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# Direct amination of poly(*p*-phenylene oxide) to substituted anilines over bimetallic Pd–Ru catalysts†

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Chemical upcycling of plastic waste offers a promising opportunity for synthesizing value-added products. Despite its potential, transforming poly(phenylene oxide) (PPO) into nitrogen-based chemicals remains underexplored. To that end, we report the direct conversion of PPO to dimethylanilines over bimetallic Pd–Ru/CNT in a mixture of octane and aqueous ammonia. In a one-pot manner, PPO is initially converted into substituted phenols, and then is aminated, yielding 30% dimethylanilines over Pd<sub>7</sub>Ru<sub>3</sub>/CNT under optimum conditions. The bimetallic catalyst outperforms its monometallic equivalents. Transmission electron microscopy (TEM), powder X-ray diffraction (XRD) and energy-dispersive X-ray spectroscopy (EDX) unveiled the features of well dispersed and small-sized metallic nanoparticles. Control experiments using deuterium indicated a high reliance of hydrogen and water for the amination step and the hydrogenolysis step, respectively. Furthermore, we demonstrated product isolation through a straightforward acid–base treatment and extraction. This work introduces a viable route for upcycling PPO into valuable nitrogen-containing compounds.

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## Introduction

Plastic production has grown rapidly since the middle of the last century. However, the generation of plastic waste is an alarming problem, with less than 10% of plastic waste being recycled.<sup>1</sup> While mechanical recycling is the dominant pathway at present, it often leads to the formation of lower-value products due to the deterioration of virgin properties through successive processes.<sup>2</sup> Chemical recycling is an alternative approach, generating small molecular compounds from plastic that could either be repolymerized or directly used as fine chemicals.<sup>3,4</sup> Current research focuses on the catalytic activation of C–C, C–O, and C–N bonds in common plastics, leading to the formation of various chemicals such as aliphatic hydrocarbons,<sup>5–19</sup> arenes<sup>20–23</sup> and oxygenates.<sup>24–26</sup>

Organonitrogen chemicals, constituting over 80% of the top 200 pharmaceuticals among others, are pivotal in the chemical sector.<sup>27</sup> To foster green chemistry, there's a push to derive these chemicals from non-fossil resources like waste polymers.<sup>28</sup> Ammonolysis and aminolysis techniques have been explored to convert polyethylene terephthalate<sup>29–35</sup> and certain polyamides<sup>36–38</sup> to amines and amides. Additionally, alanine was efficiently produced from biodegradable polylactic acid using ammonia and Ru/TiO<sub>2</sub>, with impressive purity and selectivity obtained over multiple cycles.<sup>39</sup> More recently, lignin as an aromatic polymer has been converted into phenolic amines using Ru catalyst,<sup>40</sup> emphasizing the potential of making amines from aromatic polymers.

Poly(*p*-phenylene oxide) (PPO) is a versatile engineering plastic exhibiting high heat resistance and stability. It is made *via* the oxidative condensation of 2,6-dimethylphenol, and it is broadly used spanning from electronics to automotive production. Despite its USD 1.67 billion market size in 2019, the catalytic conversion of PPO into other valuable chemicals has not been thoroughly studied.<sup>41</sup> Notably, Ru/Nb<sub>2</sub>O<sub>5</sub> has been identified for its ability to cleave PPO's C–O bonds. By adjusting the metal particle size, distinct products like *m*-xylene or 3,5-dimethylphenol have been selectively obtained.<sup>42,43</sup> Small-sized metal particles are responsible for preserving the phenolic hydroxyl group while Brønsted acid sites facilitate the cleavage of C(o)–O bonds rather than C(m)–O bonds.

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Amination of PPO was performed in a 20 mL autoclave with a magnetic stirrer. Typically, PPO (50 mg), catalyst (50 mg), solvent (4 mL) and 25% aqueous ammonia solution (1 mL) were added. The autoclave was sealed and purged with N<sub>2</sub> and H<sub>2</sub> several times. Afterwards, the H<sub>2</sub> pressure was adjusted to the desired value and the autoclave was placed into a preheated steel holder, which was heated by a hot plate and insulated by a jacket. After a certain period of reaction at the desired temperature, the autoclave was cooled down in an ice-water bath. The catalyst was separated from the liquid through filtration using a PTFE syringe filter (0.45 μm) and ethyl acetate (~15 mL). Quantitative and qualitative analysis of the liquid product was performed using a gas chromatography – flame ionization detection (GC-FID, Agilent GC 7890A) and a gas chromatography – mass spectrometry (GC-MS, Agilent 7890A GC system and 5975C inert MSD with a triple-axis detector) equipped with an HP-5 column. GC samples were prepared by mixing the reaction sample and 100 μL internal standard solution (pentadecane in octane, 0.1 g mL<sup>-1</sup>). GC-FID oven program was set to initial temperature of 60 °C, heat at 5 °C min<sup>-1</sup> until 150 °C (hold 3 min), heat at 1 °C min<sup>-1</sup> until 180 °C (hold 3 min), and heat at 15 °C min<sup>-1</sup> until 300 °C (hold 4 min). A similar oven program was used for the GC-MS.

