OXIDATIVE STRESS PARAMETERS AFTER ABDOMINAL HYSTERECTOMY AND THEIR RELATIONSHIPS WITH QUALITY OF RECOVERY

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ABSTRACT

Study aimed to investigate relationship between oxidative stress markers and postoperative recovery in woman after abdominal hysterectomy, as well as to test the hypothesis that different analgesics differently influence redox status.

The quality of recovery was evaluated with a QoR-40 questionnaire in fifty-one patients who underwent abdominal hysterectomy, preoperatively and on the 1st, 2nd, 3rd postoperative days (POD1, 2, 3). Blood samples were collected at baseline (T0), 3 (T1), 24 (T2), 48 (T3) and 72 (T4) hours after surgery. Oxidative stress markers concentrations (TBARS, NO₂⁻, H₂O₂, O₂⁻) as well as antioxidative enzymes (SOD, CAT, and GSH) were analyzed.

QoR-40 total score significantly declined on POD1 and POD2 and returned to baseline levels on POD3 (p<0.001). H₂O₂ levels decreased significantly from T0 to T3 and then, increased at T4 (p=0.011). Changes of TBARS and H₂O₂ from T0 to T3 showed significant and negative correlation (r=-0.303, p=0.046). There was no significant correlation between QoR-40 total score and any parameter of oxidative stress response (p>0.05). Changes in TBARS levels from T0 to T3 were statistically significant between the study subgroups primarily due to increase of the concentrations in patients receiving paracetamol (p=0.031). Patients age, duration of surgery and cigarette smoking status showed significant influences on and association with some oxidative stress response markers (TBARS, O₂⁻, CAT) (p<0.05).

Women who underwent hysterectomy had significant changes of H₂O₂ and TBARS activity however, those changes were not associated with changes of QoR-40 total scores during recovery.

Keywords: hysterectomy; postoperative period; oxidative stress.

SAŽETAK

Studija je imala za cilj da ispita povezanost između markera oksidativnog stresa i postoperativnog oporavka kod pacijentkinja podvrgnutih abdominalnoj histrektomiji kao i da testira hipotezu da različiti analgetici različito utiču na redoks status.

Kod 51 pacijentkinje podvrgnute abdominalnoj histrektomiji zbog benignih bolesti uterusa kvalitet postoperativnog oporavka je testiran sa upitnikom QoR-40, preoperativno, kao i prvog, drugog i trećeg postoperativnog dana (POD 1, 2 i 3). Uzorki krvi su uzeti preoperativno (T0), 3 (T1), 24 (T2), 48 (T3) i 72 (T4) sati posle operacije. Analizirane su koncentracije markera oksidativnog stresa (TBARS, NO₂⁻, H₂O₂, O₂⁻) kao i enzima antioksidacione odbrane (SOD, CAT i GSH).

Vrednosti ukupnog QoR-40 skora su značajno opale u POD1 i POD2 i vratile se na preoperativne vrednosti u POD3 (p<0.001). Koncentracije H₂O₂, su značajno opale od T0 do T3 i porasle u T4 (p=0.011). Promene od T0 do T3 u vrednostima TBARS-a i H₂O₂ su u značajnoj medusobnoj negativnoj korelaciji (r=-0.303, p=0.046). Nije bilo značajne korelacije između ukupnog QoR-40 skora i bilo kog parametra oksidativnog stresa (p>0.05). Promene vrednosti TBARS-a od T0 do T3 su bile značajno različite između studijskih podgrupa pre svega zbog porasta koncentracija kod pacijentkinja koji su primale paracetamol (p=0.031). Starost pacijenata, trajanje operacije kao i pušenje pokazali su povezanost sa pojedinim markerima stres odgovora (TBARS, O₂⁻, CAT) (p<0.05).

Kod pacijentkinja podvrgnutih histrektomiji dolazi do značajnih promena u H₂O₂ i TBARS-u, ali one nisu značajno udržane sa kvalitetom oporavka.

Keywords: histrektomija; postoperativni oporavak; oksidacioni stres.

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Received / Primljen: 02.08.2017. Accepted / Prihvaćen: 03.09.2017.

