Mechanical deformation induces depolarization of neutrophils

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The transition of neutrophils from a resting state to a primed state is an essential requirement for their function as competent immune cells. This transition can be caused not only by chemical signals but also by mechanical perturbation. After cessation of either, these cells gradually revert to a quiescent state over 40 to 120 min. We use two biophysical tools, an optical stretcher and a novel microcirculation mimetic, to effect physiologically relevant mechanical deformations of single nonadherent human neutrophils. We establish quantitative morphological analysis and mechanical phenotyping as label-free markers of neutrophil priming. We show that continued mechanical deformation of primed cells can cause active depolarization, which occurs two orders of magnitude faster than by spontaneous depriming. This work provides a cellular-level mechanism that potentially explains recent clinical studies demonstrating the potential importance, and physiological role, of neutrophil depriming in vivo and the pathophysiological implications when this deactivation is impaired, especially in disorders such as acute lung injury.

INTRODUCTION

Several biochemical agents, such as chemokines (1, 2) and bacterial chemoattractants (3, 4), induce the rapid transition of neutrophils from a resting state to a polarized and primed or fully activated state (5–7), leading to their transendothelial migration out of blood and recruitment to sites of infection and sterile injury in surrounding tissues (8). Activated neutrophils are implicated in a very broad spectrum of pathologies, especially infectious and pulmonary diseases, such as acute respiratory distress syndrome (ARDS).

Mechanical perturbations, such as shear stresses and cell stretching, can also induce activation of quiescent neutrophils, just like biochemical agents (9, 10). Furthermore, shear stresses can induce the deactivation of primed and adherent neutrophils in the presence of red blood cells or superoxide dismutase (10–12). The mechanism of shear force–induced deactivation of neutrophils remains open. Because neutrophils in the circulation are repeatedly stretched, for instance, by the multiple very narrow (5 μm) capillary constrictions present in the human pulmonary microcirculation, it is unclear why not all resting neutrophils in circulation eventually become activated.

To address this question, appropriate methods to mechanically stimulate neutrophils under near-physiological conditions (suspended while circulating and without permanent attachment; many rapid, successive deformations mimicking microvascular constrictions) are required. Most current mechanical manipulation techniques (for example, micropipette aspiration, atomic force microscopy, optical tweezers, and magnetic twisting cytometry) are not appropriate (13) largely because they require cells to be immobilized on surfaces, which results in neutrophil activation. An attractive alternative for the noncontact study of primary neutrophils in their suspended state is a microfluidic optical stretcher (OS), where the radiation pressure from two counterpropagating laser beams is used to hold and deform cells (14–17). This technique offers the unique ability to study the dynamic response of neutrophils to chemical priming without contact and at a single-cell level, which, so far, has been unachievable with bulk measurements (8, 18). The OS can also be used to apply single or continued oscillatory deformations to stationary cells, allowing a detailed study of neutrophils during priming and depriming processes.

With such an OS, we show not only that neutrophil priming can be observed directly and quantified mechanically but also that continued oscillatory mechanical deformation of primed neutrophils results in rapid depolarization. The rate at which this happens is two orders of magnitude greater than if the cells are left mechanically undisturbed. These findings are confirmed by experiments with a novel microfluidic flow device, termed microfluidic microcirculation mimetic (MMM), which leaves less time for investigating each cell in detail, but is even closer to the physiological conditions that pertain within the human pulmonary microcirculation. This work has important implications for the understanding and management of immune cell circulatory disorders, such as ARDS, which is characterized by high levels of neutrophil priming in the systemic circulation and, consequentially, high levels of neutrophil entrapment in the pulmonary microcirculation.

RESULTS

Chemical priming of resting neutrophils leads to characteristic molecular readouts and morphological changes

The known priming/activating effects of agents, such as N-formylmethionine leucyl-phenylalanine (fMLP) and granulocyte-macrophage colony-stimulating factor (GM-CSF), were accessed and quantified in the OS experiments at the single-cell level (Fig. 1A). Both GM-CSF (10 ng/ml)– and fMLP (100 nM)–treated neutrophils have significantly (P < 0.0001) higher CD11b expression (fig. S1) and lower CD62L expression (fig. S2) compared to resting neutrophils (for details of priming, see the Supplementary Materials). In addition, intracellular reactive oxygen species (ROS) production was enhanced in primed neutrophils (fig. S3).

