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SYSTEMATIC INVESTIGATION ON NOVEL NANOCOMPOSITES FOR CANCER THERAPEUTIC APPLICATIONS

A.Jeno Deva Kiruba¹, S.Anantha Lakshmi², B.Gopal Samy³, G.Sahaya Dennish Babu⁴

^{1,2} Student, Department of Biotechnology, V.S.B Engineering College, Karur
³Associate Professor, Department of Biotechnology, V.S.B Engineering College, Karur
⁴Assistant Professor Department of Physics, Chettinad College of Engineering and Technology, Karur.

ABSTRACT

Nowadays, Cancer is taken into account one amongst the leading causes of death, along side surgery actinotherapy, therapy square measure the simplest ways in which to cure and targeting the cancer cells and destroy it. However, therapy causes several adverse effects together with radiation harm of blood cells and bodies. Nano materials are often thought-about as an alternate drug delivery methodology for cancer treatment, this study investigates that the assembly and use of MoO₃ nanoparticle for cancer therapeutic applications and treatment. Synthesis of MoO₃ nanoparticles was achieved by inexpensive microwave aided solvo-thermal methodology. The ready MoO₃ nanoparticles were characterized by XRD, SEM and TEM characterization techniques. The findings instructed that the synthesized MoO₃ sized thirty five nm and had spherical form morphological options. MoO₃ induces necrobiosis and generates reactive chemical element species (ROS). This study investigated the potential utility of MoO₃ for treating cancer cells, which could be the higher thanks to treat cancer cells.

Keywords: Cancer nanotechnology, MoO3 Nanoparticles, Spherical morphology, ROS.

1. INTRODUCTION

Engineering is science, engineering, and technology conducted at the nanoscale that is regarding one to a hundred nanometers. In fact, it's not possible to ascertain with the microscopes usually utilized in a high school science classes [1]. Nanoscience and engineering square measure the study and application of extraordinarily tiny things and might be used across all the opposite science fields, like chemistry, biology, physics, materials science, and engineering, one of these new fields typically goes beneath the name "nanoscience and nanotechnology" and joins many areas of analysis as engineering, physics, chemistry, material science and biology. The analysis during this direction has been triggered by the recent convenience of latest revolutionary instruments and techniques that square measure ready to improve our investigation talent regarding the fabric properties with a resolution near to atomic scale. Such technological advances have inspirited new pioneering experiments that have unconcealed new physical properties. Associate in Nursing effects of matter at an intermediate level between atomic and bulk.

In the previous few decades, transition metal oxides have attracted the analysis community in mickle because of their diversified applications [2]. Among them, metallic element has been found to be one amongst the foremost fascinating materials because of its distinctive structural, optical, electrical, and mechanical properties and multidirectional applications like gas sensing, photocatalysis, field emission (FE), capacitors, resistive switch, star cells, lightweight diodes, hole transport materials, etc. Metallic element could be a Block D, amount five component, whereas chemical element could be a Block P, amount a pair of component within the tubular array, metallic element doesn't occur naturally as a free metal on Earth [3], numerous experimental techniques like diffraction, scanning microscope (SEM), transmission microscope and toxicity analysis of the ready MoO₃ nanoparticles on G361 cancer line results have been given during this section.

2. METHODOLOGY

The MoO₃ nanoparticles were synthesized by a microwave assisted solvothermal route, ammonium heptamolybdate tetrahydrate (AHM; $(NH_4)_6Mo_7O_{24}, 4H_2O$) and glycol were used because 2 beginning reagents. A saturated resolution of precursor compound AHM was ready at temperature and acidified to hydrogen ion concentration around five employing a pair of 2 M aqua fortis resolution, there have been 2 sorts of the acidified AHM precursor solutions, the primary one was used instantly once preparation ("fresh" solution). The second kind of acidified precursor resolution was keep at temperature month during a tightly capped meter flask before use ("aged" solution). The resultant solution was unbroken underneath a domestic kitchen appliance and permit them apply microwaves thereon it for 15min. The obtained precipitates were sublimate by fermentation alcohol and also the particles were dried at 80°C within a hot-air appliance for 8h. The dried sample was white color.

3. MODELING AND ANALYSIS

3.1X-Ray optical phenomenon technique



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To get the part additionally the also the scale of the particles diffraction, XRD, are used. The measurements are done on a Rigaku diffractometer victimization parallel beam pure mathematics with x-ray mirror and a parallel plate collimator of zero.4°. All measurements are done on powders on glass substrates so as to be ready to directly correlate optical properties with XRD-data. The angle of incidence has been 0.5° and therefore the scan has been performed from 10° to 90° . For a first set of samples the step size has been 0.02° and therefore the integration time eleven seconds, and for a later set of samples the step size has been 0.1° and therefore the integration time fifty four seconds, this suggests, that the sampling time typically is around twelve hours for every sample, that is important to urge smart statistics [4].

3.2Scanning Electron Microscope (SEM)

Imaging by secondary electrons

Secondary electrons (SE) square measure made thanks to the springless interaction of high energy electrons with valence electrons of atoms within the specimen that cause the ejection of electrons from the valence shell of the atoms. These ejected electrons any bear further scattering whereas traveling through the specimen before they emerge from the surface of the specimen. The secondary electrons have very low energies of the order of twenty to thirty electron volt, thanks to this low energy vary those SE, that leave the surface of the sample while not being absorbed return from solely regarding the highest five to fifty A.

Imaging by backscattered electrons

Back scattered electrons (BSE) square measure made thanks to the elastic interaction of the incident electrons with the nuclei of the specimen atoms, several electrons incident on the specimen will be scattered back from the specimen during this manner and therefore the scattered angle will vary up to 180°. Variety of back scattered electrons ejected from the specimen powerfully depends on the number.

