

Less Invasive Pediatric Spinal Deformity Surgery: The Case for Robotic-Assisted Placement of Pedicle Screws

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Abstract

Introduction: Pediatric spinal deformity involves a complex 3-dimensional (3D) deformity that increases the risk of pedicle screw placement due to the close proximity of neurovascular structures. To increase screw accuracy, improve patient safety, and minimize surgical complications, the placement of pedicle screws is evolving from freehand techniques to computer-assisted navigation and to the introduction of robotic-assisted placement. **Purpose:** The aim of this review was to review the current literature on the use of robotic navigation in pediatric spinal deformity surgery to provide both an error analysis of these techniques and to provide recommendations to ensure its safe application. **Methods:** A narrative review was conducted in April 2021 using the MEDLINE (PubMed) database. Studies were included if they were peer-reviewed retrospective or prospective studies, included pediatric patients, included a primary diagnosis of pediatric spine deformity, utilized robotic-assisted spinal surgery techniques, and reported thoracic or lumbar pedicle screw breach rates or pedicle screw malpositioning. **Results:** In the few studies published on the use of robotic techniques in pediatric spinal deformity surgery, several found associations between the technology and increased rates of screw placement accuracy, reduced rates of breach, and minimal complications. All were retrospective studies. **Conclusions:** Current literature is of a low level of evidence; nonetheless, the findings suggest the accuracy and safety of robotic-assisted spinal surgery in pediatric pedicle screw placement. The introduction of robotics may drive further advances in less invasive pediatric spinal deformity surgery. Further study is warranted.

Keywords

spine, pediatrics, operative treatments, mini-incision surgery, scoliosis, robotics

Introduction

Pediatric spinal deformity often involves complex 3-dimensional (3D) deformities associated with small and frequently dysplastic pedicles [37,43]. One of the goals of surgical management is to obtain a stable and solid fusion after placement of spinal anchors followed by spinal deformity correction. The mainstay of operative pediatric spinal deformity management has been open posterior spinal instrumentation and fusion (PSIF), most frequently using pedicle screws inserted with freehand techniques [17,27,53].

In children with spinal deformity, the insertion of thoracic pedicle screws presents increased risk of malpositioned pedicle screws and other complications compared with lumbar pedicle screw insertion. Usually, this is caused by altered morphology in the thoracic pedicle, including dysplastic and narrow pedicles and altered location of neurovascular structures secondary to spinal deformity [11,15,17,37,43,47,55,58]. Screw malpositioning is a

commonly documented implant-related complication; rates vary between 4.2% and 25% for placement with freehand techniques [15,17].

Multiple methods have been used to improve the accuracy and safety of thoracic pedicle screw placement, including anatomically based techniques (use of pedicle probes and pedicle wall palpation, visualization of cancellous starting points, and laminectomy with direct pedicle visualization and palpation), as well as preoperative and

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intraoperative use of various ionizing radiation techniques such as fluoroscopy, plain radiographs, and computed tomography (CT) scans. Several techniques for neuro-monitoring have gained widespread acceptance as clinical standards for improving patient safety, including transcranial motor-evoked potentials and somatosensory evoked potentials [7,23,44,49,50], spontaneous electromyography (EMG) monitoring [7,23], triggered electromyography (t-EMG) screw stimulation [40,41], and pedicle probes with electroconductive tips [36]. Issues of safety, reliability, and accuracy have been crucial in the ongoing development and improvement of computer-assisted pedicle screw navigation in spine surgery.

First used in spine surgery in the late 1990s, computer navigation is proposed as a way to allow for real-time assessment of screw trajectory accuracy [15,20]. When comparing different pedicle screw insertion methods, computer-assisted surgical navigation of pedicle screws has lower rates of malpositioned screws and unplanned return to the operating room (OR) [5,6,22,24,25,31,39,52]. Additionally, surgical navigation has been associated with significantly lower rates of medial wall breach compared to freehand techniques (0%–7.9% vs 8.6%) [5,6,22]. The incidence of an unsafe, significant medial breach (50% of the screw diameter) was 7.6 times more likely to occur using freehand screw insertion compared with surgical navigation in one study [51].

