

A flexible and fully recyclable transparent conductive organogel based on KI-containing glycerol with excellent anti-freezing and anti-drying behavior

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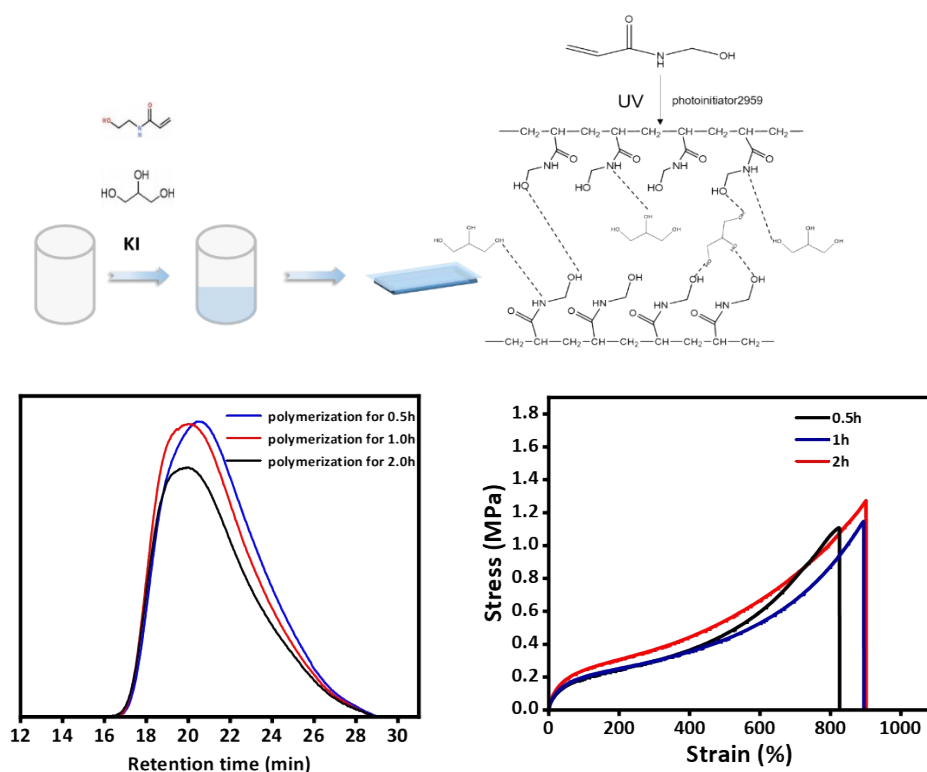


Figure S1 (a)Preparation of HEAA-KI-Gly conductive organogels. (b) GPC profile of PHEAA-KI-Gly organogels with different exposure durations under UV light. H₂O was used as the eluent. (c) The effect of different polymerization times contents on the mechanical properties of the HEAA-KI-Gly organogel.

Table 1 Proportions of different substances in the preparation of the HEAA-KI-Gly organogel.

HEAA	photo-initiator	KI	Glycerol
4.5g	0.045g	0g	0.55g
5.0g	0.05g	0.5g	0.5g
5.25g	0.0525g	1.0g	0.475g
5.5g	0.055g	1.5g	0.45g

