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Selective hydrodeoxygenation of acetophenone derivatives using a Fe₂₅Ru₇₅@SILP catalyst: a practical approach to the synthesis of alkyl phenols and anilines†

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A versatile synthetic pathway for the production of valuable alkyl phenols and anilines has been developed based on the selective hydrodeoxygenation of a wide range of hydroxy-, amino-, and nitro-acetophenone derivatives as readily available substrates. Bimetallic iron ruthenium nanoparticles immobilized on an imidazolium-based supported ionic liquid phase (Fe₂₅Ru₇₅@SILP) act as highly active and selective catalysts for the deoxygenation of the side-chain without hydrogenation of the aromatic ring. The catalytic system allows operation under continuous flow conditions with high robustness and flexibility as demonstrated for the alternating conversion of 3',5'-dimethoxy-4'-hydroxyacetophenone and 4'-hydroxynonaphenone as model substrates.

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Introduction

The development of synthetic pathways for the efficient access to alkyl phenols and anilines is attracting a lot of attention owing to the importance of these motifs as building blocks for the production of fine chemicals,¹ polymers,² and pharmaceuticals.³ Long chain alkyl phenols play a key role in the industrial production of non-ionic surfactants (*e.g.* alkyl phenol ethoxylates).⁴ Nowadays, long chain alkyl phenols are typically synthesized by direct alkylation of phenols with alkenes.⁴ However, the selectivity of this method is poor due to various side-reactions, including for example over-alkylation and isomerisation.⁴ Alkylated anilines, and in particular ethyl anilines, represent important building blocks for the production of agrochemicals⁵ and pharmaceuticals.^{3,6} While the Buchwald–Hartwig amination is generally the method of choice for the preparation of aromatic amine derivatives,⁷ it presents severe limitations including low tolerance toward functional groups (especially halides) and poor efficiency of the amination using NH₃ in the case of primary amines.^{7,8} As a result, multistep approaches involving stoichiometric

amounts of toxic reagents and waste are still commonly used.^{7d}

Thus, the development of alternative methods providing easy access to these key motifs while being at the same time potentially more sustainable and respectful of the green chemistry principles⁹ is important. In this context, we propose to access alkyl phenols and anilines through the selective hydrodeoxygenation of hydroxy-, amino-, and nitro-acetophenone derivatives (Fig. 1). These substrates are in many cases commercially available at low to moderate costs. They can be produced *via* Friedel–Crafts (FC) acylation and Fries rearrangement of phenols,¹⁰ FC acylation of nitrobenzene and aniline derivatives,¹¹ or nitration of acetophenones (Fig. 1).¹²

Unlocking the potential of this pathway requires a catalyst capable of effectively hydrodeoxygenating ketones, while leaving the aromatic ring and other functionalities untouched. Recently, promising catalysts have been reported for the selective hydrodeoxygenation of aromatic ketones to aromatic alkanes.¹³ However, catalytic systems capable of selectively hydrodeoxygenating hydroxy-, amino-, or nitro-acetophenones with H₂ are scarce due to the tendency of these substrates to undergo various competing side reactions especially when acid sites are present in the catalyst materials.¹⁴ Consequently, the development of a catalytic system possessing excellent activity, selectivity, stability, and versatility for the production of alkyl phenols and anilines from acetophenone derivatives is highly desirable, but remains a challenge.

The immobilization of metal nanoparticles on molecularly modified surfaces, and in particular supported ionic liquid phases, is attracting increasing attention to generate catalytic

