Collagen synthesis, nitric oxide and asymmetric dimethylarginine in diabetic subjects undergoing hyperbaric oxygen therapy

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Short title: Hyperbaric oxygen therapy and wound healing
The main pathological condition in patients with impaired wound healing is diabetes mellitus. These patients have significantly low circulating nitric oxide (NO) levels because the stimulatory action of insulin on NO synthesis is absent. Additionally, asymmetric dimethylarginine (ADMA), an inhibitor of NO synthase, is increased owing to the generation of oxidative stress. NO was thought to contribute to wound healing. Hyperbaric oxygen (HBO) treatment is generally used in order to accelerate the healing of wounds. The aim of this study was to determine the changes in plasma procollagen type I and III N-terminal peptides (PINP and PIIINP), total nitrite/nitrate (NOx) and ADMA levels; and to evaluate their relation to healing during the HBO treatment of foot ulcers. Data obtained from 18 diabetic patients before and after the HBO therapy were compared statistically by the Wilcoxon test. NOx was increased in 11 and ADMA was decreased in 12 patients following HBO treatment. Both PINP (32.6 ± 29.4 µg/L vs 44.3 ± 33.4 µg/L) and PIIINP (6.97 ± 3.01 µg/L vs 7.92 ± 2.49 µg/L) were significantly increased (p<0.05). Progressive reductions were observed in wound areas, as assessed by the digital wound imaging. In 12 patients, wounds healed by 50% or higher; whereas only two subjects had minimal improvements (15% or less healing). The duration of diabetes was negatively correlated with wound healing (r=-498, p<0.05). This study suggests that increased collagen synthesis is associated with wound healing during hyperbaric oxygen therapy. Nitric oxide generation may also contribute to the healing process.

**Key words**

Introduction

Wound healing is a complex process, involving inflammation, fibroplasia, neovascularization, collagen deposition, epithelialization and wound contraction. Collagen synthesis in wounds has been documented by increased concentrations of fibroblasts, type I collagen mRNA, increased production of growth factors, and decreased activity of the matrix metalloproteinases (Sheikh et al 2000, Kang et al 2004). On the other hand, elevated expression of extracellular matrix metalloproteinases, decreased proliferation rate of fibroblasts and decreased expression of growth factors have been demonstrated in nonhealing chronic wounds (Hehenberger et al 1998, Cowin et al 2001, Norgauer et al 2002). Measurements of the N-terminal propeptides of type I (PINP) and III (PIIINP) collagens in blood are commonly used as markers of collagen formation. The concentration of these propeptides reflects the synthesis rate of type I and III collagens. During wound healing, types I and III collagens which are produced and deposited by fibroblasts increase the tensile strength of the wound (Mutsaers et al 1997). It was reported that scar fibroblast selectively increases the biosynthesis of type I collagen, this abnormal metabolism results in the deposition of collagen in scar and alters the steady-state ratio of collagen type I/III in the process of wounding healing (Zhang et al 1995, Niessen et al 1999). It has been shown by animal experiments that exposure to hyperbaric oxygen increases the expression of type I collagen in healing of tendon laceration (Ishii et al 1999).

The main pathological condition in the patients with impaired wound healing is diabetes mellitus. In diabetic patients, high blood glucose hinders proliferation of cells and decreases collagen production in foot ulcers that may result in amputation of the extremities (Hehenberger et al 1998). Hyperbaric oxygen (HBO) therapy has been widely used to treat chronic wounds associated with pathological conditions compromising blood supply and tissue oxygenation (Abidia et al 2003). 20-40 exposures of HBO have been shown to exert beneficial effects on glucose metabolism in diabetic patients (Kakhnovskii et al 1982); and to improve the healing in diabetic foot ulcers (Faglia et al 1996, Kalani et al 2002).