UDK: 616.146-089.85, 616-008.9:577.334:546.21
DOI: 10.1515/SJECR-2017-0046

27
INTRODUCTION

The abdominal hysterectomy, the most common major gynecological procedure, disturb patients quality of life during postoperative period despite of modern advances in both surgery and anesthesia (1, 2). Re-establishing physical and psychological balance as well as disappearing of unpleasant symptoms, take certain amount of time during period known as postoperative recovery (3). Since postoperative recovery is nowadays recognized as a valid clinical outcome, many efforts have been done in developing instruments for its measurement as well as for exploring its pathophysiological basis (4, 5).

Surgical trauma activates several biological pathways and among them, there is increased production of reactive species causing the state known as the oxidative stress (6, 5). On the other side, the researchers have found positive association between increased oxidative stress markers and some symptoms, which participated in health-related quality of life perception like fatigue, depression, anxiety, nausea and headache (7-10). Although patients are combating with many of these symptoms during postoperative recovery the studies, which measured quality of life after surgery in details, and using validated rating questionnaires were not common.

Some additional issues increase the complexity of the matter and, consequently, the need for further investigations. For example, the synthesis of reactive species depends on the type and the extent of the intervention, anesthesia techniques and anesthetics (11-14). In addition, analgesics could modulate oxidative stress response during postoperative period either decreasing (e.g. morphine) or increasing it (e.g. ibuprofen) (15, 16). In some cases, researchers elucidated the mechanisms of stress response alterations in fine details. Paracetamol, a phenolic compound, has a large antioxidative capacity due to inhibition of myeloperoxidase, the enzyme that generates the high amounts of the pro-oxidants (17). Therefore, the studies investigating the oxidative stress response after surgery in the variety of settings and with the diversity of putative modulators are still necessary.

The current knowledge about oxidative stress markers after abdominal hysterectomy is much less than those concerning other major surgeries with more profound ischemia-reperfusion injury (e.g. vascular, cardiologic surgery, tourniquet-used interventions). Abdominal hysterectomy triggers detectable changes of oxidative stress status and studies identified several factors that contributed to production of free radicals during the intervention (e.g. peritoneal closure, retention of ovaries, hormonal changes) (18-20). However, we are not aware of any study that simultaneously investigated the relationships of oxidative stress parameters and postoperative recovery as well as the effects of therapeutic interventions.

Therefore, in this study, we hypothesized that oxidative stress injury after abdominal hysterectomy deteriorated quality of life during the postoperative recovery period and that multimodal analgesia with analgesics, that have proven or proposed antioxidant activity, will give faster recovery measured with a validated rating questionnaire.

PATIENTS AND METHODS

The study was designed as interventional, time-series, non-therapeutic trial within which there was four groups of patients according to the primary analgesic drug given during the early postoperative recovery. The cohort was formed from women who were underwent abdominal hysterectomy in Clinical Center “Kragujevac”, Kragujevac, Serbia, successively, from October 2011 to February 2013, up to the number of pre-calculated total sample size. Eligible subjects met the following inclusion criteria: females, total abdominal hysterectomy due to benign disease (leiomyoma), ASA physical status I or II and the receiving one or two of the study analgesics (morphine, ketoprofen, ketorolac, paracetamol). Exclusion criteria were: previous chronic use of anti-inflammatory drugs (i.e. steroids, NSAIDs), antipsychotics and opioids, known previous hypersensitivity to study drugs and history of medici-
cally important kidney or liver disease. The study design was comparable with similar previous published studies (1, 4). Study approval was obtained from Clinical Center Kragujevac Institutional Ethics Review Board and all study participants gave the written informed consent.

Anesthesia and analgesia

All patients were premedicated with 0.07 mg/kg i.m. midazolam and during anesthesia, standard monitoring were applied (ECG, NIBP, SpO2, EtCO2 and capnography). Induction of anesthesia was achieved with fentanyl 2-3 mcg/kg i.v., propofol 1.5-2.0 mg/kg i.v. rocuronium 0.6 mg/kg i.v. Patients were ventilated with an oxygen-air mixture (FiO2=0.4) and EtCO2 stabilized at 35-40 mm Hg. Anesthesia was maintained with sevoflurane at 1-1.5 minimum alveolar concentration (MAC), fentanyl 1 mcg/kg titrated to the dose which avoided the increase arterial blood pressure values above 20% of baseline, and additional doses of rocuronium to keep satisfactory surgical relaxation. All patients received ondansetron 4 mg prior to the end of surgery. Neuromuscular blockade was antagonized with 0.05 mg/kg i.v. neostigmine and 0.01 mg/kg i.v. atropine. Patients were awakened and extubated in the operating room and transferred to the post-anesthesia care unit (PACU) upon following simple commands.

