Forever young
Mechanisms of natural anoxia tolerance and potential links to longevity

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While mammals cannot survive oxygen deprivation for more than a few minutes without sustaining severe organ damage, some animals have mastered anaerobic life. Freshwater turtles belonging to the Trachemys and Chrysemys genera are the champion facultative anaerobes of the vertebrate world, often surviving without oxygen for many weeks at a time. The physiological and biochemical mechanisms that underlie anoxia tolerance in turtles include profound metabolic rate depression, post-translational modification of proteins, strong antioxidant defenses, activation of specific stress-responsive transcription factors, and enhanced expression of cyto-protective proteins. Turtles are also known for their incredible longevity and display characteristics of “negligible senescence.” We propose that the robust stress-tolerance mechanisms that permit long term anaerobiosis by turtles may also support the longevity of these animals. Many of the mechanisms involved in natural anoxia tolerance, such as hypometabolism or the induction of various protective proteins/pathways, have been shown to play important roles in mammalian oxygen-related diseases and improved understanding of how cells survive without oxygen could aid in the understanding and treatment of various pathological conditions that involve hypoxia or oxidative stress. In the present review we discuss the recent advances made in understanding the molecular nature of anoxia tolerance in turtles and the potential links between this tolerance and longevity.

Introduction

Some two billion years ago primitive life forms learned how to use oxygen as the final acceptor in their electron transport pathways. The high redox potential of oxygen allows it to easily accept electrons from reduced substrates, making it extremely useful for energy production. The ability to extract greater amounts of energy from organic molecules by using oxygen-linked catabolism has driven life to evolve into high complexities and has made oxygen vital to many life forms on Earth. Mammals are particularly sensitive to oxygen deprivation and situations of hypoxia or anoxia can lead rapidly to severe tissue damage or even death. In contrast, some ectothermic vertebrates are particularly well-adapted to surviving oxygen limitation. Various species of turtles have very well-developed capacities for living without oxygen (called anaerobiosis).1,2 For example, freshwater turtles living in northern regions of the US and Canada typically hibernate under water to elude freezing temperatures on land. However, bodies of water often become ice-locked, limiting the ability for animals to surface for lung breathing. Some turtle species compensate for this with a good capacity for extrapulmonary gas exchange across other epithelia whereas others have perfected strategies that allow survival without breathing oxygen for three months or more.3 Those species that tolerate extreme hypoxia or anoxia employ various biochemical and physiological mechanisms for survival, the major ones including prehibernation accumulation of huge glycogen stores to fuel anaerobic glycolysis, the use of the shell and skeletal systems for buffering lactate accumulation and minimizing acidosis, and the capacity for profound metabolic rate depression to only 10–20% of the corresponding aerobic rate. Metabolic depression allows tissues to minimize their energy needs, and therefore survive for extended periods of time using the ATP generated by anaerobic glycolysis alone. In addition to the challenges presented to tissues by severe oxygen limitation, turtles must also deal with the potential for oxidative stress occurring with the re-introduction of oxygen. The mitochondrial electron transport chain becomes reduced during hypoxia/anoxia because there is little or no oxygen to accept electrons. This makes it particularly susceptible to generating reactive oxygen species (ROS) upon reoxygenation when the oxygen tension is high. This can be extremely damaging to cells and indeed, in situations of ischemia-reperfusion, mammalian tissues show profound damage caused by ROS during the recovery phase.4,5 Situations of environmental oxygen variability, such as those experienced by diving or hibernating turtles, could present many opportunities for oxidative stress and oxidative damage, and yet, turtles emerge unharmed from repeated cycles of anoxia and reoxygenation.
Turtles are also known for their extraordinary longevity. Lifespans of over a hundred and fifty years have been documented for some species of the Chelonia, and more importantly, these animals do not appear to display the senescent phenotypes characteristic of many other species. Humans have long sought a cure for aging, and turtles present a unique model of “negligible senescence.” Criteria for “negligible senescence” include: (1) mortality rate should not increase with age and (2) reproduction should not decrease with age. Studies on painted turtles (Chrysemys picta marginata) showed that survivorship of old turtles does not differ from that of their younger counterparts and no decline in the reproductive output is apparent with age. Turtles, therefore, appear to meet the criteria necessary for “negligible senescence.” Other research has shown that telomere length, which generally decreases with age, does not appear to change between embryos and adults of European freshwater turtles (Emys orbicularis). These characteristics could make turtles a valuable model for studies that attempt to understand how cells can avoid the wear and tear of daily life.

A potential link between anoxia tolerance in turtles and their extraordinary longevity has been previously suggested. The mechanisms that turtles employ to survive anoxia might be both directly and indirectly linked to longevity. For example, in order to deal with a variable oxygen environment, turtles must have extraordinary stress-resistance mechanisms. Resistance to stress is generally correlated with longevity and organisms that are tolerant of one kind of stress are typically more likely to be tolerant to others, since many cytoprotective mechanisms operate over a wide variety of stresses. Whereas aging is often characterized by decreased stress-resistance and subsequent cellular degeneration, an animal that has robust stress-resistance mechanisms is more likely to deal better with aging. In addition, an understanding of the molecular nature of anoxia tolerance in turtles can lead to improved understanding of many age-related pathological conditions that involve oxygen restriction/deprivation (e.g., stroke or myocardial ischemia) or oxidative stress (e.g., Parkinson, Alzheimer disease). Finally, such studies might suggest ways of making cells and tissues of oxygen-sensitive animals, such as humans, more tolerant of oxygen limitation. This could lead to improved techniques for handling ischemic insults to organs, to aid in preserving organ viability during surgery, and even to enhanced technology for use in hypothermic- or cryopreservation of tissues and organs.

