Distinct mechanisms regulating mechanical force-induced Ca\(^{2+}\) signals at the plasma membrane and the ER in human MSCs

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Abstract It is unclear how subcellular organelles respond to external mechanical stimuli. Here, we investigated the molecular mechanisms by which mechanical force regulates Ca\(^{2+}\) signaling at the endoplasmic reticulum (ER) in human mesenchymal stem cells. Without extracellular Ca\(^{2+}\), ER Ca\(^{2+}\) release is the source of intracellular Ca\(^{2+}\) oscillations induced by laser-tweezer-traction at the plasma membrane, providing a model to study how mechanical stimuli can be transmitted deep inside the cell body. This ER Ca\(^{2+}\) release upon mechanical stimulation is mediated not only by the mechanical support of cytoskeleton and actomyosin contractility, but also by mechanosensitive Ca\(^{2+}\) permeable channels on the plasma membrane, specifically TRPM7. However, Ca\(^{2+}\) influx at the plasma membrane via mechanosensitive Ca\(^{2+}\) permeable channels is only mediated by the passive cytoskeletal structure but not active actomyosin contractility. Thus, active actomyosin contractility is essential for the response of ER to the external mechanical stimuli, distinct from the mechanical regulation at the plasma membrane.

Introduction Mechanical factors are known to play crucial roles in both development and tissue regeneration from stem cells. However, it remains unclear how these factors, such as mechanical forces, are converted...
eLife digest Cells receive many signals from their environment, for example, when they are compressed or pulled about by neighboring cells. Information about these ‘mechanical stimuli’ can be transmitted within the cell to trigger changes in gene expression and cell behavior.

When a cell receives a mechanical stimulus, it can activate the release of calcium ions from storage compartments within the cell, including from a compartment called the endoplasmic reticulum. Calcium ions can also enter the cell from outside via channels located in the membrane that surrounds the cell (the plasma membrane).

Kim et al. investigated how mechanical forces are transmitted in a type of human cell called mesenchymal stem cells using optical tweezers to apply a gentle force to the outside of a cell. These tweezers use a laser to attract tiny objects, in this case a bead attached to proteins in the cell’s outer membrane. The cell’s response to this mechanical stimulation was measured using a sensor protein that fluoresces a different color when it binds to calcium ions.

With this set-up, Kim et al. found that mesenchymal stem cells are able to transmit mechanical forces to different depths within the cell. The forces can travel deep to trigger the release of calcium ions from the endoplasmic reticulum. This process involves a network of protein fibers that criss-cross to support the structure of a cell—called the cytoskeleton—and also requires proteins that are associated with the cytoskeleton to contract. However, calcium ion entry through the plasma membrane due to a mechanical force does not require these contractile proteins—only the cytoskeleton is involved.

These results demonstrate that the transmission of mechanical signals to different depths within mesenchymal stem cells involves different components. Future work should shed light on how these mechanical signals control gene expression and the development of mesenchymal stem cells.

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Results and discussion

We first investigated how force regulates Ca\textsuperscript{2+} release from ER with the FRET-based Ca\textsuperscript{2+} biosensor (Figure 1A) to dissect the effect of mechanical force on each of the two Ca\textsuperscript{2+} mobilization processes (Figure 1B). The experiments were done in the absence of extracellular Ca\textsuperscript{2+} so that there was no Ca\textsuperscript{2+}...
influx through plasma membrane and the oscillations were mainly from ER release (Figure 1B). When fibronectin (Fn)-coated beads were seeded onto the HMSCs and 300 pN of mechanical force was applied by optical laser tweezers as described previously (Wang et al., 2005), Ca²⁺ oscillations were induced in HMSCs (Figure 1C-F, Video 1) but not in bovine aortic endothelial cells (BAECs) (Figure 1—figure supplement 1). In contrast, laser-tweezer-traction on bovine serum albumin (BSA)-coated beads...
Two possible mechanisms can regulate Ca\(^{2+}\) release from ER upon mechanical force application. External force can either 1) transmit deep inside the cell and mechanically alter the channels on ER for Ca\(^{2+}\) release (Himmel et al., 1993; Missiaen et al., 1996; Rath et al., 2010); or 2) trigger biochemical signaling cascades to produce IP\(_3\) that diffuses inside to activate IP\(_3\)-sensitive Ca\(^{2+}\) channels (Lehtonen and Kinnunen, 1995; Matsumoto et al., 1995). To distinguish them, we directly monitored the IP\(_3\) level using a FRET-based biosensor, LIBRAvIIs. Mechanical force did not induce any change of IP\(_3\) while ATP increased IP\(_3\) production in HMSCs (Figure 1—figure supplement 2C), suggesting that the second mechanism is unlikely. Therefore, laser-tweezers traction should transmit deep inside the cells to mechanically release Ca\(^{2+}\) from ER.

Cytoskeleton is known to transmit mechanical forces and conduct mechanotransduction (Hamill and Martinac, 2001; Orr et al., 2006; Schwartz and DeSimone, 2008), so we investigated the role of cytoskeleton and its associated proteins in the force-induced ER Ca\(^{2+}\) release. The disruption of cytoskeletal actin filaments by cytochalasin D (Cyto D) or microtubules by nocodazole (Noc) completely eliminated the force-induced Ca\(^{2+}\) oscillations (Figure 2A–B). In addition, the inhibition of actomyosin contractility by ML-7 or blebbistatin had the same effect (Figure 2C–D). Thus, the deep penetration and transmission of force inside the cell and the induction of Ca\(^{2+}\) release from ER depend on both cytoskeletal support and actomyosin contractility. This matches with the previous reports that long-distance force propagation to the deep cytoplasm depends on cytoskeleton tension (Hu et al., 2003) as well as the pivotal role of myosin light chain kinase (MLCK) and myosin II in regulating force development (Olsson et al., 2004; Herring et al., 2006; Fajmut and Brumen, 2008; He et al., 2008). There are two possible signals in ER that may trigger the Ca\(^{2+}\) development (Himmel et al., 2004). In addition, the inhibition of IP\(_3\) receptors by IP\(_3\)R channel blockers such as 2-Amino-ethoxydiphenylborate (2-APB) entirely abolished the force-induced oscillations (Figure 1—figure supplement 2A,B). These results confirmed that the ER Ca\(^{2+}\) store is the main source for the force-induced oscillations in HMSCs without extracellular Ca\(^{2+}\).

