

Studies on Nitration of Phenol over Solid Acid Catalyst



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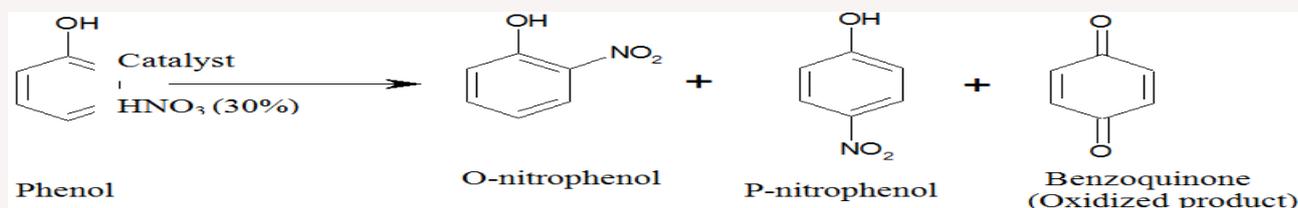
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Graphical Abstract

Phenol is nitrated selectively by dilute nitric acid over alumina and a remarkable ortho-selectivity is observed in liquid phase nitration.



Highlights

- The solid acid catalyst (Gamma-alumina) was prepared by control precipitation method.
- Phenol was selectively nitrated in liquid phase using dilute nitric acid (30%) in presence γ -alumina.
- The nitration reaction was carried out at room temperature and produce ortho-nitrophenol.
- The activation energy of the reaction was 11.272 KJ/mol.
- The kinetic study was also observed.

Abstract

Phenol was selectively nitrated in liquid phase to produce ortho-nitrophenol using dilute nitric acid (30%) at room temperature in presence of hydrochloric acid treated γ -alumina. Initially Al(NO₃)₃ and NH₄HCO₃ were reacted to prepare Al(OH)₃ which on successive calcinations at 550 °C for 5h produce γ -alumina. The γ -alumina was characterized by BET, XRD, SEM and NH₃-TPD analysis. The XRD profile confirmed the crystalline structure of the solid acid catalyst γ -alumina. The NH₃-TPD analysis showed the development of lewis acidity on the surface of hydrochloric acid treated γ -alumina. The effects of various parameters such as concentration of reactants, types of catalyst, weight of the catalyst, solvent, temperature and time of reaction have been studied. The kinetics of the reaction was also investigated.

Keywords: Gamma alumina catalyst; Nitration reaction; Ortho-nitrophenol; Rate constant

Introduction

Nitration of aromatic substances has great industrial importance because nitro-aromatics are extensively used as raw material for manufacturing dyes, pharmaceuticals, perfumes and plastics [1,2]. It is found from the previous study that the nitration of phenol using a mixture of nitric and sulfuric acid gives 2-nitrophenol and 4-nitrophenol in a ratio of 1.42 [1]. The limitation of this process lies in the use of corrosive liquid mixture of nitric and sulfuric acid which produce large amounts of wastes. The other problems are the chance of over nitration, poor selectivity and high cost of the process etc. [3]. To overcome these difficulties a new and simpler technique such as solid acid nitration should be employed. In

the solid acid nitration process the solid acid is used as a catalyst instead of sulfuric acid to produce nitronium ion. It is found from various literature that the nitration of phenol was carried out using different catalysts such as hydrous zirconia [4], tetra butyl ammonium bromide [5,6], sulfated titania [7], N₂O₅-ClNO₂ [8], p-Toluenesulfonic acid [9], ZSM-5 Zeolite [10], sulfated titania [11] and SnO₂ [12]. The selectivity of the product is increased by using proper solvent with solid acid catalyst. In the last few years various nitration procedures have been performed using different aromatic solvents such as N-nitropyrazole/ BF₃, Zn(NO₃)₂•6H₂O/2,4,6-trichloro-1,3,5-triazine in acetonitrile [13] and ionic liquid [14].

On the other hand alumina has extensive applications in organic reactions, either as an active participant or as an inert support. Alumina-based solid acid catalysts are potentially attractive due to the easy removability of substrate-product and easy recycling. The ortho-nitrophenol which is a major product of nitration reaction used for synthesis of valuable compounds. Some of these methods have contributed significantly for enhancing the scope of nitration reaction but very little work has been done on the region selectivity of phenol over alumina as a catalyst [15].