In the present review we discuss recent advances in the studies of the molecular nature of anoxia tolerance in turtles, with emphasis on the potential links between these mechanisms and the negligible senescence shown by these animals.

**Oxidative Stress and Aging**

Whereas oxygen-based metabolism has allowed life to evolve to ever new complexities, this did not come without a price. The catch is oxidative stress, the damage done by ROS generated from single electron reductions of oxygen. When ROS generation exceeds the capacity of antioxidant defenses, oxidative stress ensues and has been implicated in cellular degradation during aging as well as in a variety of disease states.

Formation of most ROS occurs at the site where the vast majority of intracellular oxygen is consumed, the mitochondria. It is here that reduced substrates donate their electrons to form an electrochemical gradient, resulting ultimately in the production of ATP, the energy currency of the cell. This occurs through a series of oxidative reactions conducted by protein complexes on the inner membrane of the mitochondria. Leakage of electrons from these complexes results in the one-electron reduction of oxygen to form superoxide, the precursor to most ROS. Dismutation of superoxide results in the production of hydrogen peroxide, whereas the subsequent interaction between superoxide and hydrogen peroxide in the Haber-Weiss reaction or cleavage of hydrogen peroxide in the Fenton reaction results in the formation of hydroxyl radicals. 15 It is estimated that approximately 1–2% of the oxygen consumed during respiration is reduced to superoxide.

The chemistry of superoxide differs dramatically based on its environment. In aqueous solutions, superoxide is a weak oxidizing agent and a much stronger reducing agent. Superoxide rapidly disappears in aqueous solutions due to its dismutation reaction, in which hydrogen peroxide and oxygen are formed:

\[
O_2^- + O_2^- + 2H^+ \rightarrow H_2O_2 + O_2
\]

Hydrogen peroxide is a weak oxidant and a weak reducing agent and is relatively stable in the absence of transition metals. However, whereas both superoxide and hydrogen peroxide have low oxidative toxicity themselves, toxicity themselves, they are both components of the net Haber-Weiss reaction that readily produces hydroxyl radicals, catalyzed by copper or iron ions:

\[
O_2^- + H_2O_2 \xrightarrow{Fe^{2+}} O_2 + \cdot OH + OH^-
\]

The hydroxyl radical is highly reactive and is responsible for most of the damage incurred to biological macromolecules by ROS. The main source of biological hydroxyl radicals is the metal-catalyzed breakdown of hydrogen peroxide by the Fenton reaction, one-half of the net Haber-Weiss reaction:

\[
M^{n+} + H_2O_2 \rightarrow M^{(n+1)+} + \cdot OH + OH^-\]

Because of their short half life (less than 1 ns), hydroxyl radicals attack molecules very close to their site of formation and their very high reactivity makes them virtually impossible to scavenge. The reaction of hydroxyl radicals with biomacromolecules produces new radicals, usually with lower reactivity. Once unleashed upon the intracellular milieu, ROS react with many cellular macromolecules, including lipids, proteins and DNA, inflicting damaging effects on multiple subcellular structures.

**ROS damage to macromolecules.** Lipid peroxidation is major source of damage causing cellular injury and death (reviewed in refs. 19 and 20). It is caused by the interaction of ROS (chiefly hydroxyl radical) with unsaturated bonds in membrane lipids.
Polyunsaturated fatty acids in membrane phospholipids are particularly susceptible to peroxidation, which can result in the rearrangement or loss of double bonds or degradation of lipid side chains. This leads to disturbance of membrane structure, causing changes in membrane fluidity, permeability, alterations in ion transport and inhibition of metabolic processes. Lipid peroxidation in the mitochondria can suppress mitochondrial metabolism, as well as affect vital mitochondrial functions such as respiration, inner membrane barrier properties, maintenance of mitochondrial membrane potential and mitochondrial calcium buffering capacity.

Almost all kinds of amino acid residues in proteins are potential targets for oxidation by ROS (reviewed in ref. 28). Oxidation of some amino acid residues (lysine, arginine, proline) can also lead to the formation of carbonyl derivatives, that are considered to be the most relevant products of free radical attack on proteins. The presence of carbonyl proteins in cell and tissue samples has become widely accepted as a marker for oxidative stress. Copper or iron binding sites on proteins are other important targets of ROS, since these sites can serve as centers for additional production of hydroxyl radicals via the Fenton reaction. These radicals can then attack side chains of the amino acid residues at the metal-binding site.

ROS appear to produce several forms of DNA damage, including modification of all bases, production of base-free sites, deletions, frame shifts, strand breaks, DNA-protein cross-links and chromosomal rearrangements (reviewed in ref. 17). Modifications of DNA can be mutagenic and might contribute to cancer, neurodegenerative diseases and aging. The hydroxyl radical is known to react with all components of the DNA molecule—the purine and pyrimidine bases as well as the deoxyribose backbone. In addition to base and sugar modifications, covalent DNA-protein cross-links can also form by exposure to free radicals. The mechanism for the formation of these cross-links involves the addition of a DNA base radical to an aromatic amino acid of a protein or the combination of a DNA base radical with an amino acid radical.

