

Characteristics of iron oxide rust prepared by peracetic acid and its removal of heavy metals in water

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Abstract. Fe₂O₃ is an especially promising material for water purification as it shows high heavy metal adsorption capacity. However, the high cost of commercial Fe₂O₃ makes it difficult to be widely used in developing countries. Herein, we probe the heavy metal removal performance of iron oxide rust. Rust was grown on iron nails in a controlled manner using peracetic acid (CH₃CO₃H), a safe and environment-friendly oxidizer. Arsenic was selected as an example of a heavy metal contaminant in this study. XRD and EDS analysis revealed that the iron oxide prepared with peracetic acid was nearly amorphous Fe₂O₃. Amorphous iron oxide is reported to show higher reactivity than crystalline iron oxide. The BET specific surface area of prepared Fe₂O₃ is 71 m²/g, which is larger than that of commercial Fe₂O₃, and the average pore diameter is 73 Å. Oxidized nails are highly effective for removing heavy metals: about 90% of 1ppm arsenic in water was removed at the residence time of 20 minutes, and the removal rate of 90% is maintained after 10 back-to-back arsenic removal experiments at the same residence time. Iron oxide prepared in this study can remove, per 1 cm², up to 0.114 mg of arsenic.

1 Introduction

Groundwater contamination by heavy metals such as Cd, As, and Cr is one of the biggest public health concerns in developing countries, e.g. Laos, Bangladesh, Cambodia, and Vietnam. Although a myriad array of materials has been studied for heavy metal removal, metal oxides such as Fe₂O₃ are especially promising as they show high heavy metal adsorption capacity [1]. However, the high cost of Fe₂O₃ makes it difficult to be widely used in developing countries. Therefore, there is a currently a growing demand for appropriate technology for heavy metal removal.

Herein, we probe the heavy metal removal performance of iron oxide rust formed on the surface of Nails. Rust was grown on iron nails in a controlled manner using Peracetic acid (CH₃CO₃H), a safe and environment-friendly oxidizer [2]. Arsenic was selected as an example contaminant in this study as it is typically present in the water of South Asian countries.

Specific research objectives are determined the optimal condition for iron nail oxidation, characterize the iron oxide formed on the nail surface, and evaluate the heavy metal removal capacity of oxidized nails in water samples.

2 Materials & Methods

2.1 Procedure for the Oxidation of Nails

Commercial iron nails were oxidized using peracetic acid, CH₃CO₃H. Peracetic acid is an environment-friendly oxidizer because it decomposes to acetic acid and hydrogen peroxide, both of which are biodegradable. Iron nails were 1 inch long and was about 0.57 g per nail. The geometric surface area of a nail was about 1.7 cm².

The surface oxide layer of the nail was removed using a sandpaper, and the nail was rinsed with deionized water. Cleaned nails were oxidized by submerging them in an aqueous solution with 10wt% peracetic acid and NaCl, which was added to improve conductivity. In order to compare different oxidation methods, nails were also oxidized with nitric acid. After oxidizing nails for a desired amount of time, nails were taken out of oxidizing solutions and rinsed with copious amounts of deionized water to stop to oxidation process, after which they were dried in air.

2.2 Characterization of rust on oxidized nails

The surface structure of oxidized iron nails was studied with an optical microscope and SEM (JEOL 7610F). EDS was used to analyse the change in atomic ratios before and after the oxidation reaction. Moreover, XRD (Panalytical Xpert) and BET (Micromeritics 2010) were used to study the crystallinity of the rust, and the specific surface area and pore diameter, respectively.

2.3 Arsenic removal using rust on oxidized nails

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Aqueous arsenic solution was prepared by dissolving sodium arsenate ($\text{Na}_2\text{HAsO}_4 \cdot 7\text{H}_2\text{O}$, Sigma-Aldrich) at 1 ppm, which is 100 times the standard for drinking water. A device for heavy metal removal was constructed using a glass column with an internal diameter of 34 mm and a length of 400 mm packed with oxidized nails. Arsenic solution was flown through the column and arsenic removal rate was measured as a function of residence time. The concentration of arsenic in the solution was measured using ICP (Leeman Labs).