In the present work a suitable solid acid catalyst was investigated for nitration of phenol to ortho-nitrophenol. The region selectivity of phenol was studied using dilute nitric acid as nitrating agent over alumina as solid acid catalyst. The selectivity of the reaction is increased by using carbon tetra chloride as a solvent with the catalyst. The effects of different process parameters such as type of catalyst, weight of catalyst, time, temperature, reactant concentration, type of solvents and HCl treatment were observed. The kinetic study was also performed.

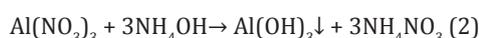
Materials and Methods

Materials

Aluminium nitrate ($\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$, 95%) of GR grade, ammonium bi carbonate (NH_4HCO_3 , 98%) of GR grade, acetone (CH_3COCH_3), carbon tetrachloride (CCl_4), methanol, sodium hydroxide (NaOH) and phenol ($\text{C}_6\text{H}_5\text{OH}$) were procured from Merck (India) Ltd., Mumbai, India. Ethanol ($\text{C}_2\text{H}_5\text{OH}$) was supplied by Jiangsu Huaxi International Trade Co. Ltd., China and nitric acid (HNO_3) was procured from S. d. fine Chem Ltd. Hydrochloric Acid (HCl) was procured from Sarabhai M Chemicals.

Catalyst preparation

Gamma alumina was prepared using control precipitation technique. Initially 15.42gm of ammonium bicarbonate and 40gm of aluminum nitrate was dissolved separately in 600ml of de-ionized water. The solutions were added drop wise to 400ml de-ionized water in a reaction vessel (Eq. 1). The reaction vessel was placed in a water bath and the temperature was maintained at 70 °C. The mixture was stirred for 30 minutes. The pH of the solution was adjusted by using HNO_3 or NaOH to precipitate Al cations in the form of hydroxide (Eq. 2). Then the precipitate was aged at 70 °C for 3hours and washed thoroughly with warm de-ionized water, ethanol followed by acetone to avoid contamination of Na ions. The washed material was air-dried at 120 °C for 12hours. After air drying the precipitate was calcined in a programmable furnace at 550 °C for 5hours. The various reactions involved in the preparation of alumina are given bellow.



The catalyst activation

Alumina powder was activated using hydrochloric acid (HCl). It was soaked for 30hours in de-ionized water and different percentages of HCl at room temperature. Then the material washed thoroughly with distilled water and dried overnight at 120 °C. The sample was then calcined at 350 °C for 2hours in presence of oxygen.

Catalyst characterization

The characterization of the catalyst was done by using the BET surface area analyzer (AUTOSORB-1C- USA), scanning electron microscope (Hitachi model SU-70), chemical composition of the catalyst, NH_3 -TPD (Quantachrome Instruments) and x-ray diffractometer (Philips PW183).

Reaction procedure

Nitration of phenol to orthonitro phenol was performed in a three necked round bottom flask type glass reactor (6.5cm I.D) with a capacity of 250cm³. The reactor was equipped with a four-leg vertical baffle (2.0cm diameter) which is located at a height of 1.5cm from the bottom of the reactor. Two gram of phenol was taken in a flask containing 3ml carbon tetrachloride. Then 1g of alumina and 4ml of dilute nitric acid (30%) were added into it. The reaction mixture was stirred at room temperature for varying period of time to study the progress of the reaction. The products were analyzed by HPLC((Column -Zorabax C-18 Reverse phase, Column Length-250mm, Column Dia-4.6mm, Pump-(Perkin Elmer, series 200, Isocratic pump), Detector-(Perkin Elmer, series 200, UV/VIS Detector), Mobile Phase- 50% Methanol / 50% Water, Flow Rate- 1.0ml/min, Wavelength-254nm).

Kinetic study

The kinetic study was performed in the same glass reactor as above. The reaction mixture was agitated for 180minutes and samples were collected at different time intervals. The experimental data were analyzed by pseudo first and second order kinetics mechanism. Although the reaction is carried out heterogeneously, but a homogeneous reaction mechanism with a second-order kinetics is assumed to interpret the experimental data. The isothermal reaction under constant-volume conditions results in the rate equation of the following form

$$-r_A = \frac{dC_A}{dt} = kC_A C_B \quad (5)$$

Where A is limiting reactant phenol and B is excess reactant nitric acid.