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There are obvious morphological consequences of priming, which can be quantified by image analysis of primed neutrophils held stationary in suspension with an OS (Fig. 1A). Resting neutrophils are spherical in shape (see Fig. 1, B to D, top), whereas activated cells form amoeboid shapes (bottom). (C) Phase contrast images of cells trapped in the OS with strong white halo simplifying contour detection. (D) Morphometry using phase contrast images from (C). Edge detection algorithms are used to fit ellipses (red line) to the contours of the cells (cyan line), giving access to the semimajor and semiminor axes, $a$ and $b$, respectively. (E) Plots of the aspect ratio, $a/b$, from the morphometry results of (D). Activated neutrophils have a significantly higher aspect ratio compared to resting neutrophils. (F) Mean circularity of F-actin cortex of cells stained with Alexa Fluor 488 Phalloidin. After fMLP and GM-CSF treatment, cellular circularity decreases significantly. ***$P < 0.001$ (significant difference). Scale bars, 5 μm. Both GM-CSF (10 ng/ml)– and fMLP (100 nM)–treated neutrophils have significantly ($P < 0.0001$) higher CD11b expression (fig. S1) and lower CD62L expression (fig. S2) compared to resting neutrophils (for details of priming, see the Supplementary Materials). In addition, intracellular ROS production was enhanced in primed neutrophils (fig. S3).
respectively. Plots of the aspect ratio, $a/b$, from the morphometric results (Fig. 1E) show that primed or activated neutrophils have significantly ($P < 0.001$) higher aspect ratios compared to resting neutrophils. Over the course of 30 to 120 min, the number of cells that are still primed decreases spontaneously (4). These morphological changes reflect the aforementioned molecular readouts (figs. S1 and S2) and can thus be seen as label-free markers of neutrophil priming. Furthermore, it is known that these morphological alterations are largely due to actin polymerization, which is, in itself, a hallmark of neutrophil activation. We therefore fixed and stained neutrophils using Alexa Fluor 488 Phalloidin (see the Supplementary Materials for details) and quantified the morphological alterations using circularity, $C$. Here, $C = P^2/(4\pi A)$, where $P$ is the perimeter and $A$ is the area. $C = 1$ for a perfect circle, and $C < 1$ for amoeboid shapes. There is a statistically significant reduction ($P < 0.001$) in the mean circularity of the F-actin cortex of cells following fMLP- and GM-CSF–induced priming (Fig. 1F).

**Chemical priming causes cell mechanical changes**

For morphological and mechanical phenotyping by the OS, cells are introduced into a microfluidic delivery system, serially trapped, and then stretched along the laser beam axis (Figs. 1A and 2A). The time-dependent strain is extracted from the video camera images, normalized by the peak stress applied and a geometric factor (21), and used to obtain the creep compliance (inverse of stiffness) for each cell. It should be noted that calculation of the stress applied requires knowledge of the refractive index (RI) of the cells, and we measured this (see the Supplementary Materials) using a digital holographic microscope (22, 23). The average RI was 1.387 ± 0.003 with no detectable difference between the resting and activated state.

Representative strain and compliance profiles (mean ± SEM) were obtained from four independent experiments for resting, fMLP-treated, and GM-CSF–treated neutrophils (Fig. 2, A and B). Reduction in strain and compliance was seen in both GM-CSF– and fMLP-primed cells within 1 s of passive mechanical testing in the OS. The fMLP- and GM-CSF–treated neutrophils were significantly ($P < 0.001$ and $P < 0.0001$, respectively) stiffer than the resting cells. Priming or activation of neutrophils is a time-dependent phenomenon, with a progressive shape change (4). We have measured the mechanical properties of fMLP- and GM-CSF–treated neutrophils during the very early stages of priming when shape alterations are still amenable to accurate elliptical fits (avoiding the few spontaneously deprimed cells). Incidentally, these early stages of priming are physiologically crucial to the sequelae of activities that neutrophils undertake following exposure to the priming signal.

