

# International Journal of Dentistry and Oral Science (IJDOS) ISSN: 2377-8075

## Awareness about Medicinal application of Copper Nanoparticles among Dental Students

Research Article

Dhanraj Ganapathy1\*, Martina Catherine2

<sup>1</sup>Professor & Head of Department, Department of Prosthodontics, Saveetha Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, Chennai, India.

<sup>2</sup>Tutor, Department of Prosthodontics, Saveetha Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, Chennai, India.

#### Abstract

**Introduction:** Nanobiotechnology is a new discipline of science that deals with nanoscale materials in fields including biotechnology, nanotechnology, physics, chemistry, and material science. For the creation of metallic nanoparticles, three primary approaches are used: chemical, physical, and biological. Copper and its compounds have also been employed as effective antibacterial, antifungal, antiviral, and molluscicidal agents, in addition to these uses. Copper nanoparticles have recently gained popularity due to their catalytic, optical, electrical, and antimicrobial properties.

Aim: This survey was conducted for assessing the awareness about medicinal application of copper nanoparticles amongst dental students.

**Materials and Method:** A cross-section research was conducted with a self-administered questionnaire containing ten questions distributed amongst 100 dental students. The questionnaire assessed the awareness about copper nanoparticles therapy in medical applications, their antibacterial properties, anti-fungal properties, anti-viral properties, anti-cancer activities, mechanism of action and toxicity effects, the responses were recorded and analysed.

**Results:** 11% of the respondents were aware of the medicinal applications of Copper Nanoparticles. 9% were aware of antibacterial properties of Copper Nanoparticles, 9% were aware of anti-fungal properties of Copper Nanoparticles, 7% were aware of anti-viral properties of Copper Nanoparticles, 5% were aware of, anti-cancer activities of Copper Nanoparticles and, 5% were aware of mechanism of action and toxicity effects, of Copper Nanoparticles.

**Conclusion:** There is limited awareness amongst dental students about use of Copper nanoparticles therapy in medical applications. Enhanced awareness initiatives and dental educational programmes together with increased importance for curriculum improvements that further promote knowledge and awareness of Copper nanoparticles therapy.

Keywords: Awareness; Copper; Nanoparticles; Students; Medicinal.

## Introduction

Nanobiotechnology is a new discipline of science that deals with nanoscale materials in fields including biotechnology, nanotechnology, physics, chemistry, and material science. For the creation of metallic nanoparticles, three primary approaches are used: chemical, physical, and biological [1]. For ages, copper has been used as a biocide. In the 1880s, copper sulphate, lime, and water (Bordeaux mixture) and copper sulphate and sodium carbonate (Burgundy mixture) were employed as potential fungicides for spraying grapes to combat mildew in the United States and France, respectively [1, 2].

Copper and its compounds have also been employed as effective antibacterial, antifungal, antiviral, and molluscicidal agents, in addition to these uses. Copper compounds, however, may be hazardous to fish and other species. It may potentially pose a threat to the ecosystem. As a result, greater doses of direct copper and copper compounds should be avoided. Copper nanoparticles, on the other hand, can be used as a replacement to prevent these problems. Copper nanoparticles have recently gained popularity due to their catalytic, optical, electrical, and antimicrobial properties [3].

Metal nanoparticles such as copper, silver, palladium, platinum, titanium, and others are technologically significant due to their

#### \*Corresponding Author:

Dhanraj Ganapathy, Professor & Head of Depa

Professor & Head of Department, Department of Prosthodontics, Saveetha Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, Chennai, India. Tel: 9841504523 E-mail: dhanrajmganapathy@yahoo.co.in

Received: September 12, 2021 Accepted: September 20, 2021 Published: September 21, 2021

Citation: Dhanraj Ganapathy, Martina Catherine. Awareness about Medicinal application of Copper Nanoparticles among Dental Students. Int J Dentistry Oral Sci. 2021;8(9):4350-4354. http://dx.doi.org/10.19070/2377-8075-21000885

Copyright: Dhanraj Ganapathy<sup>©</sup>2021. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original author and source are credited.

optical, electrical, and catalytic capabilities, as well as their applicability in a variety of fields. Due to their powerful antibacterial effects against a wide spectrum of pathogens, including multidrug-resistant species, silver and copper nanoparticles have gained prominence as innovative antimicrobial agents. Because silver is a costly metal, the cost of producing silver nanoparticles is also significant. Copper, on the other hand, is less expensive than silver and is readily available, making the synthesis of copper nanoparticles cost-effective. Copper nanoparticles have the extra benefit of oxidising to generate copper oxide nanoparticles, which are easy to mix with polymers or macromolecules and have reasonably stable chemical and physical properties [4, 5]. Our research experience has prompted us in pursuing this research [6-17]. This survey was conducted for assessing the awareness about medicinal application of Copper nanoparticles amongst dental students.

