JCI The Journal of Clinical Investigation

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H Izumo, S Lear, M Williams, R Rosa, F H Epstein

J Clin Invest. 1987;79(6):1621-1628. https://doi.org/10.1172/JCI112998.

Research Article

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Sodium-Potassium Pump, Ion Fluxes, and Cellular Dehydration in Sickle Cell Anemia

Hiroko Izumo, Stephanie Lear, Mark Williams, Robert Rosa, and Franklin H. Epstein

Charles A. Dana Research Institute, Harvard-Thorndike Laboratory of Beth Israel Hospital, Department of Medicine, Harvard Medical School and Beth Israel Hospital, Boston, Massachusetts 02215

Abstract

We studied the role of the sodium-potassium pump in erythrocytes of 12 patients with sickle cell anemia (SS). Ouabain-binding sites per cell and pump-mediated Rb/K uptake were significantly higher in SS patients than in white or black controls. Ouabain-resistant Rb/K influx was also greater than in normal controls or patients with sickle cell trait.

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These findings indicate a magnified role for the sodium-potassium pump in the pathophysiology of SS erythrocytes and suggest that its inhibition might prove useful in therapy.

Introduction

The genetic defect in sickle cell anemia (SS)¹ results in the synthesis of an abnormal hemoglobin, hemoglobin S (Hb S), which upon deoxygenation polymerizes into a gel state. This gelation process produces the characteristic sickle deformation in red cells and predisposes to microcirculatory occlusion and subsequent ischemic damage. The pathogenesis of sickling depends strongly on the intracellular concentration of deoxyhemoglobin S (1, 2). Sickling is therefore enhanced by dehydration of erythrocytes, which produces a rise in mean corpuscular hemoglobin concentration (MCHC).

In addition to the abnormality in hemoglobin, SS erythrocytes exhibit changes in membrane permeability, possibly owing to alterations in the membrane cytoskeleton (3, 4). It has been well documented that deoxygenated sickle erythrocytes exhibit increased Na and K fluxes, with net cellular Na gain and K loss (5-10). However, the effect of such changes on red cell water content and MCHC is still controversial. The contributions of the various components of ion flux during sickling, such as the sodium-potassium pump and the Na-K-Cl cotransport pathway,

Presented in part at the American Federation for Clinical Research, May 1985. (1985. Clin. Res. 33:343a.)

Received for publication 4 November 1986.

1. Abbreviations used in this paper: AA, normal erythrocyte; AS, sickle cell trait; BSKG, phosphate-buffered saline with potassium and glucose; DDC, density distribution of cells; Hb S, hemoglobin S; MCHC, mean corpuscular hemoglobin concentration; SS, sickle cell anemia.

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have not been fully elucidated, owing in part to methodological variation and to the heterogeneity of the red cell population in sickle cell blood.

We have evaluated the sodium-potassium pump in sickle erythrocytes by means of an ouabain-binding assay and ⁸⁶Rb uptake and have explored the relation of pump inhibition to cell volume. The activity of the sodium-potassium pump is increased above normal in oxygenated SS cells and further increased by deoxygenation. In deoxygenated cells, inhibition of the pump by ouabain results in a reduction of MCHC and the proportion of dense cells, consistent with cell swelling. This suggests a possible role for cardiac glycosides in anti-sickling therapy.

Methods

Subjects. Venous blood was obtained from 12 patients with SS, 6 with sickle cell trait (AS), and 40 white and 10 black normotensive control subjects, aged 15-45 yr, none of whom were taking cardiac glycosides or diuretics or had recently received blood transfusions. The blood was then analyzed by the following tests.

Ouabain binding and Rb/K uptake. 10 ml of blood was drawn from each subject into a heparinized syringe for the measurement of [³H]ouabain binding and ⁸⁶Rb uptake by previously described techniques (11). The plasma and buffy coat were separated by centrifugation and discarded; the remaining erythrocytes were washed three times with 10 vol of MgCl₂ solution (112 mM). After a final wash in ouabain-binding buffer, containing in mM: NaCl, 140; CaCl₂, 1; MgCl₂, 1; Hepes, 20; dextrose, 5%; pH 7.40, red cells were suspended in the same buffer at a hematocrit of 5–10%.

To measure ouabain binding to erythrocytes, triplicate 200-μl aliquots of the red cell suspension were added to 50 μ l of a mixture of 0.5-1.0 pmol of [3H]ouabain (New England Nuclear, Boston, MA; sp act, 17-18 Ci/mM) and unlabeled ouabain, so that the total concentration of the ligand ranged from 80 to 120 nM. This concentration of ouabain has been shown to saturate the specific-binding sites of human erythrocytes when incubated at 37°C for 1 h (11). In a single pilot experiment, SS erythrocytes incubated with the same concentration of radiolabeled ouabain also displayed saturation kinetics by 1 h at 37°C. Nonspecific binding of [3H]ouabain was measured in the presence of 0.1 mM unlabeled ouabain. After a 60-min incubation at 37°C in a shaking water bath, the cells were washed three times in 1-ml volumes of ice-cold MgCl₂, after which 200 µl of 5% TCA were added and the mixture was centrifuged for 15 min at 10,000 g. The clear supernatant containing the released radioactivity was then drained and counted in 5 ml of Biofluor in a liquid scintillation spectrometer.

