Continuous Heterogeneous Photocatalysis in Serial Micro-Batch Reactors

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A flow system for processing solids enables the efficient and scalable utilization of Heterogeneous photocatalysts for selective fluorinations.

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Continuous Heterogeneous Photocatalysis in Serial Micro-Batch Reactors

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Abstract: Solid reagents, leaching catalysts, and heterogeneous photocatalysts are commonly employed in batch processes but are ill suited for continuous flow chemistry. Heterogeneous catalysts for thermal reactions are typically used in packed bed reactors that cannot be penetrated by light and thus are not suitable for photocatalytic reactions involving solids. We demonstrate that serial micro-batch reactors (SMBRs) allow for the continuous utilization of solid materials together with liquids and gases in flow. This technology was utilized to develop selective and efficient fluoration protocols using a modified graphitic carbon nitride heterogeneous catalyst instead of costly homogeneous metal polypyridyl complexes. The merger of this inexpensive, recyclable catalyst and the SMBR approach gives access to sustainable and scalable photocatalysis.

Solid-liquid reactions are ubiquitous in chemical synthesis and are usually straightforward in batch, but they are less trivial in continuous flow devices. Packed bed reactors with heterogeneous catalysts embedded between filter units suffer from periodic deactivation or loss of catalyst due to leaching. Photochemistry on the other hand is perfectly suited for continuous flow chemistry. Heterogeneous catalysts for photocatalysis are commonly employed in batch processes, but are ill suited for continuous flow chemistry.

Difficulties associated with separation and recycling of these catalysts render heterogeneous or easily recyclable catalysts such as modified transition metal catalysts and semiconductors highly desirable for PRC. Graphitic carbon nitrides (g-C₃N₄), a class of metal-free polymers, are among the most promising materials for heterogeneous PRC.

Combining the advantages of continuous photocatalysis and g-C₃N₄ catalysis to provide an efficient and sustainable process is the most promising materials for heterogeneous PRC. Combining the advantages of continuous photocatalysis and g-C₃N₄ catalysis to provide an efficient and sustainable process is highly desirable for PRC. Graphitic carbon nitrides (g-C₃N₄), a class of metal-free polymers, are among the most promising materials for heterogeneous PRC.

Efficient irradiation is ensured by pumping a suspension through an irradiated coil reactor that is submerged in a thermostatic bath. The resulting SMBRs can be fed into an irradiated coil reactor that is submerged in a thermostatic bath.

Figure 1. Strategies for heterogeneous photocatalysis in flow. (A) Packed bed reactors are ideal for heterogeneous catalysis but suffer from insufficient light penetration. (B) A triphasic gas-liquid-solid system is used to deliver selective and efficient reactions (Fig. 1A). Small diameter packed bed reactors can overcome the irradiation problem but packing is tedious and channeling phenomena, as well as high pressure drops, are foreseeable drawbacks. Wall-coated reactors were previously utilized for continuous photocatalytic reactions however, stable and homogeneously distributed wall coatings are difficult to prepare and different for each heterogeneous catalyst. Efficient irradiation is ensured by pumping a suspension through a coil reactor but the solid material settles, leading to a heterogeneous distribution, irreproducible results, or clogging (Fig. 1B).

Dosing a catalyst suspension into a segmented gas-liquid stream should, based on suspension feeding into segmented flow regimes, provide a promising alternative for heterogeneous photocatalysis. Such triphasic systems resemble a series of small solid-liquid batch reactors (serial micro-batch reactors - SMBRs) that are separated by an inert gas spacer and “stirred” by toroidal currents (Fig. 1C).

Early solid feeding systems suffer from complex setups or costly equipment and are generally not suitable for continuous operation. To ensure straightforward access for the general community, SMBR generation was designed to be operationally simple and readily assembled from inexpensive, commercially available parts. The residence time and catalyst stoichiometry should be adjustable to enable detailed studies of all reaction parameters.

A flow setup for transformations on small scales meeting all requirements is a dosing system comprised of two parts. First, a stable gas-liquid segmented flow pattern is generated from the reaction solvent and nitrogen gas in a fluorinated ethylene propylene tube using a Y-shaped mixer (Fig. 2A). Subsequently, the catalyst suspension is added from below via a mixing Tee and a vertically mounted syringe pump that incorporates a magnetic stir bar. The resulting SMBRs can be fed into an irradiated coil reactor that is submerged in a thermostatic bath.

The residence time is adapted by changing the gas and/or liquid flow rate or the reactor volume while the catalyst stoichiometry...
can be varied by changing the rate of suspension dosing (Supporting Information). The system continuously produces identical SMBRs while all parameters can be changed during operation. The installation of a second catalyst addition unit enables a fully continuous operation by mimicking a dual syringe pump (Supporting Information).

A modified carbon nitride (CMB-C3N4) that is highly efficient for the photocatalytic production of hydrogen from water, served as heterogeneous photocatalyst for initial tests. This bench-stable semiconductor can be prepared in large quantities via a supramolecular approach from bulk chemicals and absorbs in the visible region. CMB-C3N4 shows minimal settling in viscous solvents such as ethylene glycol and 1-butyl-3-methylimidazolium tetrafluoroborate ([Bmim]BF4) a basis for the reliable preparation of a stable suspension for the accurate dosing of solid materials. Active mixing of the suspended catalyst feed via mechanical stirring improves reproducible additions, as negligible fluctuations over the course of addition were observed. The internal mixing phenomena caused by interior vortices inherent to the SMBRs result in a homogeneous and uniformly irradiated suspension of the semiconductor (Supporting Information).

