Positional Magnetic Resonance Imaging (MRI)

Effective: April 1, 2017

Next Review: February 2018
Last Review: March 2017

IMPORTANT REMINDER

Medical Policies are developed to provide guidance for members and providers regarding coverage in accordance with contract terms. Benefit determinations are based in all cases on the applicable contract language. To the extent there may be any conflict between the Medical Policy and contract language, the contract language takes precedence.

PLEASE NOTE: Contracts exclude from coverage, among other things, services or procedures that are considered investigational or cosmetic. Providers may bill members for services or procedures that are considered investigational or cosmetic. Providers are encouraged to inform members before rendering such services that the members are likely to be financially responsible for the cost of these services.

DESCRIPTION

Positional MRI (pMRI) acquires images in multiple positions in addition to conventional horizontal imaging. It allows weight-bearing and movement based images.

MEDICAL POLICY CRITERIA

Positional or upright MRI for the diagnosis and management of any condition, including but not limited to cervical, thoracic or lumbosacral back pain, is considered investigational.

NOTE: A summary of the supporting rationale for the policy criteria is at the end of the policy.

CROSS REFERENCES

None

BACKGROUND
It is theorized that imaging the body in positions related to the central loading of the spine, which occurs when standing upright or sitting or in the specific position related to the patient’s clinical symptoms, may lead to more accurate diagnosis. This is being evaluated in patients with suspected nerve root compression and in some cases of spondylolisthesis, and may be particularly relevant in cases where disease is not visible on a horizontal MRI.

One concern with positional MRI is the field strength of the scanners. Today’s clinical MRI scanners may operate at a field strength between 0.1 Tesla (T) to 3 T and are classified as either low-field (<0.5 T), mid-field (0.5-1.0 T), or high-field (>1.0 T). Low- to mid-field MRI is typically used in open scanners. Open scanners are designed for use during interventional or intraoperative procedures, when a conventional design is contraindicated (e.g., an obese or claustrophobic patient), or for changes in patient positioning.

In general, higher field strength results in an increase in signal-to-noise ratio, spatial resolution, contrast and speed. Thus, low-field scanners produce poorer-quality images compared to high-field scanners, and the longer acquisition times with low-field scanners increases the possibility of image degradation due to patient movement. However, field strength has less of an effect on the contrast-to-noise ratio, which determines the extent to which adjacent structures can be distinguished from one another. It may be possible, because of the pMRI technology to diagnose conditions not seen on a conventional or open MRIs.

**REGULATORY STATUS**

Several MRI systems have been cleared for marketing by the U.S. Food and Drug Administration (FDA) through the 510(k) process as open or total body systems for positional imaging. One such system is FONAR’s Upright® MRI. FDA product code: LNH.

**EVIDENCE SUMMARY**

In evaluating positional MRI, it is important to evaluate if this approach results in improved clinical management of the patient compared to other standard imaging.

Randomized controlled trials (RCTs) are needed to determine the following:

- Characteristics of patients who might benefit from positional MRI studies;
- Clinical benefit of basing treatment decisions, including surgery, on these additional findings; and
- How this technique might replace current diagnostic tests such as myelography.

**SYSTEMATIC REVIEWS**

Earlier systematic reviews (SR)s indicate that the literature on kinetic MRI consists primarily of examining anatomic changes in neutral, flexion, extension, and axial rotation. These do not address how anatomical changes detected by positional MRI lead to improved health outcomes. Lord et al. reviewed 16 studies using kinetic MRI on the cervical spine and identified changes in neuroforaminal size, cord compression, cord length, cross-sectional area,
ligamentum flavum thickness, and motion at the index and adjacent levels when comparing symptomatic and asymptomatic individuals\cite{1}. Lao et al. reviewed 11 studies that used kinetic MRI on lumbar and cervical spine on symptomatic individuals, but reported high study bias, since all studies were performed at the same institution.\cite{2} Both these reviews highlighted the need for studies comparing kinetic and traditional MRI in order to establish the clinical utility of kinetic MRI.

A SR of emerging MRI technologies for musculoskeletal imaging under loading stress was prepared by the Tufts Medical Center Evidence-based Practice Center for the Agency for Healthcare Research and Quality (AHRQ) in 2011.\cite{3} The review analyzed 57 studies using MRI under physiologic loading stress in an upright or sitting position or under axial load using a compression device. The majority of studies (37 cross-sectional studies and 13 case-control studies) reported on the anatomical measurements rather than patient-relevant endpoints. The most commonly imaged body region was the lumbar spine. Fifteen of 57 studies used at least two imaging tests and reported on diagnostic or patient-relevant outcomes but did not report meaningful information on the relative performance of the tests. The potential effect on image quality of low magnetic field strengths (<0.6 Tesla) in weight-bearing MRI scanners was not assessed. In 10 studies that included information on adverse effects, 5% to 15% of participants reported new-onset or worsening pain and neuropathy during MRI under loading stress. The SR concluded that, despite the large number of available studies, the evidence is insufficient to support the clinical utility of MRI under loading stress for musculoskeletal conditions.

RANDOMIZED CONTROLLED TRIALS

There are no RCTs comparing how the use of positional MRI versus other standard imaging alters patient treatment plans or health outcomes.

SECTION SUMMARY

A number of studies reported that positional MRI may show abnormalities in patients which were not seen with conventional MRI. However, no studies indicated how these results changed or affected the patient’s treatment plan thereby impacting health outcomes.\cite{4-33}

PRACTICE GUIDELINE SUMMARY

AMERICAN COLLEGE OF RADIOLOGY, AMERICAN SOCIETY OF NEURORADIOLOGY AND SOCIETY OF COMPUTED BODY TOMOGRAPHY AND MAGNETIC RESONANCE\cite{34}

In 2014, the American College of Radiology (ACR) published an updated guideline in collaboration with the American Society of Neuroradiology (ASNR) and the Society of Computed Body Tomography and Magnetic Resonance (SCBT-MR).\cite{34} This guideline states that there is insufficient supportive evidence that the use of positional MRI (referred in the guideline as kinematic or dynamic) correlates with individual patient symptoms or improves patient outcomes after therapy.

SUMMARY
There is not enough research to show that positional or upright MRI improves health outcomes for people with any condition, including but not limited to cervical, thoracic or lumbosacral back pain. In addition, no practice guidelines recommend positional or upright MRI. Therefore, positional MRI is considered investigational.

REFERENCES


### CODES

There is no specific code for positional MRI, which should be reported with an unlisted procedures code such as 76498.

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*Date of Origin: February 2006*