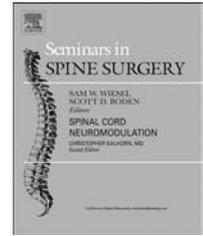


Available online at www.sciencedirect.com

ScienceDirect

www.elsevier.com/locate/semss

Evaluation and workup in revision spine surgery



Jeffrey M. Hills, Elliot Kim, Inamullah Khan, and Clinton J. Devin*

Department of Orthopaedics and Rehabilitation, Vanderbilt University Medical Center, 1215 21st Avenue South, Suite 4200, MCE-South Tower, Nashville, TN, United States

ABSTRACT

The incidence of spine surgeries performed annually continues to rise and despite advances in instrumentation and surgical techniques, reoperation rates are still high. Common reasons for revision surgery include an incorrect preoperative diagnosis, technical error, complications of procedure or implants, or a poor surgical candidate for the index procedure. Identifying the source of symptoms can present a diagnostic challenge and requires a careful and strategic approach. Given that outcomes lessen with each subsequent spine surgery, spine surgeons must be cautious and meticulous when considering a revision spine surgery. This article will review a general diagnostic approach when evaluating a patient with persistent symptoms, along with a review of the more common indications for revision spine surgery.

© 2019 Published by Elsevier Inc.

1. Introduction

Despite advances in surgical technique and fusion technology, the rate of revision spine surgery has increased.^{1–3} In 2009, the rate of revision spine fusion procedures had risen to 7.2 per 100,000 adults in the US.⁴ Reoperation rates following spine surgery vary by level and type of procedure. Lumbar 5-year overall reoperation rate (fusion and non-fusion) has been reported at 15% with only slightly higher rate in fusion surgery.⁵ Reoperation rates after cervical spine surgery have been reported up to 9% for single-level anterior cervical discectomy and fusion (ACDF), up to 10% for multilevel ACDF, 5% at 31.7 months for cervical foraminotomy and as high as 27% at 41 months after laminectomy and fusion.^{6–10} In an analysis of trends of revision spine surgery in the US, complication or failure of surgical implant was the most common indication for revision spine surgery, accounting for over 50% of cervical, thoracic, and lumbar revision surgeries.⁴ Other common diagnosis included degeneration or displacement of intervertebral disc, stenosis, kyphosis, and spondylosis.

Given the high prevalence of failed spine surgery and revision spine surgery, it is inevitable that spine practitioners will encounter patients presenting for revision spine surgery. The lack of descriptive or clinically meaningful terminology in diagnosis such as “failed back surgery syndrome” highlight the diagnostic challenge in the postoperative symptomatic spine patient.¹¹ Thus, it is crucial that a careful, systematic evaluation and diagnostic workup be performed in all patients presenting for revision spine surgery. Establishing a detailed timeline from the index surgery to onset or recurrence of symptoms is critical in developing a differential diagnosis. A patient’s symptoms should be categorized as axial back/neck pain or radicular arm/leg pain. These details should then be correlated with diagnostic laboratory and imaging evaluation to ultimately guide an effective treatment plan.

The first section of this article will review the patient evaluation including key portions of the history, exam, and diagnostic tests in developing a differential diagnosis and treatment plan when evaluating a symptomatic patient with prior spine surgery. The second section will review the incidence and diagnostic workup for the most common indications for revision surgery.

* Corresponding author.

E-mail address: Clintondevin@gmail.com (C.J. Devin).

2. Patient evaluation and developing a differential diagnosis

2.1. Patient history

A detailed patient history is critical to establish an appropriate diagnosis and treatment plan for symptomatic patients with a prior spine surgery. Patients should be asked to recall what symptoms they were having prior to the index procedure, who performed the procedure, how much relief the procedure provided, how long that relief was experienced, and if there was an inciting event that caused the pain to recur. Supplementing this history with preoperative notes and imaging from the index procedure can be helpful. Unfortunately, patients do not always recall preoperative symptoms, often discounting the extremity pain for which they underwent surgery.¹² It is helpful to consider a differential based on the patient's response to their index procedure. Patients can often be placed into 1 of 3 categories: 1) no clinical response after surgery; 2) a brief period of improvement but become symptomatic by 1 year; or 3) symptom free for an extended period (>1 year) before becoming symptomatic. By combining the presenting symptoms and response to index procedure, a working differential diagnosis can be developed.

2.1.1. No symptom improvement after index procedure

If an indicated surgery was performed, the patient should experience at least partial relief of their initial symptoms following their initial procedure. Failure to achieve any pain-free interval is likely related to an unindicated surgery (including an incorrect preoperative diagnosis or surgery directed at asymptomatic imaging findings), technically flawed procedure (e.g. unrecognized pathology, hardware malposition, and wrong-level surgery), or negative patient characteristics (high preoperative opioid burden, psychiatric distress, poorly controlled diabetes to name a few). A patient's failed outcome can be a combination of any of these factors. Identifying the specific reasons and risk factors that contributed to the poor outcome is essential to preventing a repeat failure.

Evaluation should begin with a detailed review of imaging obtained (and attention to imaging not obtained) prior to surgery and postoperative imaging provides insight into whether all areas of correlative pathology were addressed, if the intended level was decompressed, and if a diagnosis was potentially missed. Failure to recognize far lateral disc herniations or foraminal/recess stenosis in the setting of central stenosis can also lead to inadequate decompressions and persistent pain following spine surgery.¹³ The incidence of concurrent (tandem) cervical and lumbar stenosis range from less than 1% up to 20% of patients with stenosis.¹⁴ Thus, more proximal stenosis should also be considered if upper motor neuron signs exist and or imaging appears to demonstrate that a technically well done surgery was performed at the index level with no clear explanation of persistent symptoms. The incidence of wrong level spine surgery is low relative to other spine related complications, but still responsible for nearly half of the reported wrong site surgeries and occur at least once in approximately 50% of spine surgeons at some point in their career.^{15,16} Unaddressed instability can be an

additional source of worsening pain after the index procedure. Without assessing dynamic X-rays and for fluid in the facet joints on axial imaging, incidence rates up to 28% have been reported for missed degenerative spondylolisthesis.¹⁷