After integrating equation (5) the final form is represented as equation (6).

$$\ln \left[\frac{(M - X_A)}{M(1 - X_A)} \right] = C_{A0}(M - 1)kt \quad (6)$$

Where C_{A0} and C_{B0} represent the initial concentrations of reactants A and B respectively and k is the reaction rate constant.

$$M = \frac{C_{B0}}{C_{A0}} = \text{the initial molar ratio of the reactants}$$

Arrhenius equation

Arrhenius equation describes the effect of temperature on activation energy E and the frequency factor A . The value of E can be calculated from the rate constants at three different temperatures (T). The temperature dependency of reaction rate, k , is given by the Arrhenius equation,

$$k = Ae^{-E/RT} \quad (7)$$

The rate constant k can be obtained from equation (6). It is an average of values calculated at various experimental kinetic data points (i.e., reactant concentrations at various reaction times).

Result and Discussion

Characterization of the prepared catalyst

Table 1: Characteristics of different catalysts.

Sample	SBET (m ² /g)	Pore Volume (cc/g)	Pore Size(nm)
Alumina	220	0.216	219
Acid treated alumina	294.1	0.157	326

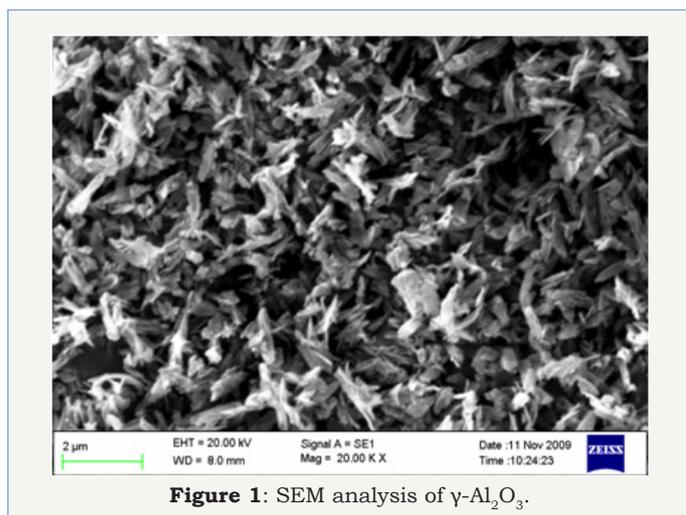


Figure 1: SEM analysis of γ -Al₂O₃.

The surface area, pore volume and the average pore size of alumina and acid treated alumina were analyzed in BET surface area analyzer. The values are shown in Table 1. The scanning electron microscope (SEM) image (Figure 1) of the alumina catalyst indicates strong agglomeration of alumina particles. The scanning electron microscope (SEM) image also confirms the formation of a gamma phase (γ -Al₂O₃) on alumina catalyst. Quantitative as well as qualitative composition analysis of prepared alumina was determined by EDS spectrum obtained from SEM. The elemental analysis of prepared alumina shows that it contains 59.42% O and 40.58% Al. The NH₃-TPD spectra of the γ -alumina powder and acid treated γ -alumina powder is shown in Figure 2. It is seen that the TPD spectra (Figure 2) of the γ -alumina powder contain intense peaks in the temperature range of 100-800 °C. These peaks can be assigned to the NH₃ desorbed from the acidic sites with medium to high strengths. Therefore, it can be concluded that the acid treated γ -alumina powder possesses strong surface acidic sites as compared to the γ -alumina powder. So the acid treatment improves the surface

area and surface acidity of γ -Al₂O₃. Thus, the resulted γ -Al₂O₃ gave more lewis acidity and acid density. The x-ray diffraction (XRD) profile of the γ -Al₂O₃ confirms spinal lattice structure of the catalyst (Figure 3). The peaks indicate the amorphous nature of γ -Al₂O₃. The crystallite sizes calculated by using the Scherrer equation were found to be in the range of 4.5-5.5nm.

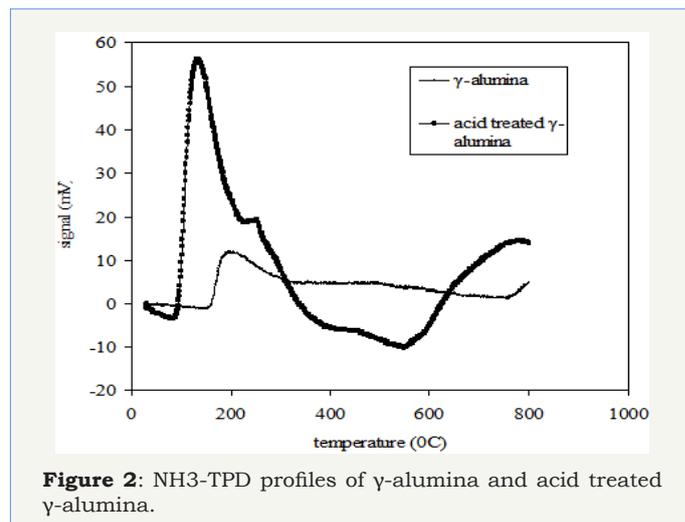


Figure 2: NH₃-TPD profiles of γ -alumina and acid treated γ -alumina.