## Materials

Petroleum

Benzene

Multiple steps

2-methylphenol + 4-methylphenol

This work

One-pot reaction

2,4-diaminobenzonitrile + 4-amino-2-methylbenzonitrile

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The Pd and Ru contents in the reaction mixture were determined using inductively coupled plasma optical emission spectroscopy (ICP-OES) on a Thermo Scientific iCAP 6000 series ICP spectrometer. For some experiments with D<sub>2</sub>, the gas phase (H<sub>2</sub>, HD, D<sub>2</sub>) was analyzed using a Hiden Analytical HPR-20 R&D quadrupole mass spectrometer.

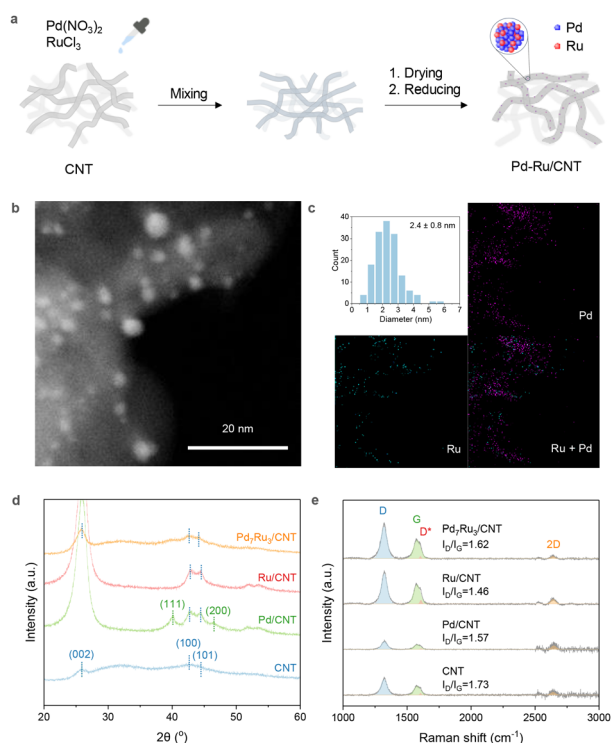
For the recycling test, the catalyst was separated from the reaction mixture by centrifugation. The liquid phase underwent quantitative analysis using GC, while the solid catalyst was sonicated with 6 mL of toluene for 1 hour to dissolve any unreacted PPO. Following this, the catalyst was washed with 15 mL of ethanol and was subsequently dried in an oven at 60 °C for 12 hours before either direct use in the subsequent run or being sent for characterization. Starting from the second cycle, due to some catalyst loss after treatment, the amount of PPO was adjusted based on the remaining catalyst to maintain a consistent PPO : catalyst mass ratio.

## Results and discussion

The bimetallic catalysts on CNT were prepared *via* the incipient wetness impregnation method (Fig. 1a). The structure of 2% Pd<sub>7</sub>Ru<sub>3</sub>/CNT is evidenced by STEM (Fig. 1b), showing metallic nanoparticles of about 2.4 nm in size (Fig. 1c) on CNT. The size of the nanoparticles is in between the sizes of the two monometallic equivalents. Specifically, 2% Pd/CNT had a

