Myotubes differentiate optimally on substrates with tissue-like stiffness: pathological implications for soft or stiff microenvironments

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Abstract
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Keywords
myofibrillogenesis, patterning, adhesion, differentiation, muscular dystrophy

Comments
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Myotubes differentiate optimally on substrates with tissue-like stiffness: pathological implications for soft or stiff microenvironments

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Contractile myocytes provide a test of the hypothesis that cells sense their mechanical as well as molecular microenvironment, altering expression, organization, and/or morphology accordingly. Here, myoblasts were cultured on collagen strips attached to glass or polymer gels of varied elasticity. Subsequent fusion into myotubes occurs independent of substrate flexibility. However, myosin/actin striations emerge later only on gels with stiffness typical of normal muscle (passive Young's modulus, $E \approx 12$ kPa). On glass and much softer or stiffer gels, including gels emulating stiff dystrophic muscle, cells do not striate. In addition, myotubes grown on top of a compliant bottom layer of glass-attached myotubes (but not softer fibroblasts) will striate, whereas the bottom cells will only assemble stress fibers and vinculin-rich adhesions. Unlike sarcomere formation, adhesion strength increases monotonically versus substrate stiffness with strongest adhesion on glass. These findings have major implications for in vivo introduction of stem cells into diseased or damaged striated muscle of altered mechanical composition.

Introduction

Injection of mesenchymal stem cells into an ever-stiffening fibrotic infarct zone in the heart leads to cellular expression of myocyte markers, but differentiation clearly stops short of myofibril assembly (Shake et al., 2002). The reason is unclear, but physical aspects of the microenvironment, especially mechanical compliance of matrix (Pelham and Wang, 1997) and perhaps adjacent cells, are becoming increasingly linked to gene expression and protein organization. Sufficient substrate stiffness appears particularly important to anchorage-dependent cells, which often rely on a finite resistance to cell-generated forces in order to induce outside-in mechanical signals. Such signals feed back into cell tension (Wang et al., 2002), cell adhesion (Choquest et al., 1997; Beningo et al., 2001), posttranslational modification (Pelham and Wang, 1997; Beningo et al., 2001), protein expression (Cukierman et al., 2001), cytoskeletal organization (Cukierman et al., 2001; Engler et al., 2004a), and even cell viability (Wang et al., 2000). It seems fair to hypothesize that substrate stiffness imparts new meaning to “anchorage dependence.” With contractile cells such as muscle cells, one might hypothesize that coupling to a soft or stiff matrix is extremely important to cell function. Myotubes ultimately transmit actomyosin contractions through their attachments to matrix. If a matrix or substrate is fully rigid, as in conventional cultures on polystyrene or coverslips, contractile efforts will clearly be isometric. If a matrix is more compliant, such as with a collagen gel, myocytes will be able to contract, but in a manner that depends on the effective stiffness of the underlying substrate (Engler et al., 2004a). Analogous to the wrinkling of films by cells (Harris et al., 1980; Oliver et al., 1995), cells generate traction forces on collagen-coated polyacrylamide (PA) gel substrates. Cells also spread in area to a greater extent on stiff, yet compliant substrates compared with softer substrates (Lo et al., 2000; Engler et al., 2004a).

A number of muscle and related pathologies involve changes in matrix properties. Beyond myocardial infarctions mentioned above, atherosclerosis generally involves smooth muscle cells (SMCs) in a matrix that is certainly...
altered and perhaps softened (Campbell and Chamley-Campbell, 1981; Glukhova and Koteliansky, 1995; Stenmark and Mecham, 1997). Muscular dystrophies are also associated with changes in matrix: the diaphragm of the dystrophin-deficient mdx mouse is more fibrotic and thus much stiffer than a normal mouse diaphragm (Stedman et al., 1991). In the same studies, muscle precursor satellite cells were found to be as abundant in dystrophic tissue as in normal tissue, but mdx did not seem successful in regenerating differentiated muscle.

The work here attempts to elucidate a role for matrix or tissue stiffness in striated muscle differentiation. Although the conventional in vivo matrix for myotubes is an ∼1-μm-thick (or less) basement membrane of laminin and collagen, tissue compliance convolves compliance of this matrix with cellular compliance. Our ex vivo studies here of normal versus pathological tissue stiffness first set the substrate stiffness scale for subsequent in vitro studies. To controllably culture skeletal muscle on suitably compliant substrates, the strong tendency of fusing myocytes to branch and overlay each other must be limited. Micropatterned, compliant substrates solve this problem and allow for isolation of mechanisms underlying cell–matrix adhesion, extending to its role in differentiation. The process typically begins with myoblast anchorage to the substrate, cell spreading, withdrawal from cell cycle, and cell fusion into nascent myotubes. Increasing expression and assembly of cytoskeleton and adhesions ultimately concludes with characteristic striations along the contractile myotube. This last step proves to be especially sensitive to substrate compliance. The results thus reveal an optimal, tissue-like matrix that is stiff, yet compliant enough to balance cell adhesion, contractility, and ultimately differentiation. Micropatterned, compliant substrates solve this problem (Fig. 1) and allow for isolation of mechanisms underlying cell–matrix adhesion, extending to its role in differentiation.