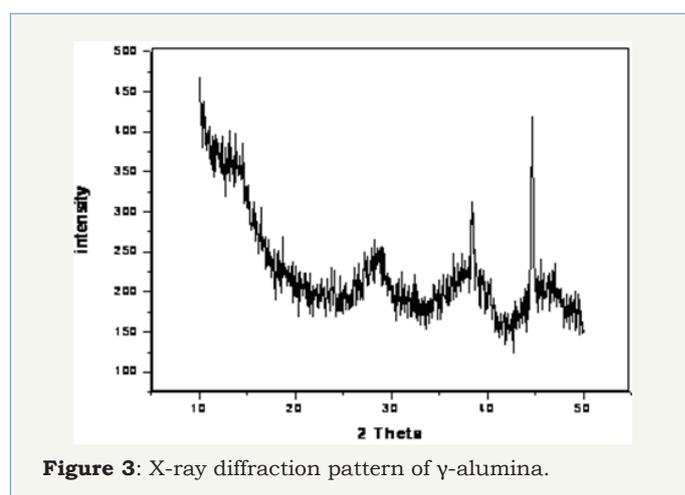


Figure 3: X-ray diffraction pattern of γ -alumina.

Selection of solvent

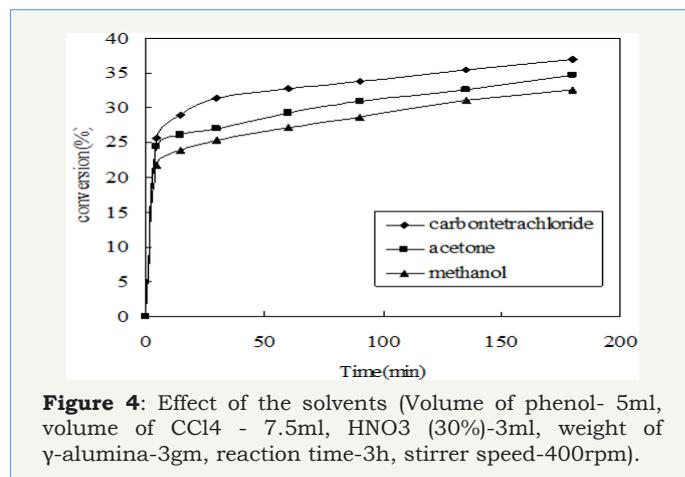


Figure 4: Effect of the solvents (Volume of phenol- 5ml, volume of CCl₄ - 7.5ml, HNO₃ (30%)-3ml, weight of γ -alumina-3gm, reaction time-3h, stirrer speed-400rpm).

To select a suitable solvent the nitration of phenol was carried out at room temperature in presence of carbon tetrachloride (CCl_4), methanol and acetone using γ -alumina catalyst. The reaction was carried out using dilute nitric acid (30%). The highest conversion of *o*-nitrophenol was obtained by using carbon tetrachloride as a solvent (Figure 4). The time required for completion of the reaction is also less in case of CCl_4 . Therefore CCl_4 is selected as a suitable solvent for the further nitration study.

Selection of catalyst

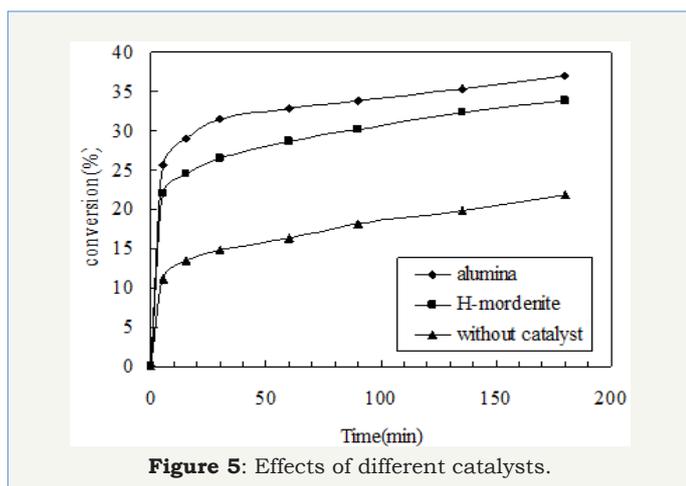


Figure 5: Effects of different catalysts.

