Case Report

Perioperative Care of an Infant With Wolf-Hirschhorn Syndrome: Is There a Risk of Malignant Hyperthermia

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Abstract

Wolf-Hirschhorn (WH) syndrome is a rare chromosomal abnormality with deletion of the short arm of chromosome 4. Associated clinical signs and symptoms of the condition include intrauterine growth restriction, severe psychomotor retardation, failure to thrive, characteristic facial features, profound developmental delays, seizures, and various congenital midline fusion anomalies. Given the associated congenital anomalies, anesthetic care may be required for various surgical interventions. We report a 3-month-old girl with WH syndrome scheduled for gastrostomy tube placement. Previous reports of anesthetic care for such patients are reviewed and potential perioperative concerns are discussed.

Keywords: Wolf-Hirschhorn syndrome; Malignant hyperthermia; Difficult airway

Introduction

Wolf-Hirschhorn (WH) syndrome is a rare chromosomal abnormality with deletion of the short arm of chromosome 4, specifically the 4p16.3 region [1]. Associated clinical signs and symptoms of the condition include intrauterine growth restriction, severe psychomotor retardation, failure to thrive, characteristic facial features, profound developmental delays, seizures and various congenital midline fusion anomalies [1-6]. The latter may include cleft lip or palate, congenital heart disease (atrial or ventricular septal defects), hypospadias and midline scalp defects [1-6]. Given the associated congenital anomalies, anesthetic care is often required for various surgical interventions. We report a 3-month-old girl with WH syndrome scheduled for gastrostomy tube placement. Previous reports of anesthetic care for such patients are reviewed and potential perioperative concerns are discussed.

Case Report

Institutional Review Board approval is not required by Nationwide Children’s Hospital for presentation of single case reports. A 3-month-old, 3 kg girl presented for gastrostomy tube placement. The patient was diagnosed with WH syndrome based on intrauterine ultrasound reports and subsequent genetic testing. Chromosomal microarray showed results consistent with the diagnosis of WH syndrome. On physical examination, the child was noted to be small for her age, quiet, and hypotonic. Dysmorphic features included wide-spaced eyes, microcephaly, microstomia, a prominent glabella, low set ears, and micrognathia. Her preoperative vital signs included a resting respiratory rate of 40 breaths/min, an oxygen saturation of 99% by pulse oximetry while receiving 0.5 L/min oxygen via nasal cannula, a heart rate of 134 beats/min, and a blood pressure (BP) of 83/62 mm Hg. Preoperative echocardiography revealed an atrial septal defect. No sedative pre-medication was required. The patient had a pre-existing 24-gauge intravenous (IV) cannula in the right saphenous vein with maintenance IV fluids infusing. The patient was held nil per os for 6 h and transported to the operating room where routine American Society of Anesthesiologists’ monitors were applied. General anesthesia was induced by the administration of IV ketamine (5 mg) and atropine (0.1 mg). Sevoflurane was administered and incrementally increased to 5%. A shoulder roll was placed beneath the patient, prior to anesthetic induction, to facilitate optimal alignment of airway structures. Because of the patient’s non-reassuring airway examination, spontaneous ventilation was maintained throughout induction and endotracheal intubation. Light cricoid pressure was applied, which brought the vocal cords into view (Cormack-Lehane score of I vs. IIa). Endotracheal intubation was easily achieved using a Miller 1 blade and a cuffed, styleted 3.0 endotracheal tube. The patient tolerated anesthetic induction and endotracheal intubation without adverse hemodynamic effects. Mechanical ventilation was initiated for the case. The surgery lasted approximately 30 min. Mild hypotension occurred during the maintenance phase.

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of anesthesia using sevoflurane (inspired concentration 2-3%),
despite the administration of a 20 mL/kg fluid bolus of 0.9% NS.
Therefore, the inspired concentration of sevoflurane was
decreased and neuromuscular blockade was provided by
the administration of rocuronium 3 mg (1 mg/kg). Fentanyl (2.5
µg/kg) was also administered. The expired concentration of
sevoflurane was decreased and neuromuscular blockade was
reversed with neostigmine 0.3 mg. Tracheal extubation
2 h later in ICU. Uneventful recovery.