Table 1. Passive elasticity of EDL muscle (mean ± SD) dissected from normal (C57) and dystrophic (mdx) mice, and probed by AFM indentation (Fig. 2)

<table>
<thead>
<tr>
<th>Mouse</th>
<th>Mean muscle stiffness, ( E ) (kPa)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C57</td>
<td>12 ± 4</td>
</tr>
<tr>
<td>mdx</td>
<td>18 ± 6</td>
</tr>
</tbody>
</table>

Results

Skeletal muscle stiffness ex vivo: normal and dystrophic

Skeletal muscle is obviously a “soft tissue” rather than hard as bone. To quantitate the transverse, passive elasticity of a representative muscle, extensor digitorum longus (EDL) muscles were excised from both normal (C57) mice and dystrophin-deficient (mdx) mice and probed locally along their lengths by atomic force microscopy (AFM; Fig. 2). The transverse direction is, of course, the direction relevant to lateral adhesion between cells and matrix. Hertz cone model fits of the indenting force yields a transverse Young’s modulus, \( E \) (Domke and Radmacher, 1998; Rotsch et al., 1999).

Data was averaged over multiple locations within each sample (\( n = 3 \)) from normal mice of different ages, where the predominant modulus was \( E_{\text{Muscle}} = 12 \pm 4 \) kPa (Table I; 85% of data). The result is in agreement with the AFM-determined elasticity of C2C12 cells (up to 1 wk): \( E_{\text{C2C12}} = 12–15 \) kPa (Collinsworth et al., 2002).

From mdx mice, EDL appeared significantly stiffer with a predominant modulus \( E_{\text{mdx}} = 18 \pm 6 \) kPa (55% of data). As with the C57 control mice, the average muscle elasticity was

Figure 2. Passive stiffness of normal and dystrophic muscle tissue. EDL muscle was dissected from normal (C57) and dystrophic (mdx) mice and probed in buffer by AFM. The inset schematic depicts the AFM probe indenting a thick tissue section on a glass support. Indentations of 1–2 μm were randomly made on ∼100-μm-thick samples at 10 or more locations per left and right EDL samples (\( n = 387 \)). Force versus indentation curves were fit with a Hertz cone model (Engler et al., 2004a) in order to determine the passive elasticity as a Young’s modulus, \( E \). Table I lists the predominant stiffness for both normal C57 muscle and mdx-derived muscle.
also a few with
curs at an elastic modulus below.
Stiff or rigid substrates, defined by
foster cell spreading and include the tissue-like stiffness,
area still tends to increase with substrate stiffness
regardless of elastic modulus (Fig. 3 C), although mean cell
stiff substrates after just 4 h. By 24 h, cells are spindle-shaped
show that myoblasts spread and elongate more on increasingly
number of indentations with $E$
were plated at low densities on homogeneous collagen I–coated
Spreading, elongation, and cooperative alignment of myo-
Myoblasts in vitro sense substrate stiffness
ent on substrate elasticity, which is consistent with the cell

time indicates that spreading (Fig. 3 B) becomes less depen-
dent on substrate elasticity, which is consistent with the cell
e elongation results (Fig. 3 C). As shown below, subsequent
fusion of spindle-shaped myoblasts into nascent myotubes
occurs on all substrates regardless of elastic modulus.
For the spindle-shaped myoblasts, Fig. 3 D demonstrates
“mechanosensitive” coalignment of adjacent cells. Effects of
cell tractions transmitted through gels to neighboring cells
were quantified here by an orientational correlation function
(OCF, Eq. 2) evaluated between any two cells
orientational correlation function (OCF, 
0.5 (indicated by a dashed line); and for anti-
correlated interactions $OCF = 0$. The fit,
$OCF = B e^{−r} + (1 − B)$, gives a decay
constant $A$ and the long range orientation
$B$ ($B = 0.5$ is random). Cells in close
proximity thus prefer to align with their
nearest neighbors, facilitating formation of
myotubes. Error bars are SEM.

