



Cite this: *Polym. Chem.*, 2020, **11**, 358

## Fractography of poly(*N*-isopropylacrylamide) hydrogel networks crosslinked with mechanofluorophores using confocal laser scanning microscopy†

Maria Stratigaki,<sup>a</sup> Christoph Baumann,<sup>a</sup> Lambert C. A. van Breemen,<sup>b</sup> Johan P. A. Heuts,<sup>c</sup> Rint P. Sijbesma<sup>c\*</sup> and Robert Göstl<sup>a\*</sup>

Due to their soft and brittle nature, the mechanical characterization of polymer hydrogels is a difficult task employing traditional testing equipment. Here, we endowed poly(*N*-isopropyl acrylamide) (PNIPAAm) hydrogel networks with Diels–Alder adducts of  $\pi$ -extended anthracenes as mechanofluorophore crosslinkers. After swelling the networks with varying amounts of water and subjecting them to force, we visualized the subsequent fluorescence caused by covalent bond scission with confocal laser scanning microscopy (CLSM) and related the intensities to the macroscopic fracture mechanics and the elastic moduli recorded with traditional uniaxial compression. The sensitivity of the mechanofluorophores allowed the analysis of low levels of mechanical stress produced *via* a hand-induced needle-puncturing process and, thus, is an alternative to conventional force application methods. The detection and precise localization of covalent bond scission *via* CLSM helps elucidating the interrelationship between molecular structure and the macroscopic properties of chemically crosslinked polymeric hydrogels. We believe that this micro-scale mechanophore-assisted fractography can establish a new paradigm for the mechanical analysis of soft matter in fields covering traditional polymer and life sciences.

Received 5th June 2019,  
Accepted 12th July 2019

DOI: 10.1039/c9py00819e

rsc.li/polymers

## 1 Introduction

Hydrogels are crosslinked polymeric networks that can absorb and retain large amounts of water. Their drastic volume changes in response to specific external stimuli, such as temperature and pH, render them promising biocompatible materials in applications for sensors, tissue engineering, or drug delivery.<sup>1</sup> However, their success as structural materials is largely impeded by their intrinsic fragility along with their extreme softness, both stemming from the low density of polymer chains in the water-swollen state. Hence, the study of their mechanical properties is of particular interest, yet very much complicated by the severe limitations regarding the utilization of conventional measurement methods and techniques.<sup>2,3</sup>

To overcome these challenges, it is important to develop more sophisticated experimental methodologies and characterization techniques. Conventional measurement methods can be adapted for the mechanical characterization of soft polymer materials to determine their elastic behavior and response, based on the principles of indentation<sup>4,5</sup> or cavitation rheology.<sup>6</sup> Concomitantly, the traditional fractography methods, such as atomic force or electron microscopy probe the fracture surface of materials, but are time-consuming and only work post-mortem. However, since the accumulation of non-reversible ruptured bonds is the dominant mechanism leading to failure in polymeric structures, it is important that imminent fracture is reported prior to the formation of any visible macroscopic cracks that will ultimately evolve into the decomposition of the polymer.<sup>7</sup> In recent years, the force-induced selective scission of covalent and non-covalent bonds was exploited to generate molecules that change their photo-physical properties upon the application of mechanical stress and strain, thus allowing the optical detection of material damage.<sup>8,9</sup> While the seminal development of such molecules changing their absorption spectra (mechanochromophores) can be regarded as elegant, their investigation on small scales was hampered intrinsically by the Lambert–Beer law.<sup>10</sup> Hence, more sensitive and scalable approaches were

<sup>a</sup>DWI – Leibniz Institute for Interactive Materials, Forckenbeckstr. 50, 52056 Aachen, Germany. E-mail: goestl@dwi.rwth-aachen.de

<sup>b</sup>Department of Mechanical Engineering, Polymer Technology, Eindhoven University of Technology, P.O. Box 513, 5600 MB Eindhoven, The Netherlands

<sup>c</sup>Laboratory of Supramolecular Polymer Chemistry, Institute for Complex Molecular Systems, Eindhoven University of Technology, P.O. Box 513, 5600 MB Eindhoven, The Netherlands. E-mail: r.p.sijbesma@tue.nl

† Electronic supplementary information (ESI) available. See DOI: 10.1039/C9PY00819E



required leading to the genesis of mechanoluminescence and mechanofluorochromism techniques.<sup>11–15</sup> Mechano-fluorophores were successfully employed to locally visualize stress concentrations in a wide range of materials, including hydrogels.<sup>16,17</sup> Yet, up to today and to the best of our knowledge, the analysis techniques employed for the study of mechanical failure in materials are limited to the macro-scale. However, a holistic description of fracture mechanisms requires to precisely resolve smaller feature-sizes on the micro-scale, as this information is valuable for tailor-making polymeric structures with distinct mechanical properties, such as biomimetic gels,<sup>18</sup> smart skin,<sup>19</sup> or muscle-like materials.<sup>20</sup>

