

Researchers report chemical reaction with potential to speed drug development

July 7 2017 a Feedstock Multifunctional allylic C-H Valuable chiral products oxidation product (such as pharmaceuticals, natural chemical products and fine chemicals) NR₂ Enantioselective Terminal olefin (1) OH R **b** This work SH R Enantioselective E/Z-selective Internal olefin (2) Regioselective (most common R class of alkenes) R'

Catalytic enantioselective oxidation of unactivated terminal and internal alkenes. Credit: *Nature* (2017). DOI: 10.1038/nature22805

Chemists have long sought to develop new reactions for the direct conversion of simple hydrocarbon building blocks into valuable



materials such as pharmaceuticals in a way that dependably creates the same chemical bonds and orientations. UT Southwestern researchers have hit upon a novel way to do that.

The ability to consistently create desirable <u>molecules</u> and avoid undesirable ones could greatly accelerate pharmaceutical production. Many molecules come in two mirror-image configurations, often labeled left-handed and right-handed. Usually, only one of the two is biologically desired, but the issue is more complicated than that, said Dr. Uttam Tambar, Associate Professor of Biochemistry and senior author of a study outlining the reaction published online in *Nature*.

"Unsaturated hydrocarbons contain only two types of bonds, carbonhydrogen and carbon-carbon. It is very challenging to selectively transform the starting material <u>hydrocarbon</u>, called the feedstock, into a product in which you have selectively transformed one carbon-hydrogen bond into a new bond in every molecule. We are able to do that," Dr. Tambar said.

"We are not only dealing with left- or right-handed configuration; we are also dealing with what position of the molecule is chemically modified," he added.

The selectivity problem challenges chemists, who often encounter it when attempting to directly transform unsaturated hydrocarbons that contain one or more carbon-carbon double bonds, resulting in difficultto-separate mixtures of compounds that require subsequent chemical steps to subdivide the mixtures into the desired molecules.

"We have developed a method for the direct conversion of a variety of double <u>bond</u>-containing materials into useful products. A specially designed chiral catalyst is used to oxidize hydrocarbons into multifunctional compounds that are obtained in high purity and are



readily converted into an assortment of molecules that will streamline the synthesis of future pharmaceutical drugs," said Dr. Leila Bayeh, a former graduate student and lead author of the study. Dr. Bayeh is now a postdoctoral researcher at the Massachusetts Institute of Technology.

The phrase "chiral catalyst" means that the catalyst that orchestrates the entire reaction has the proper orientation - a left- or right-handed configuration as the situation requires - in order to create the desired <u>chemical</u> transformation, he explained.

The orientation and proper location of all <u>chemical bonds</u> is crucial because the wrong molecule generated from a <u>reaction</u> can be ineffectual or potentially dangerous.

Hence the emphasis on consistency.

More information: Liela Bayeh et al. Catalytic allylic oxidation of internal alkenes to a multifunctional chiral building block, *Nature* (2017). DOI: 10.1038/nature22805

Provided by UT Southwestern Medical Center

Citation: Researchers report chemical reaction with potential to speed drug development (2017, July 7) retrieved 28 April 2024 from <u>https://phys.org/news/2017-07-chemical-reaction-potential-drug.html</u>

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.