Three step synthesis of benzylacetone and 4-(4-methoxyphenyl)butan-2-one in flow using micropacked bed reactors

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HIGHLIGHTS

• Three catalytic reactors and a separator were combined in a telescoped flow system.
• Telescoped flow synthesis allowed different operating conditions for each reaction.
• Telescoped flow achieved higher yields than batch for production of ketones.

GRAPHICAL ABSTRACT

The synthesis of benzylacetone from benzyl alcohol and of 4-(4-methoxyphenyl)butan-2-one from 4-methoxybenzyl alcohol, which were previously performed in a batch cascade, were successfully performed in a telescoped flow system consisting of three micropacked bed reactors and a tube-in-tube membrane to remove oxygen. The system consisted of approximately 10 mg of 1 wt% AuPd/TiO\textsubscript{2} catalyst for oxidation, 150–250 mg of anatase TiO\textsubscript{2} for C–C coupling and 10 mg of 1 wt% Pt/TiO\textsubscript{2} for reduction, operating at 115 °C, 130 °C and 120 °C respectively. Oxygen and hydrogen flowrates were 2 and 1.5 NmL/min and alcohol solution inlet flowrates were 10–80 µL/min, while the system operated at a back pressure of 5 barg. This system achieved significantly increased yields of benzylacetone compared to the batch cascade (56% compared to 8%) and slightly increased yields of 4-(4-methoxyphenyl)butan-2-one (48% compared to 41% when using the same catalyst supports). The major advantage of the telescoped flow system was the ability to separate the three reactions, so that each reaction could have its own catalyst and operating conditions, which led to significant process intensification.

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ABSTRACT

The synthesis of benzylacetone from benzyl alcohol and of 4-(4-methoxyphenyl)butan-2-one from 4-methoxybenzyl alcohol, which were previously performed in a batch cascade, were successfully performed in a telescoped flow system consisting of three micropacked bed reactors and a tube-in-tube membrane to remove oxygen. The system consisted of approximately 10 mg of 1 wt% AuPd/TiO\textsubscript{2} catalyst for oxidation, 150–250 mg of anatase TiO\textsubscript{2} for C–C coupling and 10 mg of 1 wt% Pt/TiO\textsubscript{2} for reduction, operating at 115 °C, 130 °C and 120 °C respectively. Oxygen and hydrogen flowrates were 2 and 1.5 NmL/min and alcohol solution inlet flowrates were 10–80 µL/min, while the system operated at a back pressure of 5 barg. This system achieved significantly increased yields of benzylacetone compared to the batch cascade (56% compared to 8%) and slightly increased yields of 4-(4-methoxyphenyl)butan-2-one (48% compared to 41% when using the same catalyst supports). The major advantage of the telescoped flow system was the ability to separate the three reactions, so that each reaction could have its own catalyst and operating conditions, which led to significant process intensification.

1. Introduction

Over the last decade there has been a significant increase in both the number and complexity of telescoped flow synthesis, where flow reactors and separator systems are connected together in series to enable the synthesis of complex products. A large number of telescoped reactions have already been reported in the literature in a range of research areas including active pharmaceutical ingredient (API) synthesis [1–9], biofuels [10–12] and the fine chemicals industry [13–18]. These telescoped systems offer many opportunities for process intensification due to the inherent advantages of flow reactors including increased rates of heat and mass transport, access to a wider

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range of reaction conditions, increased safety and improved control of reaction conditions [19–22]. This recent progress in telescoped systems has largely been enabled by developments in continuous flow downstream processing unit operations, such as liquid-liquid separation [23,24], extraction [18], gas-liquid separation [25] and crystallisation [1], as well as by developments in automation allowing easier control of complicated systems [26].

Despite the considerable progress made in telescoped systems, there are still challenges to be overcome, one of the most important being solid handling, including both solids formed during reaction and solids added to the reactor [27]. In flow reactors, pumping solids in slurries is difficult, especially for microreactors, and can often lead to reactor clogging and blockages making systems unreliable and preventing prolonged continuous operation [26]. While there are various solutions for lab scale slurry reactors including the use of sonication [28,29], agitation [30], droplet flow [31], gas-liquid slug flow [32] or specially designed reactors including cascade of CSTRs [33–35], these solutions increase system complexity. Furthermore, many of these solutions require the slurry to flow into and out of the reactor [28–35] instead of being retained in the reactor, which may be required in a telescoped flow system if it is desired to use different slurry catalysts in sequential reactors. Retaining the slurry in the reactor is more difficult as it requires filtration before the reactor outlet, leading to problems with clogging. For this reason most telescoped systems which require solid catalysts avoid slurry reactors and instead rely on fixed bed reactors where the catalyst is held in place, allowing for easy separation of the gas or liquid reagents from the solid catalyst [22].