Potassium influx was estimated by measuring the uptake of ⁸⁶Rb into erythrocytes (Rb/K influx). Previous studies indicate that ⁸⁶Rb can be used as an analog of potassium in human erythrocytes (12–14). After the red cells were washed with MgCl₂ and suspended in ouabain-binding buffer, 4 mM KCl, and ~ 10⁵ cpm of the K analog ⁸⁶RbCl (New England Nuclear; sp act, 1.0 mCi/mg), was added to each tube. The Na,K-ATPasemediated uptake of ⁸⁶Rb was calculated as the difference between total radioactivity taken up by triplicate aliquots with and without 0.1 mM ouabain. After a 10-min incubation at 37°C in a shaking water bath, the cells were washed with ice-cold MgCl₂. Intracellular radioactivity was released by 5% TCA and counted in the supernatant after centrif-

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ugation for 15 min at 10,000 g. The results were expressed as nanomoles of $K^+/10^9$ red blood cells per hour, using the assumption that ^{86}Rb is a tracer for the influx of stable K^+ , present at a concentration of 4 mM in the suspending medium.

The portion of ⁸⁶Rb uptake that is insensitive to ouabain and inhibited by furosemide is mediated through Na,K-cotransport present in human red blood cells (15, 16). To evaluate Na,K-pump-independent Rb/K uptake further, triplicate samples of red cells were incubated with 0.1 mM ouabain alone, or both ouabain and 1 mM furosemide, under experimental conditions analogous to those described above.

Intracellular electrolyte concentrations. In the course of the foregoing experiments, red cells from heparinized venous blood were separated from plasma and buffy coat by centrifugation and washed three times with an isosmotic solution of 112 mM MgCl₂. Before ouabain-binding and Rb/K uptake studies, 400- μ l aliquots of packed erythrocytes were removed and recentrifuged at 10,000 g to remove the remaining supernatant. A 200- μ l aliquot of packed cells was transferred in a wide-bore pipette to 9.8 ml of deionized water to induce hemolysis. Na and K concentrations were measured in triplicate by flame photometry, after centrifugation to pellet cell membrane fragments.

Ouabain-binding and Rb/K uptake after nitrogen incubation. Incubation of SS blood under nitrogen was used as a means of deoxygenating red cells, thereby inducing sickling. 30 ml of heparinized venous blood were required for this study. The erythrocytes were separated, washed, and resuspended as described earlier. After aliquots of the red cell suspension in ouabain-binding buffer were removed for baseline studies of ouabain binding, Rb/K uptake, and intracellular electrolyte concentration, the remainder of the suspension, at a hematocrit of 10%, was divided into two portions for deoxygenation. In one of these, destined for subsequent measurement of intracellular electrolytes and Rb/K uptake, KCl was added to produce a final concentration in the medium of 4 mM. Both were then exposed for 2 h in a shaking water bath at 37°C to 100% nitrogen humidified by bubbling through an isotonic solution at 37°C. Deoxygenation was confirmed by measuring Po2 with a blood gas analyzer $(P_{O_2} < 18 \text{ mmHg})$. The pH of the suspension was stable during the incubation. After 2 h of nitrogen incubation, the cell suspension was studied by ouabain binding and Rb/K assay techniques; particular care was taken to expose the red cells continuously to nitrogen during these procedures. Intracellular electrolyte concentrations were measured after deoxygenation as described above.

Separation of erythrocytes by density gradients. In a separate study, 20 ml of heparinized blood was placed in a mixture of 9% Ficoll and 33.9% Hypaque and centrifuged to separate red cells. After centrifugation, erythrocytes were resuspended in phosphate-buffered saline with K and glucose (BSKG; in mM: NaCl, 134; Na₂HPO₄, 16; NaH₂PO₄, 4; KCl, 5; glucose, 11; osmolarity, 295 mOsm; pH 7.4) to a hematocrit of 20%. Stractan gradients were prepared according to the method described by Corash et al. (17) with minor modifications (18). Stractan II (arabinogalactan) was passed through both cation and anion exchange resins. Stractan was then added to a solution containing (per 100 ml Stractan): bovine serum albumin, 3 g; glucose, 100 mg; KH₂PO₄, 2.85 mM; K₂HPO₄, 12.15 mM; MgCl₂, 112 mM. The pH and osmolality of the Stractan solution were adjusted to 7.4 and 291 mOsm. The Stractan solution was then diluted with BSKG to achieve the desired densities. For control blood, three layers of Stractan with densities of 1.110, 1.101, and 1.092 were used, whereas denser Stractan solutions were required for the blood of sickle cell patients because of the presence of heavier cells (densities of 1.150, 1.110, and 1.101 were used). Red cells suspended in BSKG were layered on top of Stractan gradients and centrifuged at 4°C at 52,000 g for 45 min. Three fractions of red cells, top, middle, and bottom, were collected and washed separately three times with MgCl₂ and once with ouabain-binding buffer. For each fraction, reticulocytes were counted in smears stained with methylene blue. Sickled forms were counted unstained. Ouabain-binding and Rb/K uptake assays were then performed on each fraction as described above.

