Review

Oxidative stress and anxiety

Relationship and cellular pathways

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High O₂ consumption, modest antioxidant defenses and a lipid-rich constitution make the brain highly vulnerable to redox imbalances. Oxidative damage in the brain causes nervous system impairment. Recently, oxidative stress has also been implicated in depression, anxiety disorders and high anxiety levels. The findings which establish a link between oxidative stress and pathological anxiety have inspired a number of other recent studies focusing on the link between oxidative status and normal anxiety and also on a possible causal relationship between cellular oxidative stress and emotional stress. This review examines the recent discoveries made on the link between oxidative status and normal anxiety levels and the putative role of oxidative stress in genesis of anxiety. We discuss the different opinions and questions that exist in the field and review the methodological approaches that are being used to determine a causal relationship between oxidative and emotional stress.

Introduction

Anxiety is an aversive emotional state, in which the feeling of fear is disproportionate to the nature of the threat. In response to threatening situations, the feeling of the emotion that constitutes the subjective feature of anxiety is accompanied by emotional stress, which involves behavioral, expressive and physiological features, such as an avoidance of the source of the danger, assuming defensive postures and an increase in blood pressure, respectively. Anxiety is a normal emotional response to a threat or potential threat. When this emotion is inappropriate, extreme and persistent, it is classified as pathological. Anxiety is implicated in a number of psychiatric disorders, such as depression, panic attacks, phobias, generalized anxiety disorder, obsessive-compulsive disorder and post-traumatic stress disorder. Anxiety disorders affect approximately 28.8% of the US population, imposing both an individual and a social burden that amounts to a total cost of $42.3 billion in the US in 1990. Anxiety disorders are the most common class of psychiatric disorders in the US and many other countries. According to an ESEMeD study including six European countries, the 12-month prevalence of inappropriate anxiety was 6.4%. In the most recent systematic review of studies conducted in 16 European countries, however, this value was estimated to be 12%. In an obese population in the UK, 56% of patients met the minimum criteria for an anxiety disorder. It is estimated that one-eighth of the total population worldwide suffers from inappropriate anxiety. Population-based studies have shown that anxiety disorders frequently go untreated. Predominantly, the research that has been performed on anxiety has focused on the regulatory systems, including gamma-aminobutyric acidergic (GABAergic) and serotoninergic systems among others. However, Kuloglu et al. recently established a link between oxidative stress and certain anxiety disorders (obsessive-compulsive disorder and panic disorder), demonstrating that other systems, such as oxidative metabolism, can affect the regulation of anxiety. These findings, which establish a link between oxidative stress and pathological anxiety, inspired a number of other recent studies focusing on the link between oxidative status and normal anxiety (Table 1) and also on a possible causal relationship between cellular oxidative stress and emotional stress.

It is well known that low/moderate concentrations of reactive oxygen species (ROS) affect a great number of physiological functions. However, when ROS concentration exceeds the antioxidative capacity of an organism, animal cells enter a state termed oxidative stress, in which the excess ROS induces oxidative damage to nucleic acids, proteins and neuronal membrane...
A link between oxidative stress and emotional stress is not surprising, since it is well accepted that oxidative damage in the brain causes an impairment of the nervous system. In living organisms, an imbalance between oxidant production and antioxidant protection that favors oxidants causes a state termed oxidative stress. In this state, there can be differences in gene expression, protein conformation and cellular signaling. This state may also alter neurotransmission, neuronal function and overall brain activity, as well as disrupting membrane integrity; even neuronal cell death may result. Oxidative stress can alter neurotransmission, neuronal function and overall brain activity. In this state, there can be differences in gene expression, protein conformation and cellular signaling. This state may also alter neurotransmission, neuronal function and overall brain activity, as well as disrupting membrane integrity; even neuronal cell death may result. 

