

# **Erythrocyte alterations and increased cardiovascular risk in chronic renal failure.**

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## **ABSTRACT**

Patients suffering from chronic renal failure have a higher burden of cardiovascular events, which increases in a dose dependent fashion as renal function worsens. Increased cardiovascular risk in these patients is thought to be mediated by the simultaneous presence of both traditional and non-traditional cardiovascular risk factors, the latter being associated with renal impairment.

Red blood cells are usually considered as carries of nutrients for tissues and respiratory gases, less so as compartments essential to vascular integrity. However, erythrocyte number, size, and integrity seem to severely influence cardiovascular morbidity and mortality as established in recent clinical studies with large patient cohorts.

In particular, the role of red blood cells in chronic renal failure tends only to be considered exclusively in relation to a change in their number. However, these cells in the uremic milieu are prone to many alterations, which may adversely affect the cardiovascular system.

In this review, we highlight the main qualitative erythrocyte alterations that may have a pathophysiologic role in the elevated cardiovascular risk of chronic renal failure

**Key words:** red blood cell, renal insufficiency, phosphatidylserine, nitric oxide, cardiovascular disease, endothelium, uremia, hemodialysis

## INTRODUCTION

Cardiovascular (CV) disease is still the main cause of morbidity and mortality in chronic renal patients (1). The risk of CV events increases when the glomerular filtrate falls below 60 ml/min/1.73 m<sup>2</sup> and gets steadily more pronounced as renal function diminishes, being especially high in patients on dialytic replacement therapy (1).

The traditional risk factors like aging, diabetes, hypertension and dyslipidemia are highly prevalent in chronic renal failure (CRF) and add significantly to CV pathology, especially at slight-moderate phases of renal dysfunction (2). Besides such factors, the higher incidence of CV events in CRF patients than in the general population is attributed to the simultaneous presence of *non*-traditional CV factors associated with renal impairment (3). The prevalence of such factors grows as renal function declines. They include disturbances of the calcium-phosphorus metabolism, malnutrition, expansion of the extracellular volume, inflammation, increased oxidative stress, hyperhomocysteinemia, a prothrombotic condition, and vascular rigidity (4).

In the case of CV disease in uremia, the role of the red blood cells (RBCs) tends only to be considered in relation to a change in their number. This typical and frequent clinical sign of uremia is rated a non-traditional CV risk factor in CRF (4). However, uremic RBCs may have a broader role in the CV system since in the uremic milieu erythrocytes are prone to many alterations which may adversely affect the CV system. The present review will comment on the main qualitative erythrocyte alterations and their potential pathophysiologic role in the uremic patient's CV risk status.

## RBC ALTERATIONS IN CHRONIC RENAL FAILURE

### ***Altered composition of the erythrocyte cell membrane***

The RBC is a fairly simple cell devoid of a nucleus; its main components are cell membrane and hemoglobin-rich cytoplasm. The cell membrane composition modulates the erythrocyte's visco-elastic properties, which are fundamental to survival of the cell in the stream of the circulation. Deformability is an intrinsic characteristic of normal RBCs, enabling them to pass down tiny capillaries and release oxygen. If they become less deformable, this triggers hemolysis in the capillaries and premature sequestration of RBCs by the reticulo-endothelial system (5), altering tissue oxygenation (6).

One of the far from negligible pathogenic features of anemia in the chronic kidney sufferer is the significant shortening of the circulating RBCs' half-life time (7). It is conjectured that this is essentially due to the toxic action of the uremic plasma environment upon the RBCs (8). It is indeed well known that when healthy donors' RBCs are re-infused into uremic patients their survival declines significantly (9). Shortening of the half-life not only means a drop in the quantity of circulating hemoglobin – the parameter normally used to measure the severity of anemia – but shows the difficulty the uremic patient's RBCs have in properly perfusing the microcirculation, precisely because of their altered rheologic properties (10).

It is of interest that recent studies have shown an altered protein composition in the uremic erythrocyte membrane which especially affects protein components in the cytoskeletal network. In particular, in RBCs from patients on maintenance hemodialysis treatment, analysis of the expression of certain key proteins of the erythrocyte membrane has especially brought to light the low level of spectrine (11). A more recent unbiased study designed to assess CRF patients' erythrocyte membrane composition by proteomics revealed some significant variations in many proteins such as beta-adducin, tropomodulin-1, ezrin and radixin (12). The cytoskeletal network is a chief factor behind RBCs' visco-elastic properties such as deformability (13). Such alterations to the membrane protein component (11, 12) might thus be part of the pathogenesis affecting erythrocyte deformability as observed in CRF (5, 14).