## Materials and Methods

A cross-section research was conducted with a self-administered questionnaire containing ten questions distributed amongst 100 dental students. The questionnaire assessed the awareness about

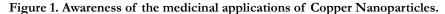
copper nanoparticles therapy in medical applications, their antibacterial properties, anti-fungal properties, anti-viral properties, anti-cancer activities, mechanism of action and toxicity effects, the responses were recorded and analysed.

#### Results

11% of the respondents were aware of the medicinal applications of Copper Nanoparticles (Fig 1). 9 % were aware of antibacterial properties of Copper Nanoparticles (Fig 2), 9 % were aware of anti-fungal properties of Copper Nanoparticles (Fig 3), 7 % were aware of anti-viral properties of Copper Nanoparticles (Fig 4), 5% were aware of anti-cancer activities of Copper Nanoparticles (Fig 5) and, 5% were aware of mechanism of action and toxicity effects, of Copper Nanoparticles (Fig 6).

## Discussion

Copper nanoparticles have been proven to be efficient against both gram-positive and gram-negative bacteria, in addition to



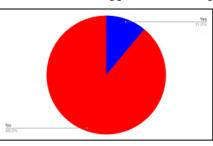


Figure 2. Awareness of anti-bacterial properties of Copper Nanoparticles.

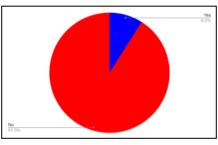


Figure 3. Awareness of anti-fungal properties of Copper Nanoparticles.

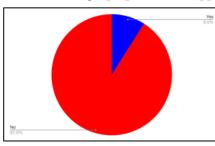


Figure 4. Awareness of anti-viral properties of Copper Nanoparticles.

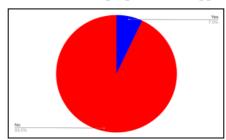


Figure 5. Awareness of, anti-cancer activities of Copper Nanoparticles.

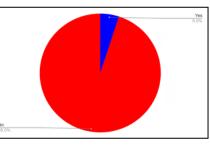
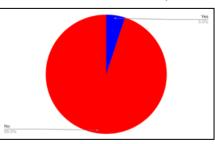


Figure 6. Awareness of mechanism of action and toxicity effects, of Copper Nanoparticles.



controlling yeast and mould growth [18]. Using the Kirby–Bauer diffusion method, Das et al. (2010) investigated the antibacterial activity of copper nanoparticles against three bacteria: Staphylococcus aureus, Bacillus subtilis, and Escherichia coli. Copper nanoparticles were discovered to be excellent growth inhibitors against these bacteria [19]. Copper nanoparticles were found to have promising antibacterial action against Micrococcus luteus, S. aureus, E. coli, Klebsiella pneumoniae, and Pseudomonas aeruginosa in a study by Ramyadevi et al. (2012). E. coli, S. aureus, M. luteus, and K. pneumoniae were the most vulnerable bacteria, while P. aeruginosa was found to be resistant to copper nanoparticles[20].

E. coli and B. subtilis susceptibility to silver and copper nanoparticles was investigated by Yoon et al. (2007). They discovered that when the concentration of nanoparticles increased, the survival rate of bacteria reduced. Silver and copper nanoparticles totally inhibited E. coli and B. subtilis at concentrations more than 70 and 60 g/mL, respectively. Copper nanoparticles were shown to be more effective than silver nanoparticles in this investigation. Copper oxides, meantime, are gaining popularity as antibacterial agents due to their ability to be synthesised with extraordinarily high surface areas and unique crystal morphologies [21]. However, gram-negative organisms were more susceptible to copper nanoparticles in time-kill studies [22].

Copper nanoparticles have also been studied for use in biotechnological applications that could help fight fungal illnesses. Researchers have tried to combine copper nanoparticles with a polymer material to create a composite that may release metal species in a controlled manner, inhibiting the growth of fungus and other pathogenic microbes [23].

Among the different species of fungi, Saccharomyces cerevisiae is said to be a model organism for studying the antifungal activity of nanomaterials [24]. Cu-based zeolites were shown to have fungicidal activity against Cladosporium cladosporoides, Phaeococcomyces chersonesos, and Ulocladium chartarum isolated from marble by Petranovskii et al. Kim et al. used a disc diffusion assay to evaluate antibacterial activity of the Cu–SiO2 nanocomposite against Candida albicans and Penicillium citrinum, and they reported promising action against both fungi [25]. In another study, the antifungal activity of hyper-branched polyamine copper nanoparticles, Cu–SiO2 nanocomposites, SiO2–Cu, and copper-doped hydroxyapatite nanopowders against C. albicans, a pathogenic fungus that causes infections in the mouth, oesophagus, gastrointestinal tract, urinary bladder, and genital tract, was investigated. The results of their research showed that colloidal hyperbranched polyamine/copper nanoparticles suppressed C. albicans development even at a low concentration of 1.4 g/100 L [26].