Robotic-assisted spine surgery (RASS) is a natural evolution from computer-assisted surgical navigation, and it has several potential advantages over freehand techniques, including improved stability and maintenance of screw trajectory during insertion, opportunity for preoperative screw trajectory planning to find ideal trajectory, and potential for screw insertion without direct visualization of bony anatomy and screw entry site. Accurate, reliable, and safe pedicle screw insertion using robotically assisted techniques opens the possibility for lesser invasive surgical approaches for pediatric spinal deformity as less invasive robotic-assisted techniques have already been adopted by adult spine surgeons to assist in the correction of adult spinal deformity [2,8,9,26,38,54].

The first robotic system cleared for use in spine surgery by the US Food and Drug Administration (FDA) was the SpineAssist (Mazor Robotics LTD, Caesarea, Israel) in 2004 [13,16]. The SpineAssist was replaced by the Mazor Renaissance (Mazor Surgical Technologies Ltd, Caesarea, Israel) in 2011, and the first reports of the use of RASS in pediatric spinal deformity utilized this system [13,16,18,28]. This system was a mechanically driven bone-mounted system with a robotic manipulator, allowing for 6 degrees of freedom, attached to a bone-mounting frame on the patient's spine.

There are now several FDA-cleared spine robots, including ROSA Spine Robot, which was subsequently replaced by the Rosa One Spine system (Zimmer Biomet Robotics,

Montpelier, France), ExcelsiusGPS (Globus Medical, Inc, Audubon, PA), and Mazor X (Medtronic, Dublin, Ireland) [13]. The increasing interest in RASS has led to further FDA clearances, including the Cirq system (Brainlab AG, Munich, Germany) [14].

The safety, efficacy, and accuracy of these systems is an active area of research [11,21,52]. Increased adoption of robotically assisted insertion of pedicle screws has largely been driven by adult spine surgeons who have pioneered single-position lateral surgery for simultaneous anterior and posterior spinal fusion in the lumbar spine, as well as minimally invasive percutaneous screw insertion in the lumbar spine [19,21].

The current generation of spine robots facilitates pedicle screw placement through an end-effector that functions as a rigid drill sleeve that obtains and maintains planned screw insertion trajectory. The current generation of robotic spine systems provides for increased stability of the robotic arm, increased stability of spine region of interest via spine stabilization clamps, and improved overall system stability via direct connection of the robot to the skeletal anatomy of the patient. Simultaneous computer-assisted surgical navigation tracking the instruments and the skeleton also provides real-time visual feedback layered on top of the robotic navigation.

We aimed to review the current literature on the use of robotic navigation in pediatric spinal deformity. We sought to provide an error analysis of these techniques to include reported breach rates and surgical and patient factors that may lead to increased breach or other complications. In addition, we aimed to provide recommendations and best practices to ensure safe placement of pedicle screws and discuss further applications of robotic surgery in pediatric spine deformity correction based on our own experience.

Methods: Search Strategies and Criteria

A narrative review was conducted in April 2021 using the MEDLINE (PubMed) database. The search strategy consisted of the operators found in Appendix A. Studies were included if they were peer-reviewed, retrospective, or prospective studies; included pediatric patients and a primary diagnosis of pediatric spine deformity such as adolescent idiopathic scoliosis, neuromuscular scoliosis, congenital scoliosis, or Scheuermann's kyphosis; utilized RASS techniques; and reported thoracic or lumbar pedicle screw breach rates or pedicle screw malpositioning.

The search identified 40 articles. Of the articles identified, 16 abstracts were screened, 12 of which were excluded, 3 for being a research design other than a prospective or retrospective study, 4 for not utilizing robotic navigation techniques, 3 for not including pediatric patients, 1 for not including thoracic or lumbar pedicle screw placement, and 1 for not providing breach rates. Four full-text articles were

Table 1. Summary of articles identified studying robotic-assisted spine surgery (RASS) in the pediatric population.

