Effect of myocardial protection and perfusion temperature on production of cytokines and nitric oxide during cardiopulmonary bypass

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ABSTRACT

Purpose: To investigate the effects of different conditions used during cardiopulmonary bypass (CPB) surgery on accompanying production of cytokine and nitric oxide (NO).

Methods: Patients undergoing CPB for the first time were prospectively enrolled and divided into two groups according to CPB parameters performed: i) normothermia (36.5-37°C) with blood cardioplegia (NB group, n=10) and ii) hypothermia (29-31°C) with crystalloid cardioplegia (HC group, n=10). Plasma samples obtained following intubation (baseline), during (5 and 30 min) and after (4 and 24 h) CPB were assayed for cytokines (ELISA) and NO metabolites (Griess reaction).

Results: Peak concentrations of interleukin (IL)-6 and IL-8 were reached at 4 h post CPB in both groups, but in the HC group those levels increased earlier and persisted for longer (24 h) compared to baseline (P<0.05). IL-10 levels also increased at 4 h compared to baseline, but only significantly so in the HC group. NO metabolites were reduced in HC group at all time points compared to baseline (P<0.05), while no significant differences were detected in the NB group.

Conclusion: The association between increased systemic levels of cytokines and reduced NO production in the HC group suggests that different myocardial protection and/or perfusion temperature used during CPB may contribute to the extent of inflammatory response.

Key words: Cardiopulmonary Bypass. Myocardium. Perfusion. Nitric Oxide. Cytokines.

RESUMO

Objetivo: Investigar a hipótese de que diferentes procedimentos durante o bypass cardiopulmonar (BCP) causa diferentes níveis de citocinas (IL) e óxido nítrico (NO). Métodos: Pacientes submetidas a BCP foram prospectivamente estudadas de acordo com bypass realizado sob normotermia (36.5-37°C) com cardioplegia sanguínea (NB grupo, n=10) ou hipotermia (29-31°C) com cardioplegia cristalóide (HC grupo, n=10). Amostras de Plasma foram obtidas após a intubação (linha de base), durante (5, 30 min) e após (4, 24 h) o BCP. Os ensaios foram realizados através de ELISA (IL) e metabólitos do NO (reação de Griess). Resultados: Os picos de concentrações de IL-6 and IL-8 estavam aumentados em 4 h pós BCP em ambos os grupos, mas no grupo HC estes níveis aumentaram precoce e persistiram aumentadas por 24 h, comparado a linha de base (P<0.05). O nível de IL-10 também teve o pico em 4 h, mas estatisticamente significante somente no grupo HC, comparado a linha de base. Os metabólitos do NO estavam reduzidos no grupo HC, em todo o tempo, comparado a linha de base (P<0.05), enquanto nenhuma diferença estatisticamente significante foi detectada no grupo NB. Conclusão: A associação entre o aumento sistêmico dos níveis de citocina e a redução da produção de NO no grupo HC sugere que o tipo de proteção miocárdica e/ou temperatura de perfusão no BCP pode ser um fator determinante na extensão da resposta inflamatória.

Introduction

Cardiac surgery with cardiopulmonary bypass (CPB) can induce systemic inflammatory response syndrome (SIRS), which is characterized by fever, increased respiratory and heart rate (or a PaCO₂ level of less than 32 mmHg), and an abnormal white blood cell count. The underlying causes of SIRS include surgical trauma, blood exposure to foreign surfaces, ischemia/reperfusion injury, mechanical shear stress, mechanical trauma, and NO in patients receiving CPB performed by two different procedures: (1) normothermia with blood cardioplegia, or (2) hypothermia with crystalloid cardioplegia.

Methods

Subjects

This study was approved by UFTM ethics committee. After giving informed consent, 20 patients undergoing elective coronary artery grafting or valve operation were prospectively enrolled. Patients were randomly divided into two groups: one that received hypothermic bypass with crystalloid cardioplegia (HC group, n=10) or normothermic bypass with blood cardioplegia (NB group, n=10). Of the ten patients in each group, 9 received revascularization and one a valve change. For all patients, the following general data were collected: age, sex, weight, height and body mass. Exclusion criteria were patients with severely impaired left ventricular function (ejection fraction <40%), previous cardiac procedures, severe systemic noncardiac disease, recent myocardial infarction (<6 wks), infectious disease before operation, or impaired lung, liver or renal function.

