Medical Policy:
Prior Authorization Criteria
Osteogenic Stimulator Low Intensity Ultrasound, Non-Invasive (Commercial)

POLICY NUMBER	EFFECTIVE DATE	APPROVED BY
M20190018	5/1/2019	MPC (Medical Policy Committee)

IMPORTANT NOTE ABOUT THIS MEDICAL POLICY:

 The treating physician or primary care provider must submit to ConnectiCare, Inc. the clinical evidence that the patient meets the criteria for the treatment or surgical procedure. Without this documentation and information, ConnectiCare will not be able to properly review the request for prior authorization. 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. The clinical review criteria expressed below reflects how ConnectiCare determines whether certain services or supplies are medically necessary. ConnectiCare 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). ConnectiCare, Inc. expressly reserves the right to revise these conclusions as clinical information changes, and welcomes further relevant information. 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. Each benefit plan 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 ConnectiCare, as some plans exclude coverage for services or supplies that ConnectiCare considers medically necessary. If there is a discrepancy between this guideline and a member's benefits plan, the benefits plan will govern. In addition, coverage may be mandated by applicable legal requirements of the State of CT and/or the Federal Government. Coverage may also differ for our Medicare members based on any applicable Centers for Medicare & Medicaid Services (CMS) coverage statements including including National Coverage Determinations (NCD), Local Coverage Determinations (LCD) and/or Local Medical Review Policies(LMRP). All coding and web site links are accurate at time of publication.

Criteria: Effective 1/1/2020

Requests will be considered using Milliman Clinical Care Guidelines (MCGs). Guidelines are developed using publications that have been assessed in terms of quality, utility, and relevance. Preference is given to publications that:

- Are designed with rigorous scientific methodology.
- Are published in higher-quality journals (e.g., journals that are read and cited most often within their field).
- Address an aspect of specific importance to the guideline in question (admission criteria, length of stay).
- Represent an update or contain new data or information not reflected in the current guideline.

On an annual basis, each guideline undergoes external review by clinically active experts (e.g., board-certified specialist physicians without stated financial conflicts of interest) to confirm the clinical appropriateness, accuracy, validity, and applicability of each guideline.

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Click Here: ConnectiCare-MCG Clinical Criteria

A-0414 AC ACG Bone Growth Stimulators, Ultrasonic

CPT code 20979

HCPC code E0760

Revision history

DATE	REVISION
07/27/2021	Updated policy with CPT/HCPC codes and MCG Guideline Codes
01/01/2020	Adopted MCG Clinical Care Guidelines