Free radical damage and aging. Aging is characterized by an increase in many detrimental changes at different levels of biological organization. The idea that ROS are among the main perpetrators of cellular damage has gained much support over the past few decades (reviewed in ref. 36 and 37). Lipid damage has been shown to increase with age, one result of this being a buildup of lipofuscin, a pigment that accumulates with age and correlates with animal senescence. Tissue damage in the form of oxidized and cross-linked proteins also increases as a function of age. For example, 30–50% of the total cellular proteins in an old animal might be oxidized, possibly accounting for the observation that the catalytic activity of many enzymes decreases with age. Evidence also exists that the rate of degradation of oxidized proteins decreases with age. Oxidative DNA damage also increases with age. Since such damage can lead to mutations and the subsequent synthesis of dysfunctional proteins, it is likely to be a major contributor to many age-associated detrimental changes.

A wide range of evidence shows mitochondrial decline as a function of aging. Since mitochondria consume >90% of the oxygen utilized in cells, and are the major site of ROS production, they appear to bear the brunt of the free radical damage observed in cells during aging. Many different studies have shown that mtDNA mutations increase with age in mammals, especially in post-mitotic highly aerobic tissues. One consequence of this damage is decreased ATP production, reducing the energy to drive many cellular processes. In addition, it has been shown that mitochondria of older animals produce significantly more ROS than mitochondria of younger animals. This can be a result of an escalating cycle, whereby damaged mitochondria leak more free radicals to inflict additional self-damage, as well as damage to the rest of the cell.

Antioxidant Defenses in Turtle Anoxia Tolerance

Anoxia tolerant turtles appear to possess excellent antioxidant defenses that aid in the damage-free survival of multiple episodes of anoxia-reoxygenation. Studies with red-eared sliders (Trachemys scripta elegans) show that these include both constitutive and inducible defenses.

Constitutive antioxidant defenses. T.s. elegans maintain high constitutive activities of various antioxidant enzymes, including catalase, superoxide dismutase (SOD) and alkyl hydroperoxide reductase. These activities are much higher than those in other ectothermic vertebrates, and are actually comparable to mammalian activities even though the metabolic rate (and oxygen consumption) of turtles is much lower than mammalian values. Tissue pools of glutathione are also higher in turtle organs compared to other ectotherms, and ascorbic acid levels in turtle brain cortex are 2–3 times higher than in mammals. High activities of antioxidant defenses also appear to serve freezing-induced anoxia/ischemia. For example, freeze-tolerant wood frogs (Rana sylvatica) that undergo cycles of ischemia/reperfusion upon freezing and thawing also show high activities of antioxidant enzymes, significantly higher than those activities in freeze-intolerant frogs.

The structure of turtle antioxidant enzymes might also be especially suitable for frequent anoxic excursions. For example, turtle glutathione reductase (GR), an enzyme that catalyzes the reduction of oxidized glutathione (GSGG), has a higher affinity for GSSG than the enzyme in most other vertebrates, allowing for better enzyme efficiency under different redox conditions. Turtle GR also showed high activity over a broad pH range (including at the low pH typical of anoxic conditions). In another recent study, two alpha classes of GSTs were found in T.s. elegans liver, a homodimer and a heterodimer. Both exhibited a major reduction in specific activity in response to anoxia and the heterodimer showed a major change in substrate preference, indicating an anoxia-responsive stable modification of the enzyme. Therefore, it appears that GST undergoes modifications in response to changing oxygen availability to optimize its function.

These mechanisms could help the turtle to avoid oxidative damage during situations of oxygen variability. Indeed, it was shown that the ratio of GSH/GSSG, which decreases under oxidative stress, actually increases during recovery from anoxia.
exposure in turtles (compared to control) suggesting that no oxidative stress occurs during reoxygenation. In addition, oxidative damage products were largely unaffected during anoxia/reoxygenation in turtle organs, as evident from minimal changes in lipid peroxidation products. Similarly, organs of freezertolerant C. picta hatchlings also showed no increase in markers of oxidative damage after exposure to freezing, supercooling or hypoxia.

The constitutive presence of such strong antioxidant defenses in turtle tissues could be directly linked to longevity. Maintenance of strong constitutive defenses throughout the turtle’s lifetime would help it to avoid the day-to-day accumulation of oxidative damage to macromolecules that is observed in mammals and, hence, extend long-term organ viability.

Inducible antioxidant defenses. In addition to the maintenance of constitutively-high antioxidant defenses, induction of proteins associated with antioxidant defense has also been observed in turtle organs in response to anoxia. Upregulation of the transcripts coding for the heavy chain of ferritin occurs in response to anoxia in liver of adult T.s. elegans. In addition, the heavy and light chains of ferritin were upregulated in response to anoxia or freezing exposures in the heart and liver of hatchling C.p. marginata. Ferritin plays a key role in sequestering iron in cells. As previously mentioned, free ferrous iron (Fe^{2+}) can participate in the Fenton reaction, catalyzing hydroxyl radical formation. Therefore, it is of benefit to increase the capacity to sequester iron during situations of oxygen variability, and thereby decrease the potential for ROS formation. The importance of ferritin upregulation under situations of anoxia/ischemia is observed across phylogeny; for example, ferritin heavy chain was upregulated in an anoxia-tolerant marine snail, whereas the light chain was freeze-responsive in wood frog heart.

