients' health parameters. It offers the advantages of being less expensive, taking less time, and using less power. The accurate measurement of a patient's heart rate and other health data is achievable, and it plays an important part in the medical monitoring system. IoT allows sensors to relay data wirelessly to a server. Basic physical characteristics are monitored regularly. It is more comfortable for the patient to utilize these wearable gadgets. This approach records values that are more exact and accurate. It saves time compared to the manual way. The doctors' workload is decreased since these values are continually recorded. In the future, machine learning and artificial intelligence algorithms will be used to analyze a component of the design. Automated diagnosis, prescription, and report analysis the system will be supplemented by the generator.

#### REFERENCES

- [1] Riazul Islam et.al. "The internet of things for health care: a comprehensive survey", *IEEE Access*, 3, 678-708 (2015)
- [2] J. Liu, N. Kato, J. Ma, and N. Kadowaki, "Device-to-Device Communication in LTE-Advanced Networks: A Survey", *IEEE Communications Surveys, and Tutorials*, 17(4), 1923-1940 (2015)
- [3] Maneesha V Ramesh et. Al, "Mobile software for health professionals to monitor remote patients. Ninth International Conference on Wireless and Optical Communications Networks", (WOCN), 13075177 (2012)
- [4] Moser, L.E., Melliar-Smith, P.M, "Personal health monitoring using a smartphone". 2015 IEEE International Conference on Mobile Services, pp. 344-351 (2015)
- [5] X. Kong, B. Fan, W. Nie, Y. Ding, "Design on mobile health service system based on Android platform". 2016 IEEE Advanced Information Management, Communicates, Electronic and Automation Control Conference (IMCEC), pp. 1683-1687(2016)
- [6] V. Tripathi, F. Shakeel, "Monitoring health care system using internet of things-an immaculate pairing", 2017 International Conference on Next Generation Computing and Information Systems (ICNGCIS), pp. 153-158 (2017)
- [7] C. Raj, C. Jain, W. Arif "Health monitoring and nous: An IoT-based e-health care system for remote telemedicine", 2017 International Conference on Wireless Communications, Signal Processing and Networking (WiSPNET), pp. 2115-2119 (2017)
- [8] G. Yang, K.Ovsthus, "The challenges of the IoT solutions in a home care project", International Conference on Computational Science and Computational Intelligence (CSCI), pp. 1771-1774 (2017)
- [9] K. Triantafyllidis, V. G. Koutkias, I. Chouvarda, et al, "A pervasive health system integrating patient monitoring, status logging, and social sharing", *IEEE Journal of Biomedical and Health Informatics*, 17(1), 30-37 (2013)
- [10] S. Gupta, G. Lipika, K.A. Abhay, " A novel framework of health monitoring systems", *International Journal of Big Data and Analytics in Healthcare (IJBDAH)*; 6.1,1-14 (2021)
- [11] M. Al Yami and D. Schaefer, "Fog computing as a complementary approach to cloud computing", International Conference on Computer and Information Sciences (ICCIS), pp. 1-5, Sakaka, Saudi Arabia (2019)
- [12] Jorge Gomez, Byron Oviedo, Emilio Zhuma, "Patient Monitoring System Based on Internet of Things", *Procedia Comput. Sci.* 90-97,83 (2016)
- [13] Rahat Ali Khan and Al-Sakib Khan Pathan,"The state-of-the-art wireless body area sensor networks: A survey", *International Journal of Distributed Sensor Networks*, 14(4), (2018)
- [14] Shaad Mahmud, Honggang Wang, "A Wireless health monitoring system using mobile phone accessories", *IEEE Internet Things J.* 4 (6), (2017).
- [15] M Ramkumar, C Ganesh Babu, A Manjunathan, S Udhayanan, M Mathankumar,"A Graphical User Interface Based Heart Rate Monitoring Process and Detection of PQRST Peaks from ECG Signal" *Lecture Notes in Networks and Systems*, 173, LNNS, pp. 481-496 (2021)
- [16] M Ramkumar, C Ganesh Babu, K Vinoth Kumar, D Hepsiba, A Manjunathan, R Sarath Kumar, "ECG Cardiac arrhythmias Classification using DWT, ICA and MLP Neural Network", *Journal of Physics: Conference Series*, 1831, issue.1, pp.01201 (2021)
- [17] C. Bhuvaneshwari, A. Manjunathan,"Advanced gesture recognition system using long-term recurrent convolution network", *Materials Today Proceedings*, 21, 731-733 (2020)
- [18] C Bhuvaneshwari, A Manjunathan, "Reimbursement of sensor nodes and path optimization", *Materials Today: Proceedings*, 45, 1547-1551 (2021)