NO, a versatile signal molecule, is the most potent endogenous vasodilator that regulates many processes including collagen synthesis and matrix remodeling. Kuboki et al (2000) found that diabetic patients have significantly lower circulating NO levels because the stimulatory action of insulin on NO synthesis is absent. Nitric oxide synthase (NOS), the enzyme that uses L-arginine and molecular oxygen as substrates for the production of NO and L-citrulline, is the key enzyme for the regulation of NO availability. Asymmetric
dimethylarginine (ADMA) is an endogenous inhibitor of NOS, and is derived from the catabolism of proteins containing methylated arginine residues. Increased oxidative stress has been shown to lead to increased levels of ADMA (Baylis 2006). In diabetic patients, increased levels of ADMA have been reported (Abbasi et al 2001), but its possible involvement in the wound healing has not been investigated.

In this study, attempts were made to determine the changes in plasma PINP and PIIINP, total nitrite/nitrate (NOx) and ADMA levels; and to evaluate their relation to wound healing during the HBO treatment of foot ulcers in diabetic patients.

**Subjects and methods**

**Subjects**

Eighteen patients (13 men, 5 women) with diabetic foot ulcers who received hyperbaric oxygen therapy were included in the study. The procedures were in accordance with the revised form of the Helsinki Declaration 2004 and all participants signed an informed consent form. The study protocol was approved by the local ethical committee. Patients were followed by the same physician responsible from diabetic control, wound care and antibiotic therapy according to the clinical and laboratory findings, and were given a diet depending on their metabolic needs without vitamin supplementation. No significant difference occurred in the glucose and HbA1c levels of patients during the experimental period. Fasting blood glucose levels ranged between 100-300 mg/dl and blood HbA1c concentrations were between 6.3-12.6 %. The clinical characteristics of patients are shown in Table 1.

**HBO treatment**

Along with the standard wound care and medical therapy, all subjects underwent HBOT according to a routine protocol for diabetic foot ulcers with healing problems, determined by the European Committee on Hyperbaric Medicine and also by the Hyperbaric Oxygen Committee of Undersea and Hyperbaric Medicine (Wattel 1998, Feldmeier 2003). HBOT was carried out in a multiplace hyperbaric chamber once a day, and six days in a week. The treatment protocol was inhalation of 3x25 min periods of 100 % oxygen at a pressure of 2.4 ATA, interspersed with 5 min periods of air breathing.

**Sample collection and wound size evaluation**

On the day of HBO treatment, the patients were admitted to the clinic between 8.00-11.00 a.m. Venous blood samples (10 ml) were collected from each patient prior to the hyperbaric
oxygen therapy in a standard sterile vacuum tube containing EDTA, and immediately centrifuged at 600 g for 15 min. Aliquots of plasma were stored at -80°C until used for the determinations of PINP, PIIINP, NOx and ADMA. After 25-30 sessions of HBO therapy, varying degrees of clinical healing were observed in the patients, and blood samples were collected for the repetitive measurements.

Clinimetric evaluation of the wounds was based on digital wound imaging, and the changes in the wound surface area were calculated with planimetric measurements using a constant reference (Kantor and Margolis 1998, Quan et al 2007).

**Analytical procedures**

PINP and PIIINP levels were measured using commercially available competitive radioimmunoassay kit (Orion Diagnostica UniQ, Espoo, Finland). Minimum detectable doses were 2 µg/L and 0.3 µg/L, the interassay CVs were <9.8 % and <6.5 %; and intraassay CVs <9.8 % and <3.0 %, respectively. Results were expressed as µg/L.

NOx (nitrite and nitrate) analyses were carried out using total nitric oxide assay kit (R&D System Europe, Abingdon, UK). Minimum detectable dose ranges from 0.09 to 0.78 mmol/L, the interassay CV was <4.6 %, and intraassay CV <2.5 %. Results were expressed as µmol/L.

ADMA levels were measured in samples by a commercial ELISA kit (DLD Diagnostika, Hamburg, Germany). The reference ranges were between 0.4 and 0.75 mmol/L; the intraassay and interassay coefficients of variation were <8% and <10%, respectively. Results were expressed as µmol/L.

**Statistics**

Results were expressed as means±SD. The significance of differences between groups was assessed using Wilcoxon test. P values less than 0.05 were regarded as significant. Correlations were estimated using the Spearman test. All analyses were performed with the Statistical Package for the Social Sciences (12.0 software version, SPSS Inc, Chicago, IL, USA).

