Introduction

Myocardial protection during cardiosurgical procedures is one of the leading problems in cardiac surgery. A lot of studies concerning this topic were published, but the optimal method wasn’t found (1,2,3). It wasn’t also unambiguously solved, which of the two main types of cardioplegia, that means blood or crystalloid cardioplegia, shows better myocardial protection. There were published a lot of studies showing better myocardial protection with the use of blood cardioplegia (4 – 12), on the other hand some authors didn’t confirm this difference (13,14). It is also difficult to compare the results of different studies because of the fact, that they are used different protocols of cardioplegia.

This study was done to compare the protective effect of blood and crystalloid cardioplegia in patients with left ventricular dysfunction undergoing coronary artery bypass grafting (CABG). Sixty consecutive patients with left ventricular ejection fraction < 35 % scheduled for CABG with the use of cardiopulmonary bypass without additional procedures were randomly divided into two groups. In the first group we used cold blood cardioplegia, in the second group cold crystalloid cardioplegia, both delivered only ortogradly. We measured hemodynamic data in early hours after operation, enzyme release and we collected other clinical data which could be influenced by perioperative myocardial protection. There was no death in either group. We also didn’t find any significant difference in incidence of perioperative myocardial infarction, arrhythmias and use of intraaortic balloon pumping between both groups. In an early hours after operation in the group with blood cardioplegia we found significantly better hemodynamic data (LVSWI, RVSWI) and significantly lower enzyme release. We conclude, that cold blood cardioplegia shows superior perioperative myocardial protection resulting in earlier restoration of myocardial function. This difference could be important in patients with high degree of left ventricular dysfunction.

Material and Methods

From January 1998 to July 1999, sixty consecutive patients with left ventricular ejection fraction < 35 % scheduled for coronary artery bypass grafting without additional procedures were randomly divided into two groups. Crystalloid cardioplegia was used in the first group and blood cardioplegia in the second one.

Blood cardioplegia was administered with the cardioplegia delivery set at a ratio of 4:1. After institution of cardiopulmonary bypass and aortic cross clamp 1000 ml of high-potassium (25 mEq/l) cold blood cardioplegia was delivered into the aortic root. After completing of the first peripheral anastomose low-potassium (16 mEq/l) cold blood cardioplegia was continuously infused through the grafts. Another 500 ml of cold low-potassium blood cardioplegia was delivered into the aortic root every 20 minutes. Neither "hot shot" nor substrate enriched cardioplegia was used. We have been using this protocol since 1995 in our institution.

In the crystalloid cardioplegia group the protocol was as followed. After institution of cardiopulmonary bypass and aortic cross clamp 1000 ml of cold crystalloid cardioplegia (St. Thomas Infusia) was delivered into the aortic root. After completing of the first peripheral anastomose low-potassium (16 mEq/l) cold blood cardioplegia was continuously infused through the grafts. The composition of the cardioplegic solutions was published in previous study (12).

All patients in this study were operated under general anaesthesia by the use of cardiopulmonary bypass. In both
groups moderate systemic hypothermia (32 °C) and topical hypothermia were used. Internal mammary artery bypasses were implanted as last anastomoses. The surgical technique and staff involved were basically the same during the period of this study.

The hemodynamic variables were measured by the thermal dilution technique with a Swan-Ganz catheter before institution of cardiopulmonary bypass and then at 1, 2, 4 and 6 hours after terminating of cardiopulmonary bypass. The last measurement was performed the first postoperative day in the morning. All the measurements were acquired when the patient’s conditions were hemodynamically stable.

Blood samples for determination of the serum levels of total creatine kinase (CK), and catalytic activity of its MB isoenzyme (CK – MB) were taken 1, 5, 7 and 11 hours after terminating of cardiopulmonary bypass and then the first and second postoperative day in the morning.

Recorded operative and postoperative clinical variables included mortality rate, perioperative myocardial infarction rate, use of intraaortic balloon counterpulsation, arrhythmias in postoperative period, defibrillation and cardioversion after unclamping the ascending aorta etc.

Comparison between groups were made with Student’s t-test when appropriate, were used to test frequencies. Values of P < 0.05 were considered statistically significant.

