XPD and hOGG1 gene polymorphisms in reperfusion oxidative stress

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ABSTRACT. Knee replacement surgery is an ischemia/reperfusion model, as it uses tourniquet applied to the knee area to stop the blood flow during the operation. Fifty patients that were undergoing elective arthroscopic knee surgery were included in our study. Human 8-oxoguanine glycosylase 1 (hOGG1) is an enzyme to repair specific DNA lesions and a good marker of hydroxyl radical damage to DNA. XPD is another DNA repair gene. We investigated the effect of hOGG1 (Ser326Cys) and XPD (Lys751Gln) polymorphisms on the oxidative stress level after reperfusion. To evaluate oxidative stress conditions, we measured 8-hydroxyguanosine and malondialdehyde (MDA) levels. Polymorphism analyses were done by PCR-RFLP; serum 8-hydroxyguanosine and MDA levels were determined by enzyme-linked immunoassay. There were no significant differences between serum MDA and 8-hydroxyguanosine levels in the samples taken before and after tourniquet application; none of these parameters were related with hOGG1 genotypes. However, we
observed increased MDA levels after tourniquet application in M allele carriers for the XPD gene; this could mean that M allele carriers are more prone to DNA damage due to oxidative activity.

**Key words:** Oxidative stress; XPD; hOGG1; Polymorphisms

**INTRODUCTION**

The metabolic effects of ischemia on skeletal muscle tissue have been extensively described in animal studies (Westman et al., 2007). Unfortunately, the underlying mechanisms are not yet understood. One model of ischemia/reperfusion conditions to be studied in humans can be knee replacement surgery, because in this case, a tourniquet is applied on the knee area to stop the blood flow through the area during the operation (Westman et al., 2007; de Groot et al., 2010). Such orthopedic surgeries are thought to be good models to evaluate the production of antioxidants and for testing antioxidant effects of anesthetic agents. In knee replacement surgery, tourniquet application has been shown to induce ischemia, anaerobic metabolism and oxidative stress, increasing the production of ROS (reactive oxygen species) (Westman et al., 2007).

Reactive oxygen species are known to attack nucleic acids and their related compounds, generating various modified nucleobases that seem to play pivotal roles in mutagenesis, and 7,8-dihydro-8-oxoguanine (8-OHdG) is one of the most abundant among these modified nucleobases (Cheng et al., 1992; Weiss et al., 2005). It is a highly mutagenic change that can occur as a consequence of the oxidation process, and the protein product of this change is frequently used to determine the level of oxidative stress (Wu et al., 2004). Human 8-oxoguanine glycosylase 1 (hOGG1) is the enzyme that repairs 8-OHdG lesions by excising 8-hydroxyguanine from damaged DNA, and it is a good marker of hydroxyl radical damage to DNA (Kohno et al., 1998; Weiss et al., 2005). The hOGG1 gene (hOGG1) is located on chromosome 3p26.2 (Weiss et al., 2005). A polymorphism of hOGG1, a guanine-to-cytosine substitution in exon 7, that causes a serine (Ser) to cysteine (Cys) amino acid change at codon 326 (Ser326Cys), has been thought to be a functional polymorphism (Vodicka et al., 2007; Yuan et al., 2010). Oxidative stress is known to cause degenerations in vital cellular components such as lipids and proteins, besides DNA, increase malondialdehyde (MDA) levels and decrease reduced glutathione activity (Goyal and Anil, 2007). MDA is a decomposition product of oxidized lipids, or it can be formed by the reaction between acrolein and hydroxyl radicals; it is a valid biomarker of oxidative stress (Lykkesfeldt, 2007).

In this study, we aimed to investigate the effect of hOGG1 (Ser326Cys) and XPD (Lys751Gln) polymorphisms on the oxidative stress levels after reperfusion by measuring serum concentrations of 8-hydroxyguanosine and MDA as oxidative stress markers.

**MATERIAL AND METHODS**

**Patients**

After obtaining approval to conduct the study from the local ethics committee and written informed consent from each patient, demographic data were collected prospectively. The patients selected for this study were 18-60 years old (N = 50) with ASA physical status...
Gene polymorphisms and oxidative stress

I-II, undergoing elective arthroscopic knee surgery with a pneumatic tourniquet time of 30-45 min (Table 1). Patients taking antioxidant drugs or those with metabolic, renal or hepatic disturbances were excluded from the study.

Anesthetic management

All operations were performed by senior surgeons and two senior residents. The surgical team and the anesthesiologist, who were responsible for the postoperative case of the patients, were blinded to the presence or absence of the hOGG1 (Ser326Cys) gene.