**Results**

Biochemical data obtained from the patients before and after 25-30 sessions of HBO are demonstrated in Table 1.
Table 1. Clinical and laboratory characteristics of the patients.

<table>
<thead>
<tr>
<th></th>
<th>Before HBOT</th>
<th>After HBOT</th>
</tr>
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<tbody>
<tr>
<td>Men/Women</td>
<td>13/5</td>
<td>---</td>
</tr>
<tr>
<td>Age</td>
<td>61 ± 8.26</td>
<td>---</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>26.1 ± 3.73</td>
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<tr>
<td>Duration of DM (years)</td>
<td>18.6 ± 10.0</td>
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<tr>
<td>PINP (µg/L)</td>
<td>32.6 ± 29.4</td>
<td>44.3 ± 33.4*</td>
</tr>
<tr>
<td>PIIINP (µg/L)</td>
<td>6.97 ± 3.01</td>
<td>7.92 ± 2.49 *</td>
</tr>
<tr>
<td>ADMA (µmol/L)</td>
<td>1.33 ± 0.5</td>
<td>1.20 ± 0.49</td>
</tr>
<tr>
<td>NOx (µmol/L)</td>
<td>76.6 ± 41.5</td>
<td>83.7 ± 38.9</td>
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*p<0.05 in comparison with the baseline values. (DM: Diabetes mellitus)

PINP and PIIINP levels were found increased in 14 patients when compared with the baseline values. These increments gave a statistical significance (p<0.05). 11 patients with increased levels of procollagen peptides had 50% or more decreased wound surface area. Wounds were evaluated by measuring the surface area twice on digital wound imaging, prior to the therapy and after 25 to 30 successive HBO treatments; and the degree of healing was expressed as the percentage of healed area. Two subjects were excluded from this evaluation due to the absence of the digital imaging data. Of 16 subjects, two had minimal improvements on wound surface (with 15% or less healing), whereas the remaining patients gave a favorable response as follows: In 7 patients, the degree of wound healing 78% or higher; and 7 patients with the healing percentages 33 to 57 (Figure 1).

When the patients were classified according to the degree of healing (<50% or >50%), the baseline NOx levels altered significantly between the groups (p<0.05; Figure 2). In addition, elevated NOx levels were seen in patients whose wounds were healed by 50% or more following HBO therapy. On the other hand, the patients who showed minor improvements (less than 50% of healing) had decreased levels of NOx after the exposure to HBO. The individual variations in NOx levels were demonstrated in Figure 3.
Figure 1. Wound surface area (cm$^2$) as estimated by the digital wound imaging in each patient. ○Before HBOT   ●After HBOT
Figure 2. Plasma NOx levels before and after 25-30 sessions of HBO therapy in diabetic patients whom were classified according to the degree of wound area change.\textsuperscript{a} \( p < 0.05 \) in comparison to basal levels (before HBO).
\textsuperscript{b} \( p < 0.05 \) in comparison to the patients with less than 50\% healing (before HBO).

Figure 3. Plasma NOx levels in diabetic patients before and after HBO therapy. Data right to the dotted line show increased NOx levels following HBO (11 patients). Patients with decreased NOx levels are placed left to the dotted line. ○ Before HBOT  ● After HBOT

Baseline plasma ADMA was lower in patients (1.25±0.5 \( \mu \text{mol/L} \)) who responded favorably (50\% or more healing) to HBO treatment when compared to that (1.59±0.56 \( \mu \text{mol/L} \)) in the others who poorly responded to the treatment (below 50\%). In 7 patients with 78\% or more healing, plasma ADMA was lowest before the therapy (1.09±0.55 \( \mu \text{mol/L} \)). No significant difference was observed between ADMA levels before and after HBOT when the patients were classified according to the degree of healing (data not shown).

The duration of diabetes was negatively correlated with wound healing (\( r = -0.498, p < 0.05 \)); but after adjusting for HbA\textsubscript{1c} by two-tailed partial correlation, this negative correlation no longer existed.
No correlation was detected between PINP and PIIINP levels. Neither ADMA nor NOx levels were correlated with the procollagen peptides. There was no significant relation between these two parameters either before or after the 25-30 sessions of HBOT. The relation between the changes in wound area and in the above-mentioned biochemical parameters was examined by the categorical analysis (the increment in the measured parameter versus >50% of healed area), the changes in NOx levels were found to be significantly associated with the degree of healing (p=0.001).