Author	Year published	Study type	Downs and Black score (x/28)	Robotic platform utilized	Number of patients	Diagnosis	Mean age at surgery (years)
Macke	2016	Retrospective review	17	Mazor Renaissance	50	AIS	N/A
Shaw	2018	Retrospective review	18	Mazor Renaissance	19	AIS	14.5 ± 1.7
Gonzalez	2020	Retrospective review	16	Mazor X Stealth Edition	40	Neuromuscular (N = 5), spondylolisthesis (N = 4), congenital scoliosis (N = 2), AIS (N = 26), other (N = 3)	14.5 ± 2.6
Morse	2021	Retrospective review	18	Mazor X Stealth Edition	19	AIS (N = 13), neuromuscular scoliosis (n = 3), Scheuermann's kyphosis (n = 1), post radiation scoliosis (n = 1), congenital scoliosis (n = 1)	14.6 ± 2.2

reviewed, 4 of which met inclusion criteria (Table 1) [16,29,34,45]. Study quality was graded using the Downs and Black checklist, a 27-question assessment with a maximum score of 28 [12]. Similar to other studies, question 27 was replaced with “Was a power analysis conducted?” with a score of 1 for yes and 0 for no [30,33].

Results

Error Analysis in Pediatric Robotic Spine Surgery

Four studies were identified that have reviewed the potential complications with the use of both second-generation and third-generation robotic navigation in pediatric spinal deformity.

Macke et al reviewed 48 pediatric patients following screw placement with the second-generation Mazor Renaissance system and analyzed 662 screws [29]. The authors performed postoperative CT scans on all patients and found a total screw medial misplacement rate of 7.2%, with the majority (4.5%) having a medial breach between 2 and 4 mm. Additionally, they reported 1.5% of screws had a medial breach between 4 and 6 mm, and 1.2% were greater than 6 mm. Despite the medially placed screws, screws were more often misplaced laterally. While the authors did not evaluate time per screw to evaluate a learning curve, they reported a 9.6% breach rate in the first third of patients and a 7.4% breach rate in the last third. They also reported that patients who underwent preoperative CT scanning in the prone position were less likely to have a breach.

Shaw et al also examined pedicle screw accuracy with the Mazor Renaissance [45]. Following screw insertion, each pedicle screw was stimulated with t-EMG, with thresholds less than 8 mA deemed abnormal. These screws were subsequently removed and the tract was probed with a ball-tip probe. If a medial wall was not palpated, a new screw

trajectory was created. The authors reviewed 49 patients and 844 instrumented pedicles and reported 28 (3.3%) screws with abnormal stimulation. All trajectories were found to have an intact medial wall and the screw was reinserted; 51% of these screws were periapical, 19 on the curve concavity, and in a logistic regression analysis, smaller pedicle width was found to be a significant predictor for t-EMG amplitude.

Recently, Gonzalez et al reviewed 40 patients with 314 screws placed robotically with the Mazor X [16]. The authors used a preoperative CT scan (12 patients) for surgical planning and then an intraoperative O-arm scan (Medtronic, Dublin, Ireland) for registration in lieu of fluoroscopy following spine exposure and facetectomies due to their belief in the difficulty of using fluoroscopy to register patients with significant apical rotation. The authors used either a spinous process clamp or a Schanz pin placed in the posterior superior iliac spine (PSIS) to attach the robotic arm. Screw position was assessed with intraoperative fluoroscopy and postoperative radiographs. They reported 4 (1.3%) screws that had difficult placement, 3 of which were lateral, and 1 which was unable to be placed due to a sclerotic pedicle. Overall, the authors reported a 98.7% screw success rate. Two patients in the series had wound drainage and returned to the OR for debridement.

Morse et al reviewed 19 patients and 194 pedicles, 168 (86.6%) of which were placed with the Mazor X [34]. A preoperative CT scan was performed for all patients and following instrumentation a 3D mobile fluoroscopic scan (Ziehm Imaging GmbH, Nuremberg, Germany) was used to assess screw position following placement in the OR. Standing biplanar low-dose whole-body slit radiographs (EOS Imaging, Paris, France), reconstructed to map individual vertebral body position, were used to assess the effect of vertebral body rotation on complications. The authors reported 29 (15.0%) screws that were

abandoned or converted to freehand; the most common reason for freehand conversion was soft tissue impaction on the end effector (48.3%). There were 15 (8.9%) total medial breaches that occurred, the majority under 2 mm. There were 2 medial breaches between 2 and 4 mm, 1 of which was associated with a durotomy, which occurred due to inadequate tightening between the bone mount bridge and the spinous process clamp, causing medial skive from excess robotic arm motion. The durotomy did not produce any adverse symptoms postoperatively. Both of these breaches occurred in the first case, and excluding breaches less than 2 mm, robotically placed screws had a 98.8% accuracy rate. Most of the breaches occurred in the first case, with the majority occurring in the thoracic spine between T4 and T12. Risk of breach was associated with small pedicle diameter and vertebral body kyphosis.