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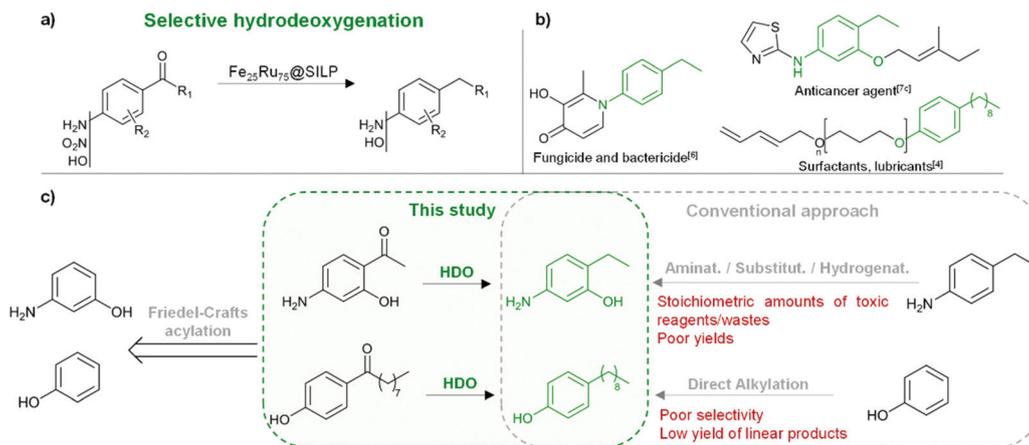


Fig. 1 Production of alkyl anilines and alkyl phenols through $\text{Fe}_{25}\text{Ru}_{75}@SILP$ -catalyzed hydrodeoxygenation of acetophenone derivatives. (a) Synthetic pathway; (b) examples of potential applications;^{4,6,7c} (c) comparison with conventional approach^{4,7d} for two selected examples.

systems with tailor-made reactivity.^{15,16} Careful tuning of the NPs composition and molecular modifier structure enables a large degree of control over H_2 activation (homolytic *versus* heterolytic) providing hydrogenation^{16a-h} and hydrodeoxygenation^{14b,16h-m} catalysts with tunable activity and selectivity. Recently, we have shown that bimetallic Fe–Ru nanoparticles immobilized on an imidazolium-based SILP possess excellent activity and selectivity for the hydrodeoxygenation of substituted hydroxyacetophenones to ethyl phenol derivatives without hydrogenation of the aromatic ring.^{16m} The acid-free $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalyst prevented the acid-catalysed side-reactions typically observed when using conventional hydrodeoxygenation catalysts. Mesomeric stabilization of the intermediates was found essential to observe high hydrodeoxygenation activity in the absence of an acidic co-catalyst.

Based on these initial findings, we decided to explore the potential of the $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalytic system for the conversion of a broad scope of long chain hydroxy-acetophenones as well as substituted amino- and nitro-acetophenones. For the first time, we report herein the remarkable versatility of the $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalyst for their selective hydrodeoxygenation and demonstrate its robustness and flexibility under continuous flow conditions.

Results and discussion

Catalyst synthesis and characterization

The $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalyst (Fig. 2) was synthesized following a recently reported procedure.^{16m} In brief, the preparation of the SILP involved the condensation of [1-butyl-3-(3-triethoxysilylpropyl)-imidazolium]NTf₂ with dehydroxylated SiO_2 . The NPs were prepared through an organometallic approach, with the *in situ* reduction of a mesitylene solution of $\{\text{Fe}[\text{N}(\text{Si}(\text{CH}_3)_3)_2]_2\}$ and $[\text{Ru}(\text{cod})(\text{cot})]$ (cod = cyclooctadiene; cot = cyclooctatriene) in the presence of the SILP material under an atmosphere of H_2 (3 bar) at 150 °C (for detailed procedure see ESI†). Scanning transmission electron microscopy with high

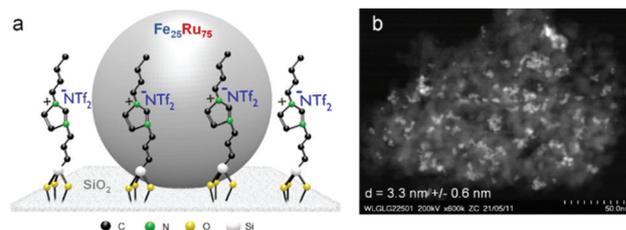


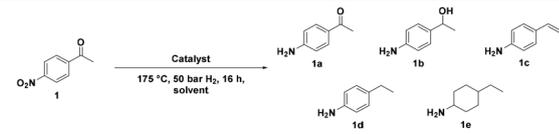
Fig. 2 (a) Schematic illustration and (b) STEM-HAADF image of the $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalyst.

