

Preparation of *para*-Aminophenol from Nitrobenzene through Bamberger Rearrangement Using a Mixture of Heterogeneous and Homogeneous Acid Catalysts

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Supporting Information

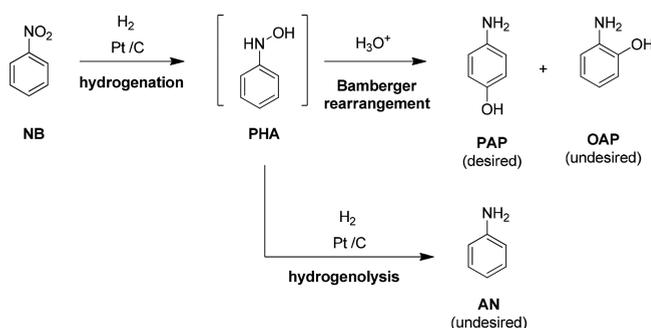
ABSTRACT: The direct preparation of *para*-aminophenol (PAP) from nitrobenzene (NB) through Bamberger rearrangement was studied in a biphasic medium using a mixture of NbO_x/SiO₂ and H₂SO₄ as an acid catalyst. After optimization of the reaction parameters, PAP was obtained with 85–88% selectivity that represents a 10% selectivity improvement compared to sulfuric acid alone. The optimized conditions were implemented in a scale-up reaction, and PAP was isolated in 84% yield (based on the recovered starting material) with 97% HPLC purity. Overall, this process requires less sulfuric acid than the traditional process, leading to a drastic reduction of the saline waste.

INTRODUCTION

para-Aminophenol (PAP) is an industrially relevant intermediate for the synthesis of drugs, pesticides, dyes, and photographic materials.¹ Whereas this last activity has almost completely disappeared since the advent of the digital age, the production of PAP is mainly driven by the manufacture of paracetamol (acetaminophen). Indeed, this analgesic and antipyretic drug is one of the most consumed drugs worldwide with an estimated production of almost 150 000 tons per year.² Recently, we have reported that paracetamol can be prepared in one step by direct amidation of hydroquinone,³ but, to date, most industrial routes rely on the acylation of PAP as the final stage, thus giving a crucial role to this intermediate. PAP can be prepared by reduction of *p*-nitrophenol using Bechamp conditions (Fe/HCl); however, this method generates a large amount of waste.⁴ Alternatively, catalytic hydrogenation of *p*-nitrophenol could furnish PAP in a much cleaner way, but the overall route is still plagued by the low selectivity of the nitration of phenol.⁵ The catalytic hydrogenation of nitrobenzene to phenylhydroxylamine (PHA) and subsequent Bamberger rearrangement⁶ in acidic aqueous conditions afford PAP with usually high selectivity for the *para* position, as only a few percent of *ortho*-aminophenol (OAP) are produced. However, aniline (AN) is the main byproduct of the hydrogenolysis of PHA (Scheme 1).⁷ Advantageously, this process occurs through a single-step operation; as a result, it provides an economical and straightforward access to PAP, which has been industrialized by Mallinckrodt Inc.⁸

A range of (supported) noble metals such as Pt, Pd, Rh, Ru, and Au have been reported for the selective hydrogenation of nitrobenzene to PAP.^{9,10} Among them, platinum is considered to be the best in terms of PAP selectivity, low loading, and high reaction rate.¹¹ Recently, nickel catalysts have also been investigated as cheaper alternatives, but these systems require high catalyst loading.¹² Sulfuric acid is classically used as a catalyst

Scheme 1. *para*-Aminophenol from Nitrobenzene through Hydrogenation/Bamberger Rearrangement



for the Bamberger rearrangement of PHA.¹¹ However, its corrosive nature is deleterious for the pressurized reactor that is essential for the hydrogenation step. Moreover, a stoichiometric amount (and more) of this “catalyst” is required, as PAP and aniline are obtained as hydrogen sulfate salts. Consequently, the reaction mixture should be neutralized by ammonia to recover free PAP thus leading to a large amount of diluted ammonium sulfate aqueous solution.

To address these issues, several attempts have been made to substitute sulfuric acid by solid acid catalysts. Mechanical mixtures of a supported Pt catalyst with ion exchange resins or heteropolyacids,¹³ SO₄²⁻/ZrO₂-Al₂O₃,¹⁴ supported aluminophosphate (SAPO),¹⁵ and SO₃H/C¹⁶ were reported. However, the selectivity in PAP is usually low to moderate. Excellent selectivity (>90%) can only be attained using zirconium sulfate.¹⁷ However, this solid acid is readily soluble in water,¹⁸ thus releasing sulfuric acid in solution. Bifunctional catalysts such as

