Accelerating metabolism and transmembrane cation flux by distorting red blood cells

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Under static conditions, mammalian red blood cells (RBCs) require a continuous supply of energy, typically via glucose, to maintain their biconcave disc shape. Mechanical distortion, in a complementary way, should lead to increased energy demand that is manifest in accelerated glycolysis. The experimental challenge in observing this phenomenon was met by reversibly and reproducibly distorting the cells and noninvasively measuring glycolytic flux. This was done with a gel-distorting device that was coupled with $^{13}$C nuclear magnetic resonance (NMR) spectroscopy. We measured [3-$^{13}$C]lactate production from [1,6-$^{13}$C]D-glucose in the RBCs suspended in gelatin gels, and up to 90% rate enhancements were recorded. Thus, for the first time, we present experiments that demonstrate the linkage of mechanical distortion to metabolic changes in whole mammalian cells. In seeking a mechanism for the linkage between shape and energy supply, we measured transmembrane cation flux with Cs⁺ (as a K⁺ congener) using $^{133}$Cs NMR spectroscopy, and the cation flux was increased up to fivefold. The postulated mechanism for these notable (in terms of whole-body energy consumption) responses is stimulation of Ca⁺-adenosine triphosphatase by increased transmembrane flux of Ca⁺ via the channel protein Piezo1 and increased glycolysis because its flux is adenosine triphosphate demand–regulated.

INTRODUCTION

Cell shape and deformability

In the human cardiovascular circulation, each red blood cell (RBC; erythrocyte) traverses two narrow capillaries, on average, every 1 min; a typical capillary in peripheral tissues and the lungs has a diameter half that of the RBC. The capillary-imposed distortions that occupy ~1% of the blood circulation time, coupled with flow-induced shape changes, mean that RBCs spend a significant fraction of their ~120-day life span being dynamically distorted. Having an experimental means of quantifying biophysical and biochemical responses of RBCs to physical deformation (1) is fundamental to understanding their energy consumption (that is poorly characterized) and metabolic interactions with the vascular endothelium in tissues (2).

Even under static conditions, RBCs require a continuous supply of free energy via glycolysis (3) to maintain their biconcave disc shape (4). The deformability of the RBC depends on metabolism, with depletion of glucose promoting a morphological transition from discocyte to echinocyte and then spherocyte (5). The phenomenon of RBC “membrane flickering” that was first described in 1890 (6) has been vigorously investigated in recent years (7–9). The viscosity of the medium affects the frequency and amplitude of the membrane oscillations that suggests a metabolic driving force (10), but a less direct effect of energy supply acts via cytoskeletal (spectrin) rearrangements (7) that alter the phosphorylation state of an interconnecting protein, 4.1R, thus changing membrane-spectrin interactions (11). There is also some evidence that RBC deformation leads to adenosine triphosphate (ATP) release from the cells (12), although it is important to state that we found no evidence of this in the current work.

Mechanosensation in cells

Mechanosensory transduction via stretch-activated (mechanosensitive) ion channels links physical stimuli such as pressure and stretch to biological responses (13). Piezo1 and Piezo2 were the first mechanosensitive ion channels to be identified in mammalian cells (14–16), and their important role has been demonstrated in mechanical nociception (17) and vascular development (18).

Most investigations of the functions of the Piezo proteins have been carried out with electrophysiological patch clamping (19), but significant differences between outcomes of patch-clamping and whole-cell experiments have been mooted (20). Here, for the first time, we present experiments that demonstrate the linkage of mechanical distortion to metabolic changes in whole mammalian cells.

Metabolism-shape duality

Using this extensive background on effectors of RBC shape, deformability, and membrane flickering, we postulated that mechanically imposed distortion would increase energy utilization in RBCs, which is associated with an autonomous drive toward restoring their natural biconcave disc shape, and that it could be studied by NMR spectroscopy. If this reciprocal relationship existed (namely, shape maintenance requires energy delivered through metabolism, and reciprocally changing shape increases metabolic rate), it would pave the way to more quantitative experiments on factors that regulate shape and volume in all cells. We aimed to mechanically distort RBCs in a reproducible manner while noninvasively measuring their glycolytic flux.

Experimentally imposing shape changes

Variably stretchable gels have been previously used in analytical NMR spectroscopy to characterize mixtures of chiral compounds (21, 22), to measure transmembrane exchange of deuterated water in RBCs (23) and to study the extent of ordering of cations inside RBCs (24). Recently, we demonstrated that RBCs are metabolically stable in carefully adjusted (with respect to pH and gelatin concentration) gels, and the cells become physically distorted in mechanically stretched/compressed gel samples (25). Here, we report the first-ever use of elastic gels to study the effects of mechanically induced shape changes on metabolic rate and cation transport in whole cells.

