INTRODUCTION

Accumulating data indicate that oxidative stress (OS) plays a major role in the pathogenesis of multiple sclerosis (MS). Reactive oxygen species (ROS), leading to OS, generated in excess primarily by macrophages, have been implicated as mediators of demyelization and axonal damage in MS. ROS cause damage to main cellular structures and components such as lipids, proteins and nucleic acids (e. g.,

CASE REPORT

Effects of the whole-body cryotherapy on a total antioxidative status and activities of some antioxidative enzymes in blood of patients with multiple sclerosis-preliminary study

Elżbieta Miller¹, Małgorzata Mrowicka², Katarzyna Malinowska², Krystian Żołyński³, and Józef Kędziora²

¹III General Hospital in Łódź, Poland, Rehabilitation Ward, ²³Medical University of Łódź, Poland, Chair of Chemistry and Clinical Biochemistry, ³Medical University of Łódź, Poland, Clinic of Orthopedition and Traumatology, Poland

Abstract : Objective. There is evidence that multiple sclerosis (MS) is not only characterized by immune mediated inflammatory reactions but also by neurodegenerative processes. Neutralization of oxidative stress and excitotoxicity, might represent a therapeutec approach to provide neuroprotection in MS. The purpose of this study was to compare changes in total antioxidative status and activities of chosen antioxidative enzymes, such as : SOD, CAT in erythrocytes of patients with MS before and after using WBCT with control group. Materials and methods. 32 patients with multiple sclerosis (ICD10-G35) and 20 healthy subjects were recruited for the study. The examined MS group (n=16) was treated with a series of 10 daily exposures in a cryogenic chamber (2-3 min, from -120°C to -110°C) and program of exercises. The control MS group (n=16) had only exercises. Plasma TAS as well as SOD and CAT activities in erythrocytes were measured. Results. The level of TAS in MS patients was distinctly reduced compared to healthy subjects. After two weeks of WBCT treatment an increase of TAS in the whole examined group (p < 0,01) were observed in relation to control MS group. There was not increase of CuZnSOD and CAT activities. Conclusion. Our results suggest positive antioxidant effects of WBCT as a short-term adjuvant treatment for patients suffered due to MS. J. Med. Invest. 57 : 168-173, February, 2010

Keywords : antioxidants, multiple sclerosis, whole-body cryotherapy

INTRODUCTION

Accumulating data indicate that oxidative stress (OS) plays a major role in the pathogenesis of multiple sclerosis (MS). Reactive oxygen species (ROS), leading to OS, generated in excess primarily by macrophages, have been implicated as mediators of demyelization and axonal damage in MS. ROS cause damage to main cellular structures and components such as lipids, proteins and nucleic acids (e. g.,
RNA, DNA), resulting in cell death by necrosis or apoptosis. In addition, weakened cellular antioxidant defense systems in the central nervous system (CNS) in MS, and its vulnerability to ROS effects may augmented damage (1).

Thus, treatment with antioxidants might theoretically prevent propagation of tissue damage and improve both survival and neurological outcome.

Hypothermia has long been known as a potent putative neuroprotectant. It delays energy depletion, reduces intracellular acidosis and ischemia, related to the accumulation of excitotoxic neurotransmitters, and attenuates the influx of intracellular calcium. Additionally, hypothermia inhibits production of oxygen free radicals involved in the secondary damage, associated with reperfusion. It also suppresses mechanisms of blood-brain barrier degeneration and postischemic remodeling (2-4).

Treatment using total immersion of the body in extremely low temperatures was first introduced in Japan towards the end of the 1970s by Yamauchi T (5) who constructed the first cryogenic chamber and successfully used cryotherapy to treat rheumatism.

We considered that the effects of whole body cryotherapy (WBCT) exposures could be associated with increasing of antioxidative system, therefore the purpose of this study was to compare the effects of WBCT on changes in total antioxidative status (TAS) of plasma and activities of the chosen antioxidative enzymes in the erythrocytes of patients with MS, such as superoxide dismutase (SOD) and catalase (CAT).

CASE REPORT

Patients presentation

Inclusion/exclusion criteria for this study were a diagnosis of MS and the ability to ambulate independently. Patients with such conditions as: circulatory or breathing insufficiency, clotting, embolism, inflammation of blood vessels, open wounds, ulcers, serious cognitive disturbances, fever, addictions, claustrophobia, and over-sensitivity to cold were excluded from the study.