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Pt/H-SZM-5,¹⁹ Pt-S₂O₈²⁻/ZrO₂,²⁰ and Pb-Pt/MgAPO²¹ were also reported as alternatives, but none of these systems can efficiently produce PAP with high selectivity. Finally, carbonic acid generated in a pressurized CO₂/H₂O system was recently used in association with platinum catalysts. This protocol is very attractive from an ecological point of view, but high PAP selectivity could only be reached at very low conversion (<20%).²²

Niobium-based catalysts have great potential in heterogeneous catalysis.²³ More particularly, niobium oxides usually exhibit acidic properties with Hammett acidities estimated to $H_0 \leq -5.6$ for Nb₂O₅·H₂O²⁴ and $H_0 \leq -8.2$ for NbOPO₄.²³ Consequently, they have been used in a range of acid-catalyzed organic transformations such as dehydration,²⁵ aldol condensation,²⁶ benzylic substitutions,²⁷ or esterification reactions.²⁸ Heterogeneous niobium-based catalysts have also been reported to catalyze similar reactions;²⁹ however, they have been largely underexploited. Nevertheless, they have recently found applications in the hydrolysis of saccharose,³⁰ the lactonization of ethyl levulinate,³¹ and in the Beckman rearrangement.³² However, to the best of our knowledge, (supported) niobium catalysts have never been employed for the Bamberger rearrangement of PHA to PAP. We report here the use NbO_x/SiO₂ as an acid cocatalyst for the direct reduction of nitrobenzene to PAP. Moreover, we describe a synergic effect between this heterogeneous catalyst and a homogeneous mineral acid that has a positive influence on the selectivity of the reaction.

RESULTS AND DISCUSSION

In order to optimize this process, it is necessary to consider both the chemical and the physicochemical aspects of this transformation. The reduction of NB to PAP is assumed to occur in a multiphasic medium where nitrobenzene droplets are dispersed in water and are acting as microreactors (Figure 1). This model has been first proposed by Augustine et al.^{11c}

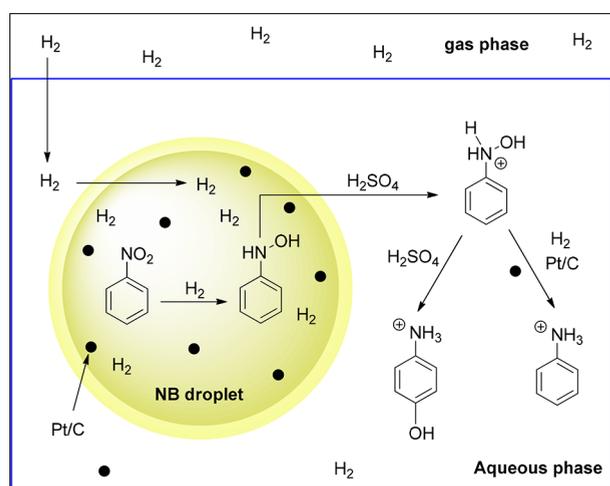


Figure 1. Reduction of NB to PAP in a multiphasic medium.

The hydrogenation of NB to PHA takes place in the NB droplets where the hydrophobic Pt/C is present and the hydrogen concentration is maximal.³³ Once produced, hydrophilic PHA migrates to the water phase through protonation and undergoes acid-catalyzed Bamberger rearrangement to give protonated PAP. From this model, one could understand that if PHA stands too long in the NB droplets, it undergoes over-

reduction to give AN as the byproduct. It is thus essential that the rate of migration of PHA is high in order to reach high selectivity. From this model, only the acid acting as catalyst for the Bamberger rearrangement could be substituted by a heterogeneous catalyst, as the presence of a Brønsted acid is essential to favor the migration of PHA in the water phase through protonation. Consequently, we envisioned that a mixture of heterogeneous and homogeneous acid catalysts could be beneficial for the selectivity of this transformation.

To test this hypothesis, NbO_x/SiO₂ was first prepared following a reported procedure^{32d} by impregnation of a Nb^V precursor [Nb(OEt)₅] onto a silica with a specific area of about 395 m²·g⁻¹.³⁴ Three batches were synthesized in order to obtain a Nb/Si molar ratio of 0.24. These catalysts were characterized by the loss at 1000 °C, ICP-EOS, and BET area (Table 1).

Table 1. Characterization of NbO_x/SiO₂ Catalysts

entry	NbO _x /SiO ₂ catalyst	loss at 1000 °C (%)	%Nb (wt %) ^a	Nb/Si molar ratio ^b	BET area (m ² /g) ^c
1	NbO _x /SiO ₂ - 1	6.5	17.2	0.15	259
2	NbO _x /SiO ₂ - 2	1.9	17.4	0.15	270
3	NbO _x /SiO ₂ - 3	4.6	15.8	0.13	264

^aDetermined by ICP-EOS. ^bCalculated based on %Nb. ^cDetermined by nitrogen adsorption at 77 K, followed by desorption at 573 K under vacuum for 5 h.