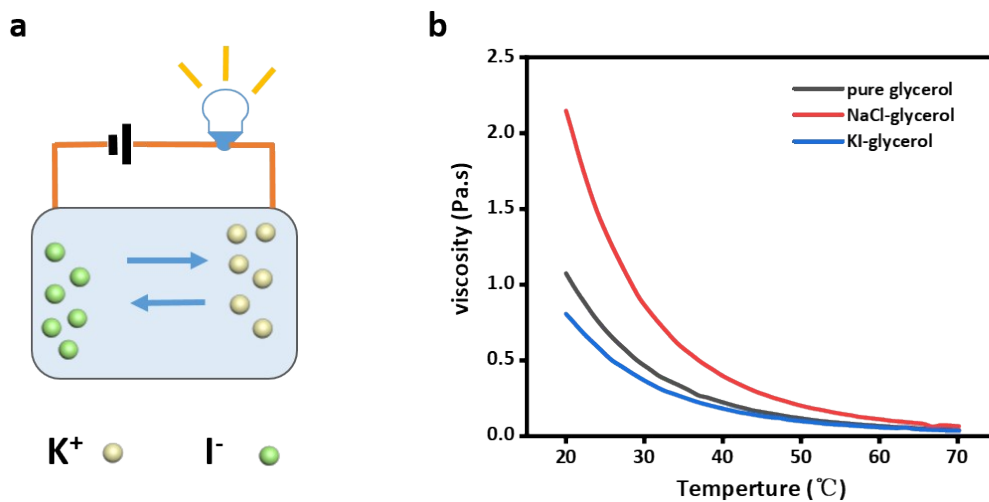


Figure S2 (a) Schematic of the conduction mechanism of the KI-Gly solution, (b) Viscosity variation of glycerol, KI-Gly solution and NaCl-Gly solution at temperatures ranging from 20.0 to 70.0 °C.

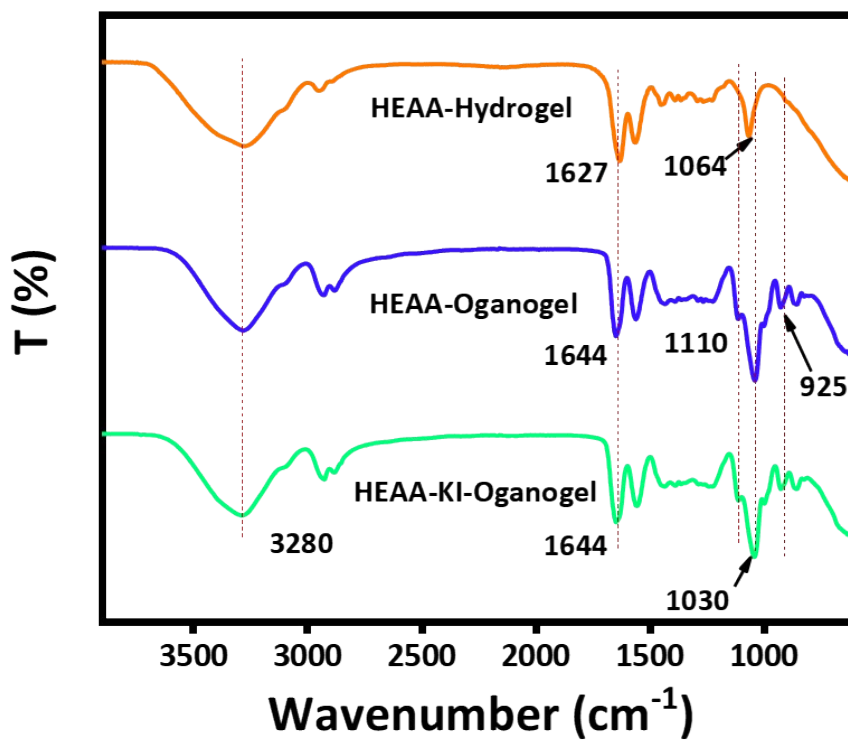
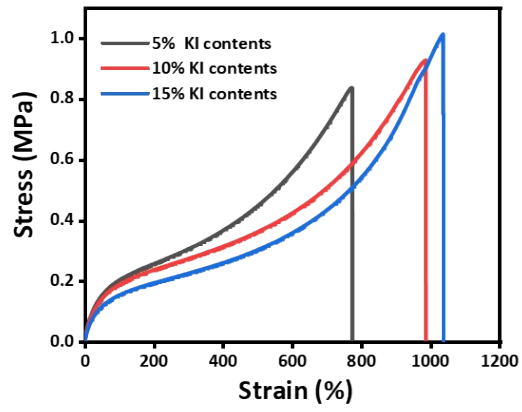
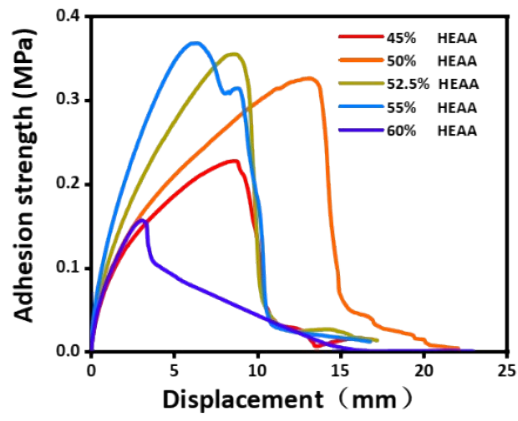


Figure S3 FT-IR spectra of a) HEAA hydrogel, b) HEAA-Gly organogel and c) HEAA-KI-Gly organogel.

