

AmeriHealth Caritas Louisiana

National Imaging Associates, Inc.	
Clinical guideline CERVICAL SPINE MRI	Original Date: September 1997
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GENERAL INFORMATION

- It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.
- Where a specific clinical indication is not directly addressed in this guideline, medical necessity determination will be made based on widely accepted standard of care criteria. These criteria are supported by evidence-based or peer-reviewed sources such as medical literature, societal guidelines and state/national recommendations.

INDICATIONS FOR CERVICAL SPINE MRI

- [†] If there is a combination request* for an overlapping body part, either requested at the same time or sequentially (within the past 3 months) the results of the prior study should be:
 - Inconclusive or show a need for additional or follow up imaging evaluation OR
 - The office notes should clearly document an indication why overlapping imaging is needed and how it will change management for the patient.

(*Unless approvable in the <u>combination section</u> as noted in the guidelines)

For evaluation of neurologic deficits 4-61-6

- With any of the following new neurological deficits documented on physical exam
 - Extremity muscular weakness (and not likely caused by plexopathy, or peripheral neuropathy)
 - Pathologic (e.g., Babinski, Lhermitte's sign, Chaddock Sign, Hoffman's) and other upper motor neuron signs); OR abnormal deep tendon reflexes (and not likely caused by plexopathy, or peripheral neuropathy)

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- Absent/decreased sensory changes along a particular cervical dermatome (nerve distribution): pin prick, touch, vibration, proprioception, or temperature (and not likely caused by plexopathy, or peripheral neuropathy)
- Upper or lower extremity increase muscle tone/spasticity
- New onset bowel or bladder dysfunction (e.g., retention or incontinence)- not related to an inherent bowel or bladder process
- Gait abnormalities (see <u>Table 1 Table 1</u> for more details)
- Suspected cervical cord compression with any neurological deficits as listed above

For evaluation of neck pain with any of the following⁷⁻⁹

- With new or worsening objective neurologic deficits (as listed above) on exam
- Failure of conservative treatment* for at least six (6) weeks within the last six (6) months 10
- With progression or worsening of symptoms during the course of conservative treatment*
- With an abnormal electromyography (EMG) or nerve conduction study (if performed) indicating a
 cervical radiculopathy. (EMG is not recommended to determine the cause of axial lumbar, thoracic,
 or cervical spine pain.)¹¹
- Isolated neck pain in pediatric population 12, 13 conservative care not required if red flags present
 - Red flags that prompt imaging should-Failure of conservative treatment* for at least six (6) weeks within the last six (6) months¹⁰
 - With progression or worsening of symptoms during the course of conservative treatment*
 - With an abnormal electromyography (EMG) or nerve conduction study (if performed)
 indicating a cervical radiculopathy. (EMG is not recommended to determine the cause of axial
 lumbar, thoracic, or cervical spine pain.)¹¹
 - Isolated back pain in pediatric population^{12, 13} conservative care not required if red flags present.

Red flags that prompt imaging include the presenceany of the following:

- Age 5 or younger—OR
- Constant pain, OR
- Pain lasting > 4 weeks, OR
- Abnormal neurologic examination, OR
- Early morning stiffness and/or gelling; OR
- Night pain that prevents or disrupts sleep; fever; OR
- Radicular pain; OR
- Fever or weight loss or malaise OR^{14, 15}
- Postural changes (e.g., kyphosis or scoliosis) OR
- Limp (or refusal to walk in a younger child)

As part of initial pre-operative / post-operative / procedural evaluation ("CT best examination to assess for hardware complication, extent of fusion and pseudoarthrosis"^{12, 16} and MRI for cord, nerve root compression, disc pathology or post-op infection)