Aims and central hypothesis

We aimed to use the gels for holding RBCs in a distorted state, thus forcing their mechanosensitive ion channels (Piezo1) to stay open for...
influx of cations (Na\(^+\), K\(^+\), Cs\(^+\), Mg\(^{2+}\), Ba\(^{2+}\), and Ca\(^{2+}\)) (26). Of these, Ca\(^{2+}\) carries the most significance, as the maximal velocity of Ca-adenosine triphosphatase (Ca-ATPase) (27) is more than twice that of Na,K-ATPase, which accounts for ~40% of the ATP turnover in the resting state of RBCs (28). Until now, this has been declared as “unused catalytic potential.” We now postulate it to be part of the “rapid homeostatic response system” of the RBC, which is called upon under mechanical distortion during rapid flow in the bloodstream and in a more extreme manner in capillaries, as noted above.

RESULTS

RBC glycolysis in gels

The metabolic stability of human RBCs in elastic gelatin gels at 20°C is demonstrated in Fig. 1. \(^{13}\)C NMR spectra were acquired from fresh cells suspended in pH-adjusted bovine gelatin gel to monitor conversion of \([1,6-{^{13}}\text{C}]\text{D-glucose}\) (initially 9.95 mM) to \([3-{^{13}}\text{C}]\text{L-lactate}\). The ongoing glycolysis in the RBCs led to the decline of the glucose resonances (α and β anomers) at chemical shift, \(d = 91.3\) and 95.1 parts per million (ppm), respectively, and the emergence of the resonances of \([3-{^{13}}\text{C}]2,3\)-bisphosphoglycerate (23BPG) and \([3-{^{13}}\text{C}]\text{L-lactate}\) at \(d = 65.6\) and 19.3 ppm, respectively. The resonance at \(d = 13.1\) ppm was from \([6-{^{13}}\text{C}]\text{L-methionine}\) that had been added as an internal peak integral standard; it remained of the same intensity throughout the 45-hour time course. These data showed that the RBCs remained metabolically active for at least 45 hours in the gel, generating \([3-{^{13}}\text{C}]\text{L-lactate}\) at a constant rate (see fig. S1 for the graph and the measured rate of glycolysis).

Stretching the silicone rubber tube to 1.7 times its original length, while the RBC-gelatin sample was still in the liquid state, and then allowing the gel to form at 10°C yielded a sample in which the RBCs were described as “70% compressed” when the tension in the silicone tube was released. When the RBC-gelatin sample was allowed to gel at 10°C in a relaxed silicone tube, it could be stretched to various extents up to twice the original length before the gel broke.

Seventy percent compression routinely yielded significant glycolytic rate enhancements like that shown in Fig. 2; however, stretching the sample by 70% always gave smaller enhancements (~60% of the corresponding compression value). We postulate that this is due to the fundamental difference in the anisotropy of the strain field in the gel imposed by stretching and compression.

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Fig. 1. \(^{13}\)C NMR (100.61 MHz) spectral time course of human RBCs metabolizing \([1,6-{^{13}}\text{C}]\text{D-glucose}\) in relaxed gelatin gel. Suspension medium was 60 mM NaOH, 110 mM NaCl, 10 mM KCl, and 10 mM CaCl\(_2\) at pH 7.4 and 20°C with 9.95 mM \([1,6-{^{13}}\text{C}]\text{D-glucose}\). Resonance assignments were as follows: 91.3 and 95.1 ppm, C1 of α and β anomers of \([1,6-{^{13}}\text{C}]\text{D-glucose}\); 65.6 ppm, C3 of 23BPG; 59.8 ppm, partially resolved C6 of α and β anomers of \([1,6-{^{13}}\text{C}]\text{D-glucose}\); 19.3 ppm, methyl carbon of \([3-{^{13}}\text{C}]\text{L-lactate}\); 13.1 ppm, methyl carbon of \([6-{^{13}}\text{C}]\text{L-methionine}\) added as an intensity reference; and 0 ppm, natural-abundance \(^{13}\)C in a silicone rubber tube used as the chemical-shift reference. A ~30° excitation pulse was used, with a recovery delay of 2 s per free induction decay (FID), to acquire 816 transients per spectrum, giving a total acquisition time of 30 min. For clarity, only every 10th spectrum is shown.
Membrane transport in gels—\(^{133}\text{Cs}\) NMR