Density distribution of cells (DDC). The DDC technique has been used in various erythrocyte disorders to determine the distribution of cells by relative density. Normal erythrocytes exhibit a narrow range of

densities. However, the DDC of SS blood reflects the heterogeneity of the red cell population including denser sickled cells, as well as less dense, younger cells containing a high proportion of reticulocytes (19).

The method, as described by Danon and Marikovsky (20), requires two phthalate esters as separating fluid, methyl phthalate (specific gravity, 1.189), and di-n-butyl phthalate (specific gravity, 1.0416), which were used to prepare a series of solutions of 20 different specific gravities, ranging from 1.062 to 1.138, in increments of 0.004. 10 ml of heparinized blood were drawn for use in these experiments. One microhematocrit tube was dipped into each of the different solutions until a column of slightly more than 5 mm was obtained. The remainder of the tube was then filled with heparinized venous blood, sealed, and spun at 12,000 g for 15 min in a refrigerated centrifuge. A set of 20 capillary tubes for each blood sample was then arrayed in order of decreasing specific gravity of the separating liquids. The results were plotted with decreasing specific gravity of the phthalate solution on the abscissa and the percentage of red cells that had passed through the separating liquid on the ordinate.

MCHC. The calculation of MCHC is used as an approximation of cell water content and is based upon measured hematocrit and hemoglobin, according to the equation: MCHC = Hb (g/dl)/packed cell volume (%) \times 100. Packed cell volume was determined by high-speed centrifugation of blood in a heparinized microhematocrit capillary tube for 6 min to minimize the potential for excess plasma trapping by sickle erythrocytes. Hemoglobin was measured spectrophotometrically by the cyanohemoglobin method, which includes all hemoglobin and hemoglobin derivatives except possibly sulfhemoglobin, using 5 ml of Drabkin's reagent for 0.02 ml of venous blood (21).

To evaluate the effects of deoxygenation on MCHC and DDC, 10 ml of heparinized blood was gassed in an Erlenmeyer flask in a shaking water bath at 37°C for 2 h with either air or nitrogen with 5% CO₂.

Statistics. Results are expressed as mean \pm SE. Comparison of two samples was done by Student's t test. Multiple samples were compared by analysis of variance and the Newman-Keuls test.

Results

Na,K transport characteristics (Table I). Binding of [³H]ouabain to human red blood cells is a high-affinity saturable process; the receptor for [³H]ouabain in the erythrocyte, as in other cells, is the Na-K-ATPase pump. In these experiments, the number of [³H]ouabain-binding sites is determined at a saturating concentration of the ligand, at which Na,K pump-mediated ion transport is completely inhibited (11).

The mean value of ouabain binding was significantly higher in sickle cell patients $(0.81\pm0.05 \text{ pmol}/10^9 \text{ cells})$ than in white controls $(0.47\pm0.02; P < 0.001)$. Black controls had an even lower value for ouabain binding (0.38 ± 0.02) than white controls. Patients with sickle cell trait had a value (0.38 ± 0.04) that did not differ significantly from that of either control group.

The mean ouabain-sensitive Rb/K uptake was significantly higher in sickle cell patients (189.6 ± 15.8 nmol/ 10^9 cells/h) than in white controls (124.8 ± 4.1 ; P < 0.001). As in the ouabain-binding assay, this value was even lower in black controls (96.9 ± 5.6) than in whites. There were no significant differences between patients with sickle trait and either control group. The ouabain-insensitive Rb/K uptake was four to five times higher in red cells of patients with SS than in either white or black controls (P < 0.001). Thus, sickle cell patients have a greater number of Na,K pump units as well as increased pump activity, as measured by ouabain-sensitive Rb/K uptake. In addition, ouabain-insensitive Rb/K uptake is increased in sickle cells as compared with normal erythrocytes.

