

Green synthesis of tin based nano medicine: Assessment of microstructure and surface property

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Abstract: The present study is about an eco-friendly and cost effective method of using five potent plant extracts and sunlight as the reaction medium and grinding as a nano technique to prepare nano composite of tin (IV) oxide from tin metal. The equivalent preparation is widely used as a drug formulation in the treatment of spectrum of infectious diseases. The intensive grinding and calcinations with the plant material lead to the formation of nano crystallite with a tetragonal phase with an average crystallite size of 27nm. The elements Ca, P, Na, Mg and Fe at the microscopic level identified at the surface of the nano crystallite by the X – ray Photo Electron Spectroscopy showed the uniqueness of the synthesis. The formation of spherical crystallites with good distribution of pores was indicated by the electron microscopic images and adsorption studies. The presence of oxygen deficiency arising due to nonstoichiometry in the crystal lattice was evident from the Electron paramagnetic Resonance study. The free radical equivalent of DPPH radical calculated for the final stage sample was suggestive of its anti-oxidant property and hence it's biological activity. The overall research study provided an alternative route of preparing nontoxic nano tin (IV) oxide composite with good biological activity apart from its already proven catalytic and sensor applications.

Keywords: Nanomedicine, Green Synthesis, Nano Tin (IV) Oxide Composite, Microstructure

1. Introduction

There is a growing demand for nano scale materials and there is a need for an integrative approach of conventional and advanced techniques of nano synthesis. Among the nano materials, those of metals and metal oxides are gaining greater significance due to their growing contribution in the chemical, electronic, pharmaceutical industries [1] Research reports on modern nano technology are many but are not cost effective and hence not affordable universally. In this context, green synthetic approach using plant extracts are widely explored owing to their simplicity and eco – friendly approach as shown by [2].

The bio molecules like alkaloids, flavones, terpenes, amino acids, carbohydrates present in the plant materials can act as reducing agents, capping agents, or can increase the rate of nano synthesis or the stability of the product.

There are many reports on the effect of plant materials playing a key role on the surface morphology and size of the metals [3].

Among the nano metal oxides, tin (IV) oxide is a versatile oxide having wider applications in the field of gas sensing and catalysis and as a transparent conducting oxide [4]. In all its above applications, its bulk structure, surface and interface properties play a key role. Apart from these applications, biological and medicinal applications have been reported recently for nano tin (IV) oxide, which gives new value addition to nano tin (IV) oxide. Triligio *et al* (2012) [5] reported the anti viral activity of nano tin (IV) oxide wires while, nano crystalline tin (IV) oxide was detected and fingerprinted in the tin based herbal formulations used for the treatment of range of chronic

diseases [6,7].

There are many synthetic methods available for nano crystalline tin (IV) oxide. Some of the important methods are Sol – gel, Solvothermal, Electro chemical, Chemical Vapour and Pulsed Laser Deposition methods. But, a very few reports are available on its green synthesis. Among the green methods of preparations of nano tin (IV) oxide and its calcium salt, there are research reports by which were in the context of drug formulations for a range of chronic diseases.

The present study was carried out to find out the physico chemical changes during the processing, nature of the intermediates if any, and the final product obtained during the successive grinding with five different plant extracts and calcinations of metallic tin. The method of preparation was in the lines of *Materia Medica* [8] which was intended for the treatment of wide range of disease, viz:- Delirium, Tuberculosis, Ulcer The bio chemical properties of the plant extracts used in the present study namely, *Tamarindus indica*, *Ficus bengalensis*, *Baringtonia acutangularis*, *Annona Squamosa Linn*, *Cyclea peltata* used in the present study are very significant [9,10,11,12,13].

The product sample at the end of each step was subjected to the analyses in terms of material science and nano science. The final product of the synthesis namely, nano tin (IV) oxide were found to exhibit all the microstructural, surface and morphological properties required for its reported application as a n type semiconductor, sensor, and catalyst. In addition, a preliminary screening for antioxidant property of the product was also carried out to relate the structure with the (re)activity of the product. The study suggested a unique synthetic process of nano composite of tin (IV) oxide which has multi dimensional applications including the bio medical ones.

2. Experimental

The precursor of the process namely, metallic tin was obtained from Sigma Chemicals, India. The plants were obtained from the local farm garden. The pure white tin metal was powdered and triturated (ground with the plant extract intensively in pestle and mortar) with the freshly prepared leaf extracts of the plants in the ratio of 1:6 till a homogenous paste was formed and then sun dried till the sample was completely free from moisture and ready for calcinations. Then it was calcined at a running temperature of 400° C. In step II, the calcined product of the step I was subjected to trituration with the second plant extract namely *Ficus Benghalensis* followed by calcinations. In a similar way the product of step II, step III and step IV were subjected to trituration with plant extract of *Baringtonia acutangularis*, *Annona Squamosa Linn*, *Cyclea Peltata* respectively. The samples were collected at the end of each step I, II, III, IV, V and labelled as S1, S2, S3, S4, and S5 respectively.