For preoperative evaluation/planning



- CSF leak highly suspected and supported by patient history and/or physical exam findings (leak (known or suspected spontaneous (idiopathic) intracranial hypotension (SIH), post lumbar puncture headache, post spinal surgery headache, orthostatic headache, rhinorrhea or otorrhea, or cerebrospinal-venous fistula))
- A follow-up study may be needed to help evaluate a patient's progress after treatment, procedure, intervention, or surgery in the last 6 months. Documentation requires a medical reason that clearly indicates why additional imaging is needed for the type and area(s) requested (routine surveillance post-op not indicated without symptoms)
- Surgical infection as evidenced by signs/symptoms, laboratory, or prior imaging findings
- New or changing neurological deficits or symptoms post-operatively^{16, 17} see <u>neurological</u> deficit section above
- When combo requests (see <u>above statement</u>⁺) are submitted (e.g., MRI and CT of the spine), the office notes should clearly document the need for both studies to be done simultaneously (e.g., the need for both soft tissue and bony anatomy is required)¹⁸
 - Combination requests where both cervical spine CT and MRI cervical spine are both approvable (not an all-inclusive list):
 - OPLL (Ossification of posterior longitudinal ligament)¹⁹¹⁹
 - Pathologic or complex fractures
 - Malignant process of spine with both bony and soft tissue involvement
 - Unstable craniocervical junction
 - Clearly documented indication for bony and soft tissue abnormality where assessment will change management (i.e., surgical approach) for the patient

For evaluation of suspected myelopathy²⁰⁻²⁴

- Does **NOT** require conservative care
- Progressive symptoms including hand clumsiness, worsening handwriting, difficulty with grasping and holding objects, diffuse numbness in the hands, pins and needles sensation, increasing difficulty with balance and ambulation
- Any of the neurological deficits as noted above

For evaluation of known or suspected multiple sclerosis (MS)^{20, 25-27}

- Evidence of MS on recent baseline Brain MRI
- Suspected or known MS with new or changing symptoms consistent with cervical spinal cord disease (focal <u>neurologic deficit</u> or clinical sign, e.g., Lhermitte sign)
- Suspected or known pediatric demyelinating diseases (MS/ADEM)

Combination studies MS²⁸²⁸

• These body regions might be evaluated separately or in combination as guided by physical examination findings (e.g., localization to a particular segment of the spinal cord), patient



history (e.g., symptom(s), time course, and where in the CNS the likely localization(s) is/are), and other available information, including prior imaging.

- Cervical and/or Thoracic MRI for evaluation of highly suspected multiple sclerosis (MS) when Brain MRI has indeterminate findings and/or does not fulfill the McDonald criteria for the diagnosis of MS²⁶
- Cervical and/or Thoracic MRI with suspected transverse myelitis with appropriate clinical symptoms (e.g., bilateral weakness, sensory disturbance, and autonomic dysfunction which typically evolve over hours or days)
- Brain MRI with Cervical and/or Thoracic MRI for evaluation of neuromyelitis optica spectrum disorders (recurrent or bilateral optic neuritis; recurrent transverse myelitis)²⁹
- Known MS, entire CNS axis (Brain, and/or Cervical and/or Thoracic spine) is approvable prior to the initiation or change of disease modification treatments and assess disease burden (to establish a new baseline)
- Known MS- Follow-up scans, including brain and spine imaging, if patients have known spine disease:
 - 6-12 months after starting/changing treatment
 - Every 1-2 years while on disease-modifying therapy to assess for subclinical disease activity, less frequently when stable for 2-3 years

For evaluation of trauma or acute injury 12,3012,30

- Presents with any of the following <u>neurological deficits</u> noted above
- With progression or worsening of symptoms during the course of <u>conservative treatment</u>*
- History of underlying spinal abnormalities (i.e., ankylosing spondylitis, diffuse idiopathic skeletal hyperostosis) (Both MRI and CT would be approvable)³¹⁻³³
- When the patient is clinically unevaluable or there are preliminary imaging findings (x-ray or CT) needing further evaluation
- When office notes specify the patient meets NEXUS (National Emergency X-Radiography Utilization Study) or CCR (Canadian Cervical Rules) criteria for imaging:
 - CT for initial imaging
 - MRI when suspect spinal cord or nerve root injury or when patient is obtunded, and CT is negative
 - CT or MRI for treatment planning of unstable spine

("MRI and CT provide complementary information. When indicated it is appropriate to perform both examinations") 31

For evaluation of known or new compression fractures with worsening neck pain 1212

- With history of malignancy
 - o To aid in differentiation of benign osteoporotic fractures from metastatic disease
 - A follow-up MRI in 6-8 weeks after initial MRI when initial imaging cannot decipher (indeterminate) benign osteoporotic fracture from metastatic disease (Kumar, 2016)