In contrast to a telescoped system, a batch cascade is where multiple reactions occur in a single reactor flask without intermediate workup. Batch cascades are a promising way to reduce the required number of unit operations in a synthesis and increase atom economy [36,37]. Batch cascades require multi-functional catalysts or else using multiple catalysts in the same flask that do not negatively interfere with each other. Additionally, all reactions must occur at the same temperature and pressure. These requirements result in a constrained design space and it is anticipated that switching from batch cascade to telescoped flow, where the reactions are separated into different reactors, could alleviate these requirements and allow for process intensification. The objective of this work is to convert a recently published batch cascade process to a telescoped flow process to demonstrate the widened design space offered by telescoped systems, as well as highlighting the difficulties in converting a multistep synthesis from batch to flow.

The reaction systems studied in this work are the multistep synthesis of benzylacetone (4) from benzyl alcohol (1) and of 4-(4-methoxyphenyl)butan-2-one (4) from 4-methoxybenzyl alcohol (1), via oxidation, aldol condensation and reduction, as shown in Fig. 1. Note that the oxidation step produces a number of side products, as multiple reactions occur in parallel including dehydrogenation, disproportionation and hydrogenolysis [38]. Two feed molecules were studied to examine the flexibility of the flow system to different substituted groups, as in some cases the batch cascade showed dramatically different yields for different substituted feed molecules [39]. These reaction systems were chosen as the products have high commercial value as food additives, insect attractants and fragrances [39,40] and because the recently developed batch cascade was more selective and produced less waste than current commercial production methods using Friedel-Crafts alkylations [40,41]. The catalysts used in the batch study were Au-Pd nanoparticles supported on MgO or TiO2, and the optimum catalyst was found to be AuPd/MgO [39]. However, it was not possible to use MgO supported catalysts in flow, as the MgO support did not retain its mechanical integrity; particles broke into smaller ones leading to blockages during prolonged use. The instability of MgO particles is attributed to the hydroxilation of MgO to Mg(OH)2 in the presence of water, which has been previously reported for this system [39,42]. Therefore only TiO2 supported catalysts were available for use in the telescoped system. All three of these reaction steps have been reported in the literature with a variety of different catalysts and reactor types, however, this work is the first to perform the telescoped synthesis in flow. Benzyl alcohol oxidation with molecular oxygen has been studied extensively [43–51] in both solvent and solvent free conditions, and the catalyst used in this work, AuPd/TiO2 is one of the most active, as it can achieve TOF greater than 10,000 h$^{-1}$ [50], reasonable selectivity [46] and its deactivation behaviour has been optimised [47]. Aldol condensation reactions are known to benefit from bifunctional catalysts which possess both acid and basic sites [52] and they have been studied with a wide range of catalysts including metal oxides (MgO, ZrO2 and TiO2) [53–55], double layered hydroxides [56] and amine-functiona-
lised SBA-15, ZrO2 and TiO2 [52]. In this work, anatase TiO2 was chosen because it was desired to use the same catalysts as in the batch study [39]. The reduction of benzaldehyde has been successfully demonstrated using Pd nanoparticles in packed beds [57] and in monolith reactors [58] and in this work both Pd/TiO2 and Pt/TiO2 were tested in packed bed reactors.