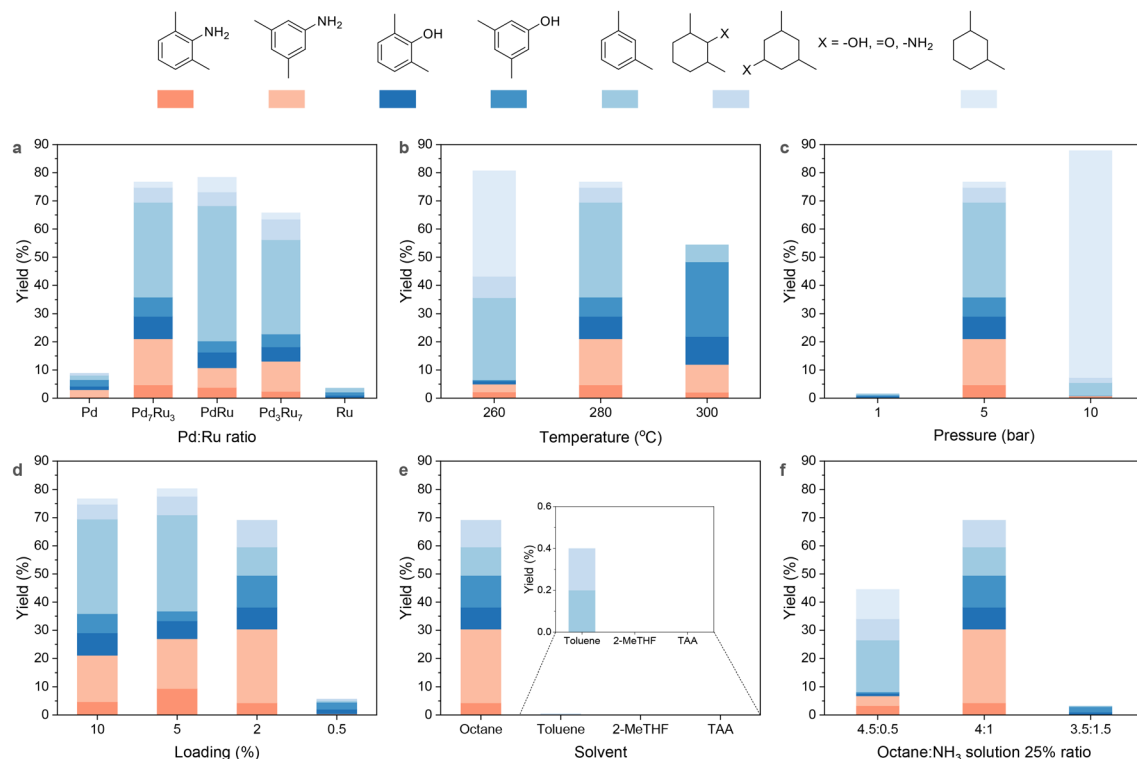
larger average size of 4.3 nm, whereas 2% Ru/CNT was smaller at 1.3 nm (Fig. S1†). Some large particles over 10 nm were seen in 2% Pd/CNT. Consistent with this, XRD analysis (Fig. 1d) showed only Pd/CNT had clear metal particle crystal phases. Elemental mapping (Fig. 1c) depicted an even distribution of Pd and Ru on the support. Further inspection (Fig. S2†) suggested Pd and Ru co-existence within particles. These observations affirm the fine dispersion and small size of Pd and Ru in the 2% loading bimetallic catalyst. Raman spectroscopy was used to study CNT structure differences among the catalysts (Fig. 1e). Characteristic CNT peaks at 1321 cm<sup>-1</sup>, 1606 cm<sup>-1</sup>, 1580 cm<sup>-1</sup>, and 2641 cm<sup>-1</sup> were identified.<sup>61</sup> Metal doping leads to a subtle I<sub>D</sub>/I<sub>G</sub> intensity ratio decrease from 1.73 to 1.46–1.62, indicating possible defect annealing during 400 °C catalyst reduction.<sup>62</sup> Bimetallic catalysts supported on CNT with varying Pd : Ru weight ratios (0 : 1, 3 : 7, 5 : 5, 7 : 3, 1 : 0) were synthesized and evaluated for PPO conversion to dimethylanilines at 5 bar H<sub>2</sub>, 280 °C for 4 h. Pd/CNT achieved a 3% dimethylanilines yield, while Ru/CNT was inactive for amination. The Pd<sub>7</sub>Ru<sub>3</sub>/CNT bimetallic catalyst significantly improved the dimethylanilines yield to 21.1% in an octane and aqueous ammonia mixture solvent (Fig. 2a). As temperatures increased, dimethylanilines yield followed a volcano trend, while dimethylphenols yield consistently increased to 36.3% (Fig. 2b). At 260 °C, hydrogenated products prevailed due to the exothermic hydrogenation of aromatic rings (low temperature favoured).<sup>55,63</sup> H<sub>2</sub> pressure also exhibited a volcano type of trend (Fig. 2c): at 1 bar H<sub>2</sub> atmosphere, hydrogenolysis and amination did not occur, while 10 bar H<sub>2</sub> led to benzene ring hydrogenation, producing nearly 90% reduced products. Previous studies indicate that aromatic compound formation from plastic and biomass depends on catalyst nanoparticle size, with smaller sizes preferred to prevent benzene ring hydrogenation and adsorption.<sup>42,43,64</sup> Thus, we assessed how the metal loading of Pd<sub>7</sub>Ru<sub>3</sub>/CNT impacts PPO amination (Fig. 2d). The yields of dimethylanilines and dimethylphenols tended to rise as metal loading decreased from 10% (3.2 nm, Fig. S3†) to 2% (2.4 nm, Fig. S3†), reaching a 30.4% amine yield. Conversely, *m*-xylene yield fell from 33.6% to 10.1%. At a further reduced metal loading of 0.5%, 6% phenolic monomers emerged due to limited metal sites for hydrogenolysis. Recognizing that a 2% metal loading is optimal, we evaluated the performance of 2% Pd<sub>7</sub>Ru<sub>3</sub> on various supports (Fig. S4†). Besides Pd<sub>7</sub>Ru<sub>3</sub>/CNT, only the MgO-supported catalyst exhibited moderate activity, yielding 19.3% of dimethylanilines. In contrast, Pd<sub>7</sub>Ru<sub>3</sub> supported on ZrO<sub>2</sub>, TiO<sub>2</sub>, and CeO<sub>2</sub> was found to be inactive.<sup>58</sup>

