

A new cross-coupling simplifies the synthesis of drug-like molecules

12 March 2018



Phil Baran, PhD, senior author of the study and Darlene Shiley Professor of Chemistry at TSRI. Credit: John D. & Catherine T. MacArthur Foundation.

Researchers at The Scripps Research Institute (TSRI) have designed a new molecule-building method that uses sulfones as partners for cross-coupling reactions, or the joining of two distinct chemical entities in a programmed fashion aided by a catalyst. The technique, described recently in the journal *Science*, paves the way toward other new chemical reactions and facilitates the synthesis of pharmaceutically-relevant molecules.

"It's already clear that this method opens the door to creating new types of compounds and new types of bonds," says Phil S. Baran, PhD, senior author of the study and Darlene Shiley Professor of Chemistry at TSRI.

This work was inspired by previous cross-coupling chemistry developed in the Baran lab, and catalyzed by discussions with pharmaceutical industry partners who view this as an area of major unmet need. Baran and his colleagues have previously studied decarboxylative cross-coupling reactions, where commonly-found carboxylic acids are transformed into many different molecules

using inexpensive metal catalysts and techniques commonly used for amide-bond synthesis. Throughout these studies, Baran and coworkers showed that decarboxylative cross-coupling can have broad applicability and facilitate the synthesis of pharmaceuticals and natural products. These reactions hinge on the transfer of one electron from the metal catalyst to an activated carboxylic acid, which allows for the cross-coupling to occur.

In this work, the authors demonstrate the new desulfonylative cross-coupling [reaction](#) by synthesizing over 60 representative molecules, including alkyl-fluorinated compounds inaccessible with earlier generation methodologies developed in the Baran group. Access to such compounds are critical for drug discovery campaigns, since fluorine atoms enhance drug-like molecular properties. Representative molecules described in the *Science* paper include some reported by Merck and Novartis in published patents.

Baran's group has already made their sulfone reagents and methods used in this study available to other chemists via Twitter who would like to use the technique. The new method is already having an impact on drug discovery programs at pharmaceutical companies with whom TSRI collaborates.

"I think the job of chemists in academia is to make things simpler. If we can have a small positive influence in making medicinal chemists lives easier, that will be a success for us," says Baran. "This chemistry is another step in that direction."

More information: Rohan R. Merchant et al, Modular radical cross-coupling with sulfones enables access to sp³-rich (fluoro)alkylated scaffolds, *Science* (2018). [DOI: 10.1126/science.aar7335](https://doi.org/10.1126/science.aar7335)

Provided by The Scripps Research Institute

APA citation: A new cross-coupling simplifies the synthesis of drug-like molecules (2018, March 12)
retrieved 25 May 2019 from <https://phys.org/news/2018-03-cross-coupling-synthesis-drug-like-molecules.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.