- With an associated new focal neurologic deficit as above³⁴³⁴
- Prior to a planned surgery/intervention or if the results of the MRI will change management

For evaluation of tumor, cancer, or metastasis with any of the following:

(MRI is usually the preferred study, but CT may be needed to further characterize solitary indeterminate lesions seen on MRI)^{12, 35-37}

Primary tumor

- Initial staging or re-staging of a known primary spinal tumor³⁸
 - Initial staging primary spinal tumor³⁸
 - Follow-up of known primary cancer of patient undergoing active treatment within the past year or as per surveillance imaging guidance for that cancer
 - Known spinal tumor with new signs or symptoms (e.g., new or increasing nontraumatic pain, physical, laboratory, and/or imaging findings)
 - With an associated new focal neurologic deficit as above³⁴

Metastatic tumor

- With evidence of metastasis on bone scan needing further clarification OR inconclusive findings on a prior imaging exam
- With an associated new focal neurologic deficit³⁴
- Known malignancy with new signs or symptoms (e.g., new or increasing nontraumatic pain, radiculopathy or neck pain that occurs at night and wakes the patient from sleep with known active cancer, physical, laboratory, and/or imaging findings) in a tumor that tends to metastasize to the spine^{12, 39}
 - ForKnown malignancy with new signs or symptoms (e.g., new or increasing nontraumatic pain, radiculopathy or neck pain that occurs at night and wakes the patient from sleep with known active cancer, physical, laboratory, and/or imaging findings) in a tumor that tends to metastasize to the spine^{12, 39}

<u>Further</u> evaluation of <u>indeterminate findings on prior imaging (unless follow up is otherwise specified within the guideline):</u>

- <u>For initial evaluation of an</u> inconclusive finding on <u>a</u> prior imaging <u>report</u> that requires further clarification
 - One follow-up exam to ensure no suspicious change has occurred in prior imaging finding.
 No further surveillance unless specified as highly suspicious or change was found on last follow-up exam¹²
- One follow-up exam of a prior indeterminate MR/CT finding to ensure no suspicious interval change has occurred. (No further surveillance unless specified as highly suspicious or change was found on last follow-up exam.)

Indication for combination studies for the initial pre-therapy staging of cancer, OR active monitoring for recurrence as clinically indicated, OR evaluation of suspected metastases



• <u>≪</u> 5 concurrent studies to include CT or MRI of any of the following areas as appropriate depending on the cancer: Neck, Abdomen, Pelvis, Chest, Brain, Cervical Spine, Thoracic Spine or Lumbar Spine

For evaluation of known or suspected infection (osteomyelitis)/abscess 1212,40

- Infection
 - As evidenced by signs and/or symptoms, laboratory (i.e., abnormal white blood cell count, ESR and/or CRP) or prior imaging findings⁴⁰⁴¹
 - Follow-up imaging of infection
 - With worsening symptoms/laboratory values (i.e., white blood cell count, ESR/CRP) or radiographic findings^{4±2}

For evaluation of known or suspected inflammatory disease or atlantoaxial instability

- In rheumatoid arthritis with neurologic signs/symptoms, or evidence of subluxation on radiographs (lateral radiograph in flexion and neutral should be the initial study)^{42, 4343, 44}
 - Patients with negative radiographs but symptoms suggestive of cervical instability or in patients with neurologic deficits MRI is indicated⁴⁴⁴⁵
- High-risk disorders affecting the atlantoaxial articulation, such as Down syndrome, Marfan syndrome with neurological signs/symptoms, abnormal neurological exam, or evidence of abnormal or inconclusive radiographs of the cervical spine⁴⁵
- Spondyloarthropathies, known or suspected
 - Ankylosing Spondylitis/Spondyloarthropathies with non-diagnostic or indeterminate xray and appropriate rheumatology workup

For evaluation of spine abnormalities related to immune system suppression, e.g., HIV, chemotherapy, leukemia, or lymphoma^{46, 4747, 48}

As evidenced by signs/symptoms, laboratory, or prior imaging findings

Other Indications for a Cervical Spine MRI

(Note- See combination requests, below, for initial advanced imaging assessment and pre-operatively)

