Oxidative Stress and Antioxidant Status in Immune Thrombocytopenia

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Abstract

Immune thrombocytopenia (ITP) is an autoimmune disorder characterized by an isolated thrombocytopenia that may manifest as skin and mucous membrane bleeding. However, the pathogenicity of this disease is much elusive. Increasing evidences demonstrate oxidative stress plays an essiential role in the pathogenesis of autoimmune diseases including ITP, which may provide a novel therapeutic approach.

Keywords: Lupus erythematosus; Reactive oxygen; Pathogenesis; Autoimmune inflammatory disease

Introduction

Immune thrombocytopenia (ITP) is an autoimmune disorder characterized by isolated thrombocytopenia due to pathogenic anti-platelet autoantibodies, T cell-mediated platelet destruction, and impaired megakaryocyte function [1]. This disorder is classified as primary, also referred to as idiopathic thrombocytopenic purpura, or as secondary to an underlying disorder such as chronic infections, including Helicobacter pylori, or other autoimmune diseases including systemic lupus erythematosus (SLE) and antiphospholipid syndrome [2]. Many aspects of immune dysregulation in ITP have widely been investigated, but its precise mechanisms are still unclear, and the exact triggering events remain elusive. Recently, several evidences have suggested that oxidative stress plays an important role in the pathogenesis of autoimmune diseases, and it is also involved in the development of ITP. These findings might provide a new hypothesis for the initiation of abnormal autoimmunity and a possible novel therapeutic approach.

Oxidative stress and antioxidant status

Reactive oxygen species (ROS) could be generated from normal cellular metabolism such as oxidative phosphorylation in mitochondria and long-chain fatty acids oxidation in peroxisomes, and be served as second messengers [3]. The ROS mainly include oxygen molecule (O2), superoxide anion radical (·O2−), hydrogen peroxide (H2O2), hydroxyl radical (·OH), and singlet oxygen (1O2) [4]. On the other hand, in maintaining the equilibrium, there also exist antioxidants and antioxidant enzymes such as superoxide dismutase (SOD), glutathione peroxidase (GPx) and catalase (CAT), which are responsible for regulating the redox state. Mitochondria and lysosomes are considered to be the two important subcellular organelles that mediate oxidative stress. In general, the forms of life maintain a reducing environment within their cells, preserved by several relevant enzymes. In general, at low to moderate concentrations, ROS act as important molecules involved in many physiological processes such as cell activation, viability, proliferation, and organ function, and they also play important roles in inflammatory response and anti-infection, as they can be used by the immune system to attack and kill pathogens [5]. Oxidative stress occurs as a result of increased activity of free radical–producing enzymes, decreased activity of free radical–removing enzymes, and insufficient levels of antioxidants [6]. It may cause irreversible changes such biomolecules as lipids, proteins and carbohydrates. On the other hand, it can also induce a variety of different types of DNA damage or replication stress [7]. It has been demonstrated that oxidative stress in a hypoxia re-oxygenation reaction in endothelial cells may induce profound damage, reflecting chromosomal aberrations and micronuclei [8]. Thus, ROS is the production of normal cells and plays an important role in the metabolism of our body. However, if it is generated excessively or unable to be eliminated, it will be harmful to our health.

