Although the efficacy of posterior decompression for symptomatic lumbar stenosis that is recalcitrant to conservative therapy is well proven, uniform agreement on the need for simultaneous arthrodesis is lacking. The variability in the rate of lumbar fusion with and without instrumentation has been attributed to a number of factors: advances in surgical technique; rapid development of instrumentation; radiographic advances in the diagnosis of disease entities of the lumbar spine; evolution in our understanding of bone healing; improved pre- and postoperative care; aggressive rehabilitation; patient compensation; hospital and surgeon reimbursement; better education of residents, fellows, and practicing neurosurgeons; and, most important, the lack of clear indications based on defined diagnostic categories. Based on review of the literature and their experience at the Barrow Neurological Institute, the authors have attempted to define indications for lumbar fusion with or without instrumentation based on defined diagnostic categories. Clear indications for fusion include trauma, tumor, or infection with two- or three-column injury, iatrogenic instability, and isthmic spondylolisthesis. Relative indications for fusion include degenerative spondylolisthesis, radiographically proven dynamic instability with pain or neurological findings, adult scoliosis, and mechanical back pain. Fusion is rarely indicated with discectomy, abnormal radiographs without appropriate findings (such as degenerative disc disease), facet joint syndrome, failed back surgery, or stable spinal stenosis.

Key Words * lumbar fusion * instrumentation * indication * instability * lumbar stenosis
extensive literature on lumbar fusion has been generated, few prospective studies have evaluated clearly defined diagnostic subgroups.

The rate of lumbar fusion doubled between 1979 and 1990.[33] Understandably, the largest increase occurred in the elderly population. Each year in this country more than 40,000 lumbar fusion procedures, which make up approximately one-fifth of all low-back operations, are performed. Approximately half of the lumbar fusions are combined with placement of instrumentation. As well as increasing the complication rate and the percentage of patients discharged to nursing homes, the addition of fusion to routine decompression increases the hospital and surgical costs by approximately 50%.[33] The addition of instrumentation to the procedure increases the total cost by approximately 100%.

Franklin and colleagues[15] evaluated the outcome of lumbar fusion in patients receiving Workers' Compensation in Washington state. Overall, the outcome of patients undergoing lumbar fusion with and without instrumentation was worse than outcomes published in the literature. Sixty-eight percent of patients reported that their back or leg pain was worse, and 56% reported that overall quality of life had not improved or was worse. However, these percentages must be interpreted cautiously. Ironically, 62% of the patients responded that they would undergo the surgery again. Most of these patients had returned to work and no longer received Workers' Compensation. Psychological factors can also undermine outcome statistics.[3,10,19,62]

A number of variables have been used to evaluate the efficacy of lumbar fusion. In our opinion, comprehensive postoperative assessment must include a thorough and long-term evaluation of radiographically confirmed fusion, complications, pain, function, and patient satisfaction. There is a consensus among most spine surgeons that the outcome of lumbar fusion can be improved with simultaneous progress on a number of fronts. Prospective studies with clearly defined diagnostic categories would probably produce the greatest improvement.[21,24,58,61,67,68] In addition, basic scientific research should address the biomechanics of fusion, the biology of bone healing, and the pathophysiological mechanisms of degenerative bone disease.[4-6,22,52]

**EVOLUTION OF UNDERSTANDING**

The development of spinal fusion is attributed to the separate work of Doctors Fred Albee and Russell Hibbs, published in 1911. Their original work involved the treatment of tuberculosis of the spine but was quickly applied to the treatment of trauma, scoliosis, and tumors. During the past 50 years, fusion has been used to treat degenerative disorders. The use of instrumentation to augment lumbar fusion is relatively recent and has generated controversy over the use of anterior fusion versus posterior fusion as well as the use of both.[9,12,16,37,44,48,54,55,60]

This controversy is typified by the recent report of Suk and coworkers.[60] They described 76 patients who underwent posterolateral fusion with pedicle screws with (36 patients) and without (40 patients) interbody fusion after decompression for spondylolitic spondylolisthesis. The mean follow-up period was 2 years. The rate of nonfusion was 8% in the posterolateral fusion with pedicle screw instrumentation only group and 0% in the posterolateral fusion with pedicle screw instrumentation group with interbody fusion. The reduction in slippage was 28% and 42% and the patient satisfaction rating was 95% and 97%, respectively. Do these data justify the extra cost and risk of adding interbody fusion? Further evaluation of the data, however, reveals that the percentage of patients with an excellent outcome increased from 45% to 75% with the addition of interbody fusion.
Most spine surgeons agree that segmental spinal instability can produce neurological symptoms such as motor weakness, sensory loss, and pain. However, consensus has not been reached about the definition of spinal instability and how to apply the concept to clinical practice. White and Panjabi[63] have defined clinical instability in the spine as "the loss of the ability of the spine under physiological loads to maintain relationships between vertebrae in such a way that there is neither damage or subsequent irritation to the spinal cord or nerve roots, and, in addition, there is not development of incapacitating deformity or pain from structural changes."