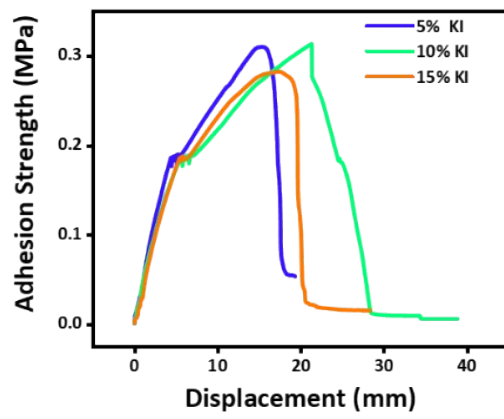
a



b



c



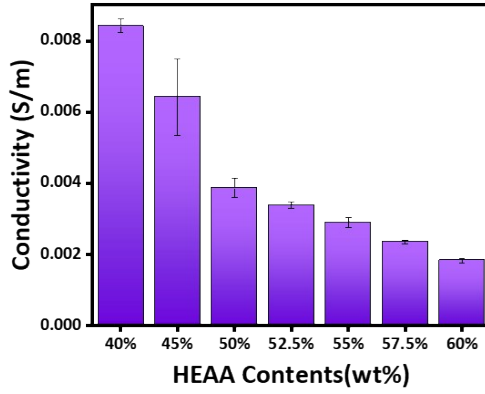


Figure S5 The effect of different HEAA monomer contents on the electrical conductivity of organogels.

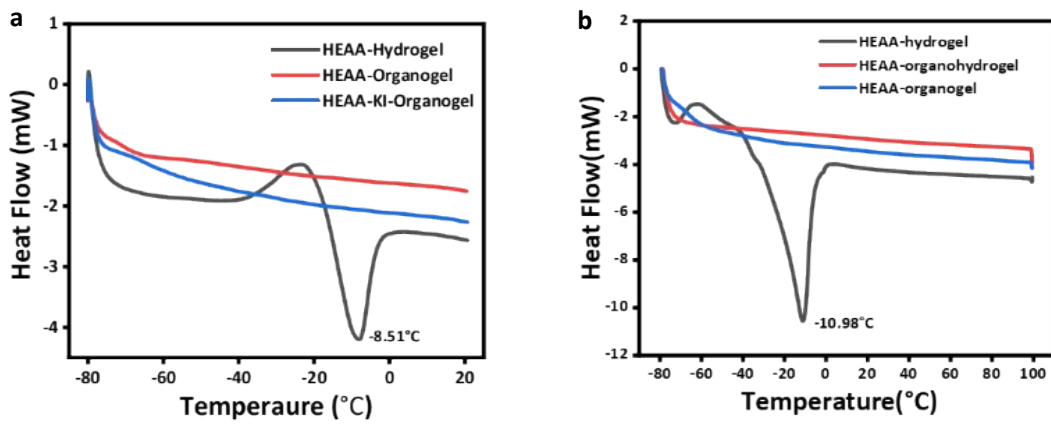


Figure S6 DSC thermograms of HEAA-KI-Gly organogels, HEAA-Gly organogels and HEAA hydrogels.