There were four analgesic drug protocols and according to that four groups: At the beginning of perioperative closure all subjects received 0.15 mcg/kg morphine (M group). Some subjects additionally received 100 mg ketoprofen (MK group) or 30 mg ketorolac (MZ group) or 1000 mg paracetamol (MP group) respectively, in i.v. infusion over 30 minutes at the beginning of surgery plus 0.075 mcg/kg at the same time point as in M group.

In the PACU, pain was assessed on regular 10-minute intervals and patients received additional 2 mg morphine boluses in order to maintain the Numeric Pain Rating Scale (NPRS) score £3. Discharge readiness from the PACU was assessed by using the modified Aldrete’s score every 15 minutes until patients met discharge criteria (score ≤9) (21). Administrations of study analgesics were continued on the ward according to the following scheme during the first 48 hours after surgery: subjects in the M subgroup received i.v. boluses of morphine 5 mg every 4 hours, in MK subgroup received ketoprofen 100 mg/8 h i.v. infusion, in MZ - ketorolac 30 mg/8 h i.v. infusion and in MP received paracetamol 1000 mg every 6 hours i.v. infusion. Additional analgesia for patients in all subgroups was performed on demand with the administration of 2 mg of i.v. boluses of morphine until achieving NPRS score of £3.

Patients’ variables and quality of postoperative recovery assessments

Perioperative data collected included subject’s age, ASA physical class, smoking habits, duration of surgery. One of the investigators not involved with patient care performed perioperative data collection. Pain intensity was measured using a 10-point Numeric Pain Rating Scale (NPRS) on 10-minute intervals in PACU and on every four hours on the ward. The postoperative recovery was the primary study outcome and it was assessed using the researcher-assisted, 40-item questionnaire which was specifically designed and validated to measure a patient’s health status after surgery and anesthesia (22). This questionnaire measures 5 dimensions of recovery: emotional state (9 items), physical comfort (12 items), psychological support (7 items), physical independence (5 items) and pain (7 items). The sum of the individual components generates an aggregate score, which we considered to be the primary study variable, ranging from 40 (the worst) to 200 (the best) points. We validated the Serbian language version internally, based on previous recommendation for clinical researchers (1).

Oxidative stress markers measurements

Venous blood samples were collected before the surgery in the preoperative room before saline infusion (T0), 3 hours (T1), 24 hours (T2), 48 hours (T3) and 72 hours (T4) after the surgery. Samples (5–10 mL) were taken in tubes. Blood samples were taken from antecubital veins into test tubes containing sodium citrate anticoagulant. Blood samples were processed and stored immediately. Blood was centrifuged to separate plasma and red blood cells (RBCs). Biochemical parameters were measured spectrophotometrically.

Index of lipid peroxidation (Thiobarbituric Acid Reactive Substances, TBARS)

The degree of lipid peroxidation in plasma was estimated by measuring thiobarbituric acid reactive substances (TBARS) using 1% thiobarbituric acid (TBA) in 0.05 M NaOH, incubated with plasma at 100°C for 15 min and read at 530 nm. Distilled water was used as a blank probe. TBA extract was obtained by combining 0.8 mL of plasma and 0.4 mL of trichloroacetic acid (TCA), and then the samples were put on ice for 10 minutes and centrifuged for 15 min at 6000 rpm (23).

Nitrite (NO2−)

Nitric oxide (NO) decomposes rapidly to form stable metabolite nitrite/nitrate products. Nitrite (NO2−) was determined as an index of nitric oxide production with Griess’s reagent: 0.1 mL 3 N perchloric acid (PCA), 0.4 mL 20 mM ethylenediaminetetraacetic acid (EDTA), and 0.2 mL plasma were put on ice for 15 min, then centrifuged for 15 min at 6000 rpm. After pouring of the supernatant, 220 μL K2CO3 was added. Nitrites were measured at 550 nm of wavelength. Distilled water was used as a blank probe (24).

