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Green synthesis of zinc oxide nanoparticles using jasmine petals

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Abstract:

The green chemistry approach for the preparation of nanoparticles is becoming more attractive as it is efficient and environmentally friendly. Zinc Oxide nanoparticles (ZnO NPs) are the second most common metal oxide, it's considered an important compound due to its properties, stability, low cost and wide usage. ZnO NPs have been synthesized based on using extract of the petals of *Jasminum officinale*. The synthesized NPs were characterized by X-ray diffraction (XRD), and they showed a distribution size with in an average of 5.05nm and a hexagonal shape, Result, showed that jasmine petal extract is an excellent choice for the green production of ZnO nanoparticles.

Keywords:

Nanoparticles, Jasmin, Zinc Oxide, Green Synthesis.

1. Introduction:

In nanotechnology can we defined as the formation of chemical compounds at a range between 1 and 100 nanometers^[1]. These particles exhibit chemical and thermal stability, which gives great them importance in our N^[2] timeanomaterials (NMs) distinctive shape, size, and have a structure that give them important properties in stability, effectiveness, and other properties This increase^[3] that can be adjusted to suit the applications used due to their large surface area material with reducing the particle size and manipulation of the the main the surface area of^[4] morphology of the nanomaterial contributed to enhancing the properties of these compounds the manufacture of these particles is using biological methods -environment friendly and non ctoxi. This has given such methods great applications in the environmental, cosmetic and Based on several studies, it was found that plant synthesis is the green.^[5] medical industries effective method compared to other methods of manufacturing nanoparticles because plants contain additional effective and protected compounds and metabolites within Specifically, ZnO NPs are used in cosmetic and medical industries, antimicrobial^[6] them ,In fact.^[7] and antimicrobial agent are remarkable and other industries ,agents, sunscreens .^[8] is an aromatic medicinal and valuable plant from the Oleaceae family *Jasmine(Jasminum)* containing approximately 600 species, The *Jasminum* species ,*Jasminum*) *angustifolium*, *umsambac* *Jasmin*, , *Jasminum pubescens* and *Jasminum flexile* *Jasminum grandiflorum...etc*)^[9]. was used in the past as Antimicrobial, anti *Jasmine* and for the treatment of breast cancer^[8] inflammatory, flavoring and Fragrance agent diarrhoea, fever, abdominal pain, asthma, ,toothache, uterine bleeding, dermatitis, headache, ...etc. It has , been known that jasmine leaves and flowers have many uses, as jasmine oil played a major role e due to its It is likely that the medicinal properties of the jasmine plant ar^[10] in aromatherapy large collection of bioactive compounds such as esters and fatty The goal of this study was to green^[11] acids, terpenoids, steols, glycosides, coumarins, Phenolics ilizations synthesize ZnO NPs using Jasmine petals extract and study distribution size and stab to benefit from propeston of both ZnO and jasmin in nano size

2. Slairtam dna dohtem:

2.1. Preparation of Jasmine flower extract :

collected from local garden and washed repeatedly with (*Jasminum officinale*) Jasmine distilled water to remove the dust particles on petal surface. The flowers were dried in an oven

at 50°C for 24h to remove the residual moisture. The dried petals were ground into Powder. The aqueous extract of the sample were prepared by soaking 5 grams of the powdered petals in 100 ml of hot distilled water for 48h with stirring every other time. Then the extract filtered using Whatman filter paper. The filtered extract was used in the experiment

2.2. Biosynthesis of ZnO nanoparticles:

The ZnO NPs were made by extracting the flowers of the jasmine. 50 ml of Jasmine petal extract was added to 50 ml of 0.45 M zinc acetate solution, and 10 ml of 0.45 M NaOH. The mixture was then continuously stirred at room temperature for 2 hours using a magnetic stirrer at 600rpm. That led to formation of yellow precipitate. The precipitate was then filtered and washed many times with distilled water and ethanol in order to remove impurities, and oven dried at 100°C for one hour. The obtained dried yellow colour powders were mashed using a mortar and pestle. Finally, the mashes yellow powders were incinerated at 400°C for 1h and fine crushed using mortar and pestle, and prepared for further characterizations.

