Nuclear and Cytoplasmic Functions of Vitamin C
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ABSTRACT: Vitamin C (ascorbic acid) is a water-soluble antioxidant and a cofactor for a large number of enzymes. It is present in all tissues and especially abundant in corneal epithelium, stem cells, and neurons. Although similar to thiols in its ability to react with many reactive oxygen species (ROS), ascorbate is much better (>100X faster) than glutathione at scavenging of primary ROS (superoxide radical and singlet oxygen). Ascorbate appears to be especially important for elimination of O$_2^*$ in the nucleus which contains little or no SOD activity. Cofactor functions of ascorbate involve the maintenance of activity of Fe(II)/2-oxoglutarate-dependent dioxygenases via reduction of Fe(III). The most prominent activity of ascorbate-dependent dioxygenases in the cytoplasm is hydroxylation of prolines in proteins involved in the formation of extracellular matrix and regulation of metabolism and hypoxia responses. In the nucleus, ascorbate is important for oxidative demethylation of 5-methylcytosine in DNA (by TET proteins) and removal of methyl groups from histone lysines (by JmjC demethylases). Differentiation and other cellular reprogramming processes involving DNA demethylation are especially sensitive to ascorbate insufficiency. High doses of vitamin C alone or in combinations with drugs produced cancer-suppressive effects which involved redox, immune, and epigenetic mechanisms. Solutions to vitamin C deficiency in cultured cells are discussed to improve the physiological relevance of in vitro models. An abundance of vitamin C in rodents limits their ability to fully recapitulate human sensitivity to adverse health effects of malnutrition and xenobiotics, including neurotoxicity, lung injury, and intergenerational and other epigenetic effects.

Vitamin C is an essential micronutrient to humans and other primates caused by loss-of-function mutations in L-gulonolactone oxidase (encoded by GULO gene) which catalyzes the final step in ascorbic acid biosynthesis from glucose. The majority of eukaryotes synthesize ascorbic acid and yeast, producing a related compound erythro-ascorbic acid. Vitamin C is found in the diet and in vivo in its reduced (ascorbic acid) or oxidized (dehydroascorbic acid) forms (Figure 1A). Due to its low pK$_a$ = 4.2 at the C3 hydroxyl, ascorbic acid is present almost completely (>99%) as ascorbate anion at physiological pH. Severe vitamin C deficiency in humans leads to scurvy, which is a fatal disease that can be cured only by a restored intake of vitamin C.\textsuperscript{1} Several overt clinical manifestations of scurvy (reopening of old wounds, bleeding, loss of teeth, gums, and skin abnormalities) are clearly linked to the long-established role of ascorbate in the production of mature collagen. Other symptoms and high-fatality rates were more difficult to explain by defects in collagen formation, pointing to a broader set of essential activities of ascorbate. Although the occurrence of scurvy is rare in the modern world, the incidence of a significant vitamin C deficiency is still common even in the wealthy countries.\textsuperscript{2–4} A recent popularity of animal product-based (ketogenic and similar) diets can potentially increase the incidence or extent of vitamin C insufficiency. Vitamin C deficiency is at least partially responsible for intergenerational epigenetic effects produced by a nutritional deprivation during pregnancy. The main physiological functions of ascorbate include its activities as an antioxidant and a critical cofactor for a growing number of enzymes that are directly or indirectly involved in multiple cellular activities. Resurgence of research interest in vitamin C is further fueled by recent discoveries of its importance in the main biochemical processes regulating epigenetics and its emerging promise in cancer treatment. These new findings indicate that the role of ascorbate in modulation of cellular responses to toxins, drugs, and other stressors extends beyond its antioxidant activities.

VITAMIN C UPTAKE AND CONCENTRATIONS IN VIVO

Ascorbate enters cells through sodium-dependent vitamin C transporters (SVCT) 1 and 2.\textsuperscript{5–8} SVCT1 (SLC23A1) is important for ascorbate absorption in the intestine and,
neurons, adrenal glands, and corneal epithelium.10,11 Ascorbate levels of ascorbate in tissues (20 mM) for most tissues, but they reach 10 mM and higher in concentrations of ascorbate are much higher and range from 1 to vitamin C, which predominantly comprises ascorbate. Cellular transporter DMT1.

Another source of oxidized vitamin C which enters cells through GLUT glucose transporters. Fe(II) is taken up by cells via the divalent metal export mechanisms of from cells are currently unknown. High levels of ascorbate in tissues (20–200× over plasma concentration) are also frequently viewed as a systemic reservoir of vitamin C in humans. The postulated reservoir role is probably not the primary biological reason for high ascorbate in human tissues, as equally high tissue concentrations of ascorbate are also present in mice and rats that continuously produce copious amounts of vitamin C (~100× over recommended intake in humans). There is now clear experimental evidence that very high ascorbate concentrations are in fact necessary for the normal functions of some cells (stem cells, for example). Although a scurvy-level vitamin C deficiency is currently rare, significantly lower ascorbate levels are frequently found among cancer patients, which in experimental models promoted cancer aggressiveness and limited responses to several chemotherapeutics.

Antioxidant Activities in Cells

Ascorbate is generally known as a water-soluble antioxidant. However, the importance of its specific protective activities and the overall role as the radical scavenger and antioxidant are not well covered or even completely ignored in some otherwise comprehensive reviews of cellular antioxidant defenses. Among small antioxidants, glutathione typically receives a lot of attention, although, as discussed below, it is clearly an inferior direct scavenger of the primary cellular oxidants in comparison to ascorbate. Underappreciation of ascorbate could be related to its (near) absence in cultured cells, which are the most commonly used biological models in studies of oxidative stress. In ascorbate-devoid cells, glutathione certainly takes on the role of the key small antioxidant in the direct detoxification of reactive oxygen species (ROS). A large portion of cellular damage by ROS frequently results from the reactive products of lipid oxidation, such as acrolein, crotonaldehyde, and 4-hydroxy-2-nonenal. Glutathione and other small thiols protect against toxicity of aldehydes, especially the most toxic group of α,β-unsaturated aldehydes, through the formation of inactive conjugates.14–16 This property is not shared by ascorbate which acts as a reducer (electron donor) and does not form covalent bonds with reactive carbonyls. However, as discussed below, ascorbate is much more effective than glutathione in the elimination of primary ROS.