The subjects (n=32) received no hormones, vitamins, minerals or any other substitutions with antioxidative properties. None of the subjects had previously used WBCT. Prior to the study, all the subjects had undergone medical check-ups including neurological and internistic examination. The subjects were randomized into two groups (16 subjects in each). Both groups were treated by program of exercises. This includes a 20-30 min program of progressive balance and muscle strengthening exercises. All sessions included 5 minutes warm-up exercises. The examined group had participated additionally to WBCT sessions. 20 healthy volunteers were chosen to the study as a control to MS.

The protocol and procedures were done according to the Helsinki Declaration and were approved by Ethics Committee of the Medical University of Łódź, Poland. The study was performed in the Department of Biochemistry and Rehabilitation Division III General Hospital in Łódź.

Experimental design

An experimental trial (WBCT) consisted of the 16 subjects treated by a cycle of 10 exposures in a cryogenic chamber carried out daily from Monday to Friday. The cryogenic chamber has two rooms: the vestibule, with the temperature of -60°C and the main chamber, with temperature between -110°C and -160°C. Liquid nitrogen is used as the coolant. Sessions in the chamber lasted 2-3 min. Gregorowicz and Zagrobelny (6) provide guidance on the appropriated duration of exposure and temperature for adult patients as well as a list of medical conditions in which WBCT is unsuitable.

The study was carried out in the spring 2008. Observations were made in 3 groups: I-MS patients treated by kinesytherapy, II-MS patients immersed by WBCT and kinesytherapy and III-healthy volunteers. MS patients were examined at 2 stages: at the beginning of the treatment and at the end. Healthy subjects had one examination.

Blood samples were collected in cooled EDTA tubes centrifuged to isolate plasma and erythrocytes. The specimens were taken one hour before the 10 days cycle of therapy and one hour after the last immersion in both MS groups.

Biochemical investigations

Total Antioxidant Status (TAS) was measured in plasma samples (healthy group and MS patients) using the kit by Randox Laboratories Ltd. (Cat. No. NX 2332). The plasma volume taken to estimation was 5 μl, a total assay volume was 305 μl. The reaction was carried out during 3 min. And measured spectrophotometrically at 600 nm.

The activities of antioxidative enzymes were determined in erythrocytes.

Superoxide dismutase activity in erythrocytes was
measured according to Misra and Fridovich (7) methods. The absorbance of the examined samples was estimated at 380 nm using Becman spectrophotometer at 37°C. The activity was expressed as U/g Hb.

Catalase activity in erythrocytes was determined according to Beers and Sizer method (8). Absorbance was measured at 240 nm using Becman spectrophotometer. Enzymatic activity was expressed as Berg Mayer units U/gHb.

Statistical analyses

Results were statistically elaborated. Due to non-parametric distribution the Wilcoxon test was used to analyze changes. Results were compare with healthy subjects.

Results of TAS measured in mM were considered statistically significant (p<0.01) whereas CAT activity decreased. The results in both groups were uniform.

Results

Our studies have shown that the level of TAS in MS patients is distinctly reduced compared to healthy subjects (Fig. 1). WBCT caused statistically significant increase of TAS in plasma of MS patients (α=0.001) (from 0.3468 mM to 0.8126 mM) compared to non WBCT MS patients (from 0.3427 mM to 0.5266 mM) (Fig. 1 Table 1). In erythrocytes of MS patients the level of SOD activity is significantly

Table 1  Effects of WBCT and kinesytherapy [II] with kinesytherapy [I] on TAS of plasma and activities of CuZnSOD and CAT in the erythrocytes patients with MS

<table>
<thead>
<tr>
<th></th>
<th>MS* Kinesytherapy [I] (n=16)</th>
<th>MS WBCT** + Kinesytherapy [II] (n=16)</th>
<th>Healthy (n=20)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Before</td>
<td>After</td>
<td>Before</td>
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<tr>
<td></td>
<td>Median</td>
<td>SD</td>
<td>Median</td>
</tr>
<tr>
<td>TAS [mM]</td>
<td>0.29</td>
<td>0.22</td>
<td>0.43</td>
</tr>
<tr>
<td>CuZnSOD [U/gHb]</td>
<td>1515</td>
<td>310</td>
<td>1479</td>
</tr>
<tr>
<td>CAT [10^4IU/gHb]</td>
<td>17.5</td>
<td>2.7</td>
<td>17.1</td>
</tr>
</tbody>
</table>

*multiple sclerosis
**whole body cryotherapy

Fig. 1  Comparison of total antioxidative status (TAS) activity in plasma patients with multiple sclerosis (MS) treated by whole body cryotherapy (WBCT) and kinesytherapy [II] and only kinesytherapy [I] with healthy subjects.
reduced compared to healthy group (Fig. 2).

