Catalytic Properties of Hydrothermally Synthesized Flower-like NiO@Fe₃O₄

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DOI: 10.29130/dubited.721970

ABSTRACT

In this study, flower-like NiO structures were synthesized by hydrothermal method. Then, Fe₃O₄ was doped on NiO that NiO structures gain magnetic properties. TEM (Transmission Electron Microscopy) and SEM (Scanning Electron Microscopy) were used to assess the morphology of the NiO structures. XRD (X-ray Diffraction) was used to evaluate the crystal structures of the NiO structures. Magnetic properties of the NiO structures were investigated using VSM (Vibrating Sample Magnetometry). Catalysis properties of the produced structures were assessed use of products as catalyst in the reduction of 4-nitrophenol (4-NP) was examined. Especially NiO and Fe₃O₄ were found to exhibit excellent catalytic performance for nitrophenol reduction, while NiO@Fe₃O₄ magnetic microflowers were found to exhibit very low catalytic activity.

Keywords: NiO, Fe₃O₄, Flower-like, 4-Nitrophenol, Hydrothermal synthesis

Hidrotermal olarak Sentezlenen Çiçek benzeri NiO@Fe₃O₄'ün Katalitik Özellikleri

ÖZET

Bu çalışmada, çiçek benzeri NiO hidrotermal yöntemle sentezlendi. Daha sonra NiO üzerine, Fe₃O₄ katkılanarak manyetik özellik kazandırıldı. Elde edilen ürünlerin kristal yapıları XRD (X-ışını kırınımı) metodu ile incelenmiştir. Üretilen yapıların morfolojik özellikleri SEM (Tarama Elektron Mikroskopisi), ve TEM (Geçirimi Elektron Mikroskobisi ) ile karakterize edildi. Ayrıca manyetik ölçümleri için VSM (Manyetik Histeresis Ölçümü) analizi yapıldı. Daha sonra ürünlerin, 4-nitrofenolün (4-NP) indirgemesinde katalizör olarak kullanılabildiği incelendi. Özellikle NiO ve Fe₃O₄'nün nitrofenolün azaltılması için mükemmel katalitik performans sergilediği görülen NiO@Fe₃O₄ manyetik mikro yapıdaki çiçeklerin çok düşük katalitik aktivite sergilediğini belirledi.

Anahtar Kellimeler: NiO, Fe₃O₄, Çiçek benzeri, 4-Nitrofenol, Hidrotermal sentez
I. INTRODUCTION

Industrial and agricultural wastewaters contain nitrophenol, which is one of the most common organic pollutants. Elimination of nitro groups can be accomplished by reducing them to amines groups. Amines groups are less toxic and biodegradable [1,2]. 4-aminophenol (4-AP) is a powerful intermediate which can be used in the synthesis of drugs. 4-aminophenol is also widely used as a corrosion inhibitor, hair colouring agent and, etc [3,4]. Therefore, conversion of nitrophenol to aminophenol is essential for wastewater management where biodegradation of toxic substance can be achieved. It is stated that the main process for the reduction of the nitro group in general can be achieved by methods such as catalytic / electrochemical reduction of nitrobenzene in the presence of strong acids or hydrogenation of nitrobenzene, which involves molecular rearrangement of the phenylhydroxylamine intermediate in the presence of iron-acid [5-8]. In the presence of sodium borohydride (NaBH₄), 4-NP can be reduced to 4-AP in aqueous media [9-12]. However, this reduction reaction mechanism works slowly in the absence of a metal catalyst. For this purpose, various metal nanoparticles have been used as catalysts to reduce nitrophenols to aminophenol.

Pt, Ag, Pd and Au nanoparticles are very effective in catalysts reactions and they are commonly used as nanocatalyst [1,13-17]. Noble nanoparticles attract the attention of researchers since they have outstanding properties such as small size, high electron affinity, quantum confinement effect, good magnetic characteristics, etc. Besides, noble nanoparticles exhibit exceptional catalysis properties. Such properties are surface related properties which can be seen on nanoparticle surface. For example, only the electrons at the surfaces of these nanoparticles are used in catalytic reactions where most of the atoms in the nanoparticles are catalytically inactive during the catalysis reaction [3]. Therefore, enhanced surface - volume ratio results in enhanced catalysis activity. Traditionally, flower-like metal oxide nanomaterials have attracted attention in a variety of fields, including catalysis, electronic and analytical sciences, due to their rough surface with large surface / volume ratios [18-21]. Many researchers are willing to use transition metal oxide nanoparticles instead of noble metal nanoparticles. They consider that transition metal oxide nanoparticles are highly reactive, they can be cheaply synthesized and can be an alternative to noble metal nanoparticles [21-23].