# Analyzing the Occurrence of Stroke using Machine Learning-A comparative Study on Supervised Learning Models

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**Abstract**—Stroke is blood coagulation or bleed in the brain which could cause permanent damage and affects mobility, intelligence, vision, and communication. Stroke is considered a health-related crisis circumstance and can cause long-term neurological damage, complication, and often death. Most of the stroke is classified as ischemic and haemorrhagic. Stroke has been observed to have abnormal ECG signals. Therefore, if the individuals have their bio-signals monitored in real-time, they can get proper treatment rapidly. Most stroke diagnosis forecast systems are based on image processing tools namely CT and MRI, which are costly and hard to use in clinical practices. Stroke is the consequent driving cause for death around the world and quite possibly the most dangerous infection for persons over the age of 65. It causes ill-effects to the cerebrum like "coronary failure" which causes ill-effects to the heart. When a stroke sickness occurs, it causes enormous clinical care and permanent disability, yet in some cases, it also results in death. Like clockwork, someone dies of a stroke every 4 minutes, yet up to 80% of stroke could be averted if the medical specialist could forecast the incidence of stroke in initial phases. In this project, we have designed an ML model for predicting stroke utilizing the KNN algorithm, SVM algorithm, and NB algorithm. Followed by comparing KNN, SVM, and NB algorithms utilizing error and accuracy. Among KNN, SVM, and NB algorithms, the Support Vector Machine algorithm has the highest accuracy of 96.66%. This project hence helps in predicting the stroke effects and provides a customized warning. Therefore, it urges clinical patients to fortify the inspiration of wellbeing prosperity and brief changes in their medical care practices.

**Keywords**—ECG signal, Stroke, MRI, CT, Machine Learning.

## I. INTRODUCTION

The first paragraph after a heading is not indented (Body text style). The World Health Organization(WHO) interpreted Stroke—"Fast-growing clinical indications of severe disruption of brain activity, having indications persisting 24 hours or greater or resulting in death, without reason apart from vascular origin"[1]. A stroke occurs when the blood dispense to some portion of the brain is hindered, preventing cerebrum tissue from obtaining oxygen and nutrients. A stroke is a health crisis and prompt treatment is vital. Early treatment can lessen the brain damage and other consequences. Stroke is one of the main dangers to general wellbeing throughout the world [15].

With the enhancements in medical services, more individuals endure stroke yet many need to adapt to the physical, mental, social environment, bringing about expanded individual and public expenses. The socio-economic effect of stroke is extensive around the world [16].

It is estimated that 4.5 million people die of stroke a year around the world and more than 9 million stroke survivors. Nearly one of every four males and almost one of five females over the age of 45 can expect to have the illeffects of stroke in case they live to the age of 85. Recurrence of stroke expands the level of inability and demise of patients [14].

Stroke is the leading cause of death in the United States, as per the Centers for Disease Control and Prevention. It is a non-communicable infection that accounts for more than 11% of all deaths worldwide [12]. Constantly, more than 795,000 persons in the US have experienced the health impacts of stroke. It is the fourth remarkable inducement for demise in India [2]. The worldwide lifetime threat for stroke in 2016 for people beginning at the age of 25 was 24.9%. The most noteworthy worldwide lifetime threat for stroke was identified in East Asia (38.8%) followed by Europe (31.7%). The most reduced stroke threat was identified in eastern sub-Saharan Africa with a percentage of 11.8.

With the enhancement of innovation in the healthcare sector, foreseeing the circumstance of stroke can be modeled by utilizing ML [11]. The algorithms in ML are valuable in creating precise predictions and providing the right examination. The obstruction with this model is that it is being prepared on literary information and not on clinical cerebrum scans [10], [18].

The dataset is selected from Kaggle with different characteristics as its features. Initially, the dataset is Pre-processed (i.e.) cleaning and preparing for the ML model to recognize. Following that Label Encoding is executed to transform the string to integer data type. Then the dataset is parted as learn and validation data. Subsequently, a model is created utilizing this new data through three supervised ML techniques. Accuracy is determined for these various techniques and analyzed to become the finest prepared model for the forecast. Succeeding legitimate examination, the paper concludes with the most appropriate algorithm for stroke analysis.

## II. LITERATURE SURVEY

To get the required knowledge about different ideas related to the stroke analysis, existing literature was taken into account. The research paper on “Computer methods and programs in Biomedicine, provides the computational analysis of stroke prediction over 10 years and classifies the patient’s stroke probability occurrence into five different categorizations [3]. The research paper [22] determines a model condition for stroke pre-analysis algorithm with dormant modifiable ill effects. The paper [23] utilizes a Decision tree algorithm to extract the attributes, a principal component analysis algorithm to lessen the dimension and the adopted back propagation neural network classification algorithm is used to establish a classification model. In the paper [24] principle component analysis algorithm is used for decreasing the dimensions and picks up the most involved features for stroke prediction. The research paper [4] suggests the forecast of thromboembolic stroke disorder by utilizing Artificial Neural Networks. Though this model has an accuracy of over 85%, neural Networks require higher processing time and to train. The research work [21] utilized the random forest algorithm to predict the ill effects of the patient suffering from a stroke. This specific work cannot be utilized on any further new stroke types later on. The article [25] provides the application interface for clinical information perception and the board for nervous system specialists in stroke grouping and expectation framework called Stroke MD. The research work [26] proposes the utilization of three various calculations to foresee the chance of stroke and inferred that the Decision tree has the most noteworthy exactness (about 75%). In any case, this model couldn't suit this present reality model.

Other paragraphs are indented (Body text Indented style). According to the journal article [27], the algorithm for stroke detection was developed using Random Forest, Decision Tree, and Multilayer Perceptron. The accuracy rates achieved for the 3 strategies were fairly similar, with just minor variations. Decision Tree had a measured efficiency of 74.31 percent, Random Forest had a determined efficiency of 74.53 percent, and multi-layer perceptron had a determined efficiency of 75.02 percent. [28] Demonstrates the application of a supervised learning method to evaluate cardiac stroke. They built the model using several machine learning approaches such as Naive Bayes, Decision Tree, and SVM and then compared their results. The methods they utilized had the highest accuracy of 60%, which is rather low. The researchers of [29] utilized several data categorization algorithms to forecast the stroke risk. The dataset was obtained first from the Region of Saudi Arabia's Hospitals. C4.5, Jrip, and MLP were the three classifiers' methods utilized (MLP). The algorithm achieved an accuracy of about 95% using these techniques. Despite the article's claim of 95 percent accuracy, the duration required for training and prediction is longer since the scientists utilized a mix of sophisticated algorithms.