<table>
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<th>Authors and reference</th>
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<tr>
<td>Chen et al [12]</td>
<td>8-month-old, 5 kg infant for cheiloplasty</td>
<td>(GA) Inhalation induction with halothane and NMB with succinylcholine and atracurium. Suspected delayed MH 2 h after surgery. Hyperventilated with 100% O₂, cooled with ice bags, dantrolene 2.5 µg/kg. Recovered uneventfully.</td>
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<td>Sammartino et al [13]</td>
<td>8-month-old, 5.2 female infant for goniotomy</td>
<td>(TIVA) Frequent respiratory tract infections related to aspiration. Premedication with IM atropine and ketamine followed by placement of IV cannula. Anesthesia induced with propofol and fentanyl. Difficult airway, requiring three attempts. Maintenance anesthesia with propofol boluses and vecuronium 0.1 mg/kg. Tracheal extubation 2 h later in ICU. Uneventful recovery.</td>
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<tr>
<td>Iacobucci et al [15]</td>
<td>5-year-old, 10 kg boy for circumcision</td>
<td>(GA) MP class 1 examination. Inhalation induction with sevoflurane. Easy endotracheal intubation. Cis-atracurium 0.1 µg/kg, alfentanil 25 µg/kg, and ketorolac 1 mg/kg administered intraoperatively. NMB reversed with neostigmine 0.3 mg. Tracheal extubation in OR. Uneventful recovery.</td>
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<tr>
<td>Mohsinuddin and Mayhew [16]</td>
<td>Case 1: 9-month-old male for open reduction of left hip</td>
<td>(GA) Previous intraoperative cancellation, at 8 months of age, due to difficult IV access and inability to place Foley catheter. Second surgery at 9 months included inhalation induction with sevoflurane and nitrous oxide. Central venous catheter was placed. Caudal block for postoperative analgesia.</td>
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<td></td>
<td>Case 2: 8-month-old male for bilateral myringotomy tubes</td>
<td>(GA) Inhalation induction with nitrous oxide and sevoflurane. Acetaminophen for postoperative analgesia. No anesthetic complications.</td>
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<td>Schmidt et al [17]</td>
<td>4-year-old, 9.9 kg female for atrial septal defect repair</td>
<td>(TIVA) No mention of respiratory infection. Premedicated with IV midazolam. RSI with propofol, fentanyl, and vecuronium (0.2 mg/kg). No airway issues. Maintained with fentanyl boluses and a propofol infusion. Arterial and central lines placed after induction. Uneventful anesthetic.</td>
</tr>
<tr>
<td>Choi et al [18]</td>
<td>33-month-old male for bilateral tympanoplasty and myringotomy</td>
<td>(TIVA) IM atropine (0.2 mg). Anesthetic induction with midazolam (0.1 mg/kg) and remifentanil infusion. Once bag-valve-mask-ventilation was established, rocuronium (0.6 mg/kg) was administered. Cormack-Lehane grade IIa view. High fever of 39 °C, 7 h after surgery with seizure activity. Fever persisted despite antipyretic agent. Suspected upper respiratory tract infection rather than delayed-onset MH. No dantrolene given. Eventual normal recovery and uneventful hospital course.</td>
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GA: general inhalational anesthesia; IM: intramuscular; IV: intravenous; MH: malignant hyperthermia; NMB: neuromuscular blockade; TIVA: total intravenous anesthesia; MP: Mallampati.

Discussion

Patients with known genetic syndromes pose a variety of challenges to the anesthesia provider [7]. WH syndrome is an extremely rare chromosomal disorder caused by partial deletion (monosomy) of the short arm of chromosome 4. It was first described in 1961 by Hirschhorn and Cooper and subsequently...
Table 2. Anesthetic Concerns in Patients With Wolf-Hirschhorn Syndrome

| 1. | Difficult airway management |
| 2. | Difficult vascular access |
| 3. | Associated congenital heart disease |
| 4. | Underlying seizure disorder |
| 5. | Hypotonia and choice of neuromuscular blocking agent |
| 6. | Controversial predisposition to malignant hyperthermia |
| 7. | Perioperative respiratory complications |
| 8. | Chronic aspiration |
| 9. | Non-MH-related febrile episodes (possibly secondary to chronic respiratory infections related to chronic aspiration) |

published in Humangenetik by both Hirschhorn and Cooper as well as Wolf et al in 1965 [8-10]. Current estimates place the prevalence at about 1:50,000 with a 2:1 female/male ratio.

