Diagnose and Treat Earlier

Comprehensive Barrett's Esophagus Test (CBEST™)

Improving the Detection of Dysplasia and Adenocarcinoma

Because patients with Barrett's esophagus (BE) are at an increased risk for esophageal adenocarcinoma (EAC), early detection and surveillance are key to optimized patient management and improved patient outcomes. However, accurately identifying and grading the varying degrees of dysplasia, particularly low grade, can be difficult using traditional testing methods.

An Innovative Solution for More Informed Treatment Decisions

- The Comprehensive Barrett's Esophagus Test (CBEST™) is unique combination of cytology, immunohistochemistry (IHC), and fluorescence in situ hybridization (FISH)
- Differentiates between low-grade dysplasia and high-grade dysplasia
- Shown to increase the detection of low-grade dysplasia by 43% compared to histopathology only²
- Aids in the prediction of response to photodynamic therapy³
- CBEST™ is performed on easy-to-collect esophageal brushing samples*

Brush Cytology Advantages



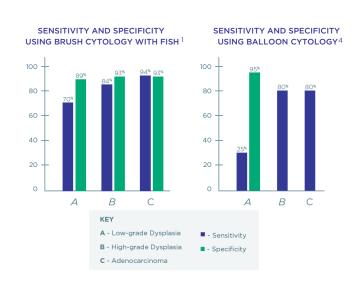
More comprehensive sampling of the esophagus compared to biopsy or cytology balloon¹



Less procedure time compared to four quadrant biopsy technique¹



Less invasive for patient



Comprehensive Barrett's Esophagus Test (CBEST™)

Test Details

The CBEST[™] panel of biomarkers have known associations with dysplastic changes of Barrett's esophagus. This multi-omics test provides an indication of progress to support treatment options.^{3,5-9}

- Cytology Imparts a characteristic range of coloration to exfoliative cells, allowing critical examination of nuclei and cytoplasmic components.
- AMACR (P504S) The concentration and activity of this protein has recently been identified in detecting dysplasia in ulcerative colitis, Crohn's disease, and Barrett's esophagus.
- p53† The study of p53 expression by IHC is of interest in Barrett's esophagus patients with a diagnosis of indefinite for dysplasia or low-grade dysplasia.
- **Ki-67**† IHC staining for MIB-1, the Ki-67 proliferation antigen, appears gradually and is an important marker for cell proliferation.
- ABPH 2.5 Stains the acidic mucin present in goblet cells.
- Feulgen Stain Used for the quantification of chromosomal material or DNA with sufficient resolution to detect the gain or loss of a single large chromosome.
- FISH Four-probe panel to detect gains and losses of MYC (8q24), p16 (CDKN2A at 9p21), HER2 (ERBB2 at 17q12), and ZNF217 (20q13) associated with higher-risk disease.

Also Available

Traditional FISH and CBEST™ for tissue specimens.† Please inquire for addtional information.

Contact Us to Get Started Today

pathnostics.com/contact 800.493.4490

INNOVATION, EXPERIENCE, AND SOLUTIONS YOU CAN TRUST

- Industry leading diagnostic solutions
- Expert board-certified pathology staff
- Partnership opportunities (TC/PC)

References

- 1. Brankley SM, Wang KK, Harwood AR, Miller DV, Legator MS, Lutzke LS, Kipp BR, Morrison LE, Halling KC. The development of a fluorescence in situ hybridization assay for the detection of dysplasia and adenocarcinoma in Barrett's esophagus. J Mol Diagn. 2006 May;8(2):260-7. doi: 10.2353/jmoldx.2006.050118.
- 2. Dong Y, Zhai J, Prunean J, Megna J, Prasad A, Moussa S, Baunoch D, Prasad A. Brushing cytology with adjunctive FISH and biomarker analyses is highly sensitive and specific in the early detection of low-grade dysplasia in Barrett's Esophagus. Poster presented at: USCAP.
- 3. Prasad GA, Wang KK, Halling KC, Buttar NS, Wongkeesong LM, Zinsmesiter AR, Brankley SM, Westra WM, Lutzke LS, Borkenhagen LS, Dunagan K.: Correlation of histology with biomarker status after photodynamic therapy in Barrett esophagus. Cancer. 2008 Aug 1;113(3):470-6.
- 4. Falk GW, Chittajallu R, Goldblum JR, Biscotti CV, Geisinger KR, Petras RE, Birgisson S, Rice TW, Richter JE: Surveillance of patients with Barrett's esophagus for dysplasia and cancer with balloon cytology. Gastroenterology. 1997;112:1787–1797.
- 5.Shi XY, Bhagwandeen B, Leong A. p16, Cyclin D1, Ki-67, and AMACR as Markers for Dysplasia in Barrett Esophagus. Applied Immunohistochemistry & Molecular Morphology. 2008 Oct;16(5):447-452. doi: 10.1097/PAI.0b013e318168598b.
- 6. Maslyonkina KS, Konyukova AK, Alexeeva DY, Sinelnikov MY, Mikhaleva LM. Barrett's esophagus: The pathomorphological and molecular genetic keystones of neoplastic progression. Cancer Med-us. 2022;11(2):447-478. doi: 10.1002/cam4.4447.
- 7. Lee C, Hayat U, Song K, Gravely AA, Mesa, H, Peltola J, Iwamoto C, Manivel C, Bilal M, Shaheen N, Shaukat A, Hanson B. A Consensus Diagnosis Utilizing Surface KI-67 Expression as an Ancillary Marker in Low-Grade Dysplasia Helps Identify Patients at High Risk of Progression to High-Grade Dysplasia and Esophaegal Adenocarcinoma. Diseases of the Esophagus. 2022;36(3). doi: 10.1093/dote/doac065.
- 8. Scott JE, Dorling, J. Differential staining of acid glycosaminoglycans (mucopolysaccharides) by Alcian blue in salt solutions. Histochemie. 1965;5:221-233. doi: 10.1007/BF00306130. 9. James PD, Atkinson M. Value of DNA image cytometry in the prediction of malignant change in Barrett's oesophagus. Gut. 1989;30:899-905. doi: 10.1136/gut.30.7.899.

†Indicates biomarkers included in CBEST™ for tissue specimens