Anesthesia and surgical techniques

Anesthetic techniques were standardized for all patients; they received midazolam 15 mg orally 1 h before operation. Central venous and radial artery cannulas were inserted under local anesthesia (1% lignocaine) and midazolam sedation (5.0 mg intravenously). Anesthesia was induced with sufentanil (0.5 mg/kg) and etomidate (0.3 mg/kg); succinylcholine (1.5 mg/kg) was used for neuromuscular blockade. Lungs were ventilated with oxygen (1 L/min) containing isoflurane (0.5 to 1.5%), to maintain anesthesia, and sufentanil (0.5-1.0 mg/kg/h) and pancuronium bromide (0.1 mg/kg). During CPB, patients received continuous infusion of propofol (4.0 mg/kg/h) and sufentanil (0.5-1.0 mg/kg/h). Patients were ventilated with a tidal volume of 6.0 - 8.0 ml/kg, aiming at normocapnia. Perioperative antibiotic prophylaxis consisted of cefazolin (2 g every 4 h during the procedure and every 6 h for 24 h thereafter). The extracorporeal circuit consisted of a roller pump, a cardiectomy reservoir, and a membrane oxygenator. The circuit was primed with 1500 ml of Ringer’s lactate solution, sodium bicarbonate and mannitol 15% (2 ml/kg). CPB was established using ascending aortic cannulation and two-stage venous cannulation in the right atrium. Intravenous heparin was administered to achieve an activated clotting time of >480 s. After the start of CPB, patients in the HC group were cooled to 29 to 31°C, whereas patients from NB group were maintained at 36.5 to 37°C. Flow rates of 2.4 L/m²/min were used. The management of myocardial protection differed between the groups, as follows: in the HC group, myocardial protection was achieved by inducing electromechanical arrest with cold, anterograde
crystalloid cardioplegia using St. Thomas’s I solution and topical cooling using normal saline solution (4°C). One liter of cardioplegia was administered initially followed by 300 ml every 30 min of cross-clamping or earlier, whenever electrical activity was seen. In the NB group, cardiac arrest was achieved by using intermittent anterograde hyperkalemic warm blood cardioplegia. At the end of bypass, anti-coagulation was reversed with protamine sulfate in both groups. After the surgery was completed, patients were transferred to an Intensive Care Unit, where standard care was followed until discharge. Extubation was performed as soon as patients were able to maintain adequate gas exchange with normal blood gases during spontaneous breathing, when they were haemodynamically stable and had satisfactory renal function.

Intraoperative measurements and plasma samples

Intraoperative variables included operation and bypass time, aortic cross-clamp time and duration of intubation. Serial blood samples (10 ml) were withdrawn from the radial artery catheter 5 min after intubation (baseline), 5 and 60 min after the beginning of CPB, and 4 and 24 h after the end of CPB. The samples were collected in EDTA-containing tubes and centrifuged at 3000 X g for 15 min. Plasma was collected and stored (-70°C) until assays for cytokines and NO were performed.

Cytokine measurements

The concentrations of IL-6, IL-8 and IL-10 in plasma samples were determined by ELISA. Briefly, flat-bottomed 96-well microtiter plates were coated with specific antibody (100 µl/well) diluted in coating buffer (1 µg/ml for IL-6 and IL-8 or 3 µg/ml for IL-10; PharMingen, San Diego, CA) and incubated overnight at 4°C. The plates were then washed and non-specific binding was blocked (120 min, 37°C) with 1% bovine serum. Non-diluted samples and standards were loaded onto plates. Recombinant human IL-6, IL-8 and IL-10 (PharMingen, San Diego, CA) standard curves were used to calculate the cytokine concentrations. The plates were thoroughly washed and the appropriate biotinylated polyclonal or monoclonal anti-cytokine antibody was added. The plates were washed 1 h later; then avenin peroxidase (diluted 1:5000) was added to each well for 15 min and each plate was thoroughly washed again. Substrate (0.4 mg of o-phenylenediamine dihydrochloride [Sigma, St. Louis, MO] + 0.4 µl of hydrogen peroxide [Merck, Rio de Janeiro, RJ, Brazil] per 1 ml of substrate buffer) was added and the reaction was stopped with H2SO4 (1 M). The optical density was measured on a plate reader (Spectra Max 250 – Molecular device, Sunnyvale, CA) at 490 nm. The optical density in the samples was compared with standard curves and results were expressed as picograms of each cytokine per 1 ml of plasma.

