

# Easy-To-Make Multi-Function Supramolecular Polymers Enabled by A Naturally Tailored Small Molecule

Qi Zhang<sup>1,2</sup>, Yuanxin Deng<sup>1,2</sup>, Chen-Yu Shi<sup>1</sup>, Da-Hui Qu<sup>1\*</sup>, Ben L. Feringa<sup>1,2\*</sup>, He Tian<sup>1\*</sup>

<sup>1</sup>Feringa Nobel Prize Scientist Joint Research Center, School of Chemistry and Molecular Engineering, East China University of Science and Technology, 130 Meilong Road, Shanghai, 200237, China

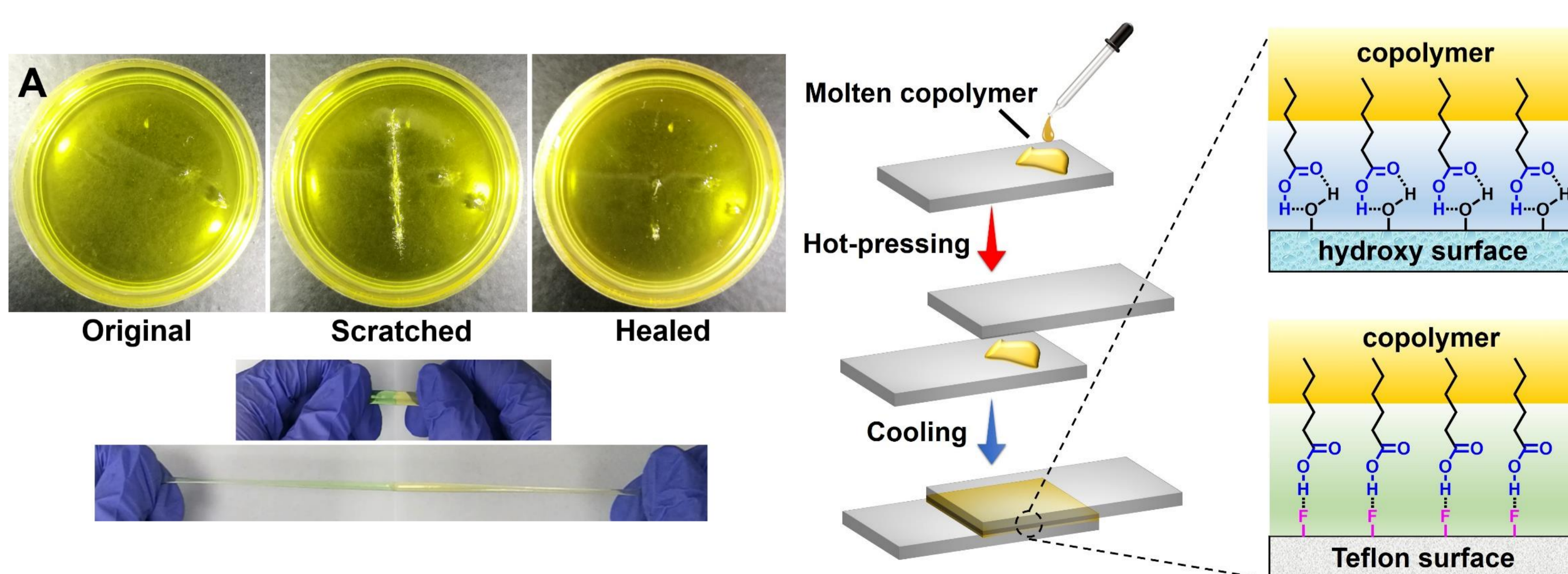
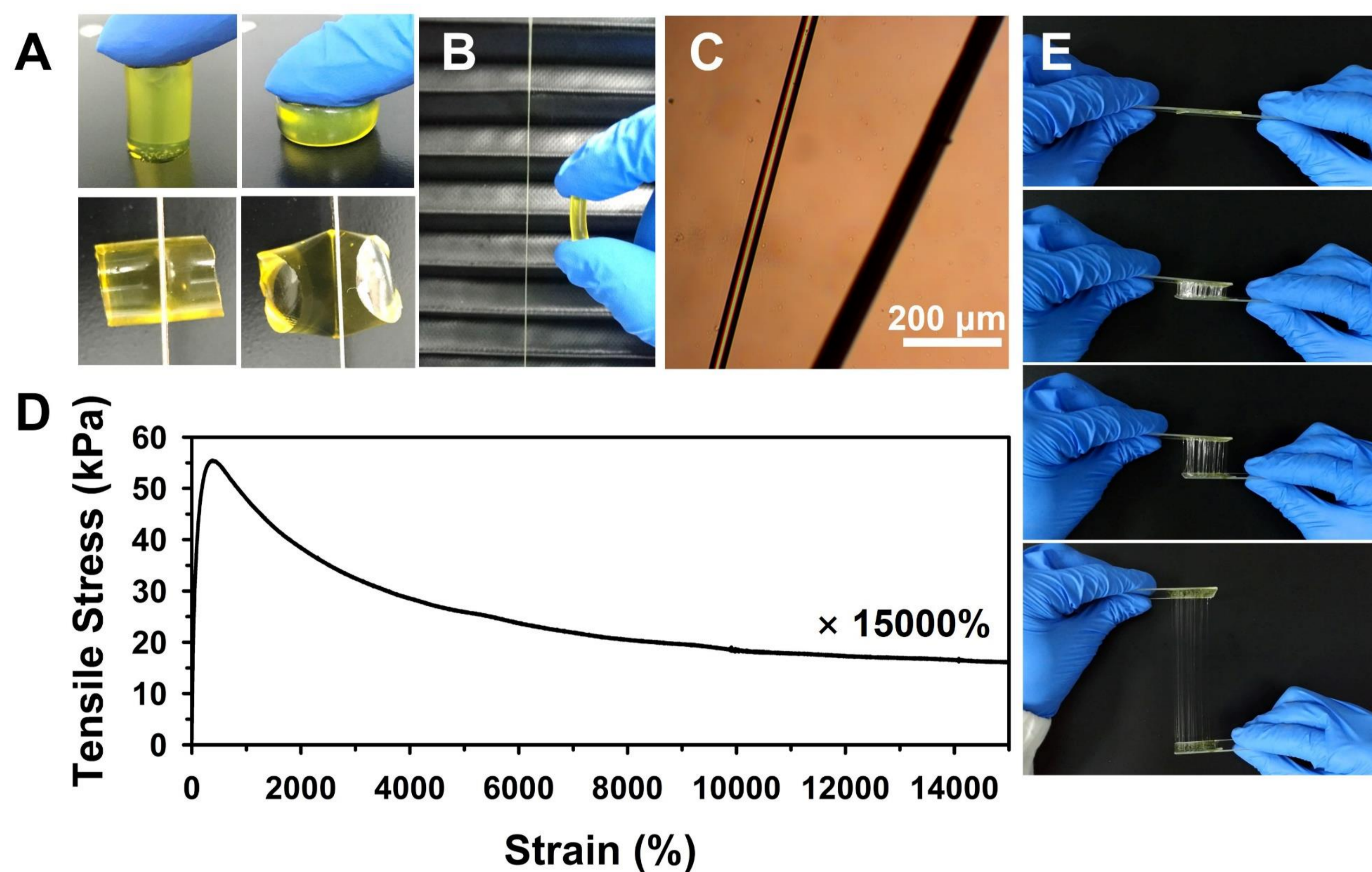
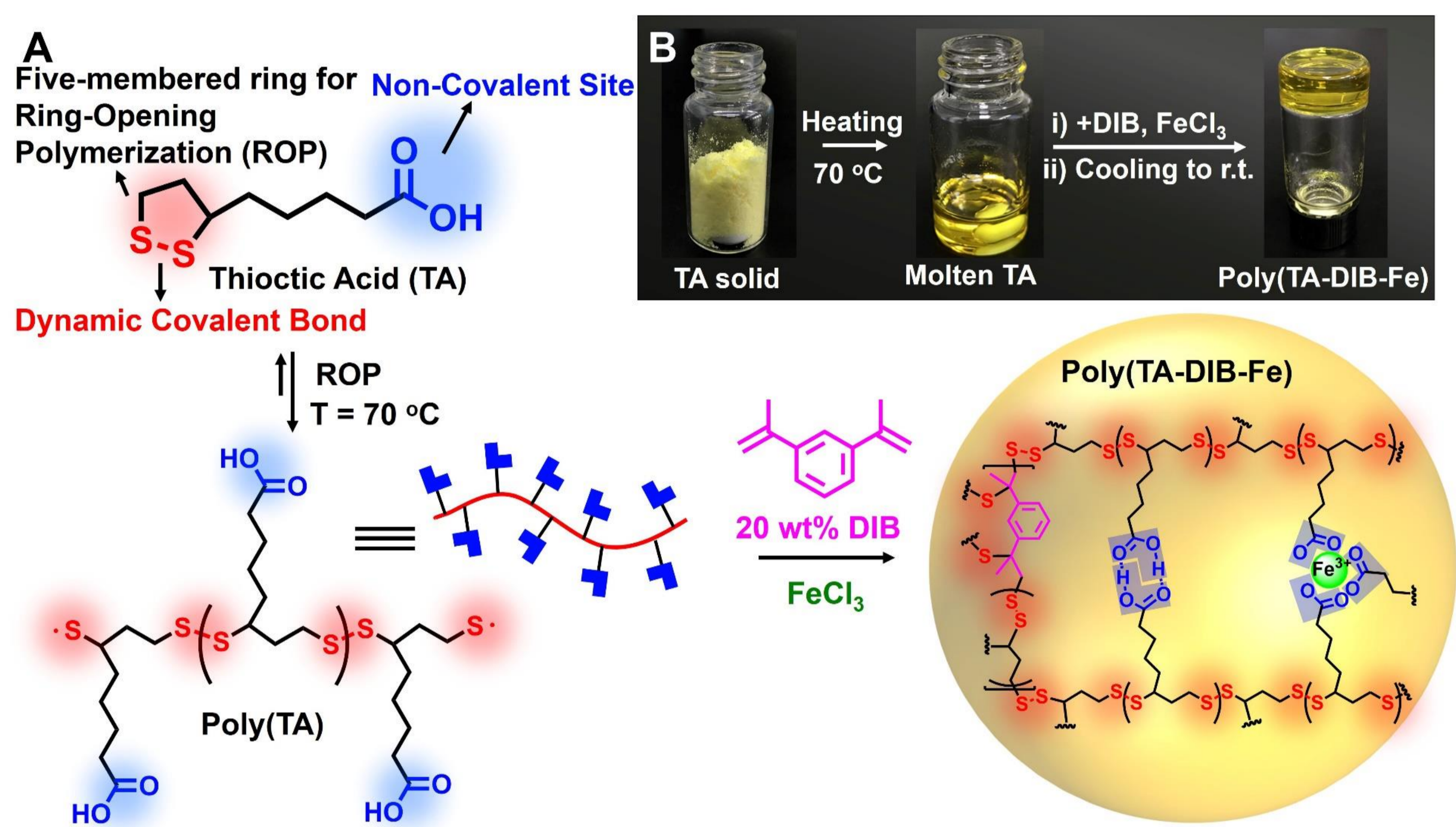
<sup>2</sup>Centre for Systems Chemistry, Stratingh Institute for Chemistry and Zernike Institute for Advanced Materials, University of Groningen, Nijenborgh 4, 9747 AG Groningen, The Netherlands

Email: dahui\_qu@ecust.edu.cn; b.l.feringa@rug.nl; tianhe@ecust.edu.cn

university of  
groningen

Polymeric materials with integrated functionalities are required to match their ever-expanding practical applications, but there is always a trade-off between complex material performances and synthetic simplification. A simple and effective synthesis route is demonstrated to transform a small molecule of biological origin, thioctic acid, into a high-performance supramolecular polymeric material, which combines processability, ultrahigh stretchability, rapid self-healing ability, and reusable adhesivity to surfaces. Meanwhile, we also disclose a strategy that directs the hierarchical self-assembly of this small molecule into a highly ordered supramolecular layered network. By combining the unique dynamic covalent ring-opening-polymerization and an evaporation-induced interfacial confinement effect, we precisely direct the dynamic supramolecular self-assembly of this simple small molecule in a scheduled hierarchical pathway, resulting in a layered structure with long-range order at both macroscopic and molecular scales.

## Supramolecular Elastomers



## Supramolecular "Glasses"