Here we report the integration of the Diels–Alder adduct of a  $\pi$ -extended anthracene and maleimide, previously synthesized in our group,<sup>21</sup> as covalent crosslinker into PNIPAAm hydrogel networks, endowing these soft materials with stress-sensing capability (Scheme 1). This mechanofluorophore is excellently photostable, exhibits a high fluorescence quantum yield (72%), and can be observed with outstanding signal-to-noise ratio due to its “turn-on” nature. We rely on PNIPAAm as a typical and well-investigated thermoresponsive hydrogel with mechanical behavior that can be fine-tuned depending on its water-absorption capability.<sup>2,22,23</sup> By unprecedented analysis relying on CLSM, we explore the fracture behavior of the PNIPAAm hydrogels on the micro-scale after employing a convenient and easy-to-implement needle-puncturing method for mechanical disruption. We make a productive correlation of material properties and mechanophore fluorescence after fracture by the precise localization of stress accumulation around a defined region of interest.



**Scheme 1** Force-induced cycloelimination of Diels–Alder adduct from 9- $\pi$ -extended anthracene and maleimide in a crosslinked polymeric hydrogel network.

## 2 Experimental section

### 2.1 Materials

The PNIPAAm hydrogel networks were synthesized by free radical polymerization. Commercially available materials were supplied by Sigma-Aldrich and used as described in Table 1, while the synthesis of the mechanophore crosslinker was based on our own previous work. The overall crosslink density was 2 mol% with a mechanophore crosslinker content of only 0.02 mol%. The prepolymer solution (NIPAAm, tetraethylene glycol dimethacrylate (TEGDMA), azobisisobutyronitrile (AIBN), and mechanophore crosslinker dissolved in dioxane and degassed with Ar flow for 30 s) was heated for 5 min to completely dissolve solid residues, and was then poured into Teflon molds to anneal for 24 h at 65 °C under N<sub>2</sub> atmosphere. The formed networks were washed in ethanol for a minimum of 24 h, and were then dried *in vacuo* for another 24 h. The controlled swelling of the networks to different degrees ( $d_s$ ) was accomplished by weighing the dry samples ( $m_d$ ) with a precision balance and then subsequently adding the required amount of water ( $m_w$ ) with a gastight microliter syringe to reach the desired swollen mass ( $m_s$ ) which was determined after 24 h:

$$d_s = \frac{m_s - m_d}{m_d} \times 100 = \frac{m_w}{m_d} \times 100 \quad (1)$$

Equilibrium swelling was achieved by immersing the networks into excess of deionized H<sub>2</sub>O for 24 h. In every case, the exact  $d_s$  for each individual sample was verified prior to imaging, compression, or other measurements.

### 2.2 Methods

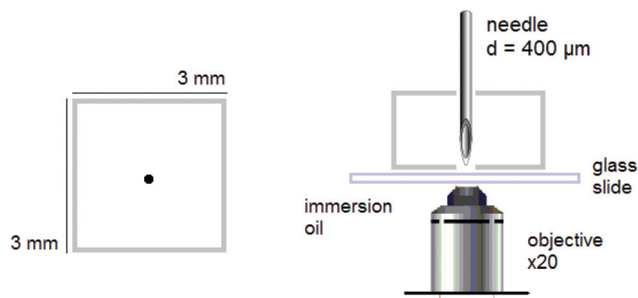
#### 2.2.1 Confocal laser scanning microscopy

**Puncture.** Dry rectangular specimens of *ca.* 3 × 3 × 3 mm<sup>3</sup> were individually and prior any further experiments swollen to different  $d_s$  varying from 0 to equilibrium. Then, they were punctured with a stainless steel needle. The diameter of the needle was purposefully chosen to a minimum, *i.e.* 400  $\mu$ m, to form a fluorescent circular trace, due to mechanofluorophore activation, that is smaller than the field of view of the used microscope, and therefore could be clearly visualized and analyzed. The beveled-tip of the needle accommodated an easier penetration into the (dry or slightly swollen) glassy materials.<sup>24</sup> The needle was inserted vertically into the center-region of the PNIPAAm networks, and penetrated the bulk of each sample until it exited the opposite surface (Fig. 1).