The nitration of phenol was carried out separately in presence of γ -alumina and H-mordenite. The reaction was carried out at room temperature for three hours. For this purpose 5ml phenol and 3ml HNO_3 (30%) were taken and 3g of catalyst was added into it. The CCl_4 (7.5ml) was used as a solvent and the highest yield of *o*-nitrophenol was obtained using alumina catalyst. The yield of *o*-nitrophenol was decreased when the catalyst was changed from alumina to H-mordenite. The conversion of *o*-nitrophenol was low in this case because of less number of acid sites as well as porosity of H-mordenite. The reaction was also carried out in absence of catalyst and the conversion was further decreased, resulting nearly equal amount of ortho- and para- isomers. The effects are shown in Figure 5.

Effect of HCl treatment on alumina catalyst

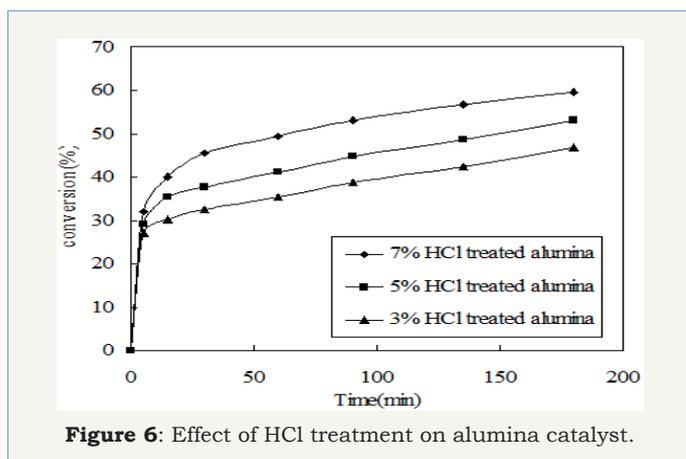


Figure 6: Effect of HCl treatment on alumina catalyst.

The catalytic activity of alumina was enhanced by treating it with different percentage of hydrochloric acid. The catalytic activity is increased with increase in surface acidity and was found to be in the following order 7% HCl treated alumina > 5% HCl treated alumina > 3% HCl treated alumina. The effects are shown in Figure 6. The other reaction conditions such as type of solvent, amount of reactants, reaction time and temperature were remain same as before.

Effect of the reaction time

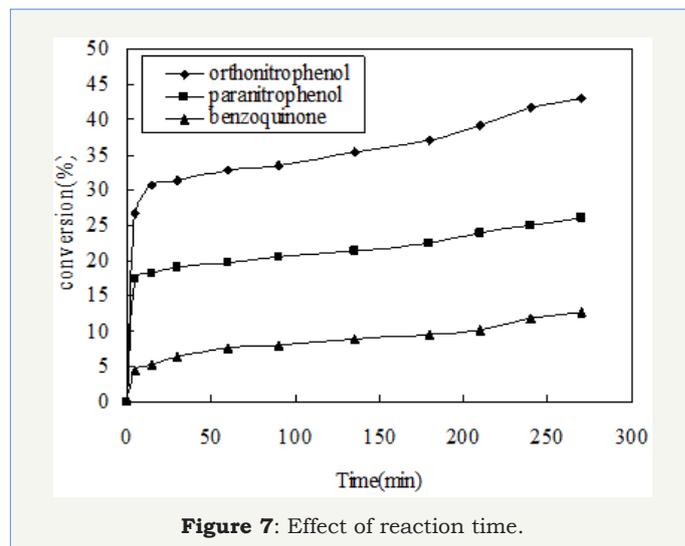


Figure 7: Effect of reaction time.

The nitration of phenol was carried out by varying reaction time in the presence of alumina catalyst (3gm) at room temperature. The reaction time was varied from 5-270min. In this case phenol (5ml) and dilute nitric acid (3ml) were used as reactants and CCl_4 (7.5ml) was used as a solvent. The results indicate that the yield of *o*-nitrophenol increases with increasing reaction time because more number of nitronium ions are formed as the time proceeds, resulting higher conversion of phenol (Figure 7). Therefore the maximum yield of *o*-nitrophenol was obtained at 4.5h.

