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Kyoto University
Protective roles of ascorbic acid in oxidative stress induced by depletion of superoxide dismutase in vertebrate cells

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SOD1, SOD2, antioxidant, DT40, conditional knockout cell
Abstract

Superoxide dismutases (SODs) are antioxidant proteins that convert superoxide to hydrogen peroxide. In vertebrate cells, SOD1 is mainly present in the cytoplasm, with small levels also found in the nucleus and mitochondrial inter-membrane space, and SOD2 is present in the mitochondrial matrix. Previously, we conditionally disrupted the SOD1 or SOD2 gene in DT40 cells and found that depletion of SOD1 caused lethality, while depletion of SOD2 led to growth retardation. Our observations from previous work showed that the lethality observed in SOD1-depleted cells was completely rescued by ascorbic acid. Ascorbic acid is a water-soluble antioxidant present in biological fluids; however, the exact target for its antioxidant effects is not known. In this study, we demonstrated that ascorbic acid offset growth defects observed in SOD2-depleted cells and also lowered mitochondrial superoxide to physiological levels in both SOD1- or SOD2-depleted cells. Moreover, depletion of SOD1 or SOD2 resulted in the accumulation of intracellular oxidative stress, and this increased oxidative stress was reduced by ascorbic acid. Taken together, this study suggests that ascorbic acid can be applied as a nontoxic antioxidant that mimics the functions of cytoplasmic and mitochondrial SODs.

Introduction

Aerobic organisms have evolutionally acquired antioxidant systems against reactive oxygen species (ROS) produced during normal metabolism, especially as a consequence of aerobic respiration in mitochondria. Although physiological amounts of ROS are necessary in aerobic organisms for cell signaling pathways and killing invading pathogens, excessive concentrations of ROS are considered to be an “oxidative stress”, damaging macromolecules such as DNA, proteins, and lipids [1-6]. The enzymes of superoxide dismutase (SOD) families are the first and most important endogenous antioxidants, converting superoxide
anions to hydrogen peroxide [7]. In mammalian cells, three types of SOD have been identified [8]. SOD1 is the major intracellular Cu/Zn-dependent SOD located in nuclei and the mitochondrial inter-membrane space. Immunocytochemical analysis showed that SOD1 is also localized in lysosomes and peroxisomes [9-12]. SOD2 is a Mn-dependent SOD localized in the mitochondrial matrix where it is involved in “dismutating” superoxide generated by the respiratory chain [13,14]. SOD3 is an extracellular Cu/Zn-dependent SOD located in the vascular extracellular space [8,15].

Oxidative stress resulting from an imbalance between superoxide and endogenous antioxidants has been widely considered to be involved in a number of diseases such as neurodegenerative disorders, cardiovascular dysfunction [16-19], and cancer development and progression [20,21]. In fact, nitric oxide, with anti-inflammatory and anti-coagulant properties, is rapidly inactivated by superoxide and converted to the strong oxidant peroxynitrite. Increasing evidence for the involvement of ROS in pathological conditions has been accompanied by searches for the antioxidant potential of natural as well as synthetic compounds.

Vitamin C is a water-soluble antioxidant and is also a cofactor for various enzymes [22]. Ascorbic acid, which is the reduced form of vitamin C, is a predominant chemical structure in biological fluids [23,24]. Although ascorbic acid is a potent antioxidant quenching potentially damaging ROS produced by normal metabolism [25,26], it also acts as a prooxidant, promoting the formation of many ROS, such as hydrogen peroxide and hydroxyl radicals [27-29].

The protective effects of ascorbic acid are believed to occur by scavenging ROS induced by various sources [22,30-32]. An increasing amount of data supports the scavenging ability of ascorbic acid; however, specific ROS species scavenged by ascorbic acid have not yet been identified. Previously, we constructed cell lines in which expression of either SOD1 or
SOD2 could be switched off by doxycycline (DOX) treatment in chicken DT 40 cells [33,34]. The availability of these cell lines allows us to specifically examine the biological effects of endogenous superoxide. We showed that SOD1 was essential for cell viability [34] and SOD2-depleted cells were viable, but exhibited slow growth [33]. Interestingly, we found that growth disturbances observed with SOD1-depleted cells were completely suppressed by ascorbic acid. These results suggested that ascorbic acid reduced elevated intracellular levels of superoxide caused by the depletion of SOD1.