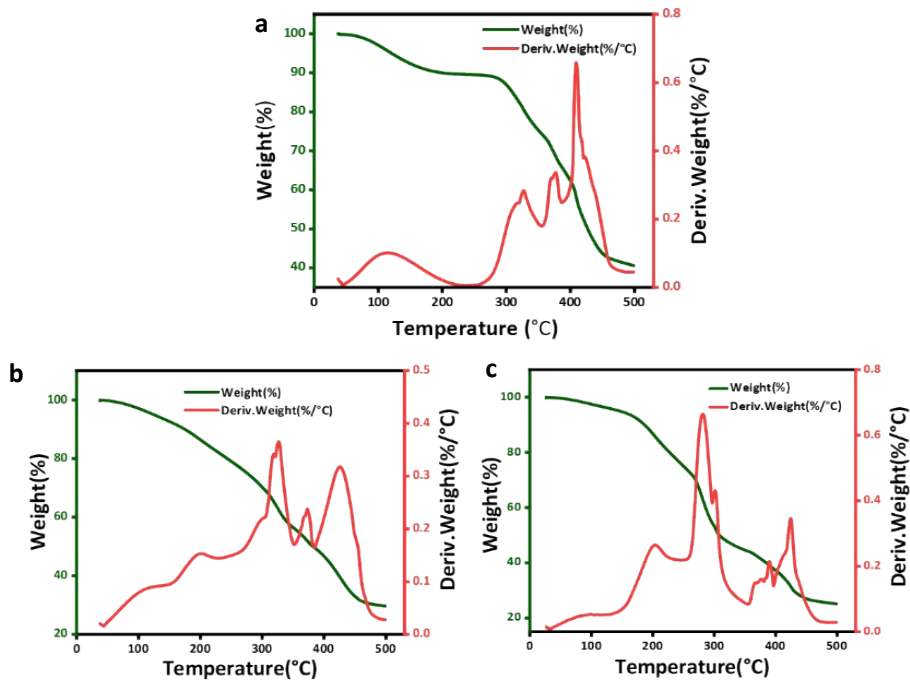


Figure S7 TGA thermograms of (a)HEAA hydrogel, (b)HEAA organohydrogel, and (c)HEAA-Gly organogel