angle annular dark field (STEM-HAADF) analysis confirmed the formation of small and well-dispersed nanoparticles with an average size of 3.3 nm (Fig. 2b). The BET surface area of 240.7 $\text{m}^2 \text{g}^{-1}$ is significantly lower than for the starting SiO_2 (342.3 $\text{m}^2 \text{g}^{-1}$) due to the chemisorption of the ionic liquid. A metal loading of 0.4 mmol g^{-1} and a Fe : Ru ratio of 25 : 75 were determined by scanning electron microscopy with energy dispersive X-ray spectroscopy (SEM-EDX), well in agreement with the theoretical values (Table S1†). The bimetallic structure of the $\text{Fe}_{25}\text{Ru}_{75}$ nanoparticles as well as the zerovalent state of Fe and Ru atoms were previously confirmed by XANES and EXAFS characterization.^{16c}

Catalytic study

Catalyst evaluation. The catalytic properties of $\text{Fe}_{25}\text{Ru}_{75}@SILP$ were first explored for the conversion of 4-nitroacetophenone (**1**), and compared to the performances of reference catalysts including $\text{Fe}_{100}@SILP$, $\text{Ru}_{100}@SILP$ and $\text{Fe}_{25}\text{Ru}_{75}@SiO_2$ (Table 1). While the use of monometallic $\text{Fe}_{100}@SILP$ resulted in no conversion (entry 1), $\text{Ru}_{100}@SILP$ lead to the hydrodeoxygenation of **1** with hydrogenation of the aromatic ring, giving **1e** in quantitative yield (entry 2). Satisfyingly, **1** was selectively hydrodeoxygenated to 4-ethylamine (**1d**) when using bimetallic $\text{Fe}_{25}\text{Ru}_{75}@SILP$ catalyst (entry 3). The suppression of aromatic ring hydrogenation on bi-



Table 1 Hydrodeoxygenation of **1** with various catalysts


Entry	Catalyst	Solvent	X [%]	Y (1a) [%]	Y (1d) [%]	Y (1e) [%]
1	Fe ₁₀₀ @SILP	Mesitylene	0	0	0	0
2	Ru ₁₀₀ @SILP	Heptane	>99	0	0	>99
3	Fe ₂₅ Ru ₇₅ @SILP	Mesitylene	>99	0	>99	0
4	Fe ₂₅ Ru ₇₅ @SiO ₂	Mesitylene	>99	0	0	>99
5	Fe ₄₀ Ru ₆₀ @SILP	Mesitylene	>99	85	15	0
6	Fe ₆₀ Ru ₄₀ @SILP	Mesitylene	>99	>99	0	0

Reaction conditions: Catalyst (10 mg, metal content: 0.004 mmol), substrate (25 eq.), solvent (0.5 mL), 175 °C, H₂ (50 bar), 16 h, 500 rpm. Conversion and product yields determined by GD-FID using decane as internal standard. X = conversion, Y = yield.

metallic Fe₂₅Ru₇₅ alloy NPs is consistent with previous reports, and is presumably due to the disruption of the arrangement of 3–4 adjacent Ru atoms necessary to hydrogenate 6-membered aromatic rings.^{16c,h} Interestingly, attempts to prepare Fe₂₅Ru₇₅ NPs on non-modified silica resulted in Fe₂₅Ru₇₅@SiO₂ materials with a reactivity typical of monometallic Ru NPs (entry 4), evidencing the importance of the SILP support as a matrix to produce small and alloyed bimetallic NPs. Gradually increasing the Fe content in Fe₄₀Ru₆₀@SILP and Fe₆₀Ru₄₀@SILP led to a sharp decrease and ultimately a suppression of the C=O hydrodeoxygenation activity, conserving only activity for the hydrogenation of the nitro functionality (entries 5 and 6).

Long chain hydroxyacetophenones

The activity of the Fe₂₅Ru₇₅@SILP catalyst was investigated for a scope of hydroxyacetophenones with various alkyl chain lengths (Table 2).