In this study, we examine the effects of ascorbic acid in SOD2-depleted cells. Our data clearly showed that ascorbic acid effectively offset cellular phenotypes caused by the depletion of SOD1 or SOD2, including increased levels of superoxide and mitochondrial dysfunction as well as growth disturbances. This data demonstrates that ascorbic acid mimics the functions of cytoplasmic or mitochondrial SOD in chicken DT40 cells.

**Materials and Methods**

**Cell culture**

Cells were cultured in RPMI 1640 supplemented with 10% fetal bovine serum, 1% chicken serum, and kanamycin at 39 °C under 5% CO2. Conditional SOD1- and SOD2-knock-out DT40 cells were established as described previously [33,34] and the expression of transgenic human SOD1 and human SOD2 was suppressed by treatment with 1 µg/ml DOX. Ascorbic acid phosphate ester magnesium salt (APM) (Wako Pure Chemical Industries Ltd., Osaka, Japan) was dissolved in phosphate buffered saline (PBS): 137 mM NaCl, 2.68 mM KCl, 8.04 mM Na2HPO4, and 1.47 mM KH2PO4. To assay the effect of various antioxidants, cells cultured in the presence or absence of DOX for the indicated periods were treated with 200 µM APM, 5 mM N-acetyl-L-cysteine (NAC, Wako, Osaka, Japan), 50 µM 6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid (Trolox,
Sigma-Aldrich), 250 µM 1,2-Dihydroxy-3,5-benzenedisulfonic acid (Tiron, Sigma-Aldrich), or 50 µM 2-mercaptoethanol (EtSH, Sigma-Aldrich). The generation of growth curves has been described previously [33,34].

**Superoxide assay**

Intracellular generation of superoxide was detected using BES-So-AM (Wako Pure Chemical Industries Ltd.), a highly specific fluorescent probe for superoxide [35]. Briefly, cells were treated with 5 µM BES-So-AM for 20 min. After washing twice with PBS, cells were suspended in PBS, and fluorescent intensity was measured using FACScan (Becton Dickinson, Franklin Lakes, NJ, USA) [35]. Mitochondrial superoxide levels were measured by flow cytometry (FACS) after staining with the mitochondrial superoxide-specific dye MitoSOX red (Invitrogen). Cells were washed with PBS and incubated with 5 µM MitoSOX red for 10 min. Measurements were performed using FACScan (Becton Dickinson, Franklin Lakes, NJ, USA). MitoSOX red was excited at 396 nm [36] and data was collected in the 580/510 nm (FL2) channel. Quantifications were performed from the mean intensity of MitoSOX fluorescence from triplicates.

**Measurement of Mitochondrial Membrane Potential**

Mitochondrial membrane potential was measured using 5, 5’, 6, 6’-tetrachloro-1, 1’, 3, 3’-tetrethyl- benzimidazoylcarbocyanine iodide (JC-1; Molecular Probes, Eugene, USA) as previously described [37]. JC-1 is a cationic carbocyanine dye that presents itself as green fluorescent monomers at a low concentration (i.e., in cells with low mitochondrial functions or membrane potential). In cells with normal mitochondrial functions, a membrane potential–driven accumulation of these dyes results in the formation of yellowish-red fluorescent J-aggregates. Briefly, cells were washed with cold PBS and incubated with 5 µM
JC-1 for 30 min at 37 °C. After washing twice with cold PBS, cells were harvested and the intensity of fluorescence was analyzed using a FACScan (Becton Dickinson, Franklin Lakes, NJ) with excitation at 490 nm and emission at 530 nm.

