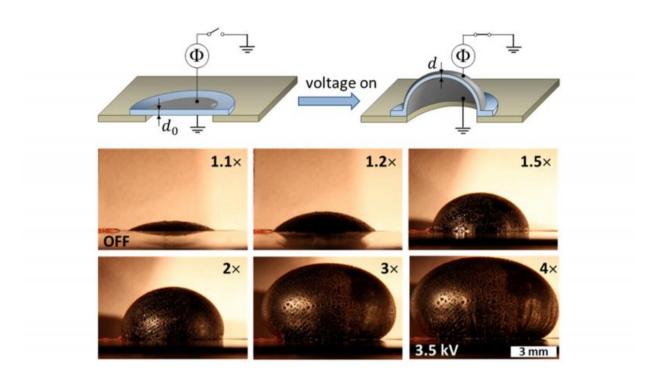


## New 'bottlebrush' electroactive polymers make dielectric elastomers increasingly viable for use in devices

November 9 2016, by Richard Spontak



Electrical response of a circular diaphragm composed of a pure bottlebrush elastomer upon electroactuation with increasing voltage and without any external pre-strain. The numbers indicate the electric field-induced area expansion under constant-volume conditions at room temperature. Credit: North Carolina State University

## A multi-institutional research team has developed a new electroactive



polymer material that can change shape and size when exposed to a relatively small electric field. The advance overcomes two longstanding challenges regarding the use of electroactive polymers to develop new devices, opening the door to a suite of applications ranging from microrobotics to designer haptic, optic, microfluidic and wearable technologies. The work was performed by researchers at North Carolina State University, the University of North Carolina at Chapel Hill, Carnegie Mellon University and the University of Akron.

"Dielectric elastomers are the most responsive electroactive polymers in terms of achievable strains, but two big hurdles have effectively prevented the smart materials community from using them in commercial devices," says Richard J. Spontak, Distinguished Professor of Chemical and Biomolecular Engineering and professor of materials science and engineering at NC State, who co-authored a paper describing the new materials. "First, previous dielectric elastomers required large electric fields in order to trigger actuation, or movement – on the order of at least 100 kilovolts per millimeter (kV/mm). With our new material, we can see actuation at levels as low as ca. 10 kV/mm."

"The second challenge is that, previously, materials had to be prestrained," Spontak says. "This would either mean using a frame to physically strain the material, or adding a second component to the polymer to retain the strain after it was applied. But our material consists of a single component that is specifically designed at the molecular level to inherently possess pre-strain. In other words, we don't need a frame or a second component – our material is ready to be used as soon as it is cross-linked into a specific shape."

The new material that has permitted this breakthrough is a "bottlebrush" silicone elastomer, which has been engineered to possess these unique properties, and it is not difficult to manufacture.



"We are working specifically with bottlebrush polymers, which are prepared by grafting long polymeric side chains to a polymer backbone," says Sergei S. Sheiko, George A. Bush, Jr. Distinguished Professor of Chemistry at UNC and corresponding author of the paper. "The resulting molecules may be viewed as filaments that are thick, yet remain quite flexible, which allows for significant reduction of the materials' rigidity and makes them more stretchable. Furthermore, the mechanical properties can be controlled by varying the bottlebrush architecture – for example, by preparing molecules with different degrees of polymerization of grafted chains and different grafting densities.

"This architectural control of mechanical properties has reduced the limit of stiffness in dry polymer materials by 1,000 times, demonstrated extensibility of up to eight times, and opened up new applications not available to stiffer materials or <u>materials</u> with liquid fractions," Sheiko says. "One of these applications – their use as free-standing <u>dielectric</u> <u>elastomers</u> – has been demonstrated, which we discuss in this paper."

"We're at the earliest stages of identifying all the potential ways in which we could use this new class of material," Spontak says. "It works better than anticipated, and now we're beginning to consider potential applications."

**More information:** Mohammad Vatankhah-Varnoosfaderani et al. Bottlebrush Elastomers: A New Platform for Freestanding Electroactuation, *Advanced Materials* (2016). DOI: <u>10.1002/adma.201604209</u>

Provided by North Carolina State University

Citation: New 'bottlebrush' electroactive polymers make dielectric elastomers increasingly viable



for use in devices (2016, November 9) retrieved 25 April 2024 from https://phys.org/news/2016-11-bottlebrush-electroactive-polymers-dielectric-elastomers.html

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