The major portion of pump-independent ouabain-insensitive Rb/K uptake can be inhibited by furosemide and its derivatives and represents Na/K/Cl cotransport (22, 23). Furosemide-sen-

Table I. Studies of Ouabain-binding Rb/K Uptake and Intracellular Electrolyte Concentrations of Erythrocytes

	Ouabain binding	Ouabain-sensitive Rb/K uptake	Ouabain-insensitive Rb/K uptake	Intracellular Na concentration	Intracellular K concentration
	pmol/10° cells	nmol/10° cells/h		mEq/liter cells	mEq/liter cells
White control	0.47±0.02	124.8±4.1	50.3±2.0	5.9±0.3	88.1±2.1
	(n = 40)	(n = 40)	(n = 40)	(n = 21)	(n = 21)
Black control	0.38±0.02*	96.9±5.6*	39.1±7.3*	7.8±0.7*	81.9±4.5
	(n = 10)	(n = 10)	(n=6)	(n = 10)	(n = 10)
SS	0.81±0.05 [‡]	189.6±15.8‡	225.6±16.9‡	19.0±1.6‡	60.0±2.9‡
	(n = 12)	(n = 12)	(n = 12)	(n = 12)	(n = 12)
AS	0.38±0.04	107.3±8.5	77.7±13.4	8.1±0.8*	76.9±2.8
	(n = 6)	(n = 6)	(n=6)	(n = 6)	(n = 6)

Values are mean \pm SE. * Significantly different from white controls (P < 0.01). \pm Significantly different from all other groups (P < 0.01).

sitive Rb/K uptake, measured in the presence of 10^{-4} M ouabain, was significantly greater in erythrocytes of patients with SS (129.2±12.3) than those of in controls (40.6±3.4; P < 0.001).

Intracellular electrolytes (Table I). Intracellular sodium concentration was significantly higher in erythrocytes of sickle cell patients (19.0 \pm 1.6 mEq/liter cells) than in those of any control group (P < 0.001). Cell sodium was significantly higher in normal black subjects (7.8 \pm 0.7) and persons with sickle trait (8.1 \pm 0.8) than in white controls (5.9 \pm 0.3) (P < 0.01). Intracellular concentration of potassium was significantly lower in SS erythrocytes than in the red cells of any of the control populations (P < 0.001).

Na,K transport characteristics of erythrocytes separated by density gradients (Tables II and III). It is known that reticulocytes, which are young red cells, have enhanced active and passive transport mechanisms for sodium and potassium as compared with mature red cells (24-28). Reticulocytes constitute the lightest fraction of the red cell population, whereas sickled erythrocytes are the heaviest cells. To evaluate the contribution of reticulocytes to the increase in ouabain binding and Rb/K uptake in the sickle cell population, erythrocytes were separated by Stractan solutions according to cell density. The ouabain binding and Rb/K uptake assays were then performed on each of the three cell fractions. Different densities of Stractan were used for sickle cell blood and for control blood because of the larger proportion of heavier cells in SS blood (see Methods). The cells of the middle fraction in sickle cell blood correspond in density to the cells of the bottom fraction in control blood because of the different densities of Stractan used.

The sodium and potassium transport parameters for control

(AA) and SS red cells are summarized in Tables II and III, respectively. In SS blood, there was a greater percentage of reticulocytes, as expected, pooled in the top fraction where significantly greater ouabain binding and pump-mediated Rb/K uptake was found than in the middle or bottom fractions (P < 0.01). In contrast, there was no difference in transport characteristics among the top, middle, and bottom fractions of control red cells.

The heaviest fraction of sickle cell blood (with only $1.4\pm0.36\%$ reticulocytes) had a ouabain-binding capacity that was not significantly different from that of the lightest layer of control blood (with $2.8\pm0.43\%$ reticulocytes) but a ouabain-sensitive Rb/K uptake that was much higher than any of the three fractions of control blood (P < 0.01). To analyze further the nature of pump-mediated Rb/K flux, the ouabain-inhibited Rb/K influx per ouabain-binding site, which represents the pump activity per pump unit, was calculated. In SS blood, this value was significantly higher for the bottom fraction of cells (with more sickled erythrocytes) than the top fraction (with more reticulocytes) (P < 0.01), implying an increased rate of turnover of each pump unit in the heaviest cells.

The effect of deoxygenation on cation transport of sickle erythrocytes (Table IV). In control erythrocytes, there was no significant change in any of the transport parameters after nitrogen incubation. In SS cells, ouabain binding, although elevated before deoxygenation $(0.69\pm0.04~\mathrm{pmol/10^9}~\mathrm{cells})$, did not change after nitrogen incubation (0.65 ± 0.05) . However, deoxygenation caused a doubling of ouabain-sensitive Rb/K uptake $(P < 0.01~\mathrm{as}~\mathrm{compared}~\mathrm{with}~\mathrm{the}~\mathrm{pre-N_2}~\mathrm{value})$. This indicates that while

Table II. Na,K Transport Characteristics of AA Erythrocytes Separated by Stractan Gradients