Heme oxygenase-1 (HO-1) is another enzyme that might have antioxidant functions during anoxia/reoxygenation episodes. It is involved in the degradation of heme, a major source of redox active iron in the cell, and its induction is also typically tied to the simultaneous upregulation of ferritin, which sequesters the iron released by the heme degradation reaction. HO-1 was upregulated in response to anoxia in turtle brain, which might be a further demonstration of the importance of minimizing redox-active iron during anoxia. In mammals, the expression of HO-1 is known to be induced in response to various forms of oxidative stress and its expression is believed to be protective in different disease states. The important antioxidant properties of HO-1 also stem from biliverdin, one of the products of heme catabolism by HO-1. Biliverdin and its catabolite, bilirubin, have antioxidant properties with demonstrated beneficial effects in a number of disease/injury states including ischemia-reperfusion, organ transplantation, graft rejection and autoimmune inflammation. The transcript levels of several other antioxidant enzymes have been shown to increase in response to anoxia in the turtle. Both manganese and copper/zinc forms of SOD increase in turtle liver in response to anoxia. In addition, array screening showed that several antioxidant genes were upregulated by anoxia exposure in heart and liver of hatchling C. p. marginata: SOD-1, glutathione peroxidase (GPX) isozymes 1 and 4, glutathione-S-transferase (GST) isozymes M5 and A2 and peroxiredoxin 1. The activity of the antioxidant enzyme, γ-glutamyltranspeptidase, also increases significantly during freezing in livers of both R. sylatica and C. picta, whereas catalase activity increased under anoxia exposure in livers of several hatching turtle species. In addition, the protein levels of the transcription factor Nrf2 and several downstream antioxidant enzymes were significantly elevated in the brains of C.p. marginata hatchlings in response to freezing exposure.

Protein levels of some members of the peroxiredoxin family were elevated in response to anoxia in several tissues of T.s. elegans (Krivoruchko and Storey, unpublished results). As in the case of ferritin and HO-1, upregulation of these enzymes in response to anoxia could serve as a preparatory step to deal with a burst of ROS production when tissue reoxygenation occurs, and might be responsible, at least in part, for the lack of oxidative damage observed in turtle tissues during aerobic recovery after anoxia exposure.

Besides their role in antioxidant defenses, the anoxia-responsive upregulation of peroxiredoxins might be of additional importance. Peroxiredoxins function in reducing H₂O₂, a molecule that has second messenger actions. Metabolic rate depression is a crucial factor in anoxia tolerance, and is expected to include reduced cell responsiveness to growth signals. This could potentially be achieved by reducing the levels of intracellular second messengers such as H₂O₂, which mediate growth factor and cytokine effects. Interestingly, upregulation of peroxiredoxins has also been shown to occur during mammalian hibernation, another situation of hypometabolism.

Other Protective Mechanisms

While strong antioxidant defenses clearly play a key role in protecting turtle cells during anoxia and reoxygenation, other protective proteins also contribute to survival under these stressful conditions. These include various molecular chaperones, stress-responsive transcription factors and pro-survival proteins.

High constitutive HSPs. Proteins that are particularly sensitive to changes in intracellular pH, redox state and ionic milieu, which occur in cells during anoxia and reoxygenation, can lose their native folded conformation to become misfolded. Proliferation of heat shock proteins (HSPs) is one of the best known cytoprotective mechanisms in response to stress. Figure 1 shows a generalized scheme for activation of the heat shock response in vertebrates. HSPs act as molecular chaperones, helping to fold nascent proteins, as well as aiding in the refolding of misfolded proteins under stress conditions. By their chaperone action, HSPs help to preserve cellular proteins and extend their functional life. Increased expression of HSPs has been shown to be a widespread response to many environmental stresses. HSPs also play a protective role in a number of pathological conditions associated with oxidative stress, including ischemia, cardiovascular disease and neurodegeneration. Studies have found significantly higher levels of Hsp60 in the hearts of anoxia-tolerant painted turtles compared to
Anoxia intolerant softshell turtles, rabbits and rats. Hsp60 is a predominantly mitochondrial chaperone involved in the folding of proteins entering the mitochondria. Hsp60 also has protective effects against oxidative stress. Levels of this protein are reduced with aging, possibly accounting for some age-associate cell damage. High constitutive levels of Hsp60 are of particular interest, since oxidative damage to the mitochondria appears to be instrumental in the escalation of age-related damage. The presence of high levels of protective HSPs, such as Hsp60, in the mitochondria could protect mitochondrial proteins from unfolding and ensure long-term mitochondrial viability as well as potentially prevent age-associated increases in ROS-production by mitochondrial proteins involved in oxidative reactions. This is certainly an interesting possibility that warrants further investigation.

High constitutive levels of Hsp72 have also been reported in turtle brain. Hsp72 is known as the inducible HSP70 family member, and is generally found at very low levels under normal conditions. The constitutive expression of Hsp72 in turtle brain suggests that it has some importance in neuroprotection in this animal. Members of the HSP70 family and their co-chaperones play a crucial role in protein sorting and quality control by selecting and directing aberrant proteins to the proteasome for degradation. A decline in protein quality control is related to many neurodegenerative diseases involving aggregation-prone proteins, including Parkinson disease, amyotrophic lateral sclerosis, prion diseases, Alzheimer disease and CAG-repeat and polyQ expansion diseases, including Huntington disease. The constitutive presence of Hsp72 might be instrumental in preventing protein aggregation and age-associated neurodegenerative diseases in the turtle brain.