A few papers on copper nanoparticle antiviral activity are published, confirming that copper nanoparticles have promising antiviral activity. Using a plaque titration experiment, Fujimori et al. examined the antiviral efficacy of nanosized copper iodide particles with an average size of 160 nm against an influenza A virus of swine origin (pandemic [H1N1] 2009). They demonstrated dose-dependent activity on virus titer, with the 50 percent effective concentration for 60 minutes of exposure duration being around 17 g/ml. SDS-PAGE examination later showed the virus's inactivation as a result of viral proteins like hemagglutinin and neuraminidase being degraded by nanosized copper iodide particles. As a result, Fujimori et al. asserted that these nanoparticles could be effective in the construction of filters, face masks, protective apparel, and kitchen towels to defend against viral attacks [27].

Ramyadevi et al. described the chemical synthesis of metallic copper nanoparticles using a polyol technique that used copper acetate as a precursor and Tween 80 as both the medium and the stabilising reagent. They also tested the anti-parasitic properties of copper nanoparticles against hematophagous malaria vector Anopheles subpictus Grassi, filariasis vector Culex quinquefasciatus, and cattle tick Rhipicephalus microplus, Canestrini larvae. Their research found that metallic nanoparticles were harmful to aquatic creatures, owing to particulate impacts rather than the release of dissolved ions [20].

Nanoparticles have a unique capability for drug loading, effective photoluminescence, and targeted administration of imaging agents and anti-cancer therapies, among many other applications. Jose et al. investigated the ability of copper nanoparticles to degrade DNA and their anti-cancer properties. They discovered that copper nanoparticles degrade isolated DNA molecules in a dose-dependent manner by generating singlet oxygen. Copper nanoparticle DNA degradation was prevented using singlet oxygen scavengers such as sodium azide and tris (hydroxyl methyl) aminomethane, showing the participation of activated oxygen species in the degradation process. They also discovered that copper nanoparticles might cause apoptosis in U937 and HeLa cells from human histiocytic lymphoma and human cervical carcinoma, respectively, through generating cytotoxicity [28].

Chang et al., proposed three pathways based on oxidative stress, coordination effects, and nonhomeostasis effects to explain why copper and zinc oxide nanoparticles cause toxicity in eukaryotic cells. Nanoparticles can enter the cell directly through the pores in the cell membrane, or they can enter through ion channels and transporter proteins on the plasma membrane, according to the researchers. Endocytosis allows certain nanoparticles to enter cells. Nanoparticles that penetrate the cell can interact directly with oxidative organelles like mitochondria. Later, redox active proteins increase the development of reactive oxygen species (ROS) in cells, and nanoparticle-produced ions (Cu2+) can cause ROS through a variety of chemical processes. ROS has the ability to cause DNA strand breaks and alter gene expression. Cu2+ ions can also form chelates with biomolecules or dislodge metal ions from certain metalloproteins, resulting in functional protein inactivation. Cu2+ produced by copper oxide nanoparticles raises local concentrations and impairs cellular metal cation homeostasis, leading to cell toxicity [29].

The effect of copper nanoparticles on the rat's dorsal root ganglion (DRG) was investigated by Prabhu et al. For 24 hours, these neurons were exposed to copper nanoparticles of varying concentrations (10-100 M) and diameters (40, 60, and 80 nm). When compared to unexposed control cultures, light microscopy, histochemical staining for copper, lactate dehydrogenase assay for cell death, and MTS [3-(4,5- dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide] assay for cell viability revealed a significant toxic effect with all sizes of nanoparticles tested. They also discovered that small-sized nanoparticles with higher concentrations had the most harmful effects [30]. PC12 cells can be employed as model cells for in vitro investigations in neuron research, according to a study by Xu et al. They found that increasing concentrations of copper nanoparticles (nano-Cu) and treatment duration reduced PC12 cell viability, showing that cell viability is related to concentration and treatment time. These findings showed that copper nanoparticles are toxic to DRG neurons in rats and PC12 cells in mice in a size and dose dependent way [22]. In another study, it has been reported that CuO nanoparticles induced cytotoxicity in HepG2 cells in a dose-dependent manner [31]. Researchers have claimed that tumor suppressor gene p53 and apoptotic gene caspase-3 were upregulated when exposed to CuO nanoparticles[32].

