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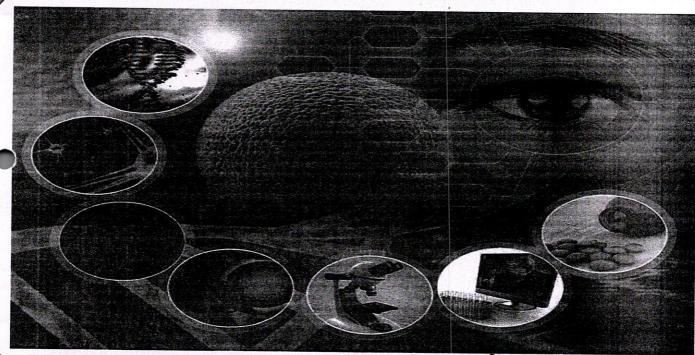
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Biosynthesis Of Zinc Oxide (Zno) Nanoparticles Using Ruta Chalepensis Sp. Plant Leaves And Its Antibacterial Activity

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Abstract: Zinc oxide is an inorganic metal compound which has wide range of application in various field. Biosynthesis of nanoparticles from plants is an ecofriendly approach alternate to conventional physical and chemical method. Recently synthesis of nanoparticles using bioactive resources gaining importance due to its least cost, simplicity and effective antimicrobial activity. Present study focus on synthesis of ZnO nanoparticles using Ruta chalepensis plant leaves extract. The prepared ZnO nanoparticles have been characterized by UV- vis absorption spectroscopy, also prepared ZnO nanoparticles determined by the antibacterial activity against standard strains of E.coli, S.typhi, S.aureus and B. subtilis. It is observed that the ZnO synthesized in the process has the efficient antimicrobial

Key words: Zinc oxide nanoparticles, UV- vis spectroscopy, Ruta chalepensis, Minimum inhibition concentration, Antibacterial activity.

Introduction:

Nanotechnology is the developed technology which deals the synthesis of nanomaterials. These nanoparticles are having the well defines shape and size (Kartiresan et al., 2009). Nanotechnology deals with the production and usage of material with nanoscale dimension. From the last decade, the metal nanoparticles have been intensively studied. The nanoparticles ranging between 1 to 100 nm size scale. Biosynthesis of nanoparticles has greatly reduced the use of physical and chemical methods. Use of green synthesis method is rapidly increasing due to usage of least toxic chemicals and environmentally safe to nature (M.Sundrajan et al., 2015). The bioactive resources involved in green synthesis of nanoparticles are plants and their derivaties, microorganisms like bacteria, fungi, yeast and algae. Inorganic metal nano particles have been of great importance due to their distinctive features such as catalytic, optical, magnetic, electronic, and antimicrobial wound healing and anti inflammatory properties. Nanoscale zinc oxide nanoparticles had been in current researches because of their potential applications like antibacterial, antifungal, wound healing, antioxidant and optic properties (Kumar et al., 2013; Happy Agarwal et al., 2017; Madhusudanan et al., 2019). Among all the metals, zinc oxide have the wide range of applications such as semiconductor, thermoelectrical, magnetic, electroluminescent material and are mostly used in the cosmetics, etc. The Zinc oxide (ZnO) nanoparticles have the good conductivity, catalytical activity, unique antibacterial and antifungal properties (Meruvu et al., 2011). Antimicrobial properties with nano sized particles are of substantial interest in the recent past. Ruta chalepensis is an aromatic angiospermic flowering plant belongs to Rutaceae family commonly called fringed rue. It is one of the frequently used plants for medicinal purposes (Enis Ben et al., 2009). Aerial parets of the plants such as leaves, stem and flowers were used for medicinal purpose. Leaveas and young stem of plant has high content of alkaloids, furocomourins, comourins, flavonoids, phenols, amino acid and saponin. Rutachalepensis has antiinflammatory, antihelmintic and spasmolytic effect. ZnO NPs have been reported in different morphologies like nanoflake, nanoflower, nanobelt, nanorod and nanowire [63-65]. Bio synthesis of zinc oxide nanoparticles by plant Ruta chalepensis has been reported. This study therefore is aimed to synthesis ZnO nanoparticles using bio method, to analyse their characteristics using spectroscopic techniques and to evaluate its antimicrobial activity.