Patients presenting with worsening or different pain since surgery should be evaluated for implant malposition, whereby iatrogenic neural injury or impingement occurs. A CT scan can better assess for malpositioned pedicle screws causing nerve root irritation (Fig. 1). The reported incidence of malpositioned pedicle screws is widely variable.^{18,19} However, it seems that the incidence of symptomatic pedicle screw breach is low, as one review reported only 2 of 45 malpositioned pedicle screws were symptomatic causing radiculopathy.²⁰

Surgery to address axial spine pain, in the absence of instability, has a higher likelihood of not responding to the intervention. Preoperative expectations certainly affect how patients perceive their response to surgery. Finally, even if indicated pathology was addressed and the procedure performed technically well, certain patient characteristics can lead to a poor response immediately following surgery. The characteristics (modifiable and not) that seem to affect immediate postoperative outcomes include a high preoperative opioid burden, poorly controlled diabetes, psychiatric disorders such as anxiety or depression, and workers' compensation status.^{21–26} Assessing a patient's psychosocial history is just as crucial as assessing spine pathology and symptoms and these characteristics should be carefully screened for and optimized before considering revision surgery.

2.1.2. Brief period of improvement, symptomatic by 1 year

The differential for patients that initially respond but become symptomatic by 1 year broadens substantially. Potential diagnoses include recurrent disc herniation, instrumentation failure and or proximal junctional kyphosis, surgical site infection, and pseudarthrosis if a fusion surgery was performed (discussed further below).¹³ The evaluation is similar to the index procedure, including a detailed documentation of the patients symptoms and exam findings, along with imaging and laboratory diagnostic workup.

2.1.3. Improvement beyond 1 year before becoming symptomatic

Patients that are over a year out from surgery prior to experiencing recurrence of symptoms can include diagnosis already discussed but is the minimum time frame needed for adjacent-level degeneration (ALD), which can present a complex diagnostic and treatment challenge. Differentiating adjacent level degeneration (radiographic findings) from adjacent level disease (symptomatic adjacent disease) is important, as radiographic findings do not necessarily correlate with clinically relevant adjacent level disease.²⁷

2.2. Presenting symptoms

A detailed discussion of the patient's symptoms is essential in developing an effective treatment plan. Differentiating axial back and neck pain from radicular arm or leg pain is critical in identifying the pain generators. Failure to carefully review the patient's symptoms risks treating asymptomatic imaging findings and another poor outcome. Patients presenting with continuing or new radicular arm or leg pain may be suffering

from either inadequate initial decompression or recurrent pathology such as a repeat disc herniation. Axial back or neck pain presents a less clear picture and could be related to physical deconditioning after surgery or the surgery itself. Axial pain may also be related to either facet arthropathy, discogenic pain, or even instability in those who present on a delayed basis.¹³ Other causes of pain that mimic spinal pathology must also be considered based on a patient's medical history including diabetic neuropathies, vascular claudication, systemic inflammatory conditions, and shoulder or hip osteoarthritis (OA). There are also reports of complex regional pain syndrome following spine surgery, which must also be considered.²⁸

2.3. Physical exam

In addition to the physical exam that should be performed during an initial evaluation, examination of patients with recurrent symptoms should pay special attention to seek out sources of pain unrelated to their spine pathology. Concurrent hip OA and degenerative lumbar spinal stenosis (DLSS), otherwise known as hip-spine syndrome, can present a challenging clinical picture. Many symptoms of hip OA and DLSS overlap including groin pain or pain radiating beyond the knee.²⁹ On exam, reproduction of symptoms with weight-bearing or hip manipulation is more consistent with hip OA. Groin pain elicited by hip internal rotation has been shown to be sensitive and specific for hip pathology.³⁰ Similarly, considerable overlap exists in symptoms of shoulder pathology and cervical spine disease. A careful shoulder exam in conjunction with adjunctive tests including selective corticosteroid injections will assist in establishing a diagnosis.³¹ When exam findings are inconsistent with anatomy, Waddell signs should be elicited, as these are known to be poor predictors of outcome

regardless of pathology.³² Finally, the surgical wound should always be evaluated for drainage, dehiscence, swelling, or erythema that may be suggestive for a post-operative infection. However, lack of these findings does not rule out infection to any degree.

2.4. Laboratory evaluation

Laboratory studies may be necessary especially in those with concern for a postoperative spinal wound infection. Initial infection workup includes white blood cell count (WBC) with differential, C-reactive protein (CRP), and erythrocyte sedimentation rate. Following spine surgery, CRP typically returns to normal within 14 days, while ESR can take 6 weeks or more.³³ Thus, CRP has largely replaced ESR due to its superior sensitivity in assessing for infection and monitoring treatment response.³⁴ While WBC in general has low sensitivity for detecting postoperative spine infection, lymphopenia in the immediate post-op period (day 4) can represent an increased susceptibility to infection.³⁵ In the setting of pseudarthrosis, a more comprehensive metabolic laboratory workup should be performed (see below).

2.5. Imaging evaluation

Imaging studies typically should begin with plain radiographs of the prior surgical site as well as site of any axial back or neck pain. Flexion-extension radiographs may provide valuable evidence in assessment of instability and to assess a prior fusion mass.³⁶ Implants should be carefully scrutinized for malpositioning, broken or loosening, and subsidence. Hardware loosening may be indicative of pseudarthrosis or hardware infection. The presence of instability as noted by haloing about the screws or movement between vertebrae (dynamic