One compound that has a beneficial effect on RBC rheology is carnitine. Especially when this important cofactor in the beta-oxidation of fatty acids is present in concentrations greater than normal, it is able to affect the biophysical and visco-elastic properties of the cell membrane (10). By techniques of ektacytometry we have observed that carnitine can increase cell membrane elasticity, an effect that is somehow linked to carnitine's interaction with some protein components of the membrane-skeleton (15). It would appear that carnitine is able to strengthen the bonds among the proteins that form the cytoskeleton, and that such action is responsible for increasing membrane elasticity, which in turn improves RBC rheology. In support of these experimental findings, it has also been observed that carnitine significantly enhances interaction among membrane-skeleton proteins within the cell membrane (16).

The finding that there is a significant inverse correlation between membrane fluidity (an index of deformability) and the erythropoietin (EPO) dosage needed by

hemodialysis patients seems to suggest that differing degrees of alteration to the uremic RBC's mechanical properties may closely affect the dosage of EPO required (14).

To date there is no direct clinical evidence to indicate that changes in the rheologic properties of the RBCs increase the CV risk run by uremic patients. However, one recent preclinical experience suggests that rat RBCs preserved for 2 weeks at 4°C are less effective in ensuring normal oxygenation in the microcirculation than non-preserved RBCs (17). Now, it is known that blood bank-stored RBCs for transfusion are prone to various alterations (18); these involve the RBC membrane-skeleton and rheologic properties to an extent similar to the defects found in uremic patients' RBCs (19). Such alterations might explain not only the above-mentioned experimental observation as to the microcirculation oxygenation, but also some potential hypoxic effects to the coronary microcirculation and attendant CV risk.

The uremic erythrocyte membrane is altered not just in its protein, but also in its phospholipid component. Human RBC phospholipids are normally distributed asymmetrically in the double layer of the cell membrane, and keeping such asymmetry is a physiologically important process for the cell. Loss of normal asymmetry indeed may have many pathophysiologic implications, especially when the aminophospholipid phosphatidylserine (PS) gets exposed on the cell surface instead of its usual location on the inner face of the membrane (20). It may generate a signal recognized by the macrophages and thereafter the cell be phagocytosed and removed from the circulation (21). This accelerated suicide mechanism on the part of the erythrocyte is termed eryptosis (22); it is thought to hold *in vivo* importance in the diminished survival of sickle-shaped RBCs (23).

The RBCs of chronic renal sufferers (whether on dialysis or on conservative management) have PS more exposed on the outer face of the cell (24). As residual renal function declines, so this alteration gradually accentuates and is boosted by compounds pathologically present in uremic plasma (24), including beta-2 microglobulin (25), acrolein (26) and indoxyl sulphate (27). Extracorporeal dialysis may reduce the ability of uremic plasma to induce erythrocytes' PS exposure, and this is even more significant when techniques are used that have a broader range of removal than conventional hemodialysis, such as hemodiafiltration (28). Again, L-carnitine has been proved *in vitro* to reduce PS exposure on uremic patients' RBCs thanks to its antioxidant action (29); this may at least partly explain the increased erythrocyte

survival observed in hemodialysed patients when treated with this compound for 24 weeks (30).

What is more, the increased exposure of PS on the erythrocytes may contribute to the anemia found in CRF (31). It has been shown in this connection that the presence of PS makes uremic RBCs susceptible to phagocytosis by human macrophages (32), which may lie behind the diminished erythrocyte survival in uremia (7). Not being fully integrated into the dosage algorithms, the diminished RBC half-life in uremia might have a role in the erythropoietic response to EPO (33), prompting the phenomenon of hemoglobin variability (34) which has been associated with increased risk of mortality in the uremic patient (35).

The exposure of PS on the outer leaflet of the RBC membrane may also play a role in the coagulation process by promoting the assembly of two coagulation factor complexes, the prothrombinase complex and the tenase complex, that lead to thrombin generation (36). Perturbations in RBC PS exposure are seen as a pathogenic mechanism for the prothrombotic state of beta-thalassemia and sickle cell disease (37). Chronic uremia is associated with an increased risk of thrombotic complications, which may represent the predominant cause of mortality particularly in dialysis patients (38). This thrombophilia is considered as being multifactorial (39), but mechanisms specific to uremia that promote hypercoagulability remain to be yet identified (40).