All patients were premedicated with 0.5 mg oral alprazolam 45 min before surgery. Heart rate (HR), non-invasive blood pressure (NIBP), peripheral oxygen saturation (SpO₂), and end-tidal partial pressure of carbon dioxide (EtCO₂) were monitored in the operating room. Anesthesia was induced with 2 mg/kg propofol and 0.5 mg/kg rocuronium bromide. Laryngeal mask airways were inserted after preoxygenation with 100% oxygen. Lungs were ventilated at a tidal volume of 10 mL/kg at a rate of 12 breaths per minute. Anesthesia was maintained with 1% sevoflurane and 30% N₂O carried in oxygen. Remifentanil, 0.25 μg·kg⁻¹·h⁻¹, was started as soon as propofol was applied. The pneumatic tourniquet was inflated after the exsanguination of the limb and the pressure was kept between 300-350 mmHg throughout the surgery.

Venous blood samples of 50 men were collected before and after tourniquet. The antecubital vein of the left arm was used to collect the baseline venous samples before tourniquet application during the intravenous line placement. Second venous samples were taken from the antecubital vein of the contralateral arm 10 min after the deflation of the tourniquet.

At the end of the surgery, all patients were extubated after reversal of neuromuscular blockade with 0.01 mg/kg atropine and 0.05 mg/kg neostigmine.

DNA isolation and genotyping

Genomic DNA was extracted from isolated lymphocytes by a standard salting out procedure (Miller et al., 1988). The extracted DNA was used for characterization of the following polymorphic DNA repair genes. Polymerase chain reaction (PCR) followed by restriction fragment length polymorphism (RFLP) was used for genotyping (Wood et al., 2001).

The hOGG1 (Ser326Cys) polymorphism, a C→G transversion in exon 7 was determined using the primers (forward) 5'-ACTGTCACTAGTCTCACCAG-3' and (reverse) 5'-GGAAGGTGCTTGGGGAAT-3'. The PCR conditions consisted of an initial denaturation step at 95°C for 2 min, followed by 35 cycles of 95°C for 1 min, 60°C for 1 min and 75°C for 1 min, and a final extension step at 72°C for 5 min. The 200-bp PCR product was digested with Fnu4HI. Genotypes were determined for the polymorphism as Ser/Ser (200 bp), Ser/Cys (100, 200 bp), or Cys/Cys (100 bp).

The XPD Lys751Gln polymorphism, an A→C transversion in exon 23 (position 35931), was determined using the primers (forward) 5'-CCTCTCCCTTTCCCTGTTC-3' and (reverse) 5'-CAGGTGAGGGGGGACATCT-3'. The PCR conditions consisted of an initial denaturation step at 95°C for 2 min, followed by 35 cycles of 94°C for 1 min, 52°C for 1 min and 72°C for 1 min, and a final extension step at 72°C for 5 min. The 734-bp PCR product was digested with PstI; the Gln allele was cut into 646- and 88-bp fragments (Lys allele not digested) (Yeh et al., 2005).
Determination of serum 8-hydroxyguanosine levels

Whole blood samples were centrifuged at 1500 g for 10 min at room temperature and serum was immediately removed and frozen at -20°C until serum 8-hydroxyguanosine levels were determined by enzyme-linked immunoassay (Cell Biolabs Inc., CA, USA).

Determination of serum MDA levels

Blood samples were centrifuged for 10 min at 1500 g at room temperature and serum was immediately removed and frozen at -20°C until serum MDA levels were determined by enzyme-linked immunoassay.

Statistical analysis

The statistical analyses were performed using the SPSS 11.0 statistical software package (SPSS, Chicago, IL, USA). P values less than 0.05 were considered to be statistically significant. Also, linear regression analysis was used to investigate genotype and DNA damage association. The Student t-test was used to investigate the serum MDA levels between the genotypes. Allele frequencies were found using the gene counting technique. To determine the relative risks, odds ratios and 95% confidence intervals were used.

RESULTS

Demographic characteristics are summarized in Table 1.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Patients (N = 50)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)*</td>
<td>50 (20-80)</td>
</tr>
<tr>
<td>Gender (female/male, N)</td>
<td>22/29</td>
</tr>
<tr>
<td>Shivering (%)</td>
<td>7.8%</td>
</tr>
<tr>
<td>Nausea (%)</td>
<td>2%</td>
</tr>
<tr>
<td>Duration of tourniquet application (min)</td>
<td>41 (15-149)</td>
</tr>
<tr>
<td>Duration of anesthesia application (min)</td>
<td>60 (25-160)</td>
</tr>
</tbody>
</table>

*Values are reported as median (interquartile range).

There was no significant difference between serum MDA (P > 0.877) and 8-hydroxyguanosine (P > 0.195) levels in the samples taken before and after tourniquet application (Table 2).

Table 2. MDA and 8-hydroxyguanosine levels before and after the tourniquet application.

