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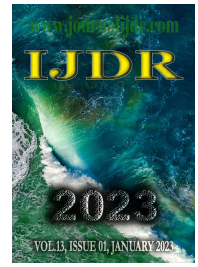
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RESEARCH ARTICLE

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THE SYSTEM MODELS AND THE BUFFERING CAPACITY OF ERYTHROCYTE MEMBRANE SURROUNDINGS IN RELATION TO PROTONS, FORMED IN THE MEMBRANE REDOXY POTENTIAL THREE STATE DEPENDENT 9 STEPPED FULL CYCLE OF PROTON CONDUCTANCE IN THE HUMAN BODY

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ABSTRACT

By our suggestion, the buffering capacity of erythrocyte membrane surroundings in relation to free protons, formed in the first 1-7 stages of proton conductance have implemented within Eight stage of 9 staged full closed cycle of proton conductance -located in the Respiring tissues -capillary blood which are existed around 87 trillion cells, Pulmonary circuit, where occurred oxygen unloading under effect of increased bicarbonate exit by bicarbonate/chloride ion shift mechanism, leading to increase of protons released from NADH, FADH, reduced KoQ resulting to entry of protons to erythrocyte membrane surroundings, all these processes described as the presence of protons from peripheral tissues favors the formation of salt bridges by protonating the terminal His residue of the beta subunits, an increase in protons causes oxygen release, while an increase in oxygen causes proton release, the hydrogen ions (protons) tend to displace the oxygen from the hemoglobin, protons promote oxygen unloading, which are inseparable parts - basic components of the System models of the Human body, including the membrane redox potential three state dependent 9 stepped full cycle of proton conductance and the four compartments, also the 10 functional systems, and four types of cells, distinguished by difference of proton conductance.

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INTRODUCTION

We described at first time the System models, including the membrane redox potential three state dependent 9 stepped full cycle of proton conductance including the four compartments, the 10 main functional systems and four types of cells, distinguished by difference of proton conductance by using the new conception as that it is existed a close relationship between following two expressions as Life has become dependent from presence of protons and electrons which were formed during the events called Big Bang 15 years ago and the presence of protons from peripheral tissues favors the formation of salt bridge in histidine residue of beta subunits (HarpersBiochemistry). The buffering capacity of erythrocyte membrane surroundings in relation to free protons, formed in the full cycle of proton and electron conductance inside the Human Body would be appeared in the 8-th stage of the full cycle as the diffusion of proton from mitochondrial matrix of all cells and metabolic water through plasma membrane of red blood cells also entry of CO₂ from all cells.

Quantity of free protons inside of erythrocyte membrane surroundings at 8- stage of cycle would make the remarkable influence to the buffering capacity of erythrocyte membrane surroundings in relation to protons, formed in the full cycle of proton and electron conductance and to diffusion speed of oxygen to 14 trillion cells that is, more protons inside of erythrocyte membrane surroundings more oxygen delivery to body cells by mechanisms as an increase in protons causes oxygen release, while an increase in oxygen causes proton release.

RESULTS AND CONCLUSION

By our suggestion, the buffering capacity of erythrocyte membrane surroundings in relation to free protons, formed in the first 1-7 stages of proton conductance have implemented within Eight stage of 9 staged full closed cycle of proton conductance -located in the Respiring tissues -capillary blood which are existed around 87 trillion cells, Pulmonary circuit, where occurred oxygen unloading under effect of increased bicarbonate exit by bicarbonate/chloride ion