Table 1  **Data establishing the link between pathological/normal anxiety and oxidative cell pathways and mechanisms**

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Expression of antioxidant genes</th>
<th>Activity of antioxidant proteins</th>
<th>Lipid peroxidation markers</th>
<th>Intracellular ROS levels assessed by using the 2',7'-dichlorofluorescin diacetate (DCFH-DA) sensor</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pathological anxiety</td>
<td>Patients with obsessive-compulsive disorder and panic disorder</td>
<td>-</td>
<td>In erythrocytes: Superoxide dismutase*, catalase, glutathione peroxidase*</td>
<td>In plasma: Malondialdehyde*</td>
</tr>
<tr>
<td>Normal trait-anxiety</td>
<td>Mouse strains with high- Vs. low-anxiety-related phenotypes (Hovatta et al.25)</td>
<td>Glyoxalase 1* and glutathione reductase 1*</td>
<td>Glyoxalase 1* and glutathione reductase 1*</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>Anxious Vs. non-anxious Swiss albino mice (Rammal et al.41,42)</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

*significantly different from controls (healthy humans, strains with low-anxiety-related phenotypes, non anxious mice).

In this review, we discuss the relationship between oxidative stress and normal anxiety by presenting the recent advances in the field and the different views that exist. We also review the methodological approach for determining the causal relationship between oxidative stress and emotional stress.

**A Link Between Oxidative Stress Metabolic Pathways and Anxiety-Related Phenotypes**

In 2005, Hovatta et al.25 identified a close relationship between antioxidative defense mechanisms and anxiety-related phenotypes in six inbred mouse strains. They found that, in the brain, the expression of glutathione reductase 1 and glyoxalase 1, which are genes involved in antioxidative metabolism, is highly correlated with anxiety-related phenotypes. Furthermore, they also found that the activity of these enzymes is highest in the most anxious mice and lowest in the least anxious strains. These authors were the first to demonstrate a link between oxidative stress metabolic pathways and normal anxiety.

Nevertheless, Hovatta et al.25 obtained surprising results in a genetic manipulation using lentivirus-mediated gene transfer: local overexpression of glutathione reductase 1 and glyoxalase 1 in the cingulated cortex of the murine brain results in an increase of anxiety-like behavior, while inhibition of glyoxalase 1 expression produces low-anxiety mice. Thus, Hovatta et al.25 were able to make a causal link between the antioxidative status of the brain and anxiety-related behavior and hypothesize that glyoxalase 1 and glutathione reductase 1 regulate anxiety in mice. It is worth mentioning that in vivo, antioxidant genes are overexpressed in response to an uncontrolled production of ROS. Indeed, in vivo, excessive ROS accumulation induces the overexpression of the glutathione redox system, including glutathione reductase, and a general overexpression of endogenous antioxidants.31,32 In the...
trait anxiety. These results are discordant with those of Hovatta et al. However, and complicate the understanding of the relationship between oxidative stress and trait anxiety. Krömer et al. and Ditzen et al. examined the expression of glyoxalase 1 in several areas of the brain areas and in red blood cells and found that this protein is expressed more in a line with a low-anxiety-related behavioral phenotype than in a line with a high-anxiety-related behavioral phenotype. It is worth noting that Hovatta et al. suggested that there is a link between glyoxalase 1 and oxidative stress but that this link is indirect. Indeed, glyoxalase 1 is an enzyme of the glyoxalase system, which protects against carbonyl stress; glutathione is a determinant cofactor for the enzymatic reaction that is catalyzed by glyoxalase 1.

A Link Between Oxidative Stress Metabolic Pathways and Anxiety-Related Behavior

The disagreements in the data of Hovatta et al., Krömer et al. and Ditzen et al. can be partly explained by the differences in the genotypes of the strains. Thus, it would be interesting to compare in the same strain the oxidative status of mice with contrasting levels of anxiety, rather than compare the oxidative status of strains that differ in their anxiety-related phenotypes. We have observed that naive Swiss albino male mice have a large heterogeneity in their anxiety levels. In first order regression analyses of the performances of mice in the light/dark choice test, which are used as a behavioral indices of anxiety, and the intracellular ROS accumulation in blood granulocytes, we found correlation coefficients (R2) ranging from 0.61 (p < 0.01) to 0.73 (p < 0.001) (Fig. 1). Our results suggest that there is a positive correlation between peripheral blood cell oxidative status and anxiety-related behavior in the light/dark box in Swiss albino male mice.