Superoxide anion radical (O2−)

The level of superoxide anion radical (O2−) was measured using nitro blue tetrazolium (NBT) reaction in TRIS-buffer combined with plasma samples and read at 530 nm. Distilled water was used as a blank probe (25).
Hydrogen peroxide (H$_2$O$_2$)

The protocol for measuring hydrogen peroxide (H$_2$O$_2$) was based on oxidation of phenol red in the presence of horseradish peroxidase. Two hundred µL sample with 800 µL phenol red solution (PRS) and 10 µL horseradish peroxidase (PD) were combined (1:20). The level of H2O2 was measured at 610 nm (26).

Determination of activities of antioxidant enzymes

Hemoglobin determination, necessary for the calculation of activity of endogenous antioxidants, was performed according to the Drabkin method. Superoxide dismutase (SOD) activity was determined by the epinephrine method of Misra and Fridovich (27). A hundred µL lysate and 1 mL carbonate buffer were mixed, and then 100 µL of epinephrine were added. Detection was performed at 470 nm. Plasma levels of reduced glutathione (GSH) are determined spectrophotometrically by Buettler’s method, based on the oxidation of glutathione (GSH) by 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB) (28). Catalase (CAT) activity was determined according to Beutler. Lysates were diluted with distilled water (1:7 v/v) and treated with chloroform-ethanol (0.6 : 1 v/v) to remove hemoglobin. Then 50 µL catalase buffer, 100 µL sample, and 1 mL 10 mM H2O2 were mixed. Detection was performed at 360 nm. Distilled water was used as a blank probe (28).

Statistical analysis

Sample size calculation assumed differences between baseline and final QoR-40 total score of at least 10 points (~5%), for pairwise comparisons (matched pairs), with a=0.05 and b=0.2, based on data from similar studies (1, 29). The preliminary calculated size was increased 1.5 times (assumption of non-parametric distribution and higher variability of study data than expected), establishing the final study population of at least 45 subjects. Statistical analysis included description methods, analysis of variance (one-way and repeated measures pairwise comparisons), Friedman test, Kruskal Wallis test, Wilcoxon Signed Rank test, and correlations (Pearson, Spearman’s rho). For the purpose of secondary analysis the patients have been divided into four subgroups according to the prescribed analgesic protocol: morphine only (M), morphine plus ketoprofen (MK), morphine plus ketorolac (MZ) and morphine plus paracetamol (MP). All statistical tests were performed two-sided and the differences were considered statistically significant at the level of p£0.05.

RESULTS

Patients’ characteristics

Study population included 51 females, average age 51.6±7.8 years (the mean ± standard deviation-SD), the youngest was 39 and the oldest was 69 years old. Among them 76.5% were ASA I (n=39), 23.5% ASA II (n=12) and 43.1% were smokers (n=22). Average duration of surgery was 74.8±26.4 minutes (minimum 35 minutes and maximum 120 minutes). There were four study subgroups according to the principal analgesic drug protocol: 19 women (37.3%) received morphine (M), 14 women (27.5%) received morphine plus ketoprofen (MK), 6 women (13.7%) received morphine plus ketorolac (MZ) and 7 women (13.7%) received morphine plus paracetamol (MP). Five patients received analgesic combinations which did not fit into abovementioned, prespecified classification and their data had been not included for analgesic subgroup analysis. There were no statistically significant differences in patients characteristics between subjects from different groups (p>0.05). In addition, exclusion criteria in our study eliminated differences in distribution of variety of other factors that could affect our outcomes.

Quality of recovery

Total QoR-40 score decreased during early postoperative period and returned at preoperative values within three days (figure 1). In general, the scores during the study changed significantly (p<0.001; ANOVA-repeated measures) with significantly lower values on the POD1 and POD2 (p<0.05; ANOVA-repeated measures, pairwise comparisons).