2.1. Analyses of Bulk and Surface Properties

The structural aspects of the samples were studied in terms of bulk and surface properties. For FT-IR spectral analysis, sample was mixed with KBr (Sigma, US) and the pellet obtained after hydrolytic press, were analyzed using Spectrum one (Perkin-Elmer Co., USA model). All measurements consisted of 500 scans and the plain KBr pellet used as background reference. FT- Raman analysis was performed using FT Raman Bruker RFS/ 27 with a He: Ne Yag laser with a 100 mW of power (with a excitation wavelength of 1064 nm) in a wavelength range of 50 – 4000 cm^{-1} using a liquid nitrogen cooled Germanium as detector and Rock – Solid interferometer.

In the present study, Perkin-Elmer 5300DV ICP-OES was used for the quantitative determination of metallic elements. The percentage composition of C, N, H and S in the sample was determined using Elemental Analyser (Euro Vector model). Photoluminescence study of the sample was performed using Cary 100 (Varian) Fluorescence spectrophotometer. Thermo gravimetric analysis of the sample was made using TGA Q50 (V20.6 Build 31) and DSC analysis using DSC Q200 (V23.10 Build 79) at nitrogen atmosphere. Powder X-ray Diffraction (PXRD) patterns of the solid calcined sample was recorded using Rigaku Powder X-ray diffractometer with Cu/30KV/15mA and a scan speed of 4deg/min in 2θ range from 10 to 80 degrees with a sampling step of 0.05 deg. EPR spectra of the sample was recorded as a first derivative using Bruker EMX computer controlled spectrometer operating at X-band frequencies (~9.5 GHz) with a rectangular cavity.

The surface area of the nano crystalline samples and their adsorption properties were studied by BET analysis. The specific pore volume and distribution of pores in the samples were determined by Barrett - Joyner-Halenda (BJH) analysis.

The surface morphology of the samples was studied using Environmental Scanning Electron Microscope (ESEM) (Hitachi S 4000 voltage of 10Kev). The relative elemental composition of the samples was computed directly with EDX software using ZAF correction. X-ray photoelectron spectroscopy (XPS) analysis was made using Kratos Axis 165 instrument with a dual anode (Mg and Al) and Mg- K_{α} anode. Morphology and the dimension of the crystalline samples were taken on a JEOL model – 200cx using an accelerating voltage of 120 kV.

2.2. Preliminary Profile of the Anti Oxidant Property

The antioxidant profile of the product was studied using EPR spectroscopy. Sample at 1mg ml^{-1} concentration in ethanol was taken and the Di Phenyl Picryl Hydrazyl free radical DPPH• concentration was fixed as 2 mM in ethanol. The sample was added to DPPH• and allowed to remain in equilibrium for 30 minutes. EPR signal of DPPH• was recorded after the equilibrium was reached and the g-value was measured and the free radical equivalent was calculated from the double integrated spectrum. The absorption intensity as obtained from the double integrated

spectrum was related to the free radical (DPPH[•]) concentration.

3. Results and Discussion

The product was collected at the end of each calcinations step and then, the physical and morphological observations were made. The intensity of the flesh color observed in the first stage of the sample S1 progressively increased with stages and reasonable red tinge in the fifth stage sample S5 was observed. The results of the analytical study can be discussed as follows.

3.1. Fourier-Transformed Infra Red Spectroscopic Analysis

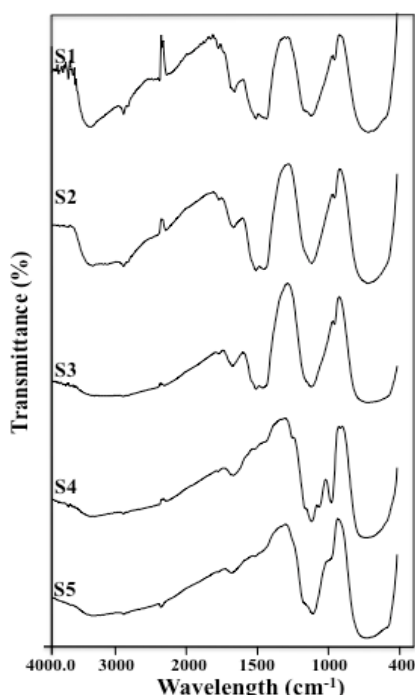


Figure 1. FT-IR spectrum of the nano tin (IV) oxide samples obtained during trituration of tin metal with five plant extracts followed by calcinations

