

Treatment of Abnormal Uterine Bleeding and Fibroids

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Approved By:	Highmark Health Options – Market Leadership
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Products:	Medicaid
Application:	All participating hospitals and providers
Page Number(s):	1 of 6

Disclaimer

Highmark Health Options medical policy is intended to serve only as a general reference resource regarding coverage for the services described. This policy does not constitute medical advice and is not intended to govern or otherwise influence medical decisions.

POLICY STATEMENT

Highmark Health Options may provide coverage under medical surgical benefits of the Company's Medicaid products for medically necessary treatment of abnormal uterine bleeding and fibroids.

This policy is designed to address medical necessity guidelines that are appropriate for the majority of individuals with a particular disease, illness, or condition. Each person's unique clinical circumstances warrant individual consideration, based upon review of applicable medical records.

The qualifications of the policy will meet the standards of the National Committee for Quality Assurance (NCQA) and the Delaware Department of Health and Social Services (DHSS) and all applicable state and federal regulations.

DEFINITIONS

Highmark Health Options (HHO) – Managed care organization serving vulnerable populations that have complex needs and qualify for Medicaid. Highmark Health Options members include individuals and families with low income, expecting mothers, children, and people with disabilities. Members pay nothing to very little for their health coverage. Highmark Health Options currently services Delaware Medicaid: Delaware Healthy Children Program (DHCP) and Diamond State Health Plan Plus members.

Abnormal Uterine Bleeding – Abnormal uterine bleeding (AUB) is bleeding from the uterus between periods. It may also be prolonged bleeding during a period or an extremely heavy period (menorrhagia).

Uterine Fibroids – Uterine fibroids are noncancerous growths of the uterus. Symptoms may include menorrhagia, pelvic pressure, or pain. Treatment options include hysterectomy, myomectomy, uterine artery embolization and fibroid ablation.

PROCEDURES

A prior authorization is required.

Treatment of uterine fibroids is considered experimental/investigational and therefore noncovered for any ONE of the following procedures/services:

- Laparoscopic and percutaneous techniques for myolysis (e.g., laser and bipolar needles, cryomyolysis); or
- Laparoscopic uterine power morcellation in hysterectomy and myomectomy; or
- MRI guidance performed in conjunction with percutaneous myolysis of uterine fibroids.

Transcatheter uterine artery embolization (UAE) of uterine arteries may be considered medically necessary for the treatment of uterine fibroids when any ONE of the following criteria is met:

- The individual is experiencing the following symptoms:
 - Menorrhagia (excessive menstrual bleeding lasting more than eight (8) days) as a direct result of the fibroid (i.e., not resulting from hyperplasia, atypia, or cancer) that interferes with daily activities or causes anemia; or
 - Pelvic pain or pressure as a direct result of the fibroid; or
 - Lower back pain as a direct result of the fibroid; or
 - Urinary symptoms (e.g., urinary frequency, urgency) related to compression of the bladder as a direct result of the fibroid; or
 - Gastrointestinal symptoms related to compression of the bowel (e.g., constipation, bloating) as a direct result of the fibroid; or
 - Dyspareunia (painful or difficult sexual relations) as a direct result of the fibroid; or
 - An abdominally palpable fibroid; or
- The individual is asymptomatic with an abdominally palpable fibroid or significantly enlarged fibroid on abdominal/vaginal examination and any ONE of the following:
 - The use of anesthesia places the individual at high surgical risk; or
 - The individual has medical contraindications to hysterectomy (e.g., morbid obesity); or
 - The use of hormonal therapy is contraindicated, or the individual is intolerant to or has previously failed a course of hormone therapy; or
 - The individual wishes to avoid hysterectomy; or
 - The individual may want to become pregnant; or
 - The individual has hydronephrosis.

One repeat transcatheter embolization of uterine arteries may be considered medically necessary to treat persistent symptoms of uterine fibroids after an initial uterine artery embolization when any ONE of the following criteria is met:

- Documentation of continued symptoms such as bleeding or pain; or
- Individual has persistent symptoms in combination with findings on imaging of an incomplete initial procedure, as evidenced by continued blood flow to the treated regions.

UAE may be considered medically necessary for the treatment of postpartum uterine hemorrhage.

UAE is considered experimental/investigational for all other indications and therefore noncovered because the safety and/or effectiveness have not been established by the available published peer-reviewed literature.

