Evaluation and treatment of spinal metastases: an overview

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Metastases to the spine are a common and somber manifestation of systemic neoplasia. The incidence of spinal metastases continues to increase, likely a result of increasing survival times for patients with cancer. Historically, surgery for spinal metastases has consisted of simple decompressive laminectomy. Results obtained in retrospective case series, however, have shown that this treatment provides little benefit to the patient. With the advent of better patient-related selection practices, in conjunction with new surgical techniques and improved postoperative care, the ability of surgical therapy to play an important and beneficial role in the multidisciplinary care of cancer patients with spinal disease has improved significantly. Controversy remains, however, with respect to the relative merits of surgery, radiotherapy, chemotherapy, or a combination of these treatments.

In this topic review, the literature on spinal column and spinal cord metastases is collated to provide a description of the presentation, investigations, indications for surgical therapy, and the role of adjuvant cancer therapies for patients with spinal metastases. In addition, the authors discuss the different surgical strategies available in the armamentarium of the neurosurgeon treating patients with spinal metastasis.

KEY WORDS • metastasis • spine • neurooncology

Cancer therapies continue to evolve and improve. As a result, in patients with many types of systemic neoplasms survival time continues to increase. Related to this increased prevalence of longer-term cancer survivors, there is likely to be an increase in the incidence of metastatic disease of the spinal column and the spinal cord. As metastases to the spine are common, neurosurgeons must be well versed in the many nuances of treatment. Indeed, the principles of clinical and surgical management of spinal metastases can serve as a paradigm for suitable and successful management of all spinal tumors.

Although surgery is now established as a viable, beneficial treatment option in the treatment of spinal metastases, controversy exists regarding the relative merits of surgery, radiotherapy, chemotherapy, and a combination of these therapies in different patient groups.

INCIDENCE OF SPINAL METASTASES

The vertebral column is the most common site of skeletal metastases. Spinal metastases are a very frequent manifestation of systemic neoplasia, with up to 70% of cancer patients harboring secondary spinal disease. Metastases are the most common group of spinal column tumors, with approximately 18,000 new cases diagnosed in North America each year. Up to 10% of cancer patients will develop symptomatic secondary spinal lesions, with multiple levels of involvement present in 40 to 70% of symptomatic cancer patients (Fig. 1). Cancers from breast, lung, prostate, hematopoietic (for example, lymphoma or multiple myeloma), and renal origins account for the vast majority of extradural spinal metastases. This reflects both the prevalence of these particular neoplasms, as well as their predilection for metastasis to bone. In addition, the spinal lesion represents the first manifestation of cancer in 12 to 20% of patients who present with symptoms related to spinal metastases.

Classic autopsy investigations have demonstrated that the distribution of extradural metastases is related to the size of the vertebrae. Thus, metastatic lesions are most commonly located in the lumbar region, less so in the thoracic spine, and least often (< 20%) in the cervical spine. Despite this distribution, symptomatic lesions occur more frequently in the thoracic region (70%) than in the lumbarosacral or cervical segments. Particular predilection for thoracic levels about T-4 and the thoracolumbar junction has been noted. The smaller size of the thoracic spinal canal in relation to the thoracic spinal cord accounts for this discrepancy.
CLASSIFICATION OF METASTASES

Spinal metastases, like all spinal tumors, are classified according to their anatomical distribution. Upwards of 95% of spinal metastases are extradural lesions. Intradural extramedullary tumors comprise the majority of the remaining lesions, whereas intramedullary metastases are rare, comprising approximately 0.5% of spinal axis metastases.

Most extradural metastases are initially osseous lesions that originate within a portion of the vertebra and subsequently impinge on the thecal sac secondary to rapid, destructive growth. Lesions located purely posteriorly (Fig. 2) or anteriorly to the thecal sac are uncommon; the majority have a predilection for lateral displacement (that is, antero- or posterolateral) with respect to the spinal cord. Despite the frequency with which neoplastic extradural compression occurs, the dura mater is a strong anatomical barrier to neoplastic invasion and is rarely breached. Lesions located purely within the epidural compartment and without bone involvement account for only a small fraction of extradural metastases, with hemopoietic malignancies noted to have a predilection for such anatomic localization.

