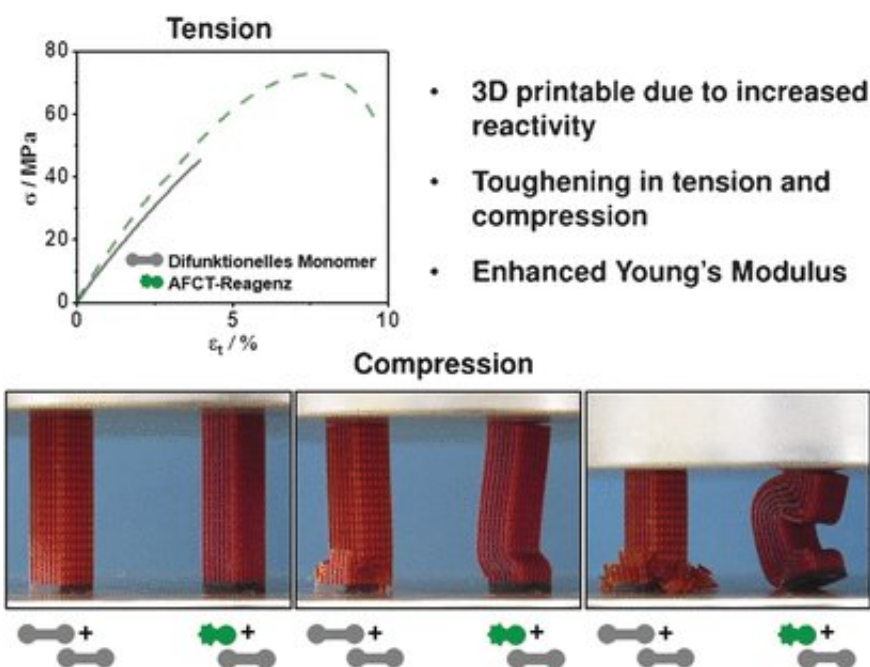


Efficient chain transfer for 3D-printing of tough photopolymers

July 6 2018



Credit: Wiley

An ever-growing number of coatings, including varnishes and printing inks, as well as tooth fillings, are cured with light. Yet, homogenous, tailored, polymer networks cannot be produced, and the materials tend to be brittle, which limits the use of photopolymers in applications like 3-D printing, biomedicine, and microelectronics. In the journal *Angewandte Chemie*, researchers present a method by which methacrylate-based, homogeneously crosslinked, tailored, tough polymers can be made – even

at high resolution for 3-D printing.

Light-curing is usually a radical chain polymerization. An initiator is split into radicals by light energy. These then attack the monomer, such as the C=C double bond in a vinyl group, which forms a new radical that becomes starting point of a growing [polymer](#) network by attacking more monomers and binding to them.

Newer methods to better control radical photopolymerization and the material properties of the products tend to slow the curing process, which is not ideal for 3-D printing. A short irradiation phase is critical for [high spatial resolution](#) and economical production times.

A new approach for the tailored production of methacrylate-based photopolymers without inhibiting the curing process has been developed by a team led by Robert Liska at the Technical University of Vienna (Austria). Their success relies on the addition of an ester-activated vinyl sulfonate ester (EVS), which acts as a chain transfer agent. It is activated because it easily splits off one portion of itself.

If the growing polymer network attacks EVS instead of the next monomer, an intermediate forms and quickly splits apart to form a terminated polymer chain in the [network](#) and a highly reactive radical (tosyl radical), which starts a new chain reaction. The more EVS is added, the shorter the average chain length in the [polymer network](#). Because shorter polymer chains remain mobile longer, the danger of shrinkage cracks during curing is significantly reduced. In contrast to conventional [chain](#) transfer agents, the polymerization is not inhibited, because there are no stable intermediates or reversible reaction steps involved. The splitting off of the tosyl radical is favored.

The researchers prepared a scaffold-like sample structure using a methacrylate copolymer. Individual layers with a thickness of 50 μm

were spatially well resolved. The material is very homogenous, solid but elastic and impact resistant with high tensile strength. These properties can be adjusted by changing the amount of EVS added. Without EVS, the material was very brittle. This new approach prepares the way for tough, photopolymers for applications in biomedicine, such as shape-memory polymers for tissue growth and as fillings for teeth.

More information: Konstanze Seidler et al. Vinyl Sulfonate Esters: Efficient Chain Transfer Agents for the 3D Printing of Tough Photopolymers without Retardation, *Angewandte Chemie International Edition* (2018). [DOI: 10.1002/anie.201803747](https://doi.org/10.1002/anie.201803747)

Provided by Wiley

Citation: Efficient chain transfer for 3D-printing of tough photopolymers (2018, July 6)
retrieved 27 April 2024 from
<https://phys.org/news/2018-07-efficient-chain-3d-printing-tough-photopolymers.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.