As some TRP channels have been shown to be mechanosensitive and have linkage to cytoskeleton

**Video 1.** A HMSC transfected with cytosolic Ca\(^{2+}\) biosensors before and after mechanical force application by optical laser tweezers on a Fn-coated bead attached to the cell (Duration of Video: 2700 s). DOI: 10.7554/eLife.04876.006
Figure 2. Cytoskeletal support, actomyosin contractility, and TRPM7 channels mediate the force-induced intracellular Ca\(^{2+}\) oscillations. The time courses represent the YPet/ECFP emission ratio of cytoplasmic Ca\(^{2+}\) in HMSCs in the absence of extracellular Ca\(^{2+}\) when these cells were pretreated with (A) 2 μM Cyto D (n = 8), (B) 1 μM Noc (n = 8), (C) 5 μM ML-7 (n = 8), and (D) 5 μM Bleb (n = 8). (E) Color images represent the YPet/ECFP emission ratio of the cytoplasmic Ca\(^{2+}\) biosensor in HMSCs transfected with NT or TRPM7 siRNA. The color scale bars represent the range of emission ratio, with cold and hot colors indicating low and high levels of Ca\(^{2+}\) concentration, respectively. (F) The time courses represent the YPet/ECFP emission ratio averaged over the cell bodies outside of nucleus treated with siRNA as indicated.

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The following figure supplements are available for figure 2:

Figure supplement 1. Stretch-activated or store-operated channels at the plasma membrane mediate force-induced Ca\(^{2+}\) release from ER.

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Figure supplement 2. TRPM7 channels as well as cytoskeletal support and actomyosin contractility mediate the force-induced intracellular Ca\(^{2+}\) oscillations.

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(Barritt and Rychkov, 2005), it is likely that TRP channels located at ER may mediate, at least in part, the force-induced ER calcium release. Notably, these two possibilities are not mutually exclusive as more than one type of channels can be responsible for the force-induced ER calcium release.

Surprisingly, blocking stretch-activated or store-operated channels at the plasma membrane by Gd³⁺, La³⁺, or streptomycin, but not L-type voltage-operated Ca²⁺ channels by Nifedipine, abolished the mechanical force-induced Ca²⁺ release from ER (Figure 2—figure supplement 1). These results suggest a possible coupling between force-induced Ca²⁺ release at ER and stretch-activated and store-operated channels at the plasma membrane. As TRPM7 is one of the major Ca²⁺ permeable mechanosensitive channels (Wei et al., 2009), we knocked down TRPM7 with targeting small interfering RNA (siRNA) to examine its role in the force-induced Ca²⁺ oscillations. The decreased expression of TRPM7 and the lower percentile of HMSCs with Ca²⁺ oscillations confirmed the effect of siRNA (Figure 2—figure supplement 2A,B). TRPM7 siRNA further abrogated the force-induced oscillations (Figure 2E–F). It is intriguing that the inhibition of TRPM7 at the plasma membrane can block the force transmission into ER in regulating calcium signals. Several possibilities may contribute to the observed results. 1) TRPM7 is functionally coupled to integrin, actomyocin contractility and cytoskeleton. As such, it may mediate and facilitate the force transmission to ER. Indeed, it has been shown that TRPM7 kinase can phosphorylate myosin II heavy chain (Clark et al., 2006) and regulate actomyocin contractility. 2) TRPM7 activity may have some downstream effect on IP₃R in ER. For example, TRPM7 can control the protease calpain (Su et al., 2006), which can regulate IP₃R degradation in ER (Diaz and Bourguignon, 2000). 3) TRPM7 may also be directly coupled to IP₃R in the ER through adaptor proteins. Indeed, another TRP channel, TRPC1 has been shown to directly couple to IP₃R in the ER through an adaptor protein Homer (Yuan et al., 2003). It becomes clear that membrane channels are not isolated entities floating in the plasma membrane. Instead, they are intimately coupled to integrins, cytoskeleton, actomyocin contractility, and ER membrane channels (Cantiello et al., 2007; Matthews et al., 2007; Deng et al., 2009). Therefore, these structural and physical couplings enable membrane channels to participate in direct force transmission to ER.

We further visualized the Ca²⁺ release from ER directly, by generating and employing an improved ER-targeting Ca²⁺ biosensor (D3ER) (Palmer et al., 2006; Ouyang et al., 2008). Mechanical force could directly induce the Ca²⁺ release from ER without extracellular Ca²⁺, as evidenced by the decrease in ER Ca²⁺ concentration (Figure 3A-B, Video 2). The measurements of direct Ca²⁺ release from ER in the presence of inhibitors or siRNA of TRMP7 were also consistent with our observations when cytosolic Ca²⁺ biosensor was used (Figure 3C).