Myoblasts in vitro sense substrate stiffness
Spreading, elongation, and cooperative alignment of myo-
blasts depend on substrate stiffness. C2C12 skeletal myoblasts
were plated at low densities on homogeneous collagen I–coated
glass as well as PA gels of varied elastic modulus $E$. Fig. 3 (A–C)
show that myoblasts spread and elongate more on increasingly
stiff substrates after just 4 h. By 24 h, cells are spindle-shaped
regardless of elastic modulus (Fig. 3 C), although mean cell
area still tends to increase with substrate stiffness $E$ (Fig. 3 B).
The stiffest gels and glass give statistically similar results for
cell area ($P = 0.4$). Interestingly, this asymptotic behavior occurs
at an elastic modulus below $E_{\text{Muscle}}$.
The strong dependence of cell spreading at low elastic
modulus, $E < E_{\text{Muscle}}$, and the asymptotic behavior of cells
spreading on stiff gels and glass substrates fits a simple hyper-
bolà (Engler et al., 2004a) that suggests a specific and sat-
nurable mechanosensitive response:

$$\text{Area} = aE/(E_{1/2}^{\text{spread}} + E)$$ (1)

where $E_{1/2}^{\text{spread}}$ is a half-max modulus for ECM analogous
to dissociation constants for biochemical interactions. Best-
fit parameters of $E_{1/2}^{\text{spread}} \sim 2.5–5$ kPa are listed in Table II.
Stiff or rigid substrates, defined by $E > E_{1/2}^{\text{spread}}$ (e.g., glass),
 foster cell spreading and include the tissue-like stiffness, $E_{\text{Muscle}}$. Soft substrates with $E < E_{1/2}^{\text{spread}}$ (e.g., 1 kPa gel)
elicit very little cellular response. A decrease in $E_{1/2}^{\text{spread}}$ with
time indicates that spreading (Fig. 3 B) becomes less depen-
dent on substrate elasticity, which is consistent with the cell
e elongation results (Fig. 3 C). As shown below, subsequent
fusion of spindle-shaped myoblasts into nascent myotubes
occurs on all substrates regardless of elastic modulus.

<table>
<thead>
<tr>
<th>Hyperbolic fit</th>
<th>4 h</th>
<th>24 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>$a$ ($\mu m^2$)</td>
<td>2,000</td>
<td>2,100</td>
</tr>
<tr>
<td>$E_{1/2}^{\text{spread}}$ (kPa)</td>
<td>5.1</td>
<td>2.4</td>
</tr>
</tbody>
</table>
ies of adhesion, inhibit branching. This is achieved by adhesively confining cells. After 2 d on collagen-micropatterned PA gels, C2C12 skeletal myoblasts recognize and align on the collagen pattern and begin to fuse into nascent myotubes. This occurs regardless of substrate elastic modulus and is sustainable in culture for weeks. Unbranched, isolated myotubes are consistently >90% of cells throughout the lifetime of all micropatterned cultures. Mouse primary cells (C57 derived) exhibit a similar behavior. (B and C) Several weeks after plating myoblasts on collagen-coated PA gels of varied stiffness or rigid glass, cells were stained for myosin (green), actin (not depicted), and the nucleus (blue) in order to assess cytoskeletal expression and organization. All substrates supported fusion into multi-nucleated myotubes, but after 2 or 4 wk only myotubes on gels of intermediate stiffness showed significant myosin striation. (D) On rigid glass, nascent myotubes showed robust arrays of stress fibers (F-actin is red) and tight substrate adhesion from vinculin by 1 wk and, as extrapolated from the PA gel results, myotubes showed no tendency to striate, even by 4 wk. Note the precision of the end-patterning on glass (see Materials and methods). Bars: (A) 100 μm; (B–D) 20 μm.

Figure 4. Myotube striation is substrate stiffness dependent, whereas myocyte patterning is not. (A) By 2 d in culture, myoblasts align on the collagen pattern and begin to fuse into nascent myotubes. This occurs regardless of substrate elastic modulus and is sustainable in culture for weeks. Unbranched, isolated myotubes are consistently >90% of cells throughout the lifetime of all micropatterned cultures. Mouse primary cells (C57 derived) exhibit a similar behavior. (B and C) Several weeks after plating myoblasts on collagen-coated PA gels of varied stiffness or rigid glass, cells were stained for myosin (green), actin (not depicted), and the nucleus (blue) in order to assess cytoskeletal expression and organization. All substrates supported fusion into multi-nucleated myotubes, but after 2 or 4 wk only myotubes on gels of intermediate stiffness showed significant myosin striation. (D) On rigid glass, nascent myotubes showed robust arrays of stress fibers (F-actin is red) and tight substrate adhesion from vinculin by 1 wk and, as extrapolated from the PA gel results, myotubes showed no tendency to striate, even by 4 wk. Note the precision of the end-patterning on glass (see Materials and methods). Bars: (A) 100 μm; (B–D) 20 μm.