**Table 1** Materials for the synthesis of PNIPAAm hydrogel networks crosslinked with mechanophore dimethacrylate

| Material           | Role                     | Amount                    | Comments   |
|--------------------|--------------------------|---------------------------|--|
| NIPAAm             | Monomer                  | 1000 mg, 8.84 mmol        | Purified by recrystallization from hexane at 65 °C                                 |
| TEGDMA             | Crosslinker              | 53 $\mu$ L, 0.175 mmol    | Purified by passing through a short column of basic Al <sub>2</sub> O <sub>3</sub> |
| AIBN               | Initiator                | 3.63 mg, 0.022 mmol       | Recrystallized from MeOH at 45 °C  |
| Diels–Alder adduct | Mechanophore crosslinker | 1.010 mg, 1.767 $\mu$ mol | Synthesized according to our previous work <sup>21</sup> (Scheme 1)                |
| Dioxane            | Solvent                  | 1 mL                      | Distilled before use   |





**Fig. 1** Schematic depiction of the experimental setup for analysis with CLSM showing the needle-punctured rectangular hydrogel sample and wet mounting onto glass slide for imaging with an inverted microscope.

**Compression.** In a second step, the punctured rectangular samples were compressed by hand with the use of laboratory tweezers, and were then subsequently visualized by CLSM. The small size of the hydrogel samples facilitated an approximately parallel placement between the tweezers (Fig. S1†). The applied force was gradually increased in order to deform the samples macroscopically, but without producing any visible failure (fracture) of the specimen. A needle-punctured hydrogel is essentially a square plate with an internal void that acts as a stress raiser upon force application.<sup>25</sup> The stress is amplified from a nominal  $\sigma_0$  to a  $\sigma_{\max}$  due to stress concentration around the tip of the flaw:

$$\sigma_{\max} = 2\sigma_0 \left( \frac{\alpha}{\rho_t} \right)^{1/2} \quad (2)$$

where  $\alpha$  is the length of a surface crack or half of the length of an internal crack, and  $\rho_t$  is the radius of curvature of the crack tip (Fig. 2). The boundaries of the samples are expected to have a negligible effect given the small region near the entrance point since their dimensions are *ca.*  $7.5d = 3$  mm, where  $d$  is the diameter of the circular hole (equal to  $400 \mu\text{m}$ ).



**Fig. 2** Surface and internal flaws in a plate and stress profile with distance along the axis. Since all materials tend to contain flaws, any externally applied force would not be uniformly distributed throughout their structure. Internal cracks and voids, thus, have the role of stress concentrators.

**Fluorescence.** A confocal laser scanning microscope (Leica LASX TCS SP8) was used to perform the analysis and record the respective images. Wet mounting of the samples onto microscope cover glasses was realized with the use of an immersion oil ( $n = 1.5180$ ). The scanning range was determined based on previous measurements<sup>21</sup> performed *via* UV-vis and fluorescence spectroscopy to determine the absorption and emission characteristics of the mechanofluorophore in solution. It can be seen in Fig. 3 that the mechanofluorophore absorbs at around 400 nm, and emits by exhibiting two characteristic peaks at 450 nm. The lower bound of the scanning range was assigned a value to maintain the minimum from the excitation laser's wavelength (Table 2).

**2.2.2 Image analysis.** An open source scientific image processing program (ImageJ)<sup>26</sup> was used to analyze the microscopy images and quantify the fluorescence intensity. A radial profile plugin was further exploited to obtain the maximum activation of the circular-shaped trace. This method produced a profile plot of normalized integrated intensities around a circular region of interest (ROI) as a function of distance from a certain point in the image (the center of the circle in this analysis). As the integrated intensity is divided by the number of the pixels in the circle, the values are normalized and comparable.

**2.2.3 Mechanical characterization.** Uniaxial compressive testing was performed at room temperature conditions with the use of a CSM micro indenter (flat punch indenter of 6 mm in diameter) to determine the mechanical response of the



**Fig. 3** UV-vis absorption spectrum (solid line) and normalized fluorescence emission spectrum (dashed line).  $\lambda_{\text{exc}} = 372$  nm.