Intradural extramedullary spinal metastases most commonly occur as tertiary drop metastases from intracranial intradural secondary lesions and are often entangled within the nerve roots of the cauda equina (Fig. 3). Melanoma, lymphoma, and medulloblastoma are common primaries. In a series reported by Chow, et al., in which the authors examined only IDEM metastases of nonneurogenic origin, 80% of patients also harbored intracranial metastatic foci, also suggesting drop metastases as the most common mechanism of spread. In this same series, the most frequent histological subtype was adenocarcinoma, and the most frequent primary source was the lung. Intramedullary spinal cord metastatic disease accounts for only 8.5% of all metastases to the central nervous system and 2.1% of all cancers in case series published before the advent of MR imaging. More sophisticated neuroimaging is likely to increase the prevalence of this difficult-to-diagnose entity. The majority of intramedullary metastases arise from the lung, with small cell carcinoma being the predominant histological subtype. Breast, colorectal, renal, melanoma, thyroid, and lymphoma have all been reported. Like IDEM metastases, intramedullary lesions are found most often in patients with concomitant brain metastases.

PATHOPHYSIOLOGICAL MECHANISMS OF SPINAL METASTASES

The majority of systemic neoplasms are thought to metastasize to the extradural spinal axis by hematogenic
Overview of spinal metastases

spread. This is believed to occur by means of one of two mechanisms: 1) via arterial emboli to the abundant bone marrow of the VBs and subsequently into the anterior or posterior extradural space through venous channels,6 or 2) via retrograde spread through the valveless extradural Batson’s venous plexus.6 Batson originally conceived the theory of retrograde venous seeding to account for the high incidence of prostatic metastases to the spinal column. More recently this mechanism has been used to account for the spread of many other systemic neoplasms.35,66 A third mechanism of extradural metastasis is direct invasion/extension of tumor into the epidural space, most characteristic of hemopoietic neoplasms such as lymphoma.82

The pathophysiological mechanism of IDEM metastases is thought to involve cerebrospinal fluid spread from intracranial secondary lesions, or arterial hemogenic routes.12,81 In a similar manner, intramedullary metastases are believed to result from leptomeningeal carcinomatosis with subsequent cerebrospinal fluid spread, or as a result of hemogenic emboli from a primary or secondary pulmonary lesion.15,18,84

CLINICAL DIAGNOSIS

A characteristic clinical syndrome is produced by spinal column metastasis.27,55,57,69,71 Pain is the initial symptom in 90 to 95% of patients. This pain is usually local and associated with tenderness elicited by palpation over the spinous process at the level of involvement. On occasion a component of radicular pain, radiating in the distribution of the nerve root at the involved level, may accompany neck or back pain. Pain of a severe, burning, dysesthetic nature is often associated with IDEM metastatic lesions,71 whereas pain that is aggravated by movement and alleviated by immobility should raise suspicion of spinal instability and pathological fracture–dislocation.70 In general, myofascial, discogenic, and spondylotic pain are more common in the cervical and lumbar spine and are relieved with rest. Spinal metastasis–induced pain most commonly localizes to the thoracic segments and is often worse with rest and at night.27 Local back or neck pain can be present for a significant duration before a correct diagnosis is reached; the median time to diagnosis in one study was 2 months.27 Unfortunately, the correct diagnosis is often not made until signs and symptoms of spinal cord compromise are manifest. It is axiomatic that back or neck pain in a cancer patient is secondary to spinal metastasis until proven otherwise.