2.3. particle size of ZnO nanoparticles :

Various methods are used to achieve the desired NP size, as one of the main applications of ZnO NPs is their use as an effective drug delivery system which is influenced by the size of ZnO nanoparticles[9]. NPs have a characteristic shape, size, and structure that give them important properties in terms of stability, efficacy, and other properties that can be tailored to the application in which they are used due to their large surface area[3]. Surface area is directly related to particle size, and this relationship affects the drug release rate, Increasing the surface area of the material reduces the particle size[4] and increases the drug release rate, Larger particles contain more drug because they are surrounded by a larger core, resulting in longer drug release[13]. Several particle size and morphology measurement techniques, such as XRD (also known as X-ray diffraction), are suitable for determining crystal structure, crystallite size, and lattice parameters [14] . SEM is widely employed as a common method for assessing NPs. The interaction between the electrons and the sample leads to the generation of a signal that provides valuable insights into the sample, including its surface morphology and chemical composition [15].

2.4. The energy dispersive x-ray (edx):

The energy dispersive x-ray (edx) is from one of techniques analysis microscopy with the microscope-mail, which depends in their work on radiology x deals with the role in detect the presence of the elements in the sample goal. The EDX Microanalysis is used in nanotechnology

By many pharmacist researchers. Edx of techniques task in detecting medications such as the delivery of medicine, where edx him do a great in studies of the investigation of the nanotechnology. Edx is used to describe metal accumulated in living tissues. From the above, EDX can be considered a useful tool in tests that require identifying elements inside or outside a cell or any other model^[16].

2.5. The X-Ray measurements (XRD)

2.5.1. XRD measurements

were performed at Room temperature, in a two-circle STOE Stadi P Powder transmission diffractometer operated in Transmission mode, working with copper $K\alpha_1$ radiation ($\lambda_{CuK\alpha_1} = 0.154060$ nm), and equipped with a curved Germanium (111) monochromator. The so-called ($\theta-2\theta$) mode was employed. The XRD spectrum of silicon reference sample (ICDD PDF2 00–027-1402) of the National Bureau of Standard (USA) were recorded in ($\theta - 2\theta$) mode for goniometer calibration in order to estimate the instrumental broadening of diffraction peaks. (Table. 1)

3. Noissucid dna stluser:

3.1. Particle size :

ray diffraction analysis (XRD) and-NPs were characterized by X-The synthesized ZnO Scanning electron microscopy (SThe XRD results revealed that the sample product was .(EM .crystalline with a hexagonal phaseThe particle sizes of the synthesized ZnO nanoparticles were found to be in the range of 1 nm to 10 nm approximately. The morphology of all synthesized ZnO nanoparticles was determined using Scanning Electron Microscopy (SEM) and investigated by analyzing Fig. (1). The SEM image depicted in Fig. () illustrates the varying sizes of ZnO nanoparticles, ranging from 34 to 123 nm.