Effect of weight of the catalysts

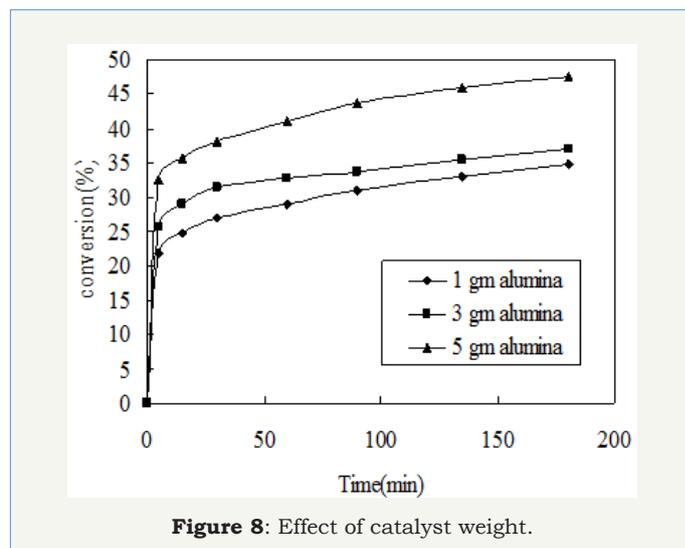


Figure 8: Effect of catalyst weight.

The conversion of o-nitrophenol also depends on the weight of catalyst. The nitration of phenol was carried out using dilute nitric acid (3ml) using different weight of alumina catalyst at room temperature. The effects are shown in Figure 8. The percent

conversion of phenol increases with increasing weight of catalyst due to availability of more surface area. The reaction speed is also enhanced with increasing catalyst dose. Therefore the highest conversion was obtained using 5gm of catalyst.

Effect of reactant concentration

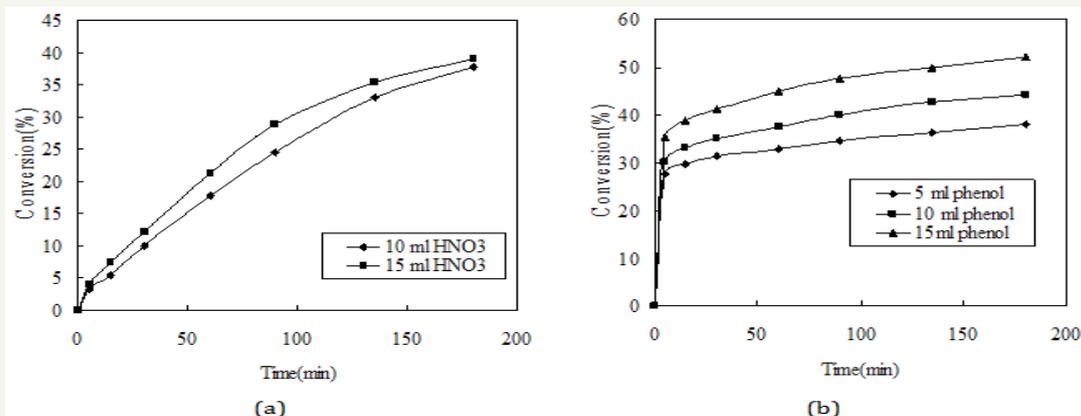


Figure 9: Effect of reactant concentration (Volume of CCl₄-7.5ml, HNO₃ (30%)-3ml, reaction time= 3h, reaction temperature-room temperature and stirrer speed-400rpm).

The effect of concentration of phenol and nitric acid is studied on the rate of reaction and the effects are shown in Figure 9. As the concentration of reactants (phenol and nitric acid) is increased, the rate of reaction also increases.