2. Materials and methods

2.1. Catalyst preparation

The 1 wt% 65.35 (weight ratio) Au-Pd/TiO2 catalyst was prepared by a modified impregnation method, where a round bottom flask was charged with HAuCl4 (0.53 mL, 12.25 mgAu/mL), PdCl2 dissolved in 0.58 M HCl (0.97 mL, 6 mgPd/mL) and water (13.5 mL) following a method described in previously published work [47]. The monometallic 1 wt% Pd/TiO2 and Pt/TiO2 catalysts were prepared by impregnation, where the metal salt PdCl2 or PtCl2 was dissolved in a small amount of
water, to which TiO₂ was added. The slurry was stirred and heated until a paste was formed, then dried (110 °C, 16 h) and later calcined at 400 °C for 3 h. For all the supported nanoparticle catalysts used, the TiO₂ support was P25 TiO₂ (Evonik). TiO₂ was used as a coupling catalyst and initially a 21 nm particle size nanopowder (Sigma Aldrich, 99.5% pure, 75% anatase and 25% rutile) was used. However, after studying the coupling reaction in isolation it was found that pure anatase TiO₂ was more active and subsequently for all telescoped experiments a 31 nm nanopowder anatase TiO₂ was used (Alfa Aesar). All of the catalysts used in this study were pelleted with 4000 N force using a pellet press, then ground and sieved to give the appropriate particle size range, typically 63–75 µm or 90–120 µm. The proportion of catalyst attaining the desired sieve fraction was 10–20%. BET studies showed only a small reduction in surface area, from 58.7 m²/g to 53.65 m²/g after pelletisation.

2.2. Reactor design & experimental set up

The reactions were carried out in both silicon-glass microreactors and in tube capillary reactors. The silicon-glass microreactors consisted of serpentine channels of dimensions 600 µm width, 300 µm height and 190 mm length with rectangular posts at the outlet to retain the solid catalyst. These reactors have been used in earlier work and their fabrication using photolithography and DRIE is described elsewhere [49]. The tube capillary reactors consisted of PTFE tubing with 1.587 mm O.D. and 1 mm I.D. (VICI Jour). The catalyst was retained in the tubing by use of a nickel mesh (Tecan, UK) of 25 µm thickness and 25 µm diameter holes which was held in place with compressive force between a PEEK union (Upchurch) and a PEEK ferrule (Upchurch). The tube reactor was used when the catalyst mass required surpassed the maximum catalyst loading of the silicon-glass microreactors (approximately 40 mg of TiO₂). The silicon-glass reactor was heated using heating cartridges in a chuck enclosed in ceramic packaging for insulation, while the tube reactor was heated in a stirred oil bath. In both cases the catalysts were loaded into the reactors by applying vacuum to the reactor inlet and introducing a known mass of catalyst through the reactor inlet. The reactors were weighed before and after loading to measure the mass of catalyst.

A number of experiments were conducted including studying the oxidation, coupling and reduction reactions independently and also in series. Additionally, in many cases when studying reactions in isolation small amounts of likely impurities (side products, unreacted reagents from upstream reactions) were added to study reaction inhibition. The experimental set-up varied depending on the reaction(s) being studied, but in all cases the same equipment was used. Liquid feeds (alcohol in acetone solution) were introduced using stainless steel 8 mL syringes (Harvard Apparatus) and syringe pumps (Harvard Apparatus, Ph.D. Ultra). The gases were fed using mass flow controllers (Brooks 5850TR) and the pressure at the reactor outlet was controlled using a back pressure regulator (Swagelok KBP series, 250 PSIG). The liquid product was collected in a custom made PEEK collection vessel, which separated the liquid from the gas via gravity. A schematic of the experimental set-up for gas-liquid-solid reactions, such as the oxidation and reduction reactions, is shown in Fig. 2. For liquid-solid experiments (coupling reaction) the gas inlet on the microreactor chip was closed and the system was pressurised with nitrogen gas flowing directly into the PEEK collection vessel.

The experimental set-up for the telescoped flow system is shown in Fig. 3, where the reactors were connected in series with a tube-in-tube membrane separator immediately after the oxidation reactor to remove the oxygen gas. The membrane separator consisted of a 65 cm long Teflon AF-2400 tubular membrane (Biogeneral, U.S.) of I.D. 0.8 mm and thickness 0.1 mm within a 3.175 mm O.D., 2.4 mm I.D. PTFE tube (VICI Jour). The tube-in-tube membrane device operated with the gas-liquid flow in the inner tube at 6 bar pressure while the outer tube was connected to a vacuum pump (KNF labs) providing −500 mbar vacuum. In the literature, such Teflon AF-2400 membranes operated satisfactorily at even higher pressure differences of 13 bar [59] or at elevated temperatures of 120 °C [45].

