Case report

Thromboelastography as a guide for transfusion therapy in a patient with Turner’s syndrome, hypoplasia of the aortic arch and aortic coarctation, undergoing aortoplasty with «sliding technique»: Case report

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A B S T R A C T

One of the most frequent lesions in congenital heart disease is hypoplasia of the aortic arch and the co-existence of aortic coarctation in older children, which is very difficult to treat. A new technique for treating this condition was recently described and it requires extensive suture lines and an accurate management of coagulation disorders associated with the use of extracorporeal circulation. We present a case of an 8 years and 9 months old girl with Turner’s syndrome, aortic coarctation and aortic arch hypoplasia who was admitted for sliding arch aortoplasty and received thromboelastography guided transfusion therapy.

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Introduction

Aortic arch hypoplasia (AAH) and coarctation of the aorta (CoA) are frequently associated with other congenital pathologies, including Turner’s syndrome. This syndrome – also known as monosomy X – is the most frequent genetic disorder in women, with one case per every 2000–2500 live births.\textsuperscript{1} This is the only life-compatible monosomy and it is characterized by small size, premature ovarian failure and congenital cardiovascular defects in a female phenotype patient. The most usual cardiovascular disorders are coarctation of the aorta, also associated with AAH, bivale aorta and mitral stenosis.\textsuperscript{2}

Patients with Turner’s syndrome exhibit a higher incidence of aortic dissection than patients with non-syndromic coarctation of the aorta, in addition to having a higher morbidity–mortality risk related to postoperative bleeding and perioperative rupture of the aorta.\textsuperscript{3–5}

The surgical treatment of patients with concomitant CoA and HHA is Aortic Advancement; nevertheless, older children are not illegible for this procedure due to the decreased age-related elasticity of the aorta.\textsuperscript{6} Other surgical interventions to correct these defects (patch-plasty and graft interposition) have limitations regarding the child’s growth because the prosthetic materials do not experience progressive growth, hence result in recurrent obstructions that can be linked to a high morbidity (enteric and bronchial fistulae, pseudoaneurysms, endocarditis, etc.).\textsuperscript{6,7}

The so-called “sliding technique” of the ascending aorta was recently described as an alternative for managing these patients, on the grounds of a potential circumferential growth as the primary benefit, preserving the normal contour and elasticity of the aortic tissue, as well as other advantages regarding the non-utilization of prosthetic material. However, this technique involves very long suture lines and thus a considerable risk of trans-operative bleeding.\textsuperscript{6}

Some of the required trans-operative surveillance factors required in highly complex procedures include the usefulness of thromboelastography to guide the transfusion therapy in heart surgery, with a subsequent reduction in the transfusion requirements and the total transfused volume, mainly during the postoperative period, probably reflecting the early correction of coagulation disorders.\textsuperscript{8} The advantages of thromboelastography include the possibility to dynamically identify hemostatic disorders, even prior to reverting the required anti-coagulation for extracorporeal circulation, allowing for early harvesting of blood by-products.

Case presentation

This is a female patient, 8 years and 9 months old, with a history of coarctation of the aorta and hypoplasia of the Aortic Arch diagnosed at 3 years of age. The patient was admitted for a sliding aortic arch aortoplasty (“sliding technique”). She was diagnosed Turner’s syndrome (45\textsuperscript{x}0) and was being followed by Endocrinology and Medical Genetics. At the time of admission, she was not receiving any medication, was never hospitalized previously and exhibited a normal psychomotor development, with no other relevant history.

The pre-anesthesia evaluation reported that patient did not experience any recent respiratory infections, her dental hygiene was good; her weight was 21.5 kg and she was 114 cm tall. Vital signs were within the normal range for her age: BP (blood pressure measured in the right arm) 94/60 mm Hg, BP (lower right limb) 82/43 mm Hg, HR (heart rate) 80 beats/min, RR (respiratory rate) 21 breaths/min.

No difficult airway predictors were identified during the physical examination and the cardiac auscultation presented enhanced second heart sound and a systolic murmur grade II/IV in crescendo – decrescendo of higher intensity at the left parasternal level, and no fremitus. The femoral and peripheral pulses in the lower limbs were decreased and no focal neurological deficit was identified.

The paraclinical, pre-surgical exams showed an Hb (hemoglobin) 13 mg/dl, HCT (hematocrit) 41%, white blood cells count 9100 with normal differential count; platelets 218,000, PT 12 s, PTT 30 s, INR 1.0. The chest X-ray showed no cardiomegaly or pulmonary infiltrates. Electrocardiogram showed sinus rhythm, left axis deviation, heart rate 106 beats/min and signs of left ventricular hypertrophy. The Echocardiogram showed situs solitus, left aortic arch.
temperature was 18°C. The perfusion period of time of 1 h and 8 min; the minimum temperature while the ischemia lasted for 1 h and 22 min, with a selective cooling and maintenance and alfa-stat during rewarming. A pH-stat strategy was used for gas control during rewarming. The ascending sliding arch aortoplasty was performed. A NIRS showing a 65% suppression. The physical condition classification was ASA 3 and the RACHS-1 surgical risk was category 4.