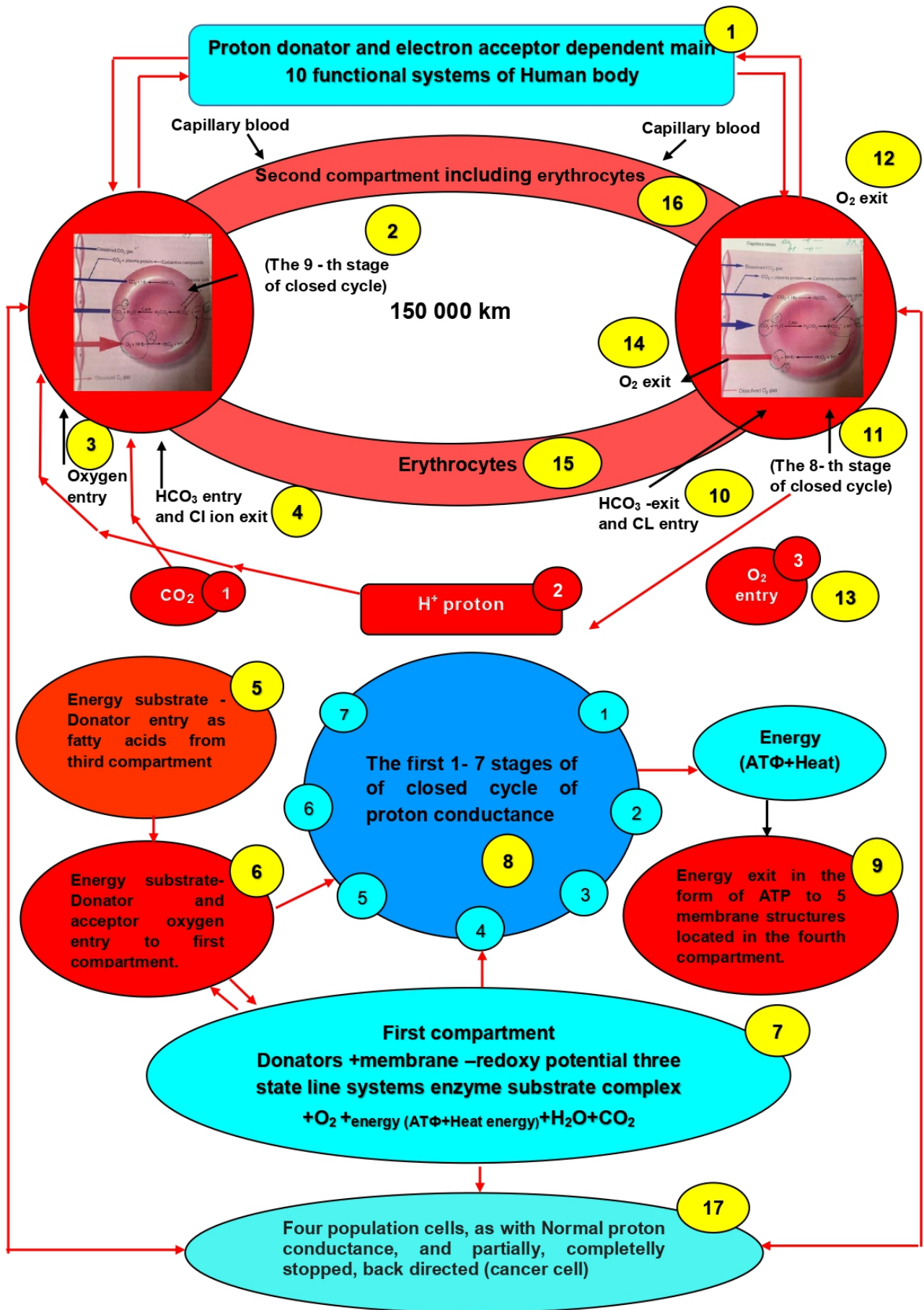


Figure 1. The system models, including the membrane redox potential three state dependent 9 stepped full cycle of proton conductance and the four compartments, the 10 functional systems