<table>
<thead>
<tr>
<th></th>
<th>MDA (pmol/mg)</th>
<th>8-Hydroxyguanosine (pmol/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before tourniquet</td>
<td>35.56 ± 9.20</td>
<td>8.60 ± 10.66</td>
</tr>
<tr>
<td>After tourniquet</td>
<td>35.82 ± 9.50</td>
<td>12.14 ± 11.84</td>
</tr>
</tbody>
</table>

The associations between genotypes, alleles and MDA and 8-hydroxyguanosine levels are shown in Table 3. We did not find any relationship between hOGG1 genotypes and serum levels of MDA and 8-hydroxyguanosine. With regard to XPD genotypes and alleles, we ob-
erved an increase in MDA levels after tourniquet application in the M allele (P < 0.046, 95% CI = 0.142-13.600) and an almost significant decrease in wild-type LL genotype patients after tourniquet application (P < 0.056, 95% CI = 0.190-13.650; Table 3).

DISCUSSION

In orthopedic surgeries, tourniquets are commonly applied to reduce blood loss and obtain a clearer surgical field; this makes it easier to develop scientific projects in the area of ischemia-reperfusion injuries. Because of DNA damage, ROS activity on membrane lipids produces peroxides such as MDA, which has been used as a marker of lipid peroxidation (Del Rio et al., 2005). In our study, we had such patients undergoing elective arthroscopic knee surgery; our aim was to investigate the changes in MDA levels before and after the tourniquet application, and we found no difference in these levels in the total patient population. Despite our results, it was previously found that ROS and MDA levels increased in the reperfusion stage after ischemia (Cheng et al., 2003; Westman et al., 2007; Lykkesfeldt, 2007). The tourniquet application durations and the times of sample collection usually differed in previous studies; there was no suggested standard duration for these applications, but the duration of the ischemic period was also found to be important in reperfusion injury (Homer-Vanniasinkam et al., 2001; Kloner and Jennings, 2001; Cheng et al., 2003). Thus, our results may differ because of such differences.

We observed an increase of MDA levels after tourniquet application in M allele carriers for the XPD gene, and contrarily, there was an almost significant decrease in LL genotype patients. This may indicate that M allele carriers for the XPD gene are much more prone to DNA damages because of oxidative damage. However, in order to correlate such parameters, we had to consider also the effects of anesthetic agents or other medications on the oxidative damage. There have been several studies about alprazolam, propofol and N-acetylcysteine, and it was previously shown that alprazolam is an effective agent in ameliorating alterations in oxidative stress and attenuating malondialdehyde levels (Goyal and Anil, 2007). Propofol was also shown to have antioxidant activity against lipid peroxidation (Murphy et al., 1993; Aldemir et al., 2001).

We did not find any significant difference in 8-hydroxyguanosine levels before and after tourniquet application. Although the polymorphism causing serine (Ser) to cysteine (Cys) amino acid change at codon 326 (Ser326Cys) was thought to be functional in some previous

<table>
<thead>
<tr>
<th>XPD genotype (N)</th>
<th>MDA levels before the operation (pmol/mg)</th>
<th>MDA levels after the operation (pmol/mg)</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>MM (3)</td>
<td>21.33 ± 17.50</td>
<td>30.66 ± 4.5</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>LM (20)</td>
<td>36.68 ± 9.17</td>
<td>39.21 ± 8.44</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>LL (18)</td>
<td>36.68 ± 5.81</td>
<td>33.26 ± 10.19</td>
<td>0.056*</td>
</tr>
<tr>
<td>M+ (23)</td>
<td>34.59 ± 11.41</td>
<td>38.04 ± 8.48</td>
<td>0.046**</td>
</tr>
<tr>
<td>L+ (38)</td>
<td>36.68 ± 7.57</td>
<td>36.23 ± 9.71</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>hOGG1 genotypes</th>
<th>8-Hydroxyguanosine levels before the operation (pmol/mg)</th>
<th>8-Hydroxyguanosine levels after the operation (pmol/mg)</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ser/Ser (n:39)</td>
<td>236.51 ± 423.15</td>
<td>239.43 ± 421.58</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Ser/Cys (n:12)</td>
<td>93.25 ± 285.61</td>
<td>95.83 ± 284.72</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Cys/Cys (n:9)</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
</tbody>
</table>

*P < 0.056, 95%CI = 0.190-13.650, **P < 0.046, 95%CI = 0.142-13.600.
studies (Vodicka et al., 2007; Yuan et al., 2010), we could not find such difference in our study.
Rather, our findings are in line with those of Kohno et al. (1998).

Our study is a rare one containing several parameters altogether, so in case of enlarging the study group, there will be more obvious results to contribute to the understanding of the molecular basis of reperfusion conditions.

REFERENCES