3.3 Transmission Electron Microscope (TEM)

TEM is that the technique that's of exploit use for analyzing the interior structure of a specimen.. STEM is one sort of TEM. Some TEM instruments square measure fitted with scan coils, which might scan a centered nonparticulate radiation across the specimen in a very formation pattern. This STEM mode is very helpful for effecting serial analysis across areas of the specimen high-resolution TEM (HRTEM) imaging could be a crystallographic imaging methodology that may be performed by the utilization of STEM. Once specimens square measure ultrathin (< a hundred nm), the elastic scattering dominates the springless scattering. The electrons interacting with the space lattice separate and kind advanced interference patterns visible at magnifications of 400k or additional, beneath some imaging conditions, the patterns correspond to atom positions, and that they will be accustomed investigate the expansion planes of a crystal structure.

3.4Cytotoxic Analysis

3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazoliumbromide (MTT) assay protocol was followed to guage the toxicity exhibited by the developed MoO₃ nanoparticles against cancer cells and traditional cells, associate equal density of fifteen× 10^3 cells/well was seeded in a very 15-well plate and left for attachment long. On the subsequent day, cells were washed with PBS and treated with totally different concentrations of MoO₃ (0.125 – 1.25 mMol) nanoparticles, once twenty four h of incubation, MTT (0.5 mg/ml in PBS) was additional and incubated for three h at thirty seven °C. The formazan advanced shaped by the live cells was dissolved victimization dimethyl sulfoxide (DMSO) and measured calorimetrically at 570/630 nm using a microplate reader.

4. RESULTS AND DISCUSSION

In this chapter, the crystallographical analysis and microstructural property of the synthesized MoO_3 nanoparticles is mentioned. Varied experimental techniques like diffraction, scanning microscope (SEM), transmission microscope and toxicity analysis of the ready MoO_3 nanoparticles on G361 cancer line results are given during this section.

4.1 Phase identification using XRD

From XRD, the crystallite size are often detected by victimisation the Scherer's formula,

$$P = \frac{0.9 \,\lambda}{\beta \,\cos\Theta}$$

Where P – crystallite size, λ – wavelength (1.54Å), β - Full maxima , θ - optical phenomenon angle. And by applying Scherer's formula the crystal size of the MoO₃ sample detected to be thirty five nm.



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The discrepancy in the relative peak intensities with the standard diffraction pattern of the MoO_3 may be attributed due to the fact that the spherical nanoparticles likely have preferred growth orientations. Moreover, the relative peak intensities of the (011) to (210) in our case is quite same with earlier report which implies that the MoO_3 nanoparticles fabricated by different methods exhibit different preferred orientations[5].

4.2 Shape and size detection using Scanning Electron Microscope



Figure 2 : FESEM images of MoO₃ nanoparticles at (a) – low magnification and (b)- high magnification

The morphology of the MoO_3 sample was observed in field emission scanning electron microscopy (FESEM) using ZEISS scanning electron microscope instrument. Figure 2 shows the morphology of the as prepared MoO_3 nanoparticles which reveals that the powder contains several spherical shaped nanoparticles of diameter ~25 – 35 nm.

4.3 Transmission Electron Microscope



Figure 3 :TEM images of MoO₃ nanoparticles

Transmission Microscopy (TEM) may be a technique that uses an associate beam to image a nanoparticle sample, providing abundant higher resolution than is fessible with light-based imaging techniques. TEM is that the most well-liked technique to directly live nanoparticle size, grain size, size distribution, and morphology, during this work, the TEM pictures were collected employing a JEOL JEM-2010 transmission microscope (JEOL, Japan) from Cochin University of Science and Technology, Cochin, Kerala. High-resolution TEM pictures were obtained employing a probe-corrected JEM-ARM200F atomic resolution. The TEM image of the MoO₃, with 3 magnification reveals the formation of nanoparticles with diameter of regarding 15–38 nm, it's perceived from the TEM pictures that the ready particles square measure slightly agglomerate and it's because of the solvent was well mixed with the particles and not dried abundant[6,7]



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editor@ijprems.com 4.4Cytotoxicity analysis



Figure 4 : % of viability of MoO3 nanoparticles on G361 cancer cells

MTT assay could be a measurement assay for testing the cell viability. MTT assay could be a common place check wide accustomed live the toxicity of the ready materials. The results of the MTT assay performed on differing kinds of cancer cells and traditional cells with varied concentrations of MoO_3 nanoparticles were illustrated in figure 4. From the figure it's clearly evident that each one cells showed weakened cell viability with a rise within the concentration of nanoparticles. The ascertained result's alright in agreement with the literature reports[8]. The percentage viability of the cells was calculated by considering the untreated management cells to be 100 percent viable and extrapolating the calculation to the treated cells. However, it absolutely was ascertained that at explicit concentrations, the synthesized nanoparticles showed a decrease in additional than five hundredth of the cell viability in G-361 cells. This apparently reveals that MoO_3 nanoparticles show selective toxicity towards carcinoma cells. The result indicated that the basis mediated ready MoO_3 NPs possess best property to neoplastic cell and may show potential property in therapy and cancer chemoprevention[9].

5. CONCLUSION

The MoO_3 nano particles were with success synthesized by microwave-assisted methodology. Their morphological structures additionally characterized by XRD, SEM, TEM. The cytotoxic check additionally distributed with success by victimisation MTT assay to check their effectualness and viability on cancer cells. Properly designed nanoparticles have the power to accumulate in tumors either by passive or active targeting and enhance the cytotoxic effects of antitumor agents. Nanoparticle drug formulations have the potential to beat the constrain of typical therapy by their ability to selection target cancer cells over healthy tissue.

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