2.3. Product analysis

The liquid product was analysed off-line using gas chromatography (Agilent, 7820A) with a FID detector and an Agilent DB-624 capillary column. Mesitylene was used as an internal standard to allow for volume change corrections associated with the generation of water and the loss of acetone through evaporation upon sample collection and depressurisation. A carbon balance where the concentration of the reactant in the feed was compared to the concentration of the reaction products in the outlet stream is shown in Eq. (1), where C represents concentration (M) and pi and fi represent the stoichiometric coefficients of the i-th product species and the alcohol feed (f = 1).

\[
\sum_{i} C_{i} \cdot f_{i} = \sum_{i} C_{i} \cdot p_{i}
\]

For the benzylacetone system the carbon balance was found to close within 95% when studying reactions in isolation, and to within 70–90% when studying the entire multistep system. All chemical species, benzyl alcohol (1), benzaldehyde (2), toluene (2b), benzene (2c), benzoic acid (2d), benzyl benzoate (2e), benzalacetone (3), dibenzalacetone (3b), benzylacetone (4), 4-phenylbutan-2-ol (5), 4-methoxybenzyl alcohol (1), 4-methoxybenzaldehde (2), 4-methylanisole (2b), anisole (2c), 4-(4-methoxyphenyl) butan-2-one (4), 4-(4-methoxyphenyl)butan-2-ol (5), acetone and mesitylene were from Sigma Aldrich.
Carbon Balance = \frac{C_{\text{Reactant In}}\left(C_{\text{Reactant Out}} + \sum_{i} f_{i} C_{\text{Product Out}}\right)}{C_{\text{Reactant In}}} \quad (1)

Reactant feed conversion, \( X \), was calculated using Eq. (2) and the selectivity, \( S_{i} \), using Eq. (3). When studying reactions in isolation the average reaction rate, \( r_{\text{avg}} \) (mol/s/g catalyst) was calculated according to Eq. (4), where \( m \) is the catalyst mass (g) and \( u \) is the inlet liquid flowrate (L/s). The reaction rate varies along the length of the packed bed, however this average reaction rate is still a useful parameter for comparison purposes. The yield of the reaction to species \( i \), \( Y_{i} \), represents the fraction of the reagent that forms species \( i \), and it was calculated according to Eq. (5) which takes into account the reaction stoichiometry. The catalyst contact time in flow was calculated by dividing the catalyst mass by the mass flow rate of alcohol, as shown in Eq. (6). In the batch system the catalyst contact time was calculated by dividing the product of reaction time \( t \) and catalyst mass by the mass of alcohol in the reactor, which is given by the product of the reactor volume \( V \), feed concentration and molecular weight of the feed alcohol \( MW_{\text{Reactant}} \) as shown in Eq. (7).

\[ X = \frac{C_{\text{Reactant In}} - C_{\text{Reactant Out}}}{C_{\text{Reactant In}}} \quad (2) \]

\[ S_{i} = \frac{f_{i} C_{\text{Product Out}}}{C_{\text{Reactant In}} - C_{\text{Reactant Out}}} \quad (3) \]

\[ r_{\text{avg}} = \frac{\left(C_{\text{Reactant In}} - C_{\text{Reactant Out}}\right) u}{m} \quad (4) \]

\[ Y_{i} = \frac{f_{i} C_{\text{Product Out}}}{C_{\text{Reactant In}}} \quad (5) \]

Flow Catalyst Contact Time = \frac{m}{u \cdot MW_{\text{Reactant}} \cdot C_{\text{Reactant}}} \quad (6)

Batch Catalyst Contact Time = \frac{t \cdot m}{V \cdot MW_{\text{Reactant}} \cdot C_{\text{Reactant}}} \quad (7)

3. Results & discussion

3.1. Batch cascade vs telescoped flow

The telescoped flow synthesis of both benzylacetone and 4-(4-methoxyphenyl)butan-2-one were successfully performed using the experimental set-up shown in Fig. 3. For both feed molecules the same catalysts were used (1 wt% AuPd/TiO₂, anatase TiO₂ and 1 wt% Pt/TiO₂) and the standard experimental conditions were 115 °C, 130 °C and 120 °C for the oxidation, coupling and reduction reactions, 2 NmL/min oxygen gas flowrate and 1.5 NmL/min hydrogen flowrate with the system back pressure regulator set to 5 barg. Variable feed concentrations, liquid flow rates and catalyst masses were used for both systems. The operating conditions were chosen based on information obtained studying the reactions in isolation by performing parametric studies (varying liquid and gas flow rates, temperature, pressure and particle sizes), which is discussed in the Supplementary Information.