To gain more insights, we then examined the effect of force on Ca²⁺ influx at the plasma membrane. ER Ca²⁺ release was blocked by an IP₃Rs inhibitor, 2-APB, in the presence of extracellular Ca²⁺ so that the only source of intracellular Ca²⁺ change is the calcium influx from extracellular space (Bootman et al., 2002). Nifedipine did not have any effect on the force-induced Ca²⁺ influx, while Gd³⁺, La³⁺, streptomycin, and TRPM7 siRNA significantly inhibited this Ca²⁺ influx (Figure 4A–B), confirming the involvement of mechanosensitive channels, especially TRPM7, which was shown to be directly activated by a membrane stretch or shear stress in various cell types (Numata et al., 2007a, 2007b; Wei et al., 2009). The disruption of actin filaments or microtubules also inhibited the force-induced Ca²⁺ influx (Figure 4C–D), suggesting that cytoskeletal integrity is essential for the membrane channels to respond to force, consistent with previous reports (Hayakawa et al., 2008). Interestingly, ML-7 or blebbistatin did not affect this force-induced Ca²⁺ influx, suggesting that active actomyosin contractility may not be involved in the mechno-regulation of membrane channels (Figure 4C–D), different from their indispensable roles in the force-induced ER Ca²⁺ release (Figure 4E).

Since mechanical stimulation such as stretch affects integrin adhesion and integrin-associated signaling, including Src and focal adhesion kinase (FAK) (Wang et al., 2005; Orr et al., 2006), we investigated whether Src and FAK mediate the force-induced Ca²⁺ signals. Our results indicated that neither PP1, an inhibitor of Src family kinases, nor PF228, an inhibitor of FAK, had a significant effect on the force-induced Ca²⁺ influx or ER release (Figure 4—figure supplement 1). Inhibition of phosphoinositide 3-kinases (PI3Ks) with LY294002 did not have any effect either (Figure 4—figure supplement 1). Although integrin-mediated Src phosphorylation and activation can regulate L-type Ca²⁺ channel functions (Gui et al., 2010), the force-induced Ca²⁺ oscillations were clearly independent of L-type Ca²⁺ channels in HMSCs (Figure 2—figure supplement 1). Mechanical stretch can induce the phosphorylation of FAK and nitric oxide formation in cardiomyocytes (Pinsky et al., 1997; Khan et al., 2003), which can modulate ryanodine receptor 2 (RYR2) and Ca²⁺ release at the
SR (Vila Petroff and Mattiazzi, 2001). However, there is minimal expression of RYRs in HMSCs (Kawano et al., 2002), which may explain the lack of FAK involvement in the mechano-regulation of Ca\textsuperscript{2+} signals. Together, the biochemical activities above have no detected involvement in the force-induced Ca\textsuperscript{2+} signals.

Further analysis of the duration between force application and the first Ca\textsuperscript{2+} signal indicated that in the absence of extracellular Ca\textsuperscript{2+}, the average delay time of ER calcium release monitored by both cytosolic Ca\textsuperscript{2+} biosensor and ER Ca\textsuperscript{2+} biosensor are around 100 s (Figure 4—figure Supplement 1C). Meanwhile, the average delay time of the calcium influx through plasma membrane in the presence of 2-APB and extracellular calcium is much shorter, around 20 s (Figure 4—figure supplement 1C), indicating faster response to mechanical force stimulation. Although the delay time of ER calcium release is longer than that of calcium influx across the plasma membrane, it does not contradict mechanical signal-mediated mechanism. Several reasons may contribute to longer time delay of ER calcium release. 1) The mechanical coupling machinery may need time for reinforcement to allow sufficient force transfer between the plasma membrane and the ER membrane. It is likely that mechanical force transmits through integrins and cytoskeleton (Wang et al., 1993; Perozo and Rees, 2003) as only Fn-coated, but not BSA-coated beads can induce ER calcium release (Figure 1C–F). However, the discrete network linkage of the existing cytoskeleton at the time of force application may not be sufficient for force focusing and transmission to ER, particularly because the apical integrins bound to the Fn-coated beads had not previously experienced force and therefore may have limited connection with cytoskeleton (Matthews et al., 2006). As force can unfold proteins, change their conformations to expose cryptic binding sites to allow the assembly of new/stronger network linkages (del Rio et al., 2009; Johnson et al., 2007), mechanical force can modulate the structural coupling of molecules and mechanical properties of the cells through cytoskeletal remodeling (Kaazempur Mofrad et al., 2005; Matthews et al., 2006; Rosenblatt et al., 2007). These modulation processes of mechanical coupling and reinforcement may need time to reach the threshold for sufficient mechanotransduction of ER calcium release; 2) the channels on ER membrane could have different kinetics.
and mechanosensitivity (Moe et al., 1998), therefore they may need larger focused and reinforced force to be developed at the site of ER to reach the threshold for physical opening. All these factors may contribute to the longer delay time for force-induced ER calcium release. Again our model does not rule out biochemical signal-mediated mechanisms. As a matter of fact, biochemical signal-mediated mechanisms, such as protein–protein interaction and cytoskeletal remodeling under mechanical tension are important to mediate mechanical signals as discussed above.

As such, our results have provided molecular insights on how mechanical force triggers intracellular Ca\(^{2+}\) oscillations through two mechanisms in HMSCs: Ca\(^{2+}\) influx at the plasma membrane and Ca\(^{2+}\) release from ER (Figure 1B). Our results showed that the deep penetration and transmission of mechanical force to regulate ER functions is dependent on not only the passive cytoskeletal support of actin filaments and microtubules, but also the active actomyosin contractility controlled by MLCK and myosin II. In contrast, the passive cytoskeletal support, but not active actomyosin contractility, is needed for the mechanotransduction at the plasma membrane levels, including the mechanoactivation of channels and Ca\(^{2+}\) influx across the plasma membrane (Figure 4E). These results hence provide direct evidence that the mechanotransduction at different depths of cell body is mediated by differential sets of mechano-sensing elements.