Oxidative stress and autoimmunity

Increased oxidative stress has been proved to be related to the development of a variety of diseases such as atherosclerosis [9], neurodegenerative diseases [10] and cancer [3]. Moreover, it also plays a vital role in the pathogenesis of autoimmune diseases. Oxidant stress could induce autoimmune dysregulation, which includes the generation of neo-epitopes and auto-antigens, leading to B- and T-cell dysregulation and the generation of autoantibodies (Figure 1). It has been reported that, in patients with autoimmune cholestatic liver diseases (AC) and autoimmune hepatitis (AIH), the levels of all lipid and protein oxidative injury products are increased, while the whole blood glutathione levels are significantly decreased. Furthermore, protein carbonyl and isoprostane levels are increased and glutathione levels decreased gradually with progression from mild fibrosis to severe fibrosis and cirrhosis in both AC and AIH patients. These findings support a major contribution of antioxidant/antioxidant imbalance in the progression of liver injury in autoimmune liver diseases [5]. SLE is an autoimmune inflammatory disease whose pathogenesis also refers to oxidative stress. Several studies have shown increased production of ROS or diminished levels of intracellular reduced glutathione in various blood components in SLE patients [11,12]. Several investigators suggested that restoration of the redox balance using antioxidant agents or diminishing effect of oxidative stress by intake of antioxidant nutrients, might attenuate the
complication of SLE induced by oxidative stress [11]. Similarly, the deficiencies of SOD and peroxiredoxin inducing oxidative stress were considered to be an essential factor that might participate in the pathogenesis of another autoimmune disease autoimmune hemolytic anemia. The SOD1-deficient mice model showed a positive correlation between autoantibodies and ROS in RBCs. RBCs are under oxidative stress constitutively, and cellular components face the risk of oxidative damage [13]. Endothelial inflammation, mitochondrial oxidative stress would be activated in the condition of high glucose levels. Therefore, type 2 diabetes (T2D) would be served as a prime example of an interplay between metabolism and immunity, making it prototypic for an in depth look into immunometabolism [14-16]. Growing evidence implies that immune cells possess an essential effect on the development of insulin resistance and then complications of T2D [17-19]. Numerous investigators have observed elevated levels of oxidative stress markers in patients with T2D [20,21]. Therefore, the imbalance between oxidant and antioxidant systems is also involved in the pathogenesis of T2D and its complications [22-24]. Recent data demonstrates that oxidative stress influences cell-cycle regulators in β cell proliferation and neogenesis [25]. Furthermore, several lines of evidence suggest that oxidative stress is implicated in the pathogenesis of diabetic neuropathy (DSPN) [26]. Many researchers have observed that oxidative stress is increased in peripheral and dorsal root ganglion nerves and vascular endothelial cells exposed to hyperglycemia [26,27]. And for T2D patients with DSPN, the total antioxidant status was decreased compared with controls [28,29].

\[\text{Environmental factors} \rightarrow \text{ROS production} \rightarrow \text{Oxidative stress} \rightarrow \text{DNA fragmentation} \rightarrow \text{Cell death} \rightarrow \text{Apoptosis} \rightarrow \text{Immunological tolerance break down (anti-antibodies generation)}\]

**Figure 1**: Oxidative stress involved in the pathogenesis of autoimmune disorders.

### Oxidative stress in ITP

ITP is an autoimmune disease due to autoantibodies against one or several platelet surface antigens, but the mechanism is complex. In the past decade, many evidences have confirmed that oxidative stress was involved in the pathogenesis of ITP. Polat et al. [30] studied lipid peroxidation, glutathione, and ascorbic acid levels in adult ITP patients, and they found that lipid peroxidation level was higher, and glutathione and ascorbic acid levels were lower in the patient group than in the control group, which implies that oxidative damage is involved in the mechanism of ITP. Other researchers also confirmed lipid peroxidation, reduced glutathione (GSH) and oxidative stress-related pathways in whole blood and/or plasma samples from ITP patients [31,32]. Recently, a more profound method of microarray analysis has been used as a tool to explore the gene expression profiles of patients with acute and chronic ITP. The expression of the gene vanin-1 was increased in patients with chronic ITP during the acute phase and patients with treatment-resistant ITP [33]. As regard to different phases of ITP, the markers present differently, the antioxidant capacity of ITP patients in the active phase was drastically reduced, while higher Gpx activity was observed in both active phase and remission in comparison to healthy controls [34]. Interestingly, investigators found that female patients with ITP demonstrated significantly higher oxidative stress than male patients, suggesting sex differences in the pathomechanisms of ITP [34]. With regard to antioxidant status and oxidative stress in chronic ITP in adult patients, Jin et al. [35] found that the number of platelets had a negative correlation with NO, oxidized glutathione(GSSG), malondialdehyde (MDA), and total oxidant status (TOS), respectively, while platelet number showed a positive correlation with SOD, CAT, Gpx and GSH. It has been found that markers of antioxidant status were significantly lower in patients with ITP compared to healthy controls. In addition, MDA was significantly higher in patients with newly diagnosed ITP compared to both healthy controls and patients with chronic ITP [31]. All these observations almost approved that oxidative stress may be partially responsible for the maintenance in thrombocytopenia.

