The Effect of Supporting Electrolyte on Ion Permeability Through Red Blood Cell Membrane

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Abstract: This study was aimed at providing information on how Na sulphate, K sulphate, Ca sulphate and Mg sulphate supporting electrolytes affect the permeation of hypotonic Na chloride, K chloride, Ca chloride and Mg chloride ions through Red Blood Cell (RBC) membrane while the RBCs haemolyse using a UV/Vis spectrophotometer. The effects of the supporting electrolytes were deduced from the changes in their respective rates of permeation through RBC membranes during haemolysis. Na₂SO₄ added to NaCl and MgSO₄ to MgCl₂ resulted in a decrease in permeation rate of the resultant solutions while the addition of K₂SO₄ to KCl and the addition of CaSO₄ to CaCl₂ resulted in an increase in permeation rate. In this research there was no regular pattern of permeation and since NaSO₄ and MgSO₄ decreased NaCl and MgCl₂ permeability respectively, then the increase in the permeability of KCl and CaCl₂ may not be as a result of the increase of K⁺ and Ca²⁺ concentration in K₂SO₄+KCl solution and CaCl₂+CaSO₄ solution respectively.

Keywords: Supporting Electrolyte, Permeability, Red Blood Cell, Membrane, Chlorides, Sulphates

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1. Introduction

The addition of supporting electrolyte to an electrolyte solution can affect the ionic strength of that electrolyte solution [1]. According to IUPAC [2], a supporting electrolyte is an electrolyte solution, whose constituents are not electroactive in the range of applied potentials being studied, and whose concentration, ionic strength, and therefore contribution to conductivity is usually much larger than that of the electroactive substance to be dissolved in it. The supporting electrolyte is a strong electrolyte in high concentration in comparison to the concentration level of other ions in the medium. The effect of added electrolyte on the equilibria of a solution with ionic strength of 0.1 M or less is independent of the chemical nature of the electrolyte but depends on the ionic strength. It also maintains constant ionic strength and constant pH [3]. Mature mammalian red blood cell (RBC) consist of a single compartment, simplifying many approaches for transport studies making

mature erythrocytes from mammals a very easy-to use model for cell membrane transport studies intracellular constant measurements (ion or metabolite concentrations, pH) and flux experiments are easier than in any other type of cell containing multiple compartments [4-6]. Ions enter and leave cells through the cell membrane which is a selective barrier and has a dynamic structure composed of two layers of phospholipid molecules interspersed with cholesterol and proteins. It regulates passage of water and ions through it [7]. The type of ions or electrolyte that is allowed to pass through a cell membrane and at what rate can be determined by certain physiological factors like tonicity and osmotic concentration [8-9]. Different electrolyte solutions can vary in the osmotic pressure they exert on a cell membrane, depending on the degree of dissociation of the particular electrolyte in the solvent. In order to understand ion permeation through red blood cell (RBC) membrane, chemical phenomenon (dissociation and ionisation); physiological phenomenon (haemolysis); physical properties

(refractive index, dispersion or absorption of light) and optical density/turbidity (light scattering), have been used to study the electrochemical effects of individual electrons and protons on cell membrane, and the relationships correlating the number and energy states of electrons in ions and electrolytes with the observable properties shown by a cell system [8-12].

In the last few decades, researchers have shown numerous interests in the application of membranes in certain areas like pharmaceuticals, biotechnology, agriculture, food, medicine, etc. [1]. RBCs have been studied extensively, to determine the permeability across cell membrane but there exists a paucity of information on ion permeation through RBC membranes in electrolytes other than KCl and NaCl [9]. Experimental studies show that supporting electrolyte concentration has a marked influence on ion transport rate [13]. Okolue *et al.* [1], studied the effect of supporting electrolyte on permeation of ions through a polymeric membrane. The result shows that there was a rapid increase in the rate of permeation of NaCl, CaCl₂ and AlCl₃ electrolytes supported by Na₂SO₄, CaSO₄ and Al₂(SO₄) electrolytes respectively. They reported that this increase was as a result of an increase in the concentration of Na⁺, Ca²⁺, and Al³⁺ ions in the resultant mixture. The observation of this research may not explain the effect of supporting electrolyte on ion migration across a biological membrane.

This study is aimed at providing information on how supporting electrolytes affect permeation of ions through RBC membrane by determining the rate at which ions of hypotonic NaCl, KCl, CaCl₂ and MgCl₂ at pH 7.4 haemolyse RBC and the rate at which ions of hypotonic mixture of NaCl+Na₂SO₄, K₂SO₄+KCl, CaCl₂+CaSO₄ and MgCl₂+MgSO₄ at pH 7.4 haemolyse RBC using haemolysis as a marker while apply UV/Vis spectrophotometry to monitor the haemolysis process. Haemolysis time is used as

an index as to the rate of osmosis of penetrating molecules into the cell [12]. The amount of time that it takes for haemolysis or crenation to occur is directly related to the rate of permeation across the cell membrane [14]. Therefore, the rate of haemolysis can also be used as an indicator of the permeability for a particular penetrating solutes [8, 10, 15]. A transient turbidity measurement from a given blood/test solution, prepared by adding blood to that test solution, can give a quantitative index of the permeability coefficient of the substance being tested. Thus, from such measurements, one can deduce relative permeability for each of the test substances [9].