Anoxia-inducible molecular chaperones. As in the case of antioxidant proteins, HSPs and other molecular chaperones also show inducibility in response to anoxia in turtle tissues. This included Hsp25/27, Hsp40, Hsp60, Hsp70, Hsc70 and Hsp90. Array screening of turtle brain comparing aerobic and anoxic animals also showed upregulation of transcripts coding for αB-crystallin, another known chaperone. In addition, the heat shock transcription factor (HSF1) that is responsible for the activation of the heat shock response was activated in response to anoxia in turtle tissues. Activation of the heat shock response in response to anoxia might help maintain protein stability under anoxic conditions as well as serve as a preparative mechanism for re-oxygenation, since increased HSP expression might also actively prevent damage following oxidative stress.

Small HSPs have been shown to have key functions in protection against protein unfolding, especially under conditions of oxidative stress. The αB-crystallin is known for its role as a molecular chaperone of cytoskeletal proteins that would otherwise aggregate and precipitate, something that has been strongly linked with neurodegenerative diseases. αB-Crystallin also responds to oxidative stress and its upregulation in turtle brain suggests an important role for this protein in neuronal protection against anoxia. Besides protecting against protein misfolding during stress, Hsp27 has been shown to possess antioxidant properties. It is able to hold glutathione in its reduced form, even under oxidative conditions, and by doing so restore redox homeostasis.

Some HSPs, including Hsp27, Hsp70 and Hsp90 can also regulate apoptosis by binding and inhibiting members of the apoptotic cascade. For example, in stressed cells, Hsp70 and Hsp90 can bind Apaf-1 (apoptosis protease activating factor-1) and by doing so inhibit events downstream of Apaf-1 activation, such as activation of caspases. Hsp70 can also sequester released AIF from the mitochondria and prevent caspase-independent cell death. Hsp27 can inhibit apoptosis by either inhibiting the release of mitochondrial cytochrome c, or by binding to cytochrome c directly.

In mammals, the HSP response declines with aging. Age-dependent waning of the HSP response has been reported in neuronal tissues, skeletal and cardiac muscle and liver. The loss of HSP inducibility with age could contribute to age-related degenerative diseases in humans. On the other hand, since maintenance of HSP inducibility is crucial to anoxic survival by the turtle, this, in conjunction with high constitutive expression of HSPs, might contribute to their negligible senescence.

NFκB. The transcriptional responses involved in anoxia-tolerance and the transcription factors involved in facilitating these responses are now becoming an active field of research. NFκB is an oxygen-responsive transcription factor, known to be activated in response to a variety of stimuli and controlling the expression of hundreds of target genes, including genes involved in the
immune response, stress response, antioxidant defenses, cell growth and differentiation and apoptosis (reviewed in ref. 97). NFκB is a dimeric transcription factor and is composed of proteins containing the Rel Homology domain. The Rel family is comprised of five members, p50 and its precursor p105 (also known as NFκB1), p52 and its precursor p100 (also known as NFκB2), p65 (also known as RelA), RelB and c-Rel. The best studied dimer, and the one frequently referred to as ‘NFκB’ occurs between p65 (RelA) and p50. Under normoxic conditions NFκB is retained in the cytoplasm through association with the inhibitor IκB, but in response to anoxia, it undergoes nuclear translocation, mediating the expression of target genes (Fig. 2). This control mechanism allows NFκB to respond rapidly to stress. In mammals, NFκB is an important mediator of transcriptional responses during ischemia/reperfusion (reviewed in ref. 98). In turtles, NFκB is activated in the turtle liver in response to anoxia. Some data on turtle liver NFκB are shown in Figure 3; protein levels of the two NFκB subunits increase in liver nuclei in response to anoxia (Fig. 3A), and the DNA-binding activity of NFκB also increases (Fig. 3B). NFκB was also shown to be anoxia-responsive in the turtle brain.99 These data suggest that NFκB has key functions in protecting these organs under anoxic conditions.

While many of the genes under NFκB control could play beneficial roles in anoxia tolerance, the vast array of genes that lie under NFκB control, as well as the lack of tools available to accurately analyze large-scale transcriptional responses in the turtle, makes it rather difficult to pinpoint all the genes that are activated by NFκB in the turtle in response to anoxia. Some potential candidates include the antioxidant genes, ferritin heavy chain and SOD, as well as the pro-survival genes Bcl-2 and Bcl-xL, which are indeed upregulated in response to anoxia in turtle liver.52

**Anti-apoptotic proteins.** Studies in mammals have shown that apoptosis is oxygen-responsive, with exposure to hypoxia promoting cell-survival and exposure to anoxia promoting apoptosis.100,101 However, tissues of *T. s. elegans* can survive anoxia for many weeks, without any apparent damage, suggesting that inhibition of apoptosis must occur on a molecular level during anaerobiosis. The promotion or inhibition of apoptosis is controlled by two conserved pathways, the death receptor pathway and the...
mitochondrial pathway (shown in Fig. 4), and the decision to initiate apoptosis often depends on the levels of pro- and anti-apoptotic proteins. The transcript levels of two major anti-apoptotic proteins, Bcl-2 and Bcl-xL, increase, possibly in an NFκB-dependent manner, in response to anoxia in turtle liver (Fig. 5). These proteins reside in the outer mitochondrial membrane and inhibit apoptosis by preventing the loss of outer mitochondrial membrane integrity, which leads to lethal downstream events such as activation of caspases. Studies in mammals have shown that enhanced expression of these proteins limits apoptosis in cells and plays a protective role in mammalian ischemic preconditioning.