In addition to reporting on errors, the authors presented a learning curve for screw time and reported a mean time per robotically placed screw of 3.6 ± 2.4 minutes. The time improved over the second half of cases and with experience. Slower screw placement times occurred in upper thoracic spine (T2–T3) and no difference was found between placement times of apical or peri-apical vertebrae and all other vertebrae [34]. The authors assessed screw accuracy, defined as a trajectory that matched preoperative templating and reported 76% of screws having a matched trajectory. Screws were most accurately placed in the lumbar spine and least accurately in the thoracic spine at the T4–T6 level. Additionally, cortically planned screws, those in which the mid-axis of the screw is placed on the pedicle cortex, were the least accurate. Accuracy improved over the second half of cases and was affected by smaller pedicle diameter, coronal rotation, and sagittal rotation. Screw insertion times are currently under 1 minute.

While the use of RASS in pediatric spinal deformity remains an underreported topic in the literature, these studies demonstrate the accuracy and safety of robotic screw placement. With the use of both second- and third-generation robotic techniques, there were no reported neurologic complications during pediatric spinal deformity correction [15,29,34,45]. These studies also highlight a limitation in assessing postoperative screw position in the pediatric population; only 1 study utilized a postoperative CT to assess screw malposition [29]. The use of imaging modalities other than CT may only detect dangerous breaches and not assess screw position, although these modalities are used to limit radiation exposure in children. Further study is required to determine the screw malpositioning rate. Of note, most reports utilized an open approach to the spine and screw placement could be visualized directly. Currently, the screw insertion technique has evolved and the transmuscular placement of proximal thoracic screws is being utilized.

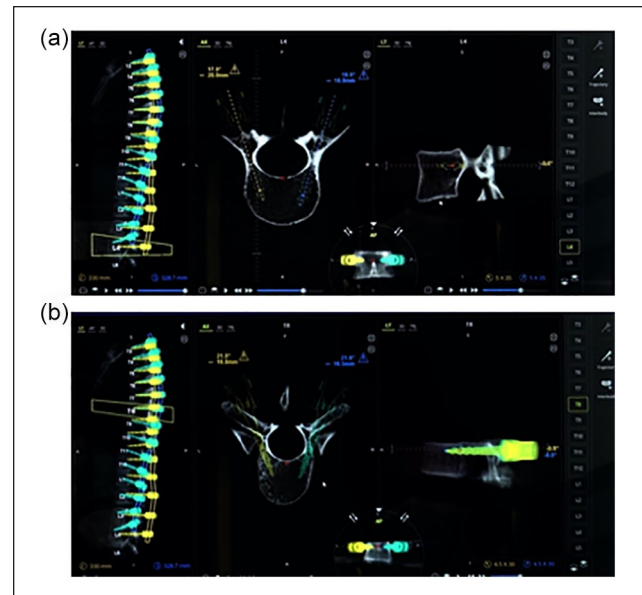


Fig. 1. Planning software showing 2 screw trajectories: (a) both screws are planning completely within the pedicle and (b) both screws are planned as in-out-in with the mid-axis of the screw entering the pedicle, exiting, and then entering the vertebral body.

Recommendations to Optimize Workflow and Ensure Safety

While RASS may offer further improved accuracy and reliability in placing pedicle screws, its adoption presents several challenges. First, the entire surgical team must learn to safely and efficiently implement a new robotically assisted workflow. This includes classroom, dry lab, cadaver lab, and mentorship-type technical training. Adequate training minimizes the duration of and id complications in the learning curve.

The use of the robot can be divided into 4 phases: preoperative planning, intraoperative setup, surgical execution, and postoperative assessment.