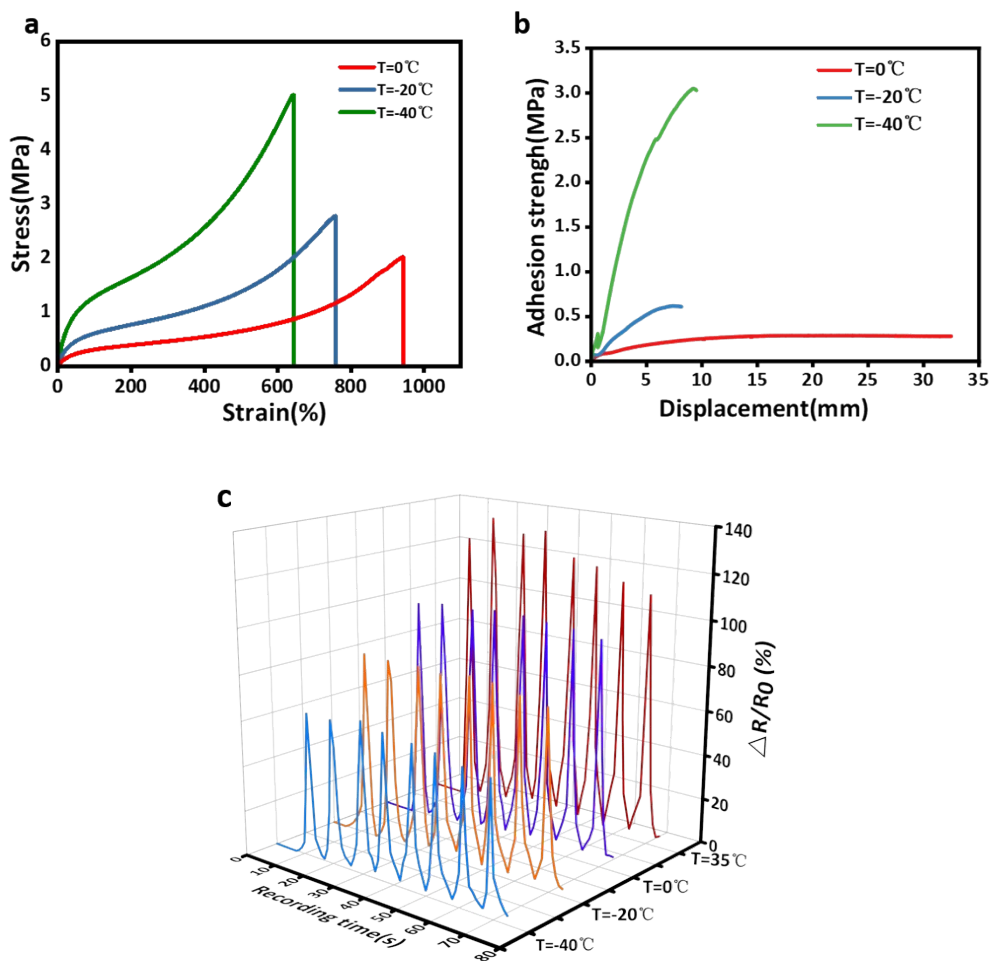


Figure S8 (a) The effect of different temperature contents on the mechanical properties of the HEAA-KI-Gly organogel. (b) Adhesion strength of organogels with different different temperature. (c) Relative resistance changes ($\Delta R/R_0$) at different temperatures under 100% tensile strain

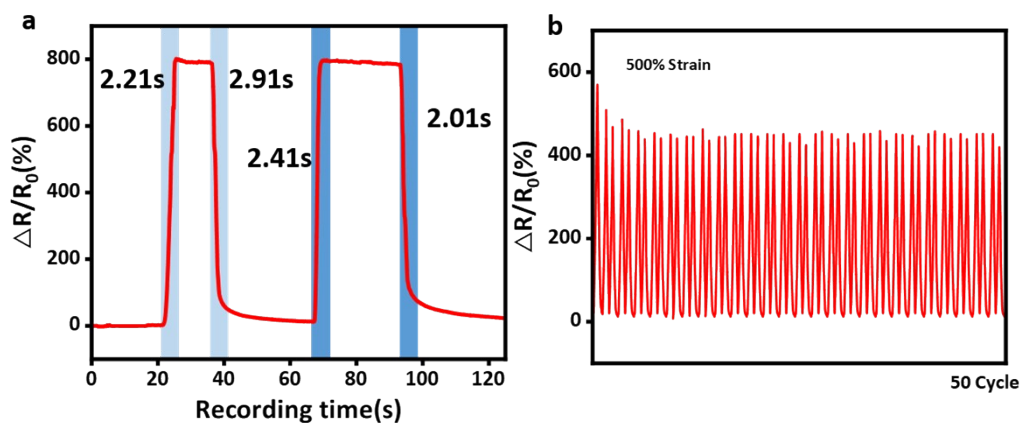


Figure S9 (a) Relative resistance changes ($\Delta R/R_0$) response time of organogels under 800% strain. (b) The durability test under repeated strains of 500% for 50 cycles.