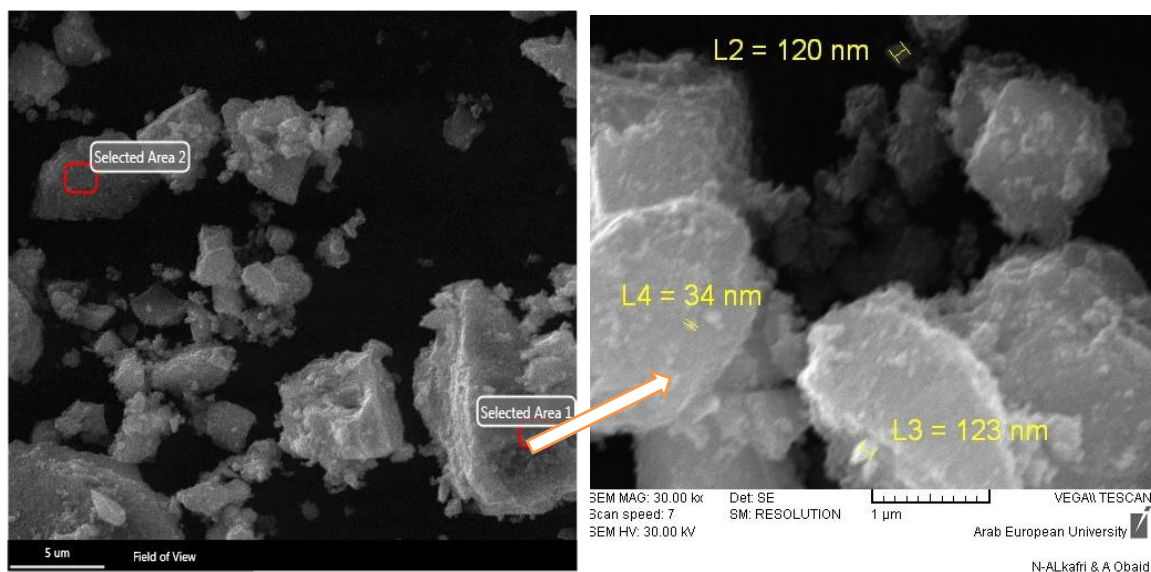


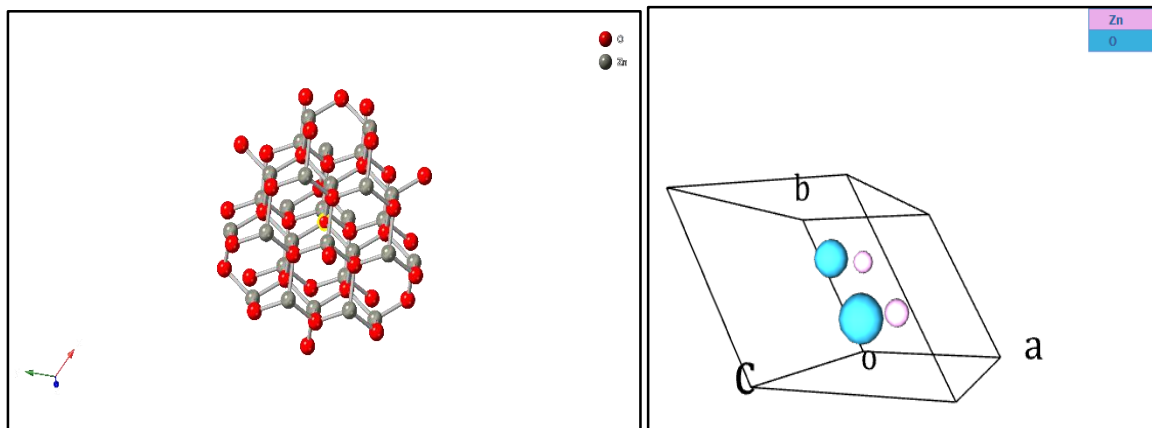
Figure. 1: SEM images of synthesized ZnO nanoparticles and their morphology and size distribution

3.2. Crystallite size:

Table. 1: Results of synthesized ZnO-NPs size

| Pos. [°2Th.] | (hkl) | Crystallite Size [nm] (Scherrer equation) |
|--------------|-------|---|
| 31.7436 | (010) | 9.381093 |
| 34.3874 | (002) | 9.036754 |
| 36.2084 | (011) | 10.02117 |
| 47.4401 | (012) | 8.332555 |
| 56.4455 | (110) | 5.759121 |
| 62.6759 | (013) | 5.637707 |
| 66.1693 | (020) | 5.767494 |
| 67.7333 | (112) | 5.064124 |
| 68.8634 | (021) | 5.101237 |
| 72.3272 | (004) | 5.061973 |
| 76.6857 | (022) | 4.75309 |
| 81.0844 | (014) | 4.804117 |
| 89.2433 | (023) | 4.203036 |
| 92.3915 | (120) | 4.098971 |
| 94.8834 | (121) | 3.586211 |

| | | |
|----------|-------|-----------|
| 98.1686 | (114) | 3.262185 |
| 102.4335 | (122) | 2.654278 |
| 103.6296 | (015) | 2.293674 |
| 106.8909 | (024) | 1.482542 |
| 109.7959 | (030) | 0.9220362 |
| 112.4603 | (031) | - |



3.3. eZAF smart quant results :

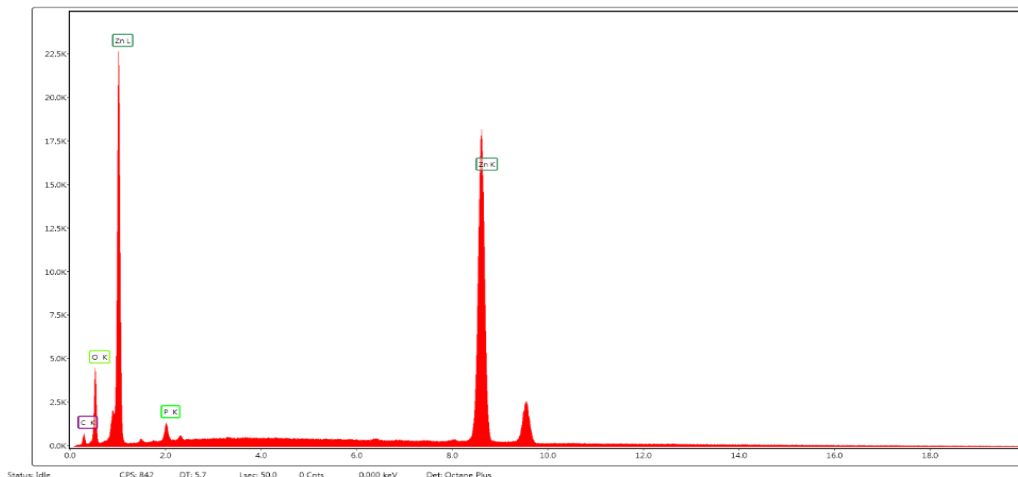
| Element | Weight % | Atomic % | Error % | Net Int. | K Ratio | Z | R | A | F |
|---------|----------|----------|---------|----------|---------|--------|--------|--------|--------|
| C K | 5.21 | 17.66 | 12.13 | 69.17 | 0.0149 | 1.3626 | 0.8508 | 0.2097 | 1 |
| O K | 10.75 | 27.36 | 7.59 | 504.86 | 0.0703 | 1.295 | 0.8764 | 0.5048 | 1 |
| Zn L | 80.22 | 49.96 | 2.45 | 2885.42 | 0.7248 | 0.9326 | 1.038 | 0.9699 | 0.9988 |
| P K | 3.82 | 5.02 | 6.37 | 184.58 | 0.0331 | 1.1158 | 0.9433 | 0.773 | 1.004 |