Fig.1 illustrates the FT-IR spectrum of the samples S1 to S5. In all the samples, peaks were observed in the regions of $625 - 665 \text{ cm}^{-1}$ and $1040 - 1060 \text{ cm}^{-1}$ corresponding to O-Sn-O bridge of SnO_2 . The peak $1040-1060 \text{ cm}^{-1}$, a characteristic of Sn=O and Sn-O, indicates the presence of adsorbed oxygen species, intermediate between O_2^- and O_2^{2-} . Also there was an additional peak of low intensity in the region $1460 - 1470$ and $865 - 870 \text{ cm}^{-1}$ in the stages of I, II, and III indicated the presence of carbonates and it disappeared in stages of IV and V. The absence of peaks at 1450 , 1088 , and 850 cm^{-1} indicated the absence of inorganic carbonates [14]. In all the samples, except the final stage sample, there was a broad peak in the region

$3200 - 3500 \text{ cm}^{-1}$ and around 1640 cm^{-1} confirming the presence of physisorbed water. In the final stage sample the peaks due to -OH stretching frequency was not intense enough. The FTIR spectra confirmed the formation of SnO_2 from the first stage of preparation itself. The sharpness and intensity of the Sn-O-Sn characteristic peak indicated that SnO_2 was the major constituent in all stages.

3.2. Fourier-Transformed Raman Analysis

The FT Raman spectrum of the final stage sample is shown in the Fig.2. The small intensity bands observed at 650 cm^{-1} in the purified tin metal was shifted to the region of $625 - 630 \text{ cm}^{-1}$ in the calcined samples. The bands in the regions $628-630 \text{ cm}^{-1}$ assigned to A_{1g} mode of SnO_2 rutile structure. There was a small hump in the region $770-790 \text{ cm}^{-1}$ assigned to B_{2g} mode. There was absence of intense and broad band at 571 cm^{-1} in the sample, which was assigned to amorphous fraction of tin (IV) oxide as observed by [15].

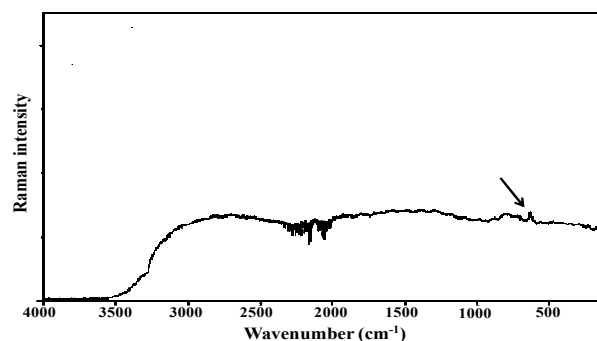


Figure 2. FT- Raman Spectrum of final stage sample of nano tin (IV) oxide obtained during the trituration of tin metal with five plant extracts followed by calcinations. (Black arrow indicates $630 A_{1g}$ mode)

3.3. ICP-OES and CNHS Analysis

Table 1 depicts the ICP-OES analysis of samples obtained from each stage (S1 to S5). It has been observed that there was a drastic reduction in the percentage of elemental tin. The samples of stage 1 (S1) showed 30% of Sn and upon progressing of stages; the final stage sample (S5) showed only 0.7% of Sn. It could be attributed to the maximum percentage of conversion of metallic tin to tin oxide during the preparations. The results further confirm that all the samples were free from toxic metals such as Pb, As, Hg and Cd. Many elements, which are therapeutically significant in relation to the targeted diseases viz:- Ca, K, Fe, Mg, Zn were introduced at the micro level. They should have been introduced by the various herbal extracts added during each of the calcinations steps. This is in contrast to the standard tin (IV) oxide or synthetically prepared nano tin (IV) oxides which are free from the above said elements.

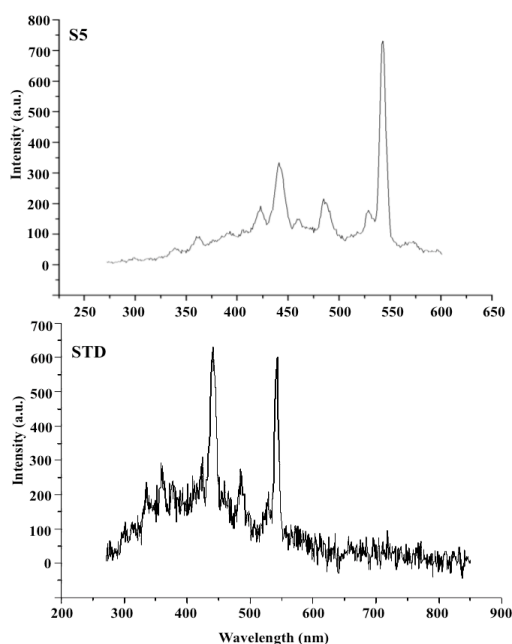
Table 1. Elemental composition of the samples of nano tin (IV) oxide obtained during the trituration of tin metal with five plant extracts followed by calcinations