Kirkaldy-Willis and Farfan[36] have defined spinal instability as "the clinical status of the patient with back problems who with the least provocation steps from the mildly asymptomatic to the severe episodic." They further define the spectrum of clinical manifestations of degenerative spine disease with three contiguous stages: 1) temporary dysfunction, 2) unstable phase, and 3) stabilization.

Based on clinical observations, analysis of pathological material, and review of the literature, Frymoyer and Selby[20] have defined a classification scheme in which there are four types of spinal instability. Type 1 is defined as axial rotational instability. Farfan and Kirkaldy-Willis[13] reported a fixed rotary deformity in some patients who present with low-back pain on twisting about the spine. Plain radiographs revealed narrowing of the disc space, facet degeneration, malalignment of the spinous processes, and a rotational deformity of the pedicles. Type 2, or translational instability,[41] manifests as episodes of recurrent back pain associated with forward translation at the affected level. On plain radiographs, the disc space is narrowed, but the alignment of the spinous processes and pedicles is normal. Lateral flexion-extension radiographs show angulatory collapse of the disc space and forward subluxation at the affected level. This type of instability typically affects women and is most common at L4-5. Type 3, or retrograde spondylolisthetic instability, is most common at L5-S1 and affects as many as one-third of the patients with low-back pain. Plain lateral radiographs demonstrate posterior translation, collapse of the disc space, and facet subluxation. Radiographic imaging of the neural canal often shows lateral stenosis of the spinal canal. Type 4, or postoperative instability, occurs after aggressive decompressive surgery of the lateral spinal canal. Removal of an entire facet, pars interarticularis, or half of each facet at the same level may produce instability. Lateral radiographs demonstrate anterior spondylolisthesis of the affected level compared with preoperative films.

Many investigators have attempted to define instability solely by using radiological methods. Dynamic imaging of the spine with flexion-extension lateral views has been used most often. In general, anterior spondylolisthesis greater than 3 to 4 mm suggests spinal instability.[18,45,51,59]

Posner and colleagues[53] used a cadaveric model to assess the normal range of motion in lumbar spines subjected to physiological flexion-extension movements. These data were then used to identify the degree of angulation or translation indicative of instability. Panjabi[49] furthered the conceptual framework of biomechanical stability of the lumbar spine by defining an experimental design based on 6ś of freedom: 3ś of translation in the x, y, and z coordinate system and 3ś of rotation in the same coordinate system. This experimental design can be used to assess the strength, fatigue, and stability of fusion and instrumentation protocols for different types of spinal instability.

White and Panjabi[63] use a point-value checklist to define clinical instability in the lumbar spine. The first category is used to assess the structural and functional integrity of the anterior and posterior elements. The second category is used to assess flexion-extension and resting radiographs; translation
greater than 4.5 mm in the sagittal plane on flexion-extension films or rotation greater than 20° in the sagittal plane at L4-5 or 25° at L5-S1 is indicative of instability. The third category is used to assess damage to the cauda equina. Finally, the fourth category is used to consider the potential for dangerous loading of the spine. Points are assessed for each of the four categories, and if the total reaches five, the spine is considered to be unstable.

LUMBAR STENOSIS

At the Barrow Neurological Institute, 206 patients with symptomatic lumbar stenosis were treated initially by decompressive surgery.[50] After a mean follow-up period of 28 months, six patients required fusion procedures. The possible common denominator for these patients was the presence of Grade 2 spondylolisthesis. Consequently, fusion is not routinely combined with lumbar decompression unless Grade 2 or worse spondylolisthesis is coexistent.

ISTHMIC SPONDYLOLISTHESIS

Patients with isthmic spondylolisthesis typically present during their teenage years. At diagnosis, the most important prognostic indicators for future low-back pain are the patient's age and the degree of slippage.[57] When the degree of spondylolisthesis is greater than 25% of the vertebral body width, the risk for developing back pain increases. The incidence of isthmic spondylolisthesis reported in autopsy studies is approximately 5%.