We also wanted to evaluate the long-term performance of the heavy metal removal device. We measured the arsenic removal rate after 10 repeated experiments at a constant residence time. To quantify the maximum arsenic adsorption per surface area of iron oxide, oxidized nails were submerged in a 10ppm arsenic solution and arsenic removal rate was measure after 12 hours.

3 Results

3.1 Change in surface structure as a function of oxidation time

The iron oxide layer at various oxidation times was observed using an optical microscope (Fig. 1). After 30 minutes of oxidation, islands of iron oxide have been formed, but after 1 hour, most of the iron surface was covered with the oxide layer but it was not uniform. After 2 hours the oxide layer has completely covered the surface and no further oxidation was observed. After 8 hours some of the oxide layer began to peel off, and after 24 hours it was severely damaged. Together, this suggests that 2 hours is the optimum duration of oxidation reaction.

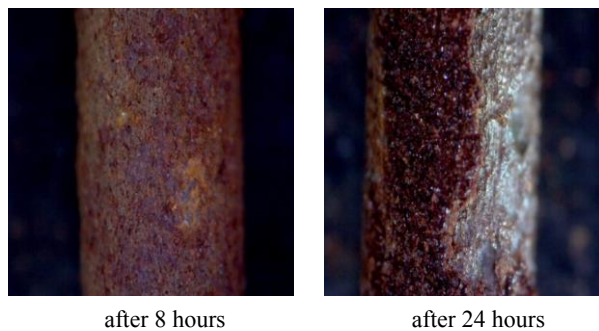
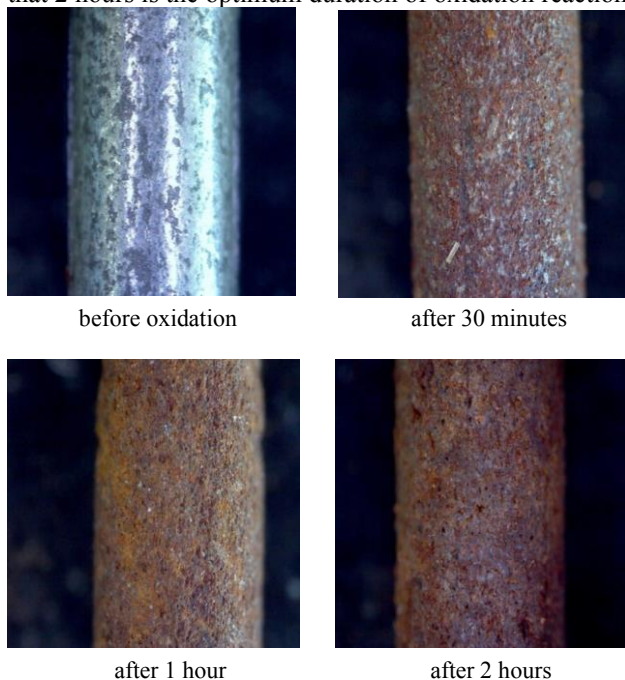


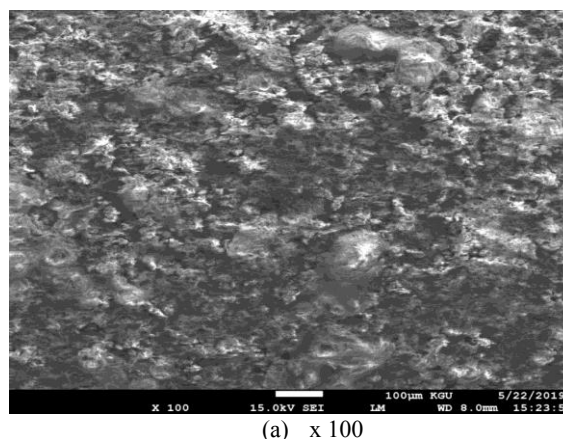
Fig. 1. Optical microscope images of nail surfaces at various oxidation times.