**Results**

There were no significant difference between the groups in the clinical profile (see Tab. 1). The ejection fraction in crystalloid and blood cardioplegia group was 28.3% and 27.3%, respectively. The clinical results are summarised in Tab. 2. There was no death in both groups and we didn’t have to use intraaortic balloon counterpulsation or left ventricular assist device in either group. The perioperative myocardial infarction rate was higher in a crystalloid group, but the difference wasn’t statistically significant. Also the plasmatic level of potassium before leaving cardiopulmonary bypass as well as total blood loss after operation and length of hospitalisation after operation didn’t show statistically significant differences between groups. There was less incidence of the attacks of the atrial fibrillations in postoperative course in the crystalloid cardioplegia group, but this difference wasn’t statistically significant. There was a rather high percentage of patients requiring defibrillation before leaving cardiopulmonary bypass in both groups, but in the crystalloid cardioplegia it was significantly higher.

Important differences between groups were found in enzymatic response in postoperative period. Some of the data included mortality rate, perioperative myocardial infarction rate, use of intraaortic balloon counterpulsation, arrhythmias in postoperative period, defibrillation and cardioversion after unclamping the ascending aorta etc.

Comparison between groups were made with Student’s t-test when appropriate, were used to test frequencies. Values of P < 0.05 were considered statistically significant.

Where applicable, data are shown as the mean ± the standard deviation.

**Tab 1:** Clinical characteristics of patients in crystalloid and blood cardioplegia group.

|                      | Crystalloid | Blood   | p Value |
|----------------------|-------------|---------|---------|
| No. of patients      | 30          | 30      | NS      |
| Age (years)          | 63 ± 9      | 67 ± 9  | NS      |
| Sex (Men / Women)    | 22/8        | 24/8    | NS      |
| BSA (m2)             | 1.87 ± 0.19 | 1.84 ± 0.15 | NS |
| LVEF (%)             | 28.3 ± 3.4  | 27.3 ± 5.1 | NS |
| No. of grafts        | 3.7 ± 0.8   | 3.4 ± 0.7 | NS |
| CPB time (min)       | 71 ± 18     | 68 ± 18  | NS      |
| Aortic cross-clamp   | 37 ± 13     | 35 ± 11  | NS      |

**Tab 2:** Clinical results.

|                      | Crystalloid | Blood   | p Value |
|----------------------|-------------|---------|---------|
| No. of patients      | 30          | 30      | NS      |
| Death (within 30 days)| 0           | 0       | NS      |
| LVAD                 | 0           | 0       | NS      |
| IABP                 | 0           | 0       | NS      |
| Perioperative infarction | 2 (6.7%)  | 1 (3.3%) | NS      |
| Spontaneous return of SR | 3 (10%)    | 11 (37%) | p<0.05  |
| Serum K⁺ before leaving CPB (mmol/l) | 4,6 ± 0.6 | 4,8 ± 0.7 | NS      |
| Blood loss (ml)      | 747 ± 221   | 737 ± 181 | NS      |
| AF in postoperative period | 8 (27%)   | 11 (37%) | NS      |
| Postoperative stay (days) | 8 ± 2      | 9 ± 2   | NS      |

Where applicable, data are shown as the mean ± the standard deviation.

LVAD = left ventricular assist device
IABP = intraaortic baloon pump
SR = sinus rhythm
CPB = cardiopulmonary bypass
AF = atrial fibrillation
NS = not significant (p > 0.05)
Discussion

This study was the first part of the study supported by the grant of IGA MH CR No. NA 5583 - 3. We suppose that presented groups are big enough to publish the results, but they are not big enough to draw common conclusions on the basis of them. Most of the investigators dealing with this topic conclude, that the difference between blood and crystalloid cardioplegia could be found only in patients with high degree of left ventricular dysfunction or in patients with metabolically compromised myocardium. To form rather compact group of such patients we included only patients with left ventricular ejection fraction < 35 % scheduled for elective coronary artery bypass grafting without additional procedures.

The difference in enzymatic response after operation was find also by others (5,9,15). The hemodynamic data measurement wasn’t performed in early hours after operations in most of the published studies and as we have showed, there is no difference between groups later after operation. The exceptions are for example studies published by Iverson (9), Jin (6) and some other, who also found better hemodynamic data in early hours after operation in blood cardioplegia group.

Conclusions

We can summarise that we didn’t find significant differences in mortality, perioperative myocardial infarction rate and need of mechanical heart support in postoperative pe-
period between blood and crystalloid cardioplegia groups. The cardiospecific enzyme response was significantly lower in the blood cardioplegia group and the LVSWI and RVS-WI was significantly higher in the blood cardioplegia group one hour after operation.

We conclude, that blood cardioplegia shows earlier improvement of myocardial function after the operation. Blood cardioplegia could be beneficial in patients with severe left ventricular dysfunction, because in these patients the left ventricular dysfunction in early hours after operation can reach the critical point leading to collapse of the circulation.

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