Wiltse, et al.,[64] presented a classification system for spondylolysis and spondylolisthesis. Type 1 (dysplastic lesion) is a congenital abnormality of the upper sacrum or arch of L-5 that allows spondylolisthesis. Type 2, which is discussed in this section, is called isthmic and defines a lesion in the pars interarticularis. Three categories of Type 2 lesions are recognized: 1) lytic-fatigue fracture of the pars, 2) elongated but intact pars, and 3) acute fracture. Type 3 (degenerative spondylolisthesis) is related to long-standing intersegmental instability and is discussed in the next section. Type 4 (traumatic) is related to fractures that are not located in the pars interarticularis. Pathological Type 4 spondylolisthesis can be caused by a generalized or localized bone disorder.

Approximately half of the patients with spondylolysis progress to spondylolisthesis. Some authors have recommended repairing the pars interarticularis defect in spondylolysis or Grade 1 spondylolisthesis. In general, an L5-S1 fusion has been used to treat Grades 1 and 2 spondylolisthesis, and an L4-S1 fusion has been used to treat higher grades of spondylolisthesis.

The indications for lumbar arthrodesis in children and adolescents with spondylolisthesis are controversial. The potential risk of aggressive spondylolisthesis has sometimes been considered an indication for surgery. The treatment of adolescents with moderate-to-severe spondylolisthesis without associated signs or symptoms is also controversial. Both Hensinger[28] and Laurent and Österman[38] recommend that all adolescents with Grade 3 or 4 spondylolisthesis undergo fusion, regardless of their symptomatology. Fusion in this group is also supported by the improved long-term prognosis when arthrodesis is performed in childhood rather than adulthood.[25] Conversely, however, some long-term follow-up studies support conservative treatment of asymptomatic children and teenagers.[2,17] We believe that fusion is indicated when 1) mechanical low-back pain persists or progresses or neurological symptoms are recalcitrant to conservative therapy, 2) spondylolisthesis is symptomatic and high grade, and 3) a child or adolescent has progressive symptoms and spondylolisthesis greater than Grade 1.

Attempted reduction of isthmic spondylolisthesis is associated with a high rate of neurological
complications and instrumentation failure. The reported rate of transient or permanent neurological sequelae has been approximately 30%. Further fueling the controversy about the appropriate surgical procedure is the issue of when to perform decompressive surgery. The literature on this topic is inconclusive.

Hanley and Levy[25] reported on 50 consecutive patients with isthmic lumbosacral spondylolisthesis. Twenty-two patients who experienced mechanical low-back pain without radicular symptoms were treated by lateral fusion of L-4 to the sacrum with autologous iliac crest bone grafting. Twenty-eight patients with both back pain and radicular symptoms underwent neural decompression and lateral fusion from L-4 to the sacrum and iliac crest bone grafting combined with the Gill procedure. No instrumentation was used. At their 40-month follow-up examination, 50% of patients in both groups reported an excellent or good outcome. This percentage did not correlate with the degree of vertebral body slippage. A trend toward unsatisfactory outcomes was present among males, middle-aged individuals, individuals who smoked cigarettes, and patients with radicular symptoms. Harris and Weinstein[27] have addressed long-term outcomes in patients with Grade 3 or 4 spondylolisthesis treated conservatively (11 patients) and surgically (21 patients). Patients in the surgical group underwent posterior interlaminar fusion. At a mean 18-year follow up in the conservatively treated group, 36% were asymptomatic, 55% had mild symptoms, and one patient had significant symptoms. Forty-five percent of patients had an abnormal finding on neurological examination, but none was incontinent. All of the patients in this group led an active life. At a mean 24-year follow up in the surgical group, 57% were asymptomatic, 38% had mild symptoms, and one had significant symptoms. Neurological examination revealed a significant finding in half of these patients. Seventy-five percent of the patients in the surgical group reported that their symptoms had improved over time compared with 36% in the conservatively treated group. The degree of spondylolisthesis was not related to outcome in either group. Radiographic measurements of the percentage and angle of slippage did not predict outcome in either group.

Johnson and Kirwan[30] followed 17 patients in whom posterior or posterolateral fusion was performed for severe spondylolisthesis. At a mean 14-year follow up, seven patients experienced occasional back pain. Nine patients believed that their spinal deformity had improved, and 16 patients rated their surgical outcome as excellent.

