Policy Name: Chemiluminescent Testing (ViziLite®) for Oral Cancer Screening
Effective Date: 1/20/2019

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
Chemiluminescent testing (ViziLite®) for oral cancer screening is investigative and therefore NOT COVERED.

Note: This policy is no longer scheduled for regular review of the scientific literature.

Description
ViziLite® (Zila Inc., Phoenix, AZ) is a chemiluminescent test, which has been proposed as an adjunct to visual examination to increase identification, evaluation, and monitoring of oral mucosal abnormalities in individuals at increased risk for oral cancer. ViziLite is a single-use product that consists of an acetic acid rinse, retractor, and light stick. The patient rinses with the acetic acid solution and expectorates. The ViziLite light stick is activated and inserted into the hollow end of the retractor. After dimming the lights, the provider examines the oral cavity using the ViziLite device. The light is purported to impart a blue hue to normal tissue, while lesions take on an “acetowhite” appearance, thus becoming clinically discernable.

FDA Approval
The ViziLite® test kit is classified by the FDA as a dental operating light and regulated as a Class II (moderate risk) device. This test system received FDA 510(k) premarket approval on January 31, 2005 (K033033). According to 510(k) approval, a standard visual examination must always be performed before use of the ViziLite® test kit.

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.

CPT Codes:
82397 - chemiluminescent assay

Original Effective Date: 3/1/2008

Re-Review Date(s): 10/18/2010
11/26/2013
10/19/2016
10/16/2019