#### Conclusion

There is limited awareness amongst dental students about use of Copper nanoparticles therapy in medical applications. Enhanced awareness initiatives and dental educational programmes together with increased importance for curriculum improvements that further promote knowledge and awareness of Copper nanoparticles therapy.

#### References

- Malathi S, Balasubramanian S. Synthesis Of Copper Nanoparticles And Their Biomedical Applications: Green Synthesis Of Copper Nanoparticles. LAP Lambert Academic Publishing; 2012. 52 p.
- [2]. Dadgostar N, Ferdous S, Henneke D. Colloidal synthesis of copper nanoparticles in a two-phase liquid–liquid system. Materials Letters. 2010 Jan 15;64(1):45-8.
- [3]. Dong L, Zhang W, Fu Y, Lu J, Liu X, Tian N, Zhang Y. Reduced Graphene Oxide Nanosheets Decorated with Copper and Silver Nanoparticles for Achieving Superior Strength and Ductility in Titanium Composites. ACS Appl Mater Interfaces. 2021 Sep 3. Pubmed PMID: 34478253.
- [4]. Peddi P, Ptsrk PR, Rani NU, Tulasi SL. Green synthesis, characterization, antioxidant, antibacterial, and photocatalytic activity of Suaeda maritima (L.) Dumort aqueous extract-mediated copper oxide nanoparticles. J Genet Eng Biotechnol. 2021 Aug 30;19(1):131. Pubmed PMID: 34460013.
- [5]. Mohamed Mowafy S, Awad Hegazy A, A Mandour D, Salah Abd El-Fatah S. Impact of copper oxide nanoparticles on the cerebral cortex of adult male albino rats and the potential protective role of crocin. Ultrastruct Pathol. 2021 Aug 30:1-12. Pubmed PMID: 34459708.
- [6]. Rbds H, Ganapathy D. Disinfection of dental impression-A current overview.
- [7]. Ramya G, Pandurangan K, Ganapathy D. Correlation between anterior crowding and bruxism-related parafunctional habits. Drug Invention Today. 2019 Oct 15;12(10).
- [8]. Anjum AS, Ganapathy D, Kumar K. Knowledge of the awareness of dentists on the management of burn injuries on the face. Drug Invention Today. 2019 Sep 1;11(9).
- [9]. Inchara R, Ganapathy D, Kumar PK. Preference of antibiotics in pediatric dentistry. Drug Invent Today. 2019 Jun 15;11:1495-8.
- [10]. Philip JM, Ganapathy DM, Ariga P. Comparative evaluation of tensile bond strength of a polyvinyl acetate-based resilient liner following various denture base surface pre-treatment methods and immersion in artificial salivary medium: An in vitro study. Contemp Clin Dent. 2012 Jul;3(3):298-301. Pubmed PMID: 23293485.
- [11]. Gupta A, Dhanraj M, Sivagami G. Implant surface modification: review of literature. The Internet Journal of Dental Science. 2009;7(1):10.
- [12]. Indhulekha V, Ganapathy D, Jain AR. Knowledge and awareness on biomedical waste management among students of four dental colleges in Chennai, India. Drug Invention Today. 2018 Dec 1;10(12):32-41.
- [13]. Mohamed Usman JA, Ayappan A, Ganapathy D, Nasir NN. Oromaxillary prosthetic rehabilitation of a maxillectomy patient using a magnet retained two-piece hollow bulb definitive obturator; a clinical report. Case Rep Dent. 2013;2013:190180. Epub 2013 Mar 4.Pubmed PMID: 23533823.
- [14]. Ganapathy DM, Joseph S, Ariga P, Selvaraj A. Evaluation of the influence of blood glucose level on oral candidal colonization in complete denture wearers with Type-II Diabetes Mellitus: An in vivo Study. Dent Res J (Isfahan). 2013 Jan;10(1):87-92.Pubmed PMID: 23878569.
- [15]. Menon A, Ganapathy DM, Mallikarjuna AV. Factors that influence the colour stability of composite resins. Drug Invention Today. 2019 Mar 1;11(3).
- [16]. Dhanraj G, Rajeshkumar S. Anticariogenic Effect of Selenium Nanoparticles Synthesized Using Brassica oleracea. Journal of Nanomaterials. 2021 Jul 10;2021.
- [17]. Ganapathy D, Department of Prostodontics, Saveetha Dental College, Saveetha Institute of Medical and Technical Sciences, – C, India. Nanobiotechnology in combating CoVid-19 [Internet]. Vol. 16, Bioinformation. 2020. p. 828–30. Available from:
- [18]. Schrand AM, Rahman MF, Hussain SM, Schlager JJ, Smith DA, Syed AF. Metal-based nanoparticles and their toxicity assessment. Wiley Interdiscip Rev Nanomed Nanobiotechnol. 2010 Sep-Oct;2(5):544-68. Pubmed PMID: 20681021.
- [19]. Das R, Gang S, Nath SS, Bhattacharjee R. Linoleic acid capped copper nanoparticles for antibacterial activity. Journal of Bionanoscience. 2010 Jun 1;4(1-2):82-6.
- [20]. Kruk T, Szczepanowicz K, Stefańska J, Socha RP, Warszyński P. Synthesis and antimicrobial activity of monodisperse copper nanoparticles. Colloids Surf B Biointerfaces. 2015 Apr 1;128:17-22. Pubmed PMID: 25723345.
- [21]. Yoon KY, Hoon Byeon J, Park JH, Hwang J. Susceptibility constants of Escherichia coli and Bacillus subtilis to silver and copper nanoparticles. Sci Total Environ. 2007 Feb 15;373(2-3):572-5. Pubmed PMID: 17173953.
- [22]. Xu P, Xu J, Liu S, Ren G, Yang Z. In vitro toxicity of nanosized copper particles in PC12 cells induced by oxidative stress. Journal of Nanoparticle Research. 2012 Jun;14(6):1-9.
- [23]. Cioffi N, Torsi L, Ditaranto N, Tantillo G, Ghibelli L, Sabbatini L, Bleve-Zacheo T, D'Alessio M, Zambonin PG, Traversa E. Copper nanoparticle/

