Anesthetic management in VACTERL syndrome

VACTERL sendromunda anestetik yaklaşım

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ABSTRACT

VACTERL Syndrome consists of many problems regarding to anesthesia management due to multiple congenital malformations which may include vertebral, cardiac, trachea-oesophageal, renal and limb anomalies. Here we present our anesthetic management - in terms of preanesthetic evaluation, premedication, induction and maintenance of anesthesia and postoperative follow up - in a newborn patient with VACTERL Syndrome undergoing tracheo-oesophageal fistula operations. J Clin Exp Invest 2014; 5 (1): 103-105

Key words: VACTERL syndrome, anesthesia, ketamin

INTRODUCTION

VACTERL syndrome contains vertebral defects (v), anal atresia (A), cardiac defects (C), tracheo-oesophageal fistula (TE), renal malformations (R), and limb abnormalities (L). All major malformations in acronym is not necessary to diagnose a newborn’s VACTERL syndrome. At least three of these congenital malformations provide the diagnosis of the VACTERL syndrome [1-3]. Due to multiple congenital anomalies of different systems, anesthesia management of these patients can be complicated. In this case report, we present our anesthetic experience in a newborn with VACTERL syndrome operated due to tracheo-oesophageal fistula.

CASE REPORT

A newborn male infant weighted 2300 gram was born by caesarean section and there was no relationship between his parents. Newborn’s operation was immediately planned due to tracheo-oesophageal fistula. VACTERYL syndrome was diagnosed in our patient because of tracheo-oesophageal fistula, esophagus atresia, pes equinovarus and posterior uretal valve patterns.

Before operation, cystostomy was opened due to posterior urethral valve. Preoperative examination was normal except rhonchus findings in chest oscillation. Laboratory evaluation revealed BUN: 17 mg/dL, creatinine: 0.7 mg/dL, glucose: 62 mg/dL, potassium: 3.1 mmol.dL-1, Na: 141 mg.dL-1, calcium: 8.5 mg.dL-1, AST: 78 U.L-1, ALT: 7 U.L-1, total protein: 5.5 g.dL-1, albumine: 2.7 g.dL-1, bilirubin: 3.2 mg.dL-1, indirect bilirubin: 2.9 mg.dL-1, WBC: 11.9 K.UL-1, hemoglobin: 18.4 g.dL-1, hematocrit: 50%, platelet: 328 K.UL-1, International Normalized Ratio (INR): 1.22, activated partial thromboplastin time (aPTT): 49.8 seconds and because of the fact that disordered coagulation parameters, 46 cc fresh frozen plasma was given. Control values was INR: 1.05, aPTT: 28 seconds. The patient was operated after giving 184 cc.24hour-1 1/3 izodeks (Eczacıbaşı-Baxter®, İstanbul) and 9.6 cc. 24hour-1 total parenteral nutritional support. Antibiotic therapy was administered by giving 35 mg amicassin once a day and 55 mg ampicillin twice a
day. Patient was operated at 24th hour of caesar-ean section. Standard anesthesia monitoring (three lead electrocardiography (EKG), SpO2, heat, non-invasive blood pressure (NIBP)) was applied to patient. It was seen that SpO2 was %65, heart rate was 130 beats.min-1, NIBP was 67/34 mm.Hg-1. Patient had multiple organ anomaly. Premedication was done by giving 100 µg atropin after then an’esthesia induction was done by giving 2 mg.kg-1 ketamine, 0,1 mg.kg-1 midazolam. Rocuronium 0,5 mg/kg was used in order to achieve rapid intubation with minimal mask ventilation.

Following anesthesia induction patient was intubated with 3.0 number endotracheal tube by the expert anesthetist. Ventilation was maintained with pressure support mode, respiration rate was 50 breath.min-1 and airway pressure was set at 15 cm -H2O. Anesthesia was maintained with 0,75 mg.kg-1 iv bolus ketamine in every 30 minutes and %60-40 O2-air gas mixture. During the operation, SpO2 was between 95-100%, heart rate was between 100-120 beats.min-1, NIBP was between 59/27-71/35 mmHg, and etCO2 was between 36-55 mmHg. Operation was ended in 130 minutes. After 10 minutes from the end of the operation, spontaneous respiration began and patient was monitored in spontaneous mode. The patient’s SpO2 was between 93-95% with 60-40% O2-air gas mixture. However, the patient was not extubated due to the decrease of saturation in room air.

Patient transferred to the neonatal intensive care unit as intubated. In postoperative second day patient was ventilated with pressure support mode followed by t tube technic. At postoperative third day all pharyngeal reflexes was fully reversed, adequate spontaneous breathing effort was established and the patient was extubated uneventfully. Postoperative 4th day patient was sent to service and patient was discharged at postoperative 7th day.

DISCUSSION

Patients with VACTERL syndrome have a combination of many different repeated operations thanks to having many anomalies. Patients’ prognosis becomes better with advances in surgical techniques [1]. However, VACTERL syndrome has risk in terms of anaesthesia due to having many anomalies. The risk of aspiration increases in VACTERL patients depending on regurgitation and tracheo-eosophageal fistula [4]. In order to prevent regurgitation and aspiration we made premedication with atropine, fast induction and intubation with rocuronium. The induction method which we used is correlated with methods used in previous reports.

There have been reported no difference between inhaled anesthetic agents in literature. Khat-avkar [4] reported that 8-year-old patient with VACTERL syndrome who had been operated more than one was operated for cataract surgery and given halothane Mariano and his friends [5] informed in their case report that a newborn baby was operated due to VACTERL syndrome and isoflurane was used in anesthesia. Also, Yildiz and et al [2] used sevoflurane to patient with VACTERL syndrome. We prefer intravenous anesthesia with ketamine combined with atropine premedication in order to prevent laryngoscopy induced vagal stimulation and bradycardia which had been showed an effective method in newborns by Barois et all [6]. Barois et all showed that immediate ketamine analgesia plus atropine for tracheal intubation was effective in terms of decreasing pain and preventing vagal stimulation induced bradycardia. Also we used atropine premedication before induction in order to prevent ketamine induced hypersalivation.

Cardiac malformations have been reported in approximately 40-80% of patients with VACTERL association [7]. The most common cardiac anomaly in VACTERL syndrome is ventricular septal defect 22.3% [8]. Patients may also present with Tetralogy of Fallot, patent ductus arteriousus and atrial septal defect [8]. In our patient there was no cardiac anomaly and patient was haemodynamically stable during operation.

As findings are so variable in patients with VAC-TERL syndrome, each patient be carefully evaluat-ed individually and anesthetic approach should be preferred according to the patient.

Our patient has no cardiac anomaly which might complicate our anesthesia management how-ever we take all cautions in order to prevent pos-sible hemodynamical disturbances, risk of regurgi-tation and aspiration. According to our experience we can conclude that anaesthesiologist should prefer anesthesia management according to patient health conditions, risk factors and type of surgical treatment. Close monitoring is also crucial in perioperative and postoperative periods.

REFERENCES