Neurological compromise in the form of weakness, senso- rity loss, and sphincter disturbance usually occurs after the onset of pain. At the time of diagnosis, sensory or motor deficits are present in 38 to 76% of patients, and 50% of patients are nonambulatory (secondary to pain and/or neurological deficit).27,67 Sphincter disturbance is also common, with 37% of patients requiring placement of a urinary catheter in one series.40 Brown–Séquard syndrome, or a variation thereof, is a common neurological finding at the time of diagnosis in patients with intramed- ullary spinal metastasis.84 The natural history of spinal metastasis is relentless progression to complete and irreversible paraplegia unless timely treatment is undertaken.7

Differential Diagnosis

Symptomatic secondary spinal lesions are a common manifestation of systemic neoplasia, and the presence of back or neck pain in a cancer patient should never be minimized. In some cancer patients presenting with axial pain and/or signs and symptoms of spinal cord dysfunction, however, another cause may account for their clinical syndrome (Table 1). A careful, thorough clinical history and physical examination coupled with appropriate laboratory and diagnostic imaging investigations, as outlined in the following sections, readily delineate metastatic spinal tumors from other more benign differential diagnostic entities.

DIAGNOSTIC INVESTIGATIONS

Laboratory investigations and diagnostic neuroimaging studies are indispensable in the evaluation of the known cancer patient with signs and symptoms suggestive of spinal cord metastasis, as well as in the patient who presents without history of systemic neoplasia.

General Investigations

A useful, logical first step is to perform routine blood studies consisting of a complete blood count, albumin, electrolyte panel, liver enzyme assays, blood urea nitrogen, and serum creatinine. Information about the immunological and nutritional status of the patient can be obtained, as can knowledge of renal dysfunction secondary to malignancy determined by increased creatinine or blood urea nitrogen levels. Alterations in liver function may suggest the presence of liver metastasis or, more rarely, a primary liver carcinoma. Hypercalcemia is a common finding in patients with metastatic bone disease and requires prompt treatment to avoid cardiac electrical dysfunction. Serum electrophoresis and urine for Bence Jones protein levels are useful tests in the diagnosis of suspected multiple myeloma. Assays for prostate-specific antigen aid in the determination of prostate carcinoma, which is a common primary source of spinal metastasis. Chest radiography and abdominal ultrasonography are useful screening tools for primary neoplasms, whereas CT scanning of the chest, abdomen, and pelvis provides detailed information about the presence of neoplasm and should be readily performed when metastatic disease is suspected.

| TABLE 1 |
| Differential diagnosis in spinal cord compression in patients with systemic cancer |

<table>
<thead>
<tr>
<th>Differential Diagnosis</th>
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<tbody>
<tr>
<td>degenerative spinal disorders (disc herniation, spinal stenosis, spondyloolisthesis)</td>
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<td>meningeal carcinomatosis/lymphomatosis</td>
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<td>infectious diseases (osteomyelitis, discitis, epidural abscess)</td>
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<tr>
<td>radiation-induced myelopathy</td>
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<tr>
<td>epidural hematoma (secondary to coagulopathy)</td>
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<tr>
<td>transverse myelitis</td>
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<tr>
<td>paraneoplastic syndromes</td>
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<tr>
<td>vascular malformations (arteriovenous malformation, dural arteriovenous fistula, cavernoma)</td>
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Diagnostic Neuroimaging of the Spinal Axis

Modern neurodiagnostic imaging studies are indispensable in the evaluation and treatment of patients with metastatic spinal tumors. These studies are used to diagnose, stage, and plan treatment in patients with metastatic lesions, as well as to follow them postoperatively.

Spinal Radiography. Plain radiographs of the spine remain a valuable and readily available initial imaging study. Anteroposterior and lateral radiographs demonstrate abnormal findings in up to 90% of patients with symptomatic spinal metastasis. Lytic lesions and vertebral collapse are common; however, both osteoblastic and -sclerotic alterations also occur, especially with breast and prostatic metastasis. Plain x-ray film findings include pedicle erosion (that is, the “winking owl” sign), paraspinal soft-tissue shadows, wedge compression, and pathological fracture–dislocation (Fig. 4). Intervertebral disc margins are invariably spared in metastatic tumor invasion, contrasting with the disc erosion commonly observed with infectious entities.

