

Medical Policy:

Vertical Expandable Prosthetic Titanium Rib

POLICY NUMBER	LAST REVIEW
MG.MM.SU.67C7	March 8, 2024

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The treating physician or primary care provider must submit to EmblemHealth, or ConnectiCare, as applicable (hereinafter jointly referred to as “EmblemHealth”), the clinical evidence that the member meets the criteria for the treatment or surgical procedure. Without this documentation and information, EmblemHealth will not be able to properly review the request preauthorization or post-payment review. The clinical review criteria expressed below reflects how EmblemHealth determines whether certain services or supplies are medically necessary. This clinical policy is not intended to pre-empt the judgment of the reviewing medical director or dictate to health care providers how to practice medicine. Health care providers are expected to exercise their medical judgment in rendering appropriate care. Health care providers are expected to exercise their medical judgment in rendering appropriate care.

EmblemHealth established the clinical review criteria based upon a review of currently available clinical information (including clinical outcome studies in the peer reviewed published medical literature, regulatory status of the technology, evidence-based guidelines of public health and health research agencies, evidence-based guidelines and positions of leading national health professional organizations, views of physicians practicing in relevant clinical areas, and other relevant factors). EmblemHealth expressly reserves the right to revise these conclusions as clinical information changes and welcomes further relevant information. Each benefit program defines which services are covered. The conclusion that a particular service or supply is medically necessary does not constitute a representation or warranty that this service or supply is covered and/or paid for by EmblemHealth, as some programs exclude coverage for services or supplies that EmblemHealth considers medically necessary.

If there is a discrepancy between this guideline and a member's benefits program, the benefits program will govern. Identification of selected brand names of devices, tests and procedures in a medical coverage policy is for reference only and is not an endorsement of any one device, test or procedure over another. In addition, coverage may be mandated by applicable legal requirements of a state, the Federal Government or the Centers for Medicare & Medicaid Services (CMS) for Medicare and Medicaid members. All coding and web site links are accurate at time of publication.

EmblemHealth may also use tools developed by third parties, such as the MCG™ Care Guidelines, to assist us in administering health benefits. The MCG™ Care Guidelines are intended to be used in connection with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice. EmblemHealth Services Company, LLC, has adopted this policy in providing management, administrative and other services to EmblemHealth Plan, Inc., EmblemHealth Insurance Company, EmblemHealth Services Company, LLC, and Health Insurance Plan of Greater New York (HIP) related to health benefit plans offered by these entities. ConnectiCare, an EmblemHealth company, has also adopted this policy. All of the aforementioned entities are affiliated companies under common control of EmblemHealth Inc.

Definitions

Vertical expandable prosthetic titanium rib (VEPTR)	Curved rod placed horizontally in the chest to shape the thoracic cavity for the treatment of spinal and thoracic deformities. In 2014, the FDA Center for Devices and Radiological Health (CDRH) cleared the VEPTR®-VEPTR II™ device for use in skeletally immature patients with severe, progressive spinal deformities and/or three-dimensional deformity of the thorax associated with, or at risk of, <u>Thoracic Insufficiency Syndrome (TIS)</u> .
Cobb angle	Measurement of the degree of spinal curvature; the Cobb angle is considered the standard measurement to quantify a scoliosis for the purpose of measuring curve progression over time. A curve is considered to be scoliosis at a Cobb angle of $\geq 10^\circ$. Any increase $\geq 5^\circ$ is regarded as a significant change; indicative of curvature progression with scoliosis considered mild at 10° – 24° , moderate at 25° – 50° and severe at $> 50^\circ$ in skeletally mature individuals. Cobb angles $> 45^\circ$ are considered severe in skeletally immature persons.
Ellis-van Creveld syndrome	Autosomal recessive genetic disorder characterized by skeletal dysplasia.
Hypoplastic thorax syndrome	Examples of the syndrome include achondroplasia, Ellis van Creveld syndrome, Jarcho-Levin syndrome and Jeune's syndrome.

Jarcho-Levin syndrome	Heritable axial skeleton growth disorder associated with malformation of the vertebral column and ribs.
Jeune syndrome	Congenital dwarfism associated with asphyxiating thoracic dystrophy.
Scoliosis	Musculoskeletal condition characterized by an abnormal lateral curvature of the spine. There are several different types of scoliosis that affect children and adolescents. The most common type is considered idiopathic but additional types of scoliosis include congenital, neuromuscular and syndromic scoliosis.
Thoracic Insufficiency Syndrome (TIS)	Rare condition defined as, "The inability of the thorax to support normal respiration or lung growth. This would include patients with progressive congenital, neuromuscular, idiopathic, or syndromic scoliosis" (FDA, 2014). TIS may include flail chest syndrome, <u>hypoplastic thorax syndrome</u> , as well as rib fusion and scoliosis.

Related Medical Guideline

Related Medical Guideline

[Surgical Correction of Chest Wall Deformities](#)

Guideline

The VEPTR is considered medically necessary in the treatment of progressive thoracic insufficiency syndrome due to rib and/or chest wall defects in infants and children between 6 months of age and skeletal maturity.

Rib/chest wall defects may be secondary to any of the following scoliosis conditions:

1. Congenital scoliosis
2. Neuromuscular scoliosis
3. Infantile and juvenile idiopathic scoliosis
4. Syndromic scoliosis

Limitations and Exclusions

1. Use of VEPTR for any condition other than those listed above (including Poland Syndrome) is not considered medically necessary due to insufficient evidence of therapeutic value.
2. Use of VEPTR as a scoliosis treatment in the absence of TIS (or risk for TIS) is not considered medically necessary.

Procedure Codes

20999	Unlisted procedure, musculoskeletal system, general
21899	Unlisted procedure, neck or thorax

Diagnosis Codes

M41.00	Infantile idiopathic scoliosis, site unspecified
M41.02	Infantile idiopathic scoliosis, cervical region

M41.03	Infantile idiopathic scoliosis, cervicothoracic region
M41.04	Infantile idiopathic scoliosis, thoracic region
M41.05	Infantile idiopathic scoliosis, thoracolumbar region
M41.06	Infantile idiopathic scoliosis, lumbar region
M41.07	Infantile idiopathic scoliosis, lumbosacral region
M41.08	Infantile idiopathic scoliosis, sacral and sacrococcygeal region
M41.11	Juvenile idiopathic scoliosis, cervical region
M41.112	Juvenile idiopathic scoliosis, cervicothoracic region
M41.113	Juvenile idiopathic scoliosis, cervicothoracic region
M41.114	Juvenile idiopathic scoliosis, thoracic region
M41.115	Juvenile idiopathic scoliosis, thoracolumbar region
M41.116	Juvenile idiopathic scoliosis, lumbar region
M41.117	Juvenile idiopathic scoliosis, lumbosacral region
M41.119	Juvenile idiopathic scoliosis, site unspecified
M41.40	Neuromuscular scoliosis, site unspecified
M41.41	Neuromuscular scoliosis, occipito-atlanto-axial region
M41.42	Neuromuscular scoliosis, cervical region
M41.43	Neuromuscular scoliosis, cervicothoracic region
M41.44	Neuromuscular scoliosis, thoracic region
M41.45	Neuromuscular scoliosis, thoracolumbar region
M41.46	Neuromuscular scoliosis, lumbar region
M41.47	Neuromuscular scoliosis, lumbosacral region
Q67.5	Congenital scoliosis NOS
Q76.3	Congenital scoliosis due to congenital bony malformation

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Revision History

Company(ies)	DATE	REVISION
ConnectiCare	Feb. 2021	ConnectiCare adopts the clinical criteria of its parent corporation EmblemHealth