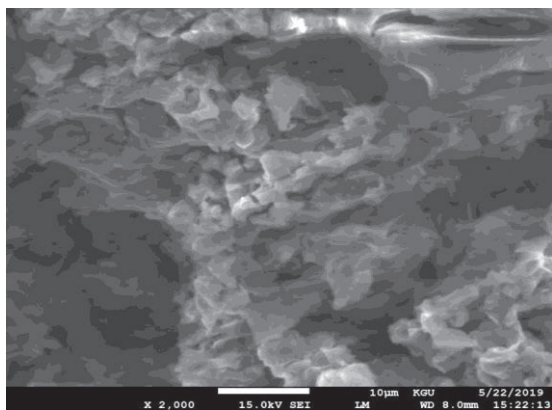
3.2 SEM analysis

SEM images of iron nails oxidized with peracetic acid for 2 hours were taken. As shown in Fig. 2, the oxide layer is uniformly form on the surface in the 100x image, and oxide crystals and pores of various sizes are observed in the 2000x image.

3.3 EDS analysis

EDS analysis results of nails before and after oxidation, as well as after arsenic adsorption are shown in Fig. 3. Before oxidation with peracetic acid (Fig. 3a), the iron nail surface contains about 11% oxidation due to oxidation in air. Carbon was detected, presumably due to coating material on iron nails designed to prevent corrosion. Figs. 3b and 3c were taken after oxidation to form iron oxide; the atomic ratio of O increased. For both sets of data, the Fe:O ratio is 1:1.47 on average, suggesting that the identity of the iron oxide formed is Fe_2O_3 (Fe:O = 1:1.5). Cl detected in Fig. 3b is from added NaCl in the oxidation solution, and arsenic can be detected in Fig. 3c due to adsorbed arsenic on iron oxide after arsenic removal experiment.

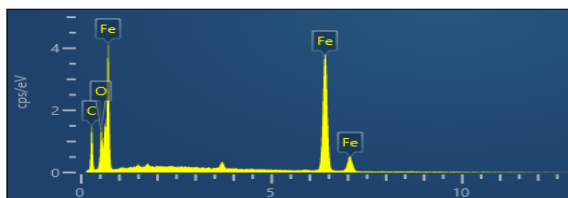




(b) x 2000

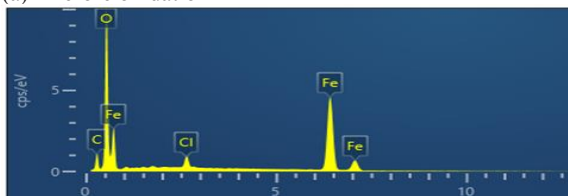
Fig. 2. SEM images of iron oxide on oxidized nail surface.

(above x 100, below x 2000)



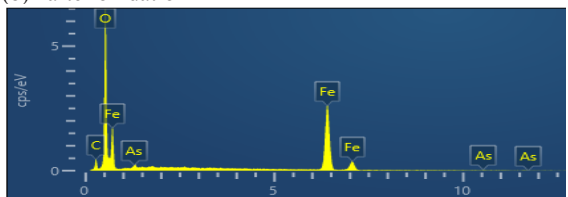
Element	Weight %	Atomic %
C	17.94	46.38
O	5.79	11.23
Fe	76.27	42.40
Total:	100.00	100.00

(a) Before oxidation



Element	Weight %	Atomic %
C	9.45	22.02
O	25.71	44.97
Cl	1.82	1.44
Fe	63.02	31.57
Total:	100.00	100.00

(b) after oxidation



Element	Weight %	Atomic %
C	6.40	15.45
O	27.88	50.56
Fe	64.54	33.53
As	1.19	0.46
Total:	100.00	100.00

(c) after arsenic removal

Fig. 3. EDS analysis of iron oxide on nail surface

To study the distribution of Fe, O and as on nails, EDS mapping was conducted on oxidized nails after arsenic removal experiments (Fig. 4) Fe and O atoms are distributed over the entire surface. Notably, O atom were the most prevalent, and adsorbed as was observed over the entire surface. Together, these suggest that iron oxide was uniformly formed on iron nails and that arsenic is removed evenly on iron nails without preference for specific sites.