Local bone destruction of 50% is required before a lesion can be detected on plain x-ray films. In a high proportion of patients with symptomatic secondary spinal lesions abnormalities are demonstrated on plain radiographs, but in up to 26% of the cases early metastatic lesions can be missed. Consequently, clinicians investigating patients with a high suspicion of metastatic spinal neoplasms should arrange expeditious CT and/or MR imaging investigations, even if normal findings are observed on plain radiographs.

Myelography. Traditionally, myelography (Fig. 5) had been the gold standard for determining the level of spinal cord compression. This status has been largely supplanted by MR imaging studies. Myelography continues to play a valuable role, however, especially in cases in which MR imaging is not readily available, is contraindicated secondary to ferromagnetic implants, or in patients unable to tolerate MR imaging. Anatomical localization and the relation of the lesion to the dura, spinal cord, and nerve roots are readily facilitated by myelography. It is desirable to visualize the entire length of the spinal axis to rule out cord compression at multiple levels. Thus, if contrast material does not readily flow rostrally past a region of spinal block after a lumbar injection, then a cisternal injection may also be required. Myelography performed via lumbar puncture in the presence of a complete spinal block carries the risk of spinal cord herniation and subsequent neurological deterioration, with deterioration reported in 14% of patients in one study. Cisternal myelography has not been associated with this risk.

Computerized Tomography Scanning. The principal use of CT scanning is in the assessment of the osseous ar-
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architecture of the spinal axis because MR imaging does not provide optimum images of bone. Computerized tomography scanning provides important information regarding the degree of tumor association with cortical bone and the extent of neoplastic destruction (Fig. 6). Thus, CT scanning is an important complement to MR imaging. In patients in whom MR imaging cannot be performed, CT scanning is particularly useful when conducted immediately following myelography. Computerized tomography myelography provides better anatomical detail of the spinal axis than either CT or myelography alone. The information acquired from CT myelography is often comparable with that obtained using MR imaging.42

**Magnetic Resonance Imaging.** Magnetic resonance imaging provides unparalleled visualization of the spinal column and spinal cord and is the neuroimaging method of choice in patients with suspected spinal tumors. It has been demonstrated to facilitate the earlier diagnosis of spinal metastases4 than other modalities. It is superior in depicting epidural and bone marrow tumor infiltration, and it delineates the extraosseous soft-tissue component of a neoplasm from the normal paraspinal soft tissue and neural structures.28 In addition to its diagnostic utility, MR imaging information is essential to the process of thorough surgical planning. Gadolinium enhancement further increases the sensitivity of MR imaging investigations, because metastases invariably enhance. Visualization of the entire vertebral axis in multiple orthogonal planes is easily accomplished, a goal that is important given the high incidence of multiple vertebral levels of tumor infiltration.19,40 Studies show that the results of MR imaging investigations alter therapeutic decisions in a significant number of patients, especially with regard to the addition or modification of radiotherapy.14

**Bone Scanning.** Given the current sensitivity of MR imaging in detecting metastases in the vertebral axis, the radioisotope bone scan is most useful as a sensitive screening or follow-up test to assess the total burden of osseous disease. Data obtained using this modality are invaluable in the process of optimum patient selection.

**MANAGEMENT**

The treatment of patients with symptomatic secondary spinal lesions is undertaken to relieve pain and to preserve or restore neurological function. Cure is not a realistic expectation in patients with metastatic spinal tumors. Palliation is the reasonable goal; life expectancy is often relatively short, with median survival ranging from 4 to 15 months in various series.82,89,100,102 These modest goals contribute immeasurably to the quality of life in cancer patients and decrease the burden of care.