Description of the samples	Percentage weight of elements												
	Sn	Ca	Cu	Fe	K	Mg	Ni	Zn	Si	C	H	N	S
S1	30	2.90	0.05	1.25	5.26	1.14	Nil	0.02	0.71	5.23	0.93	0.68	Nil
S2	30.81	2.50	0.05	1.12	2.72	0.99	Nil	0.01	0.55	2.92	0.68	0.58	Nil
S3	23.87	2.66	0.04	1.31	2.97	1.06	Nil	0.03	0.74	2.48	0.60	0.09	Nil
S5	0.72	4.25	0.04	1.40	1.9	1.32	Nil	0.07	0.24	0.15	0.14	0.3	Nil

The samples also demonstrated the presence of C and S in meagre percentage, as evidenced through C, H, N, S elemental analysis (Table 1) and as the calcination steps proceed, only a trace level of carbon, nitrogen and sulphur along with a fractional amount of hydrogen were introduced at the final stage of the processing. It is interesting to note that despite the five trituration steps of tin metal with different herbal extracts, the contribution of organic moiety was fractional and confirmed the effectiveness of the calcination steps followed.

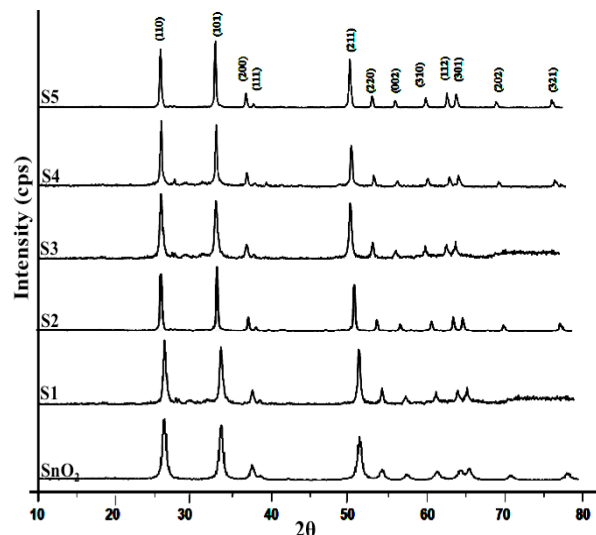
3.4. Photo Luminiscence Study

Fig. 3 depicts the photoluminescence spectrum of final stage sample. Two prominent peaks were observed at 440 and 542 nm when excited at a wave length of 295 nm. In addition, small intense peaks were also observed at 422 and 484 nm. The peak observed at 422 nm may be related to the origin of photo luminescence centre created due to oxygen vacancies. The presence of structural defect as observed by [16] due to these peaks can be well correlated with that of oxide.

**Figure 3.** Photoluminescence spectra of (S5) final stage sample of nano tin (IV) oxide obtained during the trituration of tin metal with five plant extracts followed by calcinations compared with that of (STD) commercial tin (IV) oxide

3.5. Powder X-ray Diffraction Analysis

The PXRD spectral patterns of the samples S1 to S5 were shown in Fig. 4. The intense peak positions of each sample agreed well with the reflections of tetragonal single phase of bulk SnO_2 - Casitterite -structure (ICDD 21-1250) and the absence of any other phase of tin or SnO and the absence of polymorphism in the final stage sample was confirmed. This observation suggested that SnO_2 was the major constituent in all the samples.

**Figure 4.** PXRD spectra of nano tin (IV) oxide samples obtained during the trituration of tin metal with five different plant extracts followed by calcinations

The interplanar distance 'd' values of the first and final stage sample of the nano tin (IV) oxide preparations were compared in Table 2. The formation of nano crystallites was quite evident from the broad PXRD peaks. The appearance of peaks along the phases (110), (101), (200), (220), (310), (301) and (301) with 2θ values of 26, 33, 37, 54, 61, 65, 71 degrees respectively also matched well with that of the chemically synthesized tin oxide reported by [17, 18,]. In the PXRD spectrum line broadening was observed and the crystallite size of the first stage and final stage sample (S1 to S5) in the most intense plane of 110, 101 was calculated by Sherrer equation as shown below after quadratic smoothing followed by Lorentian curve fitting of the peaks.

$$D = \frac{K\lambda}{\beta \cos \theta}$$

where K is the shape factor, λ is the X-ray wavelength (1.5406 \AA), β is the line broadening at half the maximum intensity (FWHM) in radians, and θ is the Bragg angle (Patterson, 1939), D is the mean size of the ordered (crystalline) domains, which may be smaller or equal to the grain size.