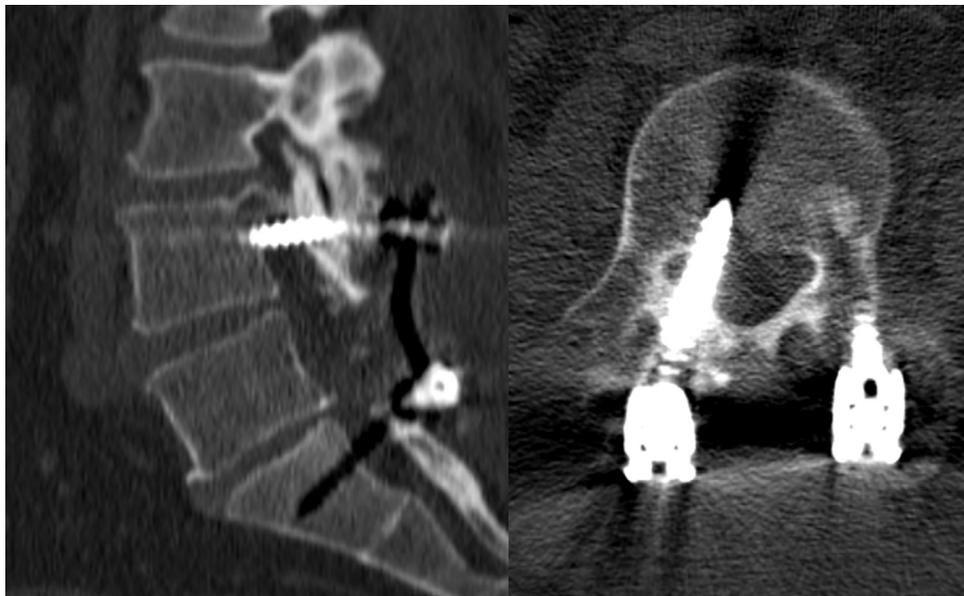


Fig. 1 – Malpositioned screw causing radiculopathy. Patient had worsening right L4 radiculopathy following an L4-S1 fusion with no pain-free interval. CT scan was obtained and showed malpositioned right L4 pedicle screw through the lateral aspect of the central canal.

pseudarthrosis) predicts that revision surgery will more likely provide benefit to the patient. A static pseudarthrosis, whereby a CT does not show solid bridging bone, yet the hardware is well engaged and no movement exists likely has scar tissue preventing movement and is less likely to benefit from revision surgery. Computed tomography (CT) is considered the gold standard for assessing fusion status and can more clearly demonstrate a bony nonunion.³⁷ Assessment of the neural elements may be indicated if there is concern for compressive pathology. Magnetic resonance imaging (MRI) with and without gadolinium enhancement is the most sensitive test in this setting given its ability to distinguish new pathology from prior epidural scar.³⁸ Furthermore, MRI with and without contrast is considered the most useful study in the evaluation for possible postoperative spinal infection. Other diagnostic modalities including electromyograms (EMGs) and nerve conduction velocity (NCV) studies may be useful to evaluate for extraspinal neural compression and exclude other etiologies such as peripheral neuropathy. While selective nerve root or facet injections can be both therapeutic as well as diagnostic, there is concern that patients who have undergone prior surgery may have soft tissue fibrosis and septations that can limit the effectiveness of such treatments.³⁹

3. Common diagnoses in revision spine surgery

3.1. Inadequate decompression or recurrent disc herniation

Recurrent leg pain requiring revision spine surgery is most commonly due to recurrent disc herniation or residual stenosis (inadequate decompression). The incidence rate for recurrent lumbar disc herniation has been estimated from 5% to 15%.⁴⁰ Rates of revision surgery for residual cervical stenosis have been reported to range from 6.6% to 9.2%.^{41,42} Patients with recurrent disc herniations will likely report an initial improvement after their index surgery followed by recurrence

of pain or new pain. In contrast, patients presenting with residual stenosis or inadequate decompression are unlikely to achieve any meaningful improvement in their symptoms after their index surgery. In either scenario, symptoms will likely be similar those that led to their initial surgery. MRI with and without gadolinium enhancement is the study of choice when evaluating for recurrent disc herniations or residual neural compression in the setting of previous spine surgery (Fig. 2).⁴³ Epidural scar will enhance with contrast, while recurrent disc herniations and hypertrophic ligamentum flavum are non-enhancing. CT myelogram can be obtained if the patient is unable to obtain an adequate MRI. Due to scar tissue and septations within the soft tissues after a spine surgery, diagnostic nerve root injections may be less reliable in the postoperative patient.

Treatment recommendations should be similar in the setting of primary pathology typically with a trial of nonsurgical therapy followed by surgical management if the patient's symptoms are refractory to non-operative modalities. In the setting of revision surgery where a repeat or more extensive decompression is needed, a fusion procedure may be considered if not already performed at the index procedure. Some studies have shown that performing spinal fusion at the time of first reoperation can improve patient outcomes.⁴⁴ However, other recent studies have shown repeat discectomy versus discectomy and fusion for the treatment of recurrent lumbar disc herniation has demonstrated similar short-term clinical outcomes with shorter operative times and length of stay and significantly lower hospital charges for the repeat discectomy group.⁴⁵

3.2. Adjacent-level degeneration & disease

Adjacent-level degeneration refers to the radiographic changes while adjacent-level disease (ALD) refers to symptomatic changes to the adjacent level following spine surgery. Spine surgery results in anatomic, biomechanical, and physiologic changes to the operative site and adjacent segments and the influence a spine fusion procedure has on the normal age-related

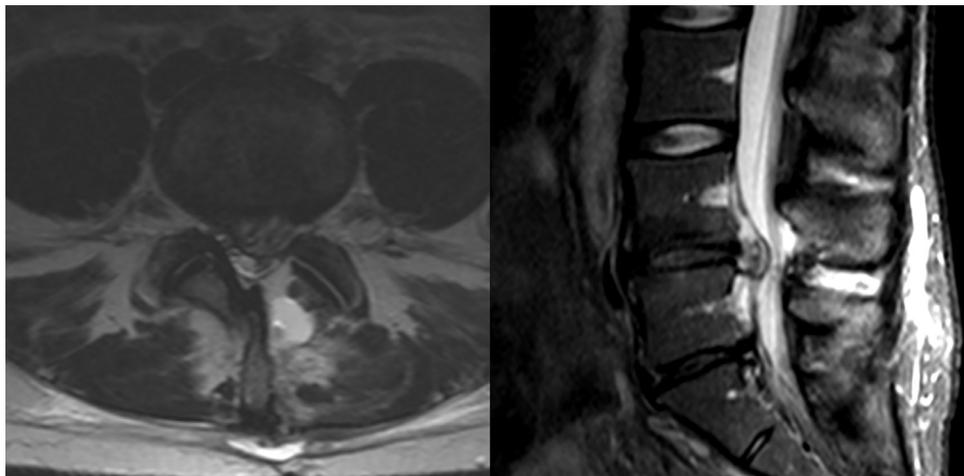


Fig. 2 – Recurrent disc herniation. Patient had undergone left L4 hemilaminectomy and microdiscectomy with some initial improvement in symptoms. However, the patient soon presented with recurrence of symptoms and MRI was obtained showing recurrent paracentral left L4-5 disc protrusion.