According to a study published in [30], three distinct algorithms may be used to forecast the risk of a stroke. Decision Trees, Nave Bayes, and Artificial Neural networks are implemented. This study revealed the Decision tree had a maximum accuracy of 75 percent. However, depending on the outcomes derived from the confusion matrix, this concept could not be applied to real-world instances. The investigators in [31] used the Cardiovascular Study database to forecast strokes. They developed a unique

automated feature selection method that picks resilient feature extraction methods on a cautious mean that presented. For more effectiveness, they coupled this technique with the SVM Classifier. However, this enabled the production of a lot of variables that gradually decrease the effectiveness of the algorithm. The article [31] suggests employing Neural Networks to forecast stroke illness. The Back-propagation technique was employed for estimation. This model obtained an accuracy of about 89 percent. However, due to the complicated design and growing neurons, NN consumes a long time to educate and even more processing time.

The article can be interpreted in the following manner: Section II goes into detail about the literature review. Section III describes the research's purpose. Section IV describes the study materials and methods. Section V explains in detail how the three machine learning techniques KNN, SVM, and NB, works. Section VI goes through the project's execution. Section VII examines the results in terms of accuracy, precision, and recall. Section VIII suggests the best approach for predicting strokes.

## III. OBJECTIVE OF THIS RESEARCH

Stroke stays as significant wellbeing trouble both for the people and for the public medical care system [17]. The objective of this study is to apply the fundamentals of ML on huge data to viably analyze the occurrence of stroke dependent on conceivably modifiable risk factors.

## IV. MATERIALS AND METHODOLOGY

This section briefly explains the Dataset, Algorithms, and Train and test datasets.

### A. Dataset Description

The dataset is collected from Kaggle and it is the documentation of around 5000 people’s information on their health status. The dataset contains 10 attributes and the labelled response.

- 1) *Age*: This attribute is numerical data and it is the age of the patient.
- 2) *Gender*: It is categorical data about the patient’s gender.
- 3) *Hypertension*: It is numerical data about whether the patient is hypertensive or not.
- 4) *heart\_disease*: It is numerical data on whether the patient had any previous cardiac issues.
- 5) *ever\_married*: It is categorical data about the patient’s marital status.
- 6) *work\_type*: It is categorical data on a patient’s employment status.
- 7) *residence\_type*: This is categorical data on the patient’s occupancy scenario.
- 8) *avg\_glucose\_level*: This is numerical data on the patient’s body glucose level.
- 9) *BMI*: It is the ratio of mass to the square of the person’s height and it is numerical data.
- 10) *smoking\_status*: It is categorical data on whether the patient smokes or not and how often he/she smokes.
- 11) *Stroke*: It is the numerical data on the stroke attacks. This is the decision:making attribute.

### B. Algorithm

The occurrence of stroke can be predicted by employing three Machine Learning algorithms namely (1) Support Vector Machine (SVM) (2) K Nearest Neighbour (KNN), (3) Naïve Bayes (NB) algorithm. The motivation behind using these algorithms is that these algorithms are very notable in modeling accurate predictors and have been utilized in few comparative research works. These models are assessed by comparing the confusion matrices [6].

### C. Data Splitting

Parting the data into various sets is a procedure usually utilized in ML. The data is generally parted into a learning and validation dataset to prepare and track down the model parameters and to evaluate the model prediction with the parameters such as accuracy or error [5]. The training dataset was utilized to fit the model and testing dataset for the evaluation.

## V. PREDICTIVE TECHNOLOGY

For predicting the stroke based on the given attributes three machine learning algorithms are used includes KNN, SVM, NB. The KNN is the laziest algorithm that predicts the stroke occurrence based on the nearest sample. The SVM predicts the stroke based on the hyper plane drawn in the plotted data. Finally, the Naïve Bayes predict the stroke by the Bayesian formula. All three algorithms used in this work are detailed with a flow chart in the upcoming sections.

### A. K Nearest Neighbor Algorithm

KNN is the least complex ML algorithm dependent on supervised learning. The KNN was first proposed by Cover and Hart in 1968 [7]. KNN algorithm deduces the analogy between new data with available datasets and places the new data into the most suitable category. KNN algorithm is commonly used in Classification problems and for Regression problems.

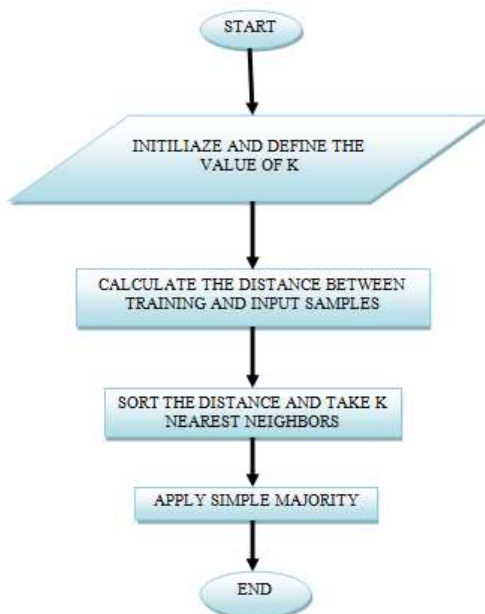


Fig. 1. Block diagram of KNN algorithm

KNN algorithm is shown in the Fig. 1, which is a non-parametric algorithm as it doesn't make any presumption on the available data. The K value is chosen to reduce the training

and validation error. The K value is the number of neighbour data available for classification. Predominantly, an odd number is chosen as the K: value. KNN algorithm has the advantage of being simple and less complex. There's no compelling reason to build a model, tune parameters and make extra assumptions. The disadvantage of the KNN algorithm is it comparatively gets slower as the number of independent variables increases. In the Fig. 2, the classification of KNN algorithm is explained.

