POLICIES AND PROCEDURE MANUAL

Policy: MP090
Section: Medical Benefit Policy
Subject: Injectable Bulking Agents for Treatment of Urinary Incontinence

I. Policy: Injectable Bulking Agents for Treatment of Urinary Incontinence

II. Purpose/Objective:
To provide a policy of coverage regarding Injectable Bulking Agents for Treatment of Urinary Incontinence

III. Responsibility:
A. Medical Directors
B. Medical Management

IV. Required Definitions
1. Attachment – a supporting document that is developed and maintained by the policy writer or department requiring/authoring the policy.
2. Exhibit – a supporting document developed and maintained in a department other than the department requiring/authoring the policy.
3. Devised – the date the policy was implemented.
4. Revised – the date of every revision to the policy, including typographical and grammatical changes.
5. Reviewed – the date documenting the annual review if the policy has no revisions necessary.

V. Additional Definitions
Medical Necessity or Medically Necessary means Covered Services rendered by a Health Care Provider that the Plan determines are:

a. appropriate for the symptoms and diagnosis or treatment of the Member’s condition, illness, disease or injury;
b. provided for the diagnosis, and the direct care and treatment of the Member’s condition, illness disease or injury;
c. in accordance with current standards of good medical treatment practiced by the general medical community.
d. not primarily for the convenience of the Member, or the Member’s Health Care Provider; and
e. the most appropriate source or level of service that can safely be provided to the Member. When applied to hospitalization, this further means that the Member requires acute care as an inpatient due to the nature of the services rendered or the Member’s condition, and the Member cannot receive safe or adequate care as an outpatient.

Medicaid Business Segment
Medical Necessity shall mean a service or benefit that is compensable under the Medical Assistance Program and if it meets any one of the following standards:

(i) The service or benefit will, or is reasonably expected to, prevent the onset of an illness, condition or disability.
(ii) The service or benefit will, or is reasonably expected to, reduce or ameliorate the physical, mental or development effects of an illness, condition, injury or disability.
(iii) The service or benefit will assist the Member to achieve or maintain maximum functional
capacity in performing daily activities, taking into account both the functional capacity of the Member and those functional capacities that are appropriate for members of the same age.

DESCRIPTION:
Periurethral injections of bulking agents such as collagen and synthetic agents (e.g. Durasphere™) are used for the management of patients with urinary incontinence resulting from intrinsic sphincter deficiency.

INDICATIONS:
The use of injectable bulking agents may be considered medically necessary when documented evidence of urinary incontinence refractory to prior non-invasive treatments (e.g., Kegel exercises and pharmacologic agents). Types of injectable bulking agents that have received FDA approval include but are not limited to the following:

1. Collagen implants
2. Carbon-coated beads
3. Ethylene vinyl alcohol copolymer
4. Spherical particles of calcium hydroxylapatite

LIMITATIONS:
A pre-treatment skin test for the bulking agent with no evidence of local bovine hypersensitivity is required prior to collagen implantation.

EXCLUSIONS:
There is insufficient evidence in the peer-reviewed published medical literature to establish the effectiveness of other agents, as periurethral bulking agents, including but not limited to Teflon®, autologous fat, autologous ear chondrocytes or silicone microimplants. The Plan does NOT provide coverage of these agents because they are considered experimental, investigational or unproven.

Note: A complete description of the process by which a given technology or service is evaluated and determined to be experimental, investigational or unproven is outlined in MP 15 - Experimental Investigational or Unproven Services or Treatment.

CODING ASSOCIATED WITH:
The following codes are included below for informational purposes and may not be all inclusive. Inclusion of a procedure or device code(s) does not constitute or imply coverage nor does it imply or guarantee provider reimbursement. Coverage is determined by the member specific benefit plan document and any applicable laws regarding coverage of specific services. Please note that per Medicare coverage rules, only specific CPT/HCPCS Codes may be covered for the Medicare Business Segment. Please consult the CMS website at www.cms.gov or the local Medicare Administrative Carrier (MAC) for more information on Medicare coverage and coding requirements.

51715 Endoscopic injection of implant material into the submucosal tissues of the urethra and/or bladder neck
L8603 Injectable bulking agent, collagen implant, urinary tract, 2.5 ml syringe
L8606 Injectable bulking agent, synthetic implant, urinary tract, 1ml syringe
95028 Intracutaneous (intradermal) tests with allergenic extracts, delayed type reaction, including reading

ICD10 Codes:
N39.3

LINE OF BUSINESS:
Eligibility and contract specific benefits, limitations and/or exclusions will apply. Coverage statements found in the line of business specific benefit document will supersede this policy. For Medicare, applicable LCD’s and NCD’s will supersede this policy. For PA Medicaid Business segment, this policy applies as written.

REFERENCES:


ECRI Hotline Response: Injectable Urethral Bulking Agents for the Treatment of Urinary Stress Incontinence. 6/4/02


This policy will be revised as necessary and reviewed no less than annually.

**Devised:** 11/93 “Contigen Implant”

**Revised:** 6/96, 2/98, 12/02 revised as “Bulking Agents for Treatment of Urinary Incontinence”; 1/04 Coding, definition; 2/06; 2/07; 2/09 (key words)

**Reviewed:** 2/05, 2/08, 2/10, 2/11, 2/12, 2/13, 2/14, 2/15, 2/16, 2/17, 2/18