**Corticosteroid Medication Administration**

Administration of an intravenous or oral corticosteroid agent serves as a vital adjuvant therapy in the treatment of patients with spinal metastases causing spinal cord compression.51 Improvement in neurological status is often documented after steroid administration. The mechanism of action of corticosteroids in this setting is not completely understood, but reduction in peritumoral vasogenic spinal cord edema has been documented experimentally and is at least partially responsible.97,98 The optimum dose of corticosteroid agent to be administered also remains open to debate; in one study the investigators recommended high-dose dexamethasone (100-mg bolus followed by 4 mg every 6 hours) because it provided better protection than more standard dosing (10 mg followed by 4 mg every 6 hours).31 No added advantage has been found with higher dosages.99 It is standard practice to administer 10 mg of dexamethasone intravenously once a diagnosis is established, followed by 4 mg every 6 hours, until definitive treatment is commenced. A tapering regimen is then instituted. Medical complications such as exacerbation of diabetes mellitus, hypertension, immunocompromise, gastric ulceration, acute delirium, and wound infection must be carefully monitored. Although it has been common practice to administer steroids prior to commencing spinal irradiation, the authors of a recent study have demonstrated that in patients without myelopathy dexamethasone need not be administered.59

**Radiation Therapy**

Radiotherapy plays an important role in the treatment of metastatic spinal tumors. Many consider it to be the mainstay of therapeutic intervention.8,46,47,60 Spinal irradiation is particularly effective in the treatment of various radiosensitive histological subtypes of metastases, such as hematopoietic tumors and prostate carcinoma. Excellent response to radiotherapy in the setting of spinal cord compression is found in 30% of all patients, including those with more radioresistant lesions such as breast carcinoma and metastatic melanoma.54

The response to radiotherapy must be considered in relation to the parameters of pain relief and functional status. Response to radiotherapy alone has been reported to be 66 to 80%,47,60 including improvement in motor dysfunction in one half of patients and stabilization of symptomatology in one third.60 Pretreatment functional status is the most important prognostic indicator. Significant correlation between ambulatory status before radiotherapy and the median length of survival after treatment initiation has been noted.51,53,60

The standard radiation dose schedule for metastatic spinal tumors consists of 20 to 30 Gy administered in five to 10 treatment sessions to the area of spinal cord compression; many variations exist and are related to the general

Fig. 6. Axial CT scan obtained through the midthoracic spine revealing extensive tumor-induced cortical bone destruction.
medical status of the patient and the extent of spinal tumor. Short-course radiotherapy (8 Gy given in two sessions) has also been shown to be as advantageous as more protracted schemes and to cause less severe side effects. Some have also advocated intraoperative radiotherapy, but this procedure is not widely conducted. In those patients presenting with complete paraplegia of several days’ duration, relief of pain should be the treatment goal; this can be accomplished by administering a single 8-Gy fraction.

Side effects are generally mild. Radiation-induced esophagitis may occur transiently 1 to 2 weeks following treatment of upper thoracic lesions. Irradiation of the lower thoracic and upper lumbar spine often produces nausea and vomiting, resulting as the radiation beam exits through the epigastrium. This side effect is most pronounced after the first two to three fractions and is generally controlled by providing standard antiemetics. Radiation-induced myelopathy is an important late-onset complication but is rare with standard dosimetry.

Surgical Therapy

The role of the neurosurgeon in the treatment of patients with metastatic spinal tumors continues to evolve. Historically, decompressive laminectomy was performed in an attempt to alleviate cord compression. Large retrospective reviews, however, have demonstrated that this surgical approach is beneficial in only a minority (<40%) of patients. Furthermore, in studies investigating the efficacy of radiotherapy alone or in combination with simple laminectomy, the authors found no difference in outcome. Such poor laminectomy-related results are not surprising, given the lateral (antero- or posterolateral) displacement of most spinal metastases in relation to the dural sac; purely posterior lesions are uncommon. Simple laminectomy thus provides inadequate decompression of the spinal cord and nerve roots in the majority of patients. Laminectomy alone is also likely to exacerbate mechanical spinal instability, especially in the case of vertebral collapse, a common finding in patients with extradural metastatic lesions. The failure to individualize surgical approach based on tumor location has likely contributed to the failure of traditional surgical methods (laminectomy) in the treatment of spinal metastases.

