Synthesis of nano-spherical nickel by templating hibiscus flower petals

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Abstract: In this study, we have represented synthesis of nano spherical nickel using porous Hibiscus rosa-sinensis petals as bio template. Synthesized Nickel nanoparticle thus characterized by SEM, TEM, XRD and wet chemical methods. Spherical Nickel particles of sized between 10nm to 200nm was observed.

Keywords: Nano Nickel, Catalyst, Bio-Template, Scanning Electron Microscopy, X-Ray Diffraction

1. Introduction

Nano-nickel has very high catalytic activity. Unprotected metal nano particle is prone to oxidation and thus conventional methods for the synthesis of metal nanoparticles are not very useful for various applications. As nano metal particle is very prone to oxidation and agglomeration, synthesis of suitably protected metal nanoparticle is found to be difficult and hence different methods are being attempted [1]. It is found that use of capping agents or coating with surfactants may resist the oxidation of nano metal particles. Synthesis of nano nickel particle (un-oxidised and non-agglomerated) is in very high demand as it has diverse applications. By changing the nickel content from 1 to 10 wt%, Lixiong Zhang et al. showed that nano nickel can be used for selective gas permeable membranes [2]. Dye separation performance of the nickel particles embedded ferromagnetic hierarchical porous carbon (FHPC) was investigated by Wang et al. [3]. Nano nickel powder exhibits a strong catalytic effect and can be used in hydrogenation of organic compounds, tail gas processing of vehicle, increase combustion efficiency of solid rocket, reduce sintering temperature etc. it is also useful as magnetically separable catalysts, catalyst supports, and gas adsorbents [4]. Although nano nickel particle has many potential applications, synthesis of these materials were mostly attempted through conventional chemical routes, which involves several steps and use of expensive chemicals including reducing agents, as a result, the widespread application of nano nickel was restricted[5]. Here in this study we have chosen hibiscus flower as soft and porous bio template as well as an in-situ reducing agent while decomposed at higher temperature. This method eventually could produce spherical nickel nano particle along with some mesoporous carbon.

There are several ways of synthesizing mesoporous carbon and nano nickel particles, however, in-situ formation of these materials are not well-established [6, 7, 8]. This paper, aims to report a very simple, cost effective and environmental hazard free green synthesis of spherical nickel nano particle by templating hibiscus flower petals as reducing material.

2. Experimental

Hibiscus flower (Hibiscus rosa-sinensis) was collected just before the dawn (when the flowers did not open up completely). Petals were separated out from rest of the flower parts and were very gently washed with distilled water and air dried at room temperature (~25°C) for ~4hrs. to create partial voids (by evaporation of moisture).

A solution of 0.5M NiCl2.6H2O, was added into partially air dried petals and allowed to soak for overnight (~16hrs.). Ni-salt soaked petals lift off from rest of the NiCl2.6H2O solution and were dried under IR lamp (~80°C) for 4hrs. The NiCl2 incorporated dried petals are placed in a graphite crucible and heated in graphite resistance furnace (ASTRO, Thermal Inc., USA) at 700°C under vacuum (~10^-5mbar) for 30mins. The rate of heating was maintained ~20°C/min. Low vacuum scanning electron microscope (TM1000,
Hitachi, Japan) was used to see the microstructure of green as well as the dried (~80°C) petals. The materials heated at 700°C were characterization by SEM-EDS, TEM, and XRD. The wt% of Ni in the sample was found out by weight chemical method.

3. Results and Discussion

In order to avoid the atmospheric contamination, flowers were plucked in the very early morning and in a fully matured bud, later it was allowed to grow (in few hours) into full flower under clean room condition. The appearance of the pyrolyzed product was black and gets attracted by magnet, thus confirms the presence of Ni particle. Metallic nickel present in the sintered product could not be separated by magnet even after the sintered product was washed, dried and vibrated ultrasonically. It was interesting to note that size of the cup like pores reduced substantially from top surface to inside of the petals, often there observed partitions within a cup (Fig. 1). As the parts of a petal were porous, the partitioning walls have micro or nano interconnected tunnels that help supply moisture, nutrients and other organic materials from the plant to all parts of the petal. The pores and tunnels beneath the pores have very high capillary action, we have utilised these pores and tunnels for adsorbing NiCl$_2$·6H$_2$O solution within it and interconnecting tunnels of the petals.

Initially when partially dried petals were dipped into the NiCl$_2$ solution, after gentle agitation (with a glass rod) for about 1-2 hrs., petals floats as submerged condition into the NiCl$_2$ solution and the colour changed from red to dirty purple.