degenerative process is unclear.⁴⁶ Adjacent-level degeneration covers many changes including adjacent disc degeneration or herniation, stenosis, listhesis, scoliosis and kyphosis, proximal junctional failure (PJK), sagittal imbalance, instability, facet degeneration, and spondylolysis. Incidence rates of ALD depend on the surgical site, approach, number of levels, and procedure performed. For example, the annual incidence of adjacent-level disease after a lumbar fusion has been reported at 1.7% for single-level fusion, 3.6% after a 2-level fusion, and 5% after 3- or 4-level fusions.⁴⁷ A review of non-fusion patients at 10-year follow-up found a reoperation rate of 11% of patients who had initially undergone laminectomy. Adding a fusion to laminectomy for the treatment of spinal stenosis appears to increase the 10-year reoperation rate.⁴⁸ ALD after ACDF ranges from 0.8% to 2.9%.^{49,50} Patients with ALD will typically report short-term positive outcomes, with progressively worsening symptoms over a year after surgery. Many risk factors have been reported including length and type of fusion, ending the fusion at L5, excessive distraction after posterior lumbar interbody fusion, laminectomy adjacent to a fusion, pelvic incidence/lumbar lordosis mismatch, sagittal alignment, and iatrogenic muscle damage to name a few. Rates of ALD and difficulty in treating ALD have prompted new techniques such as total-disc replacement (TDR) and minimally invasive surgery (MIS) in attempts to preserve biomechanics and physiology. However, while radiographic adjacent level degeneration appears to be lessened, reoperation rates between fusion and TDR and between open versus MIS appear similar.^{51,52}

Initial workup should include dynamic flexion-extension x-rays and standing full length spine x-rays to assess sagittal balance. As mentioned previously, MRI with and without gadolinium is the study of choice for evaluating adjacent neurologic compression and CT scan will best identify adjacent facet arthropathy, spondylolysis and pedicle stress fractures. Nonoperative management can be attempted but often revision and extension of the fusion is necessary. To minimize future recurrent ALD, revision surgeries for ALD should be performed by surgeons who are experts in revision techniques.

3.3. Complications of prior surgery

3.3.1. Pseudarthrosis

Pseudarthrosis remains one of the most common complications of fusion spine surgery, and one of the most common indications for revision spine surgery.^{53–55} Given that fusion surgeries have been on the rise over the last several decades, pseudarthrosis should always be considered when evaluating a patient with new or ongoing symptoms and a history of prior fusion surgery. Patients presenting with pseudarthrosis can have a variable clinical presentation with recurrent or persistent neck, back, arm or leg pain. A thorough history may reveal patient or surgical risk factors for pseudarthrosis. A detailed understanding of the index procedure including graft selection is important in understanding the patient's risk of pseudarthrosis. A systematic review of the literature concluded that iliac crest autograft appears to be the gold standard for fusion, while an interbody cage is the gold standard when strictly evaluating complication rates.⁵⁶ Other patient and surgical risk factors for pseudarthrosis include smoking, increasing body mass index, increasing ODI/NDI scores, younger age, vitamin D deficiency, history of non-instrumented fusion, non-circumferential thoracolumbar fusion and long segment fusion in spinal deformity.^{1,57–62} While the gold standard for diagnosing pseudarthrosis remains surgical exploration, noninvasive methods can help assess the fusion status. Evaluation begins with dynamic x-rays of the affected area and basic laboratory work-up. Infection and adjacent segment disease should always be ruled out. For diagnosing anterior cervical pseudarthrosis, the method reported by Riew *et al.*, has demonstrated a sensitivity and specificity similar to CT scan. By their method, flexion and extension radiographs are magnified to 150% and interspinous motion is measured, with a cutoff of ≥ 1 mm for interspinous motion indicative of a pseudarthrosis, with at least ≥ 4 mm of superjacent interspinous motion to constitute adequate dynamic motion (Fig. 3).⁶³ When evaluating the lumbar spine or cervical spine without adequate dynamic x-rays, CT remains the preferred method.³⁶ However, these two modalities should be