Refinement of surgical approaches such as the posterolateral (Fig. 7A) and anterior exposures (Fig. 7B) for spinal cord decompression, as well as the ongoing development and evolution of spinal stabilization apparatus and techniques, has greatly improved the efficacy of surgical intervention. As a first-line therapy surgery remains a desirable option—numerous studies suggest that wound dehiscence and infections are drastically increased when surgery is performed after primary radiotherapy. Radiotherapy also appears to be less effective when epidural spinal cord compression is associated with a paravertebral soft-tissue mass. In this situation, improvement in functional status has been shown when resection is used as a first-line therapy and followed by radiotherapy.
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**Indications and Contraindications to Surgery.** Optimum surgery-related outcome requires careful patient selection (Table 2). Emergency surgery is mandated in the setting of rapidly progressive or advanced paraplegia, because complete and irreversible spinal cord injury will occur if prompt decompression of the thecal sac and nerve roots is not performed. Patients harboring a pathological fracture–dislocation have an unstable spine and also require surgery for decompression and stabilization of the spinal axis. De novo radiotherapy can exacerbate this instability and cause further cord compression. In cases in which the diagnosis of spinal metastases is not certain (that is, in a cancer patient with symptoms believed to be the result of a cause other than symptomatic secondary spinal lesions, or in a patient without a history of cancer), surgery should be performed. In these cases surgery serves a diagnostic as well as therapeutic purpose. Finally, in patients in whom radiotherapy has failed to control the progression of symptoms or in those with a known radioresistant tumor, surgery should be considered the treatment of choice.

Surgical decompression cannot reverse a complete paralysis of greater than 24 hours’ duration and should not be performed. Short-course radiotherapy that provides pain relief is a more prudent course of action. Radiotherapy provides excellent local control of radiosensitive lesions such as lymphoma or multiple myeloma. Patients with a known highly radiosensitive lesion are thus best served by undergoing primary spinal irradiation. Adhering to the tenet that successful palliation is the primary goal in management of spinal metastases, those patients with a heavy burden of disease, a limited life expectancy and/or extreme medical comorbidities are not well served by surgical therapy. Spinal irradiation should be considered the first-line therapy in such cases.

**Preoperative Embolization.** Metastases originating from thyroid and renal cell carcinoma are particularly vascular. Excision can be associated with catastrophic intraoperative blood loss. Endovascular embolization is a well-established procedure that often greatly decreases the vascularity of spinal metastases (Fig. 8), decreasing intraoperative blood loss and facilitating more complete lesion resection and spinal decompression. Thus, the need for embolization should be carefully considered before beginning resection.

**Surgical Strategies**

Any successful surgical strategy must ensure both decompression of the dural sac and nerve roots as well as stabilization of the spinal column (Fig. 9). Spinal instability may be present as a result of tumor-induced bone destruction or the surgical decompression. It is important to individualize the surgical approach to the spine on a patient-by-patient basis. The surgical approach to spinal metastases may be ventral (anterio or anterolateral) or dorsal (posterior or posterolateral). An adequate posterior approach involves posterolateral (uni- or bilateral) decompression, allowing circumferential decompression of the thecal sac and nerve roots. Anterior-approach decompression involves corpectomy. In either case (invariably, following corpectomy) spinal instability will be precipitated. Consequently, the implantation of spinal instrumentation is always necessary (Fig. 10). If a longer-term survival is anticipated, use of a bone autograft is appropriate. Otherwise, an artificial structural support can be fashioned from methylmethacrylate and used in conjunction with the implanted mechanical construct. The plethora of devices and techniques available for spinal reconstruction after metastasis resection attests to the fact that no single method is supreme (Tables 3 and 4).

TABLE 2
**Indications and contraindications to surgery for spinal metastases**

<table>
<thead>
<tr>
<th>Indications &amp; Contraindications</th>
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<tbody>
<tr>
<td><strong>surgical indications</strong></td>
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<tr>
<td>rapidly progressing or far-advanced paraplegia</td>
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<tr>
<td>histological diagnosis in an unknown primary</td>
</tr>
<tr>
<td>spinal instability &amp; pathological fracture–dislocation</td>
</tr>
<tr>
<td>failure of radiotherapy/known radioresistant lesion</td>
</tr>
<tr>
<td><strong>relative contraindications to surgery</strong></td>
</tr>
<tr>
<td>longstanding complete paralysis (&lt;24 hrs)</td>
</tr>
<tr>
<td>highly radiosensitive tumor (lymphoma, myeloma)</td>
</tr>
<tr>
<td>multiple levels of involvement</td>
</tr>
<tr>
<td>poor life expectancy (&lt;3 mos)</td>
</tr>
<tr>
<td>extreme medical comorbidity</td>
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**Fig. 8.** Spinal angiography performed before (pre) and after (post) embolization of extradural dural metastasis from a primary renal cell carcinoma.

