Abstract

Crohn’s disease is a chronic inflammatory intestinal disease associated with increased of pro-inflammatory cytokines, iron deficiency and anaemia. This case report demonstrates a strategy for intravenous iron supplementation in a patient with Crohn’s disease and severe anaemia. This study found an increased in iron and haemoglobin concentrations, malondialdehyde and superoxide dismutase levels and a reduction in glutathione peroxidase (GPx), vitamin A and E. In conclusion the use of intravenous iron probably leads to increase ROS production; this effect may be reduced by using polivitaminics, and enhanced the free radical scavenger.

Keywords: Crohn’s disease; Anaemia; Iron supplementation; Iron deficiency; Oxidative stress

Introduction

Crohn’s disease (CD) is an idiopathic chronic inflammatory intestinal disease (IBD) [1] that affects gastrointestinal tract [2]. This disease is charaterized by mucosal white blood cells infiltration, proinflammatory cytokines and reactive oxygen species (ROS) production. In this context, oxidative stress control is important factor in the IBD progression and recovering [1]. Moreover, severe anaemia is a common complication of IBD, affecting 6% to 74% of the patients [3], having great impact on the mortality [4,5]. The etiology of anaemia in IBD is multifactorial [6], however, iron deficiency (ID) is the primary cause [7,8].

Case Report

In this case report, we describe a Caucasian, 18 years-old men with Crohn’s disease who is treated in an Academic Public Hospital. He was a non-smoker, non-alcoholic and denies illicit drug abuse. His family history included a father with obesity, hypertension, smoke habit and a healthy mother. His past medical history includes onset of diarrhoea for ten years, associated with abdominal pain, hypertension, smoke habit and a healthy mother. His past medical history includes onset of diarrhoea for ten years, associated with abdominal pain, hypertension, smoke habit and a healthy mother. His past medical history includes onset of diarrhoea for ten years, associated with abdominal pain, hypertension, smoke habit and a healthy mother. His past medical history includes onset of diarrhoea for ten years, associated with abdominal pain, hypertension, smoke habit and a healthy mother. His past medical history includes onset of diarrhoea for ten years, associated with abdominal pain, hypertension, smoke habit and a healthy mother.

Throughout the internment bowel habit remained 3 times a day. Blood samples were collected on alternate’s day, after 12 hours of fasting, for biochemical analysis. Haemoglobin and plasma iron values increased (Figure 1). Also, there was an increased in malondialdehyde (MDA) and superoxide dismutase (SOD) levels and a reduction in glutathione peroxidase (GPx), vitamin A and E (Table 1). The weight remained the same (29.8 kg).

<table>
<thead>
<tr>
<th></th>
<th>Day 1 (Baseline)</th>
<th>Day 3</th>
<th>Day 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDA (µ mol/l)</td>
<td>1.18</td>
<td>1.00</td>
<td>1.02</td>
</tr>
<tr>
<td>SOD (U)</td>
<td>1.05</td>
<td>0.90</td>
<td>1.21</td>
</tr>
<tr>
<td>GPx (µ mol/min/Hb)</td>
<td>20.45</td>
<td>19.29</td>
<td>16.97</td>
</tr>
<tr>
<td>Vit A (µ mol)</td>
<td>0.16</td>
<td>0.08</td>
<td>0.02</td>
</tr>
<tr>
<td>Vit E (µ mol)</td>
<td>4.31</td>
<td>3.04</td>
<td>3.81</td>
</tr>
</tbody>
</table>

Figure 1: Serum iron, Haematocrit and haemoglobin concentration before (Day 1-Baseline) and after the iron supplementation.
patients with a high degree of species may play an important role in CD. years presence of IBD, activated T lymphocytes reduce oral iron absorption. In this context, iron supplementation should be their intestinal mucosa, submucosa and muscularisserosa. 2+ GPx and transient ions such as Fe and Cu 2+, in small non-protein shielded chelates, can catalyze the production of ROS, which in turn can lead to peroxidation and radical chain reactions with molecular damage. There is direct and indirect evidence suggesting that the chronically inflamed intestines of Crohn's patients may be subjected to considerable oxidative stress. Although certain amounts of iron and reactive oxygen species may play an important role in CD.

Moreover, the balance of ROS concentrations is supported by antioxidant defense mechanisms include vitamins, the enzymes SOD, GPx and transient ions such as Fe 2+ and Cu 2+ [22]. However, Crohn's patients have low blood levels of vitamin C and E [23]. They also have low levels of Cu, Zn, SOD, GPx, catalase, vitamin A and β-carotene in their intestinal mucosa, submucosa and muscularisserosa [24]. Corroborating with this, data found in the present study showed an increased level of MDA and low levels of vitamin E, SOD and GPx at the initial of treatment and remained after iron supplementation. This coincidence of increased levels of ROS and decreased levels of antioxidant defenses results in a state of oxidative stress in Crohn's strictures [25].