Table 2. Comparison of the 'd' values (from the PXRD analysis) of the nano tin (IV) oxide samples S1 & S5 obtained during the trituration of tin metal with five different plant extracts followed by calcinations

h,k,l plane	Standard d (\AA) values (ICDD) 211250	S 1	S 5
1,1,0	3.347	3.382	3.371
1,0,1	2.642	2.661	2.690
2,0,0	2.369	2.375	2.359
2,1,1	1.764	1.770	1.764
2,2,0	1.675	1.681	1.676
3,1,0	1.498	1.478	1.484
3,0,1	1.415	1.419	1.419

In the present preparation, there was grinding with the herbal extract for each step. These steps would have contributed towards the formation of nano crystallites. In terms of modern technology, it is a top down approach of nano synthesis while according to [19], the long duration of grinding resulting in particle size reduction as propounded by the conventional method of size reduction. It was intended for increasing the bio availability of the components and therapeutic efficacy of the medicinal formulation. The absence of peaks due to metallic tin can also be correlated with the total conversion of metallic tin to tin oxide. These results agreed well with the ICP-OES detection of 0.7 % of elemental tin. The lattice parameters obtained for the final stage sample (S5), were close to the standard tin oxide. Hence, it can be concluded that there was no doping of any element into the crystal lattice which speaks for the phase (tetragonal) homogeneity and purity of the sample. During the calcinations stages, the crystallite size varied from 29.2 to 25.7nm in the 110 plane and from 34.7 to 31 nm in the 101 plane. There was preferential growth in the 101 plane as indicated by the intensity of the peak. Even though the number of calcinations steps was five, there was size stabilisation at the final stage due to the presence of calcium ions.

3.6. EPR Spectral Analysis

The EPR spectra of the final stage sample S5 measured at room temperature. Fig.5 showed a broad peak and there was an interaction of both metal and oxygen with the microwave radiation. The signal was strong and hence there is a possibility of good amount of paramagnetic species.

The ' g_{\parallel} ' value of 2.759 and g_{\perp} was found to be 2.022 and suggested the non-stoichiometry present in tin (IV) oxide with singly ionized oxygen vacancies accounting for the paramagnetic property in agreement with the observations of [20].

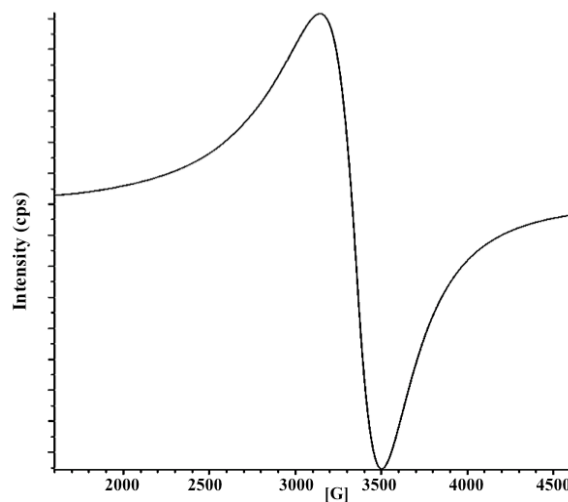


Figure 5. EPR Spectrum (at room temperature) of final sample of nano tin (IV) oxide obtained during the trituration of tin metal with five different plant extracts followed by calcinations.

3.7. Thermo Gravimetric Analysis

The thermograms of the samples S1 to S5 were given in Fig. 6. About 8% weight loss between 200 and 600 °C was observed for S1 could be attributed to the decomposition of organic matter as evidenced from the FTIR spectrum. Samples S2, S3 and S4 displayed only 4% weight loss and could be attributed to the physisorbed moisture. In the final stage sample (S5) there was no change in weight loss till 750 °C and suggested the thermal stability [21].

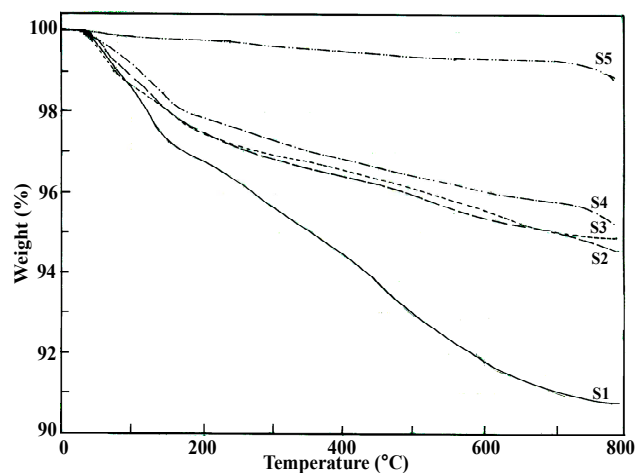


Figure 6. Thermo gravimetric analysis of samples of nano tin (IV) oxide obtained during the trituration of tin metal with five different plant extracts followed by calcinations