**Table 2** Leica SP8 TCS confocal laser scanning microscopy overview of parameters

| Parameters                   |                           |
|------------------------------|---------------------------|
| Excitation laser             | UV 405 nm                 |
| Scan format ( $x \times y$ ) | $1024 \times 1024$ pixels |
| Scan speed                   | 400 Hz                    |
| Magnification                | 20 $\times$               |
| Scanning wavelength regime   | 420–550 nm                |
| Pinhole                      | Airy 1                    |
| Line average                 | 8                         |



hydrogels with varying swelling degrees. The synthesis of the cylindrical hydrogel samples (on average  $3.2 \pm 0.5$  mm in diameter, and  $3 \pm 0.3$  mm in thickness for all  $d_s$  values) was identical to that of the rectangular specimens, but without the mechanophore crosslinker (the amount of TEGDMA was properly adjusted to compensate for this). The load ( $F$ ) – displacement ( $\Delta h$ ) data were converted into stress ( $\epsilon$ ) – strain ( $\sigma$ ) curves by calculating  $\epsilon = \Delta h \cdot h_0^{-1}$  and  $\sigma = F \cdot A_0^{-1}$ , where  $h_0$  and  $A_0$  are the initial height and surface area of the cylindrical samples, respectively. The variations in sample thickness and applied pressure yield the elastic (Young's) modulus, determined from the initial slope of the respective stress–strain curve. Two specimens were subjected to uniaxial compression to obtain an average value for the modulus of elasticity.

### 3 Results and discussion

#### 3.1 Confocal laser scanning microscopy

The mechanophore activation was initially recorded by performing a  $\lambda$ -scan from 420 to 550 nm, under a laser excitation wavelength of 405 nm (Fig. S2<sup>†</sup>). The emission spectrum with the two characteristic vibronic transitions of the anthracene at around 450 nm was in agreement with previous measurements.

The recorded images were of sub-micrometer resolution with a pixel size of 758 nm. The scale bar in all images is 100  $\mu\text{m}$ .

**3.1.1 Circular trace upon puncturing with a needle.** Fig. 4 portrays exemplary images recorded during the analysis by CLSM. A circular fluorescence trace became apparent for a punctured hydrogel sample with  $d_s = 21\%$  (Fig. 4b), while the absence of a fluorescence signal verified the lack of mechanical disruption as the particular specimen was in its pristine (non-activated) state (Fig. 4a). More quantitatively, this is visualized in Fig. 4c compared to Fig. 4d distinctly reflecting the mechanophore activation. These graphs are profile plots of the integrated fluorescence intensity as a function of distance from the center of the circle and reveal a maximum value at around 200  $\mu\text{m}$  for the punctured sample, corresponding to the radius of the needle. For this particular analysis we defined specific values for both the starting and integration angle (of  $90^\circ$ ). This method yielded fluorescence intensity values that were normalized to the number of pixels and are restricted to a cone ( $\pi/2$ ), as denoted with cyan lines in Fig. 4a and b. This process facilitated the calculation of the results as it included all maximum values since (i) the use of a needle with a beveled tip exerted forces asymmetrically on the samples rather than equally in all directions, and (ii) the needle was inserted manually with a vertical placement.



**Fig. 4** CLSM images (a and b) and fluorescence intensity analysis with ImageJ (c and d) for rectangular PNIPAAm hydrogel samples crosslinked with mechanofluorophore for a scanning regime of 420–550 nm under a 405 nm excitation laser: (a and c) before and (b and d) after needle-puncturing. The scale bar is 100  $\mu\text{m}$ . The intensity values span the 0–255 scale; 0 for black and 255 for cyan. Note here that the slightly deformed fluorescence circular shape is due to cutting of the needle tip after exiting the sample. Respective CLSM images for rectangular PNIPAAm hydrogel samples with various swelling degrees are provided in Fig. S4<sup>†</sup>.



However some deviation from an angle of  $90^\circ$  to the horizontal plane must be considered.

It should be noted that the cone chosen for the pristine sample (Fig. 4a) was identical to the one for the punctured sample (Fig. 4b) only for comparison reasons, in terms of fluorescence intensity. For every image captured with the confocal laser scanning microscope, the whole circular trace was scanned sequentially by maintaining a constant integration angle at  $90^\circ$ . The cone with the highest intensity value was finally used for the analysis each time.

The calculated maximum fluorescence intensity values for all punctured samples are shown as a function of  $d_s$  in Fig. 5. The error bars in both the  $x$ -axis and the  $y$ -axis stem from the calculation of a mean value with a standard deviation from two to five measurements. It is evident that the intensity after fracture decreases with swelling degree, *i.e.* with the further hydration of the networks. A relatively high fluorescence activation might be discerned in this graph for samples with low water contents, an intermediate activation at swelling degrees around 25%, and low to near-zero activation for higher  $d_s$ .

It is important to note that the possible effect of mechano-phore dilution upon volume expansion by water-swelling was ruled out. For this, the fluorescence intensity of swollen samples after puncturing was recorded and quantitatively compared with identical microscopy measurements after the same samples were completely dried *in vacuo* (Fig. S3†). The deviation was negligible and could therefore only explain a minor fraction of the observed decrease moreover demonstrating that the activated mechanofluorophores were not susceptible to fluorescence quenching by water.