- Tethered cord or spinal dysraphism (known or suspected), based on preliminary imaging, neurological exam, and/or high-risk cutaneous stigmata⁴⁸⁻⁵⁰⁴⁹⁻⁵¹
 - Known Arnold-Chiari syndrome (For initial imaging initial imaging (one-time initial MRI-modality assessment) see combination below)
 - Known Chiari I malformation without syrinx or hydrocephalus, follow-up imaging after initial diagnosis with new or changing signs/symptoms or exam findings consistent with spinal cord pathology⁵¹²
 - o Known Chiari II (Arnold-Chiari syndrome), III, or IV malformation
- Achondroplasia (one Cervical Spine MRI to assess the craniocervical junction, as early as
 possible, even in asymptomatic cases)⁵²³, ⁵³⁵⁴



- Syrinx or syringomyelia (known or suspected)
 - With neurologic findings and/or predisposing conditions (e.g., Chiari malformation, prior trauma, neoplasm, arachnoiditis, severe spondylosis⁵⁴)
 - With neurologic findings and/or predisposing conditions (e.g., Chiari malformation, prior trauma, neoplasm, arachnoiditis, severe spondylosis⁵⁵)
 - o To further characterize a suspicious abnormality seen on prior imaging
 - Known syrinx with new/worsening symptoms
- Toe walking in a child with signs/symptoms of myelopathy localized to the Cervical Spine
- Suspected neuroinflammatory Conditions/Diseases (e.g., sarcoidosis, Behcet's)
 - After detailed neurological exam and basic testing completed appropriate initial work up
- Initial evaluation of trigeminal neuralgia⁵⁶ not explained on recent Brain imaging

COMBINATION OF STUDIES WITH CERVICAL SPINE MR

Brain MRI/Cervical MRI

• For evaluation of known Arnold-Chiari Malformation

Cervical and Thoracic MRI

- Initial evaluation of known <u>or suspected</u> syrinx or syringomyelia
 - With neurologic findings and/or predisposing conditions (e.g., Chiari malformation, prior trauma, neoplasm, arachnoiditis, severe spondylosis⁵⁵) With neurologic findings and/or predisposing conditions (e.g., Chiari malformation, prior trauma, neoplasm, arachnoiditis, severe spondylosis⁵⁴)
 - To further characterize a suspicious abnormality seen on prior imaging
 - Known syrinx with new/worsening symptom

Any combination of Cervical and/or Thoracic and/or Lumbar MRIs

Note: These body regions might be evaluated separately or in combination as documented in the clinical notes by physical examination findings (e.g., localization to a particular segment of the spinal cord), patient history, and other available information, including prior imaging.

Exception- Indications for combination studies^{55, 56}: Are approved indications as noted below and being performed in children who will need anesthesia for the procedure

<u>Exception- Indications for combination studies^{57, 58}: Are approved indications as noted below and being performed in children who will need anesthesia for the procedure</u>

Any combination of these studies for:



- Survey/complete initial assessment of infant/child with congenital scoliosis or juvenile idiopathic scoliosis under the age of 10⁵⁷⁻⁵⁹59-61 (e.g., congenital scoliosis, idiopathic scoliosis, scoliosis with vertebral anomalies)
- In the presence of neurological deficit, progressive spinal deformity, or for preoperative planning⁶⁰
- Back pain with known vertebral anomalies (hemivertebrae, hypoplasia, agenesis, butterfly, segmentation defect, bars, or congenital wedging) in a child on preliminary imaging
 - In the presence of neurological deficit, progressive spinal deformity, or for preoperative planning⁶²
 - Back pain with known vertebral anomalies (hemivertebrae, hypoplasia, agenesis, butterfly, segmentation defect, bars, or congenital wedging) in a child on preliminary imaging
 - Scoliosis with any of the following⁶¹⁶³:
 - Progressive spinal deformity;
 - Neurologic deficit (new or unexplained);
 - Early onset;
 - Atypical curve (e.g., short segment, > 30' kyphosis, left thoracic curve, associated organ anomalies;)
 - Pre-operative planning; OR
 - When office notes clearly document how imaging will change management
- Arnold-Chiari malformations^{62, 63}64, 65
 - Arnold-Chiari I
 - For evaluation of spinal abnormalities associated with initial diagnosis of Arnold-Chiari Malformation. (C/T/L spine due to association with tethered cord and syringomyelia), and initial imaging has not been completed 50, 5751, 59
 - o Arnold-Chiari II-IV For initial evaluation and follow-up as appropriate
 - Usually associated with open and closed spinal dysraphism, particularly meningomyelocele
- Tethered cord, or spinal dysraphism (known or suspected) based on preliminary imaging, neurological exam, and/or high-risk cutaneous stigmata, when an anesthesia required for imaging (e.g., meningomyelocele, lipomeningomyelocele, diastematomyelia, fatty/thickened filum terminale, and other spinal cord malformations)
- Oncological applications (e.g., primary nervous system, metastatic)
 - Drop metastasis from brain or spine (imaging also includes brain)- see <u>Overview section</u>
 - Suspected leptomeningeal carcinomatosis (LC)⁶⁵—see Suspected leptomeningeal carcinomatosis (LC)⁶⁷ -see Overview section
 - Any combination of these for spinal survey in patient with metastases
 - o Tumor evaluation and monitoring in neurocutaneous syndromes See Overview section
- CSF leak highly suspected and supported by patient history and/or physical exam findings (leak (known or suspected spontaneous (idiopathic) intracranial hypotension (SIH), post lumbar puncture headache, post spinal surgery headache, orthostatic headache, rhinorrhea or otorrhea, or cerebrospinal-venous fistula))



