Clinical Study
Anesthesia for Pediatric Deep Brain Stimulation

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In patients refractory to medical therapy, deep brain stimulations (DBSs) have emerged as the treatment of movement disorders particularly Parkinson’s disease. Their use has also been extended in pediatric and adult patients to treat epileptogenic foci. We here performed a retrospective chart review of anesthesia records from 28 pediatric cases of patients who underwent DBS implantation for dystonia using combinations of dexmedetomidine and propofol-based anesthesia. Complications with anesthetic techniques including airway and cardiovascular difficulties were analyzed.

1. Introduction

Dystonia is a movement disorder characterized by abnormal repetitive and twisting movements resulting in painful and debilitating postures [1]. Dystonia is termed primary when it is idiopathic or may be secondary to a variety of diseases including stroke, cerebral palsy, and neurodegenerative diseases. The pathophysiology of dystonia is not well understood though secondary dystonias often involve injury to the globus pallidus (GP) [2], and electrophysiologic recordings from GP pars interna (GPI) neurons of dystonic patients show abnormal activity [3]. There is no medical cure for dystonia. Medical therapy with anti-Parkinsonian, antispasmodic, or anticholinergic drugs is targeted at symptomatic relief but is only marginally effective [4]. Surgical treatments for dystonia traditionally involved lesioning thalamic nuclei or the pallidum [5] and were associated with a high incidence of complications, including hemiparesis and dysarthria [6]. With the advent of DBS, surgeons have turned to this new surgical approach.

Deep brain stimulation (DBS) has emerged as the treatment of choice for adult movement disorders that are refractory to medical therapy. This includes DBS of the subthalamic nucleus (STN) or GPi for Parkinson’s disease (PD) [7, 8] and DBS of the ventrolateral thalamus for the treatment of essential tremor [9]. As opposed to pallidotomy, DBS is less destructive of brain tissue, is associated with fewer cognitive and motor side effects, and is reversible and adjustable. Based on the evidence involving the GPi in the pathophysiology of dystonia and based on clinical observations that pallidal DBS alleviates off-medication dystonia in PD patients, the GPi has become the target of DBS for dystonia patients with significant clinical improvements [10–12].

The anesthetic challenges during DBS include providing sedation and analgesia with minimal respiratory depression or interference with electrophysiologic brain mapping and clinical testing. Intraoperative cardiovascular hemodynamics and blood pressure control are also important, as a potential complication of DBS procedures is intracerebral hemorrhage, which may occur in 2%–4% of cases [13]. The α-2 agonist dexmedetomidine (Dex) is a unique anesthetic agent which can provide sedation, analgesia, and decreased arterial blood pressure without respiratory depression over a broad range of doses, making it a potentially ideal sedative for DBS surgery [14]. These properties have led to the current introduction of Dex-based anesthesia as an alternative to the propofol/fentanyl anesthetic technique in PD patients who require intraoperative sedation [15].

Dex has also recently been reported for use in pediatric patients during neurosurgery [16, 17]. Here we present a retrospective study of the use of Dex and propofol anesthesia during DBS surgery in 28 pediatric cases from dystonia patients ranging from 7 to 17 years of age.
Table 1: Clinical characteristics of dystonia patients undergoing deep brain stimulation. SD: standard deviation.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Mean ±/− SD</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>12.2 ±/− 2.6</td>
<td>7–17</td>
</tr>
<tr>
<td>Sex (men/women ratio)</td>
<td>18/10</td>
<td>n/a</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>42.8 ±/− 13.4</td>
<td>26.0–86.4</td>
</tr>
<tr>
<td>Length of anesthesia (min)</td>
<td>297.4 ±/− 69.5</td>
<td>125–434</td>
</tr>
<tr>
<td>Length of surgery (min)</td>
<td>167.2 ±/− 57.5</td>
<td>65–313</td>
</tr>
</tbody>
</table>

2. Materials and Methods

After obtaining IRB approval, the anesthetic records from 28 pediatric DBS cases were reviewed. The following technique was used.