We can see from the above values that we have in the sample p,c and o atoms in the k orbital and Zn in the L orbital

3.4. The energy dispersive x-ray (edx):

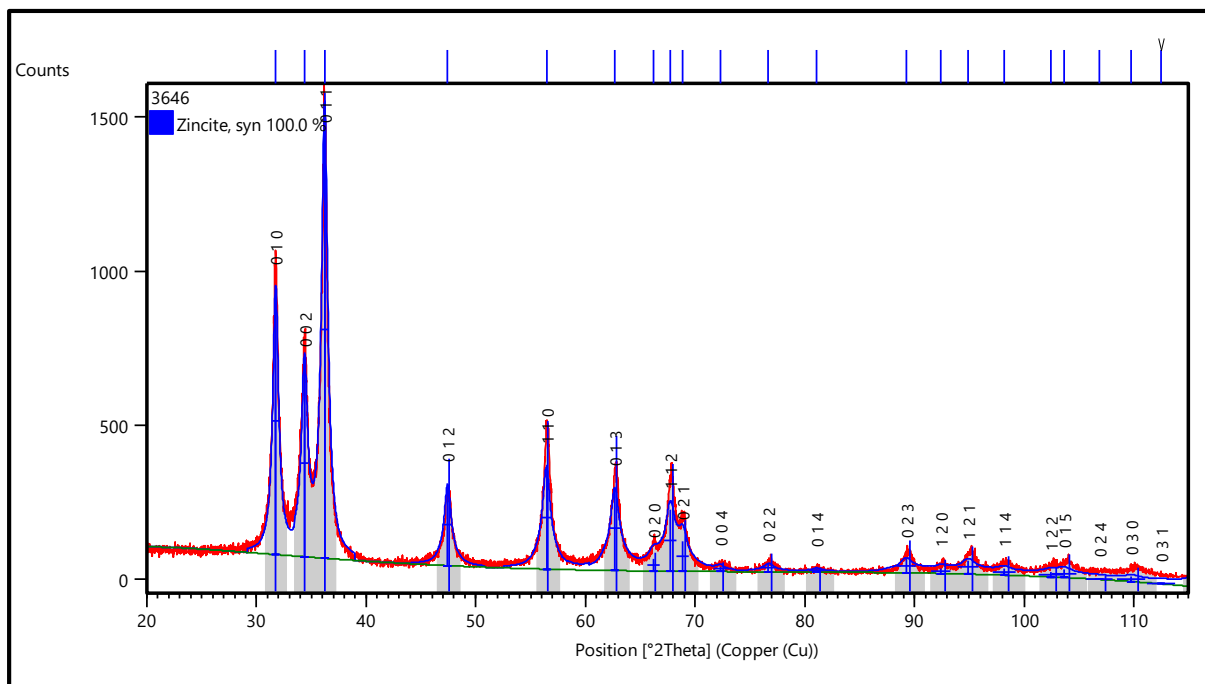
The principle of the device depends on irritating the atoms with a beam of electrons or X-rays result in a state of Instability. We have the transfer of electrons from higher to lower orbits as a result of the electronic vacancy. This transfer emits X-rays whose energy is equal to the difference in transfer between the two orbits. We notice from the figure below that there is a clear prominence at the two peaks of the oxygen in the k orbital, the zinc molecules in the k and L orbitals, and the carbon in the k orbital. These distinctive peaks In the X-ray spectrum

indicate distinct structures that contributed significantly to the detection of the elements present in the sample, which are oxygen, zinc, and carbon.