Maximum yields of 56% for benzylacetone were obtained in telescoped flow when using the most dilute feed concentrations studied (0.72 M). This is a major increase on the 8% yield obtained in batch as shown in Fig. 4 [39], however the low batch yield may be partially due to the fact that the batch system was optimised for the 4-methoxybenzyl alcohol feed and not the benzyl alcohol feed. The yield from the batch synthesis was calculated by multiplying the reported conversion by the reported selectivity and multiplying again by a factor for the unidentified species, as the reported selectivity was calculated as the fraction of desired product over the sum of all identified species (only 70–80% of species were identified) [39]. The dramatic increase in yield is attributed to the greater flexibility of the telescoped flow system being able to perform each reaction in a separate reactor, allowing the choice of different catalysts and operating condition for each reaction. It is also expected that the multistep flow system has greater potential for optimisation. In this work the choice of catalyst was limited to those previously used in batch to focus on the study of the reactor system and widened design space instead of catalyst performance. However, if more suitable aldol condensation catalysts were used, such as metal oxides (ZrO₂) [53–55], double layered hydroxides [56] and amine functionalised SBA-15, ZrO₂ and TiO₂ [52], it is expected that even better performance could be achieved.

As shown in Fig. 4, the telescoped flow system showed a higher yield of the 4-(4-methoxyphenyl)butan-2-one than the one-pot synthesis when both systems used TiO₂ supported catalysts (48% compared to 41%). However, when the batch system used the MgO catalyst, the yield exceeded that achieved in flow (48% compared to 63%). This was largely because the AuPd/TiO₂ catalyst which was used in flow, was not able to achieve the high selectivity in the oxidation reaction that the AuPd/MgO catalyst was able to achieve in batch. Unfortunately, it was not possible to use the MgO support in micropacked bed reactors, as the support particles broke into smaller particles and clogged the reactors. This demonstrates a drawback of current telescoped flow systems, in that the catalyst must be compatible with packed beds as current slurry reactors which retain the slurry in the reactor (via filtration at the reactor exit) are not reliable enough to integrate with a telescoped system. The majority of reliable slurry reactors today instead have the slurry passing out of the reactor outlet [28–35].
The main advantage of the multistep flow system was not just in yield improvement but in smaller catalyst requirements and increased productivity. Primarily, the multistep flow system allowed for the replacement of significant amounts of the expensive nanoparticle supported catalysts with the cheaper TiO$_2$ catalyst. To process a similar amount (0.01 mol) of either feed alcohol, the batch system used 500 mg of AuPd supported catalyst, while the flow system used only 20 mg of AuPd supported catalysts (approximately 10 mg of AuPd/TiO$_2$ and 10 mg of Pt/TiO$_2$) and 150–250 mg of the cheap anatase TiO$_2$. Additionally, although the catalyst contact time per gram of alcohol varied in the flow experiments due to the range of inlet liquid flowrates, feed concentrations and catalyst masses used, the experiments still suggest that the catalyst was being used more efficiently in flow than in batch. For the benzylacetone experiment shown in Fig. 4, which produced the maximum yield, the catalyst contact times were only 55 h of catalysis/galcohol and 65 h of catalysis/galcohol for the oxidation and reduction reactions and 1190 h of catalysis/galcohol for the coupling reaction. In comparison, in the one-pot system the catalyst contact time was calculated by dividing the product of reaction time (22 h) and catalyst mass (500 mg) by the mass of alcohol (1.3 g), and for the benzalacetone one-pot system this was 8460 h of catalysis/galcohol. Thus, the total amount of catalysts utilised in the benzyl alcohol multistep flow system was approximately 6.5 times lower than in the one-pot system. For the 4-methoxybenzyl alcohol system the same comparisons are harder to make, as the batch reaction reached completion before the end of the 22 h.