Boxall, et al.[7] retrospectively reviewed 43 patients with Grade 3 or greater spondylolisthesis at L5-S1. Two patients were asymptomatic at presentation, whereas 41 patients had symptoms of back pain, radicular pain, hamstring tightness, or gait disturbance. Treatment groups were as follows: four patients were treated conservatively, 11 with fusion, 18 with decompression and arthrodesis, and 10 with reduction and fusion. At a mean follow up of 6.7 years, 3% of the patients were fully active without symptoms, and 14% were fully active without limitations but experienced occasional mild pain. Of the patients with a solid arthrodesis, spondylolisthesis progressed in 26%. Nine patients had a significant increase in the angle of slippage. Of nine patients with a pseudarthrosis, five experienced progressive slipping. Kaneda and colleagues[32] performed posterolateral fusion with distraction-rod instrumentation in 53 cases of isthmic spondylolisthesis. At 3-year follow-up review, 71% had excellent, 21% had good, 6% had fair, and the remainder had unsatisfactory outcomes.

**DEGENERATIVE SPONDYLOLISTHESIS**

Degenerative spondylolisthesis (Frymoyer type 3)[20] is an acquired condition in which disc degeneration is believed to lead to facet arthropathy and segmental instability. It is seen most frequently
in elderly patients presenting with low-back pain and radicular symptoms or neurogenic claudication, whose neurological examination is often unremarkable. Radiographic evaluation demonstrates anterior spondylolisthesis, usually at L4-5 and, less frequently, at L3-4. Typically, the degree of slippage is approximately one-fourth of the vertebral body width. Dynamic lateral flexion-extension radiographs can be used to document gross instability.

The use of fusion to treat degenerative spondylolisthesis continues to be controversial. Feffer, et al., retrospectively compared degenerative spondylolisthesis in eight patients who had undergone lumbar decompression and fusion (type of fusion not reported) with 11 patients who had undergone only decompression. Preoperatively, all patients experienced leg and back pain. Radiographic studies confirmed thecal sac compression and facet joint arthrosis with anterior spondylolisthesis of at least 3 mm. The follow-up period for the fusion group was 42 months. All patients in this group were satisfied with their surgical outcomes. No spinal instability was documented on lateral flexion-extension radiographs. Among the decompression only group, with a mean follow-up period of 25 months, five patients achieved good, three patients had fair, and three patients had poor outcomes. Four patients developed radiographically confirmed instability. The degree of instability, type of instability, and surgical outcome were correlated.

For patients with degenerative spondylolisthesis at L4-5, DePalma and Rothman have recommended complete decompression of the neural elements combined with intertransverse process fusion from L-4 to L-5. Of their patients, 64% reported total pain relief, whereas 20% experienced partial relief. Reynolds and Wiltse reported a patient satisfaction level of 78% for those who underwent facet-sparing procedures. In contrast, the level of patient satisfaction was only 33% among those who had undergone complete removal of a facet.

Johnsson and coworkers studied postoperative spinal instability in 45 patients undergoing decompression for lumbar stenosis. Preoperatively, 20 patients had degenerative spondylolisthesis and 25 had acquired spinal stenosis. Postoperative slippage was seen in 18 patients. The risk of further slippage was higher for the degenerative spondylolisthesis group but it did not influence operative outcome.

Lombardi and colleagues retrospectively reviewed 107 patients who were treated for degenerative spondylolisthesis with one of three surgical protocols. No correlation was found between the degree of preoperative spondylolisthesis and surgical outcome. Follow-up periods ranged from 2 to 7 years. Among patients treated with decompressive laminectomy and aggressive facetectomy, 33% had a good-to-excellent outcome. Among patients treated with laminectomy and medial facetectomy, 80% had a good-to-excellent outcome. The third group, treated using a protocol similar to that of the second group, also underwent intertransverse process fusion between slipped levels. Among these patients, 90% had a good-to-excellent outcome.

Nasca studied the outcome of surgical decompression with fusion in patients with lumbar stenosis from multiple causes. He concluded that fusion was indicated in patients undergoing decompression for stenosis with degenerative spondylolisthesis, facet joint degeneration, degenerative disc disease with instability, and scoliosis with instability. Lehmann, et al., performed a retrospective, 33-year follow-up study in 62 patients who had undergone lumbar fusion for radiographically confirmed instability, intraoperative instability, degenerative disc disease, facet joint syndrome, spondylosis, degenerative spondylolisthesis, and chronic low-back pain. Of 33 patients available for physical examination, spinal instability was found to be present in 15 (45%), and 14 (42%) had recurrent stenosis.
The degree of spondylolisthesis and stenosis did not correlate with the patients' symptoms at follow up. The best long-term outcome was reported in patients who had undergone decompression and fusion for spondylolisthesis.