Striated myotubes on tissue-like stiffness
Fusion into nascent myotubes is a first, common path in differentiation of skeletal muscle, and to assess the progression of myotube differentiation, C2C12 cells grown for 2 and 4 wk on substrates of varying stiffness were stained for myosin (Fig. 4, B and C) as well as F-actin (not depicted). The multi-nucleated myotubes on both very soft gels (<5 kPa) and very stiff gels or glass show only diffuse actin and myosin staining after 2 wk in culture (n = 35 cells imaged). However, cells on intermediate stiffness (8 kPa and 11 kPa ~E_{(tissue)}) show actin and myosin colocalized as striations in up to 25% of the cell populations at 2 wk (Fig. 5; n = 27). After 4 wk in culture, striated myotubes double in percentage on intermediate stiffness gels. On gels with elastic moduli of 6.5 and 17 kPa, myotubes consistently show a signifi-
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...cant but lower percentage of striated cells. In contrast, very soft or very stiff gels (i.e., 1 and 23 kPa) show no more than a small subpopulation of striated myotubes (<5%) with poor striations and predominantly diffuse premyofibril myosin. Gaussian fits of the differentiation results clearly identify an optimal modulus of \( E^* \sim 12 \text{kPa} \) that maximizes myosin striations in muscle.

\( E^* \) is both well-defined and nearly constant from 2 to 4 wk. The optimum curves are also relatively narrow with a SD of 2–3 kPa. This implies of course that a majority of striated cells (63% for the Gaussian fits) are found on substrates with stiffness within just 25% of \( E^* \). The fact that \( E^* = E_{\text{Muscle}} \) is discussed below.

Cells grown on patterned glass substrates were stained after 1 and 4 wk for myosin and F-actin (Fig. 4 D), showing little to no striation (Fig. 5, circles). Stress fibers and vinculin-enriched focal adhesions are instead very prominent in these cells by 1 wk on glass. This is consistent with stress fiber assembly in cells on very stiff gels and in cell types ranging from fibroblasts (Beningo et al., 2001) to SMCs (Gaudet et al., 2003; Engler et al., 2004a). Stress fiber disassembly was seen later, but even by 4 wk on glass striations still did not appear.

**Cell-on-cell differentiation**

The initial apparent elasticity of C2C12 myotubes has been reported to be \( E_{C2C12} = 12-15 \text{kPa} \) (Collinsworth et al., 2002), which closely corresponds to the optimal gel stiffness \( E^* \) for striation. It also closely corresponds to normal skeletal muscle tissue stiffness, \( E_{\text{Muscle}} \). An initial layer of myoblasts was therefore plated on patterned glass surfaces to mimic \( E_{\text{Muscle}} \) with an underlayer of C2C12 cells. After 1–2 d, a second plating of myoblasts was made in order to generate a cell-on-cell arrangement of myotubes (Fig. 6 A). The initial, bottom cell (in red) has well-defined ends and appears locked by stress fibers and vinculin-enriched focal adhesions onto the interpenetrating polymer network (IPN) pattern (per Fig. 4 D). The second, top cell layer (with green vertical stripes) spreads over the exposed apical surface of the bottom, patterned cells. With time the top myotube striates, whereas the bottom myotube does not (Fig. 6, B and C).

Compared with cells of comparable age growing on gels of similar stiffness to that of the bottom layer of undifferentiated cells (i.e., gels \( \sim E_{\text{Muscle}} \)), the top cells in these patterned...
cell-on-cell arrangements exhibited an approximate twofold higher percentage of myosin striation. After 4 wk, the percentage of myosin-striated myotubes in the top layer approached 100% (Fig. 6 D). In contrast, cells in the bottom layer showed little indication of striation, despite their intimate contact with a soft and stably differentiated cell above.

Fibroblasts (FC7) are much softer than C2C12 cells with an AFM-determined elasticity of $E_{\text{Fibroblast}} = 2 - 5$ kPa (Wu et al., 1998; Bushell et al., 1999; Rotsch and Radmacher, 2000; Mahaffy et al., 2004). FC7’s were also used to create a first, patterned underlayer of cells (Fig. 7 A). Consistent with the softness of fibroblasts, myotubes grown on FC7’s appear unable to striate myosin (Fig. 7 B).

**Adhesion increases with substrate stiffness**

Cell adhesion after 4 wk in culture was studied on various substrates. Isolated myotubes were controllably peeled by a method of micropipette aspiration (Fig. 8, A and B) that is well-suited to elucidating adhesion along the length of single cells (Griffin et al., 2004). As individual myotubes are peeled, molecular adhesions break and limit the rate of peeling along the length of the myotubes (Gallant et al., 2002; Griffin et al., 2004).