In general, observed changes of QoR-40 total scores were similar between subgroup of patients receiving different analgesics drugs. Only on the first postoperative day the differences were statistically significant between groups (p=0.039; one-way ANOVA). The patients in MK group had a smaller drop of the scores (n=8, the mean decrease = 10.6 points, SD=6.5, min = -1, max = 18) than patients in MZ group (6, 23.2, 17.2, 7, 54) or M group (11, 25.6, 12.0, 4, 46).

Oxidative stress markers

The values of oxidative stress parameters in the whole study population were presented in the table 1. The de-
Individual variability was very high and the differences did not reach the threshold of statistical significance (table 2).

There were not statistically significant differences of reduced glutathione (GSH) values during the study (p=0.114; Fiedman test). The individual variability was high, too and notable differences particularly between T0 and T3 (almost doubling the mean values) visits did reach statistical significance (table 2).

Catalase (CAT) values across five measurements were not statistically significant (p=0.317; Fiedman test). Although the parameter, in general, decreased during the study the differences did not reach the threshold of statistical significance (table 2).

Correlation analysis of oxidative stress markers and QoR-40

There were no significant correlation between the values of parameters of oxidative stress response, except in the case of TBARS and H$_2$O$_2$. The values of changes from T0 to T3 of TBARS and the values of changes in the same period (T0 to T3) of H$_2$O$_2$ significantly and negatively correlated with each other (r=-0.303, p=0.046; Spearman’s rho). Finally, there were no significant correlation between total QoR-40 and any parameter of oxidative stress response, neither for their absolute values nor their changes between study visits.

Oxidative stress markers, analgesics and other factors

The differences of the means of TBARS changes from the baseline to T3 were statistically significant between T0 to T3 of TBARS and the values of changes in the same period (T0 to T3) of H$_2$O$_2$ significantly and negatively correlated with each other (r=-0.303, p=0.046; Spearman’s rho). Finally, there were no significant correlation between total QoR-40 and any parameter of oxidative stress response, neither for their absolute values nor their changes between study visits.

Table 1. The values of oxidative stress markers before (T0), 3 (T1), 24 (T2), 48 (T3) and 72 (T4) hours after the surgery in whole study population (n=51)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>T0</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
</tr>
</thead>
<tbody>
<tr>
<td>TBARS (µmol/ml)</td>
<td>3.23±2.26</td>
<td>3.63±2.62</td>
<td>3.74±2.43</td>
<td>3.64±2.30</td>
<td>3.49±2.17</td>
</tr>
<tr>
<td>NO$_2$ (nmol/ml)</td>
<td>10.37±7.62</td>
<td>9.58±7.88</td>
<td>10.15±7.93</td>
<td>9.38±7.56</td>
<td>10.12±7.05</td>
</tr>
<tr>
<td>H$_2$O$_2$ (nmol/ml)*</td>
<td>2.47±2.32</td>
<td>1.94±1.41</td>
<td>1.85±1.43</td>
<td>2.36±1.83</td>
<td>2.96±2.33</td>
</tr>
<tr>
<td>O$_3$ (nmol/ml)</td>
<td>6.91±4.25</td>
<td>6.54±3.83</td>
<td>8.53±5.07</td>
<td>7.48±5.27</td>
<td>6.19±4.66</td>
</tr>
<tr>
<td>SOD (U/gHgbx10$^4$)</td>
<td>584.09±1065.07</td>
<td>695.88±1298.00</td>
<td>507.21±675.36</td>
<td>570.68±692.91</td>
<td>605.77±672.69</td>
</tr>
<tr>
<td>GSH (nmol/ml)</td>
<td>5826.29±8246.42</td>
<td>6586.89±8246.42</td>
<td>6869.54±6654.74</td>
<td>8059.70±12197.24</td>
<td>7556.66±8731.99</td>
</tr>
<tr>
<td>CAT (U/gHgbx10$^4$)</td>
<td>133.45±144.76</td>
<td>103.06±87.99</td>
<td>97.27±117.51</td>
<td>115.43±123.14</td>
<td>102.15±143.04</td>
</tr>
</tbody>
</table>