Materials and methods

Gene construction and DNA plasmid

The construct of FRET-based Ca\(^{2+}\) biosensor has been described well in our previous articles (Ouyang et al., 2008; Kim et al., 2009). In brief, the fragment containing enhanced cyan fluorescent protein (ECFP), calmodulins (CaMs), and M13 was fused to YPet and subcloned into pcDNA3.1 (Invitrogen, Carlsbad, CA) for mammalian cell expression by using BamHI and EcoRI sites. The ECFP/YPet pair has allowed a higher sensitivity of FRET biosensors than those based on ECFP/Citrine pair. To generate an improved ER-targeting Ca\(^{2+}\) biosensor, the mutant peptide and CaMs regions were replaced with those regions of D3cpv and cloned between a truncated ECFP and YPet. For the ER targeting motifs, the calreticulin signal sequence MLLPVLLLGLLGAAAD was added 5′ to ECFP, and an ER retention sequence KDEL to the 3′ end of YPet. The construct of a FRET-based IP3 biosensor, LIBRAvIIs was kindly provided by Professor Akihiko Tanimura at University of Hokkaido, Japan (Tanimura et al., 2009).

Cell culture and transfection

Human mesenchymal stem cells (HMSCs) and bovine aortic endothelial cells (BAECs) were obtained from the American Type Culture Collection (ATCC, Rockville, MD). HMSCs and BAECs were cultured in human mesenchymal stem cell growth medium (MSCGM, PT-3001, Lonza Walkersville, Inc., Walkersville, MD) and in Dulbecco’s modified Eagle’s medium (DMEM), respectively, supplemented with 10% fetal bovine serum (FBS), 2 mM L-glutamine, 100 U/ml penicillin, and 100 µg/ml streptomycin. The cells were cultured in a humidified incubator of 95% O\(_2\) and 5% CO\(_2\) at 37°C. The DNA plasmids were transfected into the cells by using Lipofectamine 2000 (Invitrogen, Carlsbad, CA) according to the product instructions.

RNA interference assays

Double-stranded small interfering RNA (siRNA) sequences targeting human TRPM7 (ON-TARGETplus SMARTpool siRNA) and non-targeting control sequences were designed by Dharmaco RNAi.
Technology (Dharmacon Inc., Lafayette, CO). HMSCs were transfected with 1–2 µg siRNA specific for TRPM7 or a non-silencing control sequence according to the product instructions.

**Western blotting**

The cells transfected with TRPM7 or non-targeting siRNA were washed twice with cold phosphate buffered saline (PBS) and then lysed in lysis buffer containing 50 mM Tris, pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, and a mix of serine and cysteine protease inhibitors. Lysates were centrifuged at 10,000×g at 4°C for 10 min. Cell lysates were then applied to 15% SDS-polyacrylamide gel electrophoresis, transferred to nitrocellulose, blocked with 5% non-fat milk, and detected by Western blotting using polyclonal goat anti-TRPM7 antibody (1:100; Abcam Inc., Cambridge, MA).

**Immunostaining**

A polyclonal antibody against TRPM7 was used in both normal HMSCs and TRPM7-knockdown HMSCs. After being washed in cold phosphate buffered saline (PBS), the samples were fixed by...
4% paraformaldehyde in PBS at room temperature for 15 min. The samples were incubated with a goat polyclonal antibody against TRPM7 (1:100; Abcam Inc., Cambridge, MA) at room temperature for 2 hr, followed by the incubation with TRITC-conjugated anti-goat IgG (1:100, Jackson ImmunoResearch Lab., Inc., West Grove, PA) at room temperature for 1 hr before the mounting of anti-photobleaching reagent (Vector Lab., Inc., Burlingame, CA).

**Solutions and chemicals**

Imaging experiments were conducted with Ca\(^{2+}\) free Hanks balanced salt solution (HBSS, Invitrogen) containing 20 mM HEPES, 1 mM D-glucose, 0.5 mM EGTA, 1 mM MgCl\(_2\), and 1 mM MgSO\(_4\) (pH 7.4). During imaging experiments, the solution was kept in streptomycin free condition to prevent a possible effect on mechanosensitive ion channels. The chemical reagents 2-Amino-ethoxydiphenyl borate (2-APB), Nifedipine, Thapsigargin (TG), LaCl\(_3\), GdCl\(_3\), streptomycin, nocodazole, cytochalasin D, blebbistatin, and ML-7 were purchased from Sigma–Aldrich (Sigma, St. Louis, MO). PP1 and LY294002 were commercially obtained from Calbiochem (San Diego, CA). PF228 was obtained from Tocris Bioscience (Ellisville, MO). The amount of drug administration was based on previous publications (Kim et al., 2009; Lu et al., 2011; Seong et al., 2011).