### Antioxidant therapy for autoimmune disorders

Oxidative stress, characterized by a persistent imbalance between the production of highly ROS and reactive nitrogen species and antioxidant defence leads to an altered cellular redox status and subsequent tissue damage. A growing body of researches have demonstrated that the etiology and pathogenesis of autoimmune diseases may be closely related to the unbalance between pro-oxidation and antioxidant. Therefore, many experts hold the idea of using antioxidants to eliminate excessive oxidizing agent. Numerous therapies for oxidative stress related autoimmune disorders have been tried, including corticosteroids, biologics, antibiotics, immunosuppressive drugs, 5-aminosalicylates and vitamin D analogs [36]. As anti-oxidants serving to limit relevant damage, alpha lipoic acid (ALA) shows a clear metabolic influence, which could improve microcirculation, and has an anti-inflammatory effect, and it could contribute to the defense against oxidative stress by increasing the synthesis of anti-oxidants like glutathione, one of the most abundant intra-cellular anti-oxidants in the body. Furthermore, there is growing evidence that ALA is beneficial for improving insulin resistance for T2D patients, and for improving microcirculation [37]. However, some other experts considered these therapies are mostly directed towards providing temporary symptomatic relief. Thus many studies have been drifted towards approaches to mitigate this damage by supplementing antioxidant enzymes such as SOD, CAT and glutathione peroxidase [37,38], which could inhibit depolymerization of hyaluron of synovial fluid, inhibiting degradation of neutrophils and proteoglycans in articular cartilage, preventing collagen fragmentation by scavenging reactive oxidants, reducing elastase activity and inhibiting lipid peroxidation [38]. Nevertheless, various reports suggest they have poor pharmacokinetic profile that limits their use. The most common
therapeutic approaches for ITP are corticosteroids, immunosuppressive drugs or splenectomy. Unfortunately, certain adult patients could not remain in remission, and no other regimens are universally effective. In this condition, some physicians focused on the role of oxidant stress in the pathophysiology of ITP and study relevant drugs. There has been reported that the use of ascorbic acid in refractory ITP seems effective and might be an attractive therapeutic option for patients who were resistant to some or all of the usual therapy [39]. Elalfy et al. [40] reported adjuvant antioxidant therapy containing vitamin A, C and E and selenium yeast could significantly decrease MDA in children and adolescents with ITP, which might be considered as a complementary therapy with detectable oxidative stress to improve bleeding score as well as platelet count. However, there existed some controversial results. For instance, Steven and Bierling concluded eventually that ascorbic acid is not very effective in patients with refractory ITP [41,42]. Meanwhile, for children and adolescents who were diagnosed with ITP, Akbayram et al. [43] did not find significant differences in the oxidative stress and antioxidant capacity following steroid treatment. While Cura et al. [44] demonstrated an increase in the mean total antioxidant status level, and a decrease in the mean total oxidant status and oxidative stress index levels following one week of high-dose methylprednisolone treatment in children ITP. In addition, it has been proved an inhibitory effect of dexamethasone and prednisolone on the generation of reactive oxygen species from platelets, and this effect of both drugs was dose-dependent.

Conclusion

Oxidative modification of proteins and defect in T cell signal processing and/or antigen presentation could elicit antibodies in a variety of autoimmune diseases including ITP, in which antioxidant therapy has been demonstrated to ameliorate the oxidative stress. Standard therapy of ITP comprises corticosteroids, immunosuppressive agent, intravenous immunoglobulin, and splenectomy. However, some patients fail to respond to standard treatment or relapse. Recently, antioxidant therapeutic approach has shown to raise platelet count, indicating the value of adjunctive therapy. Further large scale prospective study will confirm its effectiveness in treating these patients with ITP.

Conflicts of Interest

We declare that we have no conflicts of interest.

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