Like many of the proteins discussed in this review, these proteins potentially have dual roles, protecting against oxidative damage as well as anoxia. Bcl-2 has been previously reported to have antioxidant effects. Transfection of cells with Bcl-2 suppressed lipid peroxidation and Bcl-2 also reduced intracellular ROS levels. In addition, studies have shown that hyperglycemia-induced lipid peroxidation and advanced glycation end products (AGE) in endothelial cells are suppressed by Bcl-2.

Brain-Specific Adaptations

The mammalian brain is especially prone to oxygen limitation and even short bouts of hypoxia can cause irreversible damage. The turtle brain, however, can survive many weeks of anoxia unharmed. Many of the mechanisms involved in brain anoxia-tolerance are also linked to age-related neurodegeneration, and as such may contribute to turtle longevity.

Channels and neurotransmitters. A robust reduction in ion channels occurs in response to anoxia in turtle brain. This includes reduction in voltage-gated Na+ channel density, decrease in mRNA of voltage gated K+ channel, decrease in K+ leak channel and silencing of the Ca2+-dependent NMDA receptor. Because damage to ion channels is involved in several age-related diseases in the brain, it has been suggested that the turtle’s ability to both maintain channel integrity while downregulating activity in response to anoxia, as well as restore those channels upon reoxygenation, could be linked to longevity.

A major characteristic of the response to oxygen limitation by mammalian brain is the uncontrolled and destructive release of excitatory neurotransmitters. The turtle brain, however, avoids such toxic release by maintaining a balance between dopamine and glutamate release and active uptake mechanisms. An increase in the inhibitory neurotransmitter GABA and GABA receptors in the brain is also observed, and likely contributes to strengthening of the inhibitory tone. Adenosine is also involved in the suppression of ion channels in response to anoxia and also plays a role in anoxia-mediated ROS suppression. An increased susceptibility to excitatory amino acids has been previously associated with age-related neurodegenerative diseases, and the turtle’s ability to avoid this could be linked to its longevity.

Figure 4. A generalized schematic for the mitochondrial apoptosis pathway. During apoptosis, the mitochondrial membrane swells and cytochrome c leaks out. This leads to binding with the pro-apoptotic protein Apaf-1, subsequent formation of the apoptosome and activation of caspase-9. Caspase-9 subsequently activates other caspases and results in cell death. Pro-survival Bcl-2 family members (such as Bcl-2 and Bcl-xL) can inhibit apoptosis by binding to the mitochondrial membrane and blocking its swelling.

Figure 5. Effect of 5 and 20 h of anoxic submergence on transcript levels of pro-survival target genes of NFκB in T. s. elegans liver. Representative RT-PCR bands and a histogram showing normalized transcript levels under control and anoxic conditions are shown. Data were normalized against α-tubulin and are shown as means ± S.E.M., n = 3 independent trials. (a) Significantly different from the corresponding control (p < 0.05). Data are taken from ref. 52.
Inosine, the deamination product of adenosine, could also have a role to play in anoxia tolerance. Recent publications suggest a potentially important role for inosine in the activation of antioxidant defenses in response to hypoxia, hyperoxia or $\text{H}_2\text{O}_2$ insult.\textsuperscript{122-124} Interestingly, the enzyme that makes inosine, 5’ nucleotidase cytosolic II isozyme (NT5C2), is upregulated in turtle brain in response to anoxia.\textsuperscript{3} As previously discussed, enhancement of antioxidant defenses is an important part of anoxic survival and an increase in inosine could be a potential trigger for this response.

Neuroglobin. Neuroglobin (Ngb) is another protein that has a potentially important protective role in turtle anoxia tolerance. Ngb is preferentially expressed in the nervous system, as well as some endocrine tissues\textsuperscript{125,126} and has been identified in the brains of rodents, fish, amphibians, birds and reptiles.\textsuperscript{127,128} Since globins generally function as either O$_2$ storage or transport proteins, Ngb is suspected of playing a role in transferring oxygen to the mitochondria in the brain and retina.\textsuperscript{127} Other possible roles for Ngb are in the decomposition of reactive oxygen and nitrogen species, acting as a terminal oxidase that regenerates NAD$^+$ under anaerobic conditions or as a detector of cellular oxygen concentrations.\textsuperscript{129-132} Recent studies have also shown that Ngb expression is reduced with increasing age\textsuperscript{133} and also plays a role in the protection against neurotoxicity.\textsuperscript{134} In mammals, Ngb also plays an anti-apoptotic role by intervening in the activation of pro-caspase 9 through interaction with cytochrome $c$ in the intrinsic mitochondrial pathway.\textsuperscript{135} A role for neuroglobin in the protection against oxidative damage in turtle brain has also been suggested and knock-out of Ngb in neuronally enriched turtle cell cultures resulted in significant increases in $\text{H}_2\text{O}_2$ release.\textsuperscript{136}

The levels of Nrb mRNA increased significantly in response to hypoxia, as well as post-anoxic reoxygenation, in $\text{T.s. elegans}$ brain with a lesser degree of upregulation in response to 1 and 4 h of anoxia.\textsuperscript{128} These results suggest that Ngb may play an important role in turtle adaptation to hypoxia, possibly by ensuring adequate supply of O$_2$ to the brain. The upregulation of Ngb during reoxygenation could also suggest a role in antioxidant defense of the brain.