Kinetic Study

Table 2

Temperature (OK), T	1/T (K ⁻¹)	k (L mole ⁻¹ sec ⁻¹)	lnk
303	0.0033	10.83333	2.382628
318	0.003145	13.91667	2.633087
333	0.003003	16	2.772589
353	0.002833	20.75	3.032546

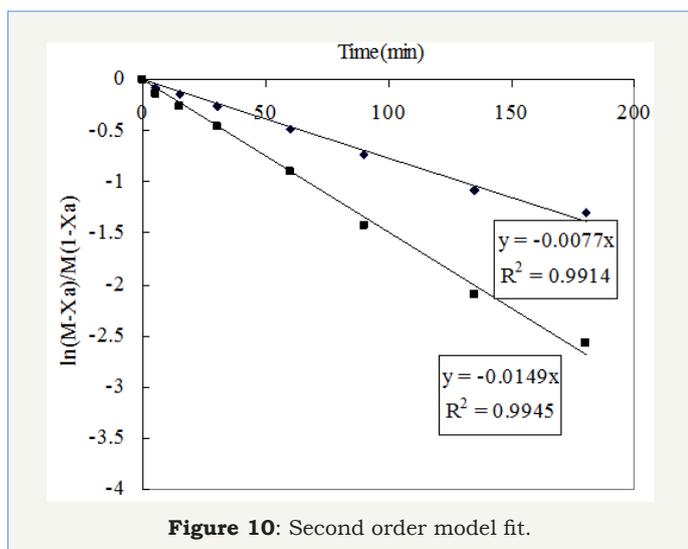


Figure 10: Second order model fit.

The pseudo second order rate expressions were taken into

consideration to correlate the kinetic data. Here the second order rate model (Equation 6) was fitted with data of nitration reaction (Figure 10). The kinetic study was carried out by varying the amount of nitric acid from 10-15ml. The other reaction conditions are kept as before for both the cases. It was observed that the kinetics of nitration for both the cases obeys the pseudo second order rate equation. The values of kinetic parameters are shown in Table 2. It can be seen from the following figures that for two different volume of nitric acid (10 and 15ml) the values of rate constant is 5.839 and 6.353L mole⁻¹ sec⁻¹ respectively.

Study of the Arrhenius Equation

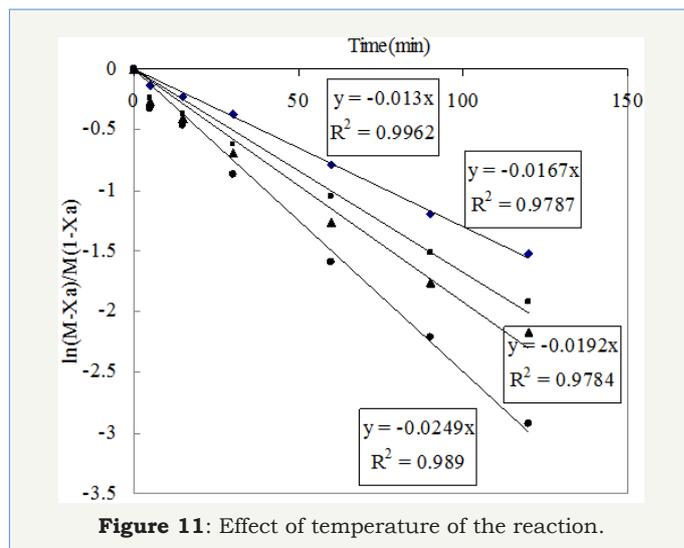


Figure 11: Effect of temperature of the reaction.

The kinetic study was also performed by varying the reaction temperature from 30-80 °C to investigate the Arrhenius Equation. The values of different parameters such as amount of phenol,

amount of solvent, weight of catalyst and the reaction time were maintained as before throughout the reaction. Fifteen ml of nitric acid was taken in this case. The values of rate constant at different temperature are obtained from Figure 11. The activation energy is calculated from the slope of Figure 12 and it is found to be 11.272KJ/mol.

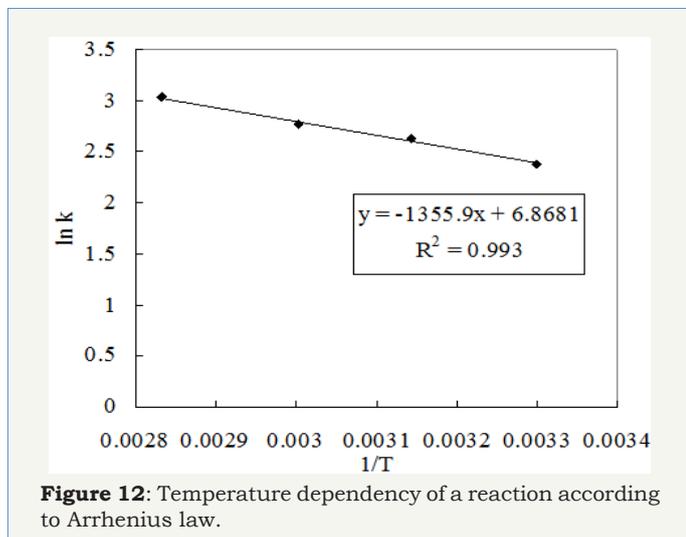


Figure 12: Temperature dependency of a reaction according to Arrhenius law.

