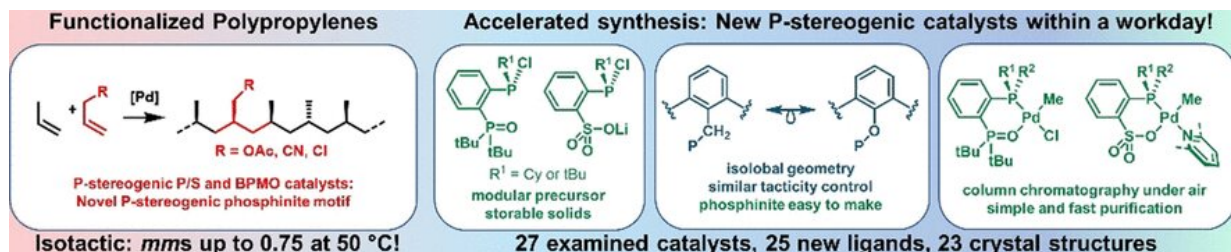


Catalysts for isotactic polar polypropylenes

October 30 2020



Credit: *Angewandte Chemie International Edition*

Polypropylene (PP) is one of the most widely used plastics in the world. By controlling the spatial orientation of the propylene building blocks and additional polar components, it should be possible to create a new generation of attractive, engineered, specialty plastics, with improved wettability or enhanced degradability, based on PP. In the journal *Angewandte Chemie*, Japanese scientists have introduced the basis for a new class of palladium catalysts for such polymerizations.

The properties of PP depend largely on the spatial orientation of the individual monomers as they are added to the growing chain (tacticity). In atactic PP (aPP) the orientation is random. In syndiotactic PP (sPP) the CH₃–side groups on the [monomer](#) alternately point toward the two sides of the polymer backbone. The most sought-after version—isotactic PP (iPP) in which all of the side groups point the same way—has particularly advantageous mechanical properties. Incorporation of additional functional, polar monomers into iPP is an important step

toward the development of novel plastics.

This type of copolymerization is heavily restricted with conventional Ziegler–Natta and metallocene catalysts because typical polar monomers first need to be 'masked.' This means they must be attached to special protective groups. With nickel and palladium catalysts, it is possible to achieve this unmasked but with significant losses in isotacticity. There has been some success with special nickel and palladium phosphine complexes (a type of phosphorus-containing organic compound), though synthesis of these catalysts is arduous and time-consuming.

Researchers working with Kyoko Nozaki at the University of Tokyo have now developed a new approach that allows more suitable catalysts to be produced much more easily. The spatial orientation of propylene monomers during polymerization is influenced by the special spatial structure (stereogenicity) at certain carbon atoms in the organic menthol substituents on the phosphine. The researchers wanted to develop phosphine compounds that have the required stereogenicity at the phosphorus atom.

To avoid the tedious synthetic challenges faced to date, they developed significantly faster synthetic protocols using storable, modular building blocks and phosphinites (a class of [organic compounds](#) containing phosphorus and oxygen). This allowed for the rapid and easy synthesis of many different phosphines and their corresponding palladium complexes. A rapid screening process successfully yielded suitable [catalyst](#) candidates.

In this way, the scientists found catalysts that polymerize propylene with polar monomers to form copolymers with particularly high isotacticity—a material they called isotactic polar polypropylene (iPPP).

More information: Falk William Seidel et al. Expedient Synthetic

Identification of a P-Stereogenic Ligand Motif for the
Palladium-Catalyzed Preparation of Isotactic Polar Polypropylenes,
Angewandte Chemie International Edition (2020). [DOI:
10.1002/anie.202009027](https://doi.org/10.1002/anie.202009027)

Provided by Wiley

Citation: Catalysts for isotactic polar polypropylenes (2020, October 30) retrieved 2 May 2024
from <https://phys.org/news/2020-10-catalysts-isotactic-polar-polypropylenes.html>

<p>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.</p>
--