In the preoperative holding area, patients received local anesthesia (lidocaine plus bupivacaine) for placement of the stereotactic head frame (Leksell Model G, Elekta Inc, Atlanta, GA), supplemented with fentanyl and propofol boluses as needed. Patients received propofol infusions for the subsequent localizing MRI. Once the MRI was completed, patients were brought to the OR, where standard anesthesia monitoring (noninvasive arterial blood pressure cuff, pulse oximetry, electrocardiogram, and bispectral index monitor (BIS)) was applied.

Sedation was continued with propofol and maintained with propofol infusions plus small boluses of fentanyl as needed until scalp incision. Local anesthesia, consisting of 0.5% bupivacaine, was injected into the supraorbital and greater occipital nerves to produce a “scalp block.” Sedation was maintained during scalp incision and burr-hole placement and until the time of intraoperative testing, using a combination of Dex infusion (0.2 to 0.9 μg/kg/hr) and propofol boluses and/or infusions (rates of 15–100 μg/kg/min). BIS monitoring was used with target values >70. Following intraoperative testing, small doses of fentanyl and/or midazolam were administered as needed for discomfort. Following surgery, each patient underwent a second MRI to check electrode placement with anesthesia provided by propofol infusion.

Our pediatric patients ranged from 7 to 17 years of age, and included both males and females suffering from primary or secondary dystonia. Demographic data and clinical characteristics are summarized in Table 1. Anesthetic records from these patients were analyzed for medications used during surgery and potential intraoperative complications.

3. Results

Our analysis is comprised of 28 consecutive cases of pallidal DBS surgery for dystonia in pediatric patients. Table 2 summarizes the medications given intraoperatively including both the average dose and range. The table displays the data for the entire procedure (MRI plus procedure) and also for the implant procedure alone. Indeed, this table describes the relative use and dose of each anesthetic agent (as bolus or infusion) during intraoperative DBS and compares it to the values used during the entire case. Propofol was used in 27 cases for the entire procedure with an average total dose of 258.6 mg, while it was used in 24 of the cases during the DBS part of the procedure with an average total dose of 115.2 mg. A scalp block was performed in 7 cases.

Table 3 displays the surgical complications in our cases. No airway difficulty, respiratory depression, hypoxemia, or unplanned conversion to general anesthesia was encountered in any of the cases. In addition, no seizures occurred, and patients remained calm during the entirety of the surgery. One patient suffered a venous air embolus following burr-hole placement. The patient started to cough and complained of chest pain. The head was lowered and bone wax applied. The patient improved but the procedure was aborted. One month later, he underwent successful DBS surgery. Our rate of venous embolism (1 out of 28 cases or 3.6%) is very similar to the rate reported in the literature for DBS surgeries [18]. Venous emboli are a known potential complication of DBS, and although rare, they should always be considered when the patient begins to cough suddenly during the operative phase [18].

4. Discussion

We here describe a retrospective study examining anesthetics used for DBS surgery in pediatric dystonia patients. As previously mentioned, awake neurosurgery presents unique anesthetic challenges. Different techniques have been developed to meet this challenge, with propofol infusion alone or in combination with fentanyl/remifentanil the method most frequently reported [19, 20]. However, the use of these anesthetic techniques may be accompanied by complications such as airway obstruction, deep sedation, and respiratory depression [21, 22].

Dex is a central α-2 adrenergic stimulant and is known to have dose-reducing effects when used with inhalational anesthetics and opiates [23]. Dex offers some unique anesthetic characteristics as it provides sedation and analgesia while minimizing respiratory depression even at large doses [24]. Therefore, Dex is ideally suited for the “asleep-awake-asleep” technique often required for awake craniotomy [25]. With this technique, patients are anesthetized or sedated for the initial craniotomy then allowed to wake up to participate in neurocognitive mapping, and the original anesthesia or sedation is resumed for the final part of the procedure [17, 25]. Sedation with Dex leaves a patient sleep but still arousable to an alert state, which is very useful for intraoperative recording [25].

The anesthetic challenges during DBS surgery are to alleviate the stress of awake neurosurgery without interfering with neurophysiological monitoring or inducing respiratory depression. We therefore employed a similar “asleep-awake-asleep” anesthesia technique for pediatric dystonia patients using a combination of propofol and Dex-based anesthesia. In this way, anesthesia was performed with a high degree of safety and minimal side effects or intraoperative complications. All patients tolerated the procedure well and appeared comfortable. BIS monitoring helped in guiding the
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References


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