Interestingly, whereas a strong increase in Ngb levels is not generally observed in response to hypoxia in rodents,\textsuperscript{132} another species that lives in environments of changing oxygen availability, the zebrafish (Danio rerio), shows strong upregulation of Ngb in response to hypoxia,\textsuperscript{137} as in turtles. These findings suggest that inducibility of this protein in response to hypoxia may be an important adaptation to low oxygen environments. Other animals that live in hypoxic environments or deal with oxygen variability show constitutively high levels of neuroglobin. This includes the subterranean blind mole rat Spalax ehrenbergi and the goldfish.\textsuperscript{132}

**The Role of Hypometabolism**

A key element to turtle anoxia tolerance is a profound metabolic rate depression to 10–20% of the corresponding aerobic resting rate at the same temperature. This suppresses, rebalances and reprioritizes the rates of ATP-utilizing and ATP-generating processes so that turtles can sustain long term viability without oxygen. Entrance into hypometabolism is not just an anoxia-specific phenomenon, but is a common survival strategy across phylogeny and is observed in phenomena including hibernation, torpor, aestivation, anaerobiosis, diapause and anhydrobiosis. Understanding the mechanisms of hypometabolism could be of potential use to medical sciences. For example, decreasing the metabolic rate of human organs meant for transplantation could lengthen the time organ explants remain viable. The topic of hypometabolism in turtles and other systems has been thoroughly reviewed elsewhere.\textsuperscript{138,139} Therefore, in the present review we will only touch on some of the key strategies associated with metabolic rate depression in the anoxic turtle.

Entrance into hypometabolism in response to anoxia in turtles is characterized by a reorganization of energy-consuming processes. For example, studies with turtle hepatocytes showed a 94% decrease in overall ATP turnover under anoxia, but found that five main ATP-consuming processes were suppressed to different degrees.\textsuperscript{140} Under anoxia, the Na$^+$K$^+$-ATPase became the dominant energy consumer of the cell, consuming 62% of total ATP turnover as compared with 28% in normoxia. Meanwhile, protein synthesis and degradation were strongly suppressed by >90%, whereas gluconeogenesis and urea synthesis were virtually halted. Similar reductions in protein synthesis were also observed in other organs.\textsuperscript{141} This reorganization of metabolic processes allows turtle cells to direct energy into the most vital processes for cellular survival under anoxia and away from less vital processes, and results in the most efficient ATP utilization under this energy-limited condition.

Reversible protein phosphorylation is considered to be the most widespread and powerful mechanism utilized by cells to make stable changes to enzyme activities in response to various stimuli. This mechanism also plays an important role in the regulation of metabolic depression in other hypometabolic systems.\textsuperscript{138,139} Reversible protein phosphorylation regulates the activities and kinetic properties of many enzymes involved in diverse cellular processes. These include membrane receptors and transporters, ion channels and ATPases, proteins involved in gene transcription, protein synthesis and degradation, cell cycle control and signal transduction cascades. For example, it was shown that reversible phosphorylation is involved in the regulation of voltage-gated ion channels (Na$^+$, Ca$^{2+}$, K$^+$) and membrane receptors (e.g., $N$-methyl-D-aspartate-type glutamate receptor) in anoxia-tolerant turtles.\textsuperscript{142-144} In addition, changes in the phosphorylation status of various ribosomal translation factors were observed in turtle tissues in response to anoxia,\textsuperscript{145} suggesting that such changes are involved in anoxia-induced suppression of protein synthesis in turtles.

It is estimated that 1–10% of cellular energy is devoted to gene transcription, depending on the tissue;\textsuperscript{146} this is also a significant metabolic cost that must be reduced in situations of hypometabolism. Suppressed rates of gene transcription have been observed in a number of hypometabolic systems in response to anoxia including brine shrimp (Artemia franciscana) and intertidal snails (Littorina littorea),\textsuperscript{147,148} as well as in mammalian
A prominent mechanism of global or localized transcriptional control is the post-translational modification of histones that are subject to a number of modifications that affect the state of transcriptional activity in cells.\(^{151,152}\) For example, histone acetylation is associated with higher transcriptional activity because this modification makes chromatin more accessible to proteins of the transcriptional apparatus.\(^{153}\) Histone deacetylases (HDACs) are a group of enzymes that catalyze the removal of acetyl groups from histones, and as such are associated with transcriptional repression. Increased HDAC expression and activity, as well as decreased histone acetylation was observed in skeletal muscle of hibernating ground squirrels.\(^{154}\) Such changes also occurred in turtle muscle in response to anoxia, with HDAC protein levels increasing by 1.7–3.5 fold, and HDAC activity increasing by 1.5-fold in response to anoxia.\(^{155}\) This suggests that histone deacetylation is an important general mode of global transcriptional suppression in hypometabolic systems.