Materials and Methods:

Plant material and preparation of extract:

Ruta chalepensis L. leaves were collected and shed dried at room temperature and powder was prepared using mixer grinder. 5 gm of powder of Ruta chalepensis L. was dissolved in 100 ml of distilled water and stirred magnetically at 70°c for 1 hour. The obtained solution was allowed to cool at room temperature and filtered through Whatman No.1 filter paper. The filtrate of aqueous extract was used as reducing agent.

Synthesis of Zinc oxide Nanoparticles:

For the synthesis of zinc oxide nanoparticles, 0.3M zinc acetate dihydrate solution was prepared by adding 4 gm of zinc acetate dihydrate, was dissolved in 60 ml of distilled water and stirred for 10 minutes. 40ml of prepared plant extract was added into the 60 ml of zinc acetate dihydrate solution. Then the solution was stirred for 20 minute. The reacted solution was dried at 80°c overnight to yield the zinc oxide nanoparticles and employed for calcination at 100°c for 2 hours. The brown colored zinc oxide nanoparticles obtained after calcination and these synthesized ZnO nanoparticles were used for antimicrobial studies.

UV-Vis spectra analysis:

The sample was measured for its maximum absorbance using UV-Vis spectrophotometry. The optical property of ZnO nanoparticles was analyzed via ultraviolet and visible absorption spectroscopy in the range of 350-450 nm.

Determination of antibacterial activity of Zinc oxide Nanoparticles:

Antibacterial activity:

Antibacterial activity of synthesised nanoparticles was carried out by agar well diffusion method against four bacterial species i.e. two Gram negative bacteria Escherichia coli, Salmonella typhi and two Gram positive bacteria Staphilococcus aureus and Bacillus subtilis. Nutrient agar plate were prepared, 100 ul of 24 hour mature broth culture of individual bacterial strain and swabbed on plate using sterile L Shaped glass rod. The well were prepared by using sterile cork borer, 4mm size well are created with the help of sterile cork borer in each Petri plate. 40 ul sample of ZnO nanoparticles was added in the well. Sterptomysin was used as positive control. Plates were kept in refrigerator for diffusion process about 30 minute, and then the plates were incubated at 37°c for 36 hours.

Determination of minimum inhibitory concentration:

Minimum inhibitory concentration of zinc oxide nanoparticles was determined using Alamer Blue Assay (Yajko et al., 1995; Rampersad, 2012) for four different bacterial species (E.coli, S. typhi, S. aureus and B. subtilis). Each well of microtitre plate was initially added with 100 ul of sterile broth. Then 100 ul of sample of zinc oxide nanoparticles was added in the first well and two fold serial dilutions of sample were made. At last, 20 ul of bacterial cell suspension was added to each well. Microtitre plate was incubated for 24 hours at 37 °C temperature for bacteria. After incubation, 30 ul of Alamer blue dye was added to each well and re-incubated for 4 hours at room temperatu

Results and Discussion:

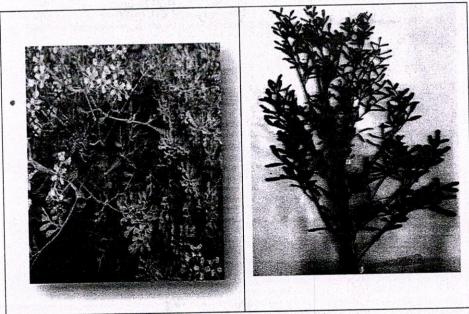


Figure.1- photoghraph of Ruta chalepensis plant

Habit and Habitat:

It is perennial herb widely distributed in Mediterranean area (Polunin and Huxley, 1987) and the Canary Islands. It grows on rocky slope. This plant was introduced in America after Spanish. It is cultivated in the tropics as a potherb or medicinal plant and has widely become naturalized. In tropical Africa it has been introduced in several countries, including the Cape Verde Islands, Sudan, Ethiopia, Somalia.

Morphometric character:

- The Ruta chalepensis belongs to class- Magnoliopsida, order- Sapindales and family-
- It is an aromatic plant characterized by glabrous, alternative bipennatisect leaves, narrow obovate segments.