Interestingly, the catalyst was active only with an octane-ammonia mixture (Fig. 2e). While toluene and TAA were effective for lignin amination,<sup>65</sup> they yielded minimal monomers from PPO. The distinctive impact of octane can be explained by its capacity to facilitate PPO dispersion even at room temperature<sup>42</sup> (Table S1†), as well as its inert nature to the Pd<sub>7</sub>Ru<sub>3</sub>/CNT under the hydrogenation condition. While toluene has the capability to dissolve PPO, its poor performance is attributed to the undesired hydrogenation of toluene



**Fig. 1** (a) Schematic diagram of the 2 wt% Pd<sub>7</sub>Ru<sub>3</sub>/CNT preparation method, (b) STEM image of 2 wt% Pd<sub>7</sub>Ru<sub>3</sub>/CNT, (c) particle size distribution and elemental mapping of 2 wt% Pd<sub>7</sub>Ru<sub>3</sub>/CNT. (d) XRD analysis and (e) Raman spectra of CNT and the catalysts at 2% metal loading.





**Fig. 2** The influence of conditions for the conversion of PPO over bimetallic Pd–Ru/CNT catalyst: (a) Pd : Ru weight ratio, (b) reaction temperature, (c) H<sub>2</sub> pressure, (d) metal loading, (e) solvent type, and (f) octane to 25% aqueous NH<sub>3</sub> aqueous solution ratio. Reaction condition: 50 mg PPO, 50 mg catalyst, 4 mL octane, 1 mL 25% aqueous NH<sub>3</sub> solution, 5 bar H<sub>2</sub>, 280 °C, 4 h. Catalyst: (a) 10% metal/CNT, (b and c) 10% Pd<sub>7</sub>Ru<sub>3</sub>/CNT, (d) Pd<sub>7</sub>Ru<sub>3</sub>/CNT, (e and f) 2% Pd<sub>7</sub>Ru<sub>3</sub>/CNT. For (f), the total volume of solvent was maintained at 5 mL. 2-MeTHF: 2-methyltetrahydrofuran, TAA: *tert*-amyl alcohol.

to methylcyclohexane, which occupies the catalytic sites and consumes hydrogen (Fig. S5†). Moreover, aromatic amine production from PPO was sensitive to the octane-NH<sub>3</sub> ratio (Fig. 2f). Change in ammonia amount dramatically affected the dimethylaniline yield, with the best ratio between octane to ammonia at 4 : 1 being identified. A lower ratio resulted in the aggregation of PPO and the catalyst (Fig. S6†), hindering the depolymerization of PPO.

We assessed the effect of D<sub>2</sub> in PPO amination using 2% Pd<sub>7</sub>Ru<sub>3</sub>/CNT by substituting H<sub>2</sub> with D<sub>2</sub>. Three trials were performed: direct PPO conversion to amines (one-pot), PPO hydrogenolysis without NH<sub>3</sub> (step 1), and substituted dimethylphenol amination (step 2), with results shown in Fig. 3. Switching to D<sub>2</sub> (Fig. 3a) led to reduced dimethylamines yields in the one-pot reaction (13.9%) and step 2 (12.5%) but left dimethylphenols yields in step 1 slightly increased, suggesting amination is more hydrogen-reliant than hydrogenolysis. In contrast, replacing H<sub>2</sub>O by D<sub>2</sub>O resulted in a significant drop of dimethylphenols yield in step 1 to 10.0% (Fig. S7†). This highlighted D<sub>2</sub>O is involved in the rate-limiting step in hydrogenolysis, likely occur *via* a previously reported route of water participated C–O bond breakage.<sup>66,67</sup> Nonetheless, the D<sub>2</sub>-H<sub>2</sub> exchange catalyzed by Pd might mitigate the isotope effect.<sup>68,69</sup>