BACKGROUND

Magnetic resonance imaging (MRI) produces high quality multiplanar images of organs and structures within the body without radiation. It is the preferred modality for evaluating the internal structure of the spinal cord, providing assessment of conditions such as degenerative disc pathology, osteomyelitis, and discitis.

OVERVIEW

*Conservative Therapy – (Spine) should include a multimodality approach consisting of a combination of active and inactive components. Inactive components, such as rest, ice, heat, modified activities, medical devices, acupuncture and/or stimulators, medications, injections (epidural, facet, bursal, and/or joint, not including trigger point), and diathermy can be utilized. Active modalities may consist of physical therapy, a physician-supervised home exercise program**, and/or osteopathic manipulative medicine (OMT) or chiropractic care when considered safe and appropriate.

**Home Exercise Program – (HEP)/ Therapy: The following elements are required to meet guidelines for completion of conservative therapy^{66, 67}68, 69:

- Information provided on exercise prescription/plan AND
- Follow-up with member with documentation provided regarding lack of improvement (failed)
 after completion of HEP (after suitable 6-week period), or inability to complete HEP due to
 physical reason- i.e., increased pain, inability to physically perform exercises. (Patient
 inconvenience or noncompliance without explanation does not constitute "inability to
 complete" HEP).
- Dates and duration of failed PT, physician-supervised HEP, or chiropractic treatment should be documented in the original office notes or an addendum to the notes.

Cervical myelopathy – Symptom severity varies, and a high index of suspicion is essential for making the proper diagnosis in early cases. Symptoms of pain and radiculopathy may not be present. The natural history of myelopathy is characterized by neurological deterioration. The most frequently encountered symptom is gait abnormality (86%) followed by increased muscular reflexes (79.1%), pathological reflexes (65.1%), paresthesia of upper limb (69.8%), and pain (67.4%).²⁴

Infection, Abscess, or Inflammatory disease

- Infection:
 - Most common site is the lumbar spine (58%), followed by the thoracic spine (30%) and the cervical spine (11%)⁶⁸
 - High risk populations (indwelling hardware, history of endocarditis, IVDA, recent procedures) with appropriate signs/symptoms



Table 1: Gait and spine imaging 69-7470-75

Gait	Characteristic	Work up/Imaging
Hemiparetic	Spastic unilateral, circumduction	Brain and/or, Cervical spine imaging based on associated symptoms
Diplegic	Spastic bilateral, circumduction	Brain, Cervical and Thoracic Spine imaging
Myelopathic	Wide based, stiff, unsteady	Cervical and/or Thoracic spine MRI based on associated symptoms
<u>Cerebellar</u> Ataxic	Broad based, clumsy, staggering, lack of coordination, usually also with limb ataxia	Brain imaging <u>- see Brain MRI Guideline</u>
Apraxic	Magnetic, shuffling, difficulty initiating	Brain imaging <u>- see Brain MRI Guideline</u>
Parkinsonian	Stooped, small steps, rigid, turning en bloc, decreased arm swing	Brain Imaging - see Brain MRI Guideline
Choreiform	Irregular, jerky, involuntary movements	Medication review, consider brain imaging as per movement disorder Brain MR guidelines
Sensory ataxic	Cautious, stomping, worsening without visual input (ie + Romberg)	EMG, blood work, consider spinal (cervical or thoracic cord imaging) imaging based on EMG
Neurogenic	Steppage, dragging of toes	 EMG initial testing; BUT if there is a foot drop, lumbar spine MRI is appropriate without EMG Pelvis MR if there is evidence of plexopathy
Vestibular	Insecure, veer to one side, worse when eyes closed, vertigo	Consider Brain/IAC MR- see Brain MRI as per GLGuideline