Ascorbate is a broad spectrum antioxidant as it can react with a range of organic radicals and ROS. One-electron reduction of ROS and other radicals by ascorbate results in the formation of the resonance-stabilized ascorbate radical, which is completely unreactive with other biomolecules and is primarily lost via disproportionation reaction with another ascorbate radical, yielding ascorbate and dehydroascorbic acid.17 A broad radical/ROS reactivity is not unique to ascorbate, as it is also found among thiols including glutathione which is present in cells at comparable concentrations to those of ascorbate. For example, both glutathione and ascorbate are very effective at removal of a highly toxic hydroxyl radical (·OH). The reactivity of ascorbate and glutathione with the most abundant cellular ROS, H₂O₂, is very limited (rate constants 2 × 10⁻¹¹ M⁻¹ s⁻¹ and 0.9 M⁻¹ s⁻¹, respectively)18 and unlikely to be physiologically significant.
The importance of glutathione in suppression of H$_2$O$_2$ toxicity stems from its role as a cofactor for glutathione peroxidases (GPXs), especially a very effective cytoplasmic enzyme GPX1 which is present in cells at ~1 μM concentration and has the rate constant of 2 × 10$^7$ M$^{-1}$ s$^{-1}$ for H$_2$O$_2$.\textsuperscript{19} Multiplication of rate constants by cellular concentrations of GPX1 and ascorbate in vivo (5 mM) gives rates of H$_2$O$_2$ elimination equal to 20 s$^{-1}$ and 0.01 s$^{-1}$, respectively. Thus, the enzymatic activity of GPX1 is expected to be 2000× more effective at detoxification of H$_2$O$_2$ in comparison to ascorbate and even higher relative to the non-enzymatic removal with glutathione. Catalase and peroxiredoxin-1 (PRDX1) represent other abundant and very effective enzymatic systems for elimination of H$_2$O$_2$ by converting it into H$_2$O, further indicating that H$_2$O$_2$ detoxification in cells is principally based on the biological mechanisms and likely includes minimal contributions from direct chemical reactions with small antioxidants. Although ascorbate is a highly hydrophilic molecule, it also suppresses oxidation of lipids by reducing the tocopheroxyl radical and thereby maintaining the levels of the main lipid-soluble antioxidant, vitamin E.\textsuperscript{20,21} Consistent with this role, addition of physiological levels of ascorbate inhibited lipid peroxide-dependent cell death (ferroptosis) in cultured cells.\textsuperscript{22} Glutathione is probably even more important for prevention of ferroptosis due to its role as a cofactor for the main lipid peroxide-eliminating enzyme GPX4.\textsuperscript{23} Thus, both ascorbate and glutathione suppress lipid peroxidation and ferroptosis but through mechanistically distinct mechanisms.

The main difference in the direct scavenging of the main ROS between ascorbate and glutathione (small thiols in general) is found in their reactivity with the superoxide anion radical (O$_2^•^−$), which is >100× higher for ascorbate than for glutathione. Rate constant for ascorbate is 2.9 × 10$^5$ M$^{-1}$ s$^{-1}$ (mean of two studies),\textsuperscript{24,25} and the highest reported rate constant for glutathione is 1.1 × 10$^5$ M$^{-1}$ s$^{-1}$.\textsuperscript{26} Other estimates of rate constant for glutathione are much lower.\textsuperscript{15,18} O$_2^•^−$ is the primary cellular ROS arising during the respiratory metabolism in mitochondria, physiological enzymatic activities (NADPH oxidases, xanthine oxidase) and redox cycling of various drugs and toxicants.\textsuperscript{15,18} Dismutation of O$_2^•^−$ by the cellular enzymes SOD1/2 produces hydrogen peroxide (H$_2$O$_2$), whereas the reaction of O$_2^•^−$ with a weak oxidant nitric oxide (NO) yields a secondary oxidant peroxynitrite (ONOO$^−$). H$_2$O$_2$ and ONOO$^−$ can consequently serve as sources of tertiary oxidants, including a very reactive OH$^•$ radical (Figure 2). Considering its abundance (10 μM or more)\textsuperscript{27} and a very high activity of cytoplasmic SOD1 in the removal of O$_2^•^−$ ($k = 2 × 10^9$ M$^{-1}$ s$^{-1}$),\textsuperscript{28} it is important to ask whether ascorbate can make a meaningful contribution to the elimination of this primary ROS. Using cellular concentrations and rate constants for calculations of their reaction rates (SOD1: 10 μM × 2 × 10$^9$ M$^{-1}$ s$^{-1}$ = 2 × 10$^8$ s$^{-1}$; ascorbate: 5 mM × 2.9 × 10$^9$ M$^{-1}$ s$^{-1}$ = 1.5 × 10$^8$ s$^{-1}$), SOD1 appears to be 13.3× more effective in the elimination of O$_2^•^−$ than ascorbate. This difference translates into a 7% contribution of ascorbate to the removal of cytoplasmic superoxide. In cells with its particularly high concentrations (stem cells, neurons, corneal epithelium), ascorbate should make a larger impact on scavenging O$_2^•^−$ and more significantly complement SOD1 activity. Although its superoxide-scavenging activity diminishes the formation of peroxynitrite, ascorbate is known to accelerate a release of NO from S-nitrosoglutathione, which is a major physiological carrier of nitric oxide. Use of specific metal chelators in cell culture found a dependence of this activity on the availability of redox-active Cu.\textsuperscript{25} Since cellular concentrations of free Cu ions are tightly controlled, ascorbate-promoted decomposition of S-nitrosoglutathione appears to occur more likely outside the cells.