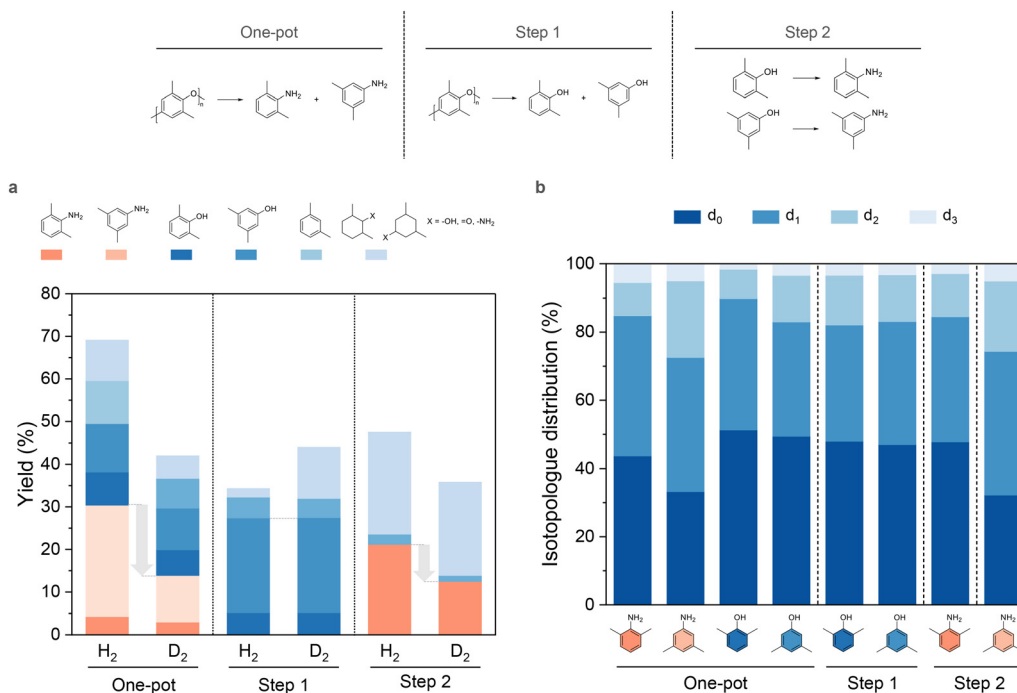
Additionally, the liquid products were analyzed by GC-MS to determine deuterium atom distribution within desired com-

pounds. Utilizing mass fragmentation patterns, we established the isotopologue distribution (Fig. 3b) by fitting the mass spectra data<sup>70</sup> with peak areas from extracted ion chromatograms. Deuterated products always account for over 50% of total product, with mono-deuterium (d<sub>1</sub>) being more abundant than di-deuterium (d<sub>2</sub>) and tri-deuterium (d<sub>3</sub>) products. Unusual adjacent fragmentations at specific *m/z* value of 77 (C<sub>6</sub>H<sub>5</sub><sup>+</sup>), 91 (C<sub>7</sub>H<sub>7</sub><sup>+</sup>), 106–107 (molecular ion losing a methyl group), and 121–122 (molecular ion) in the MS spectra (Fig. S8†) highlight deuterium substitution on the benzene ring. Generally, dimethylanilines and dimethylphenols exhibited consistent isotopologue patterns, but 3,5-dimethylaniline was exceptional with roughly 27% heavier isotopologues. One explanation is that 3,5-dimethylaniline has higher reactivity in dehydrogenation and hydrogenation, leading to multiple cycles that swap original hydrogen with deuterium.

To gain further insights into the reaction pathway of the PPO-to-dimethylanilines conversion process, a kinetic study was undertaken (Fig. 4a). The yield of dimethylphenols reached its highest value (35.7%) at 2 h and was considerably reduced with prolonged reaction time, suggesting its roles as intermediates. Meanwhile, the yield of dimethylanilines exhibited a gradual increase and reached a plateau after 4 h. Interestingly, *m*-xylene yield steadily grew, reaching 32.0% after 16 h; but the yield of 3,5-dimethylphenol displayed a reversed







**Fig. 3** (a) Product yield and (b) the distribution of deuterated and non-deuterated desired compounds. Condition for one-pot reaction (direct PPO amination): 50 mg PPO, 50 mg Pd<sub>7</sub>Ru<sub>3</sub>/CNT (2 wt% metal loading), 4 mL octane, 1 mL 25% NH<sub>3</sub> aqueous solution, 5 bar D<sub>2</sub> or H<sub>2</sub>, 280 °C, 4 h. For step 1 (PPO hydrolysis to dimethylphenols), 25% aqueous NH<sub>3</sub> solution was replaced by water. For step 2 (dimethylphenols amination), 2,6-dimethylphenol was used in (a), and 2,6-dimethylphenol or 3,5-dimethylphenol were used in two separated experiments in (b) as starting material.

trend after 4 h. This prompted us to assess dimethylphenols and dimethylanilines' activities under the reaction condition (Fig. 4b). Regardless of -OH or -NH<sub>2</sub> functional groups, the 3,5-isomer, due to less steric hindrance, was less stable than the 2,6-isomer, resulting in a greater conversion. In line with the PPO amination kinetic trend, 3,5-dimethylaniline primarily became *m*-xylene (35.7%) within 4 h. Importantly, no isomerization was seen when using 2,6- and 3,5-isomers: 2,6-dimethylphenol didn't transform into 3,5-dimethylaniline but only produced the corresponding aniline isomers.