**Preparation of beads and optical laser tweezers**

A fiber-coupled IR (infra-red) laser (1064 nm, 5W, 5 mm diameter, YLD-5-1064-LP, IPG Photonics) was used for the experiment. We used a piezoelectric system for a steering mirror and the piezo-mirror system was designed with a closed loop, and an automated shutter (LS6ZM2, Uniblitz, Rochester, NY) with a shutter controller (VCM-D1, Uniblitz). Mirrors (designed for IR), lenses (BK7, plano-convex), and other basic optics were purchased from Thorlabs (Newton, New Jersey). A hot mirror (FM01, wide band, Thorlabs) was installed inside a microscope to block the IR scattering. The piezo-mirror (S-334.2, PI Physik instrumente) was installed together with the computer interface module of E-516.13, PI. The interface module set up with the other drivers (E-503.00 and E-509.53, PI) makes it possible to control the piezo-mirror system by a computer. The laser beam passes through a laser-beam expander, a steering mirror, and a dichroic long-pass beam splitter to enter the microscope side port. Beads coated with fibronectin (Fn; 50 µg/ml) or BSA as the control were prepared as previously reported (Wang et al., 2006). The size of the beads is 10 µm and the beads were incubated for 10–20 min to allow them to adhere to cell membrane surface. Single-beam gradient optical laser tweezers with controlled 300 pN of mechanical force were applied to pull the adhered beads. A similar optical trapping system has been described in our previous report (Botvinick and Wang, 2007).

**Microscopy, imaging acquisition, and analysis**

Cells expressing various exogenous proteins were starved with 0.5% FBS for 36–48 hr before imaging experiments. All images were obtained by using Zeiss Axiovert inverted microscope equipped with a charge-coupled device (CCD) camera (Cascade 512B, Photometrics) and a 420DF20 excitation filter, a 450DFRLP dichroic mirror, and two emission filters controlled by a filter changer (480DF30 for ECFP and 535DF25 for YPet). Time lapse fluorescence images were acquired at 10 s interval by MetaFluor 6.2 software (Universal Imaging, West Chester, PA). The emission ratio of YPet/ECFP was directly computed and generated by the MetaFluor software to represent the FRET efficiency before they were subjected to quantification and analysis by Excel (Microsoft, Redmond, WA).

**Statistical analysis**

The results were expressed as the mean standard error of the mean (S.E.M). Statistical analysis of the data was performed by the unpaired Student’s t-test to determine the statistical differences between the two mean values. The statistically significant level was determined by p < 0.05.

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References
Balasubramanian L, Ahmed A, Lo CM, Sham JS, Yip KP. 2007. Integrin-mediated mechanotransduction in renal vascular smooth muscle cells: activation of calcium sparks. American Journal of Physiology Regulatory, Integrative and Comparative Physiology 293:R1586–R1594. doi: 10.1152/ajpregu.00025.2007.
Barritt G, Rychkov G. 2005. TRPs as mechanosensitive channels. Nature Cell Biology 7:105–107. doi: 10.1038/ ncb0205-105.
Belmonte S, Morad M. 2008. Shear fluid-induced Ca2+ release and the role of mitochondria in rat cardiac myocytes. Annals of the New York Academy of Sciences 1123:58–63. doi: 10.1196/annals.1420.007.
Berbey C, Weiss N, Legrand C, Allard A. 2009. Transient receptor potential canonical type 1 (TRPC1) operates as a sarcoplasmic reticulum calcium leak channel in skeletal muscle. Journal of Biological Chemistry 284:36387–36394. doi: 10.1074/jbc.M109.073221.
Bentsen MW. 2007. Optical tweezers: tethers, wavelengths, and heat. Methods in Cell Biology 82:457–466.
Berridge MJ, Lipp P, Bootman MD. 2000. The versatility and universality of calcium signalling. Nature Reviews Molecular Cell Biology 1:11–21. doi: 10.1038/35036035.
Bidaux G, Flourakis M, Thebault S, Zholos A, Beck B, Gkika D, Roudbaraki M, Bonnal JL, Mauroy B, Shuba Y, Skryma R, Prevaskarya N. 2007. Prostate cell differentiation status determines transient receptor potential melastatin member 8 channel subcellular localization and function. The Journal of Clinical Investigation 117:1647–1657. doi: 10.1172/JCI30168.
Booijman MD, Collins TJ, Mackenzie L, Roderick HL, Berridge MJ, Peppiatt CM. 2002. 2-aminoethoxydiphenylborate (2-APB) is a reliable blocker of store-operated Ca2+ entry but an inconsistent inhibitor of InsP3-induced Ca2+ release. PASEB Journal 16:1145–1150. doi: 10.1096/fj.02-0037rev.
Botvinnik EL, Wang Y. 2007. Laser tweezers in the study of mechanobiology in live cells. Methods in Cell Biology 82:497–523. doi: 10.1016/S0091-679X(06)82018-2.
Bourguignon LY, Jin HT, Iida N, Brandt NR, Zhang SH. 1993. The involvement of ankyrin in the regulation of inositol 1,4,5-Trisphosphate receptor-mediated Ca2+ release from Ca2+ storage-Vesicles in Mouse t-lymphoma cells. Journal of Biological Chemistry 268:7290–7297.
Caniatto HF, Montalbetti N, Li Q, Chen XZ. 2007. The cytoskeletal connection to ion channels as a potential mechanosensory mechanism: lessons from polycystin-2 (TRPP2). Current Topics in Membranes 59:233–296. doi: 10.1016/S1063-5823(06)59010-6.
Cheng Y, Liu A, Guo Y, Qi X. 2004. Computational models for the helix tilt angle. Journal of Chemical Information and Computer Sciences 44:1952–1960.
Clapham DE. 2007. Calcium signaling. Cell 131:1047–1058. doi: 10.1016/j.cell.2007.11.028.
Clark K, Langeslag M, van Leeuwen B, Ran L, Ryazanov AG, Figdor CG, Moelenaar WH, Jalink K, van Leeuwen FN. 2006. TRPM7, a novel regulator of actomyosin contractility and cell adhesion. The EMBO Journal 25:290–301. doi: 10.1038/sj.emboj.7600931.
del Rio A, Perez-Jimenez R, Liu R, Roca-Cusachs P, Fernandez JM, Sheetz MP. 2009. Stretching single talin rod molecules activates vinculin binding. *Science* **323**:638–641. doi: 10.1126/science.116291.

