# **Conductive Hydrogels with Superior Mechanical Properties**

Van Tron Tran<sup>1</sup>, Md. Tariful Islam Mredha<sup>1</sup>, Suraj Kumar Pathak<sup>1</sup>, Hyungsuk Yoon<sup>1,2</sup> and Insu Jeon<sup>1\*</sup>

<sup>1</sup>School of Mechanical Engineering, Chonnam National University, 77 Yongbong-ro, Buk-gu, Gwangju 61186, Republic of Korea

<sup>2</sup>Korea Research Institute of Standards and Science (KRISS), 267 Gajeong-ro, Yuseong-gu, Daejeon 34113, Republic of Korea

\*Corresponding author: i\_jeon@chonnam.ac.kr

#### 1. Introduction

Hydrogels can absorb and retain a significant amount of water, which makes them have a number of promising properties such biocompatibility and environmental friendliness [1,2], responsiveness to external stimuli [3,4], adhesion [5], and biodegradability [6]. Especially, the hydrogels that exhibit both mechanical and electrical properties have received particular attention recently. Mostly this type of hydrogel has been obtained by employing the interactions between monofunctional carboxylic groups and multivalent metal ions such as Fe3+. However, there are some limitations in their mechanical properties such as low tensile strength [7] and modulus [8], as well as low compression properties [9].

To overcome these limitations, we have developed novel hydrogels which have combination of superior mechanical and good electrical properties by exploiting the ionic interactions between bifunctional carboxylic groups and multivalent Fe3+ ions. Due to their strong ionic bondina and simultaneously breaking reforming ability upon deformation, the equilibrium hydrogels also exhibit rapid self-recovery properties at room temperature.

## 2. Hydrogel synthesis

To prepare the hydrogels, firstly the prescribed amount of *N*-acryloyl glutamic acid (AGA), acrylamide (AAm) monomers and thermo-initiator, ammonium persulfate (APS), were dissolved in deionized water by sonication to form

homogeneous solution. Then the solution was deaired for 5 min with nitrogen gas and transferred into a reaction chamber for polymerization at 60 °C for 12 h. After that the as—prepared hydrogels were cooled down room temperature, taken out of the reaction chamber, and subsequently immersed in FeCl<sub>3</sub> solution for 24 hours for Fe<sup>3+</sup> loading. Finally, the gels were kept under water for 24 h to remove excess Fe<sup>3+</sup> ions. The shape of hydrogels is shown in Fig. 1.

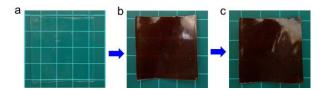


Fig.1 (a) As-prepared hydrogel; (b) Hydrogel after soaking in 0.1M Fe<sup>3+</sup> for 24 h; (c) Water-equilibrium hydrogel.

## 3. Mechanical and electrical properties

For evaluating mechanical properties, the tensile stress and strain properties of these hydrogels including as-prepared hydrogel, hydrogel after soaking in 0.1 M Fe<sup>3+</sup> solution, and equilibrium hydrogel were determined by using Universal Test Machine (Model TO-100-1C). In testing process, the initial length was set at ~12 mm and the deformation speed was set at 500% per minute. Based on the obtained results, the two best composition hydrogels were selected for further investigation such as fracture energy, compressive properties, and self-recovery properties. The best composition equilibrium hydrogels can exhibit over 30 MPa of Young's modulus, 10 MPa of strength,

~10 kJ m<sup>-2</sup> of fracture energy, ~60 MPa of compressive strength at 98 % strain, and rapid self-recovery at room temperature.

For investigating the electrical properties, the hydrogels resistance was measured. Then their resistivity and conductivity were calculated. Moreover, the dielectric constant of the hydrogels was also determined. The results indicated that, the hydrogels exhibit not only superior mechanical properties, but also good electrical conductivity and dielectric constant.

#### 4. Conclusions

With a combination of superior mechanical and good electrical properties, our developed hydrogels have great potential for applications such as load-bearing materials, soft robotics, flexible electrical devices and sensors.

### **Acknowledgment**

This work was supported by the National Research Foundation of Korea (NRF) grant funded by the Korea government (MSIP) (No. 2016R1A2B3011947).

### References

- [1] Q. Wang, J. L. Mynar, M. Yoshida, E. Lee, M. Lee, K. Okuro, K. Kinbara, and T. Aida, High-water-content mouldable hydrogels by mixing clay and a dendritic molecular binder, *Nature*, 463 (2010) 339–343.
- [2] M. C. Cushing and K. S. Anseth, Hydrogel cell cultures, *Science*, 316 (2007) 1133–1134.
- [3] I. Tokarev and S. Minko, Stimuli-responsive porous hydrogels at interfaces for molecular filtration, separation, controlled release, and gating in capsules and membranes, *Adv. Mater.*, 22 (2010) 3446–3462.
- [4] J. Hu, G. Zhang, and S. Liu, Enzyme-responsive polymeric assemblies, nanoparticles and hydrogels, *Chem. Soc. Rev.*, 41 (18) (2012) 5933–5949.
- [5] Z. Shafiq, J. Cui, L. Pastor-Perez, V. S. Miguel, R. A. Gropeanu, C. Serrano, and A. del Campo, Bioinspired underwater bonding and debonding on demand, *Angew. Chem. Int. Ed.*, 51 (2012) 4332–4335.
- [6] M. H. Park, M. K. Joo, B. G. Choi, and B. Jeong,

- Biodegradable thermogels, *Acc. Chem. Res.*, 45 (3) (2012) 424–433.
- [7] M. Zhong, X.-Y. Liu, F.-K. Shi, L.-Q. Zhang, X.-P. Wang, A. G. Cheetham, H. Cui, and X.-M. Xie, Self-healable, tough and highly stretchable ionic nanocomposite physical hydrogels, *Soft Matter*, 11 (21) (2015) 4235–4241.
- [8] P. Lin, S. Ma, X. Wang, and F. Zhou, Molecularly engineered dual-crosslinked hydrogel with ultrahigh mechanical strength, toughness, and good self-recovery, *Adv. Mater.*, 27 (2015) 2054–2059.
- [9] I. Hussain, S. M. Sayed, S. Liu, O. Oderinde, M. Kang, F. Yao, and G. Fu, Enhancing the mechanical properties and self-healing efficiency of hydroxyethyl cellulose-based conductive hydrogels via supramolecular interactions, *Eur. Polym. J.*, 105 (2018) 85–94.