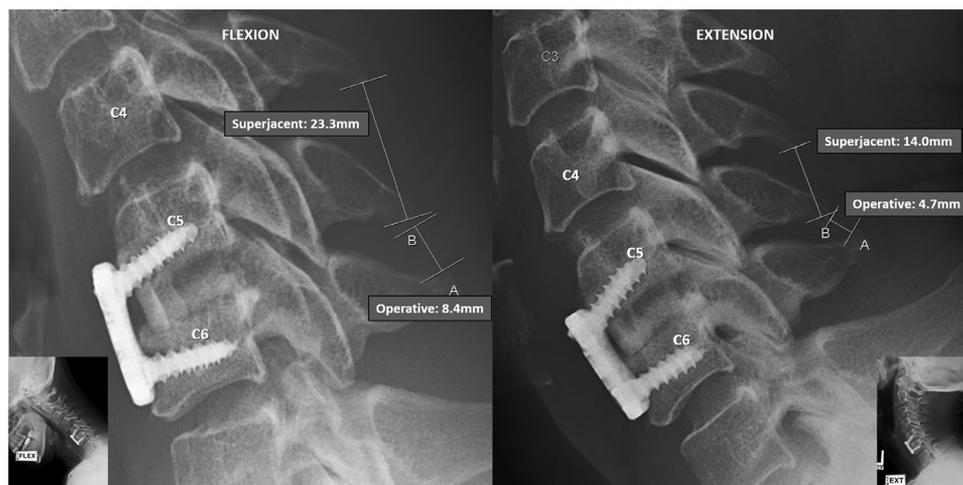


Fig. 3 – Cervical pseudarthrosis on dynamic x-rays as described by Riew *et al.* Flexion and extension cervical radiographs are magnified to 150% on the same monitor, and the interspinous motion of the superjacent level (C4-5) was measured and found to be 9.3 mm (23.3 mm – 14 mm), indicating adequate dynamic x-rays had been obtained (i.e. >4 mm of motion). The interspinous motion at the operative level (C5-6) was found to be greater than the 1 mm cutoff at 3.7 mm (8.4 mm – 4.7 mm), indicating excessive motion at the fusion site, consistent with a pseudarthrosis.

used in conjunction to differentiate a dynamic from a static pseudarthrosis.

Once a diagnosis of pseudarthrosis is suspected, a careful investigation into the patient's risk factors for pseudarthrosis should be performed and optimized prior to revision surgery. Vitamin D insufficiency (<30 ng/mL) and deficiency (<20 ng/mL) rates in all elective spine patients has been reported up to 57% and 30%, respectively.^{58,64} Additionally, Vitamin D deficiency has been identified as an independent predictor of pseudarthrosis.⁵⁷ Patients should be evaluated for osteoporosis, as recent studies have shown teriparatide to promote bone formation in osteoporotic spine fusion and osteoporotic fracture patients.^{65–67} Management of comorbidities such as diabetes mellitus, obesity, and tobacco use should be optimized. For these reasons, a bone metabolic evaluation by a specialist should be considered in patients preparing for revision surgery for pseudarthrosis.

3.3.2. Instrumentation & surgical construct failure

Instrumentation or construct failure covers a wide range of complications including screw bending, breakage, loosening, or contributing to a new neurologic complication. Rate of instrumentation or surgical construct failure have been reported from 0.5% up to 20% depending on the technique, location in spine, surgical approach, graft choice, length of construct and type of instrumentation.^{68–71} Persistent leg or arm pain in the same dermatome as the surgical site may be due to a malpositioned screw causing nerve root irritation. New or persistent neck or back pain may be due to painful or broken instrumentation. Timing can be variable, as an analysis of instrumentation failure after adult spinal deformity surgery showed a peak at 12–24 months and again at ≥ 24 months.⁷²

Patients with implant failure may be asymptomatic or present with severe pain and progressive neurological deficit, and with variable timing. Failed instrumentation is often detected on routine x-rays or for initial workup of symptoms. In symptomatic patients, other sources should be carefully investigated. Infection, pseudarthrosis, inadequate decompression, nerve root injury, physical deconditioning, incorrect diagnosis at the index operation, and adjacent segment disease should all be ruled out before attributing symptoms purely to implant failure. Reviewing the implants used in the index operation and understanding the biomechanics of the implant is important to understanding the reason for failure and for selecting appropriate revision implants. A mismatch in the modulus of elasticity between the cage and host bone will inevitably lead to cage subsidence.

A review of patients treated with anterior cervical locking plate found an implant-related complication rate of 10.7%. Complications included oblique plating, screw malposition into or excessively close to the disc space, screw or plate loosening or breaking, esophageal perforations and overly long plates causing adjacent level degeneration and ultimately revision surgery.⁷³ All of these complications are prone to pseudarthrosis. While rare, esophageal perforation is a well-known complication of anterior cervical plating and potentially fatal. If the anterior cervical plate or screws become dislodged, loosened, or symptomatic (e.g. dysphagia), the implants should be promptly removed.⁶⁹

3.3.3. Infection

Post-operative spine infections most commonly present with increased pain after a pain-free interval following their index operation, and usually within 1 year. Estimates on the incidence of late post-operative spine infections, those presenting after 12 months, are generally less than 1–2%.^{74–76} Reports of infection after anterior cervical surgery are especially low (0%–0.39%), and thus any postoperative anterior cervical spine patient presenting with new pain, fevers, wound drainage, or dysphagia with elevated inflammatory markers should have an esophageal perforation ruled out by barium esophagogram.⁷⁷ Risk factors for postoperative spine infection include patient risk factors (diabetes mellitus, smoking, malnutrition, obesity, older age, immunomodulators used in systemic arthritis's, chronic steroid exposure), and surgical risk factors (fusion surgery, posterior procedures, longer surgical time and higher blood loss).⁷⁴ The most common causative organism of postoperative spinal infection is *Staphylococcus aureus*.⁷⁶ Physical exam can be normal but peri-incisional erythema, tenderness, induration or drainage all certainly suggest infection. Diagnostic workup includes laboratory and imaging. C-reactive protein is the most sensitive indicator of postoperative spine infection, whereas white blood cell count is typically unreliable.³⁴ Plain x-rays may show signs of disc space narrowing in setting of discitis or instrumentation loosening, but advanced imaging with MRI with gadolinium contrast is the most useful modality when evaluating for infection. Findings on MRI suggestive of infection include rim enhancing fluid collections, epidural collections, bony destruction, and progressive marrow signal changes. Still, the gold standard for diagnosing surgical infection is tissue culture. A CT-guided biopsy can be considered before proceeding with revision surgery. F-fluoro-d-deoxyglucose positron emission tomography (PET) scans have shown to be a useful adjunct in postoperative infections when MRI is not an option, however other nuclear medicine studies have been less helpful.⁷⁸

When considering a revision surgery, the presentation and diagnostic workup in patients with latent infection such as from *P. acnes* may be consistent with the diagnosis of pseudarthrosis or construct failure. A 10-year review of infections after instrumented fusion surgery found 17% of CRP values, 45% of ESR, and 95% of WBC values to be normal, and found *P. acnes* to be the causative organism in just over half of all cases.⁷⁹ Thus, it is always important to consider latent infection and if intraoperative cultures are obtained, they should be incubated for a minimum 13 days.⁸⁰