*p<0.05 between study visits

Table 2. The values of oxidative stress markers before (T0), 3 (T1), 24 (T2), 48 (T3) and 72 (T4) hours after the surgery in whole study population (repeated measures analysis)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>N</th>
<th>T0</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
</tr>
</thead>
<tbody>
<tr>
<td>TBARS (µmol/ml)</td>
<td>29</td>
<td>0.13±11.21</td>
<td>0.04±10.97</td>
<td>0.71±9.70</td>
<td>9.70±9.70</td>
<td>0.26±10.24</td>
</tr>
<tr>
<td>NO$_2$ (nmol/ml)</td>
<td>29</td>
<td>1.42±23.47</td>
<td>1.46±31.44</td>
<td>1.38±26.60</td>
<td>0.04±35.20</td>
<td>1.75±23.01</td>
</tr>
<tr>
<td>H$_2$O$_2$ (nmol/ml)*</td>
<td>26</td>
<td>2.65±2.30</td>
<td>2.11±1.31</td>
<td>2.08±1.25</td>
<td>2.59±1.54</td>
<td>2.96±2.34</td>
</tr>
<tr>
<td>O$_3$ (nmol/ml)</td>
<td>29</td>
<td>6.24±3.04</td>
<td>6.52±3.77</td>
<td>7.91±4.67</td>
<td>6.56±4.67</td>
<td>6.29±4.72</td>
</tr>
<tr>
<td>SOD (U/gHgbx10$^4$)</td>
<td>31</td>
<td>642.80±1237.00</td>
<td>889.62±1522.65</td>
<td>359.47±354.45</td>
<td>626.52±791.88</td>
<td>605.77±672.69</td>
</tr>
<tr>
<td>GSH (nmol/ml)</td>
<td>31</td>
<td>5454.66±4464.90</td>
<td>6531.01±9169.65</td>
<td>6974.39±7429.45</td>
<td>9515.30±14367.34</td>
<td>7556.66±8731.99</td>
</tr>
<tr>
<td>CAT (U/gHgbx10$^4$)</td>
<td>30</td>
<td>134.95±160.69</td>
<td>84.27±76.76</td>
<td>109.14±134.52</td>
<td>117.00±141.61</td>
<td>102.15±143.04</td>
</tr>
</tbody>
</table>

*p<0.05 between study visits

Descriptive statistics included all study subjects (n=51), i.e. those with missing data at some visits.

TBARS levels values across five measurements were not statistically significant (p=0.633; Fiedman test). Although the parameter, in general, increased during the study the individual variability was high and the differences did not reach the threshold of statistical significance (table 2).

Nitrite (NO$_2$) values across five study visits were not statistically significant (p=0.633; Fiedman test). The individual variability was even higher than in other parameters (table 2).

Hydrogen peroxide (H$_2$O$_2$) values significantly decreased from baseline to T3 and then, increased at the end of the study period (table 2). Overall, the differences across the study visits were statistically significant (p=0.011; Fiedman test). Statistically significant differences of hydrogen peroxide (H$_2$O$_2$) values in patients’ serum samples have been observed between the following study visits T0 and T2 (p=0.044; Wilcoxon Signed Ranks test), T1 and T3 (p=0.040; Wilcoxon Signed Ranks test), T1 and T4 (p=0.036; Wilcoxon Signed Ranks test), T2 and T3 (p=0.003; Wilcoxon Signed Ranks test and, T2 and T4 (p=0.007; Wilcoxon Signed Ranks test).

Superoxide anion radical (O$_3$) values across five study visits were not statistically significant (p=0.293; Fiedman test). The individual variability was slight and it seems further suggesting of the stability of the production of this molecule during the study period (table 5).

There were not statistically significant differences of superoxide dismutase (SOD) values during the study (p=0.769; Fiedman test). The individual variability was rather high and notable difference particularly between T1 and T2 visit did not significantly affected overall trend of the parameter (table 2).

There were no statistically significant differences of reduced glutathione (GSH) values during the study (p=0.114; Fiedman test). The individual variability was high, too and notable difference particularly between T0 and T3 (almost doubling the mean values) visits did reached statistical significance (table 2).