Herkowitz and Kurz[29] retrospectively studied the benefit of routine arthrodesis for degenerative lumbar spondylolisthesis and stenosis. Fifty patients were equally divided into a nonfusion group and an arthrodesis group. At a mean 3-year follow-up time, the level of patient satisfaction was significantly higher in the arthrodesis group. Twenty-four patients reported a satisfactory outcome, and only one was unsatisfied. In the nonfusion group, 11 patients were satisfied with their outcome and 14 were not.

The simultaneous use of instrumentation for treating lumbar instability is also controversial. Hanley[23] and Harrington and Dickson[26] advocate the use of instrumentation to correct spinal deformities and thereby to decrease tension on neural elements. Markwalder[43] found that 91% of 100 patients achieved excellent outcomes when instrumentation was combined with fusion to treat degenerative spondylolisthesis. The results from a prospective, randomized study of 124 patients undergoing lumbar or lumbosacral fusion for degenerative conditions also support the use of instrumentation.[66] The diagnostic groups included isthmic spondylolisthesis, degenerative spondylolisthesis, degenerative disc disease, degenerative scoliosis, spinal stenosis, and failed back surgery. The rate of successful fusion for patients undergoing posterolateral fusion with autogenous bone graft was 65%, and 71% achieved a good-to-excellent outcome. The rate of successful fusion increased to 77% in patients undergoing posterolateral fusion supplemented by a semirigid pedicle screw/plate fixation system, and 89% had a satisfactory outcome. The rate of successful fusion further increased to 95% in patients undergoing fusion with a rigid pedicle screw/rod fixation system, and 95% experienced a good-to-excellent outcome.

To define the role of fusion and instrumentation further, Bridwell, et al.,[8] prospectively studied 44 patients with degenerative spondylolisthesis and stenosis. Patients were randomized to one of three groups: nonfusion (nine patients), transverse process fusion with autogenous iliac bone graft without instrumentation (10 patients), and transverse process fusion with autogenous iliac crest bone graft and instrumentation (24 patients), with a mean follow-up period of 3 years. Fusion with instrumentation significantly improved the rate of successful fusion. Furthermore, patients in the first two groups had a higher rate of postoperative spondylolisthesis: four of the nine patients in the nonfusion group and seven of the 10 patients who underwent fusion without instrumentation. Only one of the 24 patients in whom instrumentation was placed had progressive spondylolisthesis, which was defined as angulation of the motion segment greater than 10° or translation of the vertebral body greater than 3 mm on lateral flexion-extension radiographs.

Yuan, et al.,[65] studied a historical cohort of patients who underwent fusion with pedicle screw fixation of the thoracic, lumbar, and sacral spine for the treatment of degenerative spondylolisthesis or fractures. Of 2633 patients undergoing surgery for degenerative spondylolisthesis, 2177 patients were treated with and 456 patients were treated without pedicle screws. The use of pedicle screws was safe and improved the patients' pain, fusion success rate, and neurological and functional outcomes.

Mardjetko, et al.,[42] performed a metaanalysis of 25 published studies to assess the advantages of instrumentation in the treatment of degenerative lumbar spondylolisthesis. An overall satisfaction rating of 69% was calculated for the studies in which decompression without fusion had been evaluated. Fusion without instrumentation increased the satisfaction rating to 90%. The average rate of successful fusion was 86%. The level of patient satisfaction with decompression and fusion with pedicle screws was 86%,
the rate of fusion was 93%, and the complication rate was 10%. They concluded that their findings supported the use of fusion in the management of degenerative lumbar spondylolisthesis. Overall, however, it is difficult to assess the advantages of instrumentation because the study designs and treatment protocols in the literature are not uniform.

CONCLUSIONS

Based on review of the literature and our own experience at the Barrow Neurological Institute, we have attempted to define indications for lumbar fusion, with or without instrumentation, based on defined diagnostic categories. Clear indications for fusion include trauma, tumor, or infection with two- or three-column injury, iatrogenic instability, and isthmic spondylolisthesis. Relative indications include degenerative spondylolisthesis, radiographically proven dynamic instability with pain or neurological findings, adult scoliosis, and mechanical back pain. Fusion is rarely indicated in the setting of routine discectomy, abnormal radiographs without appropriate findings (such as degenerative disc disease), facet joint syndrome, failed back surgery, or stable spinal stenosis.

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