**Table 1. Major proteins/pathways discussed in this review and their roles**

| Pathway/proteins discussed | Role |
|---------------------------|------|
| **Antioxidant enzymes/proteins** | Destroy intracellular ROS and provide relief against oxidative stress (reviewed in refs. 14, 16, 181). |
| **Heat shock response** | Enhance intracellular protein folding capacity (reviewed in refs. 74, 75). |
| **NFκB** | Enhanced expression of genes involved in the immune response, stress response, antioxidant defenses, cell growth and differentiation, and apoptosis (reviewed in ref. 97). |
| **Anti-apoptotic proteins** | Inhibition of apoptosis (reviewed in ref. 102). |
| **Neuroglobin** | Enhanced oxygen delivery to the brain; Possible antioxidant effects (reviewed in ref. 127, 179). |

**Other Models for Longevity**

Other models for longevity also show remarkable stress-resistance, and from a comparative perspective, it would be interesting to investigate whether their stress-resistance mechanisms are similar to those observed in turtles. Among mammals, bats and naked mole rats are particularly known for extreme longevity. Bats are the longest-lived mammals relative to their body size.\(^{156,157}\) Their longevity often surpasses 30 years, and the longest-lived bat species on record survived for at least 41 years in the wild.\(^{158}\) Naked mole rats have a recorded lifespan that exceeds 28 years, approximately nine times greater than predicted for their body size. They display no age-related changes to body mass, basal metabolic rate, percent body fat, fat-free mass or bone density,\(^{159}\) an indication of “negligible senescence.” Both animals show remarkable tolerance to stresses. For example, primary fibroblasts isolated from the little brown bat were reported to be much more resistant to \(\text{H}_2\text{O}_2\), cadmium, UV light and heat, as compared to cells from mice.\(^{160}\) Cultured cells from this species also showed enhanced repair of \(\gamma\)-irradiation-induced DNA lesions.\(^{158}\) Hippocampal slices of mole rats are also significantly more tolerant of oxygen and nutrient deprivation than laboratory white rats\(^{161}\) and mole rat neurons maintain synaptic transmission much longer than mouse neurons and can recover from periods of anoxia exceeding 30 min.\(^{162}\) Furthermore, cultured cells from naked mole rats, including primary fibroblasts and endothelial cells, tend to be resistant to several forms of stress.\(^{163,164}\) Interestingly, no tumors have ever been found in a mole rat and they might possess extraordinary cancer-resistance mechanisms.\(^{165}\)

Bats have lower \(\text{H}_2\text{O}_2\) production\(^{166,167}\) and protein oxidation\(^{168}\) compared to other mammals, which might directly contribute to their longevity. Bats also show resistance to protein oxidation both in vivo and in vitro under conditions of acute oxidative stress.\(^{168}\) In addition, bats showed lower levels of protein ubiquitination, along with reduced proteasome activity and increased resistance to urea-induced protein unfolding compared to mice.\(^{168}\) Naked mole rats also display no increase in cysteine oxidation and ubiquitination over two decades of life.\(^{169}\) These results suggest that the longevity in these animals may be regulated by very efficient maintenance of protein homeostasis. A potential mechanism for the long-term maintenance of protein homeostasis could be via the increased expression of molecular chaperones and it would be interesting to investigate whether, as in turtles, this occurs in bats and naked mole rats. Interestingly, several species of bats are also known to hibernate, a condition characterized by strongly reduced metabolic rate, decreased body temperature and hypoxia/ischemia, as well as increased activation of various stress-tolerance mechanisms (reviewed in ref. 170). It is possible that some of the stress-tolerance mechanisms that support bat cells during long term torpor might also contribute to their enhanced longevity.

**Conclusions**

Many different molecular mechanisms appear to play protective roles in anoxia-tolerance (see Table 1 for a summary of the major pathways discussed) and a variety of these might have beneficial anti-aging properties. For example, the maintenance of constitutive protective mechanisms, while energy-expensive, would provide immediate protection against any assault on metabolic integrity as well as minimize damage accumulated throughout an animal’s lifetime. On the other hand, while the inducibility of key protective mechanisms is known to decrease with age in normal-lived animals, the turtle’s exposure to oxygen variability throughout its lifetime might put an emphasis on maintaining the inducibility of these mechanisms, making it less prone to aging-associated damage. Understanding the molecular adaptations associated with anoxia tolerance is of key importance not only from a comparative point of view, but also from a medical one, as many of these mechanisms are involved in human disease. For example, induction of a hypometabolic state in human organs could potentially prolong organ survival for transplantation,\(^{171}\) as well as prolong organ survival following trauma. Indeed, some research demonstrates that induced hypometabolism reduces the detrimental effects...
effects of hypoxic/ischemic organ injury. The natural mechanisms of anoxia tolerance are also of interest because the molecular nature of ischemic injury in mammals is not yet completely understood and the study of these responses in anoxia-tolerant animals could highlight those mechanisms that are beneficial for survival. For example, the induction of the transcription factor NFKB in turtle organs in response to anoxia suggests a beneficial role during oxygen deprivation. This transcription factor appears to play a dual role in mammalian ischemia/reperfusion injury, promoting inflammation and death, but also playing a crucial role in ischemic preconditioning. Hence, a better understanding of the role of this transcription factor and its target genes could lead to better treatment for a variety of pathogenic conditions involving ischemia/reperfusion injury, including heart disease and stroke. Similarly, other molecular mechanisms that appear to play roles in turtle hypoxia/anoxia tolerance are relevant to human disease. Neuroglobin overexpression confers neuroprotection against neuronal hypoxia/anoxia-induced damage, whereas enhanced expression of protective proteins such as heat shock proteins, antioxidant proteins and anti-apoptotic proteins has been tied with the enhanced organ protection observed during ischemic preconditioning. Despite the potential key importance of natural anoxia tolerance to medical sciences, there is still much to be learned about the mechanisms underlying this ability. It is expected that as we learn more about these processes, we will be able to better understand the molecular nature of many oxygen-associated pathological conditions in mammals and identify new treatments for these conditions, effectively improving human health and extending lifespan.

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