This simple needle-puncturing of the hydrogel networks with integrated mechanofluorophores could be further utilized for the local mapping of stress concentrations with the aid of CLSM, as shown in Fig. 6. The color spectrum allowed a distinct visualization of stress accumulation due to the asymmetrical beveled needle and its slightly diagonal insertion into the sample that maximized just at its tip (red-colored region). These fluorescence intensity maps are comparable to the frac-



Fig. 5 Maximum fluorescence intensity as a function of swelling degree. The data have been derived by calculating the mean value and standard deviation stemming from two to five measurements.



Fig. 6 CLSM images of a needle-punctured PNIPAAm hydrogel sample: (a) fluorescence; (b) bright field; (c) overlay. The intensity values span the 0–255 scale; 0 for black and 255 for red (a); 0 for black and 255 for white (b). The scale bar is  $100\ \mu\text{m}$ .

ture mechanics models employed for stress analysis in typical material-failure scenarios<sup>27</sup> and hence could be exploited in conjunction for the study of materials fracture mechanics.

### 3.1.2 Crack propagation upon compression with tweezers

*Low degree of swelling.* For  $d_s$  in the order of 10 to 22%, deformation of the hydrogels was hard to achieve as these were in a glassy state and exhibited a rigid behavior. The samples were deformed upon excess application of force and did not return to their original shape instantly, as the polymeric chains were fixed in a frozen state and were not able to rotate and translocate freely. A typical deformation structure of glassy polymers can be discerned in Fig. 7. Crazes formed due to the application of external force and appeared as a surface phenomenon with long and fine lines emanating from the circular disturbance. The lack of any visible fluorescent crazes suggests that mechanical integrity of the network remained unaffected, as these did not evolve into macroscopic cracks. We hypothesize that the lack of fluorescent crazes might be attributed to the correlation of crazing to molecular separation with disentanglement and sliding of polymer chains, rather than the scission of covalent bonds.<sup>28</sup>





**Fig. 7** CLSM images of a rectangular PNIPAAm hydrogel sample swollen to  $d_s = 10\%$  after puncturing with a needle and subsequent compression with tweezers. The images on the left (a, c and e) and the images on the right (b, d and f) compose a whole circular trace of one sample. Fluorescence (a and b); bright field (c and d); overlay (e and f). The scale bar is  $100\ \mu\text{m}$ . Note how the circular fluorescence trace falls on the needle perimeter.



**Fig. 8** CLSM images of a rectangular PNIPAAm hydrogel sample swollen to  $d_s = 28\%$  after puncturing with a needle and subsequent compression with tweezers. Annotations as in Fig. 6. The scale bar is  $100\ \mu\text{m}$ . Note how the fluorescence follows the propagation of the crack.

*Intermediate degree of swelling.* For  $d_s = 28\%$  the samples were easily deformed and returned to their original shape without any delay after removal of force application. It appears that in this regime, the polymeric chains exhibited a rubber-like elasticity. The propagation of a crack initiating from the circular hole can be seen in Fig. 8. The main crack propagated at some distance from the tip and then forked into secondary branches by distributing the stress intensity locally through bifurcation.

*High degree of swelling.* Fragmentation of the samples was effortlessly (and unintentionally) induced upon compression with laboratory tweezers for  $d_s$  from 36% up to 123%, as the highly swollen samples were very fragile. Fig. 9 portrays the formation of channels originating locally from the region where the needle was inserted into the samples. The channels (discerned by their distinct fluorescence lines) were formed upon application of force to redistribute the water throughout the hydrogel network and/or expel any excess amount. The rearrangement of  $\text{H}_2\text{O}$  related to complex fluid–solid interaction during compression and subsequent deformation of

the hydrogel samples. Given the unconfined compression that was realized here, and the anticipated variation in strain response throughout the hydrogel network, the effect of poroviscoelastic parameters needs to be considered.<sup>29</sup> However, this is beyond the scope of this study.

### 3.2 Elastic modulus

Given the biological and biomedical applications of hydrogels, it is crucial to maintain control not only over their physico-chemical response, but also their mechanical behavior over a range of swelling degrees. In this study, the elastic (Young's) modulus of the hydrogel networks was determined, as it is a fundamental and well-defined mechanical property that can reflect their integrity and can be fine-tuned upon their water absorption capability.<sup>30</sup>

Fig. 10b depicts the measured elastic moduli  $E$  as a function of  $d_s$ , calculated from the respective stress–strain curves (Fig. 10a). The non-swollen samples were relatively stiff with  $E$  exceeding the value of 100 MPa. A decreasing  $E$  with increasing  $d_s$  can be discerned, with values lying around a few MPa for





**Fig. 9** CLSM images of a rectangular PNIPAAm hydrogel sample swollen to  $d_s = 36\%$  after puncturing with a needle and subsequent compression with tweezers. The images on the left (a, c and e) and the images on the right (b, d and f) illustrate different regions of the same sample. Annotations as in Fig. 7. The scale bar is 100  $\mu\text{m}$ .