Having established that distortion-enhanced metabolism was dependent on \(\text{Ca}^{2+}\), we postulated that the effect was mediated by Piezo1 (29). Therefore, we tested whether the transmembrane cation fluxes in whole RBCs would also be sensitive to cell distortion. Figure 3 shows the reversible enhancement of \(^{133}\text{Cs}^+\) efflux from \(\text{Cs}^+\)-loaded RBCs when they were compressed in a gel, prepared in the same manner as for Figs. 1 and 2 (and fig. S1). The RBCs had been pre-loaded with \(^{133}\text{Cs}^+\) by incubating them for 2 days at 21°C in isotonic NaCl-saline, in which 100 mM CsCl replaced the equivalent NaCl. Extracellular \(^{133}\text{Cs}^+\) was removed by two cycles of centrifugation upon resuspending the cells in 154 mM NaCl. At the start of the time course in Fig. 3, the \(^{133}\text{Cs}\) NMR spectrum showed a single prominent resonance at \(\delta = 11.6\) ppm, corresponding to the \(^{133}\text{Cs}^+\) inside the cells. Over the course of 2 hours, there was growth of a second peak, corresponding to extracellular \(^{133}\text{Cs}^+\). The lower inset in Fig. 3 shows the third spectrum acquired with a substantial extracellular resonance at \(\delta = 10.55\) ppm.

After 2 hours, the tension in the silicone tube was released to compress the gel, and recording \(^{133}\text{Cs}^+\) efflux continued for the next 2 hours. Regression of a straight line onto the data at this stage showed that the rate of efflux had increased 5.4-fold (Fig. 3).

Quantifying the extracellular peak integral required special analysis because \(^{133}\text{Cs}^+\) in the anisotropic environment of the stretched gelatin gel displays quadrupolar splitting (24). A singlet in relaxed gel became a septet \((^{133}\text{Cs}^+\) has a nuclear spin of 7/2) on compression or stretching due to spatial anisotropy of the gel, causing residual quadrupolar coupling (see Supplementary Discussion) (30).

In the third stage of the experiment, the silicone tube was stretched to 1.7 times its previous length, thus returning the contents back to the relaxed state. \(^{133}\text{Cs}\) NMR spectra showed a gradual rise of the resonance from the extracellular \(^{133}\text{Cs}^+\). The slope of the line regressed from these data was substantially smaller than that in the first stage, because the system was approaching equilibrium.

The \(^{133}\text{Cs}^+\) efflux experiment in Fig. 3 followed the same sequence of relaxed-compressed-relaxed as the metabolic one in Fig. 1, but the cation flux had a much greater rate enhancement (a factor of >5).

The results of a complementary experiment with the sequence of stages in the reverse order in Fig. 3 are shown in fig. S2.

Glycolysis in RBC suspensions

Apart from mechanical stimuli, Piezo1 is activated chemically by the compound yoda1, which is known to enhance transmembrane \(\text{Ca}^{2+}\) flux (31). \(^{13}\text{C}\) NMR spectra of RBC suspensions without and with added yoda1 are shown in Fig. 4, and they display the same resonances as in Fig. 1. Because the RBC suspensions (not in gel) were incubated at 37°C, the rate of lactate production was much higher, with a 50% extent of reaction after ~4 hours. When \(2\) μl of 28.15 mM yoda1 [dissolved in dimethyl sulfoxide (DMSO)], giving a final concentration of \(19\) μmol (liter sample)\(^{-1}\), was added to an identical sample, the time course in Fig. 4B was obtained. Note the marked enhancement of lactate production over the control experiment (Fig. 4A).

Experiments carried out with differences in suspension medium, and with either \([2-^{13}\text{C}]\)acetate or \([6-^{13}\text{C}]\)-methionine as intensity references, revealed that specific intensities of the signals from the various metabolites differed between experiments. Therefore, control rates were scaled to one value, \(3.00\) mmol (liter RBC)\(^{-1}\) hour\(^{-1}\), as reported in our previous studies of human RBC metabolism (32). The rates of lactate production in the two time courses (Fig. 4) were \(3.0 \pm 0.5\) mmol (liter...
Concentration of 150 porin 1, in RBCs (proteins, including the integral membrane water transporter, aquaporin 1, in RBCs (proteins, including being an inhibitor of mitochondrial Ca\(^{2+}\) transport in human RBCs, the following experiments were carried out, with the results summarized in table S1. In NaCl-saline medium, yoda1 stimulated the rate of RBC lactate production by a factor of ~3, an effect replicated in three experiments but not in a fourth (table S1; row 1 compared with row 2, 9 with 10, 17 with 18, and 37 with 38). In addition, in separate experiments, we added eosin, p-chloromercuribenzenesulfonate (pCMBS), ammoniated ruthenium oxychloride (ruthenium red), calmidazolium, and peptide GsMTx4 (38).

Eosin is a Ca-ATPase inhibitor in RBCs (33), but even at the high concentration of 150 \(\mu\)M, it had no effect on the yoda1-stimulated glycolytic rate, whereas pCMBS reacts with exposed sulphydryl groups in proteins, including the integral membrane water transporter, aquaporin 1, in RBCs (34). Even at the high concentration of 150 \(\mu\)M, eosin had no effect on the yoda1-stimulated glycolytic rate, whereas pCMBS diminished the yoda1-enhanced response to around two-thirds of the control value, implying that exposed –SH groups are involved with the binding of the compound to Piezo1 or to a protein in close juxtaposition in the membrane.