Potential main products for this transformation include the aromatic alcohols arising from the hydrogenation of the substrates, styrene derivatives resulting from alcohol dehydration, and the desired alkyl phenols obtained after hydrogenation of the styrene-type intermediates. Standard conditions for these reactions were set to 175 °C, 50 bar and 16 h reaction time. Substrates 2–8 with increasing side chain length (from R = Me to Oct) were readily hydrodeoxygenated under these conditions, giving the corresponding alkyl phenols **2c–8c** in excellent yields and selectivities. No aromatic ring hydrogenation was observed with the bimetallic Fe₂₅Ru₇₅ NPs, in agreement with previous reports.^{14b,16m} A reaction time of 24 h was necessary to fully convert substrate **6**, presumably due to the presence of some impurities in the starting substrate. Isolation of the resulting alkyl phenols was readily accomplished by filtration of the reaction mixtures and evaporation of the solvent, providing the isolated compounds **2c–6c** and **8c** in excellent yields (80–94%). To the best of our knowledge, this is the first time that products **4c–8c** are obtained through this pathway

using molecular hydrogen. Interestingly, 4'-nonyl phenol (**8c**) is used industrially in combination with ethane or ethylene oxide to produce surfactants, technical non-ionic surfactants, wetting agents, emulsifiers and foam-stabilizers.^{4,17}

Amino- and nitro-acetophenones

The versatility of the Fe₂₅Ru₇₅@SILP catalyst was then explored for a wide range of substituted amino- and nitroacetophenones (Table 3). Under standard conditions, aminoacetophenone (**9**) and 1-(4-aminophenyl)-ethanone (**10**) were hydrodeoxygenated to the corresponding 4'-ethylaniline (**1c**) and 4'-propylaniline (**10c**) in excellent yields (92% and 97%, respectively). Interestingly, **1c** and **10c** are important building blocks for the synthesis of fungicides and bactericides.⁵ Significant amounts of 3'-methoxyanilin were observed when using 2'-methoxy-4'-aminoacetophenone (**11**) as substrate (entry 3). Increasing the reaction temperature to 200 °C and using dioxane as solvent was beneficial to avoid deacylation, and the desired product **11c** could be obtained in high yield (85%, entry 4). 2'-Hydroxy-4'-aminoacetophenone (**12**) was hydrodeoxygenated with excellent selectivity and yield (both >99%, entry 5), producing 5-amino-2-ethyl-phenol (**12c**), a building block entering in the preparation of promising anti-cancer agents.^{6c} The production of 2-bromo-4-ethyl-benzenamine (**13c**) and 2-fluoro-4-ethyl-benzenamine (**14c**) in fair to high yields under optimized conditions (57% and 89%, respectively) is particularly interesting since these products are valuable building blocks¹⁸ that cannot be obtained *via* Buchwald–Hartwig amination⁷ and that typically require multistep synthetic approaches with limited selectivity.¹⁹

A similar approach was adopted for the conversion of various nitro-acetophenones. First, a time profile was recorded using 4'-nitroacetophenone (**1**) as model substrate (Fig. 3). The conditions were adapted (120 °C, 50 bar H₂) to slow down the reaction and allow observing intermediates.

1 was very quickly (~1 h) converted to the 4'-amino-acetophenone (**1c**), which was then hydrodeoxygenated to give 4'-vinylaniline (**1b**). **1b** was then consumed over time to give a quantitative yield of **1c** after 12 h in a typical kinetic profile for sequential reaction steps. These results suggest that the Fe₂₅Ru₇₅@SILP catalyst can be used to perform HDO reactions at relatively mild temperature. The observation of **1b** as major intermediate indicates that the deoxygenation proceeds through the cleavage of the C–O bond and alcohol elimination *via* a E2-type mechanism resulting in an olefin intermediate. In addition, neither the alcohol **1a** nor its nitro-analogue could be observed, suggesting that this elimination step is considerably faster than the hydrogenation of **9** to **1a** (C=O hydrogenation) and of **1b** to **1c** (C=C hydrogenation).

