

Illuminating Progress: Recent Advances in Photoredox Catalysis

Introduction to Photoredox Catalysis

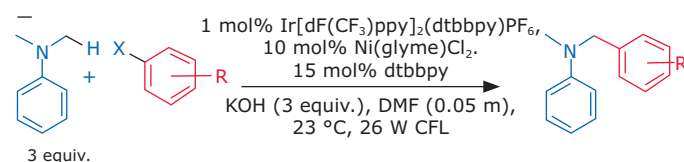
Photoredox catalysis is transforming synthetic organic chemistry, using light to drive chemical reactions with precision and efficiency. The method gained momentum after key discoveries in 2008 when researchers like David MacMillan and Dave Nicewicz showed that visible light could catalyze reactions traditionally relying on high-energy UV light, making synthesis more accessible and cost-effective (Nature Reviews Chemistry, 2019). This innovation introduced the use of visible-light-absorbing catalysts, allowing for Single-Electron Transfer (SET) processes and enabling the formation of complex carbon-carbon bonds. This capability is particularly valuable in the pharmaceutical industry, where photoredox catalysis is increasingly used to generate complex organic molecules for drug discovery.

Mechanisms of Photoredox Catalysis and SET Reactions

Photoredox catalysis harnesses light energy to facilitate single-electron transfers, forming reactive intermediates that drive a wide range of synthetic transformations (Prier et al. 2013). The open-shell pathways enabled by SET provide new avenues for bond formation, broadening the scope of organic reactions.

Zhang and Rueping (2024) explored sp³ C–H functionalizations through metallaphotoredox catalysis,

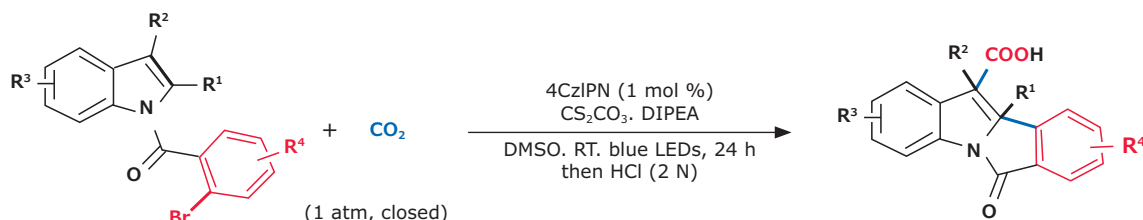
demonstrating that single-electron transfers allow for otherwise challenging modifications under milder conditions. Other notable transformations enabled by photoredox catalysis include cross-coupling, C–H functionalization, and arene functionalization, all of which are expanding the synthetic toolkit (Twilton et al., 2017).



Reaction Example: A C–H activation reaction using a photocatalyst like an iridium complex paired with a nickel catalyst. The chemical structure shows a generic aliphatic substrate (e.g., an alkane with sp³ C–H bond) reacting to form a C–C or C–heteroatom bond, with the iridium and nickel complex structures as catalysts.

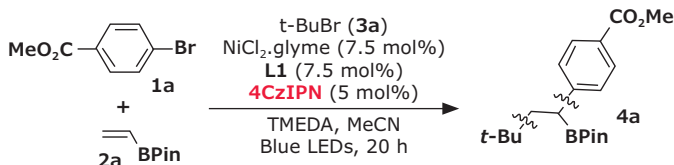
Advances in Photoredox Catalysis: Key Transformations and Applications

The ability of photoredox chemistry to form new bonds via open-shell pathways, allows for rapid and selective assembly of complex structures. Key examples include Wen-Jun Zhou's work on the reductive dearomative arylcarboxylation of indoles with CO₂ using visible-light photoredox catalysis (Nature Communications, 2020). This method presents a novel approach for CO₂ incorporation into organic compounds, offering greener and more efficient synthesis routes.



Reaction Example: CO₂ incorporation into an indole ring, with visible-light photoredox catalysis facilitating the transformation. The chemical structure shows the indole framework with the newly formed carboxylated product, along with CO₂ as a reactant and a visible-light photocatalyst like a Ru or Ir complex.

Similarly, Shang-Zheng Sun (2020) developed a dual-catalysis strategy for site-selective dicarbofunctionalization of vinyl boronates, a method that streamlines complex molecule construction and has broad applications in medicinal chemistry.



Reaction Example: A dual catalytic cycle involving both photoredox and palladium catalysis for site-selective dicarbofunctionalization. The chemical structure displays a vinyl boronate substrate undergoing C–C bond formation at the β -position, with Pd and a photocatalyst (e.g., $\text{Ru}(\text{bpy})_3^{2+}$).