3.5. XRD measurement results:

Our sample structure according to PDF ICDD 00-036-1451,



Peak List:

| Pos. [$^{\circ}2\theta$.] | Height [cts] | FWHM Left [$^{\circ}2\theta$.] | d-spacing [\AA] | Rel. Int. [%] | (hkl) |
|-----------------------------|--------------|----------------------------------|----------------------------|---------------|-------|
| 31.7436 | 872(193) | 0.6306 | 2.81659 | 58.87 | (010) |
| 34.3874 | 614(149) | 0.6530 | 2.60586 | 41.43 | (002) |
| 36.2084 | 1481(253) | 0.6702 | 2.47887 | 100.00 | (011) |

| | | | | | |
|----------|-------------|--------|---------|-------|-------|
| 47.4401 | 267(110) | 0.8053 | 1.91489 | 18.03 | (012) |
| 56.4455 | 338(161) | 0.9460 | 1.62888 | 22.81 | (110) |
| 62.6759 | 273(144) | 1.0592 | 1.48110 | 18.43 | (013) |
| 66.1693 | 38(27) | 1.1285 | 1.41112 | 2.59 | (020) |
| 67.7333 | 201(115) | 1.1610 | 1.38229 | 13.57 | (112) |
| 68.8634 | 98(64) | 1.1850 | 1.36234 | 6.63 | (021) |
| 72.3272 | 15(18) | 1.2615 | 1.30539 | 0.99 | (004) |
| 76.6857 | 27(34) | 1.3645 | 1.24169 | 1.81 | (022) |
| 81.0844 | 13(18) | 1.4768 | 1.18506 | 0.86 | (014) |
| 89.2433 | 47(59) | 1.7105 | 1.09663 | 3.16 | (023) |
| 92.3915 | 15(20) | 1.8109 | 1.06733 | 0.99 | (120) |
| 94.8834 | 43(60) | 1.8949 | 1.04576 | 2.90 | (121) |
| 98.1686 | 22(37) | 2.0127 | 1.01935 | 1.47 | (114) |
| 102.4335 | 15(29) | 2.1786 | 0.98817 | 0.98 | (122) |
| 103.6296 | 26(61) | 2.2281 | 0.98000 | 1.78 | (015) |
| 106.8909 | 4(19) | 2.3705 | 0.95893 | 0.27 | (024) |
| 109.7959 | 15(153) | 2.5072 | 0.94154 | 1.03 | (030) |
| 112.4603 | 0.000000(3) | 2.6420 | 0.92665 | 0.00 | (031) |

Pattern List:

| Visible | Ref. Code | Score | Compound Name | Displacement [$^{\circ}$ 2Th.] | Scale Factor | Chemical Formula |
|---------|-------------|-------|---------------|---------------------------------|--------------|------------------|
| * | 00-036-1451 | 78 | Zinc Oxide | 0.000 | 0.909 | Zn O |

Unit Cell

Table. 1:

| | | |
|---------------------|----------------------|-----------------------|
| ZnO | PDF ICDD 00-036-1451 | Rietveld CIF analysis |
| Crystal system: | Hexagonal | |
| Space group: | P63mc | |
| Space group number: | 186 | |
| a (Å) = b (Å): | 3.2498 | 3.263895 ± 0.000506 |

| | | |
|-----------------------|----------|---------------------|
| c (Å): | 5.2066 | 5.228758 ± 0.000975 |
| Alpha (°) = Beta (°): | 90.0000 | |
| Gamma (°): | 120.0000 | |

Atomic coordinates

| Element | X | Y | Z |
|-----------------|------------|------------|------------|
| Zn ₀ | 0.66666700 | 0.33333300 | 0.50054800 |
| Zn ₁ | 0.33333300 | 0.66666700 | 0.00054800 |
| O ₂ | 0.66666700 | 0.33333300 | 0.87976200 |
| O ₃ | 0.33333300 | 0.66666700 | 0.37976200 |

Noisulcnoc .4:

One of the most important methods for producing ZnO NPs is green synthesis, which has appeared on a large scale. Therefore, a rapid understanding of green synthesis has a major role at the level of industrial production using biological or plant extracts. So, after obtaining a jasmine plant extract from flower petals, green technology was used to effectively form zinc nanoparticles. The findings showed that the Jasmine plant’s petals extract is an excellent Option for producing ZnO nanoparticles using a green manufacturing process. Where the average size of the particles is was 5.05 nm, and the EDX test, we show the presence of zinc, oxygen, and carbon molecules? Showing distinctive results which is specific for ZNPs

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