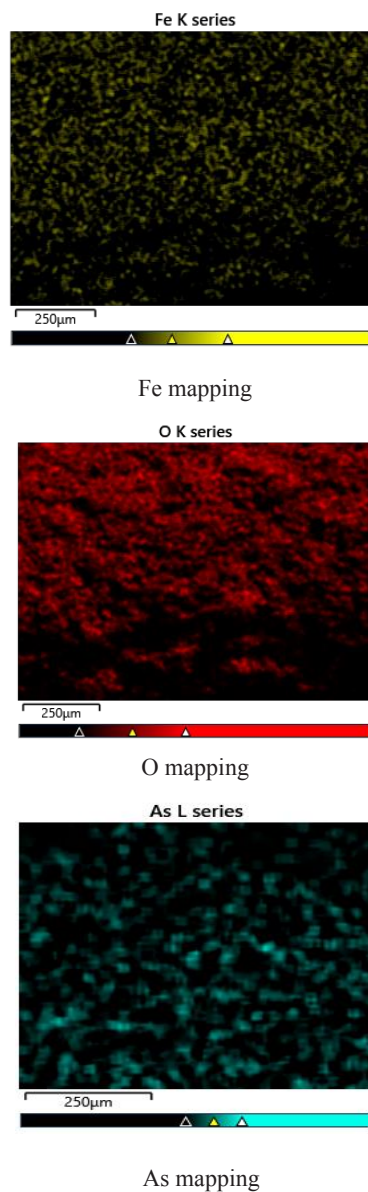


Fig. 4. EDS mapping image of oxidized iron after arsenic removal

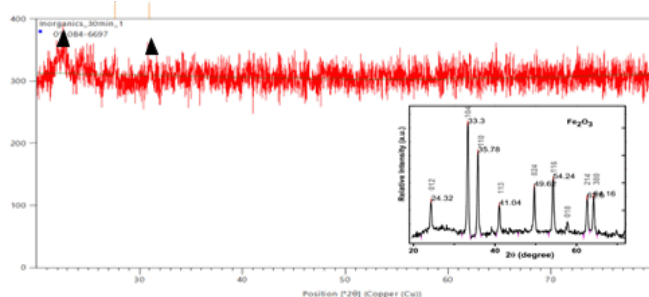
3.4 XRD analysis

Fig. 5 (a) shows the XRD analysis of iron oxide formed using peracetic acid. Iron oxide is mostly amorphous, which is to be expected since most compounds formed in aqueous solution are amorphous. However, weak crystal peaks are observed near $2\theta=24^\circ$ and 33° , which are indicated as \blacktriangle , which corresponds to the Fe_2O_3 crystal peak (see small figure).

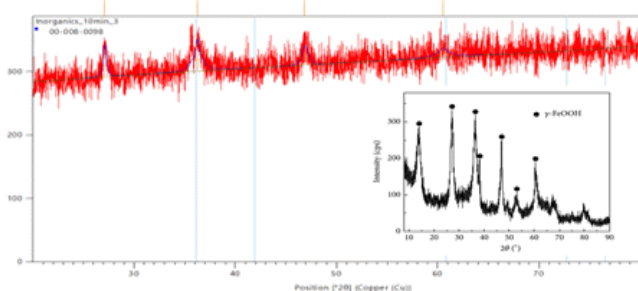
The results of the study on the removal of heavy metals of amorphous Fe_2O_3 have not been reported yet, but Pierce reported that the removal efficiency of heavy metals using amorphous $Fe(OH)_3$ was higher than that of crystalline $Fe(OH)_3$ [3]. One of the reasons for the high removal efficiency is that amorphous iron oxide has a larger specific surface area than crystalline iron oxide. made of acetic peroxide has a wide specific surface area.

Fig. 5 (b) shows the XRD pattern of iron oxide made with nitric acid for comparison. The iron oxide made with nitric acid is $FeO(OH)$ and has a low crystallinity. Saha reported that the heavy metal removal rate of amorphous $FeO(OH)$ was higher than that of other iron oxides [4].

3.5 Surface area and Pore size distribution analysis



(a) iron oxide made by peracetic acid



(b) iron oxide made by nitric acid

Fig. 5. X-ray diffraction (XRD) of iron oxide on nails.