To ascertain whether this reduction in deformability correlates with known cytoskeletal alterations following priming, we fixed and stained neutrophils using Alexa Fluor 488 Phalloidin and imaged these cells using a confocal microscope (Zeiss LSM 700) to obtain the F-actin distribution in the cells (Fig. 2C). Even qualitatively, one can notice the higher intensity of fluorescence signals indicating increased F-actin in primed cells especially within the pseudopodia, which agrees with previous reports (3). Likewise, the amoeboid shape of primed cells in phase contrast images reflects the actin distribution of primed cells (Fig. 1, E and F). Thus, the OS measurements indicate

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![Fig. 2](https://example.com/fig2.png)

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that deformability is a surrogate readout of molecular and cytoskeletal modulations characterizing neutrophil priming.

**Mechanical deformation causes priming of resting neutrophils**

Mechanical perturbations can induce activation of suspended and resting neutrophils, just like soluble agents (9). The OS technique reproduces this effect (Fig. 3, A and B), whereby repeated stretching of unprimed cells leads to a gradual decrease in deformability and change of shape to an amoeboid “polarized” phenotype (fig. S4) in about 40% of the cells, which starts within 20 to 30 s of 0.5- to 1-s pulsatile or sinusoidal stretching (movie S1). Stretching beyond 60 s eventually engenders shape changes in an even greater proportion of cells. However, one might ask, why should soluble biological agents (for example, GM-CSF, fMLP, etc.) produce the same effect in neutrophils as do mechanical perturbations? Moreover, considering the potentially deleterious effects of inadvertent priming of neutrophils in the circulation and the presence of multiple capillary constrictions, which deform and stretch neutrophils, it is surprising that the entire circulating pool of resting neutrophils does not normally become primed by these stretching forces. We therefore investigated what might happen if the already primed neutrophils were subjected to continued, physiologically relevant deformations.

**Mechanical deformations induce depolarization of activated neutrophils**

Multiple mechanical stretching with OS causes chemically (fMLP-) primed neutrophils (Fig. 3C, picture of a typical cell at $t = 0$ s) to depolarize, as evidenced by the retraction of pseudopods and decrease in aspect ratio (inset picture, right) (see also movies S2 and S3). The temporal evolution of the average aspect ratio of $n = 4$ repeats, is shown. (D) Box plots of the average aspect ratio of fMLP-treated cells in (C) at $t = 0$ s and $t = 50$ s. *$P < 0.05$ and ***$P < 0.001$ (significant difference). Scale bars, 5 μm.
depolarize, as evidenced by the retraction of pseudopods and subsequent decrease in their average aspect ratio (see also movie S2). With an onset at about $t = 15$ s, the neutrophils were observed to regain their round, resting shape by $t = 50$ s. About 60% of primed neutrophils (in $n > 6$ experiments) round up within 60 s of 0.5- to 1-s pulsatile stretching. Figure S5 shows the depolarization process of a typical single cell with a particularly strong activated phenotype in the same time interval. The decrease in aspect ratio toward that of resting unprimed cells is statistically significant (Fig. 3D). The same phenomenon also occurs for GM-CSF–primed neutrophils (fig. S6 and movie S3). Quite surprisingly, this deformation-induced depolarization occurred while the cells were suspended in the milieu of the priming agents (fMLP and GM-CSF) for the entire period in the OS; hence, mechanical depolarization was able to override the ongoing chemical priming stimulus. Oscillations in stress of 25% lower magnitude (fig. S7) elicit the same depolarization, albeit the return to round shape takes slightly longer. Critically, resting cells that were mechanically primed can also be depolarized by continued sinusoidal stretching (fig. S8).

Because there could be thermal effects due to the absorption of the OS laser light by water (at 1064-nm wavelength of the OS lasers and at the power of $2 \times 0.9$ W per fiber we used for the stretching, there should be temperature changes of about $22$°C) (24), we undertook stretching using an OS with a 780-nm laser wavelength, where the temperature increase is only a few degrees Celsius. Under these conditions, we again found identical rounding up of primed cells (see movie S4). Thus, mechanical deformation using the OS—and not thermal effects—leads to apparent depolarization of already primed neutrophils. It should also be noted that the cells were still fully viable after mechanical depolarization and could even be chemically “reactivated/primed” (see fig. S9). We can definitively rule out that the recircularization is due to the effect of the optically induced stress. Because of the physics involved, the stress can only be tensile, that is, lead to an extension or stretching of an object along the laser axis, for example, from a sphere to a prolate spheroid, hence the name optical stretcher (14, 15). If the object in the trap has already an extended, nonspherical shape, such as an activated, polarized neutrophil, it cannot be pushed into a spherical shape by the optical stresses.