Catalase (CAT) values across five measurements were not statistically significant (p=0.317; Fiedman test). Although the parameter, in general, decreased during the study the differences did not reach the threshold of statistical significance (table 2).
the analgesic study subgroups (p=0.031; Kruskal Wallis test). These alterations originated from statistically significant differences between the values of ketorolac and paracetamol subgroups (p=0.004; Mann-Whitney-Wilcoxon test) as well as between morphine and paracetamol subgroups (p=0.015; Mann-Whitney-Wilcoxon test). Overall, the largest changes was noted in paracetamol study subgroups in which there was substantial increase of TBARS activity from T0 to T3 visits and then, the sudden drop to the end of the study period (figure 2).

There were no statistically significant differences between value of other oxidative stress response markers (NO₂⁻, H₂O₂, O₂⁻, SOD, GSH, CAT) during the study period regarding different analgesics subgroups (p<0.05; Kruskal Wallis test).

Patients age, duration of surgery and cigarette smoking status showed some significant influences on and association with the oxidative stress response markers. Patients age negatively correlated with TBARS changes from T0 to T2 (r=-0.365, p=0.032; Spearman's rho) and positively with CAT changes (r=0.363, p=0.025; Spearman's rho). Duration of surgery correlated negatively with O₂⁻ changes from T0 to T1 (r=-0.388, p=0.016; Spearman's rho) and positively with CAT changes from T0 to T2 positively (r=0.327, p=0.045; Spearman's rho). There were statistically significant differences of O₂⁻ changes from T0 to T1 between the cigarette smokers and non-smokers (p=0.005; Mann-Whitney-Wilcoxon test) as well as from T0 to T2 (p=0.008; Mann-Whitney-Wilcoxon test) (figure 3).

In addition, there were statistically significant changes of SOD from T0 and T1 between cigarette smokers and non-smokers (p=0.045; Mann-Whitney-Wilcoxon test). In the smokers the median of change was -268.62 (-1554.74 and -81.40, percentile 25 and 75, respectively) and in the non-smokers the median of change was 252.34 (-32.56, 529.10)

**DISCUSSION**

The results of our study revealed that there was significant decrease of hydrogen peroxide (H₂O₂) in women after abdominal hysterectomy in all patients 24 hours after surgery. In addition, there was significant increase of index of lipid peroxidation (measured as thiobarbituric acid reactive substances - TBARS activity) during 48 hours postoperatively in the MP subgroup. Correlation analysis confirmed this type of negative relationships between these two parameters of oxidative stress response. Only a few studies published so far investigated parameters of oxidative stress response in women after hysterectomy (18-20, 30). In addition, data about H₂O₂ activity lack and, therefore, our results represent the novelty in the field.

The several enzyme cascades are involved in the synthesis and the degradation of H₂O₂ including SOD, CAT, GPx, GSH and myeloperoxidase (31). However, in our study activities of SOD and CAT were insignificantly changed which suggest that other, unmeasured, alternative pathways were involved in observed decrease of H₂O₂. It is known for example that H₂O₂ could be removed with reactions mediated with free metal ions (e.g. Fe²⁺, Cu²⁺) during which very toxic, hydroxyl radical (OH•) is formed (Fenton's reaction). Therefore, it seems that, in fact, the decrease of H₂O₂ was, in essence, rather a marker of the very pro-oxidation state then an indicator of higher activity of oxidative stress defense mechanism during early postoperative period after abdominal hysterectomy (32). Indeed, there were a plenty of proteins such as ferritin, transferrin, hemoglobin, myoglobin and other metalloproteins which could release free metal ions from the tissues during the surgical trauma, then sequestering H₂O₂ liberating (OH•) and finally making pro-oxidant state in the immediate postoperative period (32-34). Our finding about significant, negative correlation between H₂O₂ (decrease) and TBARS (increase), which is a marker of lipid peroxidation, further support that conclusion.

The effect of analgesic drugs could be additional mechanism of the decrease of H₂O₂ in our study subjects. All women received baseline morphine analgesia and the others paracetamol or a non-steroidal anti-inflammatory drug (NSAID), as well. Previous researches showed that morphine could dose-dependently decrease free radical
synthesis in some surgical patients and increase \( \text{H}_2\text{O}_2 \) degradation exploiting remarkable antioxidative activity (16, 35). It is known that NSAIDs and paracetamol could also increase synthesis of \( \text{OH}^+ \) during Fenton’s reaction which is also a mechanism of \( \text{H}_2\text{O}_2 \) sequestration, as described above (36).