Starting from substituted nitroacetophenones (**1**, **15–17**) was similarly successful, leading to the corresponding alkyl aniline derivatives in excellent yields (>99%). Notably, the conversion of 1-(3-fluoro-4-nitrophenyl)-ethanone (**16**) allowed obtaining **14c** in higher yield and milder conditions (entry 9) that when starting directly from **1**. Having the nitro functionality in *ortho* position (**17**) did not affect the catalyst's perform-



Table 2 Hydrodeoxygenation of hydroxyacetophenones with various alkyl chain length using Fe₂₅Ru₇₅@SILP

R = Me, Prop, Bu, Pent, Hex, Hept, Oct

Entry	Substrate	Product	X [%]	S [%]	Y [%]
1			>99	>99	>99 (88)
2			>99	>99	>99 (91)
3			>99	99	99 (90)
4			>99	>99	>99 (80)
5			>99	>99	>99 (82)
6 ^a			51	76	39
7			92 ^b >99	91 >99	84 >99 (94)

Reaction conditions: Catalyst (37.5 mg, metal content: 0.015 mmol), substrate (0.375 mmol, 25 eq.), solvent (0.5 mL), 175 °C, H₂ (50 bar), 16 h, 500 rpm. ^a By-product: 4'-octenylphenol. ^b 24 h. Conversion and product yields determined by GD-FID using decane as internal standard. X = conversion, S = selectivity, Y = yield. Isolated yields in brackets.

ances and gave quantitative yield of 2'-ethyl-anilin (**17c**) which can be used for the synthesis of enzyme inhibitors.¹⁸ To finish, two dialkylaminoacetophenones were investigated (1-[4-(dimethylamino)phenyl]-ethanone (**18**) and 1-[4-(diethylamino)phenyl]-ethanone (**19**). **18** and **19** were readily hydrodeoxygenated to the corresponding ethyl dialkylbenzeneamines **18c** and **19c** (>99% yield for both), highlighting the versatility of the Fe₂₅Ru₇₅@SILP catalysts as well as its tolerance toward various functional groups. In addition, product isolation was performed for several of the entries presented, giving **1c**, **10c**, **12c**, **15c**, **18c** and **19c** in good to excellent yields (79–94%) and thus confirming the good practicability of Fe₂₅Ru₇₅@SILP. To the best of our knowledge, this is also the first time that products **10c–14c** and **15c–16c** are accessed *via* the selective hydrodeoxygenation of substrates **10–14** and **15–16**.

Satisfyingly, replacing mesitylene by anisole – a solvent recommended by the CHEM21 green chemistry selection guide²⁰

– resulted also in excellent yields and selectivities, as evidenced for some selected examples (Table S2†).

Continuous flow operation

The performance of the Fe₂₅Ru₇₅@SILP catalyst was further investigated under continuous flow conditions. A solution of 4'-hydroxyacetophenone (**2**) (0.025 mol L⁻¹ in anisole) was passed over a cartridge packed with the Fe₂₅Ru₇₅@SILP catalyst (433.5 mg, 0.173 mmol metal) at a flow rate of 0.6 mL min⁻¹ under standard conditions (175 °C, 50 bar H₂) using an H-Cube Pro reactor system from ThalesNano (Fig. S1†). Anisole was chosen as solvent for the continuous flow experiments to ensure good solubility of **2** at room temperature. The reaction conditions were selected to allow the observation of intermediates, evidencing any potential changes in the catalyst reactivity over time. The evolution of the substrate conversion and products yields is given in Fig. 4. Full conversion of **2** to a mixture



Table 3 Substrate scope of amino- and nitro-acetophenone derivatives in hydrodeoxygenation using Fe₂₅Ru₇₅@SILP

R¹ = Me, Et / R² = F, Br, OH, H, OMe, Me / R³ = Me, Et

Entry	Substrate	T [°C]	Product	X [%]	S [%]	Y [%]
1 ^a		175		92	>99	92
2		175		>99	97 ^b	97 (94)
3		175		>99	55 ^c	55
4		200 175		>99 >99	85 ^c >99	85 >99 (79)
5 ^d		150		86	66 ^e	57
6 ^d		200		>99	89 ^f	89
7		175		>99	>99	>99 (93)
8		175		>99	>99	>99 (94)
9 ^d		150		>99	>99	>99
10		175		>99	>99	>99
11		175		>99	>99	>99 (80)
12 ^a		175		>99	>99	>99 (87)

Conditions: Fe₂₅Ru₇₅@SILP (10 mg catalyst containing 0.004 mmol total metal), 0.1 mmol substrate (25 equivalents), 0.5 mL mesitylene, 50 bar H₂, 16 h, 500 rpm. ^a 24 h. ^b By-product: propylbenzene. ^c By-product: 3'-methoxyanilin. ^d Solvent: dioxane. ^e By-product: 2'-bromoanilin. ^f By-product: 4'-ethylanilin. X = conversion, S = selectivity, Y = yield, determined by GD-FID using decane as internal standard. Isolated yields in parentheses.