The results show that the synthesis of the catalyst is quite reproducible as the three batches exhibit very similar Nb percentages and specific areas. However, the Nb/Si molar ratio was calculated to around 0.13–0.15 indicating that the grafting of the niobium precursor was not complete (about 60%). Considering that the acidity of NbO_x/SiO₂ is the most important parameter from a catalytic point of view, it has been determined by ammonia temperature-programmed desorption (NH₃-TPD). The NH₃-TPD profile of NbO_x/SiO₂, presented in Figure 2,

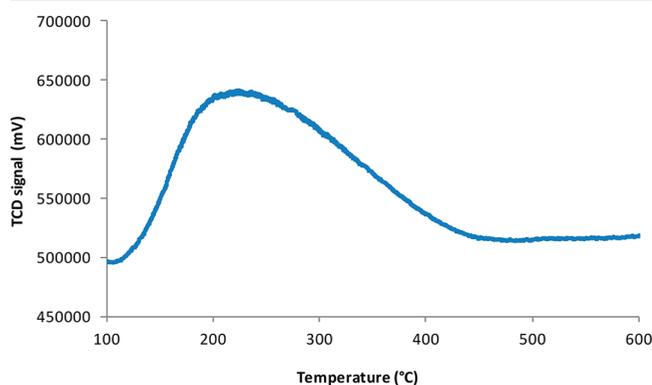


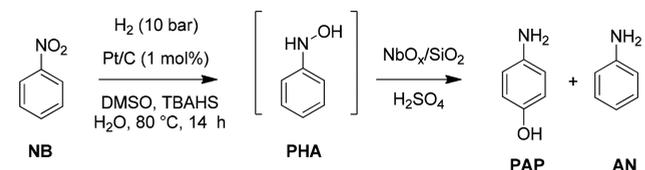
Figure 2. NH₃-TPD profile of NbO_x/SiO₂.

shows a desorption peak centered around 220 °C, indicating the presence of weak to medium acidic sites on the surface. This result is fully consistent with those of similar materials reported in the literature such as Nb₂O₅/SiO₂, Nb₂O₅/SiO₂-Al₂O₃, or Nb-MCM-41, in which the ammonia desorption peak is usually centered around 200–250 °C.^{32,32b} Moreover, considering the broadness of the signal, it could be assumed that the acidity of NbO_x/SiO₂ is widely distributed on the surface.

Finally, the quantification of acid sites of NbO_x/SiO₂ was determined to be 190 μmol·g⁻¹ after calibration.

The activity of $\text{NbO}_x/\text{SiO}_2$ was first investigated in the hydrogenation of NB to PAP under biphasic conditions at 80 °C using tetrabutylammonium hydrogensulfate (TBAHS) as a phase transfer agent (Table 2).

Table 2. Hydrogenation of NB to PAP Using Heterogeneous and Homogeneous Acid Catalysts^a



entry	H ⁺ /NB (mol %)	NbO _x /SiO ₂ (g)	conv. NB ^b (%)	sel. PAP ^b (%)	sel. AN ^b (%)
1 ^c	0* ^d	0.2	98	6 ^d	77 ^d
2	50	—	>99	54	nd
3	100	—	>99	76	24
4	100	0.2	>99	84	16
5	200	—	>99	79	21
6	200	0.2	99	86	14

^aReaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μL), TBAHS (0.6 g), H₂SO₄, NbO_x/SiO₂, H₂O (75 mL), 80 °C, 14 h. ^bDetermined by HPLC. ^cReaction run for 7 h. ^dSelectivity PHA = 6%, selectivity AOB = 9%. *The total acidity for 0.2 g of catalyst was calculated to 38 μmol of acid sites, i.e., 0.04 mol %/NB. nd: not determined.

Using 0.2 g of $\text{NbO}_x/\text{SiO}_2$, the conversion of NB reached 98% but the selectivity of PAP was only 6% (Table 2, entry 1). This result shows the ability of the catalyst to promote the Bamberger rearrangement. However, the acidity of $\text{NbO}_x/\text{SiO}_2$ is too weak to reach high selectivity. Actually, in the absence of a proton source, PHA is accumulated in the nitrobenzene phase and is mainly hydrogenated to give AN with 77% selectivity. The detection of PHA itself (6%) and azoxybenzene (AOB, 9%) also confirms the accumulation of PHA. By comparison, using 0.25 equiv of H₂SO₄ with respect to nitrobenzene (50 mol % H⁺/NB), the conversion was complete and PAP was obtained with 54% selectivity (Table 2, entry 2). The selectivity could be further improved to 76% when using 0.5 equiv of H₂SO₄ (100 mol % H⁺/NB) (Table 2, entry 3). As previously envisioned, the combination of $\text{NbO}_x/\text{SiO}_2$ with H₂SO₄ improved the selectivity to 84%, thus confirming that a Bronsted acid is definitely necessary for a rapid transfer of PHA to the water phase (Table 2, entry 4). The performance of this new system is even better than when using 1 equiv of H₂SO₄ (200 mol % H⁺/NB), giving only 79% selectivity (Table 2, entry 5). Finally, the synergetic effect was also observed in the latter case, giving 86% selectivity (Table 2, entry 6).