Determination of NO metabolites

The nitrite (NO2) plus nitrate (NO3) concentration in samples was determined by enzymatically reducing nitrate with nitrate reductase. Briefly, non-diluted plasma samples (50 ml) were incubated with the same volume of reductase buffer (0.1 M potassium phosphate, pH 7.5, containing 1 mM NADPH, 10 mM FAD and 4 U of nitrate reductase ml−1) for 20 h at 37°C. A standard nitrate curve was obtained by incubating sodium nitrate (10 to 200 mM) with the reductase buffer. The total amount of nitrite was then determined by the colorimetric Griess method. Briefly, samples were incubated with the same volume of freshly prepared Griess reagent (1% sulphanilamide, 0.1% naphthylethenediamine dihydrochloride in 5% phosphoric acid). Absorbance at 550 nm was determined using a multi-well plate reader (Multiskan MCC/340 MKII, Flow Laboratories). The results are reported as micromoles (mM) of NO3 + NO2.

Statistical analysis

Data were analyzed with SigmaStat software. General characteristics or time points were compared between groups using the Mann-Whitney test. Repeated measures analysis of variance for paired data (Friedman’s test) was used to assess changes over time in each group. If significance was found, Dunnett’s post-doc test was used for comparisons with the basal time point. Correlations between peak levels of cytokines and NO in each group were assessed by Spearman’s rank correlation. In all cases P < 0.05 was considered statistically significant.

Results

Study population

Clinical characteristics of the study groups are summarized in Table 1. Subjects in the two groups did not differ significantly in age, sex, total operation time, bypass time, cross-clamp time or intubation time. The groups also did not differ in weight, height or body mass (data not shown). One patient (representing 10% of the population) from the HC group died.
**Cytokine assay**

The systemic production of IL-6, IL-8 and IL-10 was evaluated at different time points during (5 and 30 min, T1 and T2) and after (4 and 24 h, T3 and T4) bypass and compared to baseline levels (T0). The basal concentrations of the cytokines were below the limit of detection in most patients for IL-6 (Figure 1) and IL-10 (Figure 3), but were detected in 80% patients from both groups for IL-8 (Figure 2). Thus, basal systemic levels of IL-6 and IL-10 were below the limit of sensitivity in our assay, since low levels are not likely attributable to a problem with the assay because the cytokines were detected in some patient samples, as observed by maximum values, and in the control curve. In both the HC and NB groups, plasma IL-6 (Figure 1) and IL-8 (Figure 2) reached peak concentrations 4 h after the end of bypass. In the NB group, IL-6 and IL-8 levels were significantly elevated only at 4 h compared to baseline, and then returned to control values at 24 h later (Figures 1 and 2, panel A, respectively). In the HC group, significantly increased cytokine levels were detected at 5 and 30 min after the beginning of bypass compared to baseline, and, for IL-6 and IL-8, persisted for 24 h thereafter (Figures 1 and 2, panel B, respectively). Furthermore, peak levels of IL-6 were significantly higher in HC than NB groups, and a significant correlation ($p < 0.01$) was found between peak levels of IL-6 and IL-8. Regarding IL-10, although plasma levels peaked at 4 h in both groups, statistical significance was detected only for HC group compared to baseline (Figure 3).

**Plasma nitrate concentration**

Plasma samples, obtained at the same time as blood collection for the cytokine assay, were also tested for NO$_3$ + NO$_2$ levels. In NB group these concentrations were not altered compared to baseline, but in HC group, NO metabolites were significantly reduced at all time points evaluated compared to the basal median value (Figure 4).
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FIGURE 2 - Plasma IL-8 concentrations during and after bypass. Patients were submitted to bypass under normothermia with blood cardioplegia \((n=10, \text{ panel A})\) or hypothermia with crystalloid cardioplegia \((n=10, \text{ panel B})\). Time points for the assay were: baseline (after intubation, T0), 5 and 30 min after bypass start (T1 and T2), and 4 and 24 h after the end of bypass (T3 and T4). The 25\textsuperscript{th} and 75\textsuperscript{th} percentiles are represented by a bar centered about the median; ranges are depicted by the error bars. *\(P < 0.05\) compared to baseline (Friedman test followed by Dunnett’s test)

FIGURE 3 - Plasma IL-10 concentrations during and after bypass. Patients were submitted to bypass under normothermia with blood cardioplegia \((n=10, \text{ panel A})\) or hypothermia with crystalloid cardioplegia \((n=10, \text{ panel B})\). Time points for the assay were: baseline (after intubation, T0), 5 and 30 min after bypass start (T1 and T2), and 4 and 24 h after the end of bypass (T3 and T4). The 25\textsuperscript{th} and 75\textsuperscript{th} percentiles are represented by a bar centered about the median; ranges are depicted by the error bars. *\(P < 0.05\) compared to baseline (Friedman test followed by Dunnett’s test)
Discussion