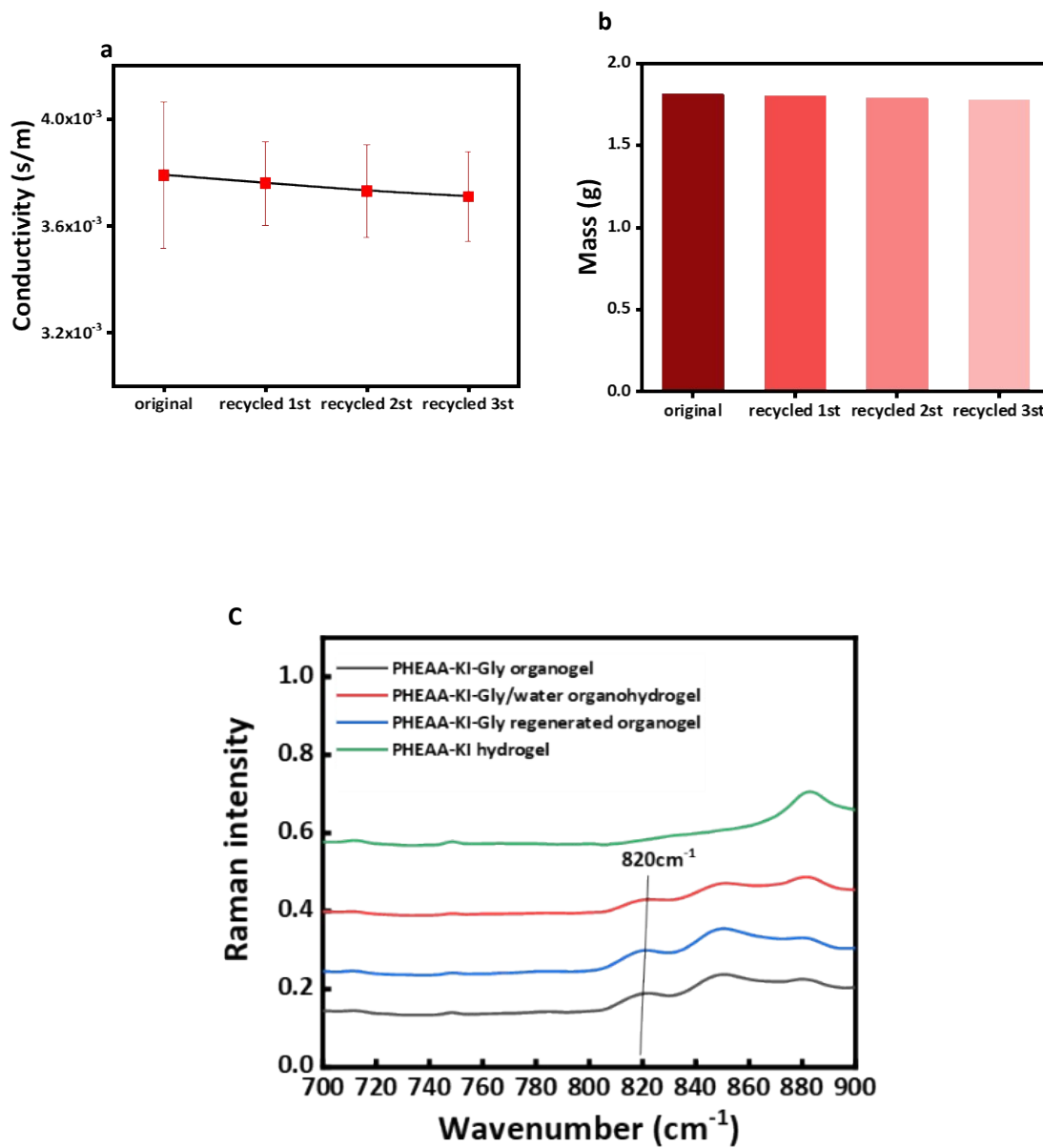


Figure S10 Comparison of quality (a) and conductivity (b) of organic gel films before and after recycling. (c) Raman Spectroscopy of PHEAA-KI-Gly organogel, PHEAA-KI-Gly/water organohydrogel, PHEAA-KI-Gly regenerated organogel and PHEAA-KI hydrogel.