Discussion

The improvement of wound healing is directly associated with the periwound-tissue oxygen tension (Smith et al 1996). The beneficial effects of hyperbaric oxygen on the healing of foot ulcers and glucose metabolism have been documented in patients with diabetes mellitus (Kakhnowskii et al 1982, Kalani et al 2002, Abidia et al 2003, Kessler et al 2003). In the present study, type I and III procollagen N-peptides were found significantly increased in sera following 25-30 sessions of HBO. This finding suggests that successive exposures to HBO increase collagen synthesis as the rate of healing is accelerated. It is well known that molecular oxygen is essential for collagen synthesis, especially for the post-translational hydroxylation of proline and lysine in procollagen during the process of collagen maturation. However, the concentrations of PINP and PIIINP measured in this study reflect the rate of synthesis rather than maturation of the procollagen fibers. It has previously been shown that HBO exposure may activate secretion of growth factors through the generation of reactive oxygen species (Tandara and Mustoe 2004, Shyu et al 2008). After HBO therapy, an increase in transforming growth factor (TGF)-β production has been observed (Kang et al, 2004). Together with the previous reports, our findings support a promoting role for oxygen on collagen synthesis rather than maturation process.

It has previously been shown that deposition of collagen types in wound area differs with regard to the stages of healing. In the earliest stage, excessive type III collagen is seen but later type I collagen predominates (Mutsaers et al 1997). Since the degree of wound healing was not similar in our subjects, a significant association between PINP and PIIINP could not be detected.

Ongoing experimental and clinical wound healing studies have presented NO as a critical mediator of tissue repair (Schwentker and Billiar 2003). NO levels were found significantly increased in wound fluid (Boykin and Baylis 2007); while it remained unaltered in plasma after a short-term HBO treatment (Chen et al 2007). Reports related to the effect of
hyperbaric oxygen on NO production in normal tissues are conflicting (Thom et al 2003, Elayan et al 2000, Akgul et al 2007). In our study, the basal levels of NOx were significantly higher in patients who exhibited a lesser degree of wound healing in response to the therapy. In this group, NOx levels were significantly decreased after the HBO exposure. In contrast, the patients who had a good response to HBO therapy had a relatively low baseline NOx, which was followed by a slight elevation after HBO. In a recent study, an improved wound healing due to increased nitric oxide generation has been demonstrated after HBO (Gallagher et al 2007). These observations led us to suggest that there is an optimum NO level for its contribution the healing process.

In our study, ADMA levels were slightly higher than those in healthy subjects, which estimated to be ~1 µmol/L (Cooke 2000). This elevation may be related to oxidative stress owing to diabetes. Some studies have shown an elevation while others demonstrated a decrease in ADMA levels of diabetic patients (Abbasi et al 2001, Paiva et al 2003), but the role of this metabolite on wound healing has not been investigated. In this study, we observed a lesser degree of healing in patients with relatively high baseline ADMA levels; while baseline ADMA levels were lowest in 7 patients who favorably responded to HBO treatment with 78% or more healing. Since our study had a limited sample size, further studies should be carried out in order to claim that determination of plasma ADMA levels may be useful to predict the prognosis of healing in diabetic patients. Supplementation of L-arginine, which is a common precursor for NO and polyamine synthesis, has enhanced wound healing in experimental animals (Shi et al 2003). Therefore, the ratio of arginine to ADMA appears to be more important than ADMA alone for the repairing process.

In conclusion, HBO treatment results in significant increases in both type I and III collagen synthesis. A slight increase observed in NO levels during this process may contribute to the wound healing. Further studies are needed to establish whether the effect of hyperbaric oxygen therapy on healing of diabetic foot lesions can be enhanced by L-arginine supplementation in humans.

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References


KUBOKI K, JIANG ZY, TAKAHARA N, HA SW, IGARASHI M, YAMAUCHI T, FEENER EP, HERBERT TP, RHODES CJ, KING GL: Regulation of endothelial