### 3.2. Catalyst inhibition

It was found that the yield of the telescoped flow system decreased with increasing feed concentration, as shown in Figs. 5 and 6. While this was partly due to the decreasing catalyst contact time, it was discovered that the coupling reaction in the multistep system suffered from extreme water inhibition due to the generation of water in both the oxidation and coupling reactions. This resulted in the requirement of using dilute feed concentrations of less than 1.3 M as well as using excess TiO$_2$ to compensate for reduced activity. The water inhibition can be seen in Fig. 5, where the amount of unreacted alcohol (1) and unreacted aldehyde (2) was significantly higher for the 3.06 M feed than the other lower concentration experiments. The result of the 3.06 M feed experiment shown in Fig. 5 suggests that both the oxidation and coupling reactions could be inhibited by high feed concentrations, as there were large amounts of unreacted benzyl alcohol (1) as well as benzaldehyde (2). However, it was later demonstrated that only the coupling reaction was inhibited by water and not the oxidation reaction, which proceeded with near 100% conversion in all cases. The benzyl alcohol (1) in the outlet stream was actually produced from the reduction of unreacted benzaldehyde (2) in the reduction reactor. This was confirmed in the 3.06 M experiment by replacing the hydrogen gas with nitrogen in the reduction reactor; the amount of unreacted benzyl alcohol then dropped to almost zero and the yield of unreacted benzaldehyde rose dramatically to 70%. Studies of the coupling reaction in isolation showed that water concentration of just 2.75 wt% resulted in a 50% drop in activity, as shown in the Supplementary Information. Assuming complete conversion of all benzyl alcohol feed, a feed concentration of only 1.3 M would produce the 2.75 wt% water necessary for water inhibition. This suggests that the 1.83 M experiment in Fig. 5 is also suffering from water inhibition, but that there is a sufficient excess of TiO$_2$ catalyst to partially compensate for this. The reduction reaction was also found to be inhibited by water. However, a higher concentration of water was required to inhibit the reduction reaction than the coupling reaction; hence this did not impose any extra design constraints on the multistep flow system. The same water inhibition was observed in the synthesis of 4-(4-Methoxyphenyl)butan-2-one as shown in Fig. 6.

The coupling reaction was inhibited by even very low concentrations of benzoic acid (2d); just 0.03 M benzoic acid in the coupling reactor feed stream led to a drop in the reaction rate of 30%. Unfortunately, trace amounts of benzoic acid are unavoidable in the oxidation reaction and even after attempting to tune the reaction conditions to minimise its formation, the concentration of benzoic acid was approximately 0.03 M resulting in some inhibition. Inhibition studies conducted for the reduction reaction in isolation also showed that the reduction reaction was inhibited by various organic species, including benzaldehyde (2) and dibenzalacetone (3b) as shown in the Supplementary Information. However, this was not found to be significant in the telescoped flow system, as the concentration of the inhibiting by-products did not reach the high values that were added to...
that used in the telescoped system (9.4 mg of 1 wt% AuPd/TiO₂ at similar conditions or even with a lower catalyst contact time than the oxidation reaction decreased. However, when the oxidation reaction was carried out at similar conditions constant. The results in Fig. 7 for the 4-(4-methoxyphenyl)benzylacetone system, show the expected trend that at lower catalyst contact times the conversion of both the oxidation and the coupling reactions respectively.

3.3. Effect of catalyst contact time

The effect of catalyst contact time was investigated for both the benzylacetone and the 4-(4-methoxyphenyl)butan-2-one system by changing the inlet liquid feed flowrate while keeping all other conditions constant. The results in Fig. 7 for the 4-(4-methoxyphenyl)butan-2-one system, show the expected trend that at lower catalyst contact times the conversion of both the oxidation and the coupling reaction decreased. However, when the oxidation reaction was carried out at similar conditions or even with a lower catalyst contact time than that used in the telescoped system (9.4 mg of 1 wt% AuPd/TiO₂ at 115 °C, 2.2 M liquid flow rate, 20 µL/min liquid flowrate of 0.96 M 4-methoxybenzyl alcohol feed at 6 bar back pressure) greater than 95% conversion was achieved. Therefore it is suspected that the unreacted alcohol (1) in the outlet is not due to lower performance in the oxidation reactor but due to unreacted aldehyde (2) being converted back to alcohol in the reduction reactor. The same trend was observed for the benzylacetone system, as shown in the Supplementary Information.