The loading of $\text{NbO}_x/\text{SiO}_2$ was next investigated (Figure 3).

From 0 to 0.2 g (0 to 0.4 mol % in Nb), the selectivity of PAP gradually increased from 76% to 84%. However, no further improvement was observed using larger amounts of catalyst (up to 2 g, i.e. about 4 mol %). In contrast, the selectivity slightly dropped to about 80%. This could be explained by the fact that the stirring of the heterogeneous system becomes less efficient when using a large quantity of solid. As a result, the catalyst loading of $\text{NbO}_x/\text{SiO}_2$ was set at 0.2 g for further optimization.

From previous literature reports, Pt/C and Pt/SiO₂ were found to be the best catalysts for the reduction of NB to PHA whereas Pt/C is more efficient for the direct conversion of NB to

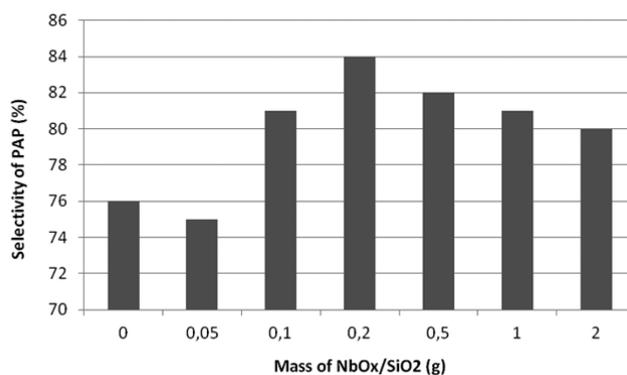
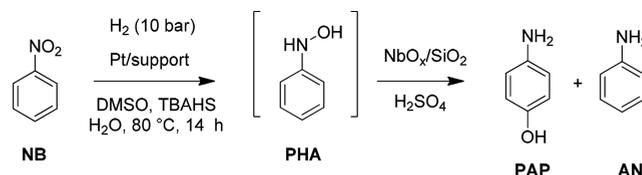


Figure 3. Selectivity of PAP versus catalyst loading. Reaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μL), TBAHS (0.6 g), H₂SO₄, (0.5 equiv, 100 mol % H⁺/NB), NbO_x/SiO₂, H₂O (75 mL), 80 °C, 14 h. The selectivity of PAP was determined determined by HPLC.

PAP. However, Figueras et al. have shown that the support sometimes has little impact on the outcome of the reaction.¹⁷ Consequently, as the role of the support is quite difficult to predict, a range of heterogeneous platinum catalysts was screened under our conditions (Table 3).

Table 3. Influence of the Support of the Platinum Catalyst^a



entry	catalyst	Pt/NB (ppm)	conv. NB ^b (%)	sel. PAP ^b (%)	sel. AN ^b (%)
1	1% Pt/C	12	>99	81	19
2 ^c	1% Pt/C	12	50	88	12
3	5% Pt/C	12	>99	80	20
4 ^d	5% Pt/C	59	42	86	14
5	5% Pt/SiO ₂	12	9	27	63
6	5% Pt/Al ₂ O ₃	12	4	63	27
7	5% Pt/ZRS	12	3	71	29
8	PtO ₂ ·H ₂ O	70	19	69	31

^aReaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), DMSO (6.5 μL), TBAHS (0.6 g), H₂SO₄ (2.5 mL), NbO_x/SiO₂ (1 g), H₂O (75 mL), 80 °C, 14 h. ^bDetermined by HPLC. ^cReaction for 2 h. ^dReaction for 5 h.

Using 1% Pt/C, the selectivity reached 81% at complete conversion and was slightly improved to 88% at only 50% conversion, respectively (Table 3, entries 1–2). Similar results could be obtained using 5% Pt/C showing that the loading of platinum on charcoal has no significant impact (Table 3, entries 3–4). Other platinum-based catalysts such as Pt/SiO₂, Pt/Al₂O₃, and Pt/ZRS were next tested. These species proved to be very poor catalysts giving very low conversion and low to moderate selectivities (Table 3, entries 5–7). Finally, Adams' catalyst (PtO₂·H₂O) was tested for comparison but it also gave moderate results despite a higher loading (Table 3, entry 8). These experiments show that the nature of the platinum support greatly impacts both the conversion and selectivity. These important disparities could be explained by the difference of polarity between supports and by the biphasic nature of the nitro-

benzene/water system. When Pt/C catalysts are used, the hydrophobic nature of the support would force the catalyst to stand in the nitrobenzene droplets, giving a high rate of hydrogenation and thus leading to high conversion of NB. In contrast, when using hydrophilic supports such as SiO₂, Al₂O₃, and ZRS, the catalyst is more likely to be dispersed in the water phase, thus giving low conversion of NB. Moreover, it could also explain why the selectivity toward PAP is low in that case, as PHA could be easily reduced to AN in the water phase.