Fraction	Density	Reticulocyte	Ouabain binding	Ouabain-sensitive Rb/K uptake	$\frac{\text{Rb/K uptake}}{\text{Ouabain-binding}} \times 10^{-3}$	Ouabain-insensitive Rb/K uptake
	specific gravity	%	pmol/10° cells	nmol/10° cells/h	h-1	nmol/10° cells/h
Тор	1.092	2.8±0.43	0.505±0.026	130.0±6.5	0.261±0.020	62.9±5.5
Middle	1.092-1.101	0.55±0.22	0.464±0.025	139.8±4.6	0.311±0.015	72.0±9.1
Bottom	1.101-1.110	0.029±0.028	0.458±0.032	132.8±10.5	0.294±0.029	61.0±12.5

n = 7. Values are mean \pm SE. Red cells suspended in BSKG at a hematocrit of 20% were layered on top of Stractan gradients of the specific gravities indicated in the table and centrifuged at 4°C for 45 min at 52,000 g. Three fractions of red cells were collected separately and washed with MgCl₂ and ouabain-binding buffer before determining their reticulocyte count, ouabain binding, and Rb/K uptake.

Table III. Na,K Transport Characteristics of SS Erythrocytes Separated by Stractan Gradients

Fraction	Density	Reticulocyte (%)	Sickled cells (%)	Ouabain binding	Ouabain-sensitive Rb/K uptake	Ouabain-sensitive Rb/K uptake/ ouabain-binding \times 10 ⁻³	Ouabain-insensitive Rb/K uptake
	specific gravity	%	%	pmol/10° cells	nmol/10° cells/h	h ⁻¹	nmol/10° cells/h
Top	1.092-1.101	15.2±4.15	0.12±0.06	1.153±0.132	311.6±28.8	0.287±0.030	384.2±62.9
Middle	1.101-1.110	4.7±1.24	1.43±0.45	0.755±0.066	260.1±33.5	0.346±0.037	172.2±38.3
Bottom	1.110-1.15	1.4±0.36	26.29±6.01	0.584±0.070	217.4±20.0	0.427±0.050	182.1±28.2

n = 10. Values are mean ±SE. Red cells suspended in BSKG at a hematocrit of 20% were layered on top of Stractan gradients of the specific gravities indicated in the table and centrifuged at 4°C for 45 min at 52,000 g. Three fractions of red cells were collected separately and washed with MgCl₂ and ouabain-binding buffer before determining their reticulocyte count, ouabain binding, and Rb/K uptake.

there was no change in the number of pump units, there was a substantial increase in pump activity associated with the sickling process. On the other hand, ouabain-insensitive Rb/K uptake decreased by an average of 42% after nitrogen incubation, from 198.7 ± 90.4 nM/ 10^9 cells/h to 116.7 ± 13.2 (P < 0.01 with paired comparison). There was a similar decrease of $\sim 35\%$ in furosemide-sensitive Rb/K uptake with deoxygenation (150.8 ± 20.6 to 98.8 ± 10.7).

After nitrogen incubation, intracellular sodium concentration in sickle erythrocytes increased significantly (12.7 \pm 1.8 to 21.2 \pm 2.1 mEq/liter cells) (P < 0.01), and intracellular potassium concentration tended to decrease (61.9 \pm 9.3 to 56.9 \pm 5.5), although the latter change did not reach the level of significance.

Effect of deoxygenation and ouabain on the density distribution of cells (Figs. 1-4). The density distribution of cells was plotted for 16 sickle cell patients and seven control subjects. Normal (AA) erythrocytes, as depicted in Fig. 1, display a narrow range of density distribution. The blood of patients with sickle cell anemia, however, may exhibit a wide range of densities, with more high-density cells representing sickled cells and more low-density cells representing reticulocytes (Fig. 1). The DDC of blood from three patients with AS displayed the same configuration as did AA (data not shown).

Density distribution curves were plotted after O_2 and N_2 to examine the effect of deoxygenation on cell density. With deoxygenation of SS blood, the curve was shifted to the left by the increased percentage of heavier cells and very high density cells which represent sickled erythrocytes (Fig. 2). The percentage of cells heavier than the median-specific gravity in air was increased from 50 to $72\pm3\%$ by deoxygenation (n=9). Incubation in O_2 did not alter the density distribution of normal or SS cells.

To evaluate the effect of inhibiting the sodium-potassium pump on the sickling process, we examined the density distribution of SS cells under nitrogen incubation in the absence and presence of 0.1 mM ouabain. In each of eight experiments, ouabain reduced the percentage of very dense cells that appeared with nitrogen exposure (Fig. 3). The percentage of cells heavier than the median-specific gravity in nitrogen alone was reduced by ouabain from 50 to $25\pm6\%$. Little or no effect was seen when ouabain was added to oxygenated SS cells (Fig. 4).

Effect of deoxygenation and ouabain on MCHC (Fig. 5). Hemoglobin and hematocrit were measured before and after nitrogen incubation. The MCHC of SS cells with deoxygenation increased in 11 of 12 cases by an average of $5.5\pm0.9\%$, from 33.7 ± 0.6 to 35.3 ± 0.9 (P<0.01), reflecting shrinkage with sickling. MCHC of control erythrocytes did not change when they were deoxygenated. Ouabain, 10^{-4} M, prevented the increase in MCHC caused by deoxygenation of SS cells. Ouabain had no significant effect on MCHC in the control population.