Although cell peeling velocity, $v_{\text{peel}}$, fluctuates with the imposed peeling tension, $T$, it is possible to separate out the effects of focal adhesion clusters for which $v_{\text{peel}} \sim 0$ (Griffin et al., 2004). Thresholding $v_{\text{peel}}$ measurements to include
only those points where cells are most rapidly detaching from the surface gives relatively smooth peeling curves (Fig. 8 C). Clearly, sufficient tension on the cell, regardless of the substrate, will increase the disruption rate of receptor-matrix bonds. Fig. 8 C shows, nonetheless, that \( \nu _{\text{peel}} \) versus \( T \) curves are increasingly shifted to the right as substrate stiffness increases.

Cell peeling curves from each stiffness are consistently of a mathematical form (Griffin et al., 2004) \( \nu _{\text{peel}} = a \log T + b \). Evaluating each curve fit at \( \nu _{\text{peel}} = 0 \) gives a zero-velocity tension \( T_v = \exp(-b/a) \) that is the minimum tension required to detach a cell from its substrate. \( T_v \) also represents the maximum contractile prestress sustainable by the cells on a given substrate. Fig. 8 D shows that \( T_v \) increases monotonically as a function of substrate \( E \), mirroring the rightward shifts to tighter adhesion with increasing substrate stiffness.

**Discussion**

Muscle differentiation in cell culture has been described for many years. However, the results here may motivate a careful reexamination of what matrix or microenvironmental compliance has been used in the past and/or whether differentiated myotubes were growing on top of less differentiated myotubes in a more three-dimensional arrangement (Figs. 6 and 7). Recent efforts with muscle tissue constructs for implantation probably fit the latter description (Borschel et al., 2004). Micropatterning used here keeps cells well separated and well defined, leading to this first set of observations of how cell differentiation, specifically myofibrillogenesis, proceeds only within a small range of microenvironmental stiffness (Fig. 5). The results also appear the first to show that membrane adhesion strength increases exponentially with substrate stiffness (Fig. 8 D).

**Tissue, matrix, and cell stiffness**

Muscle tissue is compliant as well as contractile, and many types of muscle passively resist extension with effective or average moduli of \( E \sim 5–15 \text{ kPa} \) at strains up to 10–30% (Fung, 1993). The transverse elasticity \( E_{\text{Muscle}} = 12 \text{ kPa} \) here of C57 EDL muscle from AFM indentation fits within the published range of similar measurements on cultured myotubes, \( E_{\text{C2C12}} \) (Collinsworth et al., 2002). Elson and coworkers (Pasternak et al., 1995) used a glass microprobe \( (R_p = 1 \mu\text{m}) \) to “poke” myotubes grown from C57-derived satellite cells and reported an average indentation force \( f = 12.3 \pm 0.04 \text{ nN per } \delta = 1 \mu\text{m} \); assuming a Hertz indentation model for an incompressible solid, this early method of myotube poking yields \( E_{\text{poke}} = f/(R_p^{3/2} \delta^{1/2}) = 12.3 \text{ kPa} \). Although this is exactly as reported here for \( E_{\text{Muscle}} \) the same method for partially differentiated (fused but not striated) \( mdx \)-isolated satellite cells gave \( E_{\text{poke-mdx}} \sim 3.4 \text{ kPa} \). This is significantly softer than the \( E_{\text{mdx}} \) found here by AFM, and does not represent the large heterogeneity found in vivo. The difference may reflect compensatory mechanisms in vivo such as an \( \sim 10\)-fold overexpression of filamin C at the sarcolemma in fully differentiated dystrophic muscle compared with normal sarcolemma (Thompson et al., 2000).

The passive elasticity of adherent muscle cells appears set by their contractility (Wang et al., 2002) because inhibition of myosin activity (with BDM) can soften myotubes up to 10-fold in 30 min (Collinsworth et al., 2002). Basal contractility differences might also explain the difference between \( mdx \) tissue and cell culture. Consistent with an important role for myosin, soft fibroblasts with \( E_{\text{fibroblasts}} \sim 5 \text{ kPa} \), show much less myosin compared with myotubes (Fig. 7 B). Isolated adherent myotubes grown on top of fibroblasts seem to sense this softness and thus lack striations.

The important finding that \( E_{\text{mdx}} \), \( E_{\text{C2C12}} \), and \( E_{\text{poke}} \) are all close to the optimum \( E^* \) for normal myotube differentiation (i.e., myofibrillogenesis) on compliant substrates reinforces the conclusion that substrate stiffness is a common critical factor in differentiation to striated muscle. Moreover, our finding that \( E_{\text{mdx}} \) from excised EDL muscle is significantly greater than \( E_{\text{Muscle}} \) is consistent with past findings on diaphragm (Stedman et al., 1991) that suggested fibrotic stiffening mechanisms. An increased stiffness is further suggestive of a pathologic impact of microenvironmental elasticity (be it a synthetic or natural ECM) on muscle differentiation.