The effect of the loading of DMSO was next probed (Figure 4). Indeed, this additive usually serves as a catalyst poison to avoid over-reduction of PHA to AN.

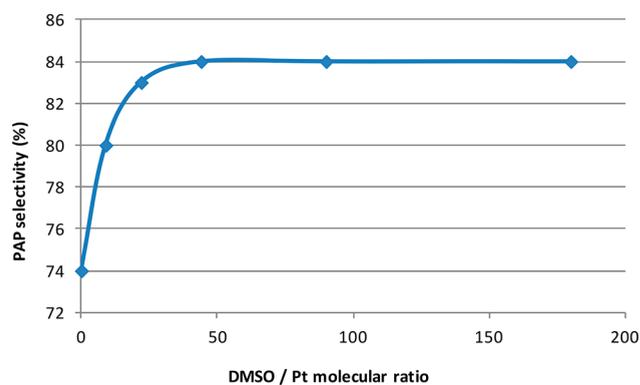


Figure 4. Selectivity of PAP versus DMSO/Pt molecular ratio. Reaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (0–13 μ L), TBAHS (0.6 g), H₂SO₄ (2.5 mL, 0.5 equiv., 100 mol % H⁺/NB), NbO_x/SiO₂ (1 g), H₂O (75 mL), 80 °C, 14 h. The selectivity of PAP was determined by HPLC.

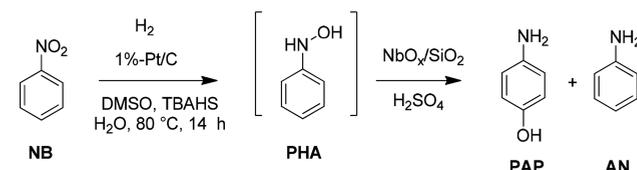
Without DMSO, the selectivity of PAP only reached 74%. Then, the selectivity increased when the molecular ratio of DMSO relative to Pt was gradually increased from 0 to 45, thus confirming the importance of this additive in the reduction process. A plateau value was reached at 84% as no further improvement could be observed using 90 or 180 equiv of DMSO. As a consequence, 90 equiv of DMSO will be used for further optimization. This value is in accordance with previous work using sulfur derivatives as additives, as values between 20 to 400 could be typically found in the literature. However, these experiments were performed with a higher platinum loading as reported by Caskey (90 to 170 ppm)^{8d,e} or Rylander (1300 ppm).^{7b}

The hydrogen pressure was next studied, and it was maintained to be constant during the experiment (Table 4).

From 2.5 to 10 bar, the selectivity was quite stable, around 83–85%, but the conversion of NB was only complete using 10 bar of hydrogen (Table 4, entries 1–3). However, the selectivity considerably dropped to 64% when increasing the hydrogen pressure to 20 bar (Table 4, entry 4). This shows that the accumulation of hydrogen should be avoided, as it favors the reduction of PHA to AN. Considering the biphasic nature of the system, the presence of a surfactant has been found to be essential to reach high PAP selectivity. Indeed, it not only could help to stabilize nitrobenzene droplets in solution but also could act as a phase transfer catalyst to facilitate the migration of PHA to the water phase. Consequently, several commercially available surfactants were tested in the model reaction (Table 5).

A blank experiment was carried out without any additive in order to establish a benchmark. Under these conditions, the

Table 4. Influence of the Hydrogen Pressure^a



entry	hydrogen pressure (bar)	conv. NB ^b (%)	sel. PAP ^b (%)	sel. AN ^b (%)
1	2.5	88	83	17
2	6	98	85	15
3	10	>99	84	16
4	20	>99	64	36

^aReaction conditions: Stainless steel reactor, H₂ (2.5–20 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μ L), TBAHS (0.6 g), H₂SO₄ (2.5 mL), NbO_x/SiO₂ (0.2 g), H₂O (75 mL), 80 °C, 14 h.

^bDetermined by HPLC.

conversion of NB is complete but the selectivity was only 74% (Table 5, entry 1). Sodium dodecyl sulfate (SDS) was first considered, as it is a cheap surfactant that is widely used in many industrial applications. However, this anionic surfactant gave lower selectivity (Table 5, entry 2). Tributylethylphosphonium ethylsulfate gave slightly improved selectivity, but the catalytic activity is very low under these conditions (Table 5, entry 3). Alkyl trimethylammonium salts bearing a benzyl or a hexadecyl chain only gave poor results (Table 5, entries 4–5). Similarly, tetrabutylammonium bromide (Br⁻), chloride (Cl⁻) or tetrafluoroborate (BF₄⁻) gave poor conversion and selectivity (Table 5, entries 6–8). These poor results could be attributed to the presence of halogen anions that could act as poisons for transition metals. The replacement of halogens by a triflate anion (OTf⁻) restored the catalytic activity of the system, and the PAP selectivity was slightly improved to 76% (Table 5, entry 9). Finally, tetrabutylammonium hydrogensulfate (TBAHS) gave full conversion and 84% selectivity (Table 5, entry 10). The difference observed between OTf⁻ and HSO₄ anions could be explained by the fact that HSO₄ is a protic anion. As a result, it is more likely to act as a phase transfer catalyst rather than a surfactant. The loading of TBAHS was also investigated (Figure 5).