Levels of ROS can be evaluated directly with sensors, such as 2',7'-dichlorofluorescin diacetate (DCFH-DA), or indirectly by measuring the levels of certain antioxidant enzymes, byproducts of lipid peroxidation or some transition metals, such as copper, zinc and iron. In our study, the intracellular redox status of the cells was evaluated using the tracer DCFH-DA, a well known sensor of ROS. To confirm the relationship between oxidative stress and emotional stress, we comparatively evaluated the peripheral oxidative status of mice with contrasting levels of anxiety (anxious and non-anxious). Mice with intermediate behaviors were eliminated. We found that high anxiety in mice in the light-dark box is accompanied by significantly high levels of intracellular ROS in lymphocytes, granulocytes and monocytes (Table 2). Our results confirm that there is the relationship between the level of intracellular ROS in peripheral blood cells and anxiety-related behavior in mice.

These results prompted us to study the oxidative status of the brain in mice with contrasting levels of anxiety. Using the same behavioral approach to distinguish between anxious and
studies are in good concordance with the initial findings of S,R-sulfoximine (BSO), an inducer of oxidative stress, induces behavior in mice. They found that treating mice with buthionine-S,R-sulfoximine, an inducer of oxidative stress, induces anxiogenic behavior in mice and established that oxidative stress leads to anxiogenic behavior. They examined the direct effect of oxidative stress on anxiety-like standing of the relationship between oxidative stress and anxiety. Hovatta et al. and show a clear link between oxidative stress and anxiety-related behavior. Recent data from Desrmaaux et al., Souza et al. and Berry et al. provide indirect evidence for the causal link between oxidative stress and anxiety-related behavior. Desrmaaux et al. showed that vitamin E deficiency in the mouse brain significantly increases the levels of central oxidative stress markers, resulting in anxiogenic behavior without abnormalities in the locomotor performance of the mice. Souza et al. demonstrated in rats that the consumption of a highly palatable diet enriched with sucrose leads to an obese phenotype, increases protein oxidation in the frontal cortex and induces anxiety-like behavior in the dark/light choice test without altering locomotion in an open field test. Berry et al. showed that mice developed anxious behavior during aging, likely due to the accumulation of oxidative damage, which is a characteristic of the aging process in animals. In addition, Berry et al. showed that a deletion of the 66Shc longevity gene in mice, which results in lower levels of oxidative stress and an extended life span, decreases anxiety-related behavior. Overall, the data presented in these reports suggest that oxidative stress can provoke anxious behavior in rodents.

### Table 2 Intracellular ROS accumulation in peripheral immune cells of anxious and non-anxious mice

<table>
<thead>
<tr>
<th>Type of cells</th>
<th>Mean fluorescent intensity (arbitrary units)</th>
<th>Anxious mice</th>
<th>Non-anxious mice</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lymphocytes</td>
<td>215 ± 31*</td>
<td>135 ± 12</td>
<td></td>
</tr>
<tr>
<td>Monocytes</td>
<td>46 ± 8*</td>
<td>25 ± 4</td>
<td></td>
</tr>
<tr>
<td>Granulocytes</td>
<td>1635 ± 375**</td>
<td>418 ± 87</td>
<td></td>
</tr>
</tbody>
</table>

The mean fluorescent intensity corresponds to fluorescence resulting from intracellular DCFH-DA oxidation by intracellular ROS. Data are expressed as mean ± SEM. (n = 10). *p < 0.05; **p < 0.01.

### Table 3 Intracellular ROS accumulation in cerebral and peripheral cells of anxious and non-anxious mice

<table>
<thead>
<tr>
<th>Type of cells</th>
<th>Mean fluorescent intensity (arbitrary units)</th>
<th>Anxious mice</th>
<th>Non-anxious mice</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neurons of cerebral cortex</td>
<td>1223 ± 232**</td>
<td>177 ± 76</td>
<td></td>
</tr>
<tr>
<td>Neurons of cerebellum</td>
<td>546 ± 128*</td>
<td>227 ± 41</td>
<td></td>
</tr>
<tr>
<td>Neurons of hippocampus</td>
<td>1749 ± 566*</td>
<td>227 ± 51</td>
<td></td>
</tr>
<tr>
<td>Glial cells of cerebral cortex</td>
<td>429 ± 48</td>
<td>310 ± 72</td>
<td></td>
</tr>
<tr>
<td>Glial cells of cerebellum</td>
<td>343 ± 36*</td>
<td>195 ± 31</td>
<td></td>
</tr>
<tr>
<td>Glial cells of hippocampus</td>
<td>468 ± 43*</td>
<td>319 ± 29</td>
<td></td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>251 ± 46*</td>
<td>152 ± 11</td>
<td></td>
</tr>
<tr>
<td>Monocytes</td>
<td>56 ± 9*</td>
<td>35 ± 2.5</td>
<td></td>
</tr>
<tr>
<td>Granulocytes</td>
<td>1979 ± 405**</td>
<td>448 ± 77</td>
<td></td>
</tr>
</tbody>
</table>

The mean fluorescent intensity corresponds to fluorescence resulting from intracellular DCFH-DA oxidation by intracellular ROS. Data are expressed as mean ± SEM. (n = 10). *p < 0.05; **p < 0.01.