Furthermore, we note that the morphological readout used in the OS is not, in itself, a formal proof of a change in other aspects of neutrophil function, including NADPH (nicotinamide adenine dinucleotide phosphate) oxidase production and degranulation responses. Unfortunately, these latter processes and other molecular marker–based metrics of depriming cannot as yet be investigated in line at a single-cell level following optical stretching because of the low intensities of luminescence-based assays and the strong photobleaching of fluorescence-based molecular marker assays in the strong laser beams of the optical trap. However, all the morphological observations during mechanically induced depolarization closely overlap with cell changes described during neutrophil depriming (4, 5). The depolarization of mechanically primed neutrophils in reaction to resolin RvE1 also showed a similar morphological phenotype (see movie S5). The possible physiological relevance of mechanical stretching on primed circulating neutrophils can hardly be overlooked, considering that the quickest time taken for a red blood cell to transit the normal pulmonary capillary bed (which consists of 40 to 100 capillary segments) is 0.8 s, with neutrophils transiting only marginally (<3 s) slower compared to red blood cells. The capillary segments in the lung have particularly narrow constrictions (5 μm), certainly smaller than the diameter of a neutrophil (25). Therefore, these cells must deform at time scales close to our stretching pulses of 0.5 to 1 s. Moreover, it takes approximately 60 to 180 s for blood to circulate fully around the body. Thus, it is possible that mechanical deformation is a physiological means of effecting depriming of primed neutrophils in the circulation.

**A microcirculation mimetic confirms OS results**

Circulating neutrophils in vivo deform substantially as they transit through capillary constrictions. Hence, the mechanical deformation that is experienced by circulating neutrophils in the body involves tactile squeezing by the capillary walls. However, the OS uses contactless optical forces. We therefore developed an MMM to more closely replicate the capillary constrictions that impose stretches on circulating cells in vivo.

The MMM is a polydimethylsiloxane-based device (see the Supplementary Materials for details) consisting of serpentine microchannels (Fig. 4A) that mimic the pulmonary microcirculation, with gently tapered inlet and outlet channels from and to reservoirs. This MMM differs from existing devices that model blood vessels in the lungs in that it does not involve branches or channel networks (26). Rather, we have maintained a serial model for two reasons. First, from the “viewpoint” of the cell, constrictions occur one after the other, branches or no branches. Second, a serial model allows us to have a very large number of constrictions (here, 187) within the field of view of the microscope objective. A constant pressure difference maintained between the inlet and the outlet, modeled on physiological pressure gradients experienced in vivo, was used to advect cells through the device, one cell at a time (see movie S5). The minimum gaps at the constrictions (5 μm) are smaller than the diameter of cells but match the size of the pulmonary capillary segments, ensuring that each cell is deformed sequentially during the advection (Fig. 4B, bottom). We used a range of physiologically relevant pressures (10, 50, and 100 mbar, that is, 10.2, 51.0, and 102 cmH2O, respectively) to advect the cells at 37°C and at room temperature (24°C).

Box plots and bar graphs of average transit times of resting neutrophils and fMLP-treated and GM-CSF–treated cells show that the primed cells (as would be anticipated) have significantly longer transit times than resting neutrophils, at 37°C (Fig. 4, C and D, with $P < 0.0001$) and at 24°C (fig. S10). Overall, all cells transit faster at 37°C compared to 24°C, but the relative differences in transit time between each cell state remain. Thus, there is a 20 to 40% increase in transit times between resting and primed cells at both temperatures. Figure 4D shows the transit time as a function of driving pressure for the various cell states. Note that, at 10 mbar, the GM-CSF (10 ng/ml)–treated cells could hardly transit through the device, hence the lack of data for this cell state at 10 mbar. Moreover, advecting resting neutrophils through MMM at very low pressures of 1 to 5 mbar, back and forth, led to pseudopod formation, indicating activation (movie S6). Succinctly, on the basis of transit times, there is a direct correlation between the increase in stiffness of primed cells in the OS within 1 s of stretching (Fig. 2, A and B) and the increase in transit times of primed cells in the MMM.