Based on the experimental observations, plausible reaction pathways for PPO conversion were proposed in Fig. 4c. Four main steps are involved, namely hydrogenolysis, hydrogenation, imination, and dehydrogenation. It is worth noting that although dimethylcyclohexanones are considered undesired products, they are key intermediates for the amination step. Previous research proposed that the amino group can only be incorporated into the products through the attack of ammonia on the ketone group of dimethylcyclohexanones and the resulting imines subsequently transfer hydrogen to phenol, leading to the formation of the desired anilines and the regeneration of the dimethylcyclohexanone.<sup>71</sup> The generation of dimethylphenols and *m*-xylene from dimethylanilines under the reaction condition suggested that this hydrogenation – imination – dehydrogenation reaction sequence is reversible. This is aligned with the proposed pathway of phenol amination over Pd/C.<sup>72</sup>

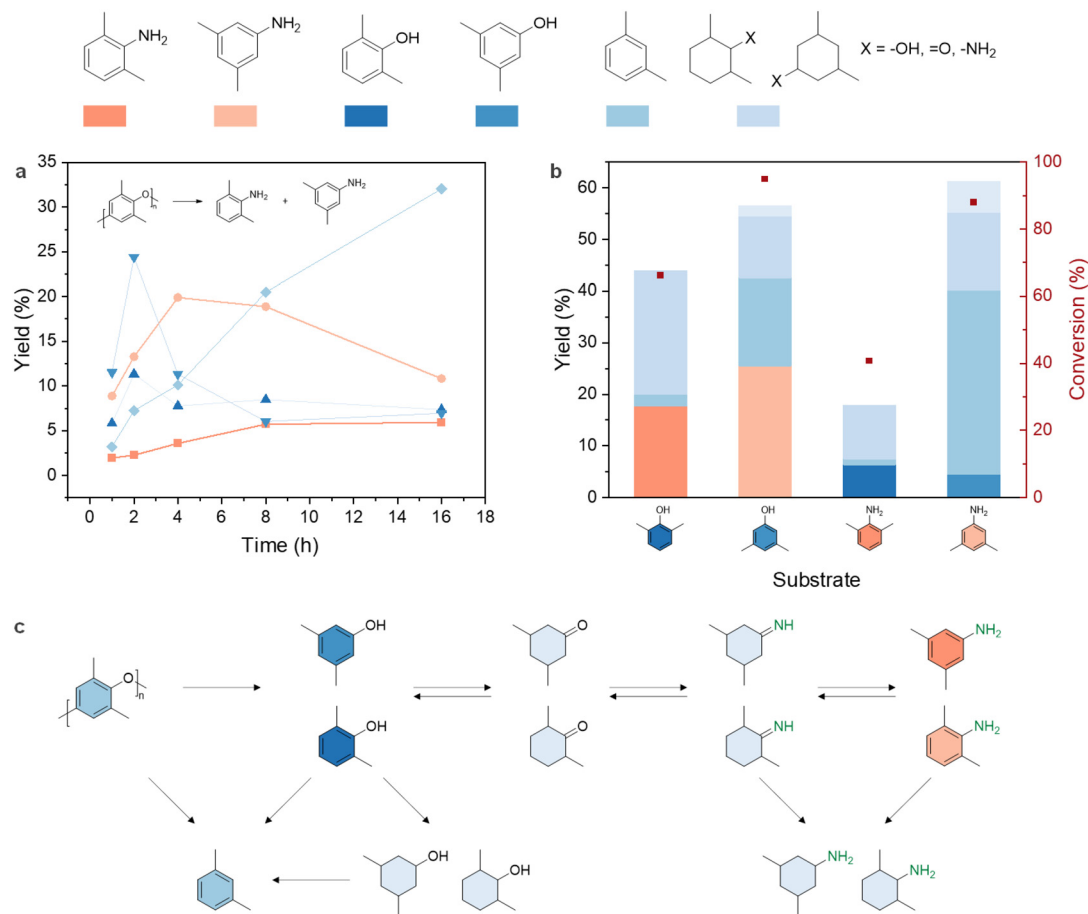
The by-product, *m*-xylene, is formed either through the C–O bond scissoring of dimethylphenols, or through the aromatiza-

tion and hydrodeoxygenation of the reduced products.<sup>72</sup> It should be noted that dimer products, such as bis(3,5-dimethylphenyl)amine and 3,3',5,5'-tetramethyl-1,1'-biphenyl, were also detected in the reaction mixture through MS analysis (Fig. S9†), indicating the occurrence of coupling reactions as side pathways.

To understand the roles of individual metal sites (Ru/Pd) and their combined effect, we examined the transformation of PPO to dimethylanilines, PPO to dimethylphenols (step 1), and 2,6-dimethylphenol to 2,6-dimethylaniline (step 2) using monometallic, bimetallic, and physically mixed catalysts (Fig. 5b–d). The GC-FID chromatograms for each route over Pd<sub>7</sub>Ru<sub>3</sub>/CNT are shown in Fig. 5a. In the one-pot transformation, the bimetallic catalyst surpassed both physically mixed and monometallic variants in dimethylaniline yield, following the order: Pd<sub>7</sub>Ru<sub>3</sub>/CNT (30.4%) > Pd/CNT (20.6%) > mixed catalyst (18.4%). This demonstrates the importance of well-dispersed and cohered active metal sites (Fig. 5b).