Ruthenium red is an inorganic dye that interacts with many proteins, including being an inhibitor of mitochondrial Ca\(^{2+}\) transport (35), and the transient receptor potential (TRP) ion channel (36) that has electrophysiological characteristics similar to those of Piezo1. At concentrations of 240 and 400 \(\mu\)M, it had no significant effect on yoda1-induced stimulation of glycolysis.

Calmidazolium binds to calmodulin and inhibits Ca-ATPase in nerve cells (37). With RBCs in NaCl-saline, it stimulated glycolysis by ~80% (table S1; row 15 versus row 9). Additionally, in the presence of yoda1, the previously attained ~3-fold enhancement was reduced to 2.4-fold (row 16/row 9). This provided further evidence of (at least indirect) involvement of Ca\(^{2+}\) in the yoda1-stimulated metabolic response. 133Cs NMR spectra of RBCs with 133Cs+ in the extracellular medium showed immediate cation influx after the addition of 19 \(\mu\)M yoda1. Of great relevance was the inhibition of yoda1-initiated 133Cs+ influx (lower curve in fig. S3B) by GsMTx4. This peptide toxin is a known inhibitor of Piezo1 (38, 39), so this result provided solid evidence of a Piezo1-mediated mechanism of the observed 133Cs+ transport.

**DISCUSSION**

Various aspects of the experimental design and data analysis are discussed in detail in Supplementary Discussion. Here, we focus on the overall, more general interpretation of the results.

**Model of RBC shape distortion responses**

Figure 5 shows a schematic diagram of the biochemical systems we postulate to be involved in the compression/stretched-induced glycolytic and 133Cs+ flux responses that were observed. The caption lists the transporters, pumps, and overall glycolytic metabolism that are interconnected in the responses.
Compression-induced glycolytic enhancement is explained by the opening of Piezo1 with the consequent influx of Ca\(^{2+}\), stimulating Ca-ATPase. This enzyme/pump reduces the concentration of ATP that, in turn, reduces the high-substrate inhibition of hexokinase, at its allosteric site, and accelerates glycolysis (32, 40–42).

Because the RBCs are held in a distorted state, Piezo1 stays open (on average) and is subject to "usage activation" (26), a phenomenon observed after repeated stimulation in patch-clamping experiments. The high maximal velocity of RBC Ca-ATPase (43), which is more than twice that of Na,K-ATPase that accounts for ~40% of the ATP turnover in normal resting RBCs (28), has been an enigma. It had been surmised that the high \(V_{\text{max}}\) of Ca-ATPase represented unused catalytic potential. However, we now postulate that this high catalytic capacity is exploited most extensively during the ~700 ms every minute when RBCs are distorted in capillaries during normal blood flow. It is part of the rapid homeostatic response system of the RBC.

We hypothesize that, once elevated at the site of entry through Piezo1, the Ca\(^{2+}\) signals locally to the cytoskeleton to bring about structural rearrangements that lead to strain in the protein network, which drives a return of the cell to its original biconcave shape. The duration of this signal (of cell distortion) is determined by the rapid ejection rate of Ca\(^{2+}\) from the RBC by Ca-ATPase.

**Future directions**

The role of the cytoskeleton in the metabolic and cation transport responses warrants further investigation. Changes in the arrangement of spectrin could be contributing to what is called mechanoprotection (26) that reduces the extent of the effects we report here. The metabolic and cation transport effects we observed were rapidly reversible (Figs. 2 and 3 and fig. S2), so this points to membrane-protein shape changes as the trigger. This is not a covalent modification or phosphorylation but simply mechanical distortion of Piezo1. Following downstream in this signaling cascade is rearrangement of the cytoskeleton on a longer time scale (minutes) based on protein phosphorylation affecting overall deformability (7, 11).

The participation of endogenous effector ligands in the reported metabolic and cation transport responses has not yet been explored/discovered. However, in view of the recent finding of lactate gating of carotid body glomus cell Ca\(^{2+}\) flux (44), this is an area worth investigating in RBCs.