Deng X, Wang Y, Zhou Y, Soboloff J, Gill DL. 2009. STIM and Orai: dynamic intermembrane coupling to control cellular calcium signals. *The Journal of Biological Chemistry* **284**:22501–22505. doi: 10.1074/jbc.R109.018655.

Diaz F, Bourguignon LY. 2000. Selective down-regulation of IP3 receptor subtypes by caspases and calcipain during TNF alpha -induced apoptosis of human T-lymphoma cells. *Cell Calcium* **27**:315–328. doi: 10.1045/caca.2000.0012.

Fajmut A, Brumen M. 2008. MLC-kinase/phosphatase control of Ca2+ signal transduction in airway smooth muscles. *Journal of Theoretical Biology* **252**:474–481. doi: 10.1016/j.jtbi.2007.10.005.

Fukatsu K, Bannai H, Zhang SB, Nakamura H, Inoue T, Mikoshiba K. 2004. Lateral diffusion of inositol 1,4,5-trisphosphate receptor type 1 is regulated by actin filaments and 4.1N in neuronal Dendrites. *The Journal of Biological Chemistry* **279**:48976–48982. doi: 10.1074/jbc.M408364200.

Gallego-Sandin S, Rodriguez-Garcia A, Alonso MT, Garcia-Sancho J. 2009. The endoplasmic reticulum of dorsal root ganglion neurons contains functional TRPV1 channels. *Journal of Biological Chemistry* **284**:32591–32601. doi: 10.1074/jbc.M109.019687.

Gui P, Chao JT, Wu X, Yang Y, Davis GE, Davis MJ. 2010. Coordinated regulation of vascular Ca2+ and K+ channels by integrin signaling. *Advances in Experimental Medicine and Biology* **674**:69–79. doi: 10.1007/978-1-4419-6066-5_7.

Hamill OP, Martincic B. 2001. Molecular basis of mechanotransduction in living cells. *Physiological Reviews* **81**:685–740.

Hayakawa K, Tatsumi H, Sokabe M. 2008. Actin stress fibers transmit and focus force to activate mechanosensitive channels. *Journal of Cell Science* **121**:496–503. doi: 10.1242/jcs.022053.

He WQ, Peng YJ, Zhang WC, Lv N, Tang J, Chen C, Zhang QH, Gao S, Chen HO, Zhi G, Feil R, Kamm KE, Stull JT, Gao X, Zhu MS. 2008. Myosin light chain kinase is central to smooth muscle contraction and required for gastrointestinal motility in mice. *Gastroenterology* **135**:610–620. doi: 10.1053/j.gastro.2008.05.032.

Herring BP, El-Mounayri O, Gallagher PJ, Yin F, Zhou J. 2006. Regulation of myosin light chain kinase and telokin expression in smooth muscle tissues. *American Journal of Physiology Cell Physiology* **291**:C817–C827. doi: 10.1152/ajpcell.00198.2006.

Himmel HM, Whorton AR, Strauss HC. 1993. Intracellular calcium, currents, and stimulus-response coupling in endothelial cells. *Hypertension* **21**:112–127. doi: 10.1161/01.HYP.21.1.112.

Horner VL, Wolfer MF. 2008. Mechanical stimulation by osmotic and hydrostatic pressure activates *Drosophila* oocytes in vitro in a calcium-dependent manner. *Developmental Biology* **316**:100–109. doi: 10.1016/j.ydbio.2008.01.014.

Hu S, Chen J, Fabry B, Numaguchi Y, Gouldstone A, Ingber DE, Fredberg JJ, Butler JP, Wang N. 2003. Intracellular stress tomography reveals stress focusing and structural anisotropy in cytoskeleton of living cells. *American Journal of Physiology Cell Physiology* **285**:C1082–C1090. doi: 10.1152/ajpcell.00159.2003.

Johnson CP, Tang HY, Carag C, Speicher DW, Discher DE. 2007. Forced unfolding of proteins within cells. *Science* **317**:663–666. doi: 10.1126/science.1139857.

Joseph SK, Samanta S. 1993. Detergent solubility of the inositol trisphosphate receptor in rat-brain membranes. Evidence for association of the receptor with ankyrin. *Journal of Biological Chemistry* **268**:6477–6486.

Kazempur Mofrad MR, Abdul-Rahim NA, Karcher H, Mack PJ, Yap B, Kamm RD. 2005. Exploring the molecular basis for mechanosensation, signal transduction, and cytoskeletal remodeling. *Acta Biomaterialia* **1**:281–293. doi: 10.1016/j.actbio.2005.02.008.

Kawano S, Shoji S, Ichinose S, Yamagata K, Tagami M, Hiraoka M. 2002. Characterization of Ca2+(+) signaling pathways in human mesenchymal stem cells. *Cell Calcium* **32**:165–174. doi: 10.1016/S0143-4160(02)001240.

Khan SA, Skaf MW, Harrison RW, Lee K, Minhas KM, Kumar A, Fradley M, Shoukas AA, Berkowitz DE, Hare JM. 2003. Nitric oxide regulation of myocardial contractility and calcium cycling: independent impact of neuronal and endothelial nitric oxide synthases. *Circulation Research* **92**:1322–1329. doi: 10.1161/01.RES.0000078171.52542.9E.

Kim T, Im W. 2010. Revisiting hydrophobic mismatch with free energy simulation studies of transmembrane helix tilt and rotation. *Biophysical Journal* **99**:175–183. doi: 10.1016/j.bpj.2010.04.015.

