

MIX BLOOD CARDIOPLEGIA (MBC) IN REDO PATIENTS UNDERGOING MITRAL VALVE REPLACEMENT SURGERY

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Abstract

Aim of Study: Redo patient undergoing valvular surgery are always a challenge for both cardiac surgeon and anaesthesiologists. Mix blood cardioplegia has had a profound impact on cardiac surgery, but there have been few studies on its use in mitral valve replacement, especially in redo patient. The purpose of this study was to determine whether mix blood cardioplegia offers any advantages in redo patient undergoing mitral valve replacement.

Patients and Methods: 20 patients, who have had a previous mitral valve surgery and were scheduled for mitral valve replacement with or without tricuspid repair, were randomized retrospectively to one of two groups of 10 with different technique of myocardial protection: group A (10 patients) had cold cristaloid cardioplegia, and group B (10 patients) had mix blood cardioplegia, a technique modified on our clinic condition (made by mixing 400-500ml oxygenated blood from oxygenator and 10ml KCL 7.5%). Systemic hypothermia was 28°C in Group A and between 32°C and 33°C in Group B. The results were primarily assessed on the basis of clinical outcome, such as hematocrit level intra and post CPB, maximum dose of inotropic support, spontaneous rhythm recovery after aortic crossclamping, length of intensive care unit stay and secondly on

postoperative blood loss and blood requirements.

Results: There were no preoperative or operative differences between the groups with regard to age, sex, diagnosis, rhythm, New York Heart Association functional class, left ventricular ejection fraction, estimated pulmonary artery systolic pressure, operation, or duration of the operation, CPB, or aortic crossclamping and the time when the patients underwent the first operation. There was one death in group A (10% mortality). The changes in hematocrit level differs significantly between the two groups on the post CPB period ($p=0.02$). There appeared to be a trend towards better spontaneous recovery of sinus rhythm after removal of the aortic crossclamp in group B compared with group A, the difference did reach statistical significance ($p=0.002$). Patients on group A required more inotropic support than Group B ($p=0.005$). There were differences even on blood requirements postoperatively, more dominant these requirement were in group A ($p=0.02$).

Discussion: Mix Blood cardioplegia had beneficial effects in clinical outcome in redo patients undergoing mitral valve replacement surgery. This may be due to its better preservation of high-energy phosphates and endogenous amino acids, less anerobic metabolic activity on reperfusion, reduced

release of cardiac troponin T, and improved post-ischemic functional recovery.

Introduction

Miocardial protection during cardiac surgery is the key point in this kind of intervention. The different techniques of myocardial protection has been evolved in the course of the years, making cardiac surgery more safe. This study is part of several years follow-up in our clinic of cardiac surgery. From many years we had made efforts to change the cardioplegia solution from cristaloid into blood, based on western clinics guidelines and modified on the conditions of our clinic. From 5 years we use sucessfully warm blood cardioplegia and the results are very good.

Blood cardioplegia has had a profound impact on cardiac surgery, especially in coronary artery bypass surgery, but there have been few studies on its use in mitral valve replacement, especially in redo patient. Warm blood cardioplegia is used to modify reperfusion injury, resulting in improved postoperative contractile function and decreased mortality.

The purpose of this study was to determine whether mix warm blood cardioplegia offers any advantages in redo patients undergoing mitral valve replacement. We designed a study to compare 2 techniques of myocardial protection: intermittent cristaloid cardioplegia, and intermittent mix warm blood cardioplegia. The results were primarily assessed on the basis of clinical outcome, and postoperative blood loss and blood requirements.

Patients and Methods

Between January 2008 and September 2012, in Cardiac Surgery Clinic in "Mother Theresa" Hospital Center, Tirana, Albania, 20

patients, who have had previous mitral valve surgery, were scheduled for MVR with or without tricuspid repair. The patients were randomized retrospectively to one of two groups of 10 with different technique of myocardial protection. In Group A (10 patients) intermittent antegrade cold cristaloid cardioplegia (4 grade C) was used. In group B, (10 patients) mix warm blood cardioplegia (MWBC) was used. Anesthesia technique were the same in both groups, general anesthesia.

All operations were performed using cardiopulmonary bypass (CPB) with ascending aortic and bicaval cannulation. Systemic hypothermia was 28°C in Group A and between 32°C and 33°C in Group B.

Antegrade cold cristaloid cardioplegia was injected immediately after aortic crossclamping at 15-20 mL·kg⁻¹, and than at 30 minutes interval. In Group B the technique MWBC was realized with our modifications on the condition of our clinic, the first dose was cold cristaloid (15-120 ml/kg/weight), after 30 minutes a second dose of cardioplegia was made by mixing 400-500 ml oxygenated blood from oxygenator and 10 ml KCL 7.5%, and a third dose of the same solution after 20 minutes. The MWBC temperature was gradually increased from 33°C to 36°C by the end of the infusion.

Electrical defibrillation was applied if ventricular fibrillation persisted beyond 2 min after aortic declamping, and a temporary pacemaker was used if there was no spontaneous rhythm or if the patient's heart rate was less than 50 beats·min⁻¹. After the operation, if systolic blood pressure was lower than 90 mm Hg and urine output less than 1 mL·kg⁻¹·h⁻¹ with central venous pressure between 10 and 12 mm Hg, inotropic support was started. Our first choice of inotropic agent was adrenaline.

