

How to free trapped radicals from carboxyl

March 29 2019



Figure: Free radical, the Monkey King is bound by the "big mountain" of the carboxyl group. The mainstream photocatalytic systems are complex and costly. The novel catalyst uses a new mechanism to combine the cheap catalyst with the carboxyl group, pushes the redox reaction cycle, and easily sets free the "free radicals." Credit: CUI Jie

The removal of carboxyl groups and the release of alkyl radical fragments from the tight binding of carboxyl groups are promising directions in organic synthesis, especially in drug synthesis. Various catalysts have been designed to solve this challenge.

In a recent study published in *Science*, researchers from University of Science and Technology of China (USTC) of the Chinese Academy of Sciences reported their newly developed [catalyst system](#), which is cheap and simple.

The free radical is versatile and controllable, representing the "Monkey King" of organic [synthesis](#), but it is pressed by the "big mountain" of the carboxyl group.

Conventional decarboxylation processes have limitations in industrialization. In recent years, the [scientific community](#) has tried to use [photocatalytic](#) reaction to achieve decarboxylation conversion, which has advantages such as simple operation, easy control, and energy saving. The photocatalytic system has been successfully applied to the synthesis of various complex functional molecules.

However, most photoredox catalysts in current use are composed of precious metal complexes such as iridium and ruthenium or are synthetically elaborate organic dyes with complicated structures. It is important to develop environmental friendly and multifunctional photocatalytic systems. The [new catalyst](#) has a new way to use the new mechanism to combine cheap [catalyst](#) with carboxyl group, push redox reaction cycle, and easily and freely rescue the "free radicals."

Based on visible light excitation for intermolecular charge transfer, scientists from USTC proposed a new concept to construct a catalytic redox cycle for [organic synthesis](#). They discovered a simple, easily available, highly efficient and non-metallic anionic composite photocatalytic system for decarboxylative reaction of carboxylic acid derivative.

The proposed catalytic system simultaneously drives a redox cycle, simplifies the photocatalytic system, and reduces the cost of the

photocatalyst. The system breaks the limitations of traditional heating method, and solves the problems of transition metals remaining in the synthesis of functional compounds and drugs.

Using this system, redox-active esters derived from various natural and unnatural amino acids successfully trigger decarboxylative coupling reactions with high efficiency and gram scale production, indicating the feasibility of industrialization.

It is also expected to promote the scale industrialization of photocatalytic technology in the production of important functional molecules, with important synthetic chemical value and good industrial application prospects.

The results may initiate a new area of research in photoredox catalysis by introducing a tricomponent system based on a salt, a phosphine and an electron- acceptor to access redox active complexes without the need for traditional transition metal or complex dye catalysts.

This study illustrated that decarboxylative alkylation is accomplished without precious transition metals or organic dyes, which may be good news for many synthetic chemists.

More information: Ming-Chen Fu et al. Photocatalytic decarboxylative alkylations mediated by triphenylphosphine and sodium iodide, *Science* (2019). [DOI: 10.1126/science.aav3200](https://doi.org/10.1126/science.aav3200)

Provided by Chinese Academy of Sciences

Citation: How to free trapped radicals from carboxyl (2019, March 29) retrieved 27 April 2024 from <https://phys.org/news/2019-03-free-radicals-carboxyl.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.