With the robust dosing system in place, the SMBR concept was illustrated for heterogeneous photocatalysis. The well documented decarboxylative fluorination of phenoxyacetic acids using Selectfluor under PRC conditions served as the model transformation, as the triphasic SMBR system is perfectly suited for reactions involving gaseous reagents and/or byproducts. Batch experiments revealed that CMB-C3N4 can substitute Ru(bpy)32+ in the decarboxylative fluorination of phenoxyacetic acid (1) and can be reused several times without losing its catalytic activity (Supporting information). Stirring is of utmost importance for an efficient, reproducible batch process, and the reaction rate drops on larger scales.

Under SMBR conditions, the drawbacks associated with stirring can be elegantly addressed while the high irradiation efficiency inherent of the small dimensions simultaneously enhances the catalytic process. Monofluorinated compound 2 is almost quantitatively formed within only 14 minutes (Fig. 3A). The generated CO2 can immediately diffuse out of the liquid phase, to form expanded gas slugs and increase the overall flow rate without disturbing the integrity of the solid-liquid SMBRs (Fig. 3B). Following the reaction, a three-step extraction protocol provides the desired organofluorine compound without chromatography and allows for both the catalyst and suspension solvent ([Bmim]BF4) to be fully recovered (Fig. 3C).
The process can be easily scaled for the gram scale synthesis of 2 using a setup with two catalyst addition units (Supporting Information). Control experiments in the absence of either the semiconducting material or light did not result in an observable gas formation and only trace amounts of 2 were identified by NMR spectroscopy (Fig. 4A). Even when the reaction mixture was not degassed, the SMBR approach still produced 2 in excellent yield, whereas no reaction occurred in batch without prior degassing of the solution product was formed, but was not degassed, the SMBR approach still produced a higher conversion than the heterogeneous PRC. Moreover, reactor fouling with the ruthenium PRC resulted in a random flow pattern over time (Supporting Information). The in situ degassing of the SMBR system may occur via diffusion of dissolved oxygen into the gas segments. When Ru(bpy)₃²⁺ was used instead of CMB-C₃N₄ employing a segmented flow pattern without prior degassing of the solution product was formed, but the homogeneous catalyst suffered from significantly lower conversion than the heterogeneous PRC. Moreover, reactor fouling with the ruthenium PRC resulted in a random flow profile over time (Supporting Information).

Homogeneous PRC fail for continuous chemistry on larger scale due to irreproducible reaction parameters and the high risk of clogging. The SMBR fluorodecarboxylation produced a range of phenoxycetic acid derivatives (Fig. 4B). The hydrophobic nature of most products necessitated an acetone:water solvent mixture to avoid the generation of an “organic film” on the surface of the FEP coil (Supporting Information). Higher catalyst loading in combination with longer reaction times (up to 48 min) ensured sufficient conversions and yields similar to those obtained by the original protocol, as well as the uncatalyzed version using UV light.

The catalytic system was amenable to other fluorinations (Fig. 4C). The decarboxylative fluorination of 4-biphenylacetic acid (3) relied previously on a strongly oxidizing iridium(III) photocatalyst ([Ir(df(CF₃)ppy)₂(dtbppy)PF₆] in presence of a base, Under SMBR conditions, 3 was converted to 4 (73% yield) within 40 min at 50 °C without additional reagents.

**Figure 4.** Application of SMBR to heterogeneous photoredox catalysis. (A) Control experiments indicate an “in situ degassing” effect of the triphasic reaction system. (B) Scope of the fluorodecarboxylation of phenoxycetic acid derivatives. (C) Decarboxylative fluorination of 4-biphenylacetic acid and direct mono-fluorination of ethylbenzene and ibuprofen methyl ester. (D) Selectivity studies for the fluorination of ibuprofen. NMR yields are given in parenthesis. 2,6-lutidine was added to dissolve the starting material. Isolated yield in parenthesis. See Supporting Information for experimental details.
The direct benzylic monofluorination of ethylbenzene (5) was achieved using the heterogeneous photocatalyst and Selectfluor (5, Fig. 4D), a transformation usually carried out using photosensitizers such as 9-fluorenone.[27] The 45% yield of the reaction without any additive was significantly increased by addition of a strong acid (65%). The fluorination of ibuprofen methyl ester (7) was carried out under water-free conditions to avoid ester hydrolysis.

Selective late-stage fluorination of ibuprofen (9) at two different positions proved possible (Fig. 4E). The decarboxylative fluorination product 10 and benzylic C-H fluorination product 11 were identified by NMR analysis. In presence of a base, only 10 was selectively obtained in moderate yield. Without water as cosolvent under acidic conditions, the decarboxylative fluorination pathway was suppressed, to produce mainly 9 (Supporting Information).

In summary, we have developed a flow chemistry system that combines the advantages batch processing offers for heterogeneous materials with the benefits of flow photochemistry. The simple and efficient continuous utilization of heterogeneous photocatalysts serves as example and can be extended to many solid catalysts and reagents where packed bed reactors are non-ideal. The system is well suited for reactions involving gases. Three different fluorination protocols utilizing a heterogeneous organic semiconductor - which holds numerous advantages over standard metal catalysts - were developed and the methodology proved efficient, flexible, and robust on various scales and allowed for the site-selective, late-stage fluorination of ibuprofen. The SMBR concept, in combination with graphitic carbon nitride photocatalysis, will contribute to the development of more sustainable and industrially relevant photoedx processes in the future.

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