Kim TJ, Seong J, Ouyang M, Sun J, Lu S, Hong JP, Wang N, Wang Y. 2009. Substrate rigidity regulates Ca2+ oscillation via RhoA pathway in stem cells. *Journal of Cellular Physiology* **218**:285–293. doi: 10.1002/jcp.21598.

Koulen P, Cai YQ, Geng L, Maeda Y, Nishimura S, Witzgall R, Ehrlich BE, Somlo S. 2002. Polycystin-2 is an endoplasmic reticulum Ca2+ channel. *Journal of Theoretical Biology* **218**:1888–197. doi: 10.1038/njcb.754.

Landsberg JW, Yuan JX. 2004. Calcium and TRP channels in pulmonary vascular smooth muscle cell proliferation. *News in Physiological Sciences* **19**:44–50.

Lehtonen JY, Kinnunen PK. 1995. Phospholipase A2 as a mechanosensor. *Biophysical Journal* **68**:1888–1894. doi: 10.1016/S0006-3495(95)80366-8.

Lu S, Kim TJ, Chen CE, Ouyang M, Seong J, Liao X, Wang Y. 2011. Computational analysis of the spatiotemporal coordination of polarized PI3K and Rac1 activities in micro-patterned live cells. *PLOS ONE* **6**:e21293. doi: 10.1371/journal.pone.0021293.

Maeda Y, Nitanai Y, Oda T. 2007. From the crystal structure of troponin to the mechanism of calcium regulation of muscle contraction. *Advances in Experimental Medicine and Biology* **592**:37–46. doi: 10.1007/978-4-311-92453-3_5.

Matsumoto H, Baron CB, Coburn RF. 1995. Smooth muscle stretch-activated phospholipase C activity. *The American Journal of Physiology* **268**:C458–C465.
Matthews BD, Overby DR, Mannix R, Ingber DE. 2006. Cellular adaptation to mechanical stress: role of integrins, Rho, cytoskeletal tension and mechanosensitive ion channels. Journal of Cell Science 119:508–518. doi: 10.1242/jcs.02760.

Matthews BD, Thodeti CK, Ingber DE. 2007. Activation of mechanosensitive ion channels by forces transmitted through integrins and the cytoskeleton. Mechanosensitive Ion Channels, Part A 58:59–85. doi: 10.1016/S1063-5823(06)58003-2.

McKinsey TA, Zhang CL, Olson EN. 2002. MEF2: a calcium-dependent regulator of cell division, differentiation and death. Trends in Biochemical Sciences 27:40–47. doi: 10.1016/S0968-0004(01)02031-X.

Missiaen L, De Smedt H, Parys JB, Sienaert I, Vanlingen S, Droogmans G, Nilius B, Casteels R. 1996. Hypotonically induced calcium release from intracellular calcium stores. The Journal of Biological Chemistry 271:4601–4604. doi: 10.1074/jbc.271.21.12287.

Miyawaki A, Llopis J, Heim R, McCaffery JM, Adams JA, Ikura M, Tsien RY. 1997. Fluorescent indicators for Ca2+ based on green fluorescent proteins and calmodulin. Nature 388:882–887. doi: 10.1038/42264.

Moe PC, Blount P, Kung C. 1998. Functional and structural conservation in the mechanosensitive channel MsCl implicates elements crucial for mechanosensation. Molecular Microbiology 28:583–592. doi: 10.1046/j.1365-2958.1998.00821.x.

Morris CE. 2011. Voltage-gated channel mechanosensitivity: fact or friction? Frontiers in Physiology 2:25. doi: 10.3389/fphys.2011.00025.

Numata T, Shimizu T, Okada Y. 2007a. Direct mechano-stress sensitivity of TRPM7 channel. Cellular Physiology and Biochemistry 19:1–8. doi: 10.1159/000091987.

Numata T, Shimizu T, Okada Y. 2007b. TRPM7 is a stretch- and swelling-activated cation channel involved in volume regulation in human epithelial cells. American Journal of Physiology Cell Physiology 292:C460–C467. doi: 10.1152/ajpcell.00367.2006.

Olsson MC, Patel JR, Fitzsimons DP, Walker JW, Moss RL. 2004. Basal myosin light chain phosphorylation is a determinant of Ca2+ sensitivity of force and activation dependence of the kinetics of myocardial force development. American Journal of Physiology Heart and Circulatory Physiology 287:H2712–H2718. doi: 10.1152/ajpheart.01067.2003.

Orr AW, Helmkne BP, Blackman BR, Schwartz MA. 2006. Mechanisms of mechanotransduction. Developmental Cell 10:11–20. doi: 10.1016/j.devcel.2005.12.006.

Ouyang M, Sun J, Chien S, Wang Y. 2008. Determination of hierarchical relationship of Src and Rac at subcellular locations with FRET biosensors. Proceedings of the National Academy of Sciences of USA 105:14353–14358. doi: 10.1073/pnas.0807537105.

Palmer AE, Giacomello M, Kortemme T, Hires SA, Lev-Ram V, Baker D, Tsien RY. 2006. Ca2+ indicators based on computationally redesigned calmodulin-peptide pairs. Chemistry & Biology 13:521–530. doi: 10.1016/j.chemb.2006.03.007.

Perozo E, Rees DC. 2003. Structure and mechanism in prokaryotic mechanosensitive channels. Current Opinion in Structural Biology 13:432–442. doi: 10.1016/S0959-440X(03)00106-4.