Table nr.1. Preoperative and Operative Data in Patients Undergoing Mitral Valve Replacement

Variable	Group A(n=10)	Group B (n=10)	P-Value
Age (years)	43+/-10	44+/-10	0.32
Sex (male/female)	7/3	8/2	
Diagnosis			
MS	5	7	
MR			
Mixed MS/MR	2	1	
MR + TR	3	2	
Rhythm			
Sinus rhythm	3	5	
Atrial fibrillation	7	5	
NYHA class			
II			
III	9	8	
IV	1	2	
PASP (mm Hg)	60+/-23	63+/-19	0.62
Ejection fraction (%)	50+/-7	52+/-5	0.15
Operation			
MVR	7	8	
MVR + tricuspid repair	3	2	
CPB time (min)	65+/-5	60+/-5	0.45
Aortic crossclamp time (min)	55+/-5	0.44	
50+/-5			
Temperature during CPB © 28°C	32-33°C		

CPB=cardiopulmonary bypass, NYHA=New York Heart Association, PASP=pulmonary artery systolic pressure, MR=mitral regurgitation, MS=mitral stenosis, MVR=mitral valve replacement, TR=tricuspid regurgitation.

Intraoperative and postoperative variables were used to assess primary clinical outcomes including hematocrit levels pre CPB, intra and post CPB period, spontaneous rhythm recovery after aortic declamping (no

requirement for electrical defibrillation or temporary pacemaker), maximum doses of inotropes, duration of inotropic support, length of intensive care unit stay, postoperative blood loss and blood requirements.

Commutative data are expressed as the mean \pm standard deviation of the mean. Analysis of the difference in clinical outcome between the two groups was performed using Student's t test.

The significance level for differences in all tests was $p < 0.05$. Univariate linear regression analysis was used to perform correlation analysis.

Table nr.2. Comparison of Clinical Outcomes

Clinical Outcome	Group A (n=10)	Group B (n=10)	p-Value
Htc/Pre CPB (%)	37.31	37.47	
Htc/in CPB/post first dosis (%)	23.9	24.3	0.1
Htc/in CPB/post second dosis (%)	21.9	24.2	0.3
Htc/post CPB (%)	26.4	31.8	0.02
Spontaneous rhythm recovery (n)	1	9	0.002
Ventricular fibrillation after aortic declamp (n)	9	1	0.002
Reperfusion time (min)	25 \pm 5	15 \pm 5	0.007
Inotropic support (adrenaline mcg/kg)	0.08 - 0.12	0.04 - 0.08	0.005
Perioperative death	1	0	10% (mortality)
Intensive care stay (days)	2.9	2.1	0.3
Blood loss (ml)	600 \pm 100	400 \pm 100	0.06
Blood transfusion	2.5	1.5	0.02
Survival		9	10

Results

There were no preoperative or operative differences between the groups with regard to age, sex, diagnosis, rhythm, New York Heart Association functional class, left ventricular ejection fraction, estimated pulmonary artery systolic pressure, operation, or duration of the operation, CPB, or aortic crossclamping the time when the patients underwent the first operation (Table nr.1). There was one death in group A (10% mortality). Clinical outcomes are shown in Table nr.2.

The changes in hematocrit level differs significantly between the two groups on the post CPB period ($p=0.02$). There appeared to be a trend towards better spontaneous recovery of sinus rhythm after removal of the aortic crossclamp in group B compared with group A, the difference did reach statistical

significance ($p=0.002$). According to our protocol for postoperative care, patients in group A required more inotropic support than Group B ($p=0.005$). There were differences even on blood requirements postoperatively, more dominant these requirements were in group A ($p=0.02$). Postoperative echocardiography showed good left ventricular ejection fractions in both groups. There were no differences between the 2 groups in duration of ICU stays.

Discussion

The basic concept of MWBC is reduction of myocardial energy demand by maintaining cardiac arrest with hyperkalemic warm blood perfusate during initial reperfusion, to restore high-energy phosphates and enhance cellular repair after ischemic cardiac arrest.

The beneficial effects of MWBC have been studied both experimentally and clinically.

They include better preservation of high-energy phosphates and endogenous amino acids, less anaerobic metabolic activity on reperfusion, reduced release of cardiac troponin T, and improved post-ischemic functional recovery. These effects facilitate coronary vasodilatation and accelerate early myocardial tissue oxygen saturation during warm reperfusion.

Tenpaku and colleagues demonstrated complete microtubule repolymerization after 10 min of reperfusion with warm blood. This mechanism may be responsible for the early and improved recovery of cardiac function associated with MWBC.

Most clinical studies of MWBC have been undertaken on coronary artery bypass surgery, with more recent investigations on MWBC in congenital heart surgery.

Most results have showed definite advantages in the use of MWBC, although there have been some that showed no beneficial effect of MWBC.

There is some evidence that a glutamate-aspartate supplement to the MWBC (substrate-enriched cardioplegia) may reduce reperfusion injury and improve both metabolic and myocardial function recovery. Glutamate and aspartate are not available at our institute, so our MWBC was not a substrate-enriched solution, and it is possible that the MWBC benefits would be more obvious if this technique were employed.

Improved spontaneous rhythm recovery has also been observed with the use of MWBC. There appeared to be a trend towards a better cardiac rhythm recovery in the MWBC group (less requirement of electrical defibrillation and temporary pacemakers), the difference did reach statistical significance when compared with patients in the non-MWBC group.

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