The isotherm graph was shown at Fig. 6. The specific surface area of rust calculated from the BET equation was $71\text{ m}^2/\text{g}$. Compared to the commercially sold Fe_2O_3 powders with a specific surface area of about $10\text{ m}^2/\text{g}$, Fe_2O_3 made of acetic peroxide has a relatively large specific surface area. The pore types from the shape of the desorption curve and hysteresis are plate or rod, and the pores of micro-size and meso-size are present together [5].

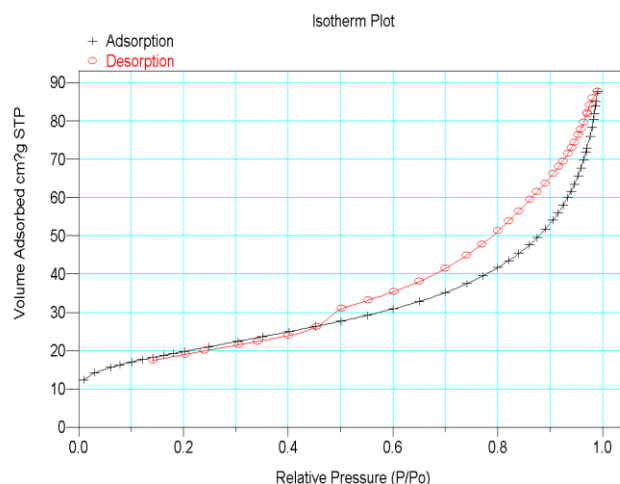


Fig. 6. Isotherm plot of Fe_2O_3 on nails.

Pore size distribution graph was shown at Fig. 7. The average pore size calculated from the BJH equation is $71\sim 75\text{ \AA}$, and from the dA/dD graph, there are few pores with a diameter of 100 \AA or more and relatively many pores with a diameter of 40 \AA .

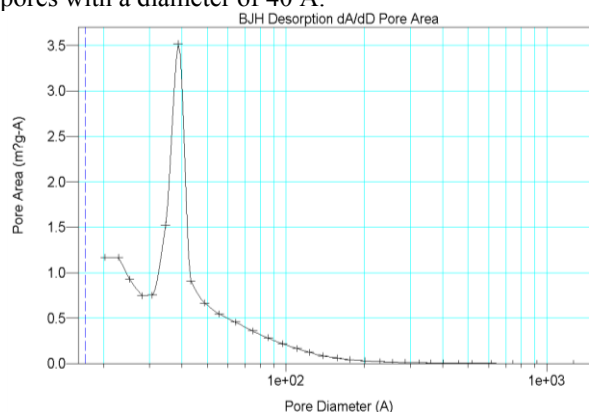


Fig. 7. Isotherm adsorption/desorption plot and BJH dA/dD plot of iron oxide rust

3.6 Arsenic removal in water using rust of oxidized nails

3.6.1 Removal of arsenic with residence time

The results of the experiment for removing arsenic with a concentration of 1 ppm using rust made of acetic peroxide and nitric acid were shown in Fig. 8.

At 5 min, the arsenic removal rate of rust made from acetic peroxide was 83% and the arsenic removal rate of rust made from nitric acid was 45% . At 10 min, the arsenic removal rate of rust made from nitric acid was 84% and 65% , at 20 min, 87.4% and 89.4% , at 30 min, 90.4% and 92.5% , at 40 min, 91.7% and 94.3% .

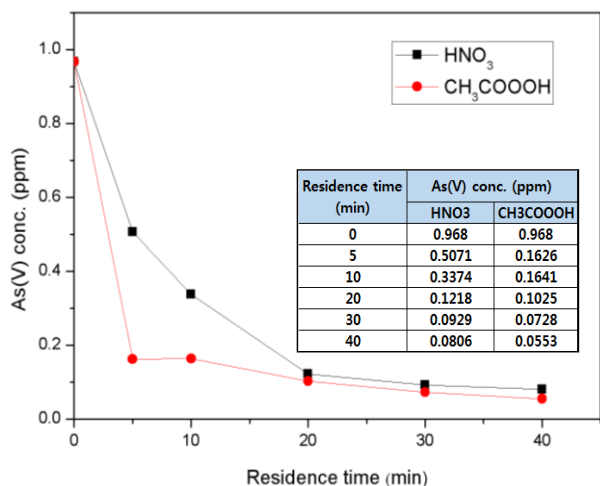


Fig. 8. Arsenic removal as a function of residence time

The arsenic removal rate of amorphous Fe_2O_3 rust made with acetic peroxide was higher than that made with nitric acid. As the residence time increase, the removal rate of arsenic also increases, but the increasing rate is low at 20 minutes or more. It is a very meaningful results that 83% of arsenic removal rate was obtained by treating 1ppm arsenic solution for 5 minutes with rusted nails.