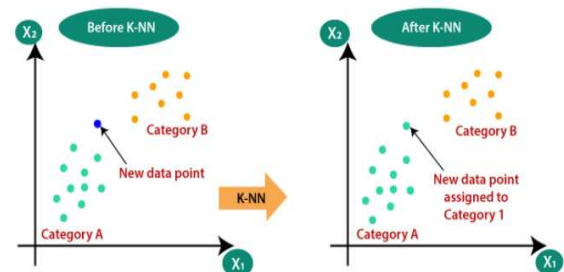


Fig. 2. Classification of KNN algorithm.

(Source: <https://www.javatpoint.com/k-nearest-neighbor-algorithm-for-machine-learning>).

### B. Support Vector Machine Algorithm

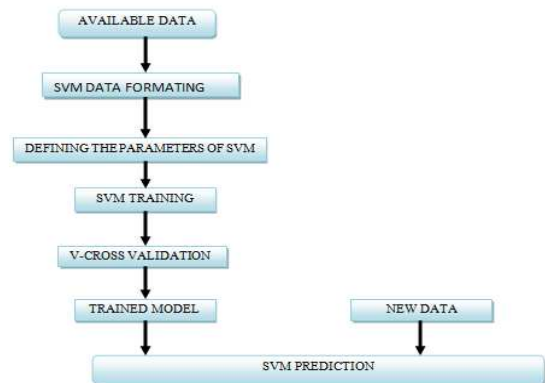


Fig. 3. Block diagram of SVM algorithm

Fig. 3 describes the SVM which is an ML algorithm based on Supervised Learning and is most commonly used for Classification and Regression problems. In 1963, the SVM algorithm was initially developed by Vladimir N. Vapnik and Alexey Ya. Chervonenkis. The target data is plotted in the n dimensional plane, where 'n' indicates the number of attributes available in the dataset [8]. After plotting, the hyper plane is being drawn. The ideal hyperplane will be the one with the greatest margin, since bigger margin guarantees, slight deviations in the data points will not influence the result of the model. The performance of SVM completely depends on the kernel selection. Yet, the ideal selection of kernel for some problems is not found. The advantage is that it functions admirably with a clear margin of partition. It is viable in high level dimensional planes furthermore in situations in which the quantity of measurement is more than samples. Fig. 4, shows the classification of SVM Algorithm. The disadvantage is that it doesn't perform well when the data has more noise and with an enormous dataset since the training time is more.

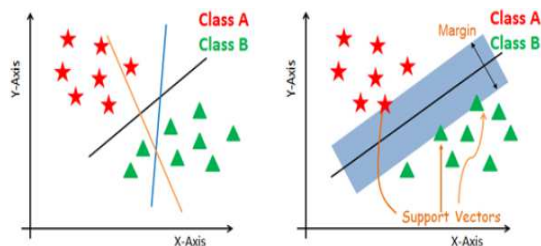


Fig. 4. Classification of SVM algorithm.

(Source:

<https://heartbeat.fritz.ai/understanding-the-mathematics-behind-support-vector-machines-5e20243d64d5>).

### C. Naïve Bayes Algorithm

Naïve Bayes based on Bayes theorem is an ML algorithm dependent on supervised learning and most commonly utilized for classification problems. It is mostly utilized in text characterization that incorporates a high dimensional training dataset. It helps in building quick ML models that can make a speedy prediction. It is a probabilistic classifier. The advantage of the NB algorithm is that it is one of the less complex and quick algorithms which is used for text classification [9]. It executes well in multi class predictions than other algorithms. The disadvantage is that it cannot learn the relationship between features as it assumes all features are independent. The types of NB algorithm are Gaussian, Multinomial and Bernoulli. It's a gathering of computations that contains an average guideline as no attribute depends on the other. NB is shown in the Fig. 5. Bayes theorem is expressed as (1):

$$P\left(\frac{A}{B}\right) = \frac{P\left(\frac{B}{A}\right)P(A)}{P(B)} \quad (1)$$

Where,  $P(A) \rightarrow$  Prior probability,  $P(B) \rightarrow$  Marginal probability,  $P(A/B) \rightarrow$  Posterior probability,  $P(B/A) \rightarrow$  Likelihood probability.

