OVERVIEW
Barrett’s esophagus is a condition in which the normal squamous epithelium is replaced by specialized columnar-type epithelium, known as intestinal metaplasia, in response to irritation and injury caused by gastroesophageal reflux disease (GERD). Intestinal metaplasia is a precursor to esophageal adenocarcinoma and may be treated with mucosal ablation techniques such as radiofrequency ablation and cryoablation.

PRIOR AUTHORIZATION
Prior Authorization review is not required

POLICY STATEMENT
BlueCHiP for Medicare and Commercial
Radiofrequency ablation may be considered medically necessary for treatment of Barrett’s esophagus with high-grade dysplasia. The diagnosis of high-grade dysplasia should be confirmed by two pathologists prior to radiofrequency ablation.

Radiofrequency ablation may be considered medically necessary for treatment of Barrett’s esophagus with low-grade dysplasia, when the initial diagnosis of low-grade dysplasia is confirmed by two pathologists, one of whom is an expert in GI Pathology. It is ideal that two experts in GI pathology agree on the diagnosis in order to confirm LGD.

Radiofrequency ablation is considered not medically necessary for treatment of Barrett’s esophagus in the absence of dysplasia due to lack of peer reviewed literature that supports efficacy.

Cryoablation is considered not medically necessary for Barrett’s esophagus, with or without dysplasia due to lack of peer reviewed literature that supports efficacy.

MEDICAL CRITERIA
None.

BACKGROUND
The esophagus is normally lined by squamous epithelium. Barrett’s esophagus occurs in the distal esophagus, may be of any length, may be focal or circumferential, and can be visualized by the endoscopist as being a different color than the background squamous mucosa. Confirmation of Barrett’s esophagus requires biopsy of the columnar epithelium and microscopic identification of intestinal metaplasia.

Intestinal metaplasia is a precursor to esophageal adenocarcinoma, and esophageal adenocarcinoma is thought to result from a stepwise accumulation of genetic abnormalities in the specialized epithelium, which results in the phenotypic expression of histologic features of low-grade dysplasia (LGD) to high-grade dysplasia (HGD) to carcinoma.

The current management of Barrett’s esophagus includes treatment of GERD and surveillance endoscopy to detect progression to HGD or adenocarcinoma. The finding of LGD typically warrants only follow-up and surveillance biopsies, whereas the finding of HGD or early-stage adenocarcinoma warrants mucosal ablation.
or resection (either endoscopic mucosal resection [EMR] or esophagectomy). EMR, either focal or circumferential, provides a histologic specimen for examination and staging (unlike ablative techniques).

Mucosal ablation techniques that are available consist of one of several thermal (multipolar electrocoagulation [MPEC], argon plasma coagulation [APC], heater probe, Nd:YAG laser, KTP-YAG laser, diode laser, argon laser, and cryoablation) or nonthermal (5-aminolevulinic acid [5-ALA] and photofrin photodynamic therapy [PDT]) techniques.

The CryoSpray Ablation™ System (formerly the SprayGenix™ Cryo Ablation System, CSA Medical, Inc.) uses a low-pressure spray for spraying liquid nitrogen through an upper endoscope. Cryotherapy allows for treatment of uneven surfaces; however, disadvantages include the uneven application inherent in spraying the cryogen.

The HALO System from BÀRRX Medical, Inc. (Sunnyvale, Calif.) uses radiofrequency (RF) energy and consists of 2 components: an energy generator and an ablation catheter.

Radiofrequency ablation (RFA) of high-grade dysplasia (HGD) in Barrett’s esophagus has been shown to be at least as effective in eradicating high-grade dysplasia as other ablative techniques with a lower progression rate to cancer and may be considered as an alternative to esophagectomy. Therefore, RFA may be considered medically necessary for patients with Barrett’s esophagus (BE) and HGD.

For patients with low-grade dysplasia (LGD), the benefit of RFA is less certain, as the rate of progression to cancer is variable in the literature. There are no high-quality trials that treat patients with an initial diagnosis of LGD and report improved outcomes. However, based on the available evidence, specialty society guidelines, and the results of clinical vetting, it is possible to define a population with a higher risk of progression by having the initial LGD diagnosis confirmed by an additional pathologist who is an expert in GI pathology. In this subpopulation of patients with LGD, it is likely that the benefit of treatment outweighs the risk. As a result, RFA of LGD may be considered medically necessary when the initial diagnosis of LGD is confirmed by an expert in gastrointestinal (GI) pathology.

For patients with non-dysplastic BE, it cannot be concluded that the benefit of RFA outweighs the risk, and therefore RFA is considered not medically necessary for this population.

Data for the efficacy of cryoablation of BE with or without dysplasia are limited. The studies consist of small numbers of patients with short-term follow-up, and therefore cryoablation of BE is considered not medically necessary.

**COVERAGE**
Benefits may vary between groups/contracts. Please refer to the appropriate member certificate/subscriber agreement for applicable surgery/not medically necessary coverage/benefits

**CODING**
BlueCHiP for Medicare and Commercial
There is no CPT code specific to radiofrequency or cryoablation of tissue in the esophagus. These procedures would likely be coded using one of the following CPT codes and would be medically necessary when filed with a covered diagnosis.

43229
43270

ICD9: 530.85

ICD10: K22.711, K22.719
REFERENCES