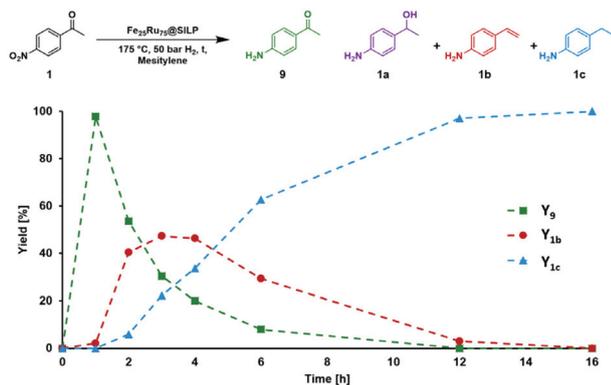


Fig. 3 Reaction/time-profile for the hydrodeoxygenation of 4'-nitroacetophenone (**1**) using Fe₂₅Ru₇₅@SILP. Reaction conditions: Fe₂₅Ru₇₅@SILP (10 mg catalyst containing 0.004 mmol total metal), 0.1 mmol **1** (25 eq.), 120 °C, 0.5 mL mesitylene, 50 bar H₂, 500 rpm. Y = yield.

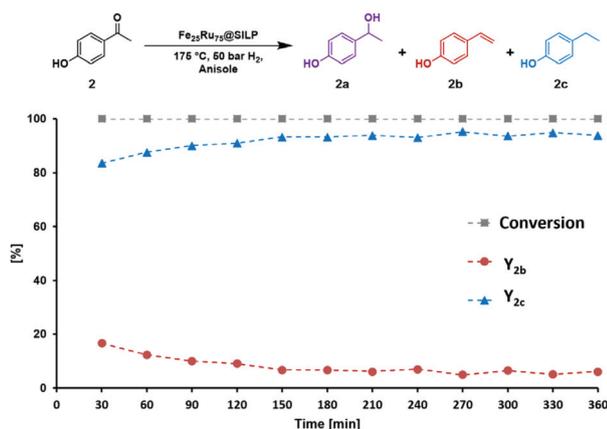


Fig. 4 Stability test of Fe₂₅Ru₇₅@SILP under continuous flow conditions. Conditions: 175 °C, 50 bar H₂, 60 NML min⁻¹ H₂ flow, 0.6 mL min⁻¹ substrate flow, substrate solution: 0.025 mol L⁻¹ in anisole, residence time: 17 s, Fe₂₅Ru₇₅@SILP (433.5 mg, 0.173 mmol metal). Product yields determined by GD-FID using decane as internal standard. Y = yield.

of 4'-ethenylphenol (**2b**) and 4'-ethylphenol (**2c**) was observed for the complete duration of the experiment, *i.e.* 6 hours time-on-stream. After a short equilibration period (0–60 min), excellent yields of the hydrodeoxygenated product 4'-ethylphenol (**2c**, 90–95%) were maintained over the remaining 5 hours without any noticeable decay. TEM characterization of the Fe₂₅Ru₇₅@SILP material after 6 hours on stream did not show any change in the size and dispersion of the FeRu NPs (3.1 nm ± 0.6 nm) (Fig. S2 and Table S1†). In addition, slight variations in metal loading and Fe:Ru ratio evidenced by SEM-EDX mapping were within the measurement error and thus insignificant (Table S1†). BET analysis showed a minor decrease in surface area after catalysis (from 240.7 to 232.1 m² g⁻¹) again within measurement error (Table S1†). These results demonstrate that Fe₂₅Ru₇₅@SILP is active, selective, and stable for the hydrodeoxygenation of **2** under continuous flow operation.