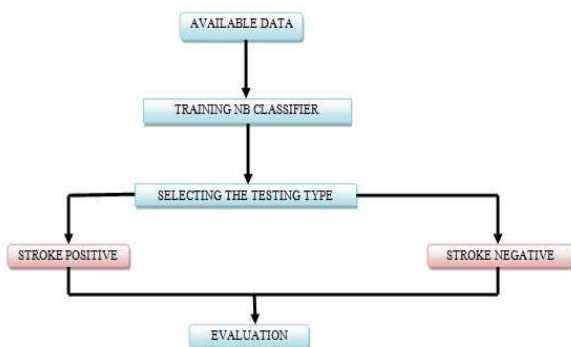


Fig. 5. Block diagram of NB algorithm

## VI. IMPLEMENTATION OF THE PROJECT

To complete this project, basic Python programming, Matplotlib, and Sklearn libraries are employed. The block diagram is depicted in Fig. 6

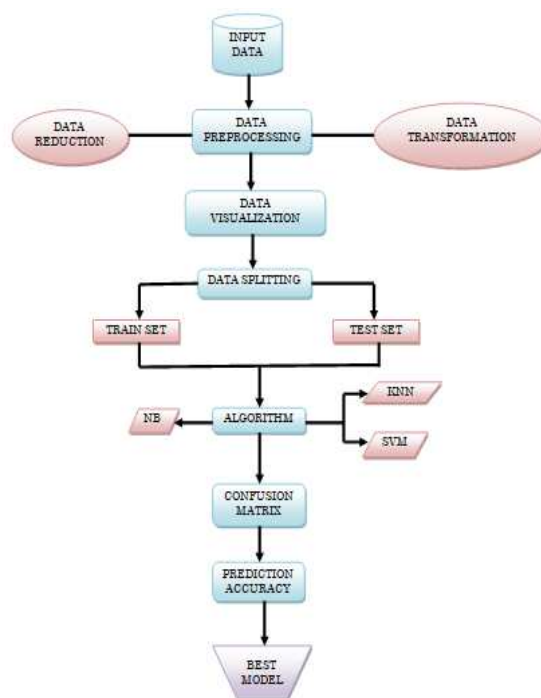


Fig. 6. Block Diagram to analyze the stroke prediction

1) *Input Data*: The data of around 5000 people have been collected from Kaggle. It contains risk factors like a lifetime, gender, hypertension, cardiac issue, marital status, employment type, habitation, Blood sugar level, BMI, and Smoking status.

2) *Data Pre:processing*: Data pre:processing is done to check the duplicate and missing values in the dataset. Followed by Label encoding, a type of data transformation is performed to convert categorical data to numerical data in the columns 'marital status', 'employment type', 'residence type', and 'smoking status. Data reduction is performed to balance the stroke positive and negative cases. Finally, the standardized data is acquired to proceed with further ML processing.

3) *Data Visualization*: Data visualization is performed to obtain a clear view of the dataset. Data visualization is the graphical representation of data.

4) *Data Splitting*: The final dataset is separated into features and labels and further into training and testing dataset in the ratio of 7:3 using Scikit learn library.

5) *Algorithm*: Three algorithms namely KNN, SVM, and NB have been utilized to train the set and to validate the dataset to predict the most appropriate model.

6) *Confusion matrix*: Confusion matrix is employed to estimate the value of Accuracy, Precision, and Recall.

7) *Best model*: The level of accuracy of the three ML models is identified and the best model for stroke prediction is obtained.

## VII. RESULT AND DISCUSSION

The dataset which is collected from Kaggle contains 10 attributes and around 5000 samples. The data is parted into two categories as training dataset and testing dataset in the ratio of 7:3. For analyzing the stroke prediction, three ML algorithms have been utilized namely KNN, SVM, and NB to develop an appropriate model. Confusion matrix elements like

True Negative, False Positive, True Positive, and False Negative have been utilized to analyze the model using performance metrics namely Precision, Recall, and Accuracy.

1) *True Positive (TP)*: The patient is affected by stroke in real:time and the model also predicts the same.

2) *False Positive (FP)*: Though, the patient is not affected by stroke in real:time the model predicts that the patient is affected by stroke.

3) *True Negative (TN)* : The patient is not affected by stroke in real:time the model predicts the same.

4) *False Negative (FN)* : Though, the patient is affected by stroke in real:time the model predicts that the patient was not affected by stroke [13].

5) *Precision*: It is the small portion of suitable illustration among the recovered illustrations [19].

6) *Recall*: Also known as Sensitivity that is the small portion of suitable illustrations that are recovered [20].

7) *Accuracy*: It is used to decide the best model. It is the percentage of correct prediction in the test.

Table I gives the formula for performance metrics employed in the machine learning algorithms like KNN, SVM, and NB. Table II compares the performance analysis of the ML algorithms.

TABLE. I. FORMULA OF PERFORMANCE METRICS

Performance Metrics	Formula
True Positive rate	TP/Actual positive
True Negative rate	TN/Actual negative
False Positive rate	FP/Actual negative
False Negative rate	FN/Actual positive
Precision	TP/(TP+FP)
Recall	TP/(TP+FN)
Accuracy.	(TP+TN)/(TP+TN+FP+FN)

The accuracy of the three algorithms is taken to analyze the best method to predict the stroke. The accuracy of the three ML algorithms is plotted in Fig. 7. The accuracy value of Naïve Bayes is 95.33%, Support Vector Machine is 96.66% and finally, the accuracy of K Nearest Neighbour is 94.0%. From this, it is identified as the accuracy is very high for the Support Vector Machine algorithm and the accuracy value is very less in the K Nearest Neighbour method. For the best model, the accuracy value should be high. The model is designed for predicting stroke. So the model accuracy is very important. From this comparison, the Support Vector Machine is found to be good.