The main cellular ROS and antioxidant mechanisms. Superoxide anion radical (O$_2^•^−$) is the primary ROS produced by various physiological and pathophysiological processes. Reduction of O$_2^•^−$ by SODs or ascorbate (Asc) produces a less reactive but more diffusible H$_2$O$_2$ which is detoxified by catalase, peroxiredoxin 1 (PRDX1), and glutathione peroxidase 1 (GPX1). Reaction of H$_2$O$_2$ with free Fe(II) and less frequently Cu(I) yields a highly damaging *OH radical. Hydroxyl and other reactive radicals are also formed during the decomposition of peroxynitrite generated in the reaction of nitric oxide radical (*NO) with O$_2^•^−$. In comparison with the biological processes, ascorbate makes no significant contribution to the detoxification of H$_2$O$_2$ and plays a minor role in the removal of cytoplasmic O$_2^•^−$ in the majority of cells. Secondary radicals (tertiary ROS) arising from H$_2$O$_2$ and ONOO$^−$ lack specialized enzymatic defenses and are primarily detoxified by small antioxidants such as ascorbate and small thiols (glutathione). Cells also lack enzymatic processes for removal of singlet oxygen (°O$_2$) which is effectively eliminated by ascorbate via a two-electron reduction to H$_2$O$_2$. In contrast to the cytoplasm, the nucleus has a minimal or no SOD activity, which elevates the importance of ascorbate in scavenging superoxide in this cellular compartment.
In contrast to H$_2$O$_2$ and O$_2$$^•$••, tertiary ROS (•OH, •CO$_2$$^−$, and •NO$_2$ radicals) (Figure 2) lack enzymatic defenses and are primarily eliminated in direct reactions with ascorbate and small thiols. Ascorbate is expected to be more important than glutathione in detoxification of nitrogen dioxide and carbonyl radicals. Both •CO$_2$$^−$ and •NO$_2$ preferentially react with the thiolate anion (RS$^−$), which constitutes only a small fraction (5%) of the total glutathione at physiological pH. The predominance of the RSH form is advantageous for the reaction of glutathione with •OH, which proceeds through hydrogen abstraction. The rate constants for the reactions of •OH with glutathione (1.6 × 10$^{10}$ M$^{−1}$ s$^{−1}$) and ascorbate (4.5 × 10$^{10}$ M$^{−1}$ s$^{−1}$) are very high and only moderately different, indicating that the relative contributions of these antioxidants to the detoxification of hydroxyl radical are largely a function of their cellular concentrations.

Another primary cellular oxidant that is effectively removed by ascorbate is singlet oxygen (1O$_2$), which is an electronically excited state of O$_2$. Although it is not a radical, 1O$_2$ is a strong oxidant and mutagen. Singlet oxygen is produced in photo-reactions of UV and visible light with biomolecules and released by certain enzymatic reactions (lipid peroxidases, for example). Cells lack specialized enzymes for detoxification of singlet oxygen and rely on small antioxidants for protection against this oxidant. The rate constant for reaction of ascorbate with 1O$_2$ is very high (k = 3 × 10$^{10}$ M$^{−1}$ s$^{−1}$) and comparable to those of specialized enzymes detoxifying superoxide or hydrogen peroxide. Unlike reactions with other ROS, ascorbate acts as a two-electron donor in the elimination of singlet oxygen:

\[ 1O_2 + \text{ascorbate} \rightarrow H_2O_2 + \text{dehydroascorbic acid} \]

Scavenging of 1O$_2$ by glutathione is also relatively fast (k = 2.4 × 10$^{10}$ M$^{−1}$ s$^{−1}$) but still approximately 100× slower than that by ascorbate. The singlet oxygen-reactive form of glutathione is the thiolate anion which is present as a minor fraction (pK$_a$ = 8.7 for glutathione). Thus, ascorbate is clearly a dominant small molecule scavenger of 1O$_2$. A biological support for its importance in detoxification of 1O$_2$ is the presence of high concentrations of ascorbate in human corneal epithelial cells (∼8 mM) which are continuously exposed to UV and visible light and, therefore, experience an extensive formation of singlet oxygen. Consistent with this interpretation, nocturnal animals have significantly lower ascorbate levels in their corneal epithelium in comparison to diurnal species. Protective effects of corneal ascorbate against UV-induced DNA damage have been confirmed in vivo. Another example where ascorbate reacts with a potent toxicant via two-electron transfer is reduction of carcinogenic chromium(VI). Reaction of Cr(VI) with ascorbate yields non-oxidizing Cr(IV). Non-catalytic reduction of Cr(VI) with ascorbate yields non-oxidizing Cr(IV). If Cr(VI) reacts with thiols, reduction proceeds through an one-electron transfer, and the resulting product, Cr(V), is a strong oxidant causing DNA breakage and phosphorylation of the transcription factor p53 by the oxidant-sensitive kinase ATM.