4. Conclusions

Outcomes following revision spine surgery are less favorable than primary surgery. Careful patient selection and accurate diagnosis for primary surgery is the best method for preventing revision surgery. When patients with a prior spine surgery present with new or recurrent symptoms, it is helpful to develop a differential based on their symptom response following their index surgery. Patients who achieved no relief after their index surgery likely had an undiagnosed surgery (including incorrect preoperative diagnosis), technically

flawed procedure, or were a poor surgical candidate. Reasons for recurrent symptoms after initial relief include recurrent disc herniation, pseudarthrosis, surgical construct or implant failure, postoperative infection, and adjacent level disease. A strategic evaluation to establish a diagnosis consistent with the patient's history, symptoms, laboratory and imaging findings, along with careful consideration of patient factors, is critical to avoid a repeat failed spine surgery.

5. Disclosures

The authors report no proprietary or commercial interest in any product mentioned or concept discussed in this article.

REFERENCES

- Fischgrund JS, Mackay M, Herkowitz HN, et al. 1997 Volvo Award winner in clinical studies. Degenerative lumbar spondylolisthesis with spinal stenosis: a prospective, randomized study comparing decompressive laminectomy and arthrodesis with and without spinal instrumentation. *Spine (Phila Pa 1976)*. 1997;22:2807–2812.
- Hacker RJ, Cauthen JC, Gilbert TJ, et al. A prospective randomized multicenter clinical evaluation of an anterior cervical fusion cage. *Spine (Phila Pa 1976)*. 2000;25:2646–2654. discussion 55.
- Snider RK. A prospective, randomized study of lumbar fusion: preliminary results. *Spine (Phila Pa 1976)*. 1994;19:109.
- Rajaei SS, Kanim LE, Bae HW. National trends in revision spinal fusion in the USA: patient characteristics and complications. *Bone Joint J*. 2014;96-B:807–816.
- Malter AD, McNeney B, Loeser JD, et al. 5-year reoperation rates after different types of lumbar spine surgery. *Spine (Phila Pa 1976)*. 1998;23:814–820.
- Singh K, Phillips FM, Park DK, et al. Factors affecting reoperations after anterior cervical discectomy and fusion within and outside of a Federal Drug Administration investigational device exemption cervical disc replacement trial. *Spine J*. 2012;12:372–378.
- Highsmith JM, Dhall SS, Haid Jr. RW, et al. Treatment of cervical stenotic myelopathy: a cost and outcome comparison of laminoplasty versus laminectomy and lateral mass fusion. *J Neurosurg Spine*. 2011;14:619–625.
- Wang TY, Lubelski D, Abdullah KG, et al. Rates of anterior cervical discectomy and fusion after initial posterior cervical foraminotomy. *Spine J*. 2015;15:971–976.
- Koerner JD, Kepler CK, Albert TJ. Revision surgery for failed cervical spine reconstruction: review article. *HSS J*. 2015;11:2–8.
- Veeravagu A, Cole T, Jiang B, et al. Revision rates and complication incidence in single- and multilevel anterior cervical discectomy and fusion procedures: an administrative database study. *Spine J*. 2014;14:1125–1131.
- Onesti ST. Failed back syndrome. *Neurologist*. 2004;10:259–264.
- Aleem IS, Duncan J, Ahmed AM, et al. Do lumbar decompression and fusion patients recall their preoperative status?: A cohort study of recall bias in patient-reported outcomes. *Spine (Phila Pa 1976)*. 2017;42:128–134.
- Guyer RD, Patterson M, Ohnmeiss DD. Failed back surgery syndrome: diagnostic evaluation. *J Am Acad Orthop Surg*. 2006;14:534–543.
- Dagi TF, Tarkington MA, Leech JJ. Tandem lumbar and cervical spinal stenosis. Natural history, prognostic indices, and results after surgical decompression. *J Neurosurg*. 1987;66:842–849.
- James MA, Seiler 3rd JG, Harrast JJ, et al. The occurrence of wrong-site surgery self-reported by candidates for certification by the American Board of Orthopaedic Surgery. *J Bone Joint Surg Am*. 2012;94(1-12):e2.
- Grimm BD, Laxer EB, Blessinger BJ, et al. Wrong-level spine surgery. *JBJS Rev*. 2014;2.
- Segebarth B, Kurd MF, Haug PH, et al. Routine upright imaging for evaluating degenerative lumbar stenosis: incidence of degenerative spondylolisthesis missed on supine MRI. *J Spinal Disord Tech*. 2015;28:394–397.
- Kowalski JM, Ludwig SC, Hutton WC, et al. Cervical spine pedicle screws: a biomechanical comparison of two insertion techniques. *Spine (Phila Pa 1976)*. 2000;25:2865–2867.
- Ludwig SC, Kramer DL, Balderston RA, et al. Placement of pedicle screws in the human cadaveric cervical spine: comparative accuracy of three techniques. *Spine (Phila Pa 1976)*. 2000;25:1655–1667.
- Abumi K, Shono Y, Ito M, et al. Complications of pedicle screw fixation in reconstructive surgery of the cervical spine. *Spine (Phila Pa 1976)*. 2000;25:962–969.
- Wick JB, Sivaganesan A, Chotai S, et al. Is there a preoperative morphine equianalgesic dose that predicts ability to achieve a clinically meaningful improvement following spine surgery? *Neurosurgery*. 2018;83:245–251.
- Lee D, Armaghani S, Archer KR, et al. Preoperative opioid use as a predictor of adverse postoperative self-reported outcomes in patients undergoing spine surgery. *J Bone Joint Surg Am*. 2014;96:e89.
- Armaghani SJ, Lee DS, Bible JE, et al. Preoperative narcotic use and its relation to depression and anxiety in patients undergoing spine surgery. *Spine (Phila Pa 1976)*. 2013;38:2196–2200.
- Scuderi GJ, Sherman AL, Brusovanik GV, et al. Symptomatic cervical disc herniation following a motor vehicle collision: return to work comparative study of workers' compensation versus personal injury insurance status. *Spine J*. 2005;5:639–644. discussion 44.
- Wong JJ, Cote P, Quesnele JJ, et al. The course and prognostic factors of symptomatic cervical disc herniation with radiculopathy: a systematic review of the literature. *Spine J*. 2014;14:1781–1789.
- Anderson PA, Subach BR, Riew KD. Predictors of outcome after anterior cervical discectomy and fusion: a multivariate analysis. *Spine*. 2009;34:161–166.
- Mannion AF, Leivseth G, Brox JJ, et al. ISSLS Prize winner: long-term follow-up suggests spinal fusion is associated with increased adjacent segment disc degeneration but without influence on clinical outcome: results of a combined follow-up from 4 randomized controlled trials. *Spine (Phila Pa 1976)*. 2014;39:1373–1383.
- Sachs BL, Zindrick MR, Beasley RD. Reflex sympathetic dystrophy after operative procedures on the lumbar spine. *J Bone Joint Surg Am*. 1993;75:721–725.
- Devin CJ, McCullough KA, Morris BJ, et al. Hip-spine syndrome. *J Am Acad Orthop Surg*. 2012;20:434–442.
- Brown MD, Gomez-Marin O, Brookfield KF, et al. Differential diagnosis of hip disease versus spine disease. *Clin Orthop Relat Res*. 2004: 280–284.
- Throckmorton TQ, Kraemer P, Kuhn JE, et al. Differentiating cervical spine and shoulder pathology: common disorders and key points of evaluation and treatment. *Instr Course Lect*. 2014;63:401–408.
- Waddell G, McCulloch JA, Kummel E, et al. Nonorganic physical signs in low-back pain. *Spine (Phila Pa 1976)*. 1980;5:117–125.
- Thelander U, Larsson S. Quantitation of C-reactive protein levels and erythrocyte sedimentation rate after spinal surgery. *Spine (Phila Pa 1976)*. 1992;17:400–404.
- Silber JS, Anderson DG, Vaccaro AR, et al. Management of postprocedural discitis. *Spine J*. 2002;2:279–287.