This indicates that, NiCl$_2$ diffused into the pores and tunnels of the petals by osmosis phenomena. Microstructure of the bottom surface of the petals, practically does not exhibit porosities, rather it exhibits cellulosic structure (Fig. 2 a-c). Pore sizes of the top surface of the dried petals vary between 1 to 12µm (Fig. 1). If we consider a cup like pore having diameter of ~10µm (as marked in Fig.1), then it could be able to accommodate 6.23x10$^{-11}$ gm NiCl$_2$·6H$_2$O equivalent to 1.52x10$^{-11}$ gm Ni, when 0.5M NiCl$_2$·6H$_2$O solution was used (density of Ni is 8.9g/cm$^3$). Considering the shapes of the cup like pores as a sphere and consequentially the nickel particle formed out of it also spherical, then the final diameter of the spherical nickel formed within the pore should be ~740µm. Similarly 1µm diameter cup should yield 0.074µm or 74nm diameter Ni particle. Since it was difficult to measure the sub-surface pore size we could not calculate lower limit of the Ni particle size. SEM micrograph and EDS analysis (Fig. 3) as well as TEM micrograph and corresponding diffraction patterns (Fig. 4.) showed that there exist good amount of Ni nano particle of size that varies from 10nm to 700nm. Figure 3 exhibits the near spherical Ni were separated from each other and the inset showed that the Ni balls were entrapped within the pores and tunnels of the pyrolyzed petals. Between the Ni spheres, porous carbon prevents them from agglomeration.

It was also observed that, even when the heat treatment was carried out at 1000°C, Ni ball did not agglomerate.

During pyrolysis of the NiCl$_2$·6H$_2$O soaked petals, various organic constituents of the petals decomposed and released. As of yet, not much information have been reported about the decomposition behavior of hibiscus flower petals. Yong Wang et.al [8] reported that the chemical constitutions of pollen contain several components along with cellulose, and hemicelluloses;
thermal decomposition of pollen in N₂ atmosphere starts ~130°C, which ends at ~450°C.

![Figure 4. TEM micrograph shows the size of Ni varies from 10 to 200nm. In the inset corresponding selected area diffraction pattern is shown.](image)

Recently, Yu-Chuan Lin et al. many other researchers reported that pyrolysis of cellulose produces several products including CO/C₂H₄ (~12.7%), CO₂ (~19.3%) and decomposition is dependent on the heating rate [9]. Lanza et al. provided good information about the nature and kinetics of gas evolution when cellulose is pyrolyzed [10, 11, 12].

This carbon is considered to be mesoporous carbon and evolved from the hibiscus petals during pyrolysis at 700°C. According to them maximum amount of CO evolved at 250°C then sequentially C₃H₆, CH₄, CO₂ and the least is H₂. In the current study thus, either CO or C₂H₄ could have reduced the NiCl₂ to Ni during the pyrolysis of NiCl₂.6H₂O soaked petals. NiCl₂ could be reduced by CO at about 180°C to form Ni (CO)₄. Subsequently the Ni (CO)₄ could produce Ni. However, Ni (CO)₄ being highly volatile should go out from the system during the pyrolysis even much before 700°C. However it was evident from the SEM-EDS analysis (Fig. 3) that good amount of Ni was present at the surface of the pyrolyzed petals. Hence CO was not the reducing agent in this case. Whereas, C₂H₄ could directly reduce NiCl₂ to Ni. In that case one should expect CH₃CHO as one of the reaction product. In order to verify this proposal we did an experiment using a simple glass apparatus, the outlet of the apparatus was dipped into Tollens’s reagent. On heating the NiCl₂ embedded hibiscus petal, the exhausted gas reduced the Tollens’ reagent to form Ag mirror; on the contrary the petals without soaked in NiCl₂ could not reduce the Tollens’ reagent. Therefore, it can be concluded that C₂H₄, which is the pyrolyzed product of cellulose present in the petals reduced NiCl₂ to form Ni, thus follows the reaction below.

\[ \text{NiCl}_2 + 
C_2H_4 + H_2O \rightarrow \text{Ni} + \text{CH}_3CHO + 2\text{HCl} \]

XRD analysis of the pyrolyzed products confirms presence of only metallic Ni and carbon (Fig. 5). Weight chemical analysis confirms that ~37wt% of metallic Ni was present in the products and rest was mesoporous/amorphous carbon. XRD was taken 20 days after the pyrolysis, while material was kept in open atmosphere that means the metallic Ni did not oxidised at least during this period. This indicates that Nano nickel particle produced by this process was coated in situ by carbon so as to prevent its oxidation.

![Figure 5. XRD analysis of the pyrolyzed product shows that the sample contains only Ni and Carbon.](image)

4. Conclusions

Porosity of the Hibiscus rosa-sinensis petals could act as a bio template as well as a reducing material for synthesizing the nano nickel particle. Current research demonstrated that the controlled atmosphere pyrolysis of NiCl₂.6H₂O soaked petals could produce nano nickel particles in situ coated with carbon. Size of nano nickel particles varies between 10nm to 200nm.

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