Well before striation, even myoblast spreading is found to be responsive to substrate compliance, with a characteristic for greatest responsiveness of \( E_{1/2\text{-spread}} = 2–5 \text{ kPa} \). This is generally less than \( E_{\text{Muscle}} \) and is suggestive of a genuine myoblast-myotube distinction in substrate sensitivity. Like myoblasts, SMCs have been found to exhibit similar stiffness-dependent spreading, \( E_{1/2\text{-spread}} = 8–10 \text{ kPa} \) (Engler et al., 2004a), near the elasticity of the SMC-rich arterial media, \( E_{\text{media}} = 5–8 \text{ kPa} \) (Engler et al., 2004b). Stem cells exhibit similar sensitivity to substrate elasticity in spreading and morphology (Fig. S1, available at http://www.jcb.org/cgi/content/full/jcb.200405004/DC1). Paxillin phosphorylation, for example, occurs in fibroblasts plated on stiff substrates for which spreading has been shown to be myosin dependent (Pelham and Wang, 1997). Recent work with gadolinium, a calcium channel inhibitor, has suggested that calcium entry through the plasma membrane strongly influences cell traction, making calcium a key second messenger in substrate sensing (Munevar et al., 2004). The results collectively suggest that, by whatever molecular mechanism, the more an attached cell pulls on its substrate without being able to deform the substrate, the more the cell and its molecules deform rather than the substrate.

**Myofibrillogenesis, membrane adhesion, and mechanosensing**

Myofibrillogenesis is a protracted process that begins at the membrane. In a nascent myotube, \( \alpha\)-actinin rich “Z-bodies” assemble with F-actin and nonmuscle myosin II in cross-linked premyofibrils (McKeena et al., 1986; Sanger et al., 2002). Premyofibrils occur along the sides and at the ends of the myotube, spanning vinculin-rich adhesive contacts with the substrate (Terai et al., 1989). Recruitment of muscle myosin II filaments and titin converts the premyofibrils into “nascent” myofibrils that lose nonmuscle myosin II and generate contractile tension, prestress, or force, \( f \). With sufficient tension, mature myofibrils assemble from nascent myofibrils as the small Z-bodies laterally fuse and align into Z-bands. Inhibitors of muscle myosin II, including BDM (and BTS, a skeletal muscle myosin inhibitor), clearly block late stage myofibril bundling and so \( f \) clearly stimulates myo-
substrates by myosin II would also tend to forcibly dissociate key interactions such as those between N-RAP and its binding partners; these interactions would otherwise be critical to extending N-RAP as a conformationally active mechanoreceptor. Such a pathway might be stimulated by phorbol esters, which have long been known to cause sarcomere disassembly without loss of vinculin- and talin-rich focal adhesions (Lin et al., 1989). Integrin ligation activates PKC in muscle cells as do phorbol esters, with FAK activation downstream of these contributing to cell spreading and survival (Disatnik and Rando, 1999). FAK regulation could also occur by calcium-dependent mechanisms (Lo et al., 2000) or stretch-activation (Nakamura et al., 2001; Munevar et al., 2004) that should also be sensitive to substrate stiffness (Wang et al., 2002). Strong and stable focal adhesions that are found for many cell types only on the most rigid matrices (Pelham and Wang, 1997; Engler et al., 2004a) and also found here (Fig. 4D) might activate the PKC pathway(s) far more strongly than is conducive for myofibrillogenesis (Fig. 9, rigid matrix).

In the absence of stiff ECM resistance, however, assembly and phosphorylation of adhesion proteins are inhibited (Pelham and Wang, 1997), and thus F-actin assembly is reduced (Engler et al., 2004a). Thus, on extremely soft substrates (1–5 kPa), it can only be observed from the studies here that cell attachment is extremely weak (Fig. 8) and, of course, striation is absent. Poor adhesion, generating only a small adhesion force, is consistent with reports of reduced mechanical loading inducing decreased myogenic expression (Torgan et al., 2000) and with other cell types grown on soft gels (Pelham and Wang, 1997; Beningo et al., 2001; Engler et al., 2004a) that prefer moderately stiff mechanical environments in vivo. Given the mechanosensitive down-regulation of focal and cytoskeletal proteins observed on soft gels, we speculate that nucleation of premyofibrils requires more mature and suitably stable adhesion structures and cytoskeletal filaments that are signaled by normal muscle stiffness (Fig. 9, soft matrix).