3.4. Oxygen separation

The removal of oxygen gas was found to be critical in the multistep flow synthesis for a number of reasons including safety and optimising the coupling and reduction reactions. While microreactors are known for their increased safety [60,61], oxygen gas removal was performed to prevent oxygen mixing with hydrogen gas downstream and creating an explosive atmosphere in the bypass vessel, which had a sufficiently large volume (200 mL) to present a hazard under the high operation pressure. It was also discovered that the rate of the coupling reaction dropped significantly if either the liquid feed was allowed to vapourise or if any gas was flowing with the liquid; this is shown in the Supplementary Information. This prevented the integration of the coupling reaction with either the oxidation or reduction reactions and it required oxygen gas removal immediately after the oxidation reaction. It is possible that the decrease in coupling reaction rate in the presence of gas was due to loss of acetone in the liquid phase by evaporation, leading to a lower concentration available for reaction and increased difficulty in removing the resultant products from the catalyst surface. The removal of oxygen gas was achieved using a tube-in-tube membrane, as complete removal of oxygen could be reliably achieved even under conditions where pressure fluctuated significantly, such as during sample taking or when changing the flowrates. In addition to oxygen, acetone also permeated through the membrane and was lost from the reaction mixture in low quantities (approximately 7% of the initial acetone was lost to evaporation, when the initial concentration of acetone was typically 9 M or 70% by mass of the feed solution), but this loss was considered acceptable.

3.5. Coupling catalyst issues and effect on telescoped reactor design

The coupling reaction was found to be a bottleneck in this system, as its reaction rate without any impurities in the feed was only 1.5 × 10⁻⁵ mol/g/s, approximately an order of magnitude lower than that of the oxidation or reduction reactions, and then its reaction rate was further reduced due to catalyst inhibition as mentioned previously. The low reaction rate led to the use of more than 150 mg of anatase TiO₂, which exceeded the 40 mg maximum capacity of the silicon-glass microreactors, requiring the use of a packed tube reactor. This larger packed bed caused a pressure drop of about 1.5 bar and due to its large catalyst mass it clogged on more than one occasion. The maximum inlet liquid flowrate was limited at 40 µL/min, as higher flowrates required even longer catalyst beds, greater pressure drops and increased the likelihood of clogging the reactor. Therefore this catalyst restricted the maximum liquid flowrate possible and hence limited the productivity of this system. Furthermore, heating the packed tube in an oil bath required long connection tubing that increased the system residence time. The study of the coupling reaction, included in the Supplementary Information, suggests that this catalyst may not be ideal for the multistep flow system, and other catalysts may provide improved performance. The key findings regarding the coupling reaction were that at temperatures > 120 °C, which are the best operating conditions for the multistep system, the reaction shows significant external mass transfer resistances and is inhibited by the product. The external mass transfer resistances are demonstrated in Fig. 8 at high temperatures of 140 °C, where the rate of reaction increased with increasing liquid flowrate for a constant catalyst contact time. At lower temperature of 100 °C external mass transfer resistances are not observed, indicating that the

Fig. 7. Yield for the 4-(4-methoxyphenyl)butan-2-one system at three different inlet liquid flowrates using a 1.1 M feed of 4-methoxybenzyl alcohol. The standard experimental conditions for temperatures, pressure and gas flowrates were used with 10 mg, 149 mg and 11 mg of catalysts for the oxidation, coupling and reduction reactions respectively.

![Graph showing yield vs. inlet liquid flowrate](image)

Fig. 8. Average reaction rate (corrected for deactivation) against inlet liquid flowrate for the coupling reaction of benzaldehyde with acetone. Experimental conditions were 6 barg, 2.2 M benzaldehyde in acetone, 63–75 µm TiO₂ catalyst. Catalyst amounts were chosen to give the desired catalyst contact times (mg/min/µL), except of the 100 °C experiment where 15 mg of catalyst was used for all experiments.

![Graph showing average reaction rate vs. inlet liquid flowrate](image)
anatase TiO\(_2\) catalyst was not very suitable for a multistep flow system. This information suggested that this catalyst did not perform well in the specific reaction conditions or was not adapted to the flow system. A side product, dibenzalacetone (3b), was added to the feed to increase the total conversion of benzalacetone (3). The reaction rate was observed to decrease as the concentrations increased, indicating product inhibition. This behavior was more pronounced at higher temperatures, as shown in Fig. 9.