Nanocatalysts are quite small hence, it is difficult to separation the nanoparticles from the reaction pot. At this point, nanocomposites are found to be promising to overcome this issue. Nanocomposites are nanoparticle clusters that have magnetic properties can be fabricated. Consequently, such nanocatalysts with magnetic properties can be separated from the reaction pot with a strong magnet. Such a case eliminate the need for physical filtration where catalysts can be lost in the filtration process. Materials with superparamagnetic characteristics are suitable candidate to fabricate such nanoparticle clusters.

In this study, NiO, Fe₃O₄ and NiO@Fe₃O₄ samples were prepared using the hydrothermal method. Various techniques were used in the sample characterization. In the presence of NaBH₄, catalysis performance of metal oxide compounds with flower-like morphology was evaluated. Flower-like NiO and Fe₃O₄ were found to exhibit high catalytic activity.

II. MATERIALS AND METHODS

A. MATERIALS, SPECTRAL DATA MEASUREMENTS, AND SYNTHESIS PROCEDURE

All the chemical reagents used in the experimental process are analytical grade pure. No purification process was applied to the chemicals. NaOH (sodium hydroxide), Ni(ac)₂.H₂O (nickel acetate tetrahydrate), urea (CO(NH₂)₂), FeSO₄.7H₂O iron(II) sulphate heptahydrate) were purchased from Sigma-Aldrich.
XRD patterns were recorded using the RIGAKU miniflex600 X-ray diffractometer has a scan range of 10 < 2θ < 90. Also, the surface morphology of the nano structures was assessed using a Field Emission Scanning Electron Microscopy (FESEM). In the confirmation of elemental composition, EDX (Energy Dispersive X-Ray) spectrometer was used. FEI Quanta 400F model device was used in FESEM-EDX investigation. TEM studies were carried out on a JEOL JEM 1220electron microscope. Cryogenic Limited PPMS VSM was used in the magnetic measurements which has the maximum magnetic field of 2±Tesla. Shimadzu UV-2600 Ultraviolet-Visible (UV-Vis) spectrophotometer was used in the catalytic study.

**A. 1. Synthesis of Flower-like NiO**

0.5 g (Ni(ac)$_2$.4H$_2$O) and 0.25 g CO(NH$_2$)$_2$ were dissolved in 30 mL ethanol. Mixture was stirred using magnetic stirrer. The homogeneous mixture was then poured in a teflon-lined stainless steel 50mL autoclave. Mixture was kept in oven for 12h at 190 °C. Teflon autoclave was cooled at room temperature. The product was centrifuged to get the precipitate. The precipitate was washed with ethanol and deionized water. The washed precipitate was dried in the oven for 24h at 60 °C. Green powder was obtained and calcination procedure was applied to the powder in an oven for 1 hour where powder was heated with a ramp rate of 10 °C / minute up to 400 °C. Result product was then cooled to room temperature [26].

**A. 2. Synthesis of Magnetic Fe$_3$O$_4$ Nanostructure**

Hydrothermal method was used in the production of magnetic nanoparticles. To this end, 0.656 g FeSO$_4$.7H$_2$O and 3.2 g NaOH were dissolved in 40 mL deionized water. The solution was mixed for 10 minutes until it homogenizes, then 50 mL of teflon autoclave covered with stainless steel was placed and left at 150 °C for 6 hours. The resulting black precipitate was collected by centrifugation at 8000 rpm for 20 minutes. The product was washed several times with deionized water and absolute ethanol. Powder like precipitate was treated in vacuum oven at 40 °C for 12 hours [27].

**A. 3. Synthesis of Magnetic NiO@Fe$_3$O$_4$ Nanoflowers**

0.1 g Ni(ac)$_2$.4H$_2$O was dissolved in 10 mL deionized water using ultrasonic bath. In a separate beaker, 0.13 g FeSO$_4$.7H$_2$O was dispersed in 10 mL deionized water. 0.64 g NaOH was added and mixed for 10 minutes. The prepared solution was then poured onto Ni(ac)$_2$.4H$_2$O. The ultrasonic bath was mixed for a further 5 minutes, a stainless steel coated 40 mL capacity teflon autoclave was taken and kept at 150 °C for 6 hours. The resulting black precipitate was collected by centrifugation at 8000 rpm for 20 minutes. The product was washed several times with deionized water and absolute ethanol. Powder like precipitate was treated in a vacuum oven at 40 °C for 12 hours.