2. Materials and Methods

Isotonic solutions of NaCl (160 mM), KCl (160 mM), CaCl₂ (110 mM) and MgCl₂.6H₂O (110 mM) at pH 7.4 and solutions of NaCl (18 mM) + Na₂SO₄ (180 mM), + KCl (18 mM) + K₂SO₄ (180 mM), CaCl₂ (26 mM) + CaSO₄ (260 mM) and MgCl₂ (26 mM) + MgSO₄ (260 mM) at pH 7.4 were prepared taking into account their molecular weight and osmotic concentration. Each of the isotonic solutions (25 mL) mentioned above were diluted with 25 mL of distilled water to give half their initial concentration (50 % hypotonicity). Blood (20 mL) collected in acid-citrate-dextrose (ACD) was washed thrice with isotonic saline and the RBC separated. Each hypotonic solution (1.5 mL) prepared was mixed with 0.05 mL of RBC suspension and their absorbance/min for 16 min were measured using a UV/Vis spectrophotometer while the RBCs haemolyse. The absorbance of each solution were recorded. The data were used to plot graphs with the time (min) on the x-axis and the absorbance (Å) on the y-axis. Logarithmic trendlines were used to analyse the graph to understand the correlation and the interrelationship of the variables of this study.

3. Results and Discussion

Figure 1. Effect of Na₂SO₄ on NaCl electrolyte permeability through RBC membrane.

Figure 2. Effect of K_2SO_4 on KCl electrolyte permeability through RBC membrane.

Figure 3. Effect of $CaSO_4 \cdot 2H_2O$ on $CaCl_2$ electrolyte permeability through RBC membrane.

Figure 4. Effect of $MgSO_4 \cdot 7H_2O$ on $MgCl_2 \cdot 6H_2O$ ions permeability through RBC membrane.

Table 1. Changes in absorbance of the RBC+Electrolyte solutions after 16 minutes of haemolysis*.

	$y=mx+c$	$y_1(x=1)$	$y_2(x=17)$	$\Delta y(y_1-y_2)$	$\% \Delta y_{1,2}$	$\% \Delta y_{a,b}$
NaCl	$y_a = -0.04 \ln(x) + 0.189$	0.189	0.076	0.113	-59.79	-60.18
NaCl + Na ₂ SO ₄	$y_b = -0.016 \ln(x) + 0.304$	0.304	0.259	0.045	-14.80	
KCl	$y_a = -0.008 \ln(x) + 0.191$	0.191	0.168	0.023	-12.04	169.57
KCl + K ₂ SO ₄	$y_b = -0.022 \ln(x) + 0.331$	0.331	0.268	0.062	-19.03	
CaCl ₂	$y_a = 0.0039 \ln(x) + 0.268$	0.268	0.279	-0.011	4.10	536.36
CaCl ₂ + CaSO ₄	$y_b = -0.017 \ln(x) + 0.231$	0.231	0.183	0.048	-20.78	
MgCl ₂	$y_a = -0.016 \ln(x) + 0.217$	0.217	0.172	0.045	-20.74	-44.44
MgCl ₂ + MgSO ₄	$y_b = -0.009 \ln(x) + 0.263$	0.263	0.238	0.025	-9.51	

*Deduced from the logarithmic trendline of the graphic interpretation of the experiment's data

From the changes in absorbance of the RBC+Electrolyte solutions after 16 minutes of haemolysis deduced from the logarithmic trendline of the graphic interpretation of the experiment's data as shown in Figure 1 to 4 and Table 1, it is observed that there was a high 59.79% change in absorption of RBC in NaCl solution and a very low 4.10% increase in absorption of RBC in CaCl₂ solution. This low increase in absorption was reversed with the addition of CaSO₄ by 536.36% though CaSO₄ was observed to be insoluble in water. The addition of Na₂SO₄ to NaCl and MgSO₄ to MgCl₂ resulted in a decrease in permeation rate of the resultant solutions while the addition of K₂SO₄ to KCl resulted to an increase in permeation rate. The increase in absorption as a result of the addition of K₂SO₄ to KCl was relatively high at 169.57%. The observation from the addition of Na₂SO₄ to NaCl and the addition of CaSO₄ to CaCl₂ is different when compared with the results from the study of the effect of supporting electrolyte on permeation of ions through a polymeric membrane carried out by Okolue *et al.* [1], which shows that there was a rapid increase in the rate of permeation of NaCl, CaCl₂ and AlCl₃ electrolytes supported by Na₂SO₄, CaSO₄ and Al₂(SO₄) electrolytes respectively which they reported was as a result of an increase in the concentration of Na⁺, Ca²⁺, and Al³⁺ ions in the resultant mixture. While there was an increase in the rate of permeation of both CaCl₂ and KCl, with the increase for CaCl₂ observed to be three (3) times higher, a decrease at almost the same rate was observed for NaCl and MgCl₂.

4. Conclusion

Supporting electrolyte concentration in the external solutions had a marked influence on the transport rate studied. Na₂SO₄ added to NaCl and MgSO₄ to MgCl₂ resulted to a decrease in permeation rate of the resultant solutions while the addition of K₂SO₄ to KCl and the addition of CaSO₄ to CaCl₂ resulted to an increase in permeation rate. In this research there is no similarity in pattern of permeation hence the effect of K₂SO₄ to KCl ion permeation and the effect of CaSO₄ on CaCl₂ ion permeation across RBC membrane may not be attributed to an increase in the concentration of the respective K⁺ and Ca²⁺ ions in the resultant solutions.

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