Laparoscopic ultrasound-guided radiofrequency ablation (e.g., Acessa™) for the treatment of uterine fibroids may be considered medically necessary when the individual is experiencing any ONE of the following symptoms:

- Menorrhagia (excessive menstrual bleeding lasting more than eight (8) days) as a direct result of the fibroid (e.g., not resulting from hyperplasia, atypia, or cancer) that interferes with daily activities or causes anemia; or
- Pelvic pain or pressure as a direct result of the fibroid; or
- Lower back pain as a direct result of the fibroid; or
- Urinary symptoms (e.g., urinary frequency, urgency) related to compression of the bladder as a direct result of the fibroid; or
- Gastrointestinal symptoms related to compression of the bowel (e.g., constipation, bloating) as a direct result of the fibroid; or
- Dyspareunia (painful or difficult sexual relations) as a direct result of the fibroid; or
- An abdominally palpable fibroid.

Laparoscopic ultrasound-guided radiofrequency ablation is considered experimental/investigational for all other indications not listed above and therefore noncovered because the safety and/or effectiveness have not been established by the available published peer-reviewed literature.

Endometrial ablation with or without hysteroscopic guidance, using an FDA-approved device, may be considered medically necessary in women who would otherwise be considered candidates for hysterectomy when any ONE of the following criteria is met:

- In women with menorrhagia who are not candidates for hormone therapy; or
- Decline hormone therapy; or
- Who are unresponsive to hormone therapy.

Endometrial ablation with or without hysteroscopic guidance for all other indications is considered not medically necessary.

POST-PAYMENT AUDIT STATEMENT

The medical record must include documentation that reflects the medical necessity criteria and is subject to audit by Highmark Health Options at any time pursuant to the terms of your provider agreement.

PLACE OF SERVICE: INPATIENT/OUTPATIENT

Experimental/investigational (E/I) services are not covered regardless of place of service.

Treatment of Uterine Fibroids is typically an outpatient procedure which is only eligible for coverage as an inpatient procedure in special circumstances, including, but not limited to, the presence of a comorbid condition that would require monitoring in a more controlled environment such as the inpatient setting.

CODING REQUIREMENTS

CPT code	Description
77022	Magnetic resonance imaging guidance for, and monitoring of, parenchymal tissue ablation.
36245	Selective catheter placement, arterial system; each first order abdominal, pelvic, or lower extremity artery branch, within a vascular family.
36246	Selective catheter placement, arterial system; initial second order abdominal, pelvic, or lower extremity artery branch, within a vascular family.
36247	Selective catheter placement, arterial system; initial third order or more selective abdominal, pelvic, or lower extremity artery branch, within a vascular family.

36248	Selective catheter placement, arterial system; additional second order, third order, and beyond, abdominal, pelvic, or lower extremity artery branch, within a vascular family (list in addition to code for initial second or third order vessel as appropriate).
37243	Vascular embolization or occlusion, inclusive of all radiological supervision and interpretation, intraprocedural road mapping, and imaging guidance necessary to complete the intervention; for tumors, organ ischemia, or infarction.
37244	Vascular embolization or occlusion, inclusive of all radiological supervision and interpretation, intraprocedural road mapping, and imaging guidance necessary to complete the intervention; for arterial or venous hemorrhage or lymphatic extravasation.
75894	Transcatheter therapy, embolization, any method, radiological supervision, and interpretation.
58674	Laparoscopy, surgical, ablation of uterine fibroid(s) including intraoperative ultrasound guidance and monitoring, radiofrequency.
58353	Endometrial ablation, thermal, without hysteroscopic guidance.
58356	Endometrial cryoablation with ultrasonic guidance, including endometrial curettage, when performed.
58563	Hysteroscopy, surgical; with endometrial ablation (e.g., endometrial resection, electro-surgical ablation, thermoablation).

COVERED DIAGNOSIS CODES FOR PROCEDURE CODE 58674

Codes						
D25.0	D25.1	D25.2	D25.9			

COVERED DIAGNOSIS CODES FOR PROCEDURE CODES 75894

Codes						
O43.211	O43.212	O43.213	O43.221	O43.222	O43.223	O43.231
O43.232	O43.233	O44.30	O44.31	O44.32	O44.33	O44.50
O44.51	O44.52	O44.53	O72.0	O72.1	O72.2	

NONCOVERED DIAGNOSIS CODES FOR PROCEDURE CODES 58578, 77022

Codes						
D25.0	D25.1	D25.2	D25.9			

COVERED DIAGNOSIS CODES FOR PROCEDURE CODES 58353, 58356, AND 58563

Codes						
N92.0	N92.1	N92.4				

REIMBURSEMENT

Participating facilities will be reimbursed per their Highmark Health Options contract.

Reference

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POLICY UPDATE HISTORY

10/27/2021	Approved in Medical Policy Committee
11/2021	Approved in QI/UM
10/26/2022	Annual review; approved in Medical Policy Committee
11/2022	Approved in QI/UM