Planning may be performed in the OR with a plan and scan workflow or outside the OR with preoperative high-resolution spine imaging ahead of surgery. Preoperative imaging allows for optimal planning of screw trajectories, especially with abnormal or highly dysplastic anatomy. The preoperative scan is uploaded to the planning software and the surgeon can choose screw trajectory optimizing safety and screw purchase. The planning software maximizes the surgeon's ability to select the correct trajectory and screw size as the screw's trajectory, diameter, and length can be viewed in the axial, coronal, and sagittal planes. During templating, hypoplastic, sclerotic, and small diameter pedicles are noted, as pedicle morphology differences may suggest extra-pedicular (in-out-in) trajectories (Fig. 1). Decreased screw placement accuracy was noted when preoperative plan included screw placement directly overlying the pedicle cortex [34]. Additionally, when pedicle diameters were

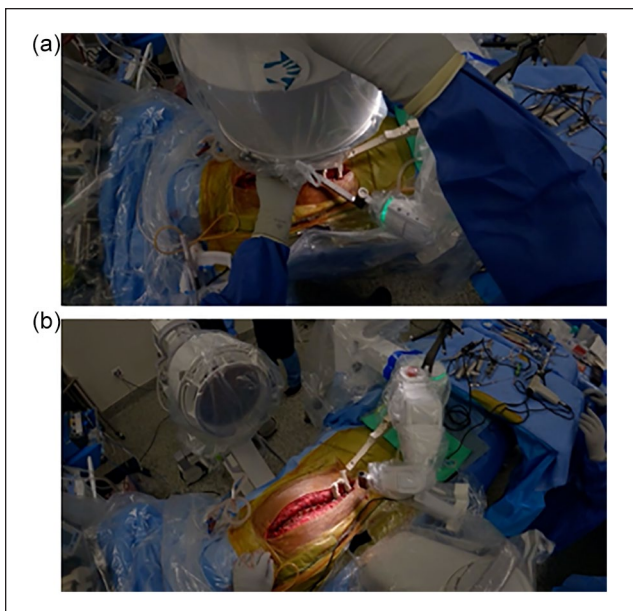


Fig. 2. Intraoperative positioning of the fluoroscopic C-arm during registration. The surgeon guides the fluoroscopy technician to ensure appropriate placement during an anteroposterior (a) and a lateral (b) image.

less than the smallest pedicle screw diameter, there was a significantly increased risk of breach [34,45]. Vertebral body kyphosis also correlated with increased rates of breach felt to be associated with soft-tissue pressure on the robotic end effector resulting in decreased accuracy of screw placement. Patient factors including height, weight, and length of deformity also impact deformity planning, because more than 1 registration may be required when more than 7 spinal levels are involved.

Setup of the robotic system can create ergonomic challenges for the surgeon and OR staff. We recommend placing the camera at the head of the table with a direct line of sight of the robotic arm and the operative field. Staff must be familiar with draping the robot, and all assistants must vigilantly maintain a sterile field. Monitors and the base station should be placed to allow visualization by the surgeon and assistants. The radiology technician must manipulate the fluoroscopic C-arm to capture the appropriate registration views while avoiding contact between the C-arm and the robot (Fig. 2). Failure to manipulate the C-arm correctly can result in registration failure requiring the registration to be completed again. Prior to registration, when obtaining the topographical optical scan, a blue towel is placed over the spine and the lights are turned away to best define the field of view. During registration, the targeting device can be brought close to the patient to obtain the largest field of view when taking fluoroscopic images. When moving between the anteroposterior (AP) view and the oblique view, the targeting device must be removed so that it does