_Determination of intracellular ROS levels_

Intracellular ROS were measured using 2’,7’-dichlorodihydrofluorescein diacetate (DCFH-DA; Molecular Probes, USA). Intracellular peroxide-dependent oxidation converts DCFH-DA to the fluorescent compound 2’,7’-dichlorofluorescein (DCF), as previously described [37]. After washing twice with PBS, cells were incubated with DCFA-DA (20 µM) for 30 min at 37 °C. After incubation, cells were harvested and resuspended in 50 mM HEPES buffer (5 mM HEPES, pH 7.4, 5 mM KCl, 140 mM NaCl, 2 mM CaCl₂, 1 mM MgCl₂, and 10 mM glucose). The fluorescence intensity was determined using FACScan (Becton Dickinson, Franklin Lakes, NJ) with excitation at 485 nm and emission at 530 nm.

_Results and Discussion_

Ascorbic Acid relief of growth perturbations in both SOD1 and SOD2 depleted cells

Previously, we generated conditional SOD1⁻/⁻ and SOD2⁻/⁻ cells in which the human SOD1 or SOD2 trans-gene was expressed under control of the tetracycline inducible promoter. Expression of each SOD trans-gene can be switched off by the addition of DOX leading to the complete depletion of exogenous SOD proteins [33,34]. These cells enable us to characterize cellular phenotypes caused by endogenous increases in superoxide due to the absence of SOD activity. Our previous work showed that SOD1 was essential for cell viability whereas SOD2 was not, but it was required for normal cell growth. Our observation from previous work is that the lethality observed in SOD1-depleted cells was completely rescued in the presence of ascorbic acid, Ascorbic acid phosphate ester magnesium salt
Ascorbic acid in aqueous solution can be oxidized easily and can be stabilized by phosphorylation [38]. Ascorbic acid phosphate has been used as a non-autooxidizable form of vitamin C in several cell culture studies including our previous studies [32,39]. To examine the effects of ascorbic acid on SOD2-depleted cells, we analyzed the mode of cell growth of SOD2-depleted cells in the presence of ascorbic acid. In spite of the different cellular localizations of SOD1 and SOD2, we found that ascorbic acid fully recovered the growth delay observed in SOD2-depleted cells (Figure 1B). We confirmed that SOD2−/− hSOD2 cells without DOX treatment proliferated normally in the presence of ascorbic acid (Supplemental Fig. S1). To examine other antioxidants for the rescue of growth deficiencies observed in SOD1-depleted or SOD2-depleted cells, we compared the growth properties of these cells in the presence of several antioxidants, NAC (N-acetyl-L-cysteine), a nontoxic dietary glutathione precursor, Tiron (4,5-dihydroxy-1,3-benzene disulfonic acid), a widely used antioxidant, and Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid), a cell-permeable, water-soluble vitamin E derivative. 2-Mercaptoethanol is a strong reducing agent for disulfide bonds and has antioxidant capacity by scavenging hydroxyl radicals. As shown in Figure 1, none of these antioxidants, except for ascorbic acid, rescued both SOD1-depleted cells and SOD2-depleted cells (Figure 1A and 1B). We should mention that original wild type cells proliferated normally in the absence or presence of DOX and/or ascorbic acid (Supplemental Fig. S2) or other antioxidants (data not shown).

Figure 1

Ascorbic acid suppresses increased mitochondrial or cellular superoxide levels in SOD1 or SOD2 depleted cells.

Ascorbic acid is known to be a non-enzymatic antioxidant exerting anti-mutagenic effects
by scavenging organic radicals [39,40]. We previously measured intracellular levels of superoxide using BES-So-AM and found that the levels of superoxide were increased in SOD1-depleted cells. These elevated superoxide levels were completely suppressed by ascorbic acid without changing the efficacy of suppression of the expression of hSOD1 by the tet-off promoter [34]. Even though BES-So-AM is a more practical superoxide probe than others, it should not rule out the possibility that BES-So-AM exhibited a fluorescent response to hydroxyl radicals in addition to superoxide [35]. In contrast to SOD1-depleted cells, we found that superoxide levels in the cytoplasm were not altered in SOD2-depleted cells. We speculated that it was probably due to the exclusive localization of SOD2 in the mitochondrial matrix (Figure 2A).

Next, we measured superoxide levels in mitochondria by the Mito-SOX red oxidation method. Superoxide levels in mitochondria were significantly elevated in SOD2-depleted cells, while there was no change in the level of superoxide in the cytoplasm (Figure 2B). Superoxide levels in mitochondria were also elevated in SOD1-depleted cells (Figure 2B). This data demonstrated that a SOD deficiency elevates the level of superoxide in mitochondria.