Recent evidences indicate that uremic erythrocytes may display a pathological pro-coagulant phenotype (40, 41). A role for surface-exposed PS in the increased pro-coagulant activity of uremic RBC is suggested by significant correlations between PS levels and (i) prothrombinase activation, (ii) plasma levels of thrombin generation markers, (iii) plasma levels of fibrinolysis markers, and (iv) thrombotic events in a 3-years retrospective analysis (41). Furthermore, preincubation of uremic RBCs with annexin V (41) or with lactadherin (40), which have a propensity for binding to PS rendering it unavailable for PS-mediated processes, strongly inhibited RBC pro-coagulant activity. Erythrocyte-promoted hypercoagulability may also result from the cell release of vesicles called microparticles, which expose PS and express membrane antigens on their surface. It has been recently shown that circulating levels of RBC-derived pro-coagulant microparticles are significantly increased in uremic patients compared to healthy subjects (40).

Based on these findings, one may reasonably assume that abnormal RBC PS exposure might have a part in the induction of a hypercoagulable state in uremic

patients. Alternatively, RBC PS exposure could represent in these patients a new marker or a predictor of clinical thrombosis.

### ***Altered erythrocyte enzymatic activity***

#### *Diminished erythrocyte activity in the intracellular antioxidant system*

The term “oxidative stress” covers a set of alterations affecting tissue, cells and macromolecules once a disequilibrium sets in between excessive production of anti-oxidant agents and an inadequate antioxidant defense mechanism. That these systems should be correctly balanced (so-called ox-redox equilibrium) is essential to a whole range of physiological functions in our organism.

Pro-oxidant substances are highly reactive compounds having a half-life of a few seconds. When these hydroxyl radicals interact inside the cell environment with the chief biological macromolecules (lipids, proteins, carbohydrates and nucleic acids) they permanently modify the structure, function of these last, and are hence potential agents for CV damage (42).

When renal function is diminished, there is increased circulation of free oxygen radicals (reactive oxygen species, ROS) to an extent proportional to the level of glomerular filtrate (43). A rise in ROS oxidizes the oxyhemoglobin and forms hydrogen peroxide ( $H_2O_2$ ) and methemoglobin, which are responsible for tissue hypoxia (44). The presence of too many ROS also causes lipid peroxidation, resulting in permanent oxidative alterations to the lipoproteins (45). Some of these are intensely atherogenic such as malonyldialdehyde, the final products of advanced glycation or the protein products of advanced oxidation; they trigger the atherosclerotic process and damage the intimal wall of the arteries, causing a vascular inflammatory response and expression of leucocyte adhesion molecules, followed by migration of inflammatory cells into the sub-endothelial space. Oxidized LDL, in its turn, stimulates the formation of foamy cells and releases into circulation a great number of pro-inflammatory cytokines, which may destabilize plaque and increase the risk of ischemic cardiomyopathy (46, 47).

The lipid damage brought about by ROS via peroxidation may alter the integrity of the erythrocyte membrane, reducing cell half-life (48). Notably, RBCs may themselves contribute to uremic oxidative stress via alterations to the cellular enzymes that act upon the redox equilibrium. In CRF one notes, significantly, a marked increase in the activity of glutathione peroxidase and glutathione reductase (pro-oxidant enzymes), as

well as a significant reduction in superoxide dismutase and catalase (enzymes with antioxidant properties) (49).

Again, in uremia the erythrocyte enzyme activity may be affected by the hemodialysis treatment. Interestingly, it has been observed that using vitamin E-enriched membranes significantly improves the erythrocyte activity of superoxide dismutase (50) and the deformability of RBCs (51), which reduces the need for EPO. By using such filters along with vitamin C (250 mg i.v.) for two months it also proved possible to reduce intradialytic hemolysis and oxidative damage to the RBCs (52). What is more, glucose levels in the dialyzer fluid may play an important role in regulating the erythrocyte antioxidant capacity. Bober and coll. have shown that a glucose concentration of 5.6 mmol/l increases the exose monophosphate cycle in the erythrocytes, which benefits the antioxidant system and reduces the risk of hemolysis (53).