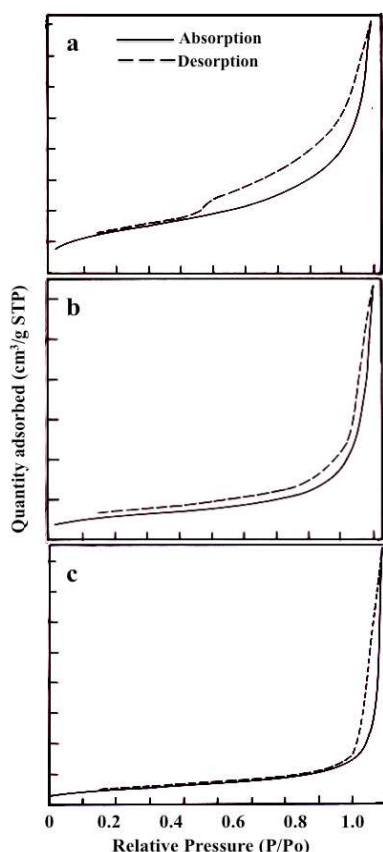


Figure 7. BET isotherms of a: Stage 1 (S1) sample. b: Stage 5 (S5) sample of nano tin (IV) oxide obtained during the trituration of tin metal with five different plant extracts followed by calcinations and c: Standard tin oxide sample

Table 3. Comparison of grain size and crystallite size (from the PXRD analysis) for sample S1 & S5 of nano tin (IV) oxide obtained during the trituration of tin with five different plant extracts followed by calcinations

Sample	Average crystallite size in nm	BET surface area m ² /g	Grain size in nm	Cumulative pore volume cm ³ /g
S1	29	9.1665	94.181	0.0253
S5	37	3.8862	222	0.0193
Standard SnO ₂ (present study)	>50	8.529	101	0.0100

The BJH pore size distribution as shown in Fig. 8a-8b. The cumulative pore volume and pore area were found to be 0.015 cm³/g and 0.0253 m²/g for S1 and for S5 sample. The pore distribution was wide as can be seen in the figure which reflects in the adsorption capacity of the final sample.

BJH pore radius varied over a wide range from 9.7 to 203 Å and 10.4 Å to 174 Å for S1 and S5 samples respectively. This technique characterises pore size distribution independent of external area due to particle size of the sample. The surface morphology as predicted by the above techniques was correlated with Transmission Electron Microscopic (TEM) images of the samples.

3.9. Scanning Electron Microscopic Images

The shape, size, morphology of the nano crystallites

3.8. BET and BJH Pore Analyses

BET isotherm studies were carried out for the initial and the final stage samples S1 and S5. Table 3 compares the stage I and stage 5 of the nano tin (IV) oxide samples and Fig. 7a and Fig. 7b illustrates the BET isotherm of stage 1 sample and final stage sample respectively. For S1 sample, BET isotherm corresponds to Type IV isotherm based on the IUPAC recommendations (1985) and there was a hysteresis loop associated with the capillary condensation taking place in mesopores. This type of isotherm can also be correlated with a higher BET surface area and nano crystallites compared to standard bulk tin (IV) oxide.

For the final stage sample S5, Type II BET isotherm was observed. The curve was first concave to the P/P₀ axis and then almost linear and finally convex to the P/P₀ axis. It indicated the formation of an adsorbed layer whose thickness increases progressively with increasing relative pressure until P/P₀ → 1. Similar observations were made with commercially available standard tin oxide (Fig. 7c).

Table 3 illustrated the BET surface area and grain size of samples of S1 and S5 and standard tin oxide. The average particle diameter in nanometre was calculated from the following formula;

$$6000 / (\text{BET surface area in m}^2/\text{g}) \times (\text{density in g/cm}^3)$$

There was a steep decrease in the BET surface area of the sample from first stage to the final stage in relation to their crystallite size calculated from PXRD. The presence of aggregates with inclusion of other elements getting into the surface as can be correlated with the SEM.

determine their applications in the field of electronics, medicine, chemical industry.[22]; The surface morphology of the samples S1 to S5 as depicted in Fig. 9. In the SEM images, the initial stage sample S1, showed small particles with grain boundaries, which also correlated well with the BET surface area report. Further stages of calcinations brought more agglomeration leading to bigger particle size in stages 2, 3, and 4 with relatively lesser crystallinity. In the final stage sample S5, there was disintegration of bigger crystallite particles so that there was distribution of various sizes of particles. The grain size of the samples as shown in the figure is in the order of micrometer suggestive of the agglomeration of nano sized crystallites formed