**CONCLUSIONS**

It is a fundamental property of all cells that they sense mechanically and physicochemically induced changes in their shape. We report here a new noninvasive approach to quantifying perturbations of cellular metabolism and transmembrane cation flux that are brought about by altering the shape of cells. This was done by first suspending the cells in a gel and then mechanically distorting the whole sample. Another challenge when developing this study was to inspect inside the cells and record changes in their metabolite composition and transmembrane cation distributions. This was achieved using NMR spectroscopy with its
Fig. 5. Schematic representation of the Ca²⁺-mediated linkage between activation of Piezo1 by mechanical distortion of the RBC and stimulation of its glycolysis. The various participants in this complex response are as follows: GLUT1, which is the glucose transporter required to deliver this fuel molecule to the cytoplasm; Ca-ATPase, which responds to an increase in [Ca²⁺] by catalyzing the hydrolysis of one molecule of ATP per Ca²⁺ ion ejected from the cytoplasm; and Na,K-ATPase, which constitutively pumps three Na⁺ ions from the cell while simultaneously importing two K⁺ ions with the concomitant hydrolysis of one molecule of ATP; in the physiological operation of the RBC, this reaction consumes ~40% of the free energy derived from glycolysis (28, 32). Capnophorin (also called Band 3) catalyzes the one-for-one exchange of the anions, Cl⁻ and HCO₃⁻, and is central to the attainment of bulk electroneutrality. The monocarboxylate transporter (MCT) mediates the facilitated diffusion of lactate across the RBC membrane. Its operation was evident from the splitting of the [3-¹³C]-lactate resonance that indicated two compartments (inside and outside the RBCs) from the ¹³C NMR spectral time courses. The Gárdos channel (also called KCa3.1 or KCNN4) mediates the efflux of K⁺ under the control of Ca²⁺. The membrane protein glycoporphin is linked to the cytoskeleton, as is capnophorin; these linkages are made in as yet to be defined ways, but they are affected by Ca²⁺. Piezo1 mediates the exchange of both monovalent and divalent cations, with permeabilities (P) in the order of P₁⁺ > P₂⁺ > P₃⁺ > P₄⁺ > P₅⁺ (26). The free energy of covalent bond cleavage of glucose and its subsequent metabolites is captured as anhydride bond energy in ATP. This “energy currency” molecule is “spent” on driving the two ATPases shown in the diagram, as well as other kinase reactions not shown, producing adenosine diphosphate (ADP). Glycolysis is ATP demand–regulated so increased influx of Ca²⁺ stimulates Ca-ATPase that increases ATP hydrolysis and hence glycolytic flux. The positive feedback of extracellular lactate to Piezo1 is only speculated to occur in RBCs based on the findings on glomus cells from the carotid body (44).

almost unique capability to record signals from ions and metabolites in cells.

Drugs affecting the pathways involved in the connection between cell shapes and metabolism will potentially lead to the ability to manipulate biochemical and energetic steady states in the whole body. The experimental approaches described here are ready for application to other more phylogenetically primitive cells and to highly evolved ones, including those in pathological states like neoplasia.

### MATERIALS AND METHODS

#### Chemicals and solutions

Analytical reagent chemicals were purchased from Sigma-Aldrich, unless otherwise indicated. The saline used for the preparation of all RBC samples (154 mM NaCl) was filtered three times through a 0.2-μm membrane filter (Millipore) to remove potentially infective particulate matter. The osmolality of each solution was adjusted to 290 ± 5 mosmol kg⁻¹ guided by a vapor pressure osmometer (model 5520, Wescor Instruments).

#### Red blood cells

Blood was obtained by venipuncture from the cubital fossa of consenting healthy donors under the approval of the University of Sydney Human Ethics Committee (Institutional Review Board) for the project. Heparin (15 U ml⁻¹) was added as the anticoagulant, and the blood was centrifuged at 3000g for 5 min at 4°C. The plasma and buffy coat were removed by vacuum pump aspiration. The RBC pellet was washed three times in five volumes of saline, with repeated centrifugation and supernatant aspiration. Before the last wash, the RBC suspension was bubbled with carbon monoxide for 10 min to convert the Fe(II) in hemoglobin to its stable diamagnetic form in carboxyhemoglobin. For prolonged time courses, the final washing medium consisted of 60 mM NaOH, 110 mM NaCl, and 10 mM KCl.

Gelsatin (1.75 g, grade 20N, Gelita) was dissolved in 5 ml of “neutralizing saline” consisting of 60 mM NaOH, 110 mM NaCl, and 10 mM KCl. This particular amount of NaOH yielded pH 7.4 in the otherwise acidic gelatin solution. The mixture was heated for ~20 min at 60°C in a water bath until dissolution occurred and then was cleared of air bubbles by centrifugation at 2000g for 30 s.

RBC suspension (2.0 ml, Ht ~ 0.85) was added to the gelatin solution at 42.0°C to give a final Ht ~ 0.20. The sample was gently mixed (so as not to introduce new bubbles) with a spatula, withdrawn into a silicone rubber tube of 5.0-mm inside diameter (ID) and 6.9-mm outside diameter (OD) (Sims Portex) with a 5-ml disposable syringe, and sealed with a Delrin plug. The time of exposure of the RBCs to higher than physiological temperatures was kept to ~2 min to avoid...
potential metabolic and/or morphological changes. The samples were inserted into bottomless thick-walled glass NMR tubes of 8.0-mm ID and 10.0-mm OD (New Era Enterprises) and then stored at 10°C for 30 min to promote gelation. Stretching the gel was achieved in the previously described apparatus (further information about the apparatus is given in the Supplementary Materials) (21, 45). For gels destined for later compression, the silicone rubber tube was stretched inside the outer glass tube while the gelatin was still in the liquid state. After gelation, the sample was compressed by releasing the thumbscrew. The extent of stretching or compressing the gels (and RBCs), ε, was defined as

$$\varepsilon = \frac{|l_l - l_0|}{l_0} \times 100\% \quad (1)$$

where $l_l$ is the length of the elastic tube in the stretched/compressed state and $l_0$ is the original length of the elastic tube up to the rim of the outer glass tube.