Concerning the use of antioxidant agents, Usberti and colleagues demonstrated how intravenous administration of 1200 mg of reduced glutathione (GSH) at the end of each hemodialysis session over a span of not less than nine months enabled 28 dialysis patients (14 of them on EPO treatment) to significantly improve erythrocyte survival and enhance the uremic picture (54). Oral administration of 600 mg/die over a period of 14 months enabled 36 dialysis patients to increase the erythrocyte activity of superoxide dismutase vis-à-vis healthy controls, suggesting it had a protective role (55). When uremic patients on replacement therapy (34 on hemodialysis, 13 on peritoneal dialysis) took 300 mg/die of vitamin E for 20 weeks it proved to be associated with improvement of the erythrocyte osmotic fragility (56). However, other studies have failed to confirm the efficacy of vitamin E in modifying the erythrocyte antioxidant activity in dialysis patients (57).

#### *Altered erythrocyte Nitric Oxide production and bioavailability*

Nitric oxide (NO) is a gaseous free radical produced by the Nitric Oxide Synthase (NOS), which is constitutively expressed in endothelial (eNOS) and neuronal (nNOS) tissues and is inducible (iNOS) in several tissues and cells in specific pathophysiological conditions. NO is known to be involved in the regulation of many intracellular signal pathways and exerts a wide pleiotropic action (58). At the vascular level, NO is a key molecule in the regulation of endothelium-mediated vasodilatation (59); also, it regulates several vascular mechanisms such as platelet aggregation,

leukocyte adhesion to the endothelium and smooth muscle cells proliferation/migration, whose alterations represent crucial events in the onset and development of the atherosclerotic process (60).

In this regard, endothelial dysfunction is considered one of the main pathological mechanisms that may contribute to increased CV risk in CRF (61). In fact, reduced endothelium-dependent vasodilation and increased plasma levels of the endothelium-derived mediators have been reported in uremic patients (61-63). The impaired endothelium-dependent vasodilation in CRF patients suggests a reduction of the NO bioavailability (61). However, the mechanisms underlying this phenomenon, which is triggered by the uremic state, remain not well defined.

To date, circulating levels of NO were mainly attributed to that produced by the endothelium and RBCs had been considered merely transporters of this gas. In fact, it is well known that erythrocytes bind to and inactivate the NO produced in the endothelium through a rapid reaction with oxyhemoglobin to form methemoglobin, S-nitrosohemoglobin and nitrates (64).

Under hypoxic conditions, the RBCs play a key role in the release of NO resulting in a NO-dependent vasodilation (65, 66), while under normal-oxygen or in various pathological conditions, the NO production/release by RBCs still needs to be better characterized. In this regard, the localization and function of the NOS in RBCs and in platelets were described for the first time in 2006 (67), although prior studies suggested the presence of this enzyme in RBCs (66, 68). In particular, in 2006, Kleinbongard and colleagues demonstrated the presence of a NOS-erythrocyte isoform, thus extending the role of RBC from merely transporters of NO to producers of such gas (67). Afterwards, the same authors hypothesized a RBC-NOS regulating mechanism comparable to that of endothelial NOS, although mature RBCs lack cellular organelles which play a key role in the regulation of endothelial NOS (69). More recently, it has been demonstrated that RBCs are able to produce NO even in normal-oxygen condition through the activation of their endothelial isoform of NOS (70). Moreover, eNOS expression levels and activity in the RBCs of patients with coronary artery disease are significantly reduced as compared to RBCs of healthy subjects (71).

Interestingly, our group has recently published a study describing the possible mechanisms involved in the NO synthesis and bioavailability in RBCs from uremic patients (72). In particular, we assessed the enzymatic NO production in erythrocytes of patients on hemodialysis and compared it with that of healthy individuals.

Our study demonstrated that, in basal conditions, although RBC-NOS expression was lower in uremic RBCs, its phosphorylation levels in Serine-1177, NO production and bioavailability (levels of cyclic guanosine monophosphate, cGMP) were significantly higher as compared to control RBCs. On the contrary, following RBCs stimulation with insulin or ionomycin (known to activate eNOS through a phosphorylation or calcium dependent mechanisms, respectively), the NO levels and its bioavailability were significantly higher in RBCs from healthy subjects compared to RBCs from hemodialysis patients, suggesting that uremic condition might reduce the response of RBC-NOS to further stimuli.