**Fig. 9.** Schematic diagram outlining the factors to be considered in planning surgery for spinal metastases.
rapid evolution in the surgical management of spinal metastases. The introduction of new surgical techniques and reconstructive instruments has brought with it a move to more extensive and radical surgical procedures. These modern surgical methods have been conceived with the goal of affording maximum therapeutic benefit to the patient. Their exact utility, however, remains to be proven.

The decision to perform anterior or posterior procedures depends on numerous interrelated factors. Each approach has a specific role, and no single procedure is universally more advantageous than another.

**Tumor Location.** Whereas laterally situated lesions are most common, extradural metastases can present in all manners of spatial relation to the thecal sac. Generally, lesions involving the VB and causing anterior or anterolateral compression are best approached via anterior or anterolateral routes in which vertebral corpectomy is performed a biomechanically sound stabilization construct is placed. Laterally situated lesions can often be adequately resected via a posterolateral approach. Only a minority of cases in which isolated posterior compression is present are best served by laminectomy. Intradural extramedullary and intramedullary lesions are almost always approached via a wide posterior laminectomy.

**Spinal Level.** Extreme rostrally and caudally located lesions pose a specific challenge. Although extensive and high morbid–related procedures are available that allow access to the craniocervical junction, most patients with metastatic lesions are not suitable candidates for these interventions, because such surgical therapies are contrary to the basic philosophy of successful palliation and preservation of quality of life. Additionally, anterior stabilization and instrumentation is very difficult at both the craniocervical and lumbosacral junctions. As such, in most cases, the best initial intervention is posterior decompression and stabilization in cases in which these extreme levels are affected.

**Extent of Involvement.** Extradural metastases involving one or two contiguous VB levels is best approached via anterior or anterolateral routes, because these allow for direct decompression and the possibility of a construction of a sound fusion mass. If three or more contiguous levels require attention, however, decompression is best achieved using a posterolateral approach combined with posterior fusion and placement of instrumentation, or a combined anterior–posterior procedure.

**Cortical Bone Integrity.** Multiple levels of tumor infiltration are commonly found. Thus, it is important to determine the integrity of the VBs adjacent to the region in which the decompression is performed. If adjacent bone is compromised, anterior fusion after anterolateral decompression may not be possible. In such cases, the surgeon should consider posterolateral decompression and placement of posterior instrumentation as a safer alternative.

### TABLE 3
Selected anterior spinal fixation methods*

<table>
<thead>
<tr>
<th>Original Proponents &amp; Yr</th>
<th>Method</th>
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</thead>
<tbody>
<tr>
<td>Cloward, 1958</td>
<td>Cloward</td>
</tr>
<tr>
<td>Robinson &amp; Southwick, 1960</td>
<td>Smith–Robinson</td>
</tr>
<tr>
<td>Cross, et al., 1971</td>
<td>MMA</td>
</tr>
<tr>
<td>Ono &amp; Tada, 1975</td>
<td>metal prosthesis</td>
</tr>
<tr>
<td>Fielding, et al., 1979</td>
<td>corpectomy/iliac crest</td>
</tr>
<tr>
<td>Sundaresan, et al., 1984</td>
<td>MMA/double K-wire</td>
</tr>
<tr>
<td>Perrin &amp; McBroom, 1988</td>
<td>Wellesley wedge</td>
</tr>
<tr>
<td>Matsui, et al, 1994</td>
<td>ceramic prosthesis</td>
</tr>
<tr>
<td>Gokaslan, et al., 1998</td>
<td>MMA/locking plate</td>
</tr>
<tr>
<td>Schulte, et al., 2000</td>
<td>bioglass/polyurethane prosthesis</td>
</tr>
</tbody>
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* MMA = methylmethacrylate.