Radical-scavenging activities of antioxidants are recognized as being important for all physiological functions of cells, including preservation of genome stability. The main ROS-detoxifying enzymes exhibit a clear intracellular compartmentalization. SOD1, PRDX1, and GPX1 are primarily cytoplasmic, SOD2 is a mitochondrial enzyme, and catalase resides in peroxisomes. The nucleus is the second largest compartment in the majority of cells, and it does not appear to contain significant levels of these antioxidant enzymes under normal conditions. A severe oxidative stress can cause a nuclear translocation of SOD1, which is linked to the regulation of transcription of the antioxidant genes. The paucity of enzymatic defenses makes the nucleus more dependent on the anti-ROS activity of small antioxidants. Nuclear concentrations of ascorbate have not been specifically established yet, but they are expected to be similar to those in cytoplasm since nuclear pores allow a free diffusion of molecules with molecular weight <1000 Da. The molecular weight of the ascorbate anion (175.1) is much smaller than this threshold, which should promote a near equilibrium between nuclear and cytoplasmic pools. Immunomicroscopy-based measurements in human cultured cells found that nuclear ascorbate levels were higher than those in the cytoplasm. However, it is unclear whether the employed aldehyde fixation for immunomicroscopy can be effective for retention and detection of the small molecule ascorbate. In the (near) absence of SODs, ascorbate is expected to play a dominant role in the removal of superoxide inside the nucleus. The apparent exclusion of SOD1 from the nucleus could be linked to the avoidance of the deleterious genotoxic effects of this enzyme which causes strand breaks when it comes in contact with DNA. Similar to cytoplasm, ascorbate should also act as a main reducer of mutagenic singlet oxygen. H$_2$O$_2$, the product of 1O$_2$ and O$_2$$^•$•• reduction by ascorbate, is less reactive and more diffusible and can be enzymatically eliminated upon its entry into the cytoplasm. The effectiveness of ascorbate in the removal of nuclear O$_2$$^•$•• can be judged from the extent of the nuclear oxidant-sensitive signaling. The redox-active drug bleomycin causes DNA breaks and releases superoxide, causing activation of the nuclear kinase ATM via DNA damage-dependent and direct oxidation-mediated mechanisms, respectively. Physiological concentrations of ascorbate produced a strong suppression of the direct oxidation-mediated activation of ATM and phosphorylation of its nucleoplasmic target CHK2. These observations experimentally support the notion that ascorbate plays a major role in detoxification of O$_2$$^•$•• in the nucleus. Stem cells contain much higher concentrations of ascorbate in comparison to their differentiated progeny, which was linked to the need for the cofactor activity of ascorbate in DNA demethylation activity and epigenetic reprogramming of gene expression in stem cells. It is unclear why reduction of Fe(III) to Fe(II) in DNA demethylases (TET proteins) would require >10× higher ascorbate concentrations than in other members of the Fe(II)/2-oxoglutarate-dependent dioxygenases. Another possibility is that high ascorbate concentrations are not needed as much for the Fe(III) reduction in TET proteins as for the elimination of O$_2$$^•$•• produced during aborted catalytic cycles, which otherwise can inactivate the enzyme. The active intermediate in TET enzymes is Fe(II)-O$_2$ which, if not used catalytically, will form Fe(III)-O$_2$$^•$•• with a potential to produce damage locally or upon the release of O$_2$$^•$••. A similar mechanism is responsible for the release of superoxide and self-inactivation of bleomycin by its Fe(II)-bound O$_2$. The absence of the external substrate, the uncoupling between oxygen activation and substrate oxidation occurs with a relatively high frequency in Fe(II)-dependent microsomal CYP450 monoxygenases. These enzymes release superoxide as a result of the decay of the initial Fe(II)-O$_2$ oxycomplex. For many substrates, the coupling efficiencies are below 10%, indicating that a large fraction of Fe-activated oxygen in CYP450s is released as ROS. The uncoupling frequency in oxidative demethylation of DNA by TET proteins is currently unknown.
Ascorbate is a soluble cofactor for more than 60 cellular enzymes, which is indicative of its broad impact on physiological processes and consistent with multiple clinical symptoms in scurvy and developmental abnormalities arising as a result of maternal vitamin C deficiency during pregnancy. The main class of enzymes engaging ascorbate as the unbound cofactor are Fe(II)/2-oxoglutarate-dependent dioxygenases. 2-Oxoglutarate acts a cosubstrate and Fe(II) as the O2-binding cofactor. The catalytic reaction of these enzymes involves a transfer of one oxygen atom from Fe(II)-bound O2 onto the substrate and the resulting formation of stable hydroxylated proline derivatives.

Cofactor for Cytoplasmic Enzymes

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Cysteine is a sulfur-containing amino acid that plays a crucial role in cellular processes. It is involved in various biological functions, including detoxification, redox regulation, and signal transduction. The balance of cysteine levels in the body is regulated by the methionine cycle, which involves the conversion of methionine to cysteine through the action of enzymes such as cysteine desulfhydrase. This balance is important for maintaining cellular homeostasis and tissue integrity.

Cysteine is also a precursor of other important molecules. It is converted into glutathione, a powerful antioxidant that protects cells from oxidative stress. Cysteine is also involved in the synthesis of taurine, a sulfur-containing amino acid that plays a role in various physiological processes, including bile formation, neurotransmission, and energy metabolism. Additionally, cysteine is a precursor of cysteic acid, a metabolite that is involved in the detoxification of certain chemicals.

Cysteine is also crucial for cell proliferation and differentiation. It is involved in the regulation of cell cycle progression and can influence the expression of growth factors and their receptors. Cysteine levels have been linked to cancer risk and progression, as deregulation of the methionine cycle can lead to alterations in cell metabolism and proliferation.

In summary, the balance of cysteine levels is critical for maintaining cellular health and homeostasis. Disruptions in the methionine cycle can lead to various health issues, including oxidative stress and altered cell proliferation. Therefore, understanding the regulation of cysteine levels and the role of the methionine cycle in these processes is essential for developing strategies to maintain cellular health and prevent disease.