**A. 4. Catalytic Test**

In a typical test, freshly prepared NaBH$_4$ solution (0.2 M, 0.3 mL) was added to 3 mL of 0.1 mM aqueous 4-NP solution. It was scanned on the UV spectrometer between 250nm - 550 nm wavelength. 3 mg of catalyst (NiO, Fe$_3$O$_4$ and NiO@Fe$_3$O$_4$) were added to the mixture and stirred for 1 min at room temperature (25 °C). Reduction of 4-NP to 4-AP was tracked using UV-Vis spectrophotometer.
III. RESULTS AND DISCUSSION

A. CHARACTERIZATION OF NiO, Fe₃O₄ AND NiO@Fe₃O₄

In Figure 1, XRD pattern of flower-like NiO, Fe₃O₄ NiO@Fe₃O₄ structures were presented. In the XRD pattern of NiO, three strong peaks with high density were seen which illustrates (111), (2 0 0) and (2 2 0) crystal orientation. Peaks indicate that prepared NiO is cubic crystal structure [28]. When XRD patterns are examined, sharp peaks prove the pure and well crystallized NiO products that are obtained by hydrothermal synthesis method. When the XRD pattern of Fe₃O₄ magnetic nanoparticles was examined, it was seen that the characteristic peak positions found for the spinel structures in the Fe₃O₄ structures. It has been found that Fe₃O₄ samples are pure, in other words, they are not contaminated, and they do not contain trace of other elements [27,29]. NiO@Fe₃O₄ magnetic nanoflowers were obtained by NiO with Fe₃O₄. XRD pattern of NiO@Fe₃O₄ magnetic nanoflowers revealed that the diffraction peaks that both nano structures have remained. Both peaks observed at the pattern of NiO and peaks observed at the XRD pattern of Fe₃O₄ were seen at the XRD pattern of NiO@Fe₃O₄. Such a case is an indication that both NiO and Fe₃O₄ structures protect their structures which indicate that NiO@Fe₃O₄ structure is in composite form.

![Figure 1. XRD patterns of NiO, Fe₃O₄ and NiO@Fe₃O₄](image)

The morphology and microstructure of the samples were analysed by field emission scanning electron microscopy (FESEM). Figure 2 shows low and high magnification FESEM images of flower-like NiO, Fe₃O₄ and NiO@Fe₃O₄ structures. NiO's nano-bowls and nano leaves (nanosheets) were found to turn into well-packed and self-organizing rose-like flowers. Figure 2(e) shows that the Fe₃O₄ magnetic nanoparticles have relatively uniform octahedral morphology. The average diameter of the flower-like NiO was found to be approximately 2.5-3 μm. In addition, FESEM images of NiO@Fe₃O₄ was also obtained. Images confirm that the flower-like structure was preserved. Figure 2(i) shows the high resolution TEM micrograph of single nanosheet of NiO@Fe₃O₄. TEM results are quite consistent with FESEM observations.

Figure 3(c) shows the EDX spectrum of the prepared samples. The presence of Fe atoms in the EDX spectrum of NiO@Fe₃O₄ is a proof that Fe₃O₄ is bound to the NiO microparticle.
Figure 2. Low and high magnification FESEM images of as-synthesized NiO (a,b), Fe$_3$O$_4$ (d, e), NiO@ Fe$_3$O$_4$ (g,h) and TEM images of the flower-like NiO (c), Fe$_3$O$_4$ (f), NiO@ Fe$_3$O$_4$ (i)
Figure 3. EDX spectrum of NiO, Fe₃O₄, NiO@Fe₃O₄

Figure 4 shows the magnetization curves of the Fe₃O₄ and NiO@Fe₃O₄ nanoparticles. It was determined that the Fe₃O₄ magnetic nanoparticles obtained after six hours had a superparamagnetic character at room temperature. At this temperature, the saturated magnetization values for the Fe₃O₄ nanoparticle were determined to be 89.8 emu / g. Due to the high crystalline properties of Fe₃O₄ nanoparticles, saturated magnetization value was found to be quite high. Sahin et al. reported the saturated magnetization value for Fe₃O₄ nanoparticles as 52.2 emu g⁻¹[30]. Saturated magnetization values of NiO@Fe₃O₄ o found as 21.5 emu / g. They also show super paramagnetic characteristics. Mahmoud et al synthesized Fe₃O₄@NiO core-shell nanocomposite by precipitation method to investigate Cr (VI) adsorption efficiency. The saturated magnetization value of the synthesized nanocomposites was found to be 11.5 emu / g for their samples [31]. It is understood that our samples have higher magnetic saturation values than that of results reported in the literature.
B. CATALYTIC ACTIVITY OF NiO, Fe₃O₄ AND NiO@Fe₃O₄