We have observed that activity of CuZnSOD in erythrocytes of MS patients after WBCT treatment was not statistically significant change (Fig. 2 Table 1).

Contrary to activity SOD, activity of CAT was distinctly (2-fold) higher in erythrocytes of MS patients than in healthy volunteers and exercise therapy had no effect on CAT activity (Fig. 3 Table 1).

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**Fig. 2** Comparison of superoxide dismutase (CuZnSOD) activity in erythrocytes patients with multiple sclerosis (MS) treated by whole body cryotherapy (WBCT) and kinesytherapy [II] and only kinesytherapy [I] with healthy subjects.

**Fig. 3** Comparison of catalase (CAT) activity in erythrocytes patients with multiple sclerosis (MS) treated by whole body cryotherapy (WBCT) and kinesytherapy [II], and only kinesytherapy [I] with healthy subjects.
DISCUSSION

Oxidative stress is a hallmark of neuregenerative disorders MS symptomatology and complications of its negative symptoms. OS could be caused by prolonged altered metabolism, exposure to exogenous oxidizing agents or compounds and it has been tied to inflammatory response.

The antioxidant defense system in the organism comprises enzymatic and non enzymatic components. The enzymatic antioxidant system includes a variety of antioxidant enzymes such as SOD, CAT, GPx. We observed that MS patients the activities of erythrocyte antioxidative enzymes, specially SOD is distinctly reduced (contrary to CAT) (Fig. 2, 3).

Antioxidants, whether synthesized endogenously or exogenously administered are reducing agents that neutralize the oxidative compounds (ROS) before they can cause damage to different biomolecules.

The measurement of TAS in plasma represents the body redox status better than does measure of the single circulating antioxidant.

After strenuous physical exercise, a significant increase in TAS may occur (9). We observed that after exercise MS patients TAS in plasma was increased compared to untreated patients.

The present data demonstrate that plasma TAS was found to be significantly lower in MS patients than in healthy group (Fig. 1). This findings indicate that an impaired antioxidant defense system that may be dependent partly on lower activity of SOD. It seems that the observed low level of TAS in plasma of MS patients may be also dependent on the low concentrations of endogenous antioxidants, mainly uric acid. In humans, over half the antioxidant capacity of blood plasma comes from uric acid. It is known that the lower values of uric acid (~194 μmol/l) have been associated with MS, whereas serum uric acid in healthy subjects was ~290 μmol/l. MS patients in remission reach the level of uric acid about 230 μmol/l. Uric acid like ascorbic acid are strong reducing agents and potent antioxidant (10).

It should be emphasized that despite lowering activities of CuZnSOD and CAT the total antioxidative status significantly increased from 0.3468 mM to 0.8126 mM. Maybe that other antioxidantive enzymes or non enzymatic substances like GSH, glutamate reductase, vitamins, microelements that can also affect this phenomenon.

There are evidences that both cryotherapy and moderate exercises cause increase in antioxidant status (9-12). Similar results were found in the study performed by Duque, et al. (13) in which mean TAS value significantly increased 2 minutes after the exposure to cold in the healthy man group treated with WBCT.

Siems, et al. (14) investigated the effects of the acute cold stimuli (winter swimming) on a number of oxidative stress markers and showed that these stimuli induced a decrease in the concentration of several major plasma antioxidants (i.e. ascorbic acid, uric acid) (14). In addition, erythrocyte level of oxidized glutathione and the ratio of oxidized glutathione/total glutathione also increased following the cold exposure (15). The authors also indicated the higher enzymatic protection (i.e.changes in the activities of red blood cells catalase, glutathione peroxidase, superoxide dismutase) in healthy individuals regularly practicing winter swimming activities compared with control subjects (16).

If protection against oxidative stress due to WBCT treatment develops, increase in the values of TAS observed after the exposures to cold stress should reach the values observed in healthy subjects.

The obtained results suggest that cryotherapy may play an important role in the process of activation of antioxidative properties in MS patients, since the increase in the TAS was considerably greater after using WBCT than after physical exercise.

WBCT is worth of particular concern in treatment of patients with MS because of safety of used procedure, minimal side effects.

CONCLUSION

Our observations showed that WBCT resulted in the increase of TAS ten days after the exposure.

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