**TABLE 4**
Selected posterior spinal fixation methods

<table>
<thead>
<tr>
<th>Original Proponent &amp; Yr</th>
<th>Method</th>
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<tbody>
<tr>
<td>Rogers, 1942</td>
<td>interspinous wiring</td>
</tr>
<tr>
<td>Robinson &amp; Smith, 1955</td>
<td>posterolateral fusion</td>
</tr>
<tr>
<td>Harrington &amp; Dickson, 1976</td>
<td>Harrington rods</td>
</tr>
<tr>
<td>Roy-Camille, et al., 1976</td>
<td>Roy-Camille plates</td>
</tr>
<tr>
<td>Livingston &amp; Perrin, 1978</td>
<td>MMA/sublaminar wires</td>
</tr>
<tr>
<td>Perrin &amp; Livingston, 1982</td>
<td>Halifax clamp</td>
</tr>
<tr>
<td>Holness, 1984</td>
<td>Luque rectangle</td>
</tr>
<tr>
<td>Luque, 1986</td>
<td>variable spine plating</td>
</tr>
<tr>
<td>Steffee, et al., 1986</td>
<td>Hartshill rectangle</td>
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<tr>
<td>Mirovsky, et al., 1999</td>
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Overview of spinal metastases

Patient Debility. Local or systemic debility is often a factor when contemplating surgery in the patient with spinal metastases. Anterior cervical approaches through a radiation-exposed neck can be fraught with difficulty and complication. Patients with primary or secondary pulmonary lesions are often unable to tolerate the operative stresses associated with transthoracic or retroperitoneal exposures. Many cancer patients have systemic comorbidities that make consideration of the less debilitating of two possible spinal procedures the more reasonable option. To subject individuals with limited life expectancy to lengthy procedures and subsequent long convalescence periods is to betray the primary goal of treatment—successful palliation.

OUTCOME

The prognosis of the patients treated for symptomatic secondary spinal lesions is variable, given the pretreatment heterogeneity of the patient population. By the same token, results obtained in published studies defy comparison and the drawing of universal conclusions. In general, the authors of recent surgery-based series report 80 to 90% relief of pain, as well as up to 90% stabilized or improved neurological status.11,29,79,100,102 The outcome after treatment of symptomatic secondary spinal lesions depends on a number of variables (Table 5), the most durable of which is the pretreatment neurological status; intervention has little chance of restoring useful motor function in individuals in whom complete paralysis has developed prior to treatment.

SUMMARY

Metastatic spinal neoplasms are the most commonly encountered tumors of the spine. All neurosurgeons will be involved with the evaluation and treatment of these patients. Successful treatment requires recognition of the characteristic clinical presentation of this entity, combined with appropriate investigations and treatment planning. The management of patients with spinal metastases has evolved greatly over the last decade. Radiotherapy can no longer be considered the universal first-choice treatment, because the advent of new surgical strategies has greatly increased the benefit of surgical interventions. In this drive to expand our surgery-related armamentarium, rapid technical advances have been made. Patients with spinal metastases now frequently undergo radical procedures in an attempt to achieve maximum local disease control. Neurosurgeons, however, must not lose sight of the primary goal of treatment for these patients: successful palliation and improvement in quality of life. Application of this principle in a setting of multidisciplinary cooperation among neurosurgeons, orthopedic surgeons, and radiation oncologists will provide the maximum benefit for this large group of cancer patients.

TABLE 5
Prognostic factors for surgery for spinal metastases

<table>
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<tr>
<th>Prognostic Factor</th>
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<tr>
<td>extent of pretreatment deficit</td>
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<tr>
<td>duration of symptoms</td>
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<tr>
<td>histology of metastatic tumor</td>
</tr>
<tr>
<td>location of tumor</td>
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<tr>
<td>extent of primary tumor control</td>
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</tbody>
</table>

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