The mechanical properties of the substrate or matrix play a critical role in cell motility, cytoskeletal organization, gene expression, and stem cell differentiation (1–5). Designing materials for in vitro cell culture and tissue engineering applications therefore requires careful attention to both mechanical and biochemical properties. The model material system most commonly employed for studies of substrate stiffness effects is covalently cross-linked hydrogels of polyacrylamide (pAAm), which can be formulated with moduli covering a wide range of physiologically relevant values, ~0.010–50 kPa (pAAm), which can be formulated with moduli covering a wide range of physiologically relevant values, ~0.010–50 kPa (5). Thin layers (~10–500 μm) are typically used, yielding flexible and often fragile gels that are coated on a rigid substrate, e.g., a glass coverslip, to facilitate handling.

A commonly encountered, but poorly characterized, problem arises when the modulus of the gels is reduced too far: sharp lines resembling cracks may appear on the surface of the gel upon immersion in aqueous media (2,6–8). These features result from attachment of the gel to a rigid substrate. When the gel is immersed in an aqueous environment, it swells due to the osmotic pressure difference with the surrounding medium, and equilibrium is attained when this osmotic pressure difference is matched by the elastic stress experienced by the surface-attached gel. If an unconstrained gel swells by a factor $l_f$ in each dimension, then the effective strain $\varepsilon$ experienced by an identical gel grafted to a rigid substrate is defined by $\varepsilon = 1 - \frac{1}{l_f}$. Our previous study of pAAm gels, with estimated shear moduli $G$ of ~0.6–24 kPa and thicknesses of 3 μm–1 mm, yielded creases beyond a critical strain $\varepsilon_c$ ranging from 0.30 to 0.37 with an average of 0.33 (7).

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The insensitivity of $\varepsilon_c$ to material properties and length scale agrees with theoretical predictions (11), and suggests that the key to preventing creases is to tune the balance between modulus and osmotic stress to maintain a level of swelling below $\lambda_f \approx 1.5$. We first consider how these two factors depend on gel composition, then validate this approach using the model system of pAAm gels containing varying contents of AAm monomer $w_{AAm}$ and bisacrylamide (Bis) cross-linker $w_c$ (Table 1).
According to classical theories of network elasticity, the shear modulus is proportional to the density of elastically active strands times thermal energy $k_B T$. For the affine model (12) of a network with $N$ Kuhn steps of length $b$ between junctions, and polymer volume fraction $f_0$, 

$$G = \frac{\phi_0}{b^3} k_B T. \quad (1)$$

Thus, the modulus scales linearly with the concentration of polymer in the gel, but inversely with the length of the chains. In the idealized case that each cross-linker is active and every monomer is incorporated into the network, $N$ is proportional to the ratio of monomer to cross-linker concentrations. Because $f_0$ is also proportional to monomer concentration (assuming the amount of cross-linker is small), we may expect from Eq. 1 that $G$ should be proportional to $w_x$ and independent of $w_m$. In reality, nonidealities in gel structure generally make $G$ an increasing function of both monomer and cross-linker concentration.

The osmotic stress $\pi$ driving swelling of a polymer gel is roughly equal to that of a semidilute polymer solution of the same composition. When immersed in a good solvent, the initial driving force for swelling (12) is given by

$$\pi \sim \frac{\phi_0 k_B T}{b^3}. \quad (2)$$

This is a strongly-increasing function of polymer concentration, but does not depend on $N$, thus it is nearly independent of cross-linker concentration. Because swelling is determined by a balance between $G$ and $\pi$, comparing Eq. 1 and Eq. 2 suggests that $\lambda_f$ can be reduced while maintaining fixed $G$ by decreasing $\phi_0$ (or $w_m$) and simultaneously increasing $w_x$.

To test this assertion, we studied pAAm gels where at each composition (Table 1) two identical gels were polymerized between coverslips; however, in one case the bottom coverslip was treated with an adhesion promoter, whereas in the other case it was treated with a release coating allowing unconstrained swelling. Equilibrium values of $\lambda_f$ for unconstrained gels are plotted in Fig. 3 A, with open symbols representing compositions where surface-attached gels formed creases and solid symbols representing those that did not.

Two important points should be noted. First, whereas $\lambda_f$ increases as $w_x$ is lowered for all values of $w_m$, there is no simple trend in the dependence of swelling on $w_m$. This is at first surprising, because the prediction for a covalently cross-linked network swelled in a good solvent (13) is

$$\lambda_f \sim \phi_0^{0.25} A^{0.2}, \quad (3)$$

### TABLE 1 Compositions and moduli of pAAm gels

<table>
<thead>
<tr>
<th>AAm* content $w_m$ (wt %)</th>
<th>Bis† content $w_x$ (wt %)</th>
<th>$G$ (Pa)</th>
</tr>
</thead>
<tbody>
<tr>
<td>3.0%</td>
<td>0.02–0.2%</td>
<td>12–160</td>
</tr>
<tr>
<td>5.0%</td>
<td>0.02–0.2%</td>
<td>60–1100</td>
</tr>
<tr>
<td>7.5%</td>
<td>0.02–0.2%</td>
<td>—</td>
</tr>
<tr>
<td>10.0%</td>
<td>0.02–0.2%</td>
<td>400–9600</td>
</tr>
<tr>
<td>12.0%</td>
<td>0.02–0.2%</td>
<td>—</td>
</tr>
</tbody>
</table>

*Acrylamide.
†Bisacrylamide.