35. Takahashi J, Shono Y, Hirabayashi H, et al. Usefulness of white blood cell differential for early diagnosis of surgical wound infection following spinal instrumentation surgery. *Spine (Phila Pa 1976)*. 2006;31:1020–1025.
36. Gruskay JA, Webb ML, Grauer JN. Methods of evaluating lumbar and cervical fusion. *Spine J*. 2014;14:531–539.
37. Ghiselli G, Wharton N, Hipp JA, et al. Prospective analysis of imaging prediction of pseudarthrosis after anterior cervical discectomy and fusion: computed tomography versus flexion-extension motion analysis with intraoperative correlation. *Spine*. 2011;36:463–468.
38. Ross JS, Masaryk TJ, Schrader M, et al. MR imaging of the post-operative lumbar spine: assessment with gadopentetate dimeglumine. *AJNR Am J Neuroradiol*. 1990;11:771–776.
39. Rahimzadeh P, Sharma V, Imani F, et al. Adjuvant hyaluronidase to epidural steroid improves the quality of analgesia in failed back surgery syndrome: a prospective randomized clinical trial. *Pain Phys*. 2014;17:E75–E82.
40. Lee JK, Amorosa L, Cho SK, et al. Recurrent lumbar disk herniation. *J Am Acad Orthop Surg*. 2010;18:327–337.
41. Liu G, Buchowski JM, Bunmaprasert T, et al. Revision surgery following cervical laminoplasty: etiology and treatment strategies. *Spine (Phila Pa 1976)*. 2009;34:2760–2768.
42. Bydon M, Mathios D, Macki M, et al. Long-term patient outcomes after posterior cervical foraminotomy: an analysis of 151 cases. *J Neurosurg Spine*. 2014;21:727–731.
43. Ross JS, Masaryk TJ, Schrader M, et al. MR imaging of the post-operative lumbar spine: assessment with gadopentetate dimeglumine. *AJR Am J Roentgenol*. 1990;155:867–872.
44. Osterman H, Sund R, Seitsalo S, et al. Risk of multiple reoperations after lumbar discectomy: a population-based study. *Spine*. 2003;28:621–627.
45. Guan J, Ravindra VM, Schmidt MH, et al. Comparing clinical outcomes of repeat discectomy versus fusion for recurrent disc herniation utilizing the N2QOD. *J Neurosurg Spine*. 2017;26:39–44.
46. Hilibrand AS, Robbins M. Adjacent segment degeneration and adjacent segment disease: the consequences of spinal fusion? *Spine J*. 2004;4:190S–194S.
47. Sears WR, Sergides IG, Kazemi N, et al. Incidence and prevalence of surgery at segments adjacent to a previous posterior lumbar arthrodesis. *Spine J*. 2011;11:11–20.
48. Katz JN, Lipson SJ, Chang LC, et al. Seven- to 10-year outcome of decompressive surgery for degenerative lumbar spinal stenosis. *Spine (Phila Pa 1976)*. 1996;21:92–98.
49. Hilibrand AS, Carlson GD, Palumbo MA, et al. Radiculopathy and myelopathy at segments adjacent to the site of a previous anterior cervical arthrodesis. *J Bone Joint Surg Am*. 1999;81:519–528.
50. Wu JC, Liu L, Wen-Cheng H, et al. The incidence of adjacent segment disease requiring surgery after anterior cervical discectomy and fusion: estimation using an 11-year comprehensive nationwide database in Taiwan. *Neurosurgery*. 2012;70:594–601.
51. Zigler JE, Glenn J, Delamarter RB. Five-year adjacent-level degenerative changes in patients with single-level disease treated using lumbar total disc replacement with ProDisc-L versus circumferential fusion. *J Neurosurg Spine*. 2012;17:504–511.
52. Radcliff KE, Kepler CK, Maaieh M, et al. What is the rate of lumbar adjacent segment disease after percutaneous versus open fusion? *Orthop Surg*. 2014;6:118–120.
53. Hu RW, Jaglal S, Axcell T, et al. A population-based study of reoperations after back surgery. *Spine (Phila Pa 1976)*. 1997;22:2265–2270. discussion 71.
54. Martin BI, Mirza SK, Comstock BA, et al. Reoperation rates following lumbar spine surgery and the influence of spinal fusion procedures. *Spine (Phila Pa 1976)*. 2007;32:382–387.
55. Raizman NM, O'Brien JR, Poehling-Monaghan KL, et al. Pseudarthrosis of the spine. *J Am Acad Orthop Surg*. 2009;17:494–503.
56. Jacobs W, Willems PC, Kruyt M, et al. Systematic review of anterior interbody fusion techniques for single- and double-level cervical degenerative disc disease. *Spine (Phila Pa 1976)*. 2011;36:E950–E960.
57. Ravindra VM, Godzik J, Dailey AT, et al. Vitamin D levels and 1-year fusion outcomes in elective spine surgery: a prospective observational study. *Spine (Phila Pa 1976)*. 2015;40:1536–1541.
58. Stoker GE, Buchowski JM, Bridwell KH, et al. Preoperative vitamin D status of adults undergoing surgical spinal fusion. *Spine (Phila Pa 1976)*. 2013;38:507–515.
59. Brown CW, Orme TJ, Richardson HD. The rate of pseudarthrosis (surgical nonunion) in patients who are smokers and patients who are nonsmokers: a comparison study. *Spine (Phila Pa 1976)*. 1986;11:942–943.
60. Hilibrand AS, Fye MA, Emery SE, et al. Impact of smoking on the outcome of anterior cervical arthrodesis with interbody or strut-grafting. *J Bone Joint Surg Am*. 2001;83-A:668–673.
61. Kim YJ, Bridwell KH, Lenke LG, et al. Pseudarthrosis in long adult spinal deformity instrumentation and fusion to the sacrum: prevalence and risk factor analysis of 144 cases. *Spine (Phila Pa 1976)*. 2006;31:2329–2336.
62. Bono CM, Lee CK. Critical analysis of trends in fusion for degenerative disc disease over the past 20 years: influence of technique on fusion rate and clinical outcome. *Spine (Phila Pa 1976)*. 2004;29:455–463. discussion Z5.
63. Song KS, Piyaskulkaew C, Chuntarapas T, et al. Dynamic radiographic criteria for detecting pseudarthrosis following anterior cervical arthrodesis. *J Bone Joint Surg Am*. 2014;96:557–563.
64. Ravindra VM, Godzik J, Guan J, et al. Prevalence of Vitamin D deficiency in patients undergoing elective spine surgery: a cross-sectional analysis. *World Neurosurg*. 2015;83:1114–1119.
65. Chaudhary N, Lee JS, Wu JY, et al. Evidence for use of teriparatide in spinal fusion surgery in osteoporotic patients. *World Neurosurg*. 2017;100:551–556.
66. Lecoultre J, Stoll D, Chevalley F, et al. [Improvement of fracture healing with teriparatide: series of 22 cases and review of the literature]. *Rev Med Suisse*. 2015;11:663–667.
67. Ebata S, Takahashi J, Hasegawa T, et al. Role of weekly teriparatide administration in osseous union enhancement within six months after posterior or transforaminal lumbar interbody fusion for osteoporosis-associated lumbar degenerative disorders: a multicenter, prospective randomized study. *J Bone Joint Surg Am*. 2017;99:365–372.
68. Clarke MJ, Ecker RD, Krauss WE, et al. Same-segment and adjacent-segment disease following posterior cervical foraminotomy. *J Neurosurg Spine*. 2007;6:5–9.
69. Lowery GL, McDonough RF. The significance of hardware failure in anterior cervical plate fixation. Patients with 2- to 7-year follow-up. *Spine (Phila Pa 1976)*. 1998;23:181–186. discussion 6–7.
70. Lonstein JE, Denis F, Perra JH, et al. Complications associated with pedicle screws. *J Bone Joint Surg Am*. 1999;81:1519–1528.
71. McAfee PC, Weiland DJ, Carlow JJ. Survivorship analysis of pedicle spinal instrumentation. *Spine (Phila Pa 1976)*. 1991;16:S422–S427.
72. Daniels AH, Bess S, Line B, et al. Peak timing for complications following adult spinal deformity surgery. *World Neurosurg*. 2018.
73. Ning X, Wen Y, Xiao-Jian Y, et al. Anterior cervical locking plate-related complications; prevention and treatment recommendations. *Int Orthop*. 2008;32:649–655.