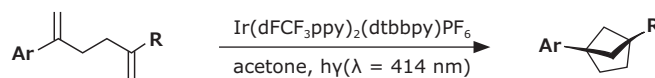
Photoredox Catalysis: Challenges and Innovations in Scaling Up

Scaling photoredox reactions from laboratory to industrial settings presents several technical hurdles, especially in achieving consistent light penetration in larger reaction vessels. Ken Schwieter, Innovation Manager at MilliporeSigma, highlights the Beer-Lambert law's implications for light travel in solutions, explaining that light typically only penetrates around 2 mm from the vessel wall, limiting reaction efficiency in larger vessels. This limitation can cause uneven irradiation and slower reaction rates, leading to inefficiencies in scale-up.

Flow chemistry offers a compelling solution, utilizing narrow microchannels — typically less than 1 mm in diameter — to ensure uniform light exposure across the reaction mixture. Sambiagio and Noël (2020) emphasize that this continuous-flow approach improves yield and consistency by maximizing photon flux and enhancing heat and mass transfer. Harper et al. (2019) demonstrated that laser-driven platforms can address light penetration issues in larger reactors, significantly enhancing reaction rates and productivity. These flow chemistry innovations are critical in pharmaceutical and agrochemical industries, where consistent scaling is essential.

Technological Innovations: Flow Photochemistry and High-Throughput Experimentation

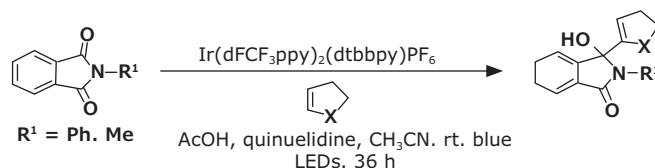
Innovations in flow photochemistry, high-throughput experimentation, and photoreactor design have catalyzed the industrial potential of photoredox reactions. Standardized photoreactors now offer consistent temperature control, mixing efficiency, and optimized light exposure, which is crucial for complex reactions that require precise conditions (Bulioni et al., 2021). For example, Rigotti (2022) developed an enantioselective crossed intramolecular [2 + 2] photocycloaddition mediated by a chiral Lewis acid, an application that highlights how precise photoreactor setups allow for improved selectivity in stereochemically complex reactions.



Reaction Example: Enantioselective [2 + 2] cycloaddition facilitated by a chiral Lewis acid under photoredox conditions. The chemical structure shows two alkenes undergoing cycloaddition to form a four-membered ring, with the Lewis acid catalyst structure influencing enantioselectivity.

Enhanced Reaction Control and Selectivity

Recent technological advances have addressed challenges in reaction control and selectivity, particularly through innovative photoreactor designs and controlled photon delivery. Using high-throughput experimentation and tailored reactor configurations, chemists can fine-tune parameters to optimize reaction conditions for scale-up. For example, the Morita–Baylis–Hillman reaction for non-electron-deficient olefins has been adapted for photoredox catalysis by Long-Hai Li (2022), demonstrating how precise control over light and reaction kinetics enables efficient reactions with challenging substrates.



Reaction Example: The photoredox-activated Morita–Baylis–Hillman reaction, where an electron-deficient alkene and aldehyde form a new C–C bond. The chemical structure shows a substrate with a β -position alkene reacting with an aldehyde under photoredox conditions, resulting in the addition product.

Standardized photoreactors eliminate the need for improvised setups, providing researchers with all-in-one platforms that ensure reproducible conditions. This consistency is essential for scaling photoredox catalysis to industrial levels. For instance, purpose-built photoreactors incorporate advanced temperature control, optimized light distribution, and improved mixing, which collectively enhance reaction efficiency and product yield (Sambiagio and Noël, 2019).

Applications and Industrial Impact of Photoredox Catalysis

As photoredox catalysis continues to advance, its applications in industry grow. Pharmaceutical and agrochemical companies are increasingly turning to photoredox methods to generate complex molecules more sustainably. This technique's ability to drive bond-forming reactions under mild conditions has opened doors to more eco-friendly and cost-effective synthetic pathways, particularly for compounds with multiple chiral centers or sensitive functional groups (Moschetta et al., 2017). High-throughput experimentation enables rapid screening of reaction conditions, expediting the discovery and optimization of new compounds for medicinal applications.

Additionally, photoredox catalysis offers a greener alternative to traditional thermally-driven processes by reducing the need for harsh reagents and high temperatures. This aligns with industry-wide goals to minimize environmental impact. For example, recent studies have demonstrated the effectiveness of photoredox catalysis in producing agrochemicals with fewer by-products, thereby enhancing sustainability across the chemical industry (Harper et al., 2018).

Future Directions: Expanding the Boundaries of Photoredox Catalysis

With ongoing advancements in photoredox catalysis, researchers are exploring new frontiers in organic synthesis. The integration of dual-catalysis strategies, such as those combining photoredox with transition metal or organocatalysis, is expanding the range of achievable transformations.

As equipment becomes more accessible and technology continues to mature, researchers can expect significant improvements in reproducibility and scalability, which will further expand the chemical space available for exploration. Advances in photoredox catalysis promise not only to refine synthetic methodologies but also to pave the way for new discoveries in drug development and materials science.