Furthermore, in the MMM, primed cells, following their advection (at 50 mbar) through 187 constrictions, round up in about 5 to 10 min, showing a significant increase in circularity, compared to their counterparts at the inlet reservoir (Fig. 5, A and B). This increase in circularity potentially functions as a morphological indicator of depriming since resting neutrophils are spherical. In tandem with shape change,
we also found molecular evidence that the primed neutrophils, following stenotic advection in MMM, become depolarized. Hence, there was a significant decrease in the average fluorescence intensity of the CD11b signal in the postconstriction (outlet) cells compared to the preconstriction cells (inlet) (Fig. 5, C and D).

Higher CD11b expression in primed cells compared to resting cells is one of the hallmarks of priming. Thus, the decrease in CD11b expression provides further support for the view that the repeated mechanical deformation of primed neutrophils can induce their return to a resting state. As a second complimentary approach, the MMM strengthens the findings observed for single cells in the OS by also providing a link between CD11b fluorescence and depolarization, which is not possible to assess in the OS as a result of photobleaching during the laser-based trapping and manipulations.

Note that cell viability remained high after advection through the MMM in both resting and primed states (fig. S9). Cells could even be chemically reprimed after mechanical depolarization.

Chemically (GM-CSF–) primed neutrophils, which were mechanically depolarized either by optical stretching or in the MMM and then kept in the medium still containing GM-CSF, became primed again within 5 to 15 min, as evidenced by a strongly polarized cell shape. In addition, by adding fMLP, neutrophils that had been mechanically depolarized in the MMM could be activated again.

These additional observations are instructive because they show that our cells remain viable after mechanical depolarization. Apoptotic or otherwise dying/dead cells would not be able to show such a physiological response. We did not observe differences of mechanically depolarized neutrophils in response to GM-CSF and fMLP. Our results therefore indicate that mechanical stimuli (deformation) can modulate the neutrophil state by priming resting cells and depolarize and/or potentially even deprime already primed or activated neutrophils in suspension. The mechanical depolarization result, which, in itself, is quite astonishing, suggests that depolarization may well happen in vivo because deformation-imposing constrictions abound in the microcirculation, especially within the lung. Accumulation of primed neutrophils is a major feature of several blood/immune system dysfunctions, such as atherosclerosis (27) and chronic obstructive pulmonary disease (28). Can it be that these dysfunctions result, inter alia, from a failure of neutrophil depolarization?

**DISCUSSION**

Neutrophils are crucial for immunity against infection, but research on them has been hampered by their experimentally difficult nature (8). Even slight perturbations with experimental tools alter the neutrophil state. Our use of nontactile optically generated forces in the OS enables
detailed quantification of neutrophil morphology and mechanical properties as a new contact- and label-free method to assess priming. The size of resting neutrophils that we report here (10.39 ± 0.19 \(\mu\)m in diameter) is larger than measurements made over two decades ago (7, 29) but agrees with more recent measurements, such as those of Ruban et al. (30), who report human granulocyte diameters of 9.6 ± 0.5 \(\mu\)m. Moreover, our measurements have been undertaken on neutrophils in suspension, which is the physiological state of these cells while in the circulation. All previous neutrophil morphometries were, as far as we are aware, carried out on cells either chemically fixed or confined to a surface or to some tactile support with the attendant obfuscation of results (31). The stiffening of primed compared to resting neutrophils, which we found in OS measurements, fits well with their delayed transit in the MMM. These results agree with previous reports on mechanical properties of activated neutrophils (7, 32, 33) and the delayed transit of primed neutrophils through the lungs in humans (34). However, when Gossett et al. deformed neutrophils at very high strain rates (\(\sim 105 \text{ s}^{-1}\)) using deformability cytometry (DC) (35), they found the opposite: Primed cells were more deformable. These strain rates are known to fluidize F-actin (36), and DC apparently probes nuclear mechanics rather than the cytoskeleton (35). Otto et al. (37), using real-time DC (RT-DC), also found agonist-activated neutrophils to be more deformable than resting neutrophils. Both techniques measure cell mechanics at very short time scales: microseconds in DC and milliseconds in RT-DC. Perhaps, activated neutrophils are stiffer than resting neutrophils at longer time scales of seconds and minutes (as seen in OS and micropipette aspiration), which are physiologically more relevant to neutrophils in vivo. Functionally, increased stiffness following priming of suspended neutrophils might be useful in the tethering and rolling processes that precede the extravasation of activated neutrophils out of the vasculature into the surrounding tissues.