Samples used for $^{13}$C NMR measurements were prepared as described previously (25), with the addition of 9 to 15 mM [1,6-$^{13}$C]d-glucose; they were incubated for up to 45 hours in either relaxed or compressed states ($\varepsilon = 50$ to 70%) at 20°C.

Calculation of solute and ion concentrations in RBC-gelatin samples

Because the gelatin concentration used in making the RBC suspensions (350 g liter$^{-1}$, w/v; see additional comment on this below) was comparable to that of the hemoglobin inside the cells, we calculated concentrations of ions and solutes on a per-aqueous volume basis as follows.

1) In a sample prepared as described above, 1.75 g of gelatin was dissolved in 5 ml of neutralizing saline. The partial specific volume of gelatin, $v = 0.7417$ ml g$^{-1}$ (46), implied that its volume was $1.75 \times 0.7417 = 1.3$ ml. This, coupled with the volume of neutralizing buffer (5 ml) and RBC suspension (2 ml), inferred a total sample volume of 8.3 ml. Two milliliters of this preparation was drawn into each silicone rubber tube and sealed at the bottom with a Delrin plug.

2) In 2 ml of RBC suspension of $H_t = 0.865$, the aqueous volume is made up of the water in the extracellular space, viz, $2.0 \times (1 - H_t)$ = 0.27 ml, and the water inside the cells. The latter was calculated as $2.0 \times H_t \times \alpha = 1.24$ ml, where $\alpha = 0.717$ is the volume fraction of an iso-osmotic human RBC, that is occupied by water (47).

3) The volume of neutralizing saline was 5.0 ml, so the total extracellular aqueous volume was 5.0 + 0.27 + 1.24 = 6.51 ml.

4) To make the Ca$^{2+}$ concentration of 10 mM in the extracellular space at the start of an experiment, the volume of added 1 M CaCl$_2$ was $53$ ml.

5) With [1,6-$^{13}$C]d-glucose and [6-$^{13}$CH$_3$]-methionine, the amounts added were chosen to give concentrations as if they were averaged over the total aqueous volume of the sample. For example, for the sample used to generate Fig. 1, 11.86 mg of [1,6-$^{13}$C]-glucose yielded a concentration of 9.95 mM in the aqueous space, whereas 5.2 mg of [6-$^{13}$CH$_3$]-methionine gave a concentration of 5.32 mM.

6) The effective $H_t$ across the whole sample volume was 2.0 $\times H_t$ = 0.865/8.3 = 0.208. Therefore, when the amount of [1-$^{13}$C]-lactate was computed during a spectral time course, based on relativity with the peak integral of the known concentration of [6-$^{13}$CH$_3$]-methionine, it was rescaled to give the rate per liter of whole RBCs by dividing by 0.208 (for example, as for Fig. 2).

Calculation of $^{133}$Cs$^+$ concentrations in RBC-gelatin samples

The rationale for estimating the concentration of $^{133}$Cs$^+$ inside the RBCs in a compressed gel experiment was as follows (using the particular numerical values associated with the samples used for Fig. 3 and Fig. S2).

1) A typical $^{133}$Cs$^+$ loading protocol involved suspending 15 ml of saline-washed RBCs (called the RBC sample volume below) of $H_t = 0.8$ (subscript 1 denotes the first $H_t$ measured) in 35 ml (denoted below as “Cs saline volume”) of Cs$^+$-saline: 100 mM CsCl, 44 mM NaCl, 5 mM K$_2$HPO$_4$, 15 mM d-glucose, 10 mM inosine, 0.075 mM penicillin G, and 0.075 mM streptomycin. This gave a total water volume in the incubated sample of 0.8 ($H_t$) $\times 0.717$ [this is the $\alpha$ value or aqueous volume fraction of RBCs (47)] $\times 15$ ml (“RBC sample volume”) + 35 ml (Cs saline volume) = 43.6 ml and $H_t = 0.24$ (subscript 2 denotes the second $H_t$ value).

2) At the start of the incubation, the $^{133}$Cs$^+$ concentration outside the RBCs was 100 mM (denoted below as [Cs$^+$]$_{start}$), and yet, if it was averaged over the whole aqueous volume, it was 30.0 $\times 100/43.6 = 68.8$ mM. After incubation for 4 days at 4°C, or for 2 to 3 days at 21°C with one change of incubation medium, the sample was centrifuged (3000g for 5 min at 10°C) and the pellet plus some residual supernatant, gave the final $H_t = 0.785$ (subscript 3 denotes the third $H_t$ value). A $^{133}$Cs NMR spectrum was recorded from 3 ml of this suspension: In the particular example discussed here, it gave relative peak integrals of intracellular/extracellular = 0.41:0.59.