$d_s \sim 20\text{--}30\%$ , while dropping as low as  $<1$  MPa for  $d_s$  up to 180%. Given that native biological materials cover multiple orders of magnitude regarding their elastic modulus,<sup>31</sup> this

swelling-induced softening of the PNIPAAm networks can serve as a good model to investigate the mechanical properties of bio-analogous hydrogels.

As the mechanical testing for the determination of the elastic modulus was based on an unconfined configuration, the cylindrical hydrogel samples were allowed to have a non-restricted expansion in the radial direction during compression. Therefore, the enhanced variation in moduli values for samples with relatively high water content, *i.e.*  $d_s > 30\%$ , (Fig. 10b) could be attributed to the cumulative effect of visco-elastic response of the polymeric chains along with the poro-elastic behavior due to the  $\text{H}_2\text{O}$  molecules in the networks, as it is apparent that for these swelling degrees the presence of water becomes significant.<sup>32,33</sup>

### 3.3 Correlation between macroscopic observations and fluorescence intensity

We exerted low levels of mechanical disruption to the networks with the use of a needle, which was employed before in the study of puncture-resistant hydrogels<sup>34</sup> and the fracture behavior of soft gels and solids.<sup>24,35</sup> Alongside, the compression of the samples with the use of common laboratory tweezers provided valuable information over the macroscopic behavior and deformation response of the hydrogels. The optical transparency of the samples allowed us to correlate the subsequent micro-scale analysis and visualization of bond scission by fluorescence intensity to the macroscopic structure and properties.

Based on the macroscopic observations of the fracture behavior of the hydrogels, we anticipated that upon hydration of the gels with excess of water, the fluorescence intensity would maximize as a result of extensive bond scission within the brittle and fragile materials. However, we found that for high  $d_s$  the fluorescence intensity leveled off to near-zero.

These macroscopic observations on the deformation response of the hydrogels corroborated the presence of a glass-to-rubber transition upon hydration. It can be argued that



**Fig. 10** Mechanical behavior of cylindrical PNIPAAm hydrogel samples determined by uniaxial compression: (a) stress–strain curve of a hydrogel sample swollen to  $d_s = 27\%$ . The linear fit of the curve for the determination of the elastic modulus is also shown with a solid line. Respective curves and linear fits for all measured samples are provided in Fig. S5.† (b)  $E$  as a function of  $d_s$ . The error bars are the result of two uniaxial compression measurements performed on individual hydrogel specimens.



beyond a critical  $d_s$ , the interaction between the solid and the liquid phase of the hydrogel was altered, also affecting the fracture mechanism. Indeed, it became apparent that beyond  $d_s \sim 25\%$  the fluorescence signal leveled off and reached a plateau (Fig. 5). An analogous trend was recorded for  $E$  as a function of  $d_s$  (Fig. 10b). Although the low level force applied to puncture a dry sample (in its glassy state) and a swollen sample (in a rubbery state) possibly varied, this did not affect the fracture process, since the insertion of the needle through the sample translated into the cutting of a finite number of polymer chains.

We ascribed this supposed glass-to-rubber transition to the reduction of physical crosslinks between the polymer chains from a certain  $d_s$  on, due to their saturation with water. We hypothesized that the hydrophilic NIPAAm moieties were initially hydrated upon exposure of a dry sample to  $H_2O$ , as the completely hydrated state of PNIPAAm relies on the formation of three hydrogen bonds per repeating unit.<sup>36</sup> These water molecules were directly attached to the polar groups and constituted the “bound” water that was strongly associated with the polymer. Additionally absorbed water existed in a “free” state that could be easily removed, as it filled the free volume in the polymer mesh, and in a “semi-bound” state relating to the physically trapped water. These reduced intersegmental attractions between the polymer chains due to their preferential hydrogen bonding to water were proposed to lead to a significant reduction in tensile strength in UV-cured crosslinked polymers upon water absorption when exposed to high humidity conditions.<sup>37</sup> Indeed, the secondary hydrogen bonds that were formed between the chains might contribute to the rigidity and toughness of the polymer,<sup>38</sup> and lead to a reduction in elastic modulus with increasing water content as they become fewer.<sup>39</sup> The increased presence of physical crosslinks in the form of intermolecular H-bonds between the polymeric chains was also found to play a dominant role in the behavior of PNIPAAm gels, as their elastic moduli in the collapsed state were significantly higher than those in the swollen state.<sup>40</sup> Generally, H-bonds are mechanochemically weak and can be readily disrupted. However, our results corroborate a synergistic effect between physical and chemical crosslinks, as a mechanism for stress distribution among the chains and effective transfer of load to the mechanophores, which is known to be influenced by parameters such as chain entanglements and interactions among them.<sup>41</sup>