MRI for Evaluation of Discitis — Discitis is a known complication of cervical discography. Postoperative discitis in the cervical spine does not occur frequently but can result from accidental inoculation of bacteria into the disc space intra-operatively by a contaminated spinal needle being used as a radiological marker. There may be other causes for postoperative discitis, e.g., esophageal perforation, hematogenous spread, inoculation of bacteria during surgery. Patients with an alteration in the nature of their symptoms after cervical discectomy and fusion may have discitis. Symptoms may include

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complaints of mild paresthesia in extremities and neck pain. MRI may be performed to reveal feature of discitis with associated abscesses and may help to confirm the diagnosis and decide on further management.

MRI for Cervical Radiculopathy — MRI is a useful test to evaluate the spine because it can show abnormal areas of the soft tissues around the spine; in addition to the bones, it can also show pictures of the nerves and discs and is used to find tumors, herniated discs, or other soft tissue disorders. MRI has a role both in the pre-operative screening and post-operative assessment of radicular symptoms due to either disc or osteophyte.

Table 2: MRI and Cutaneous Stigmata ***

Risk Stratification for Various Cutaneous Markers		
<u>High Risk</u>	<u>Intermediate Risk</u>	<u>Low Risk</u>
 Hypertrichosis Infantile hemangioma Atretic meningocele DST Subcutaneous lipoma Caudal appendage Segmental hemangiomas in association with LUMBAR‡ syndrome 	Capillary malformations (also referred to as NFS or salmon patch when pink and poorly defined or PWS when darker red and well-defined)	 Coccygeal dimple Light hair Isolated café au lait spots Mongolian spots Hypo- and hypermelanotic macules or papules Deviated or forked gluteal cleft Nonmidline lesions

myelopathy, bony defects, anorectal malformations, arterial anomalies, and renal anomalies.

MRI and Multiple Sclerosis (MS) — MRI is a sensitive method of detecting the white matter lesions of MS. These plaques on MRI generally appear as multiple, well-demarcated, homogeneous, small ovoid lesions which often lack mass effect and are oriented perpendicular to the long axis of the lateral ventricles. Sometimes they present as large, space occupying lesions that may be misinterpreted as tumors, abscesses, or infarcts.

MRI and Neck Pain — Neck pain is common in the general population and usually relates to musculoskeletal causes, but it may also be caused by spinal cord tumors. When neck pain is accompanied by extremity weakness, abnormal gait, or asymmetric reflexes, spinal MRI may be performed to evaluate the cause of the pain. MRI may reveal areas of cystic expansion within the spinal cord. Enhancement with gadolinium contrast may suggest that the lesion is neoplastic.



Ossification Posterior Longitudinal Ligament (OPLL)¹⁹¹⁹ – Most common in cervical spine (rare but more severe in thoracic spine)

Neck and Back Pain with Cancer History – Bone is the third most common site of metastases after the liver and the lungs, and approximately two-thirds of all osseous metastases occur in the spine. Approximately 60–70% of patients with systemic cancer will have spinal metastasis. Radiographic (x-ray) examination should be performed in cases of back pain when a patient has a cancer history, but without known active cancer or a tumor that tends to metastasize to the spine. This can make a diagnosis in many cases. This may occasionally allow for selection of bone scan in lieu of MRI in some cases. When radiographs do not answer the clinical question, then MRI may be appropriate after a consideration of conservative care.