Our findings confirmed the presence of the eNOS isoform in erythrocytes (topic still highly debated and controversial) and identified, for the first time, an altered mechanism of eNOS activation in uremic RBCs (72). Interestingly, our data are supported by previous studies, which showed a marked increase of NO production in both erythrocytes and platelets from uremic patients on hemodialysis (73, 74). Moreover, a study published in 1982 (75) showed increased cGMP levels in RBCs from uremic patients compared to those from healthy subjects. This was confirmed in our recent study (72) where we demonstrated accumulating levels of cGMP in RBCs of subjects suffering from uremia, which was associated with a defective transport to the bloodstream of this nucleotide. In particular, we proved that this effect was associated with an impaired activity of erythrocyte cGMP membrane-transporter (multidrug-resistance-associated protein-4, MRP4), whose levels of nitration and nitrosylation were significantly increased under uremic conditions (72).

These data allow us to hypothesize that the erythrocyte intracellular cGMP levels might be a useful biomarker to monitor the uremic pro-oxidant and/or pro-inflammatory state associated with reduced vascular NO bioavailability in CRF.

### ***Abnormal heterotypic cell-cell interaction***

#### ***RBC-platelet interaction***

Activated platelets may engage in dynamic interplay with other blood cellular elements such as erythrocytes (76, 77). By the use of flow cytometry and transmission electron microscopy, we first reported direct evidence of uremic platelet-RBC adherence *in vitro*, and an increase in the number of circulating platelet-erythrocyte aggregates in chronic uremic patients (78). This observation seems attributable to both renal failure and hemodialysis treatment. Indeed, levels of platelet-RBC aggregates were higher in

CRF patients not on dialysis than in healthy subjects but lower than in dialysis patients (78). In the latter, platelet-erythrocyte aggregates could rise still further during the hemodialysis session, mainly depending on the membrane material contained in the hemodialyzer (78, 79). Fibronectin might serve as a molecular bridge between adjacent platelets and RBCs (78).

Interactions between platelets and erythrocytes markedly enhance *in vitro* several aspects of platelet reactivity (80, 81), and may be implicated *in vivo* in vascular abnormalities found in sickle cell disease (76, 77). Although atherosclerosis and vascular events are common in uremic patients, the significance of platelet-RBC aggregates in these patients remains yet to be elucidated.

#### *RBC-endothelial cell interaction*

We have shown that RBCs from patients on hemodialysis have an increased propensity to adhere to human endothelial cell monolayers, possibly via a direct interaction between PS exposed on the external leaflet of the erythrocyte plasma membrane and thrombospondin in the subendothelial matrix (82). Interestingly, we observed that adhesion of RBCs from uremic patients to the vascular endothelium may modulate NO release by endothelial cultures (83), as previously demonstrated for sickle RBC (84). In cultured human endothelial cells, erythrocytes from uremic patients inhibited the eNOS expression (mRNA and protein), activity and, consequently, NO production, whereas erythrocytes from normal subjects had no such role (83). Defective eNOS is thought to represent a key parameter characterizing endothelial cell dysfunction, and in uremia it might have several consequences. Reduced NO production may be crucial to functional and structural vascular changes, promote oxidative stress and increased expression of adhesion molecules resulting in endothelial damage and atherosclerosis (85), and play a role in elevated blood pressure (86).

The enhanced interaction between uremic RBC and the endothelium, besides the pro-inflammatory effects stemming from the reduced NO bioavailability, may also directly induce a vascular inflammatory phenotype (87). In fact, adherence of uremic RBCs to human endothelial cells induced a time-dependent up-regulation of vascular cell adhesion molecule-1 and of intercellular adhesion molecules-1 (87), an early step in atherogenesis (46). Such increased vascular adhesion molecule expression was associated with mitogen-activated protein kinase activation and with impairment of

protein kinase Akt phosphorylation, which are pro-inflammatory, NO-independent, signal transduction pathways (88, 89). In our experimental model (87), the increase in endothelial adhesion molecule expression was supported by the remarkable enhanced adhesion of human monocytoïd cells to endothelial cells cultured with uremic RBCs. These findings suggest that therapeutic strategies aimed at limiting the action of these pathways may have the potential to prevent and treat uremic vascular disease, a concept that deserves specific investigation.

## **CONCLUSION**

The current review of the literature has focused on the potential role of erythrocyte qualitative alterations in cardiovascular complications in CRF. On the basis of the aforementioned findings, it is possible to suggest that erythrocytes in the uremic milieu undergo structural and functional impairments such as reduction of antioxidant activity and alteration of the production of nitric oxide, increased pro-coagulant activity and endothelium adhesion, modification in the composition of plasma membrane, all aspects which may jeopardize their properties.