Multiple stretching of resting neutrophils in the OS and advection through MMM at very low pressures led to the activation of a subset of these cells, indicating that in vivo–like stretching forces can activate neutrophils. Reports of this mechanically induced priming are found in the literature (9). Our earlier discovery that neutrophil priming is spontaneously reversible (4), the ongoing search for depriming mechanisms (38, 39), and the direct involvement of primed neutrophils in so many pathological states accentuate the importance of our novel finding that primed neutrophils can be depolarized by mechanical stimuli. From a broader biophysical perspective, mechanical deformation is a potent stimulus for cellular processes, such as growth, differentiation, migration, remodeling, and gene expression (40). We report here for the first time that whole-cell deformation depolarizes preprimed neutrophils. This is distinct from previous reports of depriming by fluid shear stress, which was insufficient to cause passive deformation of cells, acted via surface receptors only, and required the presence of erythrocytes or superoxide anion scavengers (10). Our single-cell investigations also provide unprecedented real-time insight into the kinetics of mechanical depolarization, which occurs within

![Fig. 5. Depolarization in the MMM: shape change and molecular readouts.](image-url)
cleavage of and depolymerization of F-actin, which, together with proteolytic receptor down-regulation with concomitant deactivation of Rac, suggests such an in vivo mode of depolarization. Succinctly, although it may reflect an in vitro artifact and would need to be coupled reagents, how does the healthy body deprime neutrophils? Because chemical stimuli prime and seemingly deprime neutrophils, our results extend the trend with mechanical stimuli also priming and depolarizing/depriming neutrophils. Circulating neutrophils have to repeatedly squeeze through microvasculature constrictions smaller than their diameter while being advected through the pulmonary capillary bed at high speed (25). Although it may reflect an in vitro artifact and would need to be substantiated in vivo, our observed in vitro depolarization by deformation suggests such an in vivo mode of depolarization. Succinctly, the physiological and pathological implications of our discovery—that in vivo—mimicking deformations can depolarize activated and suspended neutrophils—may explain why circulating neutrophils do not all eventually get activated. Our finding presents neutrophil mechanotransduction as a biophysical challenge with clinical promise. Neutrophils are notoriously difficult to work with, which could be one reason why less is known about potential mechanosensitive mechanisms than for many other cell types. The main body of work on mechanical depriming of neutrophils by fluid shear stress has been contributed by the group of Schmid-Schönbein (10–12, 43, 44). The physical stimulus seems to be transduced by heterotrimeric guanine nucleotide–binding protein–coupled receptor down-regulation with concomitant deactivation of Rac and depolymerization of F-actin, which, together with proteolytic cleavage of β3 integrin, facilitates membrane detachment and lamellipodia retraction. In other cell types, Rac controls the cooperative relationship between myosin II and the actin cross-linker cortezillin I, where both proteins are essential for cellular mechanosensory responses (45). Rho and CDC42 do not seem to be involved (46). The sensitivity to fluid shear stress can also be modulated by the amount or organization of cholesterol, which changes the membrane’s physical properties (47). Recently, other membrane receptors, such as the formyl peptide receptor, have also been implicated in the depolarizing mechanism (48), whereas the activation with PAF can also be increased by fluid shear stress (49). However, the situation that pertains to the phenomena we report appears substantially different for two main reasons: (i) The fluid shear stresses used to elicit the mechanisms reported and discussed previously are “between about 1 and 10 dyn/cm²”, a range that is below the level to achieve a significant passive, viscoelastic response” (43). Stresses even in the OS are between 1 and 10 Pa (10 to 100 dyne/cm²), an order of magnitude larger and even greater in the MMM. Thus, in our case, the entire cell is being deformed, and with it the plasma membrane, the actin cortex, the microtubule cytoskeleton, and also the nucleus. Any of these structures, which bear tensile or compressive stresses during deformation, could be involved in mechanotransduction in principle so that it is possible that mechanisms other than those described for the sensing of fluid shear stress might contribute (50). (ii) Cells in our case are fully suspended, with no chance to adhere to anything. This rules out any mechanism involving integrin-mediated mechanosensing involving focal adhesion complexes—likely the most important and best-studied mechanism of mechanosensing in adherent cells (51–53). The molecular mechanism behind the whole-cell deformation—induced depolarization (19) is now an open question, as is whether the observed depolarization actually represents depriming.

