

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide**  
**Mouse Monoclonal Antibody [Clone Cocktail ]**  
**Catalog # AH11055**

**Specification**

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Product Information**

Application	IHC, IF, FC
Primary Accession	<a href="#">P06731</a>
Other Accession	<a href="#">1048</a> , <a href="#">634</a> , <a href="#">709196</a>
Reactivity	Human
Host	Mouse
Clonality	Monoclonal
Isotype	Mouse/IgG s
Calculated MW	80-200kDa KDa

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Additional Information**

**Gene ID** 1048

**Other Names**

Carcinoembryonic antigen-related cell adhesion molecule 5, Carcinoembryonic antigen, CEA, Meconium antigen 100, CD66e, CEACAM5, CEA

**Application Note**

IHC~~1:100~500  
IF~~1:50~200  
FC~~1:10~50

**Storage**

Store at 2 to 8°C. Antibody is stable for 24 months.

**Precautions**

Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide is for research use only and not for use in diagnostic or therapeutic procedures.

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Protein Information**

**Name** CEACAM5 ([HGNC:1817](#))

**Function**

Cell surface glycoprotein that plays a role in cell adhesion, intracellular signaling and tumor progression (PubMed: [10864933](http://www.uniprot.org/citations/10864933), PubMed: [10910050](http://www.uniprot.org/citations/10910050), PubMed: [2803308](http://www.uniprot.org/citations/2803308)). Mediates homophilic and heterophilic cell adhesion with other carcinoembryonic antigen-related cell adhesion molecules, such as CEACAM6 (PubMed: [2803308](#))

href="http://www.uniprot.org/citations/2803308" target="\_blank">2803308</a>). Plays a role as an oncogene by promoting tumor progression; induces resistance to anoikis of colorectal carcinoma cells (PubMed:<a href="http://www.uniprot.org/citations/10910050" target="\_blank">10910050</a>).

**Cellular Location**

Cell membrane; Lipid-anchor, GPI-anchor. Apical cell membrane. Cell surface Note=Localized to the apical glycocalyx surface