A first illustrative estimate of the force needed for myofibrillogenesis might be made from the minimum tension, $T_o$, needed to peel myotubes off of the gels of tissue-like stiffness (Fig. 8). The maturation force per premyofibril, $f^*$, when suitably calculated across the cell width $w$ should not exceed $T_o$ otherwise the cell will self-peel. If $N$ is the average number of premyofibrils across $w$, then $f^* < T_o \times w/N$. Using the focal adhesion staining in Fig. 4D to estimate $N = 10$, $w = 20$ µm, and from Fig. 8D $T_o = 1–2$ nN/µm, the optimal force for differentiation on an $E^*$ substrate is $f^* \leq 1$ nN. This is much less than past estimates of cell forces on glass (~5–10 nN), and suggests a need for lability in premyofibril formation and myogenetic progression (Fig. 9).

**Conclusion**

Muscle has well-recognized active and passive components, and so it is not surprising that striation requires more than cell anchorage. The studies here show that differentiation in culture depends intimately on optimal outside-in signaling of matrix elasticity. Most surprising, however, is the narrowness of the optimum substrate compliance, which has important implications for muscle disease and stiffened as well as softened tissue. This is particularly relevant to attempts to introduce stem cells into dystrophic skeletal muscle or infarcted regions of myocar-
dium (Mueller et al., 2002; De Bari et al., 2003; Kobinger et al., 2003). Our results suggest that both dystrophic and infarct processes induce tissue stiffening, implanted stem cells are unlikely to fully differentiate into contractile muscle. Indeed, stem cells injected into fibrotic, dystrophic skeletal muscle have been shown to differentiate into connective tissue rather than muscle (Huard et al., 2003), which could be a function of mechanical signaling as well as well-accepted cytokine signaling to the cells. On the other hand, injection of stem cells into an ever-stiffening infarct zone of the heart resulted in the cells beginning to express myocyte markers, but stopping differentiation short of myofibrillogenesis, highly analogous to our cells on a stiff matrix (Shake et al., 2002). Instruction of stem cells for tissue repair will undoubtedly require an understanding of the signaling pathways involved. Our results suggest that it may also require modification of tissue compliance if tissue repair is ultimately going to be successful.

### Materials and methods

#### AFM on skeletal muscle tissue samples
Left and right whole EDL muscles were excised from C57 normal (3, 13, and 21 mo old) and mdx dystrophic mice (3, 9, and 15 mo old). Muscle was carefully divided into thinner bundles with a razor and wet mounted onto covergloss. Tissue section thickness was confirmed by light microscopy to be ~100 μm, and samples were kept in DME plus antibiotics. Samples were probed within 8 h of dissection to minimize rigor. Samples were placed on an Asylum 1-D AFM (Asylum Research) and indented perpendicular to the myocyte axis by a pyramid-tipped probe (Veeco) with a thermally determined spring constant, $k_p = 60$ pN/nm. Force-indentation profiles were obtained at >10 locations of each muscle section, and each profile was fit up to a 2-D Hertz model with a Hertz cone model (Domke and Radmacher, 1998; Rotsch et al., 1999) to determine the elastic modulus. Histograms of all elastic moduli were plotted, and peak moduli were reported.

#### Cell culture
C2C12 cells (<10 passage; American Type Culture Collection) were maintained in polystyrene flasks in a 37°C, 5% CO2 incubator, and cultured in growth media: DME, supplemented with 20% of FBS, 0.5% chicken embryo extract, plus antibiotics. 1–2 d after plating on various substrates detailed below, cell media was switched to differentiation media, containing DME with 10% horse serum, 1% rat brain extract (Portier et al., 1999), and antibiotics. FC7 primary human dermal fibroblasts were derived from a normal 21-yr-old female donor (anonymouse) and used between passage numbers 6 and 10. FC7 human fibroblasts were cultured in identical media conditions as C2C12 cells. Mesenchymal stem cells used in the supplemental information section were obtained from Osiris Therapeutics and cultured in low glucose DME with 10% FBS.

Cell multilayers were created by plating an initial layer of cells on patterned glass substrates detailed below. A subsequent layer of C2C12 cells was plated on top of the initial layer 1–2 d after the initial plating and the media was switched to differentiation media 1–2 d later. The bottom cell layer consisted of either C2C12 cells or FC7 fibroblasts cultured in identical conditions to C2C12 cells.

Cell culture products were purchased from Invitrogen Life Technologies, whereas all other products were purchased from Sigma-Aldrich unless noted. Statistical image analyses were performed with Scion Image and displayed as cells averages (±SD) with paired t tests to determine significance. Live cell experiments were performed for no more than 1 h at RT, with frequent media renewal.