The selectivity progressively increased and reached 84% when using 2 mol %. However, further increasing the quantity of TBAHS did not have a significant effect on the selectivity. This result is in accordance with the typical surfactant concentration found in the literature.

The concentration of NB in the water suspension was next investigated (Table 6). However, considering that nitrobenzene and water are not miscible at room temperature, it is quite inappropriate to refer to a concentration. In that case, we have preferred to use the NB/water (v/v) ratio.

All experiments led to a complete conversion of NB whatever the ratio used (Table 6, entries 1–4). However, slight variations of selectivity could be observed depending on the NB quantity. The selectivity was found to be optimal for a 0.12 NB/water ratio that would correspond, if nitrobenzene were soluble in water, to a 1.05 M concentration (Table 6, entry 2). Under these conditions, the NB droplets should be well dispersed in water.

Considering that the transformation of NB to PAP involves two distinct processes, namely, hydrogenation and Bamberger rearrangement, the optimal temperature is quite difficult to

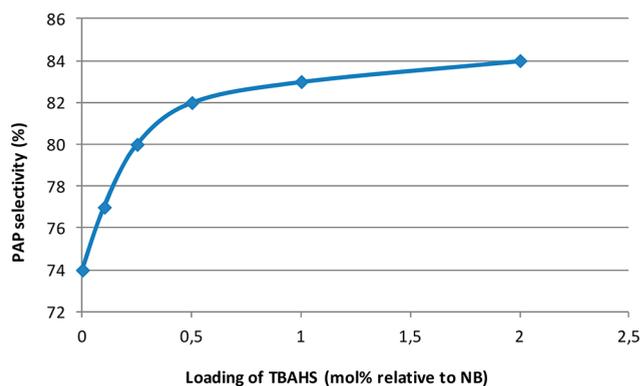


Figure 5. Selectivity of PAP versus loading of TBAHS. Reaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μL), TBAHS (0–2 mol %), H₂SO₄ (2.5 mL, 0.5 equiv, 100 mol % H⁺/NB), NbO_x/SiO₂ (1 g), H₂O (75 mL), 80 °C, 14 h. The selectivity of PAP was determined by HPLC.

Table 6. Influence of the Nitrobenzene Concentration^a

entry	NB/H ₂ O ratio (v/v)	conv. NB ^b (%)	sel. PAP ^b (%)	sel. AN ^b (%)
1	0.08	>99	81	19
2	0.12	>99	84	16
3	0.17	>99	78	22
4	0.23	>99	77	23

^aReaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μL), TBAHS (0.6 g), H₂SO₄ (2.5 mL), NbO_x/SiO₂ (0.2 g), H₂O (75 mL), 80 °C, 14 h. ^bDetermined by HPLC.

Table 7. Influence of the Temperature^a

entry	temp (°C)	conv. NB ^b (%)	sel. PAP ^b (%)	sel. AN ^b (%)
1	60	>99	82	18
2	80	>99	81	19
3	100	>99	82	18
4	120	27–37	28–38	62–72

^aReaction conditions: Stainless steel reactor, H₂ (10 bar), NB (9 mL), 1 wt % Pt/C (20 mg), DMSO (6.5 μL), TBAHS (0.6 g), H₂SO₄ (2.5 mL), NbO_x/SiO₂ (0.2 g), H₂O (75 mL), 60–120 °C, 14 h. ^bDetermined by HPLC.

However, at higher temperature, the results were seriously altered and the reaction presents a lack of reproducibility (Table 7, entry 4). This phenomenon has been attributed to the fact that the solubility of NB in water increases with the temperature,³⁵ thus increasing the homogeneous character of the system.

All the parameters that could influence the outcome of the reaction have been studied, but the PAP selectivity plateaued at about 84%. In order to obtain further information, the reaction

was followed over time. However, given the heterogeneous nature of the mixture, taking samples would necessarily bias the results. Consequently, separate experiments were carried out and stopped at different times (Figure 6).