3.6.2 Change of arsenic removal rate with repeated experiment

The results of the experiment were shown at Fig. 9 after 10 times of arsenic removal experiment with the initial arsenic concentration at 1.014ppm for 20 minutes. The removal rate of arsenic was between 87.9% and 93.3%. This shows that the efficiency of arsenic removal has not been reduced even in 10 repeated experiments considering the experimental error, and the nail can still adsorb arsenic.

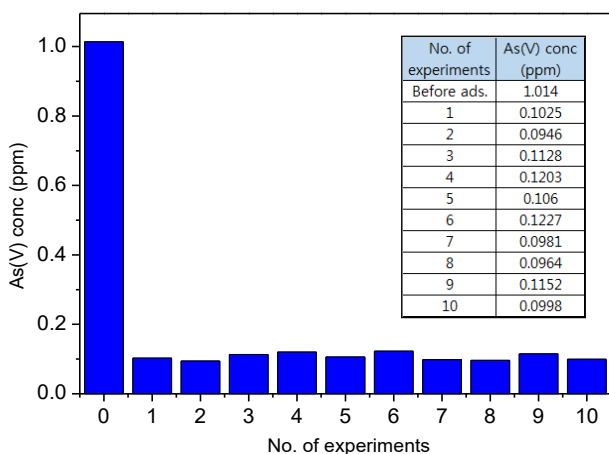


Fig. 9. Ten repeated experiments for Arsenic removal

3.6.3 Maximum arsenic adsorption measurement

To design arsenic removal device, it is necessary to know the maximum amount of arsenic adsorption per unit area of the rusted nail. To achieve this, arsenic concentration was measured after 12 hours with two oxidized nails with peracetic acid in a 50 ml aqueous solution of 8.91 ppm arsenic concentration. The arsenic concentration was decreased from 8.91ppm to 1.17ppm (Fig. 10). From this result, the maximum amount of arsenic that can be removed per rusted nail is 0.194mg, and the amount of arsenic that can be removed per unit area is 0.114mg.

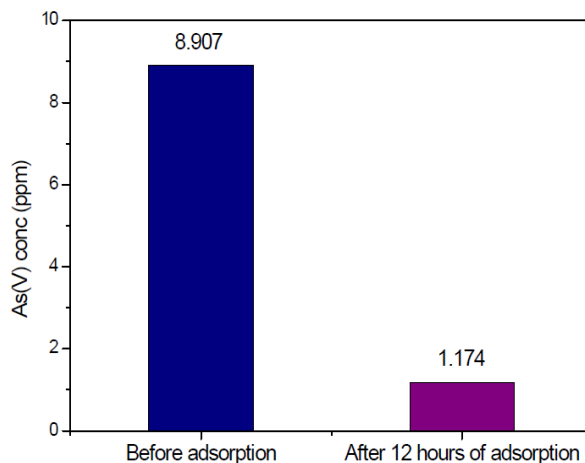


Fig. 10. Maximum amount of arsenic adsorption.

4 Conclusions

- (1) Fe_2O_3 was easily synthesized on iron nails using peracetic acid. As prepared Fe_2O_3 was amorphous and had a large specific surface area of $71 \text{ m}^2\text{g}^{-1}$ and a average pore diameter of 73 \AA .
- (2) Rust of oxidized nails are highly effective for removing heavy metals: 90% of arsenic in water was removed at the residence time of 20 minutes.
- (3) Oxidized iron nails are robust and maintain their removal rate of 90% after 10 back-to-back arsenic removal experiments at the residence time of 20 minutes.
- (4) Each oxidized nail removed up to 0.194 mg of arsenic, sand iron oxide prepared in this study can remove, per 1 cm^2 , up to 0.114 mg of arsenic.

References

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