3) At the final “steady state,” the concentration of $^{133}$Cs$^+$ inside the cells, [Cs$^+$]$_{ss}$, was related to the concentration in the extracellular space, [Cs$^+$]$_{ss}$, through the relative signal intensities $\text{Sig}_{ss}/\text{Sig}_{ss} = \text{Sig}_{ss}/\text{Sig}_{ss}$ and the relative compartment volumes (1 $-$ $H_t$)/$H_t$. Thus,

$$[\text{Cs}^+]_{ss} = \text{Sig}_{ss} (1 - H_t)/H_t [\text{Cs}^+]_{ss} \quad (2)$$

4) The conservation of mass condition for total $^{133}$Cs$^+$ in the incubation suspension was

$$H_t \times \text{RBC sample volume} \times [\text{Cs}^+]_{ss} + ((1 - H_t) \times \text{RBC sample volume} + \text{Cs saline volume}) [\text{Cs}^+]_{ss} = \text{original number of moles of} \ [^{133}\text{Cs}^+] (\text{viz., } 30 \text{ ml } \times 100 \text{ mM} = 0.003 \text{ mol}) \quad (3)$$

5) Rearranging Eq. 3 to obtain an expression for [Cs$^+$]$_{ss}$ in terms of the other parameters and the initial concentration, [Cs$^+$]$_{start}$, we obtained

$$[\text{Cs}^+]_{ss} = \text{Cs saline volume } \times [\text{Cs}^+]_{start} (1 - H_t) \times \text{Sig}_{ss} \quad \text{Cs saline volume } H_t + \text{RBC sample volume} (H_t \times \text{Sig}_{ss} - H_t) (1 + \text{Sig}_{ss}) + H_t \quad (3)$$

and then using the numerical values stated above

$$[\text{Cs}^+]_{ss} = \frac{0.035 \times 0.1 \times (1 - 0.785) \times (0.44/0.59)}{0.035 \times 0.785 + 0.015 \times (0.80(0.44/0.59) - 0.785(1 + 0.44/0.59) + 0.785) = 16.5 \text{ mmol (liter RBC)}^{-1} \quad (4)
Solving for \([Cs^+]^{15}\) gives its concentration as 86.9 mmol [liter extracellular solution]^{-1}.

6) RBCs with this concentration of Cs⁺ were centrifugally washed twice (as described above) in four volumes of 154 mM NaCl containing 15 mM D-glucose, thus removing Cs⁺ from the extracellular medium. Four milliliters of these RBCs (Ht = 0.86) was added to the gelatin solution, as described in the “Gels” section, at 42°C, giving a final Ht of 0.2.

**Calculation of [3-13C]L-lactate concentrations in RBC suspensions**

1) Use of calibration standards: Known amounts of [1,6-13C]D-glucose were added to RBC suspensions to measure the rate of glycolysis in the absence and presence of yoda1. To calibrate the resonance intensities of glucose and lactate to their concentrations, and hence calculate the glycolytic rate, a known amount of [6-13C]L-methionine was added to the sample as the peak intensity reference; its chemical shift was δ = 13.1 ppm. However, to improve the signal-to-noise (S/N) ratio in the NMR spectra, a rapid repetition rate of radio frequency (RF) pulses was used. This led to a steady state of partial saturation of the magnetization of the various spin populations in the sample, and these differed between the solute species. Thus, integrals (and amplitudes) of different peaks from molecules at the same concentration had different values. The following analysis was used to calibrate the methyl resonance intensity of [3-13C]L-lactate to its underlying concentration.

A known amount of pure [3-13C]L-lactate was added to a sample of RBCs in a suspension, or a known amount of [1,6-13C]D-glucose was allowed to be fully metabolized by the RBCs to therefore yield a known amount of [3-13C]L-lactate. A known amount of [6-13C]L-methionine was also added to the sample, having confirmed that its resonance remained at the same intensity throughout many hours of incubation (more than 40 hours at 20°C or 24 hours at 37°C).

