Policy Name: Synthetic Ceramic-Based and Bioactive Glass Bone Substitutes/Fillers
Effective Date: 12/16/2019

Important Information – Please Read Before Using This Policy

These services may or may not be covered by all Medica plans. Please refer to the member’s plan document for specific coverage information. If there is a difference between this general information and the member’s plan document, the member’s plan document will be used to determine coverage. With respect to Medicare and Minnesota Health Care Programs, this policy will apply unless those programs require different coverage. Members may contact Medica Customer Service at the phone number listed on their member identification card to discuss their benefits more specifically. Providers with questions about this Medica coverage policy may call the Medica Provider Service Center toll-free at 1-800-458-5512.

Medica coverage policies are not medical advice. Members should consult with appropriate health care providers to obtain needed medical advice, care and treatment.

Coverage Policy

Synthetic ceramic-based and bioactive glass bone substitutes/fillers, used singly or in combination with other grafts*, are investigative and therefore NOT COVERED.

* Note: Medica considers the following to be investigative and therefore not covered for orthopedic applications: (1) Autologous blood-derive biologics (e.g., platelet-rich plasma, autologous conditioned serum, autologous whole blood), (2) stem cell therapy (e.g. AlloStem®, Cellentra™ VCBM, Osteocel® Plus, Trinity® Evolution™), and (3) OsteoAmp™ allogeneic morphogenic protein.

Note: This policy does not apply to dental applications.

See related Medica Utilization Management policies Lumbar Spine Surgeries and Cervical Spine Surgeries for specific medical necessity criteria.

See also related Medica coverage policies: (1) Autologous Blood-Derived Products (Platelet-Rich Plasma, Autologous Conditioned Serum, Autologous Whole Blood; (2) Stem Cell Therapy for Orthopedic Applications; (3) Recombinant Human Bone Morphogenic Protein-2 (rhBMP-2)/InFUSE and Allogeneic Morphogenic Protein (e.g., OsteoAMP™); (4) Motion Preserving Posterior Interspinous/Interlaminar Decompression/Stabilization Devices (e.g., X-Stop, Coflex, Dynesys, DIAM spinal stabilization, Wallis system, total facet joint replacement system); (5) mild® Procedure (mild® Device Kit); (6) Laser Spine Surgery, (7) Sacroiliac Joint Fusion, Open and Minimally Invasive, (8) Percutaneous Disc Decompression Procedures (Manual, Automated or Laser Discectomy, and Plasma Disc Decompression [PDD]), and (9) Percutaneous Radiofrequency and Laser Ablation/Denervation Procedures for Facet and Sacroiliac Joints.

Description

Ceramics and bioactive glass fillers are synthetically produced bone substitutes/extenders and void fillers used to fill voids and gaps in bone structure. These may be located in the extremities, spine, pelvis, or cranium. These products can be obtained as injectables, pastes, putties, solid matrices, and granules. Ceramics are made by a process called sintering - a process using high temperatures to extract individual crystals that fuse into grains of varying sizes.
Ceramic-based bone grafts are made up of collagen, calcium phosphate (CaP), calcium sulfate and one or more of the following products:
1. Synthetic hydroxyapatite (HA), a component in bone and teeth
2. Beta-tricalcium phosphates (β-TCP)
3. Biphasic calcium phosphate (BCP), which consists of both HA and β-TCP
These have been proposed for use as stand-alone products or in combination with other bone substitutes and/or enhancement products (e.g., platelet rich plasma, bone morphogenic protein, demineralize bone matrices).

**FDA Approval**
Multiple synthetic ceramic-based and bioactive glass bone substitutes/fillers have received FDA approval, mostly through the 510(k) clearance process. To locate marketing clearance information for a specific device or manufacturer, search the Center for Devices and Radiological Health (CDRH) 510(k) database or the Premarket Approval (PMA) database by product and/or manufacturer name, located at: [https://www.accessdata.fda.gov/scripts/cdrh/devicesatfda/](https://www.accessdata.fda.gov/scripts/cdrh/devicesatfda/).

Examples of synthetically produced beta tricalcium phosphate bone fillers include, but are not limited to:
1. Conduit TCP Granules
2. FM-O2
3. Formagraft Bone Graft
4. GenerOs
5. GranOS
6. Integra Mozaik
7. Integra OS
8. IsoTis Mozaik
9. OSferion
10. Osteoconductive Scaffold
11. OsteoStrux
12. OsSatura TCP
13. Osteomatrix
14. OsteoVation B-TCP
15. Synthes ChronOS
16. Vitoss
17. Vitoss Bioactive Foam-2X.

Examples of synthetic bioactive glass bone fillers include, but are not limited to:
1. Bi-Ostetic Bioactive Glass
2. BioSphere Flex
3. BioSphere Putty Bone Graft
4. BonAlive
5. CLM BioActive Scaffold
6. FIBERGRAFT BG Morsels
7. Interface
8. NanoFUSE
9. NovaBone
10. NovaBone BIOACTIVE Strip
11. NovaBone-C/M
12. Osteofuse Bioactive
13. PerioGlas
Examples of synthetic nanocrystalline hydroxyapatite bone fillers include, but are not limited to:
1. Beta-BSM (injectable form)
2. CarriGen
3. Cem-Ostetic
4. Gamma-BSM moldable putty
5. NanOss Bioactive.

Prior Authorization
Prior authorization is not applicable. Claims for this service are subject to retrospective review and denial of coverage, as investigative services are not eligible for reimbursement.

Coding Considerations
Use the current applicable CPT/HCPCS code(s). The following codes are included below for informational purposes only, and are subject to change without notice. Inclusion or exclusion of a code does not constitute or imply member coverage or provider reimbursement.

HCPC Codes:
- **C9359** - Porous purified collagen matrix bone void filler (Integra Mozaik Osteoconductive Scaffold Putty, Integra OS Osteoconductive Scaffold Putty), per 0.5 cc
- **C9362** - Porous purified collagen matrix bone void filler (Integra Mozaik Osteoconductive Scaffold Strip), per 0.5 cc

Original Effective Date: 12/16/2019

Re-Review Date(s):