In the present study, the patient did a treatment with a sulfasalazine (SAZ) that is a prodrug composed by a molecule of 5-aminosalicylic acid (5-ASA), linked by an azo bond, which has been shown to be effective in the therapy of IBD such as ulcerative colitis and CD. The point is well known that the oral administration of SAZ is capable of reducing the mucosal injury and inflammation associated with Crohn's disease, where 5-ASA is the pharmacologically active moiety of SAZ [25]. It has been suggested that 5-ASA protects the gut mucosa by inhibiting cyclooxygenase and lipoxygenase activities [25]. Nielsen et al. [26] demonstrated a high rate of lipid peroxidation in the gut mucosa obtained from patients with IBD. They also reported a significant inhibition in lipid peroxidation among patients who had been treated with SAZ for 5 weeks [27]. In our study the use of sulfasalazine for 10 days did not change the antioxidants and MDA levels, maybe because the iron treatment had a potential prooxidant effect.

In conclusion the use of intravenous iron probably leads to increase ROS production; this effect may be reduced by using polivitamins, and enhanced the free radical scavenger. More work is needed in this area with anaemia treatment and prevention the effects of ROS in Crohn’s disease.

Table 1: Pro and anti-oxidant profile before and after iron supplementation in Crohn’s patient.

<table>
<thead>
<tr>
<th></th>
<th>Day 7</th>
<th>Reference values*</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDA: Malondialdehyde</td>
<td>4.81</td>
<td>1.44</td>
</tr>
<tr>
<td>SOD: Superoxide dismutase</td>
<td>17.49</td>
<td>1.64-2.40</td>
</tr>
<tr>
<td>GPx: Glutathione peroxidase</td>
<td>0.08</td>
<td>&gt;27.00</td>
</tr>
<tr>
<td>Vitamin A</td>
<td>2.02</td>
<td>0.35-1.75</td>
</tr>
<tr>
<td>Vitamin E</td>
<td>16.00-36.00</td>
<td></td>
</tr>
</tbody>
</table>


Discussion

A recent meta-analysis of European studies showed an overall anaemia prevalence of 27% for CD [9]. Two recent Scandinavian studies showed a significant decrease of the prevalence of anaemia in IBD during the disease course [10,11]. Indeed, Wiskin et al. [12], showed that 30% of pediatric IBD patients had persistent anaemia 2 years after diagnosis. In IBD cases, the intestinal blood loss and the reduced iron absorption due to inflammatory bowel, may lead to a negative iron balance [6]. However, causes of iron deficiency also could be self-imposed dietary restrictions in an attempt to improve or relieve gastrointestinal symptoms [13]. On the other hand, the inflammation results in interleukin-6 secretion that causes an increase in hepatic hepcidin synthesis, which potentially decreases iron duodenal absorption [14]. In this context, iron supplementation should be considered in cases of ID with or without anaemia [15]. Moreover, in presence of IBD, activated T lymphocytes reduce oral iron absorption. Thus, evidences indicate that less than half of the patients receiving oral iron supplementation tolerate the dose required to normalize iron and haemoglobin levels [16]. In addition, recent evidences showed that patients with a high degree of inflammation evidenced by elevated C-reactive protein levels may take longer to respond to the oral treatment of iron [17]. In contrast, responsiveness to intra-venous treatment appears to be independent of the inflammatory status of IBD patients [18]. Thus, intravenous iron supplementation should then be employed, showing a more effective response, promoting faster improvement of haemoglobin [8,19].

The only treatment option for ID is iron supplementation with metal ions. Iron frequently have unsaturated electrons, and is involved in the Fenton reaction, generating the highly reactive hydroxyl radical (-OH). Furthermore, iron is incorporated into and is essential for the function of ROS-producing enzymes such as nicotinamide adenine dinucleotide phosphate hydride (NADPH) oxidases [20]. The chemical properties of iron render it a potential hazard within the organism in that ferrous ion (Fe 2+), in small non-protein shielded chelates, can catalyze the production of ROS, which in turn can lead to peroxidation and radical chain reactions with molecular damage [21]. There is direct and indirect evidence suggesting that the chronically inflamed intestines of Crohn's patients may be subjected to considerable oxidative stress. Although certain amounts of iron and reactive oxygen species may play an important role in CD.

Moreover, the balance of ROS concentrations is supported by antioxidant defense mechanisms include vitamins, the enzymes SOD, GPx and transient ions such as Fe 2+ and Cu 2+ [22]. However, Crohn's patients have low blood levels of vitamin C and E [23]. They also have low levels of Cu, Zn, SOD, GPx, catalase, vitamin A and β-carotene in their intestinal mucosa, submucosa and muscularisserosa [24]. Corroborating with this, data found in the present study showed an increased level of MDA and low levels of vitamin E, SOD and GPx at the initial of treatment and remained after iron supplementation. This

References

countries: a systematic review and individual patient data meta-analysis. Inflamm Bowel Dis 20: 936-945.