Conversion was observed to be less than 40%, which explained why the catalyst contact time was a critical factor in determining the overall yield of the product. A large catalyst bed would be needed, and high flowrates would cause an unreasonable pressure drop. However, despite these challenges, the telescoped flow system was still able to produce high yields of the final product, demonstrating its potential for industrial application.

Deactivation was found to be a serious problem in the batch cascade study. Reusing catalyst from one batch to the next led to a fall in conversion from 100% to 46% after a single use in hydrogen atmosphere. Deactivation has also been observed in flow for the oxidation of benzyl alcohol, where it was shown to be related to the catalyst formulation and preparation method [47]. Similarly, in this multistep flow system deactivation was also found to be a significant problem. Deactivation of up to 50% in 8 h of operation was observed when studying the coupling and reduction reactions in isolation, where a colour change of the catalyst was observed, forming a front that travelled down the length of the packed bed (shown in the Supplementary Information). This colour change provides further evidence to the hypothesis originally suggested in the batch study that deactivation is linked to adsorption of carbon species on the catalyst surface [39].

### 3.6. Catalyst deactivation

Deactivation was found to be a serious problem in the batch cascade study, where reusing catalyst from one batch to the next led to a fall in conversion from 100% to 46% after a single use in hydrogen atmosphere and from 96% to 0% in nitrogen atmosphere [39]. Deactivation has also been observed in flow for the oxidation of benzyl alcohol, where it was shown to be related to the catalyst formulation and preparation method [47]. Similarly, in this multistep flow system deactivation was also found to be a significant problem. Deactivation of up to 50% in 8 h of operation was observed when studying the coupling and reduction reactions in isolation, where a colour change of the catalyst was observed, forming a front that travelled down the length of the packed bed (shown in the Supplementary Information). This colour change provides further evidence to the hypothesis originally suggested in the batch study that deactivation is linked to adsorption of carbon species on the catalyst surface [39]. The problem of using deactivating catalysts in telescoped systems could be partially overcome by using an excess of catalyst. This was demonstrated for the benzalacetone telescoped system, as the yield of benzalacetone remained stable after 6.5 h of continuous operation, as shown in Fig. 10. While the duration of this flow experiment (6.5 h) is far shorter than the batch (22 h), the amount of alcohol processed (0.01 mol in flow and 0.009 mol in batch) is comparable, hence showing that the effect of deactivation in flow was reduced. Despite the relative stability of the system over the 6.5 h experiment, catalysts with higher stability would be needed for industrial application.

### 4. Conclusions

The multistep synthesis of benzalacetone and 4-(4-methoxyphenyl)butan-2-one using AuPd, Pd and Pt supported catalysts was successfully converted from batch cascade to a telescoped flow system demonstrating both the advantages and challenges of flow systems. The most critical advantage of the telescoped flow system compared to the batch cascade was the ability to separate the three reactions, oxidation, coupling and reduction, hence allowing more freedom to choose different catalysts and operating conditions without the necessity of finding a compromise among reactions. In this case this freedom enabled process intensification, reducing catalyst contact requirements by a factor of 6.5 for the benzalacetone system and allowing the replacement of significant amounts of expensive nanoparticle supported catalysts with cheaper anatase TiO\(_2\) catalyst. Additionally, the flow system attained higher yields of 4-(4-methoxyphenyl)butan-2-one than the batch system when both systems used TiO\(_2\) supported catalysts (48% compared to 41%). However, the drawback of the telescoped flow system was that it was not possible to use the MgO supported catalyst that was found to be better for this reaction in the batch cascade, due to the MgO not having suitable mechanical properties to be used in a micropacked bed. This resulted in the telescoped flow system not being able to achieve the highest yield of 4-(4-methoxyphenyl)butan-2-one of 63%, which was achieved in batch with the MgO supported catalyst. Despite this drawback, the telescoped flow system with TiO\(_2\) supported catalysts was able to outperform the batch flow system with the preferred MgO supported catalysts for the synthesis of benzalacetone (56% compared to 8%). While in this work the selection of catalysts for the telescoped flow system was restricted to those used in batch to focus on the effect of reactor configuration (batch vs flow), it is expected that the
performance of the telescoped flow system could be improved if the choice of catalysts is extended beyond the nanoparticle supported catalysts previously used in the batch system. This would be taking full advantage of the telescoped flow system’s extended design space and may overcome some of the challenges encountered in this work, including water and product inhibition.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/jcej.2018.09.137.

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