

COMMENTARY

Designing toughness and strength for soft materials

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Soft materials, such as hydrogels, elastomers, and plastics, are pervasive in nature and society. For example, except for teeth, nails, and bones, all other components of the human body are hydrogels, and we eat, wear, and use soft materials as foods, clothes, shoes, and car tires, just to name a few, in our daily life. Although amorphous polymer chains endow soft materials with high flexibility or “softness,” they can also lead to inferior mechanical performances, such as low fracture toughness and low strength, frequently hampering applications and innovations of soft materials. Over recent years, intensive efforts have been devoted to the development of soft materials that possess extraordinary mechanical properties, especially by seeking inspiration from nature and biology. For instance, synthetic hydrogels can now be made much tougher than articular cartilages (1, 2) and more adhesive than mussel glues (3, 4). In PNAS, Wu et al. (5) report the fabrication of bioinspired fibers with both high toughness and high strength by drawing from a hydrogel at ambient temperatures and pressures. This nascent progress in the field raises a generic question. What are the fundamental principles for the design of soft materials to achieve certain mechanical properties in nature and engineered systems? Compared with hard materials such as steels and ceramics, which have been explored over centuries, such principles are still elusive in the field of soft materials.

The goal of this commentary is to elucidate the design principles for soft materials that are both tough and strong, such as natural silks (6) and the hydrogel fibers synthesized by Wu et al. (5). I will show that three design principles are required for soft materials to achieve both high toughness and high strength: (i) high mechanical dissipation and stretchability, (ii) below flaw-sensitive length scale, and (iii) simultaneous stiffening and fracture of significant polymer chains at failure. By drawing from a multiphase self-assembled hydrogel, Wu et al. (5) exploited all three principles to fabricate fibers that were both tough and strong. In control cases, when any principle was violated, the resultant fibers turned out to be brittle and/or weak (5). In the following sections, I will elaborate on these principles and propose possible designs

for future soft materials with extraordinary mechanical properties and beyond.

High Mechanical Dissipation and Stretchability

The intrinsic fracture toughness of polymer networks, defined as the energy required to fracture a single layer of polymer chains (7), is on the order of $1\sim 100\text{ Jm}^{-2}$, which is much lower than the fracture toughness of articular cartilage (over $1,000\text{ Jm}^{-2}$). The intrinsic fracture toughness alone cannot give a tough soft material. To design tough soft materials, one needs to induce significant mechanical dissipation in the process zone around the crack tip (Fig. 1A). The total fracture toughness of the soft material can be expressed as (8, 9)

$$\Gamma = \Gamma_0 + \Gamma_D,$$

where Γ_0 is the intrinsic fracture toughness and Γ_D is the contribution from dissipation in the process zone. The mechanical dissipation manifests as a hysteresis loop in the nominal stress-stretch curve of the material (Fig. 1B). Although the failure strengths of soft materials are typically much lower than the failure strengths of hard materials, one can design high stretchability of a soft material to achieve a large area of the hysteresis loop, as shown in Fig. 1B. The mechanisms for dissipating mechanical energy in soft materials include fracture of polymer chains, reversible cross-links, and domain transformations, and the mechanisms for achieving high stretchability include interpenetration of long-chain networks, hybrid physical and chemical cross-links, and high-functionality cross-links (8). A combination of mechanisms for high dissipation and stretchability can give various designs of tough soft materials (8). For example, in Wu et al.'s work (5), reversible cross-links and high-functionality cross-links have been used to give dissipative, stretchy, and thus tough hydrogels (5).

Below Flaw-Insensitive Length Scale

The high fracture toughness of a material does not guarantee its high fracture strength. For one thing, the fracture strength can be significantly affected by flaws

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Author contributions: X.Z. designed research, performed research, contributed new reagents/analytic tools, analyzed data, and wrote the paper.

The author declares no conflict of interest.

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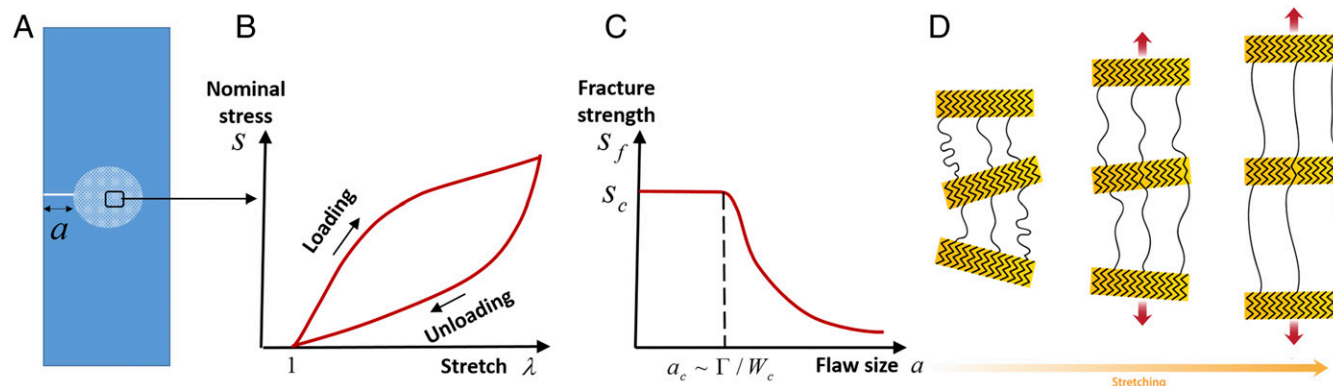