Discussion

Abnormalities of cation transport have been described in association with sickling but the role of the sodium-potassium pump and associated changes in red cell water content have not been firmly established. Tosteson, Shea, and Darling first reported in 1952 (5) that oxygenated erythrocytes from patients with SS had slightly higher intracellular concentrations of sodium and lower concentrations of potassium than normal red blood cells. Incubation of SS cells in nitrogen, so as to produce sickling, resulted in a further gain in sodium and loss of potassium, whereas anoxia did not affect the cation content of normal erythrocytes. Follow-

Table IV. Effect of Deoxygenation on Na,K Transport Characteristics and Intracellular Electrolyte Concentrations of Erythrocytes

	Ouabain binding	Ouabain-sensitive Rb/K uptake	Ouabain-insensitive Rb/K uptake	Furosemide-sensitive Rb/K uptake	Intracellular Na concentration	Intracellular K concentration
	pmol/10° cells	nmol/10° cells/h	nmol/10° cells/h	nmol/10° cells/h	mEq/liter cells	mEq/liter cells
Control (AA), $n = 7$						
Pre-N ₂	0.42±0.02	112.9±7.7	54.0±5.2	50.14±5.00	6.4±0.9	84.2±2.3
Post-N ₂	0.42 ± 0.02	127.5±11.9	57.4±7.9	48.3±3.08	6.6±0.8	82.1±1.8
Sickle cells (SS), $n = 11$						
Pre-N ₂	0.69±0.04	150.5±9.3	198.7±90.4	150.83±20.6	12.7±1.8	61.87±9.32
Post-N ₂	0.65±0.05	303.0±19.1*	116.7±13.2*	98.8±10.7*	21.2±2.1*	56.98±5.53

Values are mean \pm SE. * P < 0.01 as compared with pre-N₂ values. Red cells suspended in Hepes buffer at pH 7.4 were deoxygenated by exposure to nitrogen in a shaking water bath at a temperature of 37°C for 2 h.

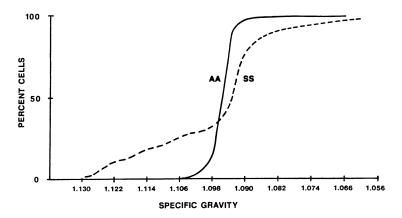
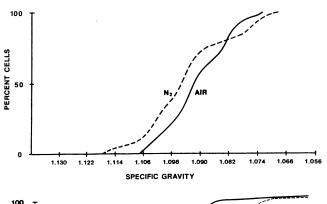
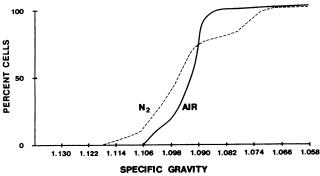


Figure 1. Density distribution of red cells from a representative sample of AA (solid line) and of SS (dashed line) blood under aerated conditions. Specific gravity of the phthalate ester is plotted on the abscissa and the percentage of cells heavier than each phthalate reference is plotted on the ordinate. Curves were drawn by eye through the data points obtained at intervals of 0.004 specific gravity. Normal erythrocytes display a narrow range of densities, whereas SS erythrocytes are distributed over a wider range.

up studies of ion fluxes with isotopic tracers showed that sickling, again induced by exposure to nitrogen, greatly accelerated bidirectional fluxes of sodium and potassium in SS erythrocytes and increased lactate production, implying active transport (6, 7). Because these experiments were carried out just before the description of the Na-K-ATPase pump and its inhibition by ouabain in human red blood cells, the contribution of Na-K-ATPase-mediated ion fluxes to these phenomena was not specifically examined. Subsequently, the potassium efflux from deoxygenated red cells containing Hb-S has often been used as an in vitro measure of red cell sickling, though it is now known to vary with pH (9). Clark, Morrison, and Shohet (18) reported both active and passive Rb/K influx to be within the normal range in oxygenated SS cells, though the passive efflux rate of internal potassium was increased. Their experiments, it should be noted, were carried out on red cells that had been loaded with sodium by overnight storage at 4°C, a procedure that would tend to minimize any difference in ouabain-sensitive K influx between SS and AA cells if a difference in internal sodium contributed to this. They did not assess the effect of sickling induced by deoxygenation. They noted an increase in Na-K-ATPase activity in sickle cell membranes, especially in those separated cell fractions rich in reticulocytes, and ascribed the increase in enzymatic activity to the increased percentage of young erythrocytes, a conclusion reached also by Luthra and Sears (29). Berkowitz and Orringer recently reinvestigated the passive fluxes of Na⁺ and Rb⁺ in SS erythrocytes exposed to ouabain to eliminate active transport. While no difference was found in the influx of Na⁺ or Rb⁺ between oxygenated sickle cells and those of normal subjects, deoxygenation of sickle cells produced a substantial increase in the passive influx of Na⁺ and Rb⁺ that was not blocked by furosemide (30).