TABLE. II. PERFORMANCE SURVEY OF THE MACHINE LEARNING ALGORITHMS

Algorithm	Accuracy	Precision	Recall
Naïve Bayes	95.3333	93.58974	97.3333
Support Vector Machine	96.66667	97.36842	96.1039
K:Nearest neighbor	94.0	93.50649	94.7368

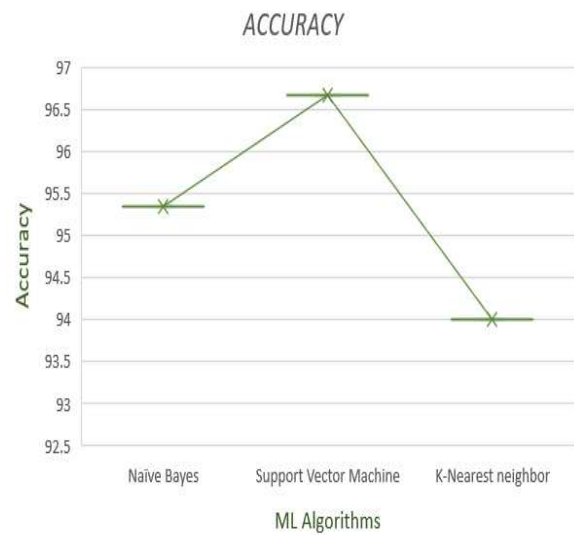


Fig. 7. Comparison of 3 ML algorithms based on accuracy.

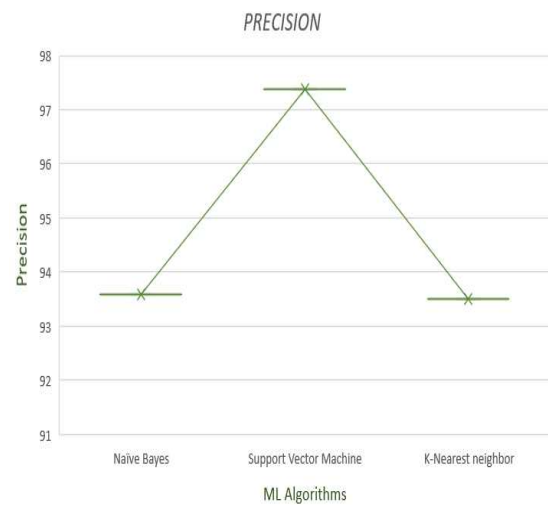


Fig. 8. Comparison of 3 ML algorithms based on precision

The precision of the three algorithms is taken to analyze the best method to predict the stroke. The precision of the three ML algorithms is plotted in Fig. 8. The precision value of Naïve Bayes is 93.58%, Support Vector Machine is 97.36% and finally, the precision of K Nearest Neighbour is 93.50%. From this, it is identified as the precision is very high for the Support Vector Machine algorithm and the precision value is very less in the K Nearest Neighbour method.

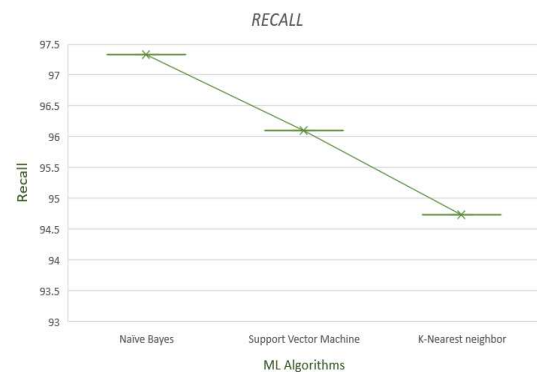


Fig. 9. Comparison of 3 ML algorithms based on recall.

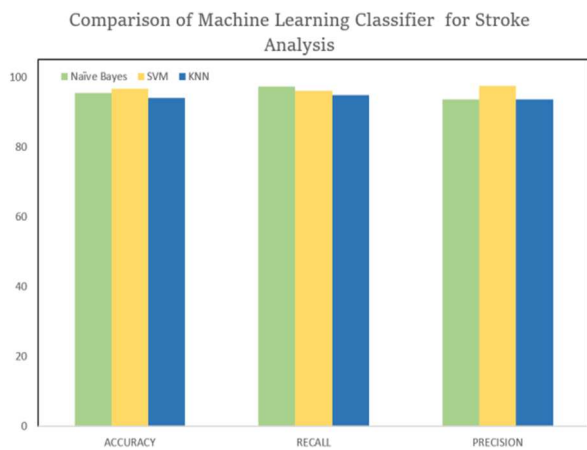


Fig. 10. Bar graph for analyzing the performance of Machine learning Algorithms.

The recall of the three algorithms is taken to analyze the best method to predict the stroke. The recall of three ML algorithms is plotted in Fig. 9. The recall value of Naïve Bayes is 97.33%, Support Vector Machine is 96.10% and finally, the accuracy of K Nearest Neighbour is 94.73%. From this, it is identified as the recall is very high for the Naïve Bayes algorithm and the recall value is very less in the K Nearest Neighbour method. Fig. 10 shows the Bar graph for analyzing the performance of Machine Learning Algorithms.

#### VIII. CONCLUSION

This project can be anticipated to develop a mobile application to give a customized message based on every patient's degree of effects from stroke and a lifestyle remedy message on the hazard factors of stroke. Further advancements in this research can incorporate foreseeing different infections separated from a stroke. This model completely predicts the occurrence of stroke and might have been used in guiding patients and their families for planning clinical preliminaries. This methodology gives a steady advantage beyond current clinical practice that can convert into significant advantages for populace wellbeing and work with the adoption of ML based danger predictors in clinical practice.