Conclusion

Phenol is selectively nitrated to ortho-nitrophenol in high yields with dilute nitric acid in carbon tetrachloride solvent over gamma-alumina catalyst. The gamma-alumina catalyst could be easily separated from the reaction mixture by filtration for recycle and the workup procedure is simple as compared to the other conventional process. The region selective nitration of phenol using dilute nitric acid over solid acid catalyst without any use of acetic anhydride/acetyl nitrate, metal nitrates or sulfuric acid is a comparatively clean and environmental friendly process.

References

- Sharda P Dagade, Vijay S Kadam, Mohan K Dongare (2002) Regioselective nitration of phenol over solid acid catalysts. *Catalysis Communications* 3(2002): 67-70.
- Majid M Heravi, Tina Benmorad, Khadijeh Bakhtiari, Fatemeh F Bamoharram, Hossein H Oskooie (2007) $H_3+xPMo_{12-x}VO_{40}$ (heteropolyacids)-catalyzed regioselective nitration of phenol to o-nitrophenol in heterogeneous system. *Journal of Molecular Catalysis A: Chemical* 264(1-2): 318-321.
- T Esakkidurai, K Pitchumani (2002) Zeolite-mediated regioselective nitration of phenol in solid state. *Journal of Molecular Catalysis A: Chemical* 185(1-2): 305-309.
- Sujata Mallick, KM Parida (2007) Selective nitration of phenol over silicotungstic acid supported zirconia. *Catalysis Communications* 8(10): 1487-1492.
- Nitin S Nandurkar, Mayur J Bhanushali, Sachin R Jagtap, Bhalchandra M Bhanage (2007) Ultrasound promoted regioselective nitration of phenols using dilute nitric acid in the presence of phase transfer catalyst. *Ultrason Sonochem* 14(1): 41-45.
- Ashutosh V Joshi, Mubeen Baidoosi, Sudip Mukhopadhyay, Yoel Sasson (2003) Nitration of phenol and substituted phenols with dilute nitric acid using phase-transfer catalysts. *Org Proc Res Dev* 7(1): 95-97.
- KR Sunajadevi, S Sugunan (2005) Sulfated titania mediated regioselective nitration of phenol in solid state. *Catalysis Communications* 6(9): 611-616.
- Mathew R Heal, Mark AJ Harrison, J Neil Cape (2007) Aqueous phase nitration of phenol by N_2O_5 and $ClNO_2$. *Atmospheric Environment* 41(17): 3515-3520.
- V Anuradha, PV Srinivas, P Aparna, J Madhusudana Rao (2006) p-Toluenesulfonic acid catalyzed regioselective nitration of phenols with metal nitrates. *Tetrahedron Letters* 47(28): 4933-4935.
- M Arshadi, M Ghiaci, A Gil (2010) Nitration of phenol over a ZSM-5 zeolite. *Ind Eng Chem Res* 49(12): 5504-5510.
- KR Sunajadevi, S Sugunan (2006) Selective nitration of phenol over sulfated titania systems prepared via sol-gel route. *Materials Letters* 60(29-30): 3813-3817.
- AS Khder, Al Ahmed (2009) Selective nitration of phenol over nanosized tungsten oxide supported on sulfated SnO_2 as a solid acid catalyst. *Applied Catalysis A: General* 354(1-2): 153-160.
- Firouzeh Nemati, Hossein Kiani (2010) $Zn(NO_3)_2 \cdot 6H_2O/2,4,6$ -trichloro-1,3,5-triazine (TCT) a mild and selective system for nitration of phenols. *Chinese Chemical Letters* 21(4): 403-406.
- R Rajagopal, KV Srinivasan (2003) Ultrasound promoted para-selective nitration of phenols in ionic liquid. *Ultrason Sonochem* 10(1): 41-43.
- SM Kemdeo, VS Sapkal, GN Chaudhari (2010) TiO_2 - SiO_2 mixed oxide supported MoO_3 catalyst: Physicochemical characterization and activities in nitration of phenol. *Journal of Molecular Catalysis A: Chemical* 323(1-2): 70-77.



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