For step 1 (Fig. 5c), Ru/CNT demonstrated decent hydrogenolysis activity, producing 13.5% dimethylphenol yield, in line with prior studies highlighting Ru's efficacy in aromatic C–O bond cleavage in lignin.<sup>65,73</sup> The bimetallic catalyst showed even higher C–O linkage cleavage activity, yielding 27.4% dimethylphenols. On the other hand, step 2 (Fig. 5d) witnessed Pd/CNT outperforming Pd<sub>7</sub>Ru<sub>3</sub>/CNT in amination of 2,6-dimethylphenol, yielding 30.3% of 2,6-dimethylaniline. The physically mixed Ru and Pd catalysts resulted in a moderate 2,6-dimethylaniline yield (13.7%), but Ru/CNT showed no





**Fig. 4** (a) Kinetic study for the PPO amination, (b) reactions starting from different substrates, and (c) proposed reaction pathway. Reaction condition: 50 mg substrate, 50 mg Pd<sub>7</sub>Ru<sub>3</sub>/CNT (2 wt% metal loading), 4 mL octane, 1 mL 25% aqueous NH<sub>3</sub> solution, 5 bar H<sub>2</sub>, 280 °C, 4 h.

activity. Pd/C's effectiveness in phenol's liquid-phase amination to aniline, in contrast with Ru/C's inertness in the step, demonstrates Pd's selectivity in the hydrogenation/dehydrogenation of aromatic rings.<sup>52,72</sup> Combined, these control experiments demonstrate a synergistic effect in the bimetallic catalyst in PPO amination, especially in step 1, while step 2 is predominantly Pd-dependent. Pd<sub>7</sub>Ru<sub>3</sub>/CNT resulted in the production of more 3,5-dimethylphenol compared to Pd/CNT and Ru/CNT, and 3,5-dimethylphenol possessed a higher activity compared to 2,6-dimethylphenol in the amination reaction (Fig. 4b). We believe that these factors induce the improved formation of dimethylanilines with Pd<sub>7</sub>Ru<sub>3</sub>/CNT.

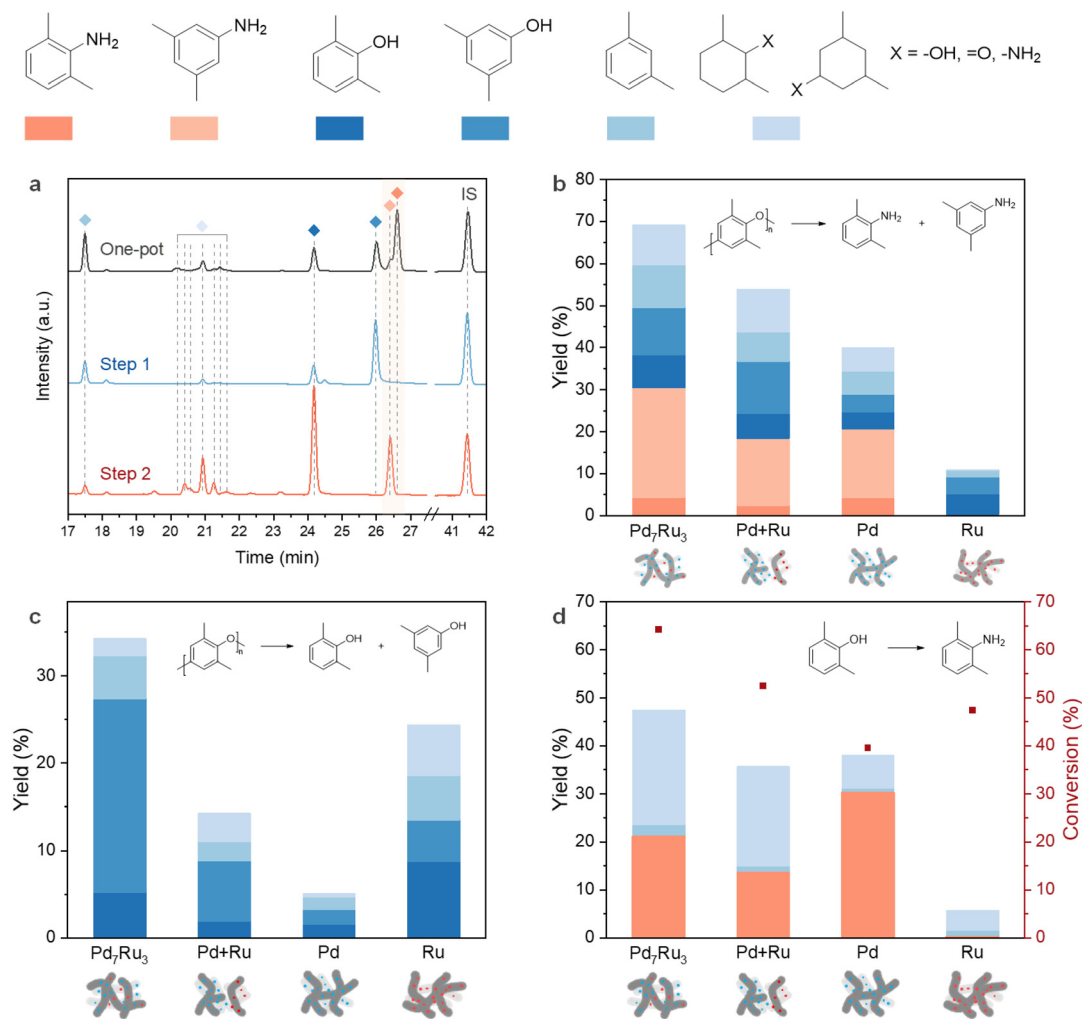
Considering that dimethylanilines are generated *via* dimethylphenols amination, it might be expected that the total dimethylanilines and dimethylphenols yield in the one-pot reaction would be equal to or lower than the dimethylphenols yield in step 1. However, the results showed the opposite trend, where step 1 only generated 27.4% of dimethylphenols compared to 49.5% of the total dimethylanilines and dimethylphenols yield in the one-pot reaction. Given that the only difference between the one-pot reaction and step 1 is the pres-