"Neoplasms causing VCF (vertebral compression fractures) include: primary bone neoplasms, such as hemangioma or giant cell tumors, and tumor-like conditions causing bony and cellular remodeling, such as aneurysmal bone cysts, or Paget's disease (osteitis deformans); infiltrative neoplasms, including and not limited to, multiple myeloma and lymphoma, and metastatic neoplasms." ⁷⁶

Most common spine metastasis involving primary metastasis originate from the following tumors in descending order: breast (21%), lung (19%), prostate (7.5%), renal (5%), gastrointestinal (4.5%), and thyroid (2.5%). While all tumors can seed to the spine, the cancers mentioned above metastasize to the spinal column early in the disease process. Spinal metastasis is more commonly found in the thoracic region, followed by the lumbar region, while the cervical region is the least likely site of metastasis. 3939

Cervical Spine Trauma Imaging³⁰ — The National Emergency X-Radiography Utilization Study (NEXUS) and the Canadian Cervical Rules (CCR) represent clinical criteria used to help determine the presence of significant cervical spine injury. Although the criteria are highly sensitive (99.6% for NEXUS), specificity is low (12.9% for Nexus).

A patient not meeting any of the NEXUS criteria of focal neurologic deficit, midline spinal tenderness, altered consciousness, intoxication, or distracting injury is unlikely to have a significant cervical spine injury. Imaging evaluation of the cervical spine in these patients is not necessary. In the CCR criteria, a patient without any high risk factors (Age >65 years; paresthesias in extremities; dangerous mechanism; falls from ≥3 feet/5 stairs; axial load to head; motor vehicle crash with high speed, rollover, or ejection; bicycle collision; motorized recreational vehicle accident) is next evaluated for low risk factors (Simple rear-end motor vehicle crash, patient in sitting position in emergency center, patient ambulatory at any time after trauma, delayed onset of neck pain, absence of midline cervical spine tenderness). If the patient meets a low-risk criteria, they are asked to move their head 45 degrees from midline in both directions. If the patient can accomplish this, the spine is cleared, and imaging is not necessary.



MRI and Neurocutaneous Syndromes

- In NF-1, clinical evaluation appears to be more useful to detect complications than is screening imaging in asymptomatic patients. Imaging is indicated in evaluation of suspected tumors based on clinical evaluation and for follow-up of known intracranial tumors.⁷⁷ and intraspinal tumors.⁷⁶
- Conversely in NF-2, routine MR imaging screening is always indicated, given the high prevalence of CNS tumors, especially vestibular schwannomas. In patients with NF-2, routine screening brain/IAC imaging is indicated annually starting from age 10, if asymptomatic, or earlier with clinical signs/symptoms. Most individuals with NF2 eventually develop a spinal tumor, mostly commonly schwannomas, but meningioma and ependymomas are also seen. Spinal imaging at baseline and every 2 to 3 years is also advised with more frequent imaging, if warranted, based on sites of tumor involvement.
- In patients with Tuberous Sclerosis, Brain MRI should be obtained every 1-3 years up until age 25 for surveillance for CNS abnormalities.⁷⁹
- In Von Hippel Lindau Syndrome, imaging of the brain and spinal cord for hemangioblastomas is recommended every 2 years. 80
- In Sturge Weber Syndrome, Brain MRI can rule out intracranial involvement after only age 1 and is recommended in patients <1 year old only if symptomatic.⁸¹
 - Drop Metastases⁸²In patients with Tuberous Sclerosis, Brain MRI should be obtained every 1 3 years up until age 25 for surveillance for CNS abnormalities.⁷⁸
 - In Von Hippel Lindau Syndrome, imaging of the brain and spinal cord for hemangioblastomas is recommended every 2 years. 79
 - In Sturge Weber Syndrome, Brain MRI can rule out intracranial involvement after only age 1 and is recommended in patients <1 year old only if symptomatic.⁸⁰

<u>Drop Metastases</u>⁸¹ – Drop metastases are intradural extramedullary spinal metastases that arise from intracranial lesions. Common examples of intracranial neoplasms that result in drop metastases include pineal tumors, ependymomas, medulloblastomas, germinomas, primitive neuroectodermal tumors (PNET), glioblastomas multiform, anaplastic astrocytomas, oligodendrogliomas, and less commonly choroid plexus neoplasms and teratomas.