Fig. 1. Schematic illustration of the design principles for soft materials to be tough and strong. (A) Soft material contains a crack of size a and a process zone around the crack tip. (B) Toughening. High mechanical dissipation in the process zone, manifested as a hysteresis loop in the curve of nominal stress s vs. stretch λ , toughens the material. (C) Flaw-insensitive strength. The fracture strength of a material increases with the decrease of flaw size until the flaw reaches a critical length scale a_c below which the fracture strength is flaw-insensitive s_c . (D) High strength. High-functionality cross-links that accommodate hidden polymer chains can promote simultaneous stiffening and fracture of significant polymer chains, which gives soft material high fracture strength.

presented in the material, such as cavities, cracks, and impurities. A critical length scale has been defined as (10, 11)

$$a_c = \beta \frac{\Gamma}{W_c},$$

where W_c is the work to rupture of a material with no flaw or only flaws that do not affect its fracture strength and β is a dimensionless parameter that accounts for geometrical and nonlinear effects. If the size of any flaw in the material is larger than a_c , the fracture strength decreases significantly with the flaw size, but if all flaws are smaller than a_c , the fracture strength reaches a plateau value s_c insensitive to flaws (Fig. 1C). This flaw-sensitivity principle applies to both hard and soft materials, and the critical length scale ranges from tens of nanometers for ceramics and glass (11) to millimeters for elastomers and gels (10). Although eliminating flaws from materials is difficult, one can design materials to make all possible flaws smaller than a_c to achieve s_c . In general, enhancing the fracture toughness of a material tends to increase its a_c ; for example, toughening of ceramics (12), elastomers (13), and hydrogels (1, 2) makes them more flaw-tolerant. Another simple approach to achieve flaw-insensitive strength is by making the characteristic length of a material smaller than a_c , such as drawing fibers with diameters equal to or less than a_c (5).

Simultaneous Stiffening and Fracture of Significant Polymer Chains

A flaw-insensitive fracture strength still does not mean a high fracture strength, particularly in soft materials. For instance, the flaw-insensitive fracture strengths of tough hydrogels are usually lower than 1 MPa (1, 2). On the other hand, the ideal fracture strength of soft materials can be extremely high. Let us consider an ideal polymer network with uniform chain lengths and no defect or entanglement (14). By assuming all chains reach extension limit and rupture simultaneously, we can evaluate the ideal fracture strength of the network (in terms of nominal stress) to be

$$s_{ideal} = N \cdot \sqrt{n} \cdot b \cdot f,$$

where N is the number of polymer chains per unit volume, n is the number of Kuhn monomers per chain, b is the length of a

Kuhn monomer, and f is the rupture force of a polymer chain. For a polymer network with a shear modulus of 100 kPa, where N is around $2.5 \times 10^{25} \text{ m}^{-3}$ and n , b , and f are taken as typical values of 100, 1 nm (15) and 4 nN (16), the ideal fracture strength can be estimated to be 1 GPa. (Note the ideal fracture strength in terms of true stress is $\sqrt{n} \cdot s_{ideal}$, which is even higher.) This theoretical limit indicates that it is possible to design extremely strong soft materials, but such design requires a significant amount of polymer chains to reach the extension limit and to rupture simultaneously. However, the polymer networks in real soft materials usually have nonuniform chain lengths and defects; thus, the polymer chains tend to stiffen and fracture in a sequential manner, leading to low fracture strength. One method to promote simultaneous stiffening and fracture of multiple polymer chains is to introduce high-functionality cross-links that accommodate hidden chains (17), such as crystalline domains and nanoparticles with segments of polymer chains embedded in or attached on them. Multiple polymer chains with different lengths bridge two adjacent high-functionality cross-links (Fig. 1D). Under deformation, the shorter chains can pull hidden segments from the cross-links, and eventually reach the extension limit and fracture simultaneously with longer chains (Fig. 1D). In Wu et al.'s hydrogel (5), both silica nanoparticles and crystalline domains were used as high-functionality cross-links. Another possible method for simultaneous stiffening and fracture of polymer chains is to use slidable cross-links that automatically adjust chain lengths during deformation to achieve uniform chain forces throughout the network (18).

Although this commentary has discussed the principles for designing soft materials that are both tough and strong, more and more questions are emerging in this rapidly evolving field. For example, it is equally important to elucidate the design principles for soft materials to achieve other mechanical properties, such as antifatigue, high adhesiveness (3, 4), and low surface friction (19). Beyond mechanical properties, intensive efforts have been devoted to making soft materials responsive to external stimuli, such as temperature, pH, light, and electromagnetic field. What are the design principles for responsive soft materials to achieve high actuation strain and stress, high energy efficiency, and fast response? Furthermore, as soft materials are enabling future electronics,

medicine, robots, and machines (20), how can we harness the designed properties of soft materials most effectively to help address societal challenges in relevant areas, such as health care, food, water, and energy?

Acknowledgments

I thank Hyunwoo Yuk for help in drawing the schematic figure. My research is supported by National Science Foundation Grant CMMI-1661627, Office of Naval Research Grant N00014-14-1-0619, and the MIT Institute for Soldier Nanotechnologies.

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