This study investigated the effect of two conditions during CPB, namely perfusion temperature and cardioplegia type, on the systemic production of cytokines and NO. Patients undergoing CPB under either of these conditions showed elevated levels of each cytokine studied, however the time course of these increases differed. Cytokine levels were elevated earlier and longer in patients with hypothermic bypass with crystalloid cardioplegia. Furthermore, NO production was reduced under these latter conditions. Thus, perfusion temperature and cardioplegia type significantly affected the systemic inflammatory response. Our results show that cardiac surgery with CPB increased the systemic production of cytokines and that pro- and anti-inflammatory systems can be activated simultaneously. These results reinforce previous reports. We also showed that peak levels of all cytokines occurred 4 h after the end of bypass, again concordant with other studies. It is important to note that the groups in our study presented similar general characteristics, intraoperative parameters, and mortality rates. Increased systemic levels of IL-6 and IL-8 in CPB patients have been associated with myocardial reperfusion injury and inferior clinical outcome in adults. They have also been correlated with impaired cardiovascular and respiratory function in children. Moreover, transcardiac IL-6 and IL-8 are negatively correlated to cardiac index suggesting that reducing cardiac inflammatory reaction improves post-ischemic cardiac function. Importantly we found that systemic production of pro-inflammatory cytokines increased regardless of the perfusion temperature or cardioplegia type, but the time course of these elevations differed between groups. Patients with hypothermic perfusion and crystalloid cardioplegia had elevated plasma concentrations of IL-6 and IL-8 that were detected earlier and persisted longer, up to 24 h after CPB, than those with normothermic perfusion and with blood cardioplegia. In the former patients, peak levels of IL-6 correlated with IL-8 suggesting that, in addition to a temporal association, production of one cytokine could have triggered manufacture of the other to mediate the systemic inflammatory response. IL-10 production was only significantly increased in patients with hypothermic perfusion and crystalloid cardioplegia. This increase most likely indicates a counter-regulatory mechanism for the intense pro-inflammatory response observed in these patients. The effect of perfusion temperature and cardioplegia type during bypass on the development of SIRS has been investigated previously, but results are unclear. Some reports did not find differences in serum cytokine levels between patients with normothermic and hypothermic bypass, while others found a more rapid decline in IL-8 plasma levels in the former. The use of blood cardioplegia, compared to crystalloid cardioplegia, has been mainly associated with increased myocardial protection. In the former, reduced IL-6 levels were observed during and after bypass with superior cardiac index and lower expression of TGF-b associated with decreased endothelial damage. Although increased levels of inflammatory cytokines were observed, which could induce iNOS activity, the systemic production of NO was reduced in HC patients compared to baseline. These results seem to be contradictory, however, increased concentrations of IL-10, which inhibits iNOS, could have been responsible, at least in part, for the reduced levels of NO. Reduced NO production during bypass has been reported previously. In patients submitted to CPB under
hypothermia, respiratory changes indicative of pulmonary dysfunction coincided with decreased exhaled NO\textsuperscript{15}. Another study in patients that received bypass with different degrees of hypothermia showed reduced plasma NO levels until the first post-operative day followed by recovery\textsuperscript{16}. This result is in accord with our findings in that reduced NO concentrations were found in HC patients until 24 h after the end of CPB. Treatment of patients with inhaled NO at low concentrations during and after bypass\textsuperscript{17}, or with NO donor during reperfusion\textsuperscript{13}, reduced myocardial injury\textsuperscript{17} and the cardiac inflammatory reaction, as assessed by IL-6, IL-8 and TNF-a levels\textsuperscript{13}. In line with this report, our study showed an association between systemically reduced NO production and increased duration of elevated cytokine levels in patients with hypothermia and crystalloid cardioplegia, although no significant differences were detected in clinical outcome between these patients and those with normothermia and blood cardioplegia. Blood cardioplegic solution supplemented with L-arginine, the substrate for NO synthesis, was associated with reduced release of biochemical markers of myocardial damage, suggesting improved myocardial protection\textsuperscript{18}. A limitation of the present study is the small number of patients. These low numbers do not allow an accurate assessment of clinical outcome. Also, the study design precludes evaluation of the individual role of perfusion temperature or cardioplegia type. In addition, considering that the definition of hypothermia and normothermia has been heterogeneous between authors\textsuperscript{14}, and that additives in cardioplegic solutions would provide added protection, this could hinder an accurate comparison of results between various groups.

Conclusions

These results describe an association between reduced systemic production of NO and increased IL-6, IL-8 and IL-10 levels in CPB patients with hypothermia and crystalloid cardioplegia. Our findings suggest that the temperature perfusion and/or cardioplegia type are determining factors for the intensity of the systemic inflammatory response. Further studies with a larger patient population are needed to determine whether these factors affect the clinical outcome.

References


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