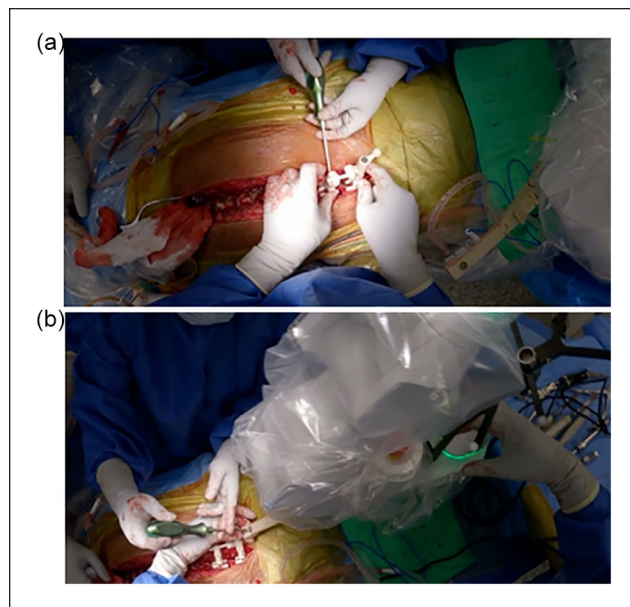


Fig. 3. Direct attachment of Mazor X to the patient. Intraoperative imaging demonstrating adequate tightening of both the dual spinous process clamp (a) and connection to the bone mount bridge (b).

not inadvertently strike the patient. Following the registration, the surgeon must confirm that the registration produces an anatomic match to the preoperative imaging.

Although most robotic spine systems are freestanding on the OR floor, the Mazor X Stealth robotic platform is attached to the operating table and directly to the patient via threaded skeletal pins and/or spinous process clamps (Fig. 3). Direct patient attachment decreases the risk of inadvertent malalignment from patient or robot motion.

We advocate following traditional freehand pedicle screw steps including visual confirmation of starting points, palpation of pedicle tracks with ball-tip probe, and intraoperative fluoroscopy to confirm screw placement (Fig. 4). The surgeon must visually confirm that the anti-skive pin is properly seated on the pedicle. Occasionally, the anti-skive pin will not seat properly onto the pedicle and skive can occur. To prevent this, a burr can be used to create a pilot hole so that the pin does not slide off the pedicle and the trajectory can be maintained for drilling [16,34]. In addition, we found that the use of intraoperative 3D fluoroscopic scans is helpful to assess accuracy and modify screw planning.

Future Applications: Minimally Invasive Pediatric Spinal Deformity Surgery

Minimally invasive surgery (MIS) has become an acceptable surgical approach for traumatic, neoplastic, degenerative,