The effect of deoxygenation on the volume of SS red cells has been a subject of controversy, probably because of the varied experimental conditions employed by different investigators. Deoxygenation of sickle cell blood in vivo or in vitro produces a fall in the water content of erythrocytes (5, 31, 32) but this has not always been observed (4, 6), especially when red cells have been separated and suspended in artificial media, from which calcium or magnesium may be omitted (30, 32, 33). It is now clear that the effect of sickling on red cell volume is strongly influenced by pH (9, 32) in that red cell shrinkage is retarded





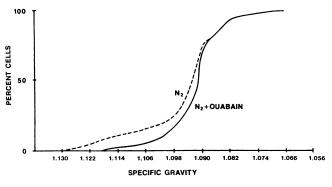
N₂/ AIR

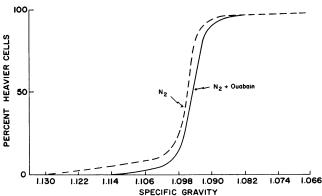
N₂/ AIR

1.130 1.122 1.114 1.106 1.098 1.090 1.082 1.074 1.066 1.05

SPECIFIC GRAVITY

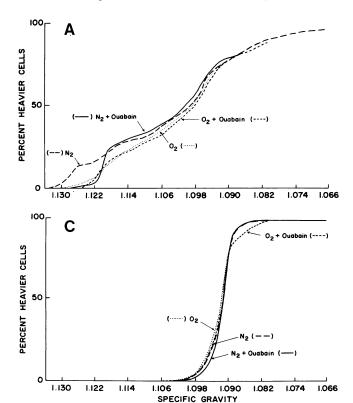
Figure 2. Density distribution of SS cells, aerated (solid line) and after nitrogen incubation (dashed line) from three representative patient studies. Specific gravity of the phthalate ester is plotted on the abscissa. The shift to the left with deoxygenation represents a larger percentage of heavier cells.





by pH greater than 7.4 and accelerated by acidic pH. Sodium and potassium transport across red cell membranes is also affected by the presence or absence of divalent cations in the suspending medium (34).

The present studies demonstrate clearcut differences in transport characteristics between sickle cell erythrocytes and those of control patients, both related to and independent of the



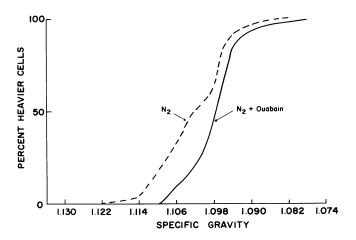


Figure 3. The effect of ouabain on the density distribution of SS cells deoxygenated by exposure to nitrogen for 2 h. Three representative studies are shown. Incubation with 0.1 mM ouabain during nitrogen exposure results in fewer of the heaviest (i.e., sickled) cells than with deoxygenation alone.

Na-K pump. Firstly, there are significantly more ouabain-binding sites in freshly drawn, aerated sickle red cells than in normal erythrocytes, together with an increase in Na-K-ATPase-mediated Rb/K uptake. When cells are separated on density gradients, the fraction of SS cells containing the most reticulocytes also contains the highest number of ouabain-binding sites, suggesting that the increase in Na-K pump units in sickle cell blood

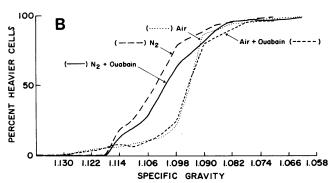
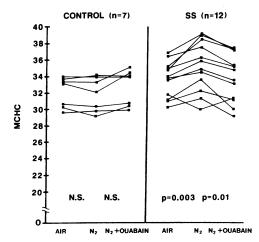


Figure 4. The effect of ouabain on the density distribution of SS cells exposed to nitrogen for 2 h is contrasted in (A) and (B) with its effect on aerated cells in two representative patients. (C) illustrates the lack of effect of ouabain on oxygenated or deoxygenated cells in a control patient with SA. Results similar to this were obtained with AA erythrocytes.