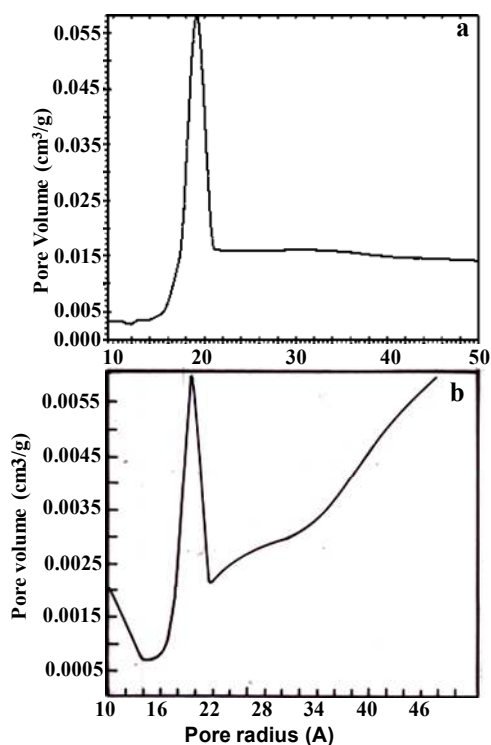


Figure 8. BJH pore volume distribution of sample S1 (a) and S5 (b) of nano tin (IV) oxide obtained during the trituration of metallic tin with five different plant extracts followed by calcinations

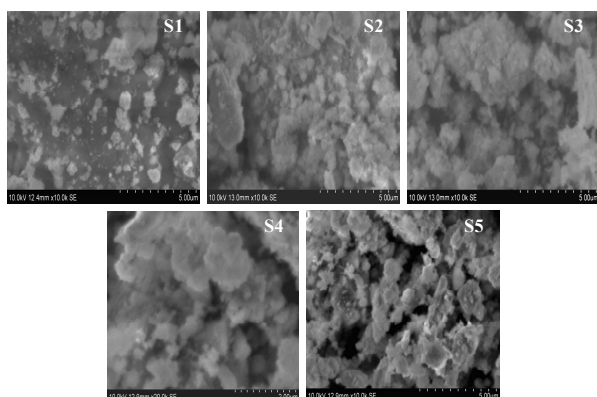


Figure 9. SEM images of S1, S2 S3, S4 and S5 stage samples of nano tin (IV) oxide obtained during the trituration tin metal with five different plant extracts followed by calcinations

3.10. Transmission Electron Microscopic Image

The TEM image of the fifth (final) stage sample Fig.10 indicated the presence of particles being composed of aggregates of irregular spherical crystallites. It was also indicative of the presence of bigger grains composed of the nano crystalline domains due to repeated calcinations. The surface composition as can be seen from the TEM image of the final stage sample showed the presence of elements of Calcium and Phosphorous, even though presence of Iron, Potassium, Magnesium were indicated by the ICP-OES results. There was oxygen deficiency indicative of non-stoichiometric defect in the crystal lattice.

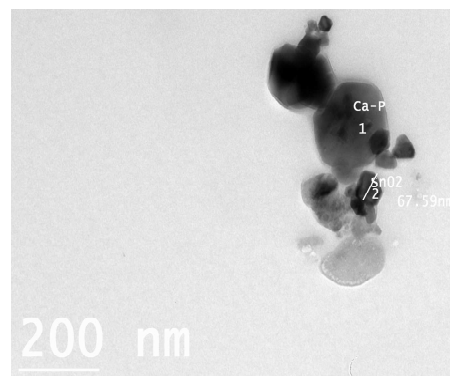


Figure 10. TEM image of S10 stage (Final stage) sample of nano tin (IV) oxide obtained during the trituration of tin metal with five different plant extracts followed by calcinations

3.11. X-ray Photoelectron Spectroscopic Analysis

According to [23], the surface state of the particles is determined by the type of calcinations process viz: presence of air, oxygen or any other carrier gas and the size of the crystallite is affected by the presence or absence of carrying gas. In the present study, the chemical state of tin and oxygen at the surface were examined by XPS. Fig.11 shows Sn 3d regions, O (1s) and wide scan of the samples S1, S3, S5 respectively.