#### Gel substrates
Cells were plated on PA gel substrates of variable stiffness and observed for up to 4 wk. PA gels were made with a micropatterned glass stamp (Fig. 1 B). A mask with 100-μm-wide reflective lines spaced 20 μm apart was designed and applied to a photoresist-coated glass slide. The photoresist (Shipley) was excited by 365 nm light, binding it to the glass, and then baked to ensure attachment. When placed in an HF glass-etch for 10 min, glass in the 20-μm spaces without photoresist was removed, whereas photoresist-protected glass was not etched, thus giving 20-μm-wide grooves spaced 100 μm apart along the length of the glass (Fig. 1 B'). Residual photoresist was then removed by acetone. Collagen solution (1 mg/ml) was drawn into the grooves in this glass “stamp”, and the stamp was inverted onto the PA gel in the presence of the sulfo-SANPAH cross-linker. The sample was kept at pH 8.5 in a 37°C incubator overnight to allow the collagen to react with the gel. Due to the confinement of collagen, collagen bound solely to PA in the grooved areas of the stamp (Fig. 1, B, C, and C'), thus promoting cell attachment only within these 20-μm strips (Fig. 1, C' and D).

#### Mechanics of gels
The elastic modulus of various gels was determined by AFM as well as macroscopic methods (Dimitriadi et al., 2002; Engler et al., 2004a). AFM measurements were averaged over several trials and multiple locations per gel; collagen attachment had no detectable effect.

#### Glass micropatterns
Cells were also grown on glass, patterned by an IPN to produce isolated myotubes on glass (Griffin et al., 2004). In brief, clean glass was patterned with phototoughs and the exposed glass was bonded to allyltrimethoxysilane. The photoresist was then stripped by acetone and the allyltrimethoxysilane-covered glass was bound to the IPN. The remaining clean glass was bonded to 3-aminopropyltriethoxysilane and a layer of collagen. The final pattern was a series of adhesives collagen-adsorbed glass strips, 20-μm wide and 200–1,000-μm long, surrounded by the antithetical IPN.

#### Myoblast spreading and alignment
Myoblasts were first observed on unpatterned PA gel of various stiffnesses for initial spreading and alignment tendencies (as a ratio of the cell’s major axis to its minor axis equals “elongation ratio”) after 4 and 24 h in culture. An orientational correlation function (OCF, Eq. 2) between the reference and incident cell (Bates and Frenkel, 2000), was used to assess the spontaneous alignment of unpatterned cells.

#### Immunofluorescence
Patterned cells after 1, 2, and/or 4 wk were labeled with rhodamine-phalloidin, Hoechst 33342 nuclear stain (Molecular Probes), and immunofluorescent antibodies to myosin (Zymed Laboratories) or vinculin. Cells were fixed with formaldehyde and blocked in 5% BSA for 1 h at 37°C. Patterned cells were permeabilized with 0.5% Triton X-100 (Fisher Scientific) and incubated at 4°C for 8–10 h in antomyosin (Zymed Laboratories) or vinculin, diluted 1:100 in PBS. Cells with nonspecific binding properties of N,N,N’,N’-tetramethylethylenediamine (1/20,000 vol). 25 μl of the polymerizing solution was placed on an aminosilanized glass slide or coverslip, or covalently bound the gel samples to a rigid support; coverslips coated with dichlorodimethylsilane were placed on top to ensure a smooth gel surface and easy detachment for the final PA gel bound the aminosilanized coverslip. When polymerized, PA gels were ~70 μm thick as confirmed by microscopy. Their elastic modulus was set by the acrylamide and bis-acrylamide concentrations (Engler et al., 2004a).

#### Micropipette aspiration
Isolated myotubes were controllably peeled from gel and IPN-coated glass substrates to determine the dependence of minimal peeling tension on

\[
O C F = 1/2 [\cos(2 \theta) + 1] \tag{2}
\]
substrate stiffness. In brief, a fluid shear stress with Poiseuille pipe flow was created in a 75-μm-aspirating pipette by a syringe pump with known flow rates (Griffin et al., 2004). When positioned over a detached cell end, the local shear force induced cell peeling at the leading edge of a patterned myotube, aspirating the cell into the pipette. Peeling tension was calculated from the shear stress integrated along the length of the cell in the pipette, given corrections for pipette entrance and gel displacement effects. The resultant peeling velocities from these experiments were very heterogeneous due to randomly distributed focal adhesion within numerous cells tested; thus velocity-tension plots were only made using binned averages of active peeling (and not focal adhesion), defined as a velocity at least 50% of the previous point.

Online supplemental material

Fig. S1 shows that mesenchymal stem cells exhibit myoblast morphology on muscle-like stiffness. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200405004/DC1.

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