In the presence of NaBH₄, the catalytic activities of the samples were investigated for the reduction of nitrophenol to aminophenol. For all experiments, initial concentrations were reacted with 0.1 mM 4-NP and 0.2 M NaBH₄, respectively. The bounded peak was seen at UV-Vis spectra at 317 nm which is the indication of 4-NP. After the addition of NaBH₄, the peak then shifted to 400 nm region. This peak illustrates the formation of the p-nitrophenolate ions which was affected from the NaBH₄ [32, 33]. Addition of 3 mg of NiO, Fe₃O₄ and NiO@Fe₃O₄ nanoparticles result in the simultaneous growth of new peaks which were appeared around 295-300 nm. Such appearance confirms the existence of 4-AP which results in a sudden diminish in the intensity of the peak at 400 nm (See Fig. 5).

When 3 mg of NiO catalyst was added to the medium, the reduction process was observed to occur in 55 minutes, while it was found that this time was reduced when using Fe₃O₄ (38 minutes). Mandlimath and Gopal were found to be completed in about 4 hours in the presence of 0.8 mg commercial NiO catalyst [20].

The efficiency of the reduction reaction using our catalytic samples was investigated where produced samples were added to the reaction medium. To evaluate the reaction efficiency, linear graphs of ln (At / A₀) against time were obtained. The reaction rate was assessed using the diminish in the intensity of 4-NP peak in the UV-Vis spectrum. Figure 5 (d) shows the relation between the absorption peak density and time where absorption peak density was illustrated in natural logarithm form. The slope of the plot was used to calculate the rate constant of the reduction reaction. The rate constant (k_app) of the NiO, Fe₃O₄ and NiO@Fe₃O₄ compounds were found to be 1.2x10⁻³, 1.4x10⁻³, 4x10⁻⁵ s⁻¹, respectively. Jiang et al. synthesized nickel nanoparticles in varying sizes and morphologies that was used in the reduction process. All the nickel samples were compared to commercial RANEY Ni. When using RANEY Ni catalyst, the reaction rate constant (k_app) was found to be 0.32x10⁻³ s⁻¹ and showed lower value to other Ni nanoparticles [34].

In the presence of NaBH₄, NiO and Fe₃O₄ exhibited the best catalytic activity for the reduction of 4-NP to 4-AP. The catalytic performance of the NiO@Fe₃O₄ was found to be low compared to NiO and Fe₃O₄ nanostructures. It was also seen that NiO@Fe₃O₄ managed to reduce 20% of product to 4-AP in 70 min.
Table 1. Results obtained for the reduction of 4-NP ([4-NP] = 0.1 mM, [NaBH₄] = 0.2 M)

| Catalyst (3 mg) | Time (min) | K_{app} (s⁻¹) | Conversion (%) | R²  |
|----------------|-----------|---------------|---------------|-----|
| NiO            | 55        | 1.2x10⁻³      | 97.55         | 0.9981 |
| Fe₃O₄          | 38        | 1.4x10⁻³      | 98.76         | 0.9748 |
| NiO@Fe₃O₄     | > 70  <   | 4x10⁻⁵        | 20.17         | 0.9824 |

Figure 5. UV-Vis spectra obtained in the p-nitrophenol reduction in the presence of NiO (a), Fe₃O₄ (b) and NiO@Fe₃O₄ (c) and rate constants of the reaction (d).

IV. CONCLUSION

NiO, Fe₃O₄ and NiO@Fe₃O₄ samples were successfully produced using the hydrothermal techniques. The chemical and physical characteristics of the samples were illuminated using various spectroscopic and imaging methods. Flower-like NiO specimens were found to be cubic crystalline, while Fe₃O₄ specimens were in face centred cubic (fcc) crystal form. It was seen that NiO nanoparticles gained magnetic properties with Fe₃O₄ doping. The use of the samples as a catalyst in the reduction process of 4-NP was examined. It is known fact that noble metals are frequently used in this reduction process. It was found that the metal oxide catalysts samples, which were used in this work, showed as high performance as noble metals. The electron transfer mechanism existing on the surface of the noble metals affect the performance of the reduction process. Since our samples has flower like architecture,
electron transfer to can be existed in the inner surface samples. As a result, it was illustrated that NiO, Fe$_3$O$_4$, NiO@Fe$_3$O$_4$ nanostructures showed outstanding catalytical properties. Their catalytic performance is as good as noble metal nanostructures reported in literature.

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