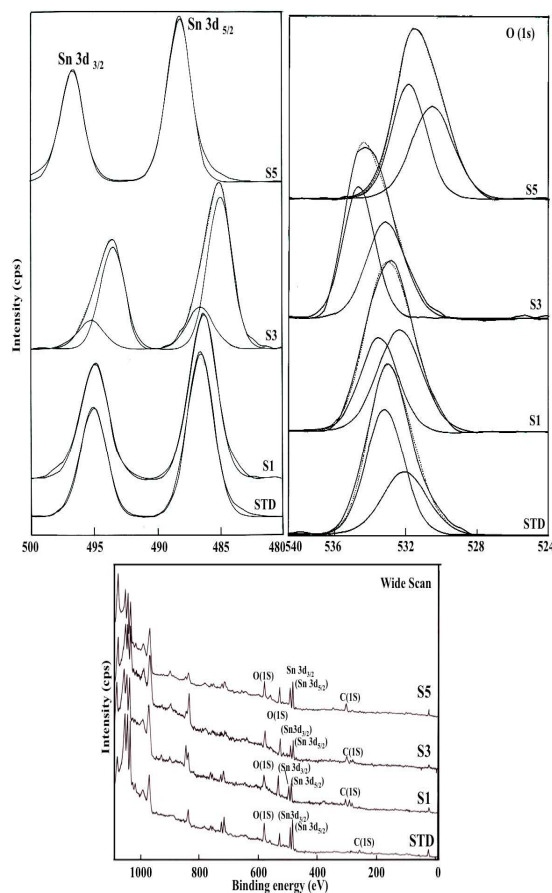


Figure 11. XPS spectrum, Binding energy curve of Sn(3d3/2 & Sn 3d 5/2 O(1s) and wide scan of S1, S2, S3, S4 & S5 stage samples of nano tin (IV) oxide obtained during the trituration of tin metal with five plant extracts followed by calcinations

Usually, O (1s) peak has been observed in the binding energy region of 529 to 535 eV with the peak from 529 to 530 eV which is attributed to lattice oxygen [24] attributed the peak in the region 530.7 - 531.6 eV to oxygen in non – stoichiometric oxides and the peak around 533 eV to a weakly bound surface oxygen.

Table 4. Elemental composition and binding energy values observed from XPS spectrum for stage1(S1) and stage5 (S5) samples of nano tin (IV) oxide obtained during the trituration of tin metal with five different plant extracts followed by calcinations

Sample	Element	Binding energy (eV)
Bulk SnO ₂	Sn(3d _{5/2} : 3d _{3/2})	486.6 : 495.0 (Δ=9.4)
S1	Sn(3d _{5/2} : 3d _{3/2})	486.4 : 494.8 (Δ=8.5)
S3	Sn(3d _{5/2} : 3d _{3/2})	486.0 : 495.2 (Δ=10.2)
S5	Sn(3d _{5/2} : 3d _{3/2})	488.2 : 496.8(Δ=(8.6)
Bulk SnO ₂	O(1s)lattice: adsorbed	530.8 and 531.9
S1	O(1s)	530.5 and 531.6
S3	O(1s)	529.9 and 530.9
S5	O(1s)	532.2 and 533.5

The binding energy values are sensitive to the covalent character of the metal oxygen bond and we observed a increase in both the Sn (3d) and O (1s) binding energy levels compared to the bulk tin oxide indicating smaller covalency and the absence of more electropositive elements.

3.12. Antioxidant Profile Using EPR Spectra

For antioxidant profiling of the sample, the free radical quenching behaviour was assessed using EPR spectroscopy. The free radical equivalent of DPPH[•] of the sample was calculated. The free radical equivalent of DPPH per mg of the sample was calculated as 0.8026. The result can be correlated with the capability of the product for medical purposes. In the present study the anti oxidant property of tin (IV) oxide present with the free radical quenching factor could play a major role in the antimicrobial property. It can also be correlated with the anti oxidant property of titanium oxide (also having tetragonal structure similar to tin oxide), which was responsible for the degradation of Reactive Oxygen Species (ROS) with immediate inflammatory response as observed by ^[25] Contreras *et al* (2007). The plant extracts provided alkaloids, cyclopeptides, flavanoids, phenols, tannins and minerals like Cu, Fe, Mn, Zn, Se which could have contributed towards therapeutics or a secondary effect for the anti oxidant property.

4. Conclusion

The research study provided a new top down approach for the synthesis of nano composite of tin (IV) oxide with multi dimensional applications. There was a natural

inclusion of significant elements like Ca, Mg giving a unique surface of the nano crystalline tin (IV) oxide in the parlance of modern nanotechnology. The impact of grinding with plant extracts repeatedly not only to reduce the particle size but also improved the bio availability. The analytical results of the study also clearly indicated the formation of nano crystallites of tin oxide with increasing crystallinity and preferred orientation in the (101) plane with proceeding steps of grinding and calcinations. The process provided a technique of preparing a multifunctional nano tin (IV) oxide with unique bulk and surface property.

Competing Interests

The authors declare that they have no competing interests.

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