Figure 3. Role of ascorbate in the activity of Fe(II)/2-oxoglutarate-dependent dioxygenases. (A) General mechanism of oxidative hydroxylation by Fe(II)/2-oxoglutarate-dependent dioxygenases. Fe(II)-bound O2 serves as a source of oxygen atoms in oxidative hydroxylation of a substrate and the cosubstrate 2-oxoglutarate. Ascorbate (Asc) acts as a reductant of Fe(III) to Fe(II), which restores activity of dioxygenases after each catalytic cycle. Constitutive degradation of hypoxia-inducible transcription factors HIF-1α and HIF-2α in normoxic cells. HIF-1α/2α are continuously produced and then immediately targeted for degradation in normoxia via O2-dependent hydroxylation of specific Pro residues (Pro402 and Pro564 in HIF-1α). Hydroxylated prolines are recognized by VHL, which triggers polyubiquitination (Ub+) of both HIFαs and their subsequent proteolysis by proteasomes. Ascorbate acts as a preferred reductant of Fe(III) to Fe(II) to maintain activity of HIF-targeting prolyl hydroxylases PHD1–3. Another hydroxylation site in HIF-1α in normoxic conditions is Asn803, which regulates binding of the transcription coactivators p300/CREB (not shown). Asn803 hydroxylation is mediated by the Fe(II)/2-oxoglutarate-dependent dioxygenase FIH, whose activity is promoted by ascorbate through reduction of Fe(III) to Fe(II). Ascorbate acts as a preferred reductant of Fe(III) to Fe(II), which restores activity of dioxygenases after each catalytic cycle. (B) Removal of genomic 5-methylcytosine (5-mC) by a sequential oxidation of the methyl group by TET enzymes. Ascorbate is required for reduction of Fe(III) to Fe(II) to restore TET activity after each catalytic cycle. BER, base excision repair; 5-hmC, 5-hydroxymethylcytosine; 5-fC, 5-formylcytosine; and 5-caC, 5-carboxycytosine. (C) Role of ascorbate in demethylation of histone lysines. Jumanji C (JmjC) domain-containing histone demethylases catalyzes the removal of a methyl group from tri-, di-, and monomethylated ε-amino groups of lysine. Hydroxymethyl group, the product of demethylase activity, is unstable and spontaneously released in the form of formaldehyde. As with other dioxygenases, ascorbate is involved in the reduction of Fe(III) to Fe(II) to restore activity of JmjC demethylases after each reaction cycle.
HIF1/2 are the best known, but not the only, targets of prolyl hydroxylases PHD1–3. A recently discovered novel substrate of PHD2 is a serine-threonine protein kinase AKT which regulates a metabolic branch in mitogenic signaling from growth factor receptors. In response to growth factor stimulation, AKT is activated by phosphorylation at Thr308 which is reversed by protein phosphatase 2A. The recruitment of this phosphatase occurs in response to Pro hydroxylation of AKT by PHD2.62

Another biochemical process in the cytoplasm that has long been considered as dependent on ascorbate is carnitine production. The first step in carnitine biosynthesis is catalyzed by 6-N-trimethyllysine hydroxylase which is also a member of the Fe(II)/2-oxoglutarate-dependent family of dioxygenases. Carnitine is necessary for the transport of long-chain fatty acids into the mitochondria and ATP production via β-oxidation. Surprisingly, mice with a severe deficiency in tissue vitamin C (<2% relative to normal animals) did not show any significant changes in carnitine levels in serum and multiple organs.63 It remains unclear whether 6-N-trimethyllysine hydroxylase has a very low Km for ascorbate or is more promiscuous in the choice of the reducers for restoration of catalytic Fe(II) from Fe(III).

### ASCORBATE IN EPIGENETIC REGULATION

Reprogramming of gene expression is the most recently discovered fundamental biological process that is strongly affected by cellular vitamin C. The lateness of this discovery for ascorbate is related to the fact that biochemical mechanisms for how two critical steps in the control of gene expression, demethylation of 5-methylC in DNA and lysine demethylation of histones, have been defined only relatively recently. Cell culture and in vivo studies have clearly shown that ascorbate was important for the erasure of the epigenetic memory in embryonic stem cells or differentiated cells by acting as a cofactor for the family of ten–eleven translocation (TET1–3) proteins that remove cytosine methylation in DNA.64 Cytosine methylation occurs at its C5 position in CpG dinucleotides and usually acts as a transcription-repressive mark. TET enzymes are Fe(II)-dependent dioxygenases that catalyze a series of consecutive oxidations of 5-methylcytosine by first hydroxylating it to 5-hydroxymethylcytosine, which is further oxidized to 5-formylcytosine and 5-carboxycytosine. The last two oxidation products are then removed during base excision repair by thymine DNA glycosylase and replaced with the unmethylated cytosine (Figure 3C). In addition to in vitro assays with recombinant TET proteins, there is clear evidence for the importance of ascorbate in demethylation of 5-methylcytosine in various cultured cells and in vivo.65–67 Addition of ascorbate to human embryonic stem cells produced a dramatic impact on their epigenome, involving a widespread and specific DNA demethylation of 1847 genes.68 The ascorbate-demethylated group of genes included the gene sets that experienced a loss of 5-methylcytosine during differentiation of human embryonic stem cells and contained bivalent marks. A high ascorbate content of hematopoietic stem cells was necessary for their DNA demethylation, TET2-dependent gene expression signature and stimulation of differentiation and suppression of leukemia development.12,69 Erasure of epigenetic memory to create induced pluripotent stem cells was dependent on active DNA demethylation by TET proteins which were shown to be activated by ascorbate through reduction of Fe(III) to Fe(II).70 In addition to its differentiation effects in the developing brain, TET-dependent regional demethylation of DNA continuously occurs in neurons in the adult brain, which is important for neuronal activity and memory formation.72–74 In a mouse model of neurodegeneration, vitamin C deficiency impaired brain cognition and increased amyloid accumulation and deposition.75 Efficient DNA demethylation in hematopoietic stem cells required higher ascorbate concentrations than other ascorbate-promoted enzymatic processes.12,69 This property of TET-mediated DNA demethylation could be a reason for the unusually high ascorbate concentrations maintained in hematopoietic stem cells and neurons in vivo. These cells are also particularly sensitive to oxidative stress, as evidenced by the residence of hematopoietic stem cells in the hypoxic niche of the bone marrow and numerous neurodegenerative conditions associated with a compromised ability of the brain cells to deal with oxidants or their damage to proteins or DNA. Neurons appear to be continuously producing large amounts of superoxide that in addition to SOD activity requires normal levels of ascorbate for its efficient elimination.76