#### REFERENCES

[1] Ralph L.Sacco, Scott E Kasner, Joseph P.Broderick, Louis R.Caplan, J.J (Buddy) Connors, Antonio Culebras, 2013 : AHA Journals.,44(7).

[2] Jeyaraj DP and Paulin S. 2013 in India:Journal of stroke15(3).

[3] Gangavarapu Sailasya, Gorli L Aruna Kumari, 2021Int. J. Adv. Comput. Sci. Appl.,12(6).

[4] Minhaz Uddin Emon, Maria Sultana Keya, Tamara Islam Meghla, Md. Mahfujur Rahman, M Shamim Al Mamun, and M Shamim Kaiser., 2021 Conference Paper. DOI: 10.1109/ICECA49313.2020.9297525.

[5] Shichao Zhang, IEEE Trans. Knowl. Data Eng., DOI: 10.1109/TKDE.2021.3049250.

[6] Yongli Zhang,2012 ICICA 2012: Information Computing and Applications 308pp.179:186.

[7] Pouria Kaviani, Sunita Dhotre,2017 IJARCSMS., 4(11).

[8] Soodamani Ashokan, Suriya G.S Narayanan, Mandresh S, Vidhyasagar BS, Paavai Anand G 2020 Int. J. Eng. Res. Technol., 07(03).

[9] Achala Vagal, Heidi Sucharew, Christopher Lindsell, Dawn Kleindorfer, Kathleen Alwell, Charles J. Moomaw, Daniel Woo, Matthew Flaherty, Pooja Khatri, Opeolu Adeoye, Simona Ferioli, Jason Mackey, Sharyl Martini, Felipe De Los Rios La Rosa F., 2018Journal of Behav Brain Sci08(10).

[10] Krishna Mohan, C. and Yegnanarayana, B., 2010. Classification of sport videos using edge-based features and autoassociative neural network models. Signal, Image and Video Processing, 4(1), pp.61-73.

[11] Yangguang Liu, Yangming Zhou, Shiting Wen, Chaogang Tang., 2014 Int. J. Mob. Comput. Multimed. Commun.,6(4).

[12] Ramesh, G.P. and Mohan Kumar, N., 2018. Radiometric analysis of ankle edema via RZF antenna for biomedical applications. Wireless Personal Communications, 102(2), pp.1785-1798.

[13] Hager Ahmed, Sara F. Abd:el ghany, Eman M.G.Youn, Nahla F.Omran, Abdelmgeid A.Ali., 2019Int. J. Adv. Sci.28(15).

[14] Eric S. Donkor., 2018Article ID 3238165:doi: 10.1155/2018/3238165.

[15] Shasha Wang, Ying Li, Jishu Tian, Xiaoqiong Peng, Ling Yi, Cuiping Du, Changmei Feng, Chunmei Liu, Rong Deng, Xianju Liang., 2020 Cardiovascular diagnosis and therapy.,10(5).

[16] Sarfraz N. Brohi, Thulasyammal Ramiah Pilla, Sukhminder Kaur, Harsimren Kaur, Sanath Sukumaran, David Asirvatham.,2019 ACM J. Emerg. Technol. Comput. Syst. DOI: 10.1007/978-3-030-23943-5.

[17] Peter Horvath, Thomas Wild, Ulrike Kutay., 2011 J. Biomol. Screen., 16(9).

[18] Parameshchhari, B.D. and Panduranga, H.T., 2022. Medical image encryption using SCAN technique and chaotic tent map system. In Recent Advances in Artificial Intelligence and Data Engineering (pp. 181-193). Springer, Singapore.

[19] Vamsi Bandi, Debnath Bhattacharyya, Divya Midhunchakkravarthy., 2020IIETA Journal., DOI: https://doi.org/10.18280/ria.340609.

[20] Seung Nam Min , Se Jin Park , Dong Joon Kim , Murali Subramaniyam , Kyung:Sun Lee., 2018 Eur Neurol., DOI: 10.1159/000488366.

[21] M. Sheetal Singh, Prakash Choudhary., 2017 8th Annual Industrial Automation and Electromechanical Engineering Conference (IEMECON)., DOI: 10.1109/IEMECON.2017.8079581.

[22] A.Sudha, P.Gayathri, N.Jaisankar., 2012 Int. J. Comput. Appl.(0975 – 8887) 43(14).

[23] Medical software user interfaces, stroke MD application design: Elena Zamsa., 2016 IEEE Explore., DOI: 10.1109/EHB.2015.7391403.

[24] Kansadub T., Thammaboosadee S., Kiattisins S., Jalayondeja C., 2015 8th Biomedical Engineering International Conference (BMEiCON): DOI:10.1109/BMEiCON.2015.7399556.

[25] Nwosu, C.S., Dev, S., Bhardwaj, P., Veeravalli, B., John, D., 2019in41st Annual International Conference of the IEEE Engineering in Medicine and Biology Society IEEE.

[26] Fahd Saleh Alotaibi: 2019 inInt. J. Adv. Comput. Sci. Appl. (IJACSA).

[27] Ohoud Almadani, Riyad Alshammari.,2018 in Int. J. Adv. Comput. Sci. Appl. (IJACSA).