The patient was brought to the OR, basic monitoring was performed and the patent venous access in the upper left limb was confirmed. The patient was pre-medicated with midazolam 2 mg IV. IV induction was performed with fentanyl 150 mcg, propofol 50 mg and cisatracurium 6 mg followed with the administration of a dose of cephalotin and ε-aminocaproic acid (75 mg/kg/dose); successful orotracheal intubation was then accomplished and adequate positioning was verified with auscultation, capnography and radiographic control.

Invasive monitoring was possible through the placement of a three lumens (5.5F) central venous catheter in the right internal jugular artery and the right radial artery was catheterized (22G). Additionally, brain oximetry was performed (NIRS: near infrared spectroscopy) showing a 65% baseline. Maintenance was achieved with a 10 mcg/kg/h infusion of sevoflurane 1 CAM and fentanyl, in addition to an infusion of ε-aminocaproic acid (75 mg/kg/h). The innominate artery was cannulated for selective cerebral perfusion, before the start of perfusion.

The cardiopulmonary bypass was initiated with no events reported, maintaining perfusion pressures between 30 and 43 mm Hg. The heart was protected with an antegrade crystalloid cardioplegia with appropriate electrical activity suppression. The ascending sliding arch aortoplasty was performed. A pH-stat strategy was used for gas control during cooling and maintenance and alfa-stat during rewarming.

The cardiopulmonary bypass lasted for 2 h and 29 min, while the ischemia lasted for 1 h and 22 min, with a selective perfusion period of time of 1 h and 8 min; the minimum temperature was 18°C. The levels of brain oximetry (NIRS) were kept at 70–95%. Milrinone was started with a loading dose of 50 mcg/kg in 30 min and the infusion was continued between 0.5 and 0.75 mcg/kg/min.

The thromboelastography protocol included an initial baseline sample, then a sample for thromboelastography (TEG) with heparinase during rewarming and another heparinase-free sample following the correction, based on the previous sample, in order to ascertain adequate reversal of the coagulation disorders.

Fig. 1 – The thromboelastography protocol included an initial baseline sample, then a sample for thromboelastography (TEG) with heparinase during rewarming and another heparinase-free sample following the correction, based on the previous sample, in order to ascertain adequate reversal of the coagulation disorders.
started on dexmedetomidin 0.5–0.7 mcg/kg/h, no loading dose. The patient was transferred with assisted ventilation for programmed extubation, and her vital signs were BP 82/49 mm Hg, MBP 58 mm Hg, HR 132 bpm, RR 23 rpm, SPO2 100%, NIRS 73%. The intraoperative bleeding was estimated at 600 ml.

Nitroglycerin requirements persisted during the postoperative period for controlling high blood pressure and the patient developed hypophosphatemia that required corrective action. There was no need for transfusion of blood byproducts in the postoperative period. Orotracheal extubation was successful on the second day post-op and the patient was discharged from the cardiovascular ICU upon weaning from the inodilator (Milrinone) and adequate blood pressure control with Captopril and Amlodipine. The patient was discharged from hospital on the sixth day after surgery.

Discussion

This case illustrates the anesthetic management of a Turner’s syndrome patient, with coarctation and hypoplasia of the aortic arch of the aorta, for a surgical intervention as recently described in the literature, associated with extensive vascular suture lines and with the use of profound hypothermia, that leads to significant disorders in hemostasis. Thromboelastography was used to guide the transfusion therapy with evidence of rapid correction of the coagulation disorders following the transfusion and heparinase to neutralize heparin in anticoagulated patients.10–12 It has also been observed that the TEG variables that are more closely related to bleeding following extracorporeal circulation are the alpha angle (α) and the maximum amplitude (MA).13,14

In addition to the inclusion of kaolin to promote clot formation and heparinase to neutralize heparin in anticoagulated patients, some changes have been introduced to enable a quick rendering of TEG curves under critical circumstances. Examples of these changes are the addition of Celite and tissue factor for faster results and hence rapid administration of treatment to the patient.15 The use of each activator may result in specific alterations in the TGE values. For this reason, there have been some studies in pediatric populations to assess these changes. These studies have indicated that the kaolin-activated TEG reference values do not differ significantly in children between 1 month and 16 years of age and healthy adults.16 The main changes seen following tissue factor activation in patients less than 2 years old are shortening of the R & K times and increased alpha angle (α) and maximum amplitude, enabling an even faster interpretation of the results since the alpha angle value can be obtained in 4–6 min.17

**Fig. 2** – The second TGE sample following the transfusion showed values within the normal limits.
In this particular case, despite the use of an institutional algorithm to correct any TGE changes, a decision was made to transfuse cryoprecipitates because of the persistent layered bleeding even after the appropriate corrections were made; the TGE curve was reassessed and the presence of hypofibrinogenemia was suspected due to the prolongation of time k and the reduced maximum amplitude, despite a normal alpha angle. Then, the second thromboelastography showed a normal curve and thus, no additional blood products were administered, except for packed red blood cells.

It must be said, however, that TEG is not widely available as are the standard laboratory tests and this is a limitation for its routine use as a guide for transfusion protocols. Nevertheless, research in this area will help in assessing the impact of using thromboelastography over the morbidity–mortality associated with transfusions and bleeding of pediatric patients undergoing heart surgery, for improved decision-making.

Currently there are no literature reports on the specific use of TEG-guided transfusion protocols in patients in whom this surgical technique is used.

Conflict of interest

None.

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None.

REFERENCES