These erythrocyte molecular transformations certainly may play an important role per se in understanding RBC role in the pathogenesis of vascular complication of uremia and, additionally, through them, it might be possible to identify potential erythrocyte biomarkers useful both for prevention and for the monitoring of CV risk in CRF patients.

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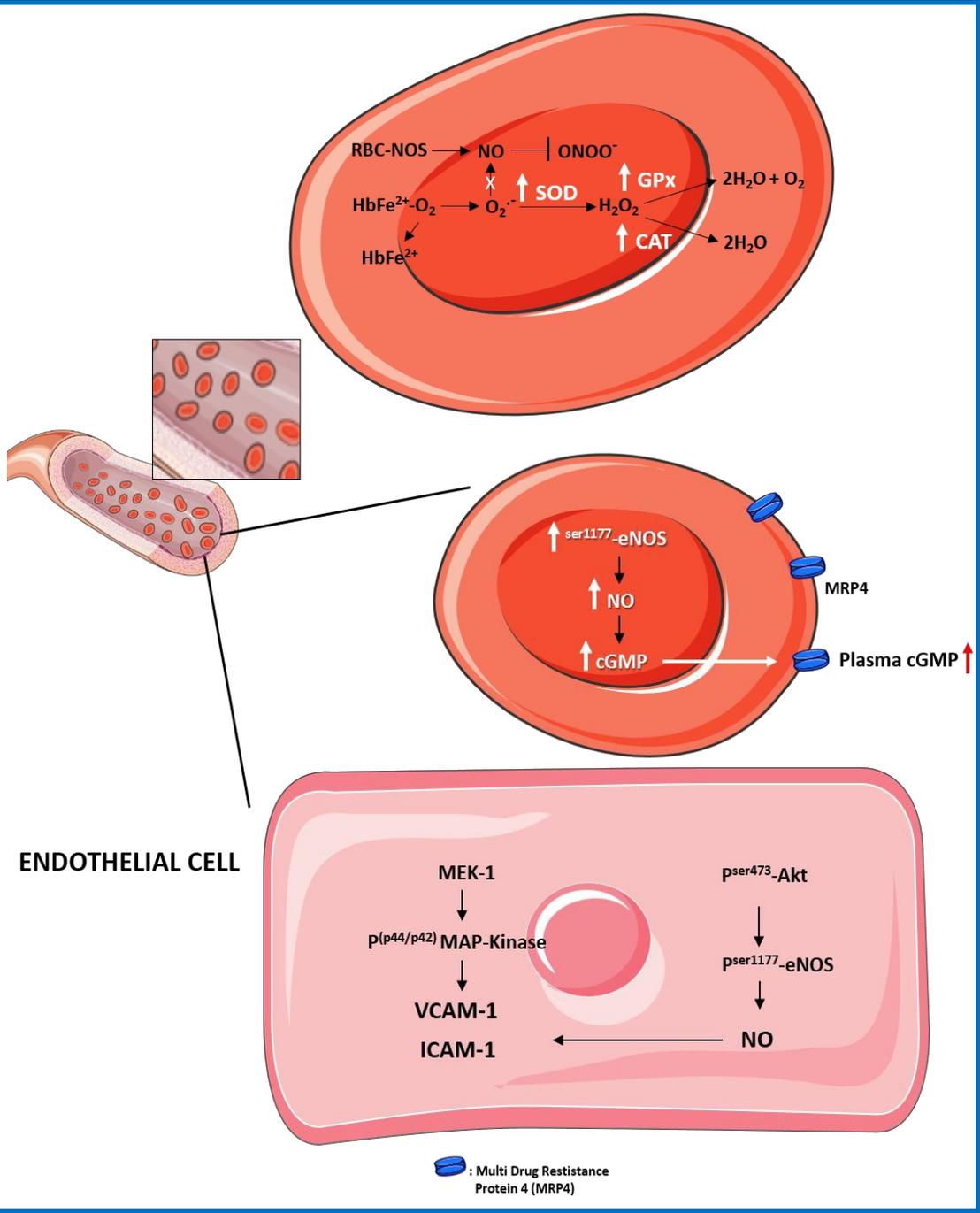
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# HEALTHY RBC



# UREMIC RBC