Besides the existence of different states of water in the hydrogel, the effect of local inhomogeneities should also not be ruled out.<sup>42</sup> In heterogeneous polymeric systems, such as the one under investigation, multiple length scales are present with both short and long chain segments throughout the formed network. Upon force application, stress concentrates on the shortest chain which must already be in its fully stretched state for failure to occur. Therefore, a “pre-stretched” state of several small segments can also be considered to cause a type of random-scission along the network leading to near-zero mechanophore scission.

## 4 Conclusions

We showed that the integration of mechanofluorophores into PNIPAAm hydrogel networks in the form of chemical crosslinks facilitated the investigation of their molecular structure – macroscopic property interrelationships *via* the mapping of mechanical stresses and the local imaging of forces with the use of confocal laser scanning microscopy with  $\mu\text{m}$ -resolution. With this unprecedented analysis method, we elucidated the mechanical behavior of these PNIPAAm hydrogels with different water contents. We found that the dissolution of H-bonds upon swelling of the materials resulted in gradually reduced mechanical performance and completely altered their respective fracture mechanisms. Contrarily to our expectations, the easy fracture of highly swollen samples led to the scission of fewer mechanofluorophores compared to the lowly swollen states, though covalent bond scission along the network backbone was abundant. These results suggest that H-bonds significantly contribute to stress-distribution in PNIPAAm hydrogels and assist to scatter covalent bond scission throughout the material. The latter may function as a sacrificial element warranting the mechanical integrity of PNIPAAm hydrogels with low to moderate water content. This might also be the case for other hydrogels, other acrylamides for instance, with analogous H-bonding behavior. Concludingly, we believe that this unique methodology will be useful for the investigation of a wide range of mechanical phenomena in soft matter, even in a directly quantitative context, that proved to be difficult to analyze with traditional mechanical characterization techniques.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The Erasmus<sup>+</sup> framework is acknowledged for supporting the mobility of M. S. as a recent PhD graduate to DWI. R. G. and C. B. are grateful for support by a Freigeist-Fellowship of the Volkswagen Foundation. Parts of the analytical investigations were performed at the Center for Chemical Polymer Technology CPT, which was supported by the European Commission and the federal state of North Rhine-Westphalia (Grant No. 300088302). Financial support is acknowledged from the European Commission (EUSMI, 731019).

## References

- 1 E. Caló and V. V. Khutoryanskiy, *Eur. Polym. J.*, 2015, **65**, 252–267.
- 2 M. A. Haq, Y. Su and D. Wang, *Mater. Sci. Eng., C*, 2017, **70**, 842–855.