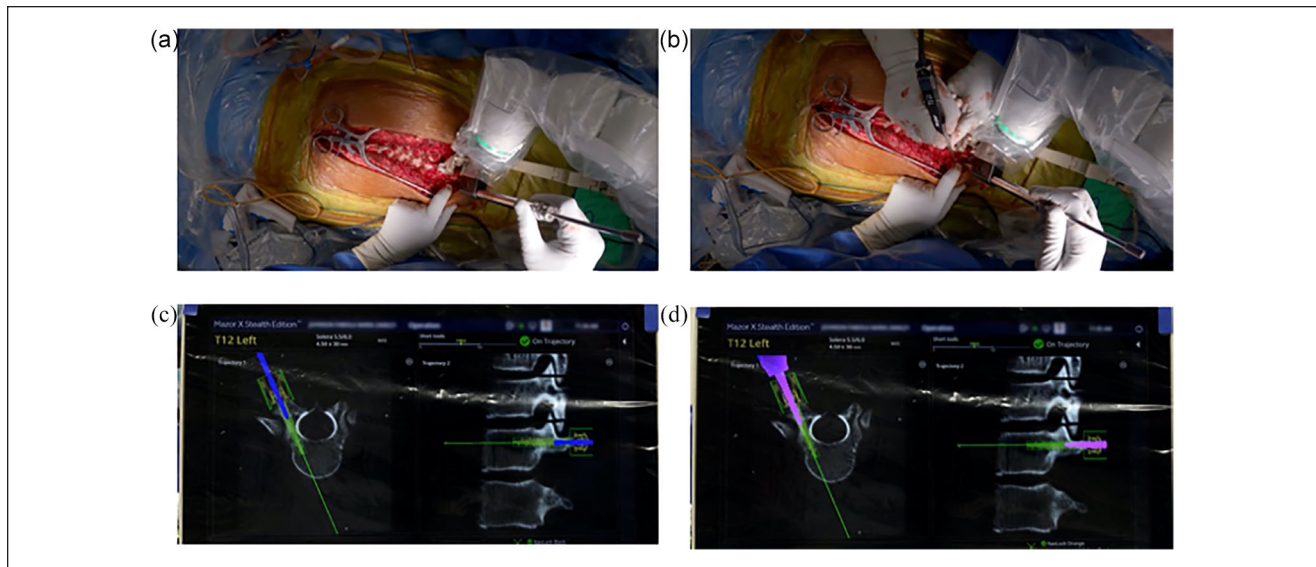


Fig. 4. Intraoperative screw placement. Direct visualization of a starting point (a). Given the potential of medial skive for this pedicle, a pilot hole was created for the start point with a burr (b). The simultaneous navigation showing drilling is occurring on the templated screw trajectory (c). Screw placement follows on the templated screw trajectory (d).

and even deformity spinal conditions in adults. These methods have demonstrated similar outcomes, with lower morbidity and mortality [3,4,35,48]. More recently, several authors have suggested that MIS has a role in the management of pediatric spinal deformity surgery [1,10,32,42,46,56,57]. When compared to open posterior spinal fusion (PSF), MIS has consistently been associated with lower estimated blood loss with no differences from open surgery in complication rates but with longer operative times [1,32,42,46,57].

Robotic surgical techniques facilitate a less invasive access to the pedicles via percutaneous or transmuscular approaches, without direct open exposure of the spine. Avoiding dissection of the paraspinal musculature may maintain better vascularity to the bone, thereby decreasing risk of infection, enhancing healing, and lessening pain. Compared to adults, pediatric patients have more flexible spines and excellent fusion rates. Less invasive approaches coupled with lower profile implant systems may result in similar safety and complication profiles as traditional open surgery. Rapid advancement in robotic techniques has resulted in faster operative times, more accurate pedicle screw placement, and equivalent results to computer-assisted freehand surgical navigation. Further research is required to confirm the use of these technologies in maintaining curve correction and measuring screw malpositioning.

Appendix A

Search Terms

“adolescences”[All Fields] OR “adolescence”[All Fields] OR “adolescent”[MeSH Terms] OR “adolescent”[All

Fields] OR “adolescence”[All Fields] OR “adolescents”[All Fields] OR “adolescent s”[All Fields]) AND (“idiopathic”[All Fields] OR “idiopathically”[All Fields] OR “idiopathics”[All Fields]) AND (“scoliosis”[MeSH Terms] OR “scoliosis”[All Fields] OR “scolioses”[All Fields]) OR ((“paediatrics”[All Fields] OR “pediatrics”[MeSH Terms] OR “pediatrics”[All Fields] OR “paediatric”[All Fields] OR “pediatric”[All Fields]) AND (“spine”[MeSH Terms] OR “spine”[All Fields] OR “spines”[All Fields] OR “spine s”[All Fields])) OR ((“paediatrics”[All Fields] OR “pediatrics”[MeSH Terms] OR “pediatrics”[All Fields] OR “paediatric”[All Fields] OR “pediatric”[All Fields]) AND (“spine deform”[Journal] OR (“spine”[All Fields] AND “deformity”[All Fields]) OR “spine deformity”[All Fields])) OR (“Neuromuscular”[All Fields] AND (“scoliosis”[MeSH Terms] OR “scoliosis”[All Fields] OR “scolioses”[All Fields])) OR ((“congenital”[MeSH Subheading] OR “congenital”[All Fields] OR “congenitally”[All Fields]) AND (“scoliosis”[MeSH Terms] OR “scoliosis”[All Fields] OR “scolioses”[All Fields])) AND (“robot”[All Fields] OR “robots”[All Fields] OR “robotically”[All Fields] OR “robotics”[MeSH Terms] OR “robotics”[All Fields] OR “robotic”[All Fields] OR “robotization”[All Fields] OR “robotized”[All Fields] OR “robots”[All Fields]).

Declaration of Conflicting Interests

The author(s) declared the following potential conflicts of interest with respect to the research, authorship, and/or publication of this article: Roger F. Widmann, MD, reports a relationship

with Medtronic. The other authors declared no potential conflicts of interest.

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Human/Animal Rights

All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975, as revised in 2013.

Required Author Forms

Disclosure forms provided by the authors are available with the online version of this article as supplemental material.

Supplemental Material

Supplemental material for this article is available online.

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