74. Meredith DS, Kepler CK, Huang RC, et al. Postoperative infections of the lumbar spine: presentation and management. *Int Orthop*. 2012;36:439–444.
75. Kuo CH, Wang ST, Yu WK, et al. Postoperative spinal deep wound infection: a six-year review of 3230 selective procedures. *J Chin Med Assoc*. 2004;67:398–402.
76. Weinstein MA, McCabe JP, Cammisa Jr. FP. Postoperative spinal wound infection: a review of 2,391 consecutive index procedures. *J Spinal Disord*. 2000;13:422–426.
77. Ghobrial GM, Harrop JS, Sasso RC, et al. Anterior cervical infection: presentation and incidence of an uncommon postoperative complication. *Global Spine J*. 2017;7:12S–16S.
78. Gemmel F, Rijk PC, Collins JM, et al. Expanding role of 18F-fluoro-D-deoxyglucose PET and PET/CT in spinal infections. *Eur Spine J*. 2010;19:540–551.
79. Collins I, Wilson-MacDonald J, Chami G, et al. The diagnosis and management of infection following instrumented spinal fusion. *Eur Spine J*. 2008;17:445–450.
80. Butler-Wu SM, Burns EM, Pottinger PS, et al. Optimization of periprosthetic culture for diagnosis of *Propionibacterium acnes* prosthetic joint infection. *J Clin Microbiol*. 2011;49:2490–2495.