Leptomeningeal Carcinomatosis ⁸³⁸² – Leptomeningeal carcinomatosis is complication of cancer in which cancerous cells spread to the membranes (meninges) that covers the brain and spinal cord. The most common solid tumors that involve the leptomeninges are breast, lung, melanoma, gastrointestinal, and primary central nervous system tumors.



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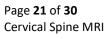


POLICY HISTORY

Date	Summary
May 2023	 Updated references
	 Updated background section
	 Clarified pathological reflexes
	Added trigeminal neuralgia
	 Added "Further evaluation of indeterminate or questionable findings
	on prior imaging":
	 Clarified cerebellar ataxia in gait table
	Removed Additional Resources
	 General Information moved to beginning of guideline with added
	statement on clinical indications not addressed in this guideline
	 Added statement regarding clinical indications not addressed in the
	guideline.
March 2022	Added
	Combination request for overlapping body part statement
	Clarified muscle weakness no related to plexopathy or peripheral
	neuropathy
	 Clarified bowel and bladder dysfunction – not related to an inherent bowel or bladder problem
	Clarified isolated neck pain in pediatric patient
	 Clarified combination MS for cervical and/or thoracic spine combination requests
	 Added subsection for cervical and thoracic spine section for syrinx and syringomyelia
	Descriptions for tethered cord
	Background section of Drop Metastases
	Background section of Leptomeningeal Carcinomatosis
	Clarified toe walking in pediatric patient with myelopathy for cervical
	spine spine
	Removed
	Removed from combination section syrinx and syringomyelia and added
	subsection for cervical and thoracic spine section
	Removed pediatric back pain from the total spine combination section



November 2021	• Added +0698T
April 2021	Added/modified
	 Modified section on neurological deficits
	 Back pain in a child added/modified red flags
	→ Gait table in background
	 Post-surgical modified/clarified surgical criteria for
	combination exams and surgeon preference for exam type
	 Removed myelopathy combination studies
	→ Updated/added MS Criteria
	 Combination section for initial imaging and follow up
	- Added pediatric MS
	Modified known tumor imaging into primary and metastatic
	disease
	→ Added toe walking for pediatric patients
	→ Modified Combination exam wording
	→ Added Achondroplasia to criteria
May 2020	• Added:
,	→ For evaluation of neurologic deficits are new
	Added Imaging of Ossification of the Posterior Longitudinal
1	Ligament (OPPL)
	Added imaging in high risk patients predisposed to spinal
	injury
	 Added imaging in high risk patients for atlantoaxial injury
	Added transverse myelitis
	 Modified Initial imaging of new or increasing non-traumatic
	neck pain or radiculopathy or neck pain that occurs at night
	and wakes the patient from sleep with known active cancer
	and a tumor that tends to metastasize to the spine
	Added to background of imaging of infection
	Added Osteopathic Manipulative medicine to conservative
	care therapy
luna 2010	a Addada
June 2019	Added: - Added:
	new or worsening objective neuro deficits for chronic and
	acute back pain
	○ CSF leak
	 last 6 months for allowable post op f/u period and removed
	EMG-comment
	red flags specifically for peds back pain and pain related to
	malignancy, infection, inflammation





- new sections: pars defect; compression fractures; congenital abnormalities including section on scoliosis and vertebral anomalies in children w/back pain;
- For combination studies cervical/thoracic/lumbar added drop metastasis, tumor evaluation for neurocutaneous syndromes, and abnormalities associated w/Arnold Chiari, as well as separate indication for tethered cord or spinal dysraphism
- Improved section for evaluation of multiple sclerosis including NMO disorders and recurrent transverse myelitis; Lhermitte sign
- Modified section on evaluation of neurologic deficits; added specific pathologic findings; spasticity, sensory, or motor level changes
- Included signs in section on myelopathy including hyperreflexia and pathologic reflexes
- Enhanced sections on trauma; rheumatoid arthritis; back pain in cancer patients with known active cancer in tumors that tend to metastasize to spine
- Expanded on tethered cord in Other Indications for imaging and added section on sacral dimple
- For combination studies Brain/Cervical Spine added suspected MS with new or changing symptoms and follow up to initiation of treatment with known MS



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ADDITIONAL RESOURCES

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Reviewed / Approved by NIA Clinical Guideline Committee

GENERAL INFORMATION

It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.

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