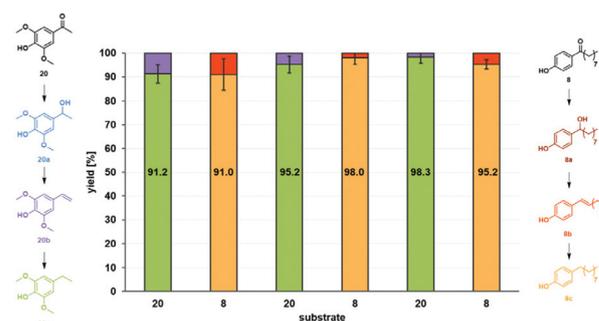


Fig. 5 Substrate switch under continuous flow conditions. Conditions: 175 °C, 50 bar H₂, 60 NML min⁻¹ H₂ flow, 0.5 mL min⁻¹ substrate flow, substrate solutions: 0.015 mol L⁻¹ 3',5'-dimethoxy-4'-hydroxyacetophenone (**20**) in anisole and 0.015 mol L⁻¹ 4'-hydroxynonanophenone (**8**) in mesitylene, residence time: 18 s, Fe₂₅Ru₇₅@SILP (455.9 mg, 0.182 mmol metal). Between each substrate switch, the system was washed with anisole, mesitylene or heptane (for details see ESI†). Product yields determined by GD-FID using decane as internal standard.

Based on these encouraging results, the versatility of Fe₂₅Ru₇₅@SILP was further explored under continuous flow conditions. Solutions containing biomass-derived 3',5'-dimethoxy-4'-hydroxyacetophenone (**20**)²¹ in anisole and 4'-hydroxynonanophenone (**8**) in mesitylene were alternately passed over the catalyst bed using the continuous flow set-up described previously. Each substrate solution was applied with a flow rate of 0.5 mL min⁻¹ under 50 bar H₂ at 175 °C for 45 minutes without any catalyst make-up or changes in the set-up between the reactions. The substrate switch was performed three times in a row to study the impact of real-time repeated substrate changes on the catalyst's performances (Fig. 5). Under optimized reaction conditions (Table S3†), the substrate feed could be switched back and forth between substrate **20** and **8** to alternatively produce **20c** and **8c** in more than 90% yield. For both substrates, the styrene-type compounds **20b** and **8b** were observed as only intermediates, outlining once again that the deoxygenation step (C–O bond cleavage) is significantly faster than hydrogenation of the C=O and C=C bonds.

The experiment represents a total of 4.5 h time-on-stream without any change in activity, selectivity, or stability of the catalytic system for either of the substrates, highlighting its robustness and versatility in the selective hydrodeoxygenation of hydroxyacetophenone derivatives.

Synthetic approach evaluation

To finish, the synthetic approach we propose here (FC acylation + hydrodeoxygenation) was systematically compared to conventional methods for the preparation of 2'-hydroxy-4-ethylaniline (**12**) and 4'-nonylphenol (**8**), the two products highlighted in Fig. 1b and c. Five parameters related to the green chemistry principles were taken into consideration to rank the pathways, *i.e.* the overall reaction yield (Y), the atom economy (AE), the number of steps (steps), the hazardous nature of the reagents (safety), and the economical aspect (difference of



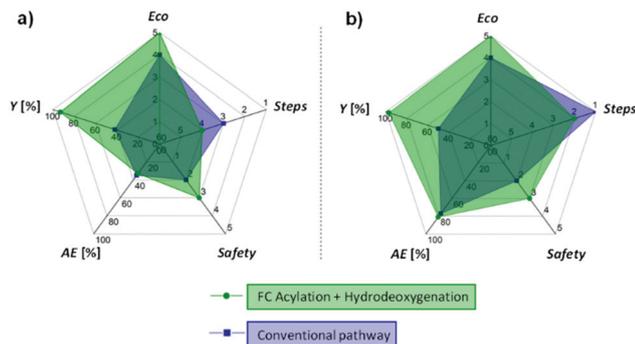


Fig. 6 Comparison between our approach and conventional synthetic pathways for the production of (a) 2'-hydroxy-4-ethylaniline (**12**), and (b) 4'-nonylphenol (**8**). Area proportional to the green chemistry metrics (large area = good metrics). Details concerning the different synthetic pathways and corresponding rankings are available in the ESI.†

