Management of Anesthesia in a Child with a Large Neck Rhabdoid Tumor

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Introduction

Extrarenal malignant rhabdoid tumor (MRT) is a rare, highly aggressive childhood tumor with an extremely poor prognosis [1]. It represents less than 1% of all pediatric soft tissue malignancies, typically involving infants under the age of 1 year [2]. Extrarenal MRT frequently involves locations deep in the neck, abdomen, paranasal and orbital regions [3]. Due to its location, the neck tumor requires delicate surgery and meticulous anesthesia care.

Case Report

A 9-month-old female patient presented with giant rhabdoid neck mass (fig. 1). She underwent general anesthesia for neck tumor excision after three cycles of chemotherapy using Cooperative Weichteilsarcoma Studiengruppe (CWS) guidance, which resulted in reduction of the tumor mass. A computed tomography (CT) scan revealed a significant large protruding mass on the right lateral contour of the neck, which dislocated the trachea to the left and narrowed its lumen in its cranial half (fig. 2). Noticeable intratumor bleeding was an indication for urgent surgery. Her preoperative hemoglobin level was 8.5 g/dl; other laboratory tests were
normal. The patient had a Mallampati score of III. After atropine and midazolam premedication and preoxygenation, anesthesia was deepened with 1.5–2% sevoflurane in 100% oxygen, titrated to maintain spontaneous respiration. Oral laryngoscopy was performed using a Macintosh blade while the patient was breathing spontaneously, and revealed the airway to be Cormack-Lehane grade 3. The trachea was intubated with an armored tube No. 3.5 on the third attempt. After confirming the correct position of the tracheal tube, neuromuscular blockade was achieved with vecuronium bromide. A central venous catheter was placed in the left internal jugular vein, and blood pressure was continuously measured invasively. Anesthesia was than maintained with 1.5% sevoflurane in air and oxygen (50:50). Analgesia was maintained with intravenous fentanyl, administered after securing the airway. Awake fiberoptic bronchoscope-assisted intubation was planned as an alternative in case mask ventilation and intubation failed. During the operation, which lasted for almost 4 h, the patient received 370 ml of concentrated red blood cells, 400 ml of fresh frozen plasma and 1,200 ml of Hartmann’s solution. The intraoperative period was uneventful and macroscopically 95% of the tumor was removed (fig. 3). After the operation, the patient was transferred to the intensive care unit and extubated 24 h later.

Discussion

This case illustrates potential difficulties and challenges in airway management during major surgery in infants with a large neck tumor and the importance of prediction and adequate preoperative preparation. Head and neck masses in children are classified as developmental, inflammatory or neoplastic. Neoplasms of the head and neck account for approximately 5% of all childhood malignancies. Rhabdoid tumors were originally termed ‘rhabdo-
myosarcomatoid tumors’ due to their similar appearance to muscle-based tumors [4]. In the original 1978 Wilms tumor study [5] they were described as very aggressive renal neoplasms. Treatment of MRT is based on chemotherapy, radiotherapy and early surgical resection of the primary tumor if feasible [6]. Unfortunately, limited use of radiotherapy in these very young children leads to poor prognosis, with the survival rate being 17–30% [7]. Most children with a large neck mass have severe airway compromise (50% of cases). Airway management in infants with a large neck mass is a challenge for the anesthesiologist in consideration to displaced airway, difficulties in visualizing the airway and risk of sudden complete airway occlusion resulting in hypoventilation, hypoxemia, bradycardia or asystole [8]. Severe hemodynamic effects can occur due to the proximity of magistral blood vessels and decompression induced by surgical removal of a large neck tumor. Providing a safe airway to our patient was a crucial step. Preparation involved careful planning as well as the presence of two experienced anesthesiologists and equipment for a difficult airway that included a fiberoptic bronchoscope [9], laryngeal mask airway, oral and nasal airway, intubation stylet, and tube exchanger. Inhalation induction of anesthesia with sevoflurane was planned to avoid the risk of apnea associated with intravenous agent administration. The intraoperative period was uneventful and after the surgical removal of the tumor, the patient was transferred to the intensive care unit where we used a protocol for difficult airway extubation. Although there were no complications in establishing the airway in our case, it is important to evaluate the airway before the induction of anesthesia and to be prepared for possible difficult airway management.

**Conclusion**

Our case shows that the careful preoperative evaluation and anticipation of a difficult airway in an infant with a giant neck mass can minimize the possibility of intraoperative complications. Establishing a safe airway, inhalation induction of anesthesia with sevoflurane, intubation during spontaneous breathing and hemodynamic monitoring to prevent blood loss were crucial factors in the anesthetic management of this case.

**Disclosure Statement**

The authors report no conflicts of interest.

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