[28] Kansadub, T., Thammaboosadee, S., Kiattisins, S., Jalayondeja, C., 2015 in8th Biomedical Engineering International Conference (BMEiCON) IEEE.

[29] Sivaramkrishnan, M., Varma, P. S., Kaliappan, S., Saleem, R., Lal, A., Sumana, B. K., & Ramkumar, M. S., 2021In 2021 2nd International Conference on Smart Electronics and Communication (ICOSEC) IEEE., pp.726:732.

[30] Aditya Khosla, Yu Cao, Cliff Chiung:Yu Lin, Hsu:Kuang Chiu, Junling Hu, Honglak Lee., 2010 in Proceedings of the 16th ACM SIGKDD international conference on Knowledge discovery and data mining.

[31] Shanathi, D., Sahoo, G., Saravanan, N., 2009 Int. J. Biometric Bioinform. (IJBB).



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# Applications of Deep Learning (DL) Techniques in Detecting Breast Cancer and Malignant Cells

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According to the most recent statistics, the most common type of cancer globally is the breast carcinoma and kills close to 900,000 people annually. Early and accurate diagnosis of the illness can increase the likelihood of successful treatment and lower the mortality rate. In fact, an early diagnosis can help stop it from spreading and prevent the premature victims from getting it. Researchers who study cancer have a number of difficulties when attempting to differentiate between benign and malignant tumors as well as attempting to make judgments about benign and metastatic breast carcinoma. Examine the effectiveness of automated deep learning algorithms at identifying malignant cells in women's breasts and cancer stage. This paper suggests applying deep learning algorithms to whole-slide pathology images in order to possibly increase diagnostic efficacy and accuracy. The convolutional neural networks (CNN), sparse auto encoders (SAE), and stacked sparse auto encoders are illustrations of techniques of deep learning were used in this research work. There are numerous public mammographic databases available. The methods discussed in this paper are put to the test using the mini-MIAS mammographic database. The stacked sparse auto encoder performs better, this method has to be tested in a clinical setting before being used. It has higher accuracy and precision as compared to CNN and SAE. Better diagnostic performance was achieved by several deep learning methods. Deep learning algorithms are used to better reliably identify tiny tumors while detecting breast cancer via a mammogram.

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 **Contents****I. Introduction**

According to a reputable source like the Centers for Disease Control and Prevention (CDC), breast carcinoma is one of the frequently causing type of cancer in women. Depending on a number of variables, there are wide variations in the probability of healing breast cancer and the nature of a woman's tumour. 2 of the biggest factors are the disease's stage when she receives her diagnosis and aspects. This cancer develops in the breast cells. Most of the time one of the two breast regions named as the lobules or the ducts is where the cancer first appears. A cancerous condition called adipose tissue, which is found in your breast, is also possible both the fibrous connective tissue and tissue. Unchecked cancer cells could potentially frequently spread to unaffected tissue of breast and can reach to the lymph nodes situated below the arms [1].

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# Retinal blood vessel segmentation using AI

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Retinal microvascular is a dependable marker of abnormalities in vessel morphology, that have been linked to a variety of clinical disorders, both in ocular and metastatic disease. However, accurate vessel segmentation, which would be intricate- and time-intensive, is required for objective and statistical evaluation of the retinal blood vessels. In terms of segmenting retinal vessels, artificial intelligence (AI) has shown a significant amount of promise. In this study, the fundus images retinal blood vessel is segmented using deep learning methods. The data set required for this study is collected from the Kaggle website and pre-processed using various techniques to make it compatible with the deep learning models. The pre-processed images are then segmented using deep learning models such as LadderNet and UNet. The efficiency of the deep learning models are validated using performance metrics such as Intersection of Union (IoU), accuracy and F1 score. This study shows an accuracy of 0.98% using the UNet deep learning model and it is deemed to be an efficient model than the pre-existing models.

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## Contents

### I. Introduction

Retinal blood vessels, classification of the blood vessels and their structural details is crucial for the computer-aided identification and treatment of various ailments. Numerous research has focused at how these eye abnormalities relate to the properties of the retinal blood vessels. Some fatal systemic ailments, such as cardiovascular conditions and neurological conditions, can also be shown in fundus imaging. It is hypothesized that a number of retinal blood vessels morphological features are connected to illness susceptibility and advancement. The very first step to scientifically evaluating the fundus image vasculature and statistically understanding the morphological features is retinal blood vessels segmentation, which is the segmentation of visible vessels from a fundus image. These methods enable for the collection of statistical data that is noninvasive or in vivo. The primary goal of recognising and pinpointing retinal vessels is to differentiate the various vascularization structure tissues of the retina from the perspective of the fundus image. Scientist's attention has been drawn to the recognition of retinal vessels due to the availability of non-invasive fundus imaging technology and the key details obtained from of the vascularization framework for the identification and prognosis of a broad range of retinal pathogenesis. Using fundus camera equipment, retinal blood vessels can be imaged efficaciously and non-invasively. Developing scientific proof encompassing translational evidence suggests that morphologies in retinal blood vessel sections are initial markers of cardio-metabolic risk and outcome similar to any illness. As a result, data from large population-based-related appears to work is required to determine the essence of these morphometric leanings. Numerous processes have been used to investigate retinal images. While these offer a variety of retinal vessel lists, they are frequently restricted in terms of research and numerical value, and have constrained processing, including the ability to distinguish between venous system and capillaries. As a result, developing a reliable method for retinal image examination technique and generating a rich quantification of retinal vasculature in large numbers of fundus cases.

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