We examined whether ascorbic acid suppresses the increased superoxide levels in mitochondria as efficiently as that in the cytoplasm. As shown in Figure 2B, increased superoxide levels in mitochondria, both in SOD1-depleted and SOD2-depleted cells were completely suppressed by ascorbic acid. These results indicate that ascorbic acid scavenges superoxide in mitochondria effectively and helps maintain a physiological level of superoxide.

Figure 2
Increased mitochondrial superoxide levels could induce dysfunctions in mitochondria. The mitochondrial membrane potential (ΔΨ) is an important criterion for assessing mitochondrial function and has been used as an indicator of mitochondrial integrity. To determine the impact of elevated mitochondrial superoxide levels on mitochondrial function, the mitochondrial membrane potential (ΔΨ) was analyzed using a unique cationic dye, JC-1 (5,5′,6,6′-tetrachloro-1,1′,3,3′- tetraethylbenzimidazolylcarbocyanine iodide). Significantly, abnormal up-transitions in mitochondrial permeability were observed in SOD2-depleted cells. While similar elevated superoxide levels in mitochondria were detected in SOD1−/−hSOD1-depleted cells (Figure 3), no significant change in the mitochondrial membrane potential was observed in SOD1-depleted cells (Figure 3). Elevations in ΔΨ or mitochondrial hyperpolarization are an indication of abnormal mitochondrial status. For example, systemic lupus erythematosus is characterized by abnormal T-cell activation and cell death, which depend on the controlled production of ROS intermediates and ATP in mitochondria; lupus T-cells exhibit mitochondrial dysfunction and persistent elevations in mitochondrial membrane potential that predisposes lupus T-cells to cell death [41]. It has been assumed that increased levels of superoxide, due to the depletion of SOD2, generate cellular circumstances similar to those in lupus T-cells. Because SOD1 and SOD2 are localized in different mitochondrial fractions, the mitochondrial inter-membrane and matrix respectively, these results suggest that increased superoxide in the mitochondrial matrix is more harmful to mitochondrial function than that in the mitochondrial inter-membrane. SOD2 may play a critical role in maintaining superoxide levels in the mitochondrial matrix, thereby protecting mitochondria. The abnormal mitochondrial membrane potential observed in SOD2-depleted cells was completely recovered by treatment with ascorbic acid. Ascorbic acid scavenges superoxide effectively in mitochondria and maintains superoxide in mitochondria at physiological levels, keeping the mitochondrial membrane potential at
normal levels in the absence of SOD2 in DT 40 cells.

Figure 3

*Ascorbic acid suppresses overall increased oxidative stress in SOD1 or SOD2 depleted cells*

Inadequate removal of reactive oxygen species (ROS) could result in overall oxidative stress in cells. However, superoxide itself is not highly reactive and impermeable [42,43,44]. An increased level of superoxide triggered by a SOD deficiency could generate other secondary ROS, leading to elevations in overall oxidative stress. It is known that secondary ROS derived from superoxide, such as hydrogen peroxide and hydroxyl radicals, are more reactive and penetrate biological membranes more efficiently than superoxide [42,44]. In addition, superoxide reacts with nitric oxide (NO), produced as a consequence of oxidative stress, and strong oxidants such as peroxynitrite (ONOO\(^-\)) are produced. To examine the overall levels of oxidative stress, we stained SOD1-depleted or SOD2-depleted cells with dichlorodihydrofluorescein (DCFH). DCFH is a cell-permeable fluorescence dye and reacts with a broad spectrum of cellular ROS. Overall oxidative stress was increased both in SOD1-depleted and SOD2-depleted cells. Unexpectedly, a higher level of oxidative stress was detected in SOD2-depleted cells than in SOD1-depleted cells, even though a similar level of superoxide was detected (Figure 4). It is not clear what accounts for this difference in the two cell-lines; we would like to suggest the possibility that potential molecules that become ROS after interacting with superoxide may be more abundant in the mitochondrial matrix than those in the mitochondrial inter-membrane space. This data may also indicate that secondary ROS generated by superoxide, not superoxide itself, causes the abnormal mitochondrial membrane potential observed in SOD2-depleted cells (Figure 3).