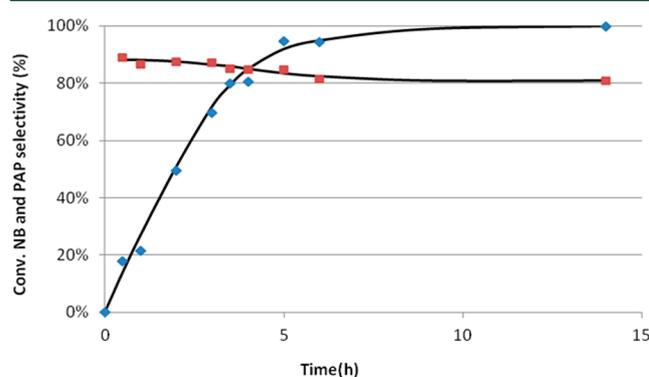


Figure 6. Conversion of NB and PAP selectivity over time.

After only 30 min, the conversion was 20% and the PAP selectivity almost reached 90%. Then, the selectivity was still 88% at 50% conversion. Finally, the PAP selectivity progressively decreased to 81% after reaching complete conversion.

These results are perfectly consistent with the “droplet” model described by Augustine et al.,^{11c} and three regimes could be envisioned (Figure 7).

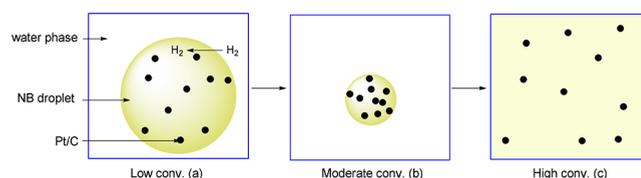


Figure 7. Different regimes of the “droplet” model.

At low conversion, the droplets of NB would be well dispersed in the water phase, thus giving high selectivity (Figure 7a). At moderate conversion, the size of the droplets would diminish, thus concentrating the catalyst in the droplets, but the selectivity should remain quite high (Figure 7b). However, at high conversion, the solubility limit of NB in water would be reached and the system would become homogeneous. This would lead to over-reduction of PHA in AN, thus reducing the selectivity (Figure 7c).

From this mechanistic rationale, it can be concluded that it is important to carry out the reaction at incomplete conversion in order to preserve the droplets and to maintain high PAP selectivity. Actually, this is the strategy that is currently employed at the industrial scale. For comparison purposes, a scale-up reaction was carried out under our optimized conditions using 174 mmol of nitrobenzene (Scheme 2).

The reaction was stopped after 3.5 h to give a 50% conversion. The catalysts were filtered, and 95% of unreacted NB was recovered from the filtrate by decantation. The remaining water phase was basified using aqueous ammonia until pH 8 in order to precipitate PAP that was filtered and washed with water. Following this procedure, PAP was obtained in 84% isolated yield (based on the remaining starting material). HPLC of the crude product revealed that the purity was about 97%, the main byproduct being *ortho*-aminophenol (OAP) (about 3%) (Figure 8).

Scheme 2. Scale-up Reaction on the Laboratory Scale

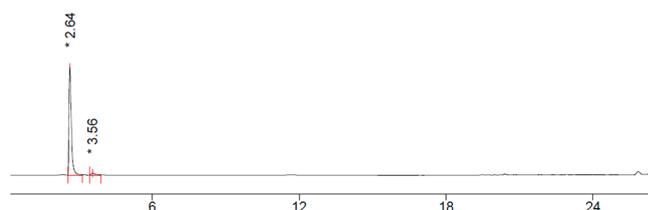
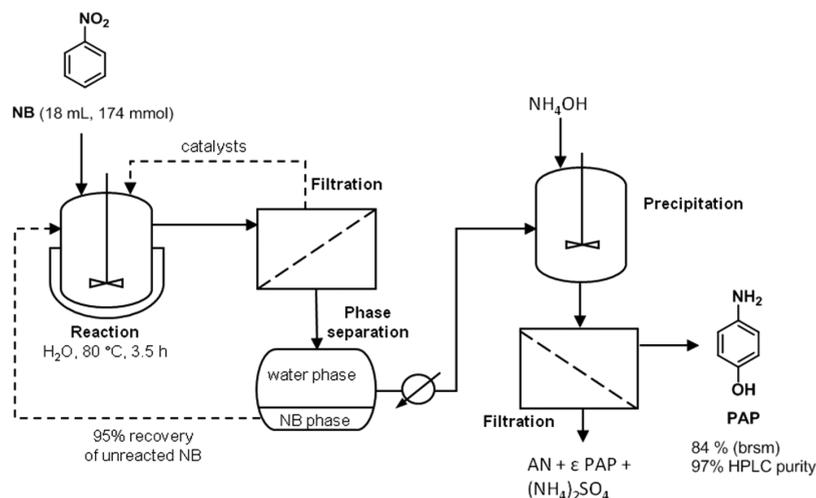


Figure 8. HPLC chromatogram of crude PAP. Retention times: 2.64 min (PAP, 97%), 3.56 min (OAP, 3%). See [Supporting Information](#) for full details.

If necessary, the purity of PAP could be further improved to >99% by recrystallization. The filtrate was also analyzed and mainly contains $(\text{NH}_4)_2\text{SO}_4$, AN, and some traces of PAP.