Pinsky DJ, Patton S, Mesaros S, Broukovich V, Kubaszewski E, Grunfeld S, Malinski T. 1997. Mechanical transduction of nitric oxide synthesis in the beating heart. Circulation Research 81:372–379. doi: 10.1161/01.RES.81.3.372.

Rath AL, Bonewald LF, Ling J, Jiang JX, Van Dyke ME, Nicollella DP. 2010. Correlation of cell strain in single osteocytes with intracellular calcium, but not intracellular nitric oxide, in response to fluid flow. Journal of Biomechanics 43:1560–1564. doi: 10.1016/j.jbiomech.2010.01.030.

Rong Y, Distelhorst CW. 2008. Bcl-2 protein family members: versatile regulators of calcium signaling in cell survival and apoptosis. Annual Review of Physiology 70:73–91. doi: 10.1146/annurev.physiol.70.021507.105852.

Rosenblatt N, Hu S, Suki B, Wang N, Stamenović D. 2007. Contributions of the active and passive components of the cytoskeletal prestress to stiffening of airway smooth muscle cells. Annals of Biomedical Engineering 35:224–234. doi: 10.1007/s10439-006-9228-z.

Schug ZT, Da Fonseca PC, Bhanumathy CD, Wagner L II, Zhang X, Bailey B, Morris EP, Yule DI, Joseph SK. 2008. Molecular characterization of the inositol 1,4,5-trisphosphate receptor pore-forming segment. The Journal of Biological Chemistry 283:2939–2948. doi: 10.1074/jbc.M706645200.

Schwartz MA, DeSimone DW. 2008. Cell adhesion receptors in mechanotransduction. Current Opinion in Cell Biology 20:551–556. doi: 10.1016/j.ceb.2008.05.005.

Seong J, Ouyang M, Kim T, Sun J, Wen PC, Lu S, Zhuo Y, Llewellyn NM, Schlaepfer DD, Guan JL, Chien S, Wang Y. 2011. Detection of focal adhesion kinase activation at membrane microdomains by fluorescence resonance energy transfer. Nature Communications 2:406. doi: 10.1038/ncomms1414.

Su LT, Agapito MA, Li M, Simonson WT, Hutterlocher A, Habas R, Yue L, Runnels LW. 2006. TRPM7 regulates cell adhesion by controlling the calcium-dependent protease calpain. The Journal of Biological Chemistry 281:11260–11270. doi: 10.1074/jbc.M512855200.

Sun S, Liu Y, Lipsky S, Cho M. 2007. Physical manipulation of calcium oscillations facilitates osteo-differentiation of human mesenchymal stem cells. FASEB Journal 21:1472–1480. doi: 10.1096/fj.06-7153com.

Taninuma A, Morita T, Nuez A, Shitara A, Hashimoto N, Tojyo Y. 2009. Use of fluorescence resonance energy transfer-based biosensors for the Quantitative analysis of inositol 1,4,5-Trisphosphate dynamics in calcium oscillations. The Journal of Biological Chemistry 284:8910–8917. doi: 10.1074/jbc.M805865200.

Turvey MR, Fogarty KE, Thorn P. 2005. Inositol (1,4,5)-trisphosphate receptor links to filamentous actin are important for generating local Ca2+ signals in pancreatic acinar cells. Journal of Cell Science 118:971–980. doi: 10.1242/jcs.01693.
Vila Petroff MG, Mattiazi AR. 2001. Angiotensin II and cardiac excitation-contraction coupling: questions and controversies. *Heart, Lung & Circulation* **10**:90–98. doi: 10.1046/j.1444-2892.2001.00083.x.

Walker DS, Ly S, Lockwood KC, Baylis HA. 2002. A direct interaction between IP3 receptors and myosin II regulates IP3 signaling in C-elegans. *Current Biology* **12**:951–956. doi: 10.1016/S0960-9822(02)00868-0.

Wang N, Butler JP, Ingber DE. 1993. Mechanotransduction across the cell-surface and through the cytoskeleton. *Science* **260**:1124–1127. doi: 10.1126/science.7684161.

Wang Y, Botvinick EL, Zhao Y, Berns MW, Usami S, Tsien RY, Chien S. 2005. Visualizing the mechanical activation of Src. *Nature* **434**:1040–1045. doi: 10.1038/nature03469.

Wang Y, Jin G, Miao H, Li JY, Usami S, Chien S. 2006. Integrins regulate VE-cadherin and catenins: dependence of this regulation on Src, but not on Ras. *Proceedings of the National Academy of Sciences of USA* **103**:1774–1779. doi: 10.1073/pnas.0510774103.

Wehrens XH, Lehnart SE, Marks AR. 2005. Intracellular calcium release and cardiac disease. *Annual Review of Physiology* **67**:69–98. doi: 10.1146/annurev.physiol.67.040403.114521.

Wei C, Wang X, Chen M, Ouyang K, Song LS, Cheng H. 2009. Calcium flickers steer cell migration. *Nature* **457**:901–905. doi: 10.1038/nature07577.

West AE, Chen WG, Dalva MB, Dolmetsch RE, Kornhauser JM, Shaywitz AJ, Takasu MA, Tao X, Greenberg ME. 2001. Calcium regulation of neuronal gene expression. *Proceedings of the National Academy of Sciences of USA* **98**:11024–11031. doi: 10.1073/pnas.191352298.

Yuan JP, Kiselyov K, Shin DM, Chen J, Shcheynikov N, Kang SH, Dehoff MH, Schwarz MK, Seeburg PH, Muallem S, Worley PF. 2003. Homer binds TRPC family channels and is required for gating of TRPC1 by IP3 receptors. *Cell* **114**:777–789. doi: 10.1016/S0092-8674(03)00716-5.