polymer composites with antifungal and bacteriostatic properties. Chemistry of Materials. 2005 Oct 18;17(21):5255-62.

- [24]. Longano D, Ditaranto N, Cioffi N, Di Niso F, Sibillano T, Ancona A, Conte A, Del Nobile MA, Sabbatini L, Torsi L. Analytical characterization of laser-generated copper nanoparticles for antibacterial composite food packaging. Anal Bioanal Chem. 2012 May;403(4):1179-86. Pubmed PMID: 22262051.
- [25]. Kim YH, Lee DK, Jo BG, Jeong JH, Kang YS. Synthesis of oleate capped Cu nanoparticles by thermal decomposition. Colloids and Surfaces A: Physicochemical and Engineering Aspects. 2006 Aug 15;284:364-8.
- [26]. Mahapatra SS, Karak N. Hyperbranched polyamine/Cu nanoparticles for epoxy thermoset. Journal of Macromolecular Science<sup>\*</sup>, Part A: Pure and Applied Chemistry. 2009 Jan 26;46(3):296-303.
- [27]. Fujimori Y, Sato T, Hayata T, Nagao T, Nakayama M, Nakayama T, Sugamata R, Suzuki K. Novel antiviral characteristics of nanosized copper(I) iodide particles showing inactivation activity against 2009 pandemic H1N1 influenza virus. Appl Environ Microbiol. 2012 Feb;78(4):951-5. Pubmed PMID: 22156433.
- [28]. Jose GP, Santra S, Mandal SK, Sengupta TK. Singlet oxygen mediated DNA

degradation by copper nanoparticles: potential towards cytotoxic effect on cancer cells. J Nanobiotechnology. 2011 Mar 25;9:9. Pubmed PMID: 21439072.

- [29]. Chang YN, Zhang M, Xia L, Zhang J, Xing G. The toxic effects and mechanisms of CuO and ZnO nanoparticles. Materials. 2012 Dec;5(12):2850-71.
- [30]. Prabhu BM, Ali SF, Murdock RC, Hussain SM, Srivatsan M. Copper nanoparticles exert size and concentration dependent toxicity on somatosensory neurons of rat. Nanotoxicology. 2010 Jun 1;4(2):150-160. Pubmed PMID: 20543894.
- [31]. Laha D, Pramanik A, Maity J, Mukherjee A, Pramanik P, Laskar A, Karmakar P. Interplay between autophagy and apoptosis mediated by copper oxide nanoparticles in human breast cancer cells MCF7. Biochim Biophys Acta. 2014 Jan;1840(1):1-9. Pubmed PMID: 23962629.
- [32]. Siddiqui MA, Alhadlaq HA, Ahmad J, Al-Khedhairy AA, Musarrat J, Ahamed M. Copper oxide nanoparticles induced mitochondria mediated apoptosis in human hepatocarcinoma cells. PLoS One. 2013 Aug 5;8(8):e69534. Pubmed PMID: 23940521.