- 3 C. Creton and M. Ciccotti, *Rep. Prog. Phys.*, 2016, **79**, 046601.
- 4 D. Lee, M. M. Rahman, Y. Zhou and S. Ryu, *Langmuir*, 2015, **31**, 9684–9693.
- 5 J. T. Pham, F. Schellenberger, M. Kappl and H.-J. Butt, *Phys. Rev. Mater.*, 2017, **1**, 015602.
- 6 S. Kundu and A. J. Crosby, *Soft Matter*, 2009, **5**, 3963–3968.
- 7 A.-D. N. Celestine, N. R. Sottos and S. R. White, *Strain*, 2019, **55**, e12310.
- 8 D. A. Davis, A. Hamilton, J. Yang, L. D. Cremar, D. Van Gough, S. L. Potisek, M. T. Ong, P. V. Braun, T. J. Martinez, S. R. White, J. S. Moore and N. R. Sottos, *Nature*, 2009, **459**, 68–72.
- 9 R. Göstl, J. M. Clough and R. P. Sijbesma, *Mechanochemistry in Materials*, Royal Society of Chemistry, 2017, pp. 53–75.
- 10 J. Li, C. Nagamani and J. S. Moore, *Acc. Chem. Res.*, 2015, **48**, 2181–2190.
- 11 Z. Chi, X. Zhang, B. Xu, X. Zhou, C. Ma, Y. Zhang, S. Liu and J. Xu, *Chem. Soc. Rev.*, 2012, **41**, 3878–3896.
- 12 B. R. Crenshaw and C. Weder, *Macromolecules*, 2006, **39**, 9581–9589.
- 13 Y. Chen and R. P. Sijbesma, *Macromolecules*, 2014, **47**, 3797–3805.
- 14 J. W. Kim, Y. Jung, G. W. Coates and M. N. Silberstein, *Macromolecules*, 2015, **48**, 1335–1342.
- 15 C. Calvino, L. Neumann, C. Weder and S. Schrettl, *J. Polym. Sci., Part A: Polym. Chem.*, 2017, **55**, 640–652.
- 16 K. R. Fitch and A. P. Goodwin, *Chem. Mater.*, 2014, **26**, 6771–6776.
- 17 F. Cellini, L. Block, J. Li, S. Khapli, S. D. Peterson and M. Porfiri, *Sens. Actuators, B*, 2016, **234**, 510–520.
- 18 M. Fernández-Castano Romera, R. Göstl, H. Shaikh, G. Ter Huurne, J. Schill, I. Voets, C. Storm and R. Sijbesma, *J. Am. Chem. Soc.*, 2019, **141**, 1989–1997.
- 19 X.-Q. Wang, C.-F. Wang, Z.-F. Zhou and S. Chen, *Adv. Opt. Mater.*, 2014, **2**, 652–662.
- 20 T. Matsuda, R. Kawakami, R. Namba, T. Nakajima and J. P. Gong, *Science*, 2019, **363**, 504–508.
- 21 R. Göstl and R. P. Sijbesma, *Chem. Sci.*, 2016, **7**, 370–375.
- 22 A. Halperin, M. Kröger and F. M. Winnik, *Angew. Chem., Int. Ed.*, 2015, **54**, 15342–15367.
- 23 F. Afroze, E. Nies and H. Berghmans, *J. Mol. Struct.*, 2000, **554**, 55–68.
- 24 S. Das and A. Ghatak, *J. Mater. Sci.*, 2011, **46**, 2895–2904.
- 25 W. Callister and D. Rethwisch, *Materials Science and Engineering: An Introduction*, Wiley, 8th edn, 2009.
- 26 C. A. Schneider, W. S. Rasband and K. W. Eliceiri, *Nat. Methods*, 2012, **9**, 671–675.
- 27 A. Mohammadipour and K. Willam, *J. Appl. Mech.*, 2016, **83**, 071003.
- 28 R. N. Haward, H. E. Daniels and L. R. G. Treloar, *J. Polym. Sci., Part B: Polym. Phys.*, 1978, **16**, 1169–1179.
- 29 D. Caccavo, S. Cascone, G. Lamberti and A. A. Barba, *Chem. Soc. Rev.*, 2018, **47**, 2357–2373.
- 30 K.-i. Hoshino, T. Nakajima, T. Matsuda, T. Sakai and J. P. Gong, *Soft Matter*, 2018, **14**, 9693–9701.
- 31 R. Lanza, R. Langer and J. Vacanti, *Principles of Tissue Engineering*, Elsevier Science, 2011.
- 32 J. Delavoipière, Y. Tran, E. Verneuil and A. Chateauminois, *Soft Matter*, 2016, **12**, 8049–8058.
- 33 M. L. Oyen, *Int. Mater. Rev.*, 2014, **59**, 44–59.
- 34 Y. Liu, W. Zhou, Q. Zhou, K. Peng, A. Yasin and H. Yang, *RSC Adv.*, 2017, **7**, 29489–29495.
- 35 S. Fakhouri, S. B. Hutchens and A. J. Crosby, *Soft Matter*, 2015, **11**, 4723–4730.
- 36 L. Tavagnacco, E. Zaccarelli and E. Chiessi, *Phys. Chem. Chem. Phys.*, 2018, **20**, 9997–10010.
- 37 D. A. Bolon, G. M. Lucas, D. R. Olson and K. K. Webb, *J. Appl. Polym. Sci.*, 1980, **25**, 543–553.
- 38 N. Lakhera, K. E. Smith and C. P. Frick, *J. Appl. Polym. Sci.*, 2013, **128**, 1913–1921.
- 39 A. Hamouda, *J. Mater. Process. Technol.*, 2002, **124**, 238–243.
- 40 T. Takigawa, T. Yamawaki, K. Takahashi and T. Masuda, *Polym. Gels Networks*, 1998, **5**, 585–589.
- 41 C. Lee, B. Beiermann, M. Silberstein, J. Wang, J. Moore, N. Sottos and P. Braun, *Macromolecules*, 2013, **46**, 3746–3752.
- 42 M. N. Silberstein, L. D. Cremar, B. A. Beiermann, S. B. Kramer, T. J. Martinez, S. R. White and N. R. Sottos, *J. Mech. Phys. Solids*, 2014, **63**, 141–153.