Post-translational histone modifications represent a second and more dynamic level of epigenetic regulation of gene expression in chromatin. Methylation of lysines located in the N-terminal tails is one of the well-characterized modifications that occurs in chromatin-deposited histones. The most frequent sites of lysine methylation are K4, K9, K27, K36, and K79 in histone H3 and K20 in histone H4. Lysine can be mono-, di-, and trimethylated at its N-6 position. Trimethylation of histone H3 at K4, K36, and K79 is associated with actively transcribed genes, whereas the presence of trimethylated K9 and K27 is associated with gene repression.77,78 Methylation status of histone lysines is dynamically regulated in response to transcriptional needs of cells. Although methylation of histone lysines was known for a long time, lysine demethylases have been discovered relatively recently. There are two classes of histone lysine demethylases that differ in their catalytic mechanisms and substrate specificity. The majority of lysine demethylases belong to the class of JmjC domain-containing Fe(II)/2-oxoglutarate-dependent dioxygenases (>20 enzymes) that collectively can demethylate lysines at any position in histones and are capable of demethylating mono-, di-, or trimethylated N-6. A pair of flavin adenine dinucleotide-dependent amine oxidases LSD1 (KDM1A) and LSD2 (KDM1B) represent a second class of histone lysine demethylases.79,80 LSD1 and LSD2 can remove a methyl group only from mono- or dimethylated lysines and act specifically on methylated lysine-4 in histone H3. Amine oxidation in the methylated lysine by LSD1/2 generates an unstable imine which is spontaneously hydrolyzed, releasing formaldehyde and yielding unmethylated lysine.

JmjC histone demethylases catalyze hydroxylation of a methyl group in lysines, which results in its dissociation in the form of formaldehyde (Figure 3D).81 Similar to other Fe(II)/2-oxoglutarate-dependent dioxygenases, JmjC histone demethylases require ascorbate for their optimal catalytic activity. In vitro histone lysine demethylation by JmjC hydroxylases was inhibited when ascorbate was omitted from the assay.82,83 In cellular models, ascorbate was identified as a critical factor for histone demethylation and epigenetic changes during reprogramming of somatic cells into induced pluripotent stem cells by acting as the activator of H3K36 and H3K9 demethylases.84,85

Ascorbate-induced demethylation of H3K9me2 in stem cells resulted from a specific activation of histone demethylases Kdm3a and Kdm3b. Importantly, vitamin C-stimulated Kdm3a/b-mediated H3K9me2 demethylation and TET-mediated DNA demethylation were independent processes.86 It is expected that nuclear ascorbate is a major stimulatory factor for activity of all
JmjC histone demethylases, although the degree of their sensitivity to its levels relative to other reducers of Fe(III) may vary.

**VITAMIN C IN CANCER THERAPY**

Two open-label studies in the 1970s led by Nobel prize-winning biochemist Linus Pauling and conducted in terminal cancer patients using intravenous administration of large doses of vitamin C have found significant beneficial effects on survival. However, follow-up double-blind, placebo-controlled clinical trials did not show any survival benefits of vitamin C in cancer patients, which effectively shut down for a long time the whole notion that vitamin C could work as an anticancer treatment. What was not appreciated until recently is the difference in the delivery of pharmacological doses of vitamin C between early positive studies and the later negative trials. In the follow-up trials, vitamin C was given orally, which is now known to provide only a limited increase in systemic ascorbate which is tightly regulated via the rates of absorption and urinary excretion. Therapeutic effects of large doses of vitamin C have been linked to a preferential accumulation of dehydroascorbic acid by cancer cells due to their common overexpression of GLUT transporters. In colorectal cancers, the overexpression of GLUT1 glucose transporter and hyperaccumulation of dehydroascorbic acid were specifically associated with mutations in KRAS and BRAF oncogenes. GLUT1 is also strongly upregulated by hypoxic signaling, and human clear cell renal carcinomas lacking the HIF1α/2α-targeting ubiquitin ligase VHL were more sensitive to killing by high doses of vitamin C through its hyperaccumulation. The cause of death in cancer cells that hyperaccumulated dehydroascorbic acid was linked to the exhaustion of energy resources and glutathione which were consumed by the reduction of dehydroascorbic acid to ascorbate. Overexpression of GLUT1 is associated with a metabolic reprogramming of cancer cells toward aerobic glycolysis (Warburg effect), which makes them more dependent on the inefficient ATP generation in glycolysis and, therefore, more vulnerable to energy depletion by dehydroascorbate reduction. Several studies have also linked a hyperaccumulation of ascorbate in cancer cells with the increased production of oxidants as a result of elevated levels of labile iron which, unlike its protein-bound form, readily catalyzes the Fenton reaction with the resulting generation of a highly toxic "OH radical. Oxidative and energy stresses by high vitamin C doses were associated with the induction of DNA damage and PARP hyperactivation-dependent cytotoxicity. A mechanistically distinct process based on the increased differentiation of stem cells operates in the suppression of leukemia by vitamin C. Patients with myelodysplastic syndrome and acute myeloid leukemia frequently carry inactivating mutations in TET2, which leads to a DNA hypomethylation phenotype and impaired differentiation of hematopoietic stem cells. Treatment with vitamin C mimicked TET restoration by enhancing 5-methylcytosine oxidation and promoting differentiation of stem cells and suppression of leukemia in mouse models and in primary patient-derived xenografts. A combination of vitamin C with clinically used drugs or ionizing radiation enhanced the killing of cancer cells of different histological origins. For many combined regiments, the increased killing of cancer cells can be attributed to their stressed energy metabolism and elevated ROS, diminishing their ability to survive chemotherapeutic or radiation treatments. In the case of bleomycin, a drug using Fe(II)-activated oxygen to oxidize and break DNA, ascorbate enhanced its DNA-damaging activity by acting as a very effective reducer of Fe(III) to Fe(II). The administration of high doses of vitamin C or possibly just restoration of its levels in ascorbate-deficient patients can also promote epigenetic and transcriptomic changes, making cancer cells more susceptible to specific cancer treatments, as it was found for ascorbate-induced sensitization of melanoma to BET inhibitors. Vitamin C has also diminished the aggressiveness of many cancers by inhibiting their growth, invasiveness, and metastasis. These beneficial responses were promoted by a decreased hypoxic signaling produced by a more efficient proline hydroxylation of HIF-1/HIF-2 and, consequently, their more rapid degradation in ascorbate-enriched cells. These transcription factors are known to promote motility, invasiveness, and metastasis of cancer cells and are markers of aggressive tumors. In ascorbate-deficient tumors, vitamin C supplementation could also boost the production of mature collagen and consequently, strengthen cell–cell adherence and the extracellular matrix, which would exert suppressive effects on the ability of cancer cells to migrate and invade surrounding tissue and reach distant sites. Another broad mechanism which has been overlooked so far in practically all preclinical models is the impact of vitamin C on antitumor immune responses. Humor tumor xenografts are grown in immunodeficient mice in order to avoid their immune rejection. Thus, this commonly used *in vivo* model excludes any contribution of the immune cells to antitumor responses. A recent study of lymphoma growth in the immunocompetent mice showed that vitamin C administration was synergistic with a clinically used anti-PD1 immune checkpoint treatment via increased DNA demethylation in tumor and immune cells. The combination therapy strongly increased intratumor infiltration of CD8⁺ T lymphocytes, granzyme B production by cytotoxic T cells and natural killer cells, and interleukin-12 biosynthesis by antigen-presenting cells compared with anti-PD1 alone. These findings raise a possibility that vitamin C may act as a general promotor of immune responses against many cancers that are responsive to anti-PD1/PD-L1 therapy.