FIGURE 1 (A) Phase-contrast optical image of a creased pAAm gel surface (1 kPa; 10 wt % AAm, 0.01 wt % sodium acrylate, 0.02 wt % Bis. Scale bar, 200 μm. (B) A confocal fluorescence image (x,y,z scale 1:1:3) after swelling reveals sharp folds in the gel surface (inset: x-y slice 32 μm below the surface).

FIGURE 2 (A) Phase-contrast images indicate that adult neural stem cells sensed the surface creases and projected neurites along the creases on the unstable pAAm formulation. (B) Immunostaining for cell phenotypes (10).

FIGURE 3 (A) Linear swelling $\lambda_f$ versus Bis content $w_x$ for different AAm contents $w_m$ (see legend). Open symbols represent compositions where surface-attached gels formed creases. Dotted and dashed lines at $\lambda_f = 1.5$ and $w_x = 0.028$ wt %, respectively, delineate creased from stable gels. (B) Shear moduli $G$ of pAAm gels. (Open symbols) Compositions where surface-attached gels formed creases. (Solid line) Predicted modulus if all cross-links were elastically active. Estimated uncertainties correspond approximately to marker sizes in both plots.
and raising $w_m$ at fixed $w_s$ should increase both $\phi_0$ and $N$. However, nonidealities in network structure can again lead to significant deviations from this prediction, and prior reports have even shown nonmonotonic dependence of swelling on monomer concentration for similar systems (14).

The second point concerns the transition from the flat to the creased state for surface-attached gels. The dotted line at $\lambda_f = 1.5$, corresponding to the previously-established criterion of $\epsilon_c = 0.33$ (7), captures the onset of creasing with the exception of 3 wt % AAm gels (red circles in Fig. 3A) where the transition occurs above $\lambda_f = 1.6$. Because swelling is largely insensitive to $w_m$, the transition from flat to creased states may also be described by the dashed line at $w_s = 0.028$ wt %, which cleanly divides the data over the full range of $w_m$ studied. The guideline $\lambda_f \approx 1.5$ remains useful, because it is general to other chemistries and can thus be easily translated to different material systems.

The values of $G'$ measured by oscillatory shear rheology (which are nearly equal to $G$ for a predominantly elastic material) for gels with 3, 5, and 10 wt % AAm, plotted in Fig. 3B (full data in Table S1 in the Supporting Material), are in good agreement with a previous study (5). The dashed line at $w_s = 0.028$ wt % again demarcates the transition from flat to creased supported gel samples. The solid line represents the predicted shear modulus if every cross-linker were elastically active: although the data follow the same trend, they fall below the predicted curve, also in agreement with the earlier study (5). Unlike the swelling data in Fig. 3A, the modulus is quite sensitive to monomer concentration. For 10 wt % AAm, the modulus can be reduced to ~600 Pa by lowering Bis concentration, beyond which further reductions in $w_s$ lead to creasing. By lowering the AAm concentration to 3 wt %, however, the modulus can be reduced below 100 Pa without triggering the surface instability. We have not studied the rheological properties of 12 wt % gels, but their moduli should be tunable from ~50 kPa at high cross-linker concentration down to ~2 kPa before the onset of creases (5). Thus, appropriate choices of monomer and cross-linker content allow the modulus to be tuned from 50 kPa to below 100 Pa without encountering creases.

We offer one caveat: even if the composition is chosen to avoid creases, substantial swelling of the surface-attached gel (up to ~2-fold) (7) may occur. Thus, the modulus of the swelled gel will be slightly different than determined by rheology for the unswellled gel, and in general, there may be significant anisotropy between the in-plane and out-of-plane incremental moduli due to the lateral prestress in the swollen surface-attached gel. For gels close to the instability, traction forces applied by the cell may even exceed the critical compression and cause local folding. Rather than managing the balance between osmotic stress and gel modulus, an alternative approach that avoids these complications is to eliminate swelling stresses altogether by allowing an unconstrained gel to swell freely and subsequently attaching it to a rigid support. Although we have found this method to be effective for preparing gels with moduli of ~0.5 kPa and above, it is difficult especially for soft gels and likely not suitable for high-throughput cell-based studies.

In summary, surface creasing of pAAm substrates can significantly influence cell behavior and must be carefully considered for the soft substrates increasingly used in mechanotransduction experiments. We anticipate that this work can guide the synthesis of both smooth and creased soft cell substrates for cell engineering efforts.

**SUPPORTING MATERIAL**

Materials and methods, complete modulus and swelling data for Fig. 3B, and references are available at http://www.biophysj.org/biophysj/supplemental/S0006-3495(10)01195-1.

**ACKNOWLEDGMENTS**

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**REFERENCES and FOOTNOTES**


10. Neuronal differentiation conditions (retinoic acid and forskolin) were used during culture. $\beta$-Tubulin III (red), GFAP (blue), and Nestin (green) mark neurons, astrocytes, and immature stem cells, respectively.