Conclusion

The progress made in photoredox catalysis over the past decade has transformed this field from a niche research area into a powerful tool in synthetic chemistry. As technological innovations continue to solve challenges in scaling and reproducibility, photoredox catalysis will play an increasingly vital role in industrial applications. Its potential for green, efficient synthesis aligns well with sustainability goals in pharmaceuticals, agrochemicals, and other sectors, marking it as a crucial tool in the future of organic synthesis.

Photoredox Chemistry Offerings

Explore our extensive portfolio of acridinium, iridium, and ruthenium catalysts, and other metal-free organic **photocatalysts** for any reaction design.

The **Penn PhD Photoreactor M2** is an advanced device with variable light sources (365 nm, 395 nm, 420 nm, 450 nm) and user-controlled temperature (30–50 °C) and light-intensity (max. 3.4 W) parameters.



The **SynLED Parallel Photoreactor 2.0** and **UV** are entry-level devices to run 16 reactions in parallel at fixed wavelength (450 and 365 nm, respectively), fixed temperature (30 °C), and light-intensity (1 W per LED).



References

- Buglioni L, Raymenants F, Slattery A, Zondag SDA, Noël T. 2021. Technological innovations in photochemistry for organic synthesis: flow chemistry, high-throughput experimentation, scale-up, and photoelectrochemistry. *Chem Rev.* 122(2): 1832–1885.
- Corcoran EB, McMullen JP, Lévesque F, Wismer MK, Naber JR. 2020. Photon equivalents as a parameter for scaling photoredox reactions in flow: translation of photocatalytic C–N cross-coupling from lab scale to multikilogram scale. *Angew Chem Int Ed.* 59(19): 7356–7361. <https://doi.org/10.1002/anie.201915412>.
- Harper KC, Moschetta EG, Bordawekar SV, Wittenberger SJ. 2019. A laser-driven flow chemistry platform for scaling photochemical reactions with visible light. *ACS Cent Sci.* 5(1): 1–7.
- Li LH, Wei HZ, Wei Y, Shi M. 2022. The Morita–Baylis–Hillman reaction for non-electron-deficient olefins enabled by photoredox catalysis. *Chem Sci.* 13(5): 1478–1483. <https://doi.org/10.1039/d1sc06784b>.
- Moschetta EG, Richter SM, Wittenberger SJ. 2017. Heuristics, protocol, and considerations for flow chemistry in photoredox catalysis. *ChemPhotoChem.* <https://doi.org/10.1002/cptc.201700128>.
- Prier CK, Rankic DA, MacMillan DWC. 2013. Visible light photoredox catalysis with transition metal complexes: applications in organic synthesis. *Chem Rev.* 113(7): 5322–5363.
- Rigotti T, Schwinger DP, Graßl R, Jandl C, Bach T. 2022. Enantioselective crossed intramolecular [2+2] photocycloaddition reactions mediated by a chiral chelating Lewis acid. *Chem Sci.* 13, 2378–2384 (2022). <https://doi.org/10.1039/d2sc00113f>.
- Sambiagio C, Noël T. 2019. Flow photochemistry: shine some light on those tubes! *Trends Chem.* 2(2): 92–106. <https://doi.org/10.1016/j.trechm.2019.09.003>.
- Sun SZ, Duan Y, Mega RS, Somerville RJ, Martin R. 2020. Site-selective 1,2-dicarbofunctionalization of vinyl boronates through dual catalysis. *Angew Chem Int Ed.* 59(8): 2135–2139. <https://doi.org/10.1002/anie.201916279>.
- Twilton J, Le C, Zhang P, et al. 2017. The merger of transition metal and photocatalysis. *Nat Rev Chem.* 1, 0052. <https://doi.org/10.1038/s41570-017-0052>.
- Zhang J, Rueping M. 2024. Metallaphotoredox catalysis for sp³ C–H functionalizations through single-electron transfer. *Nat Catal.* 7:963–976.
- Zhou WJ. 2020. Reductive dearomative arylcarboxylation of indoles with CO₂ via visible-light photoredox catalysis. *Nat Commun.* 11:3263. <https://doi.org/10.1038/s41467-020-17166-3>.
- Nature Reviews Chemistry. 2019. The brilliant history of photoredox catalysis. *Nat Rev Chem.* <https://doi.org/10.1038/d42473-019-00033-7>.

To place an order or receive technical assistance

In Europe, please call Customer Service:

France: 0825 045 645	Spain: 901 516 645 Option 1
Germany: 069 86798021	Switzerland: 0848 645 645
Italy: 848 845 645	United Kingdom: 0870 900 4645

For other countries across Europe, please call: +44 (0) 115 943 0840

Or visit: MerckMillipore.com/offices

For Technical Service visit: MerckMillipore.com/techservice

MerckMillipore.com

We have built a unique collection of life science brands with unrivalled experience in supporting your scientific advancements.

Millipore® Sigma-Aldrich® Supelco® Milli-Q® SAFC® BioReliance®

Merck KGaA
Frankfurter Strasse 250
64293 Darmstadt, Germany