**VITAMIN C DEFICIENCY IN CULTURED CELLS AND ITS REMEDIATION**

Various tissue culture media have been historically formulated to provide robust growth of cells by abundantly supplying them with energy-generating and macromolecule-building nutrients. Much less attention has been given to micronutrients, most of which are provided via the addition of fetal bovine serum to growth media. Vitamin C is a such micronutrient that is absent in the majority of cell culture media. The typical addition of 10% fetal bovine serum to culture media theoretically should provide the 1/10th of the physiological concentration of vitamin C, which in practice is much lower due to its loss during storage of serum and serum-supplemented media. Overnight fed cells typically contain low micromolar concentrations of vitamin C (5–20 μM), which is approximately 1% of its levels in cells in *vivo*. Vitamin C usually becomes undetectable in cultured cells at 2 days postfeeding. Primary and other cells growing in growth factor-supplemented synthetic media without addition of serum are completely devoid of vitamin C. Although rodents produce their own ascorbic acid, its synthesis occurs in the liver and all nonhepatic rodent cells in culture are also vitamin C deficient. It is unclear how much ascorbic acid biosynthesis remains in primary rodent hepatocytes which exhibit major...
Changes in their metabolic activities within a few days in culture. A monolayer of rodent hepatocytes would also need to continuously generate massive amounts of ascorbic acid to establish and maintain its physiological concentration in a very large volume of media relative to the number of cells in the dishes. In the absence of extracellular vitamin C in culture, primary and other cells rapidly lose their cellular ascorbate. Thus, it is likely that even rodent hepatocytes in culture are vitamin C deficient.

A full restoration of vitamin C levels in cells can be achieved by the addition of dehydroascorbic acid that readily enters cells via ubiquitously expressed GLUT transporters. Glucose competitively inhibits the uptake of dehydroascorbic acid, which hinders restoration of cellular ascorbate when high-glucose growth media are used. A switch to a low-glucose growth medium or a balanced salt solution (Krebs buffer) supplemented with low glucose (0.5–1 mM) and a regular serum percentage can result in the full restoration of physiological levels of ascorbate in cells after only 1–2 h of incubation. Expression of GLUT transporters is impacted by many factors, and the optimal concentrations of dehydroascorbic acid should be determined for each cell line. A more gradual delivery of vitamin C, but usually only to subphysiological levels, can be accomplished by the supplementation of culture media with ascorbate (daily additions of 50–200 μM). Even when ascorbate is used as a source of vitamin C supplementation, its easy oxidation in media can lead to a significant formation and uptake of dehydroascorbic acid. This is especially common for cells grown in serum-free media which contain large amounts of iron to compensate for the absence of transferrin-mediated iron uptake from serum. The addition of ascorbate-2-phosphate, a more stable derivative of ascorbate, typically leads to higher levels of cellular vitamin C, and it has a lower risk of the formation of ROS in the growth media. Ascorbate-2-phosphate is not redox-active, and the release of ascorbate occurs after its dephosphorylation in cells.

Methodological challenges associated with vitamin C supplementation in culture include its relatively rapid loss from cells. Consequently, cells should be used either shortly after restoration of ascorbate concentrations or periodic additions of a freshly prepared stock of ascorbic acid in cold H2O are employed for a prolonged maintenance of vitamin C levels in cells. Even though oxidation of ascorbate to dehydroascorbic acid still yields a biologically active form of vitamin C, it is relatively unstable and undergoes irreversible hydrolysis and further rearrangements. Incubations with excessive amounts of vitamin C can cause cytotoxicity, which can result either from the overloading of cells with dehydroascorbic acid when its reduction to ascorbate consumes important cellular reducers or from excessive amounts of ascorbate in O2-rich media where it reacts with iron and catalyzes the production of toxic ROS. Cancer cells are characterized by a strongly elevated uptake of glucose and overexpression of GLUT transporters, which makes them more sensitive to killing by high doses of dehydroascorbic acid.