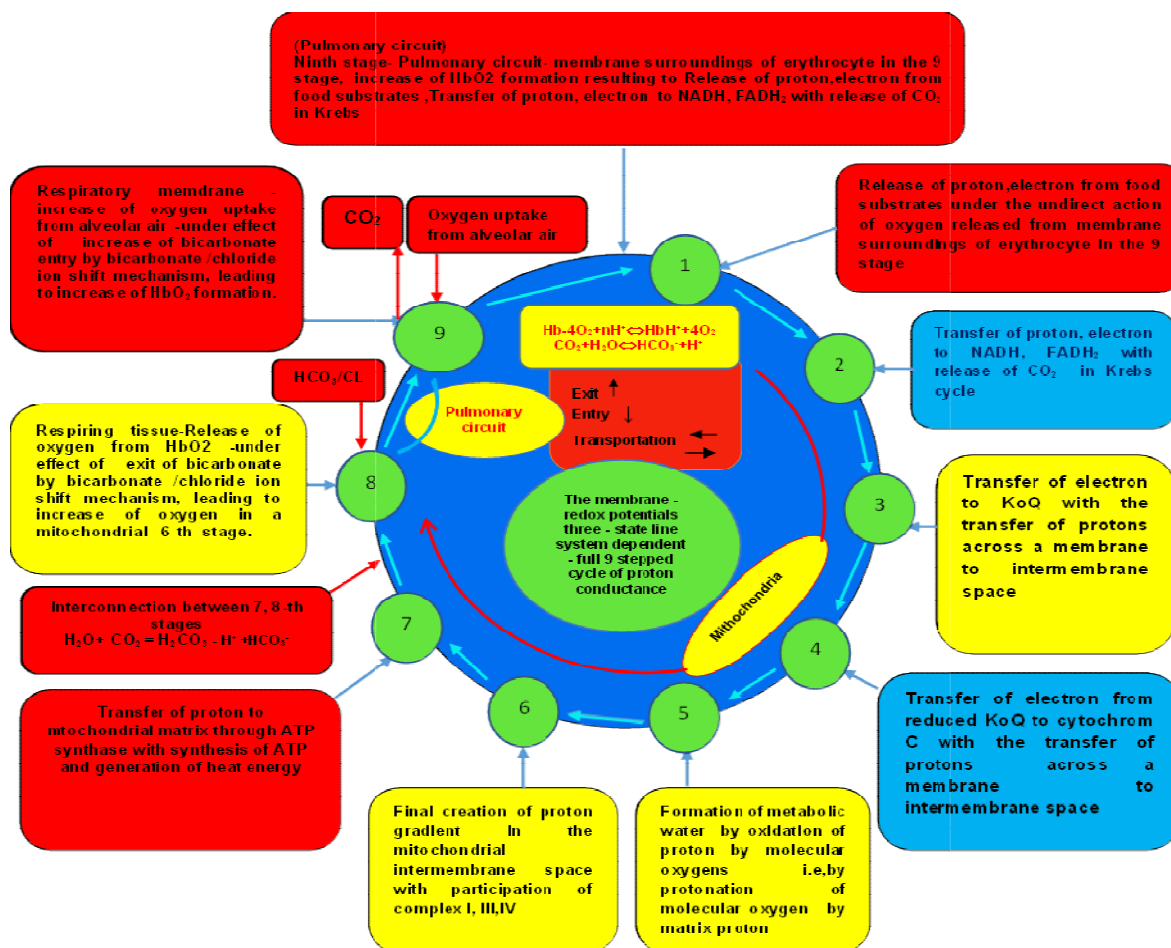


Figure 2. The membrane redox potential three state dependent 9 stepped full cycle of proton conductance

formed free proton, metabolic water, carbon dioxide and ATP, at second: following to this have been created the possibility to start the 9-th stage of closed cycle, located in the Respiratory membrane - Pulmonary circuit-increase of oxygen uptake from alveolar air -under effect of increased bicarbonate entry by HCO₃⁻ entry and CL ion exit (bicarbonate / chloride ion shift mechanism), Oxygen entry leading to increase of HbO₂ formation and the 8-th stage of closed cycle, located in the Respiring tissues - Pulmonary circuit -oxygen unloading by HCO₃⁻ -exit and CL entry-O₂ exit -Release of oxygen from HbO₂ under effect of exit of bicarbonate by bicarbonate exit / chloride ion entry shift mechanism, leading to increase of oxygen in a mitochondrial - 6-th stage of proton conductance, which have been conditioned the Energy substrate -Donator entry as fatty acids from third compartment to second compartment, which have been followed by Energy substrate-Donator and acceptor oxygen entry from second compartment to first compartment, where have been formed ATP, owing to formation of ATP in the first compartment have been created the condition to functioning of fourth compartment parameters, as 5 membrane structures-5 function systems, where conducted the normal genetic-cell division, information-response, biosynthetic, bioenergetic, biotransformation functions by using of high energy phosphate - ATP, high energy electrons NADPH, which have existed in the level of all cells of the 10 main systems of Human body as proton donator and electron acceptor delivering. In this connection that we had been established that it is existed a close relationship between following two expressions as Life has become dependent from presence of protons and electrons which were formed during the events called Big Bang 15 years ago and the presence of protons from peripheral tissues favors the formation of salt bridge in histidine residue of beta subunits (Harpers Biochemistry) we are trying to describe the system models, including the membrane redox potential three state dependent 9 stepped full cycle of proton conductance including the four compartments, the 10 functional systems.

The 10 main systems of Human body as proton donator and electron acceptor delivering have been consisted of First functional system is the system of delivering electron-proton donators as foods to living cells to maintain the normal level of donators as carbohydrate, aminoacids, fatty acids within membrane redox potentials 3 state line system as very important place of conducting of protons, electrons, starting from cyanobacteria formed during last 4,4 billion years, identical to Gastroenterological system, Second functional system is the system of delivering electron-proton acceptors as oxygen to living cells to maintain the normal level of acceptors as oxygens within membrane redox potentials 3 state line system as very important place of conducting of protons, electrons, starting from cyanobacteria formed during last 4,4 billion years, identical to Respiratory system, Third functional system is the system delivering electron-proton acceptors as oxygens and electron-proton donators as foods together to 87 trillion living cells to maintain the normal level of acceptors as oxygens and donators within membrane redox potentials 3 state line system as very important place of conducting of protons, electrons, starting from cyanobacteria formed during last 4,4 billion years, identical to Cardiovascular system, Fourth functional system is the system eliminating and neutralizing toxic metabolites and carbon dioxide, protonated carbon dioxide, also free protons formed during the functioning of the energy making system as "Donators (glucose, aminoacids, fatty acids) + membrane redox potentials three - state line system + acceptor as O₂ + ADP + Pi + H⁺ + nH⁺_{memb.space} = (ATP + heat energy) + H₂O + nH⁺_{matrix} + CO₂"-reaction medium, identical to Renal - urinary and Acid-Base controlling system, Fifth functional system is the system of converting some toxic metabolic products to normal metabolic products and conducting the synthesis and resynthesis of saturated and unsaturated fatty acids as main components of all membrane structures belong to membrane - redox potential 3 state line systems included to "Donators (glucose, aminoacids, fatty acids) + membrane redox potentials three - state line

system + acceptor as $O_2 + ADP + Pi + H^+ + nH^+_{memb.space} = (ATP + \text{heat energy}) + H_2O + nH^+_{matrix} + CO_2$ -reaction medium, identical to Hepato-biliary system, these all functional systems have functioned owing to 5 membrane structures-5 main functions as the normal genetic-cell division, information-response, biosynthetic, bioenergetic, biotransformation functions and four types of cells, distinguished by difference of proton conductance.

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