In agreement with our recent findings, observed a significant relationship between trait anxiety and ROS formation in monocytes of hypertensive individuals. Recently, Masood et al. published work that contributes to the understanding of the relationship between oxidative stress and anxiety. They examined the direct effect of oxidative stress on anxiety-like behavior and established that oxidative stress leads to anxiogenic behavior in mice. They found that treating mice with buthionine-S,R-sulfoximine (BSO), an inducer of oxidative stress, induces anxious behavior through the NADPH oxidase pathway. The anxiogenic behavior due to BSO treatment was observed in several mouse models of anxiety, including elevated plus maze, hole-board and open field tests. Masood et al. induced oxidative stress in mice by depleting glutathione with BSO inhibition of gamma-glutamylcysteine synthetase. Glutathione depletion causes a myriad of cellular stresses, including oxidative, nitrosative and carbonyl stresses, as glutathione is an important determinant of the oxygen, nitrogen and dicarbonyl metabolisms.

Excessive production of ROS induces oxidative damage of cellular structures, production of reactive nitrogen species triggers nitrosylation reactions, which can alter the structure of proteins to inhibit their normal function. excessive accumulation of reactive dicarbonyl compounds leads to damage of protein and nucleotides by dicarbonyl glycation.

### Indirect Evidence for the Causal Link Between Oxidative Stress and Anxiety

Recent data from Desrmaaux et al., Souza et al. and Berry et al. provide indirect evidence for the causal link between oxidative stress and anxiety-related behavior. Desrmaaux et al. showed that vitamin E deficiency in the mouse brain significantly increases the levels of central oxidative stress markers, resulting in anxiogenic behavior without abnormalities in the locomotor performance of the mice. Souza et al. demonstrated in rats that the consumption of a highly palatable diet enriched with sucrose leads to an obese phenotype, increases protein oxidation in the frontal cortex and induces anxiety-like behavior in the dark/light choice test without altering locomotion in an open field test. Berry et al. showed that mice developed anxious behavior during aging, likely due to the accumulation of oxidative damage, which is a characteristic of the aging process in animals. In addition, Berry et al. showed that a deletion of the 66Shc longevity gene in mice, which results in lower levels of oxidative stress and an extended life span, decreases anxiety-related behavior. Overall, the data presented in these reports suggest that oxidative stress can provoke anxious behavior in rodents.

### Conclusion

This review summarizes the data to support a link between oxidative stress and anxiety. While all of the data demonstrate that there is a link between oxidative stress and high-anxiety-related behavior, a cause-effect relationship has yet to be completely established. Some of these studies suggest that oxidative stress causes anxiety-related behaviors but do not explain the underlying mechanisms. While there are some limits in the approach to establish the anxiogenic effect of oxidative stress, the available data are consistent with this causal relationship. The potential causal role of oxidative stress on anxiety may generate interest in antioxidants. Masood et al. were able to show that oxidative stress-related anxiety can be reversed in mice upon inhibition of NADPH oxidase or phosphodiesterase-2, enzyme that is indirectly implicated in oxidative stress mechanisms. Surprisingly, they found that diazepam, which is a well known anxiolytic, does not fully reverse the oxidative stress-related anxiety. These results point to a possible use for antioxidants.
in the prevention or reduction of high anxiety. Further research will be necessary to show whether anxious subjects need more antiOXidants than non-anxious subjects. Recent work52-53 has shown that some dietary polyphenols have both anxiolytic and antioxidant effects, which may be beneficial to anxious subjects.

References

32. Bouayed J, Rammal H, Younos C, Soulmaini R. Chlorogenic acid, a polyphenol from Prunus domestica (Mirabelle), with coupled anxiolytic and antioxidant effects.