MEAN ±SE 32.2±0.7 31.9±0.8 32.7±0.8 33.7±0.6 35.3±0.9 34.1±0.8

Figure 5. The effect of nitrogen incubation and ouabain on MCHC of control and SS erythrocytes. MCHC is expressed as grams per

is at least in part due to the large population of young red cells. These changes appear even more pronounced when a comparison is made with normal black control patients rather than with Caucasian controls, because black control subjects had diminished values of ouabain binding and Rb/K uptake as compared with whites. The average levels of ouabain binding and Rb/K uptake in SS erythrocytes was about twice those of red cells in normal black controls. These findings are consistent with those of Lasker and associates, who described a significantly decreased activity of Na-K-ATPase in erythrocytes of black subjects as compared with those of whites (35). The average red cell sodium content is significantly greater in normotensive blacks with AA hemoglobin than in their white counterparts (35-37). The present data suggest that this is the consequence of a reduction in the number of Na/K pump units per cell. In SS erythrocytes, on the other hand, there appears to be an increase in sodium-potassium pump activity owing to two factors: an increase in ouabain-binding sites, largely due to the younger population of cells, and an increase in cell sodium. The combination of an increase in pump activity, together with an increase in cell sodium and a decrease in cell potassium, is consistent with an increased passive leak of Na+ and K+ in the red cells of patients with sickle

The influx of Rb/K mediated by pump-independent, ouabain-resistant mechanisms is also substantially increased in aerated erythrocytes from sickle cell patients. About two-thirds of this flux could be inhibited by furosemide, implying that it was mediated by Na-K-Cl cotransport. In our experiments, ouabainresistant Rb/K flux decreased when SS cells were deoxygenated. These findings differ from those of Berkowitz and Orringer, who reported no difference in "passive" 86Rb influx between oxygenated red cells from sickle cell patients and normals, though in SS cells this parameter increased when cells were deoxygenated (30). Surprisingly, they found no evidence of Na-K-Cl cotransport in either oxygenated or deoxygenated SS cells. The reasons for these differences in results deserve further investigation.

The major finding of the present experiments is that the increase in Rb/K influx produced by deoxygenation of sickle cells was entirely accounted for by marked acceleration of the Na-K-ATPase pump. The increase in ouabain-inhibitable Rb/ K influx seen in SS cells exposed to nitrogen was, in fact, greater

in every case than the increase in total Rb/K influx, so that when exposed to nitrogen, pump-independent uptake of 86Rb decreased, including that portion sensitive to furosemide. The number of pump units per cell, as assessed by ouabain binding, did not change. The rise in internal concentrations of sodium seen in deoxygenated sickle cells is best interpreted as a consequence of the increase in passive influx of sodium first noted by Tosteson (7). This presumably stimulated the turnover of Na-K-ATPase, as reflected in the ratio of Rb/K uptake to ouabain binding. At the same time, the reduction in sodium gradient across the red cell membrane would be expected to reduce the driving force for coupled entry of rubidium or potassium via the furosemide-sensitive Na-K-Cl transporter. The result of the deoxygenation of sickle red cells is therefore a massive increase, in absolute as well as proportionate terms, in ion transport driven by Na-K-ATPase. Because the pump extrudes 3 Na⁺ ions for every 2 K⁺ that enter the cell, the increase in pump-mediated transport activity tends to reduce cell volume. A dehydrating effect of the red cell Na⁺/K⁺ pump was demonstrated by Clark and her associates in normal red blood cells that were loaded with sodium by exposing them to nystatin (38). These authors also suggested that the pump might contribute to cell shrinkage in sickle cells in which intracellular sodium had been increased by deoxygenation. Inhibition of Na-K-ATPase in sickle cells might therefore be expected to increase cell volume, thereby reducing mean corpuscular hemoglobin concentration.

These predictions were borne out in the present experiments in which samples of sickle cell blood were deoxygenated by exposure to nitrogen. When sickle cells were deoxygenated, their MCHC increased, in 11 of 12 instances, by an average of 5.5%. The addition of ouabain to SS cells under nitrogen prevented cell shrinkage and caused a decrease in MCHC. Density distribution studies were confirmatory. Nitrogen incubation increased the percentage of high-density cells in SS blood, and ouabain tended to reverse this change, reducing the number of highdensity cells.

These findings have an important implication for therapeutic strategies in sickle cell anemia. The predominance of pumpmediated active transport in SS erythrocytes and its magnification when these cells are deoxygenated suggest the possibility that inhibiting the Na-K-ATPase pump with cardiac glycosides might be useful in patients with sickle cell anemia by causing some degree of red cell swelling and thereby reducing the concentration of deoxyhemoglobin S. Ample evidence, both in vitro and in vivo, indicates that an increase in cell water reduces the tendency for SS cells to sickle (39-42) and thus may ameliorate certain clinical manifestations of sickle cell anemia (42). Furthermore, concentrations of cardiac glycosides that are nontoxic and can be achieved by therapeutic dosages of these drugs have been shown in subjects without sickle cell anemia to exert an inhibitory action on red cell Na-K-ATPase associated with an increase in red cell sodium and a fall in red cell potassium (43, 44) and, over several days, a small decrease in MCHC (45). Even a small decrease in the cellular concentration of deoxyhemoglobin S might exert a salutary effect in patients with sickle cell anemia because of the exponential relationship between MCHC and the delay time for gelation of sickle hemoglobin (1).

Acknowledgments

This research was aided by United States Public Health Service grants HL-15157, AM-18078, and RR-01032.

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