

# MEDICAL POLICY

MEDICAL POLICY DETAILS	
Medical Policy Title	<b>Lumbar Traction: Vertebral Axial Decompression and Home Lumbar Traction Devices</b>
Policy Number	<b>1.01.50</b>
Category	<b>Technology Assessment</b>
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Product Disclaimer	<ul style="list-style-type: none"> <li>• <i>If a product excludes coverage for a service, it is not covered, and medical policy criteria do not apply.</i></li> <li>• <i>If a commercial product (including an Essential Plan or Child Health Plus product), medical policy criteria apply to the benefit.</i></li> <li>• <i>If a Medicaid product covers a specific service, and there are no New York State Medicaid guidelines (eMedNY) criteria, medical policy criteria apply to the benefit.</i></li> <li>• <i>If a Medicare product (including Medicare HMO-Dual Special Needs Program (DSNP) product) covers a specific service, and there is no national or local Medicare coverage decision for the service, medical policy criteria apply to the benefit.</i></li> <li>• <i>If a Medicare HMO-Dual Special Needs Program (DSNP) product DOES NOT cover a specific service, please refer to the Medicaid Product coverage line.</i></li> </ul>

## POLICY STATEMENT

Based upon our criteria and assessment of the peer-reviewed literature, lumbar traction by any method has not been medically proven to be effective and, therefore, is considered **investigational** for all indications, including low back pain.

*Refer to Corporate Medical Policy #1.01.00 Durable Medical Equipment – Standard and Non-Standard.*

*Refer to Corporate Medical Policy #11.01.03 Experimental or Investigational Services.*

## DESCRIPTION

Lumbar traction, a widely used treatment for low back pain, often in combination with other modalities, can be provided manually by a therapist or by mechanical means and may be self-administered using portable devices.

Vertebral axial decompression has been investigated as a technique to reduce intradiscal pressure and relieve low back pain associated with herniated lumbar discs or degenerative lumbar disc disease. Manufacturers of the VAX-D state the device provides traction without the accompanying abdominal muscular contractions that other types of traction elicit, leading to a condition where there is negative intradiscal pressure. Negative intradiscal pressure is speculated to help heal the annulus in a variety of ways. The patient lies on a powered traction table wearing a pelvic harness and grasping hand grips. The table is slowly extended, and a distraction force is applied via the pelvic harness until the desired tension is reached, followed by a gradual decrease of the tension. The cyclic nature of the treatment allows the patient to withstand stronger distraction forces, compared to static lumbar traction techniques. An individual session typically includes 15 cycles of tension, and 10 to 15 daily treatments may be administered. A number of vertebral axial decompression devices have been marketed.

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Pneumatic lumbar traction devices have been developed for home use. The patient, wearing pelvic and lower rib cage harnesses, lies on a two-piece surface. Pneumatic power provided by a hand pump moves the sections apart, applying traction through the harnesses. Examples are the Saunders Lumbar HomeTrac, Saunders STx, Orthotrac, and Comfort Trac.

**RATIONALE**

A number of manufacturers have received Section 510(k) approval from the FDA to market devices characterized by the FDA as powered traction devices. Examples include, but not limited to: the VAX-D Therapeutic Table in 1996, Decompression Reduction Stabilization (DRS) in 1998, Accu-SPINA System in 2000, DRX9000 in 2003, Lordex Power Traction in 2003, SpineRx-LDM in 2003, Healthstar Elite in 2004, Antalgic-Trak in 2005, and SpineMED S200b/S200c Decompression Table in 2005.

There is little published scientific evidence regarding vertebral axial decompression, and the available studies do not provide evidence that the technology is equal or superior to static lumbar traction or to other conservative treatments or surgical interventions for low back pain (LBP). A single case of sudden progression of lumbar disk protrusion during VAX-D therapy, requiring urgent microdiscectomy, has been reported in the literature (Deen et al., 2003).

There is only limited-quality evidence from studies with small sample sizes and moderate to high risk of bias. The effects shown by these studies are small and are not clinically relevant. Published studies of traction as a treatment for LBP lack methodological rigor and reach conflicting conclusions. A 2013 update of the Cochrane review of traction for LBP, with or without sciatica, included a review of 37 randomized, controlled trials. The authors concluded that traction, either alone or in combination with other treatments, has little or no impact on pain intensity, functional status, global improvement, or return to work among people with LBP. For people with mixed-symptom patterns (acute, subacute, or chronic LBP with or without sciatica), there was low-to-moderate-quality evidence that traction may make little or no difference in pain intensity, functional status, global improvement or return to work, when compared to placebo, sham traction or no treatment. For chronic LBP without sciatica, there was moderate-quality evidence that traction probably makes little or no difference in pain intensity when compared with sham treatment. No studies reported on the effect of traction on functional status, global improvement or return to work. Adverse effects were reported in seven of the 32 studies.

The American College of Physicians (2017) published a clinical practice guideline indicating that evidence was low-quality and insufficient to determine the effectiveness of traction for LBP.

The Agency for Healthcare Research and Quality (AHRQ) technology assessment concluded that currently available evidence is too limited in quality and quantity to allow for formulation of evidence-based conclusions regarding the efficacy of decompression therapy as a therapy for chronic back pain when compared with other nonsurgical treatment options (Jurecki-Tiller, 2007).

**CODES**

- *Eligibility for reimbursement is based upon the benefits set forth in the member’s subscriber contract.*
- **CODES MAY NOT BE COVERED UNDER ALL CIRCUMSTANCES. PLEASE READ THE POLICY AND GUIDELINES STATEMENTS CAREFULLY.**
- *Codes may not be all inclusive as the AMA and CMS code updates may occur more frequently than policy updates.*
- *Code Key: Experimental/Investigational = (E/I), Not medically necessary/ appropriate = (NMN).*

**CPT Codes**

<b>Code</b>	<b>Description</b>
No specific code	

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## HCPCS Codes

Code	Description
E0830 (E/I)	Ambulatory traction device, all types, each
S9090 (E/I)	Vertebral axial decompression, per session

## ICD10 Codes

Code	Description
	Investigational for all codes

## REFERENCES

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\*Key Article

**KEY WORDS**

Home lumbar traction device, Lumbar traction, VAX-D, DRS System, DRX, HomeTrac, Orthotrac, ComforTrac, Saunders, Motorized Spinal Traction, Intervertebral Differential Dynamics.

**CMS COVERAGE FOR MEDICARE PRODUCT MEMBERS**

There is currently a National Coverage Determination (NCD) for Lumbar Traction. Please refer to the following NCD websites for Medicare Members:

Vertebral Axial Decompression: [<http://www.cms.gov/medicare-coverage-database/details/ncd-details.aspx?NCDId=124&ncdver=1&bc=AgAAgAAAAAAA&>] accessed 5/19/23

Traction Equipment: <http://www.cms.gov/medicare-coverage-database/details/ncd-details.aspx?NCDId=190&ncdver=2&NCAId=3&ver=5&NcaName=Air-Fluidized+Beds+for+Pressure+Ulcers&bc=ACAAAAAIAAA&>] accessed 5/19/23