In summary, we have used several biophysical tools specifically designed to study neutrophil priming and depolarization by deformations mimicking the passage through the pulmonary microcapillary bed and have obtained results that provide novel insights into neutrophil function, with putative pathophysiological consequences. Optical stretching has afforded us a contact-free assessment of neutrophil mechanics in suspension, mimicking the circulatory environment of neutrophils. We have shown that multiple stretching of primed neutrophils in suspension leads to their rapid depolarization if not even to depriming. Because the MMM, from a pressure and size perspective, models the pulmonary capillary bed, the mechanical depolarization we have uncovered presents itself as a cellular-level mechanism for recent clinical inference that the pulmonary vascular operates as a site for physiological depriming, with obvious pathological implications when such a depriming mechanism is impaired.

Materials and Methods

Study Design

This study tested the response of resting and activated neutrophils from healthy adults to in vivo–like mechanical stimuli, imposed during controlled laboratory experiments. Institutional review board guidelines of the University of Cambridge (UK) and the Technische Universität Dresden (Germany) were followed.

Cell Preparation

Primary human neutrophils were isolated from peripheral blood taken from more than 100 healthy donors at the Department of Medicine, Addenbrooke’s Hospital, University of Cambridge and...
The resting neutrophils were immobilized using poly-D-lysine (Materials) using a digital holographic microscope (DHM) (Optical stretching and determination of creep compliance). The forces that trap and deform the cell outwardly along the surface arise from the change in the RI at the cell medium interface and the conservation of momentum. The axial strain during optical stretching is given by

\[ \varepsilon(t) = \frac{a(t) - a_0}{a_0} \]  

where \( a_0 \) is the semimajor axis of the nonstretched cell and \( a(t) \) is the time-varying semimajor axis measured (Fig. 1F). The optical stress on the cells was computed using an electromagnetic wave model (55), which requires knowledge of the cells’ average RI. We measured the RI of resting neutrophils (see the Supplementary Materials) using a digital holographic microscope (DHM) (22, 23). The resting neutrophils were immobilized using poly-D-lysine-coated slides. Because surfaces can easily activate neutrophils, we checked the DHM results by simultaneously trapping suspended resting neutrophils in the OS and measuring their RI using a quantitative phase camera (Phasics). There was no statistically significant difference in the average RI measured using DHM or the Phasics camera. Thus, the mechanical properties reported here are decoupled from the dielectric characteristics of cells.

The strain was normalized by the peak value of the calculated optical stress, \( \sigma_{th} \) and a geometric factor, \( F_g \), to give the creep compliance

\[ J(t) = \frac{\varepsilon(t)}{\sigma_{th} F_g} \]  

The geometric factor was calculated as described elsewhere (21) to account for cell shape and stress distribution.

Fluorescence staining and confocal microscopy
Alexa Fluor 488 Phalloidin was used to treat cells fixed with 4% formaldehyde and permeabilized with 0.1% Triton X-100. For immunofluorescence, fluorescein isothiocyanate–conjugated monoclonal CD11b antibody and Pacific Blue–conjugated monoclonal CD62L were used on live cells. Nuclear staining to verify lobulation was undertaken using Syto 61 dye, whereas MitoTracker Orange was used to identify cytoplasmic regions. A confocal microscope, LSM 700 (Zeiss), was used for imaging and fluorometry, with all settings (gain, percentage transmission, etc.) kept constant for all samples of a given experiment for valid comparisons. Further details are given in the Supplementary Materials.

Statistical analysis
For each cell treatment condition, the number of cells per OS experiment was usually \( n > 30 \). Strain and creep compliance data are presented as mean ± SEM. Representative compliance data were chosen from a total number of \( n \) independent experiments, as indicated in the figures and figure legends. Overall, the number of individual cells analyzed totaled about 4000 for OS and about 2000 for MMM. For each cell state or treatment condition, the number of cells per advection experiment (MMM) was \( n > 40 \). Advection times are representative of three independent experiments and are presented as mean ± SEM. Statistical comparisons for all experiments were performed using one-way analysis of variance (ANOVA) in Origin (OriginLab Corporation) to account for multiple groups. Within ANOVA, significant differences were reported only where at least three different mean comparison tests (Tukey, Bonferroni, and Dunn–Šidák) simultaneously showed these differences. All box plots include whiskers at the 5 to 95% range, horizontal box lines at the 25 to 75% range, the median line, and the mean (inset box).

**SUPPLEMENTARY MATERIALS**

Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/3/e1602536/DC1

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