ence of NH<sub>3</sub>, it is clear that NH<sub>3</sub> has enhanced the hydrogenolysis of PPO.

We conducted a recycling test to assess the reusability of the 2% Pd<sub>7</sub>Ru<sub>3</sub>/CNT. The yield of dimethylanilines decreased after each run, reaching 7% at the third run (Fig. S10a†). Elemental analysis of the reaction mixture using ICP-OES indicated that the leaching of Pd and Ru from the catalyst was minimal, accounting for less than 0.2% (Fig. S10b†). Hence, metal leaching is unlikely to be the primary pathway for deactivation. Raman spectroscopy revealed no significant alteration in the I<sub>D</sub>/I<sub>G</sub> ratios of the catalysts after each run, implying the stability of the CNT support (Fig. S10c†). TEM analysis on the used catalyst revealed the aggregation of metal particles (Fig. S10d†), suggesting that metal aggregation may be the primary pathway for catalyst deactivation. These findings align with our observations regarding the impact of catalyst loading on product yields, where a smaller particle size was found to be more beneficial in terms of dimethylaniline yields (Fig. 2d and Fig. S3†).

Finally, we conducted a demonstration of isolating dimethylanilines from the product mixture. Successful separation was achieved through a straightforward procedure involving treatment with HCl and NaOH, followed by extraction





**Fig. 5** (a) GC-FID chromatogram and products yield of (b) one-pot reaction, (c) Step 1 and (d) Step 2. Condition for one-pot reaction: 50 mg PPO, 50 mg catalyst (2 wt% metal loading), 4 mL octane, 1 mL NH<sub>3</sub> aqueous solution 25%, 5 bar H<sub>2</sub>, 280 °C, 4 h. For step 1, 25% aqueous NH<sub>3</sub> solution was replaced by water. 2,6-dimethylphenol was used as starting material for Step 2. Pd + Ru: physical mixture of 2% Pd/CNT and 2% Ru/CNT at 7 : 3 weight ratio. IS: internal standard (pentadecane).

with ethyl acetate (Fig. S11†). This eliminates the necessity for column chromatography or distillation in isolating dimethylanilines.

## Conclusion

In conclusion, we have developed a heterogeneous catalytic process using a 2% Pd<sub>7</sub>Ru<sub>3</sub>/CNT bimetallic catalyst for producing substituted anilines from PPO in an aqueous NH<sub>3</sub>-octane mixture. This method yields primarily dimethylanilines (30%), accompanied by dimethylphenols (19%) and *m*-xylene (10%) in a one-pot amination. Based on the control experiments using deuterium and the kinetic studies, a reaction pathway from PPO to dimethylanilines involving hydrogenolysis, hydrogenation, imination and dehydrogenation steps was proposed. The synergy of Ru and Pd significantly enhances the dimethylanilines production in the one-pot transformation, particularly

during the hydrogenolysis, whereas the amination relied dominantly on Pd. Notably, dimethylanilines can be isolated from the reaction mixture through a straightforward acid-base treatment and extraction, eliminating the need for column chromatography or distillation. This strategy not only leverages PPO waste for valuable product generation but also potentially diminishes the dependence on petroleum feedstocks for dimethylanilines synthesis.

## Author contributions

N. Y. conceived and supervised the project. N. Y., G. G. and P. T. T. N. designed experiments. P. T. T. N. conducted experiments, catalyst synthesis and characterization. J. M. participated in some experiments. B. Y. and Q. H. conducted TEM analysis of catalysts. N. Y., G. G. and P. T. T. N. wrote the paper.



## Conflicts of interest

There are no conflicts to declare.

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