The time-limited and paracetamol-dependent increase of TBARS (marker of lipid peroxidation) detected in our study could be consequences of at least two circumstances: methodological specificity of the TBARS assay and effects of analgesic drugs. The assay used for determination of TBARS is based on measurement of malonyldialdehyde (MDA) concentration. The synthesis and degradation of this metabolite is variable depending on many factors such as the types and content of polyunsaturated free fatty acid in lipid membrane and the contribution of both non-enzymatic and non-lipid pathways (37).

Morphine, during experimental oxidative stress conditions, which included the presence of adenosine diphosphate (ADP) and ferrous ion, prevented the lipid peroxidation due to inhibition of oxygen consumption and \( \text{H}_2\text{O}_2 \) generation (38). However, in other methodological settings this effect was either absent or achieved with high doses of morphine (39, 40). The use of paracetamol, on the other side, is associated with the consumption of a powerful antioxidant molecule, GSH, due to conjugation of its metabolites during the phase 2 of the drug’s biotransformation (41, 42). This fact support our results that the subgroup of women, which received paracetamol, was more susceptible for development of pro-oxidant state than the others.

In our study, the concentration of other parameters of oxidative stress (SOD, CAT, NO, \( \text{O}_2^- \)), as well as GSH did not differ in successive plasma samples, either in whole patients population or in individual ones subgroups. Although the results of other, previously published studies in the field are not fully consistent and some of them support our findings this part of our study could be interpreted bearing the mind several its limitation (43). Firstly, our study sample was not very large, being underpowered for detecting significant differences but of smaller magnitudes. Secondly, the influence of other confounders, not included in our analysis could not be excluded. Finally, it is difficult to extrapolate our results to the patients treated in other clinical setting, such as receiving other type of anesthetic and analgesics drugs. Obviously, future research with more focused designs is needed in order to provide additional valid evidences.

The effects of age, duration of surgery and cigarette smoking in our study are further evidences about the disturbances of pathways of oxidative stress response at the time of abdominal hysterectomy and during postoperative period. We noted disturbances of concentrations/activities of TBARS, \( \text{O}_2^- \), \( \text{CAT} \) and SOD pointing to activation of some pro-oxidant mechanisms and compensatory synthesis of some enzymes of antioxidative defense. All these factors are known triggers of oxidative stress response cascades but small numbers of cases in our research as well as the parameter- and time-dependent changes precluded more detailed analysis of their influence on the main study outcomes (44, 45).

The quality of recovery in women included in our study did not correlate with any of measured oxidative stress response parameter. The use of quality of recovery with QR-40 instrument had been validated in many studies with surgical patients in general anesthesia including women after abdominal hysterectomy (1). However, according to our knowledge, there were no studies, that particularly explored association between postoperative quality of recovery (e.g. QR-40 scores) and pro-oxidant reactive species or biomarkers of oxidative stress defense. In one recent research both MDA and quality of recovery score were assessed during 24 hours after Gynecologic Laparoscopic Surgery but no direct relationships between the two variables were established and reported (46). In general, studies dealing with postoperative recovery more often investigated other important biological pathways like cytokine response than oxidative stress parameters (4).

In conclusion, our findings suggest that during the hysterectomy women experienced a detectable degree of damage of cell membranes by lipid peroxidation type and that the subgroup that received paracetamol is under the highest risk. Of all the analgesics used, paracetamol has the highest potential for exhaustion of antioxidant defense mechanisms particularly in the presence of other stressors, which liberate reactive, pro-oxidant species as if it is a surgical intervention. Our study is, as we are aware, the first one, which in a comparative design, by the same method, examined the differences between different analgesics on parameters of lipid peroxidation and other indicators of oxidation stress during abdominal hysterectomy.

ACKNOWLEDGMENT

The authors thanks to Faculty of Medical Sciences of University of Kragujevac, Kragujevac, Serbia, for supporting the research with the Junior Project, JP 09-12, “Analysis of factors associated with postoperative recovery in patients after elective abdominal hysterectomy”.

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