The concentration of [3-13C]L-lactate was inferred from the ratio of its integral to that of [6-13C]L-methionine. For a given NMR experiment with its specified relaxation delay(s), RF pulse angle(s), and acquisition time, a known concentration of lactate, \(C_{1\text{Cal}}\), gives a signal \(S_{1\text{Cal}}\)

\[
C_{1\text{Cal}} = \epsilon_1 S_{1\text{Cal}}
\]

where the subscript 1 denotes lactate 13C-methyl, \(\epsilon_1\) is the “extinction coefficient” [mol liter⁻¹ (unit of signal)]⁻¹, and \(S_{1\text{Cal}}\) is the numerical value of the signal. Similarly, for the standard [6-13C]L-methionine (denoted by the subscript 2), we had

\[
C_{2\text{Cal}} = \epsilon_2 S_{2\text{Cal}}
\]

The ratio of the two signals, \(\frac{S_{1\text{Cal}}}{S_{2\text{Cal}}} = \frac{S_{1\text{Cal}}}{S_{2\text{Cal}}}\) (where \(r\) denotes ratio), is the most convenient way to record the data, so the ratio of the extinction coefficients, \(\epsilon_1\), becomes

\[
\frac{\epsilon_1}{\epsilon_2} = \epsilon_r = \frac{C_{1\text{Cal}}}{C_{2\text{Cal}}} S_{1\text{Cal}} = \frac{C_{1\text{Cal}}}{S_{1\text{Cal}}}
\]

Then, the calculated concentration of [3-13C]L-lactate, \(C_1\), in any 13C NMR spectrum in the presence of a known concentration of [6-13C]L-methionine, \(C_2\), is given by

\[
C_1 = \epsilon_r S_r C_2
\]

This analysis was performed on the spectra shown in Fig. 2 and fig. S1 to calculate the effective glycolytic rates.

2) An alternative method to estimate the actual concentration of metabolites in RBC suspensions was to use the conservation of mass condition for 13C-labeled species. A known amount of [1,6-13C]D-glucose was added to the sample, typically 10 mM in 3 ml of RBC suspension, and at any time, all this 13C-label gave resonances effectively only from [1,6-13C]D-glucose, [3-13C]2BPG, or [1,3-13C]L-lactate. Approximately 5% of the labeled metabolites were glycolytic and pentose-phosphate pathway intermediates that were in such low concentrations (32, 48) that they were undetected in the 13C NMR spectra (hence the word “effectively” above). Because progressive saturation affects the resonance intensity differently for each solute, it was necessary to calculate an effective extinction coefficient, \(\epsilon_i\) (where \(i\) is the substrate label/abbreviation), that relates resonance intensity to concentration. This was solved using the conservation of mass (concentration) expression

\[
\frac{1}{\epsilon_{\text{Glc}}} (\text{Sig}_{\text{GlcClβ}} + \text{Sig}_{\text{GlcClα}} + \text{Sig}_{\text{GlcClβ+α}} + \text{Sig}_{\text{GlcClα}}) + \frac{1}{\epsilon_{\text{BPG}}} \text{Sig}_{\text{BPG}} + \frac{1}{\epsilon_{\text{Lact}}} \text{Sig}_{\text{Lact}} = [\text{Glc}]_0
\]

where \(\epsilon_i\) denotes the resonance integral or amplitude (as appropriate), assuming equal extinction coefficients for different atoms and anomers of glucose.

The value of each \(1/\epsilon_i\) was estimated from all, or a subset, of the spectra in a time course such as that shown in Fig. 2 and fig. S1. Thus, the system of simultaneous algebraic equations was overdetermined and was therefore amenable to statistical analysis by using the function LinearModelFit in Mathematica (49). In addition, the concentration of [3-13C]L-lactate was given by \(\frac{1}{\epsilon_{\text{Lact}}} \text{Sig}_{\text{Lact}}\) and so forth for the other species.

**NMR spectroscopy**

13C NMR and 133Cs NMR spectra were recorded at 100.61 and 52.48 MHz, respectively, on a Bruker Avance III spectrometer that has a 9.4-T vertical Oxford Instruments wide-bore magnet using a 10-mm broadband observe probe. Temperature was calibrated (Bruker script) using a sample of neat methanol; for gel-based experiments, it was set to 20°C to ensure persistence of the gel state, with its attendant elastic properties. A simple delay-pulse-acquire RF pulse sequence with CYCLOPS phase cycling was used to obtain the FIDs that were summed and then smoothed by exponential multiplication before Fourier transformation.

**Computing**

All spectra were phased and baseline-corrected in Bruker TopSpin 3.5 before they were imported into Mathematica (49) for postprocessing. The definitive metabolic experiments used [6-13C]L-methionine as the internal intensity reference. Even for this compound, the calibration of peak intensities was necessary (see above) to account for progressive saturation in the rapid repetition spectral acquisition that was used to obtain the highest possible S/N ratio in a given time.

Progress curves of NMR spectral intensity were fitted by straight lines or exponential curves using the Mathematica (49) function
Kuchel and Shishmarev, fig. S2. Efflux of Cs+ from human RBCs suspended in gelatin gel. Supplementary material for this article is available at http://advances.sciencemag.org/cgi/.

**SUPPLEMENTARY MATERIALS**

Supplementary material for this article is available at http://advances.sciencemag.org/cgi/.

**REFERENCES AND NOTES**

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