value between the product and the starting substrate, eco), giving the spider web-type graphs shown in Fig. 6 (details concerning the different synthetic pathways and corresponding rankings are available in the ESI, Fig. S3–S6 and Table S4†). While the hydrodeoxygenation-based approach necessitates in both cases an additional step as compared to the conventional pathways, the other parameters are either equal (AE), or clearly in favour (Y, eco, safety) of the alternative we propose, highlighting the potential benefits associated to the production of alkyl phenols and anilines *via* selective hydrodeoxygenation of acetophenone derivatives.

Conclusion

In conclusion, the catalytic hydrodeoxygenation of hydroxy-, amino-, and nitro-acetophenone derivatives was established as a broadly applicable synthetic pathway for the synthesis of alkyl phenols and anilines. The bimetallic $\text{Fe}_{25}\text{Ru}_{75}\text{@SILP}$ catalytic system proved highly effective for the selective hydrodeoxygenation of a broad range of substrates comprising various functionalities while conserving the aromaticity in the products. The activity, selectivity and stability of $\text{Fe}_{25}\text{Ru}_{75}\text{@SILP}$ were demonstrated for the continuous flow hydrodeoxygenation of 4-hydroxyacetophenone (**1**) for over 6 h time-on-stream. In addition, the alternating hydrodeoxygenation of 3',5'-dimethoxy-4'-hydroxyacetophenone (**20**) and 4'-hydroxynonanophenone (**7**) under continuous flow conditions was achieved in real-time over a single catalyst cartridge, producing the corresponding phenols alternatively in excellent yield and outlining the versatility of the $\text{Fe}_{25}\text{Ru}_{75}\text{@SILP}$ catalytic system. The potential benefits of the proposed pathway with regards to sustainability were evidenced for two selected examples. This approach provides a promising alternative strategy for the effective synthesis of long chain alkyl phenols and ethyl anilines that are key building blocks for the production of important fine chemicals, agrochemicals, and pharmaceuticals.

Experimental section

Synthesis of $\text{Fe}_{25}\text{Ru}_{75}\text{@SILP}$

A solution of $\{\text{Fe}[\text{N}(\text{Si}(\text{CH}_3)_3)_2]_2\}_2$ (18.0 mg, 0.05 mmol Fe) in mesitylene (2 mL) was combined with a solution of $[\text{Ru}(\text{cod})(\text{cot})]$ (47.0 mg, 0.15 mmol Ru) in mesitylene (2 mL) in a Fischer–Porter bottle (70 mL). The SILP (500 mg) was added to the solution of the metal precursors along with mesitylene (1 mL) and the reaction mixture was stirred under argon at room temperature for 30 min. The Fischer–Porter bottle was evacuated and backfilled with H_2 and the suspension was stirred under H_2 (3 bar) at 150 °C for 18 h. Under this reducing environment a black powder was obtained indicating the immobilization of the NPs onto the SILP. Mesitylene was decanted and $\text{Fe}_{25}\text{Ru}_{75}\text{@SILP}$ were washed with fresh toluene (3×3 mL) and dried *in vacuo* at rt for 1 h.

Autoclave reactions

The substrate (25 eq., 0.1 mmol), catalyst (10 mg, containing 0.004 mmol metal) and solvent (mesitylene, 0.5 mL) were weighed inside of a glovebox into the autoclave. The autoclave was closed, transferred out of the glovebox and pressurized with 50 bar hydrogen. After heating the autoclave under stirring (500 rpm) to 175 °C for 16 h, it was cooled down in a water bath and depressurized. The GC samples were prepared by adding acetone (250 mg) to the reaction mixture and taking a sample through a syringe with a filter. To the sample, decane (internal standard, 20 mg) and acetone (200 mg) were added. For all amines, methanol/dioxane (1/1) was used as solvent for the GC samples.

Conflicts of interest

There are no conflicts to declare.

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