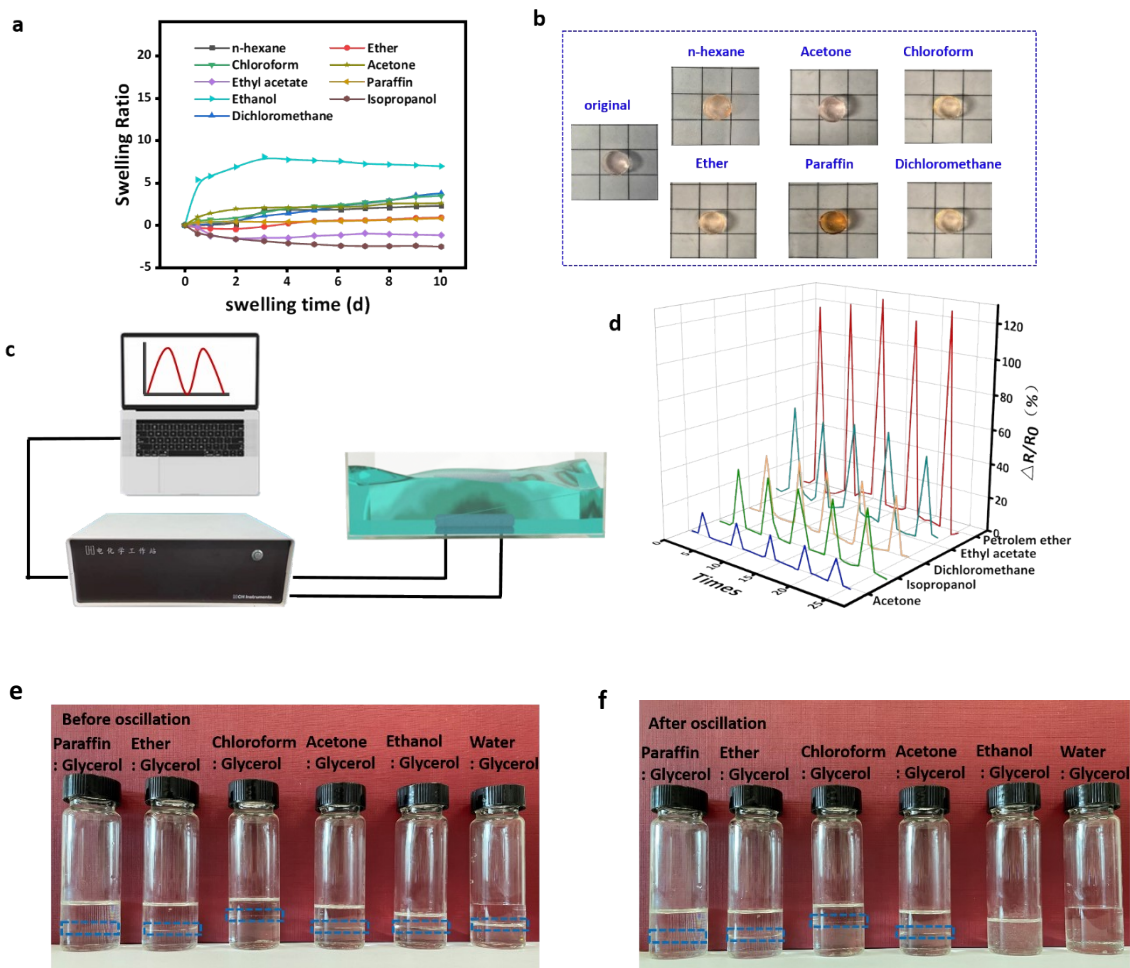


Figure S11 Swelling pictures(a) and swelling ratios(b) of organogel before and after swelling in different low polarity solvents. (c) Illustration of hydrogel sensors applied in solvents. (d) Real-time relative resistance variation of sensors with stretching (under 50% strain) in various solvents. Mutual solubility of solvents with different polarities and glycerol (e)before oscillation and (f) after oscillation

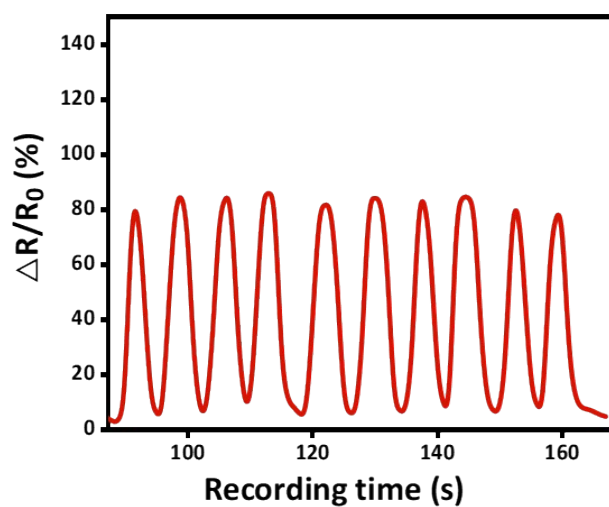


Table 2 Comparison of the performance of glycerol organogels and glycerol water organohydrogels

Materials(G/W organohydrogel)	Temperature range of application	Stretchability	Recyclability	Adhesiveness
PDA-CNTs-PACA ¹	-20°C to 60°C	600%	no	60KPa
PVA-HEC ²	-30°C to 65°C	400%	no	no
PVA-PANI ³	-20°C	472%	yes	no
PEDOT:PSS-SF ⁴	-40 °C to 60 °C	100%	no	no
P(AM-MAANa-DMC) ⁵	-20°C	1500%	no	no
PDMAPS-SA ⁶	-25°C to 25°C	50%	no	no
PAA-CNF ⁷	-46°C	980%	no	25KPa
P(SBMA-HEAA) ⁸	-30°C to 60°C	600%	no	21KPa
This work	-60°C to 70°C	1200%	yes	300KPa

Notes and references

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