WH syndrome is characterized by a spectrum of observable abnormalities, the most important of which for the anesthesia provider include congenital heart disease and facial structural abnormalities, which may lead to difficulties with airway management. Congenital heart disease includes primarily atrial and ventricular defects. Phenotypic features include ocular hypertelorism (wide-set eyes), low-set malformed ears, micrognathia (small, recessed mandible), microcephaly (small head), prominent glabella (a “Greek helmet” appearance), broad or beaked nose, cleft lip or palate, small down turned “fish like” mouth, hypotonia, intrauterine growth restriction (IUGR) with a subsequent failure to thrive. Pitt-Rogers-Danks syndrome is now grouped together with WH syndrome, since both are the result of deletions on the same portion of chromosome 4, although “Pitt” syndrome generally results in milder physical and congenital abnormalities [1, 6].

To date, there are a limited number of reports in the literature regarding the anesthetic care of patients with WH syndrome (Table 1) [11-18]. As with all anesthetic care, appropriate preoperative preparation begins with a thorough history and physical examination. Of primary concern to the anesthesia provider is the potential for difficulties with airway management and endotracheal intubation related to micrognathia, microstomia, and associate cleft lip/palate. The appropriate equipment for dealing with these difficulties should be readily available prior to anesthetic induction, as recommended by Engelhardt and Weiss [19]. General anesthesia can be induced by the incremental inhalation of sevoflurane in 100% oxygen with the maintenance of spontaneous ventilation until the airway is secured. In our patient, we administered a small dose of ketamine to induce anesthesia and maintain spontaneous ventilation followed by the administration of incremental concentrations of sevoflurane until adequate bag-valve-mask ventilation was demonstrated. Endotracheal intubation was accomplished without the use of neuromuscular blocking agents. Airway management may be further compromised by the risk of aspiration as noted in some of the case reports which have led some to use rapid sequence intubation (RSI). However, the use of RSI must be weighed against the risks of potential difficulties with airway management and endotracheal intubation.

While the early reports in the anesthesia literature suggested a correlation between the administration of inhaled anesthetics and the development of malignant hyperthermia (MH), more recent articles have questioned this association as there are several published reports showing that volatile agents have been used safely and without complications. Both of the patients in the two previous reports of potential MH-like symptoms had current upper respiratory infections and received high doses of atropine, muddying the picture as to whether or not it was MH, anticholinergic crisis, or active infection. When we questioned the MH hotline regarding a possible correlation between WH syndrome and the development of MH, we were told: “Although there is one report alleging an association, that has not been proven, nor do we consider that disorder a risk for MH.”

Given the associated hypotonia, another perioperative concern is the choice of neuromuscular blocking (N MBA), especially with regard to the safety of using succinylcholine [20]. Anecdotal reports have demonstrated the safety and efficacy of succinylcholine while other reports have failed to demonstrate significant perioperative issues such as prolonged duration of action following non-depolarizing agents including vecuronium, rocuronium, and cis-atracurium [21]. Alternatively, endotracheal intubation can also be accomplished without the use of an NMBA using a combination of propofol and remifentanil or as noted in one case report, midazolam and remifentanil [22].

Seizures may be a comorbid condition in patients with WH syndrome. Preoperative management strategies to limit the potential for perioperative seizures include optimizing and confirming therapeutic anticonvulsant levels prior to the surgical procedure. Routine anticonvulsant medications should be administered the morning of the procedure despite concerns of the patient’s nil per os status with subsequent intraoperative dosing as needed. Alternative routes of delivery (IV or rectal) may be required when enteral administration is not feasible. Consultation with the neurology or pharmacology service may be helpful to determine dosing conversion from enteral to IV administration or to guide intraoperative redosing.

In summary, WH syndrome is a rare chromosomal abnormality with deletion of the short arm of chromosome 4. The potential perioperative implications of WH syndrome are summarized in Table 2. While initial reports suggested a predisposition to the development of MH, subsequent reports have failed to confirm this association. Other comorbid conditions which may impact perioperative care include facial anomalies which lead to difficult airway management, associated congenital heart disease, central nervous system involvement including hypotonia and underlying seizure disorder, and the potential for chronic aspiration and perioperative respiratory problems.

References