In order to fully evaluate the potential of this method, the recovery and recycling of catalysts was also studied over five runs ([Figure 9](#)).

For each reaction, the conversion was stopped at around 50% conversion (the hydrogen uptake was monitored). Then, the mixture of 1% Pt/C and $\text{NbO}_x/\text{SiO}_2$ was filtered, washed with toluene and MeOH, and dried under vacuum. The resulting mixture was directly reused for the next batch. From these experiments, the catalytic system proved to be efficient over the first four runs with about 48–52% conversion of NB and 85–86% PAP selectivity ([Figure 9](#), top). However, the reaction time should be progressively increased to reach about 50% conversion ([Figure 9](#), bottom). This could be explained by partial deactivation of the catalytic system but also by the mass loss of catalysts during each filtration, representing about 15% between run 1 and run 5. Indeed, the adjustment of the quantity of both Pt/C and $\text{NbO}_x/\text{SiO}_2$ to the original amount restored the catalytic properties of the mixture.

Finally, the results obtained were compared with those of the Mallinckrodt process employing excess sulfuric acid for the Bamberger rearrangement and giving the best selectivity ([Table 8](#)).

The conditions employing $\text{NbO}_x/\text{SiO}_2$ as an acid cocatalyst allow the reduction of the quantity of H_2SO_4 by more than 3-fold while keeping similar performances ([Table 8](#), entries 1–3). Consequently, the quantity of $(\text{NH}_4)_2\text{SO}_4$, produced by neutralization, was drastically reduced from 2.2 to 0.76 kg per kg of PAP ([Table 8](#), entry 4). This highlights the potential of

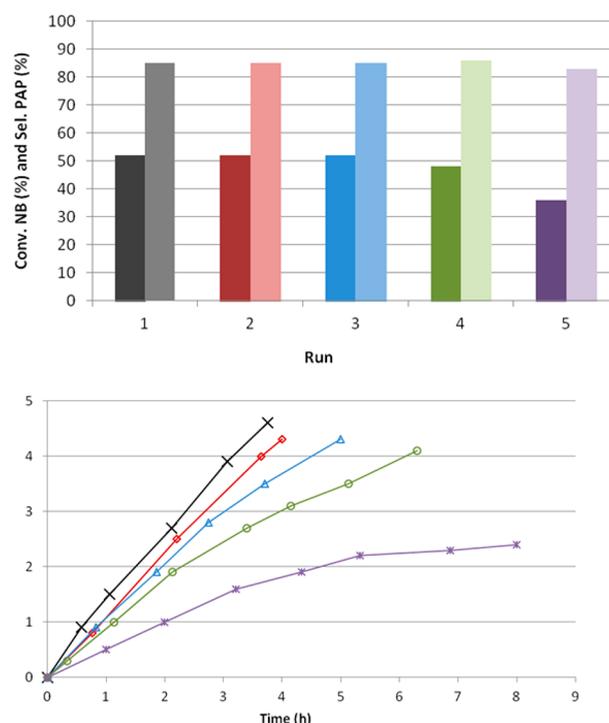


Figure 9. Recycling of the catalysts. Top: ■ Conversion of NB (%); ■ Selectivity of PAP (%). Bottom: Uptake of hydrogen over time.

Table 8. Comparison with the Mallinckrodt Process

entry	parameter	Mallinckrodt process ^{8c}	this work
1	H^+/NB ratio	1.8	0.5
2	conv. NB (%)	59	50
3	sel. PAP (%)	85	85–88
4	kg of $(\text{NH}_4)_2\text{SO}_4$ per kg of PAP	2.2	0.76

using a mixture of heterogeneous and homogeneous acid catalysts.

CONCLUSION

In conclusion, we have demonstrated that the association of a heterogeneous niobium-based catalyst with a mineral acid could be beneficial for the preparation of *para*-aminophenol (PAP)

from nitrobenzene through Bamberger rearrangement. Under these conditions, PAP was obtained with 85–88% selectivity that is representative of a 10% selectivity improvement compared to sulfuric acid alone. Moreover, this system allowed the reduction of the quantity of sulfuric acid by 3. As a consequence, less ammonia was required for neutralization, thus reducing the generation of saline waste of the traditional process. Further studies will focus on applying this strategy to other acid catalyzed transformations.

■ ASSOCIATED CONTENT

● Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.oprd.7b00354.

HPLC method, general procedures, ^1H and ^{13}C NMR spectra of *para*-aminophenol (PDF)

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Notes

The authors declare no competing financial interest.

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■ ABBREVIATIONS

NB, nitrobenzene; PAP, *para*-aminophenol; PHA, phenyl hydroxylamine; AN, aniline; OAP, *ortho*-aminophenol; AOB, azoxybenzene; DMSO, dimethyl sulfoxide; TBAHS, tetrabutylammonium hydrogensulfate

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