Vitamin C restoration in cells can be assessed by its direct measurements and/or by a functional test such as a suppression of ATM-dependent phosphorylation of its targets KAP1 and/or CHK2 by chromate.

ASSAYS FOR VITAMIN C FORMS

Irrespective of the use of HPLC or microplate readers, the majority of the currently used assays for vitamin C in biological samples can be divided into two groups based on whether they measure its reduced (ascorbic acid) or oxidized (dehydroascorbic acid) forms. Ascorbic acid is measured by electrochemical detectors coupled with HPLC or by spectrophotometric methods detecting Fe(II) formation from Fe(III) (known as the FRASC assay - ferric reducing/antioxidant and ascorbic acid). Fluorescence-based assays (HPLC or microplate reader versions) detect the formation of dehydroascorbic acid conjugates with dyes such as o-phenylenediamine or 1,2-diamino-4,5-dimethoxybenzene. For measurements of total vitamin C by electrochemical detection, samples are treated with a reducer to convert dehydroascorbic acid into ascorbic acid. In the FRASC assay, dehydroascorbic acid is not detectable, which may result in a significant underestimation of total vitamin C due to its ready autoxidation in samples ex vivo. Assays based on the conjugation of dyes with dehydroascorbic acid use pretreatments of samples with ascorbate oxidase or TEMPOL to convert all vitamin C into its oxidized form. A detection of both oxidized and reduced forms of vitamin C has been challenging due to a very small percentage of the dehydroascorbic acid that is typically present in biological samples (tissues, cells or plasma/serum). Dehydroascorbic acid values were typically calculated from the differences in HPLC-electrochemical measurements between samples treated with reducer (total ascorbate) and untreated samples (ascorbate only). These estimates are not very precise or accurate as they derive from subtraction of a large number from only a slightly larger number. In assays utilizing dehydroascorbate conjugation with dyes, omission of ascorbate oxidase or TEMPOL still results in a very significant reaction of probes with ascorbate through catalysis of its oxidation. LC-MS is currently the only method for the direct determination of both abundant ascorbate and scarce dehydroascorbic acid.

CONCLUDING REMARKS: VITAMIN C IN MECHANISTIC TOXICOLOGY/PHARMACOLOGY

The multitude of antioxidant and biochemical functions of ascorbate makes it important that biologically relevant concentrations of vitamin C are included in studies of toxic and other stress responses in cultured cells. Vitamin C addition is routinely used in the field of stem cell biology, but this practice has not yet been commonly accepted in toxicology/pharmacology or stress biology in general even when epigenetic changes or ROS effects are specifically examined. For oxidants, cellular ascorbate was found to exert major effects on the type and magnitude of genotoxic events and epigenetic alterations. In addition to the changes in the magnitude of cytotoxicity or other general responses, ascorbate can dramatically alter the relative importance of specific signaling mechanisms activated in cells by carcinogens and DNA-damaging cancer drugs. The importance of extracellular ascorbate should also be considered when a chemical is known to react with ascorbate directly or with Fe(II) that is abundantly produced in ascorbate-supplemented media. Histone lysine demethylation is a dynamic process that occurs continuously during transcription cycles and in response to promoter binding by transcription factors. The important function of ascorbate as a cofactor for histone demethylases raises a possibility that transcriptomic and, consequently, downstream biological responses to many redox-inactive xenobiotics and stressors can also be affected to some extent by nuclear ascorbate.

The importance of ascorbate in the regulation of cellular reprogramming and gene expression via its cofactor functions for the DNA-demethylating TET enzymes and histone lysine...
demethylases raises a possibility that populations with vitamin C insufficiency could be particularly susceptible to adverse health effects of epigenetic toxicants. Mice and rats continuously produce large amounts of vitamin C, which may limit their sensitivity in capturing intergenerational and other epigenetic consequences of human malnutrition and its combination with other stressors or drugs. Clearly established roles of ascorbate in neurodevelopment, memory formation, and the development of neurodegenerative processes 72 raise a possibility that individuals with low systemic levels of vitamin C could be especially vulnerable to neurotoxic effects of chemicals and/or other stressors. This vulnerability would be difficult to detect in standard rodent models, although the use of Gal−/− mice or similarly deficient rats can be readily adapted to assess the impact of vitamin C insufficiency. Vitamin C levels in rodent and human tissues are generally similar, but concentrations of ascorbate in the lung lining fluids of laboratory rodents are more than 10× higher than those in humans. 110–113 This difference accounts for a rapid extracellular inactivation of soluble chromate in rodent lungs, making them resistant to carcinogenesis by this Cr(VI) form in contrast to its carcinogenicity in human lungs. Slowly solubilizing particles of chromium(VI) escape this detoxification by extracellular ascorbate and induce genotoxicity upon their intracellular dissolution.12 Oxidative stress in the lung is a common adverse effect for many inhaled toxicants. The use of standard laboratory rodents containing very high levels of ascorbate in the lung lining fluid represents an additional uncertainty in their use for the mechanistic assessment of human health risks by airborne contaminants.

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Biography

Anatoly Zhitkovich is Professor of Medical Science in the Department of Pathology and Laboratory Medicine at Brown University. He received B.Sc. with distinction in biochemistry from Belarusian State University and a Ph.D. in biochemistry from Belarusian Academy of Sciences. Prof. Zhitkovich has long-standing research interests in mechanisms of DNA damage by carcinogens and anticancer drugs and activation of stress signaling networks engaging DNA repair, cell cycle checkpoints, and cell death programs. One of the research directions in his lab has been the elucidation of the role of ascorbate in redox metabolism and the biological activity of carcinogenic metals and metal-containing drugs.

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