Importantly, ascorbic acid suppressed increased levels of oxidative stress and maintained
oxidative stress at physiological levels both in SOD1-depleted and SOD2-depleted cells (Figure 4). Ascorbic acid suppresses the formation of secondary ROS by scavenging superoxide and lowers the level of overall oxidative stress in SOD depleted cells.

Figure 4

Oxidative stress and mitochondrial dysfunction have been linked to neurodegenerative disorders, such as Parkinson’s and Alzheimer’s diseases [45,46]. Recently, it was reported that Sirt3-dependent mitochondrial SOD2 activity is an important factor in preventing cancer permissive circumstances [47]. Accumulated evidence for the involvement of oxidative stress in various pathological conditions has encouraged the search for natural and synthetic antioxidants as well as the antioxidant potentials of natural products. Thus, developing screening systems that specifically identify molecules/chemicals that can replace SOD functions is vital. In conclusion, we showed that ascorbic acid scavenges superoxide in the cytoplasm and mitochondria effectively and restores physiological levels of superoxide and oxidative stress in SOD-deficient cells, thereby “mimicking” the enzymatic functions of SODs. Our SOD1-depleted and SOD2-depleted cell lines should be powerful tools for screening and identifying molecules/chemicals that mimic the functions of SODs.

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Japan.
References


1998;21:45-54.


Figure legends

Figure 1. Effect of antioxidants on the growth curves of SOD1- or SOD2-deleted cells. Indicated knockout cells were cultured with or without 1μg/ml DOX and/or various antioxidants.

Figure 2. Suppression of increased superoxide levels in SOD1- or SOD2-depleted cells by ascorbic acid. (A) Cytoplasmic superoxide levels in SOD2-depleted cells. Levels of superoxide were measured with BES-So-AM. (B) Mitochondrial superoxide levels in SOD1- or SOD2-depleted cells. Mitochondrial superoxide levels in the indicated cells were measured with Mito SOX red. In either case, cells were treated with or without 1μg/ml DOX and 200μM ascorbic acid for 96 hours. Data from flow cytometric analyses are presented as mean fluorescence intensity (MFI). The MFI of cells not treated with DOX was used as the standard (arbitrary unit [a.u.] = 1). Error bars indicate standard errors based on three independent experiments.

Figure 3. Normalization of up-shifted mitochondrial membrane potential in SOD1- or SOD2-depleted cells. Cells were treated with or without 1μg/ml DOX and 200μM ascorbic acid for 96 hours. The mitochondrial membrane potential (ΔΨm) was assessed with JC-1. The ratio of red to green fluorescence is a semiquantitative indicator of ΔΨm. Mean green fluorescence (FL-1 channel) and mean orange-red fluorescence (FL-2 channel) were quantified using flow cytometry. Ψm was expressed as the mean red to mean green fluorescence ratio. Ψm of cells not treated with DOX was used as the standard (a.u. = 1). Error bars indicate standard errors based on three independent experiments.

Figure 4. Suppression of increased cellular oxidative stress in SOD1- or SOD2-depleted cells
by ascorbic acid. Cells were treated with or without 1μg/ml DOX and /or 200μM ascorbic acid for 96 hours. Cellular oxidative stress generated within cells was detected using the fluorescent probe DCFH oxidation. Data from flow cytometric analyses are presented as MFI. The MFI of cells not treated with DOX was used as the standard (a.u.=1). Error bars indicate standard errors based on three independent experiments.
Figure 1

(A) SOD1<sup>−/−</sup>:hSOD1

(B) SOD2<sup>−/−</sup>:hSOD2
Figure 2
Figure 3
Figure 4

![Bar chart showing relative fluorescence levels for different conditions. The chart compares control, DOX, Control, DOX, APM, DOX+APM and control, DOX, APM, DOX+APM for both Wild Type and SOD1^{+}\text{hSOD1} and SOD2^{+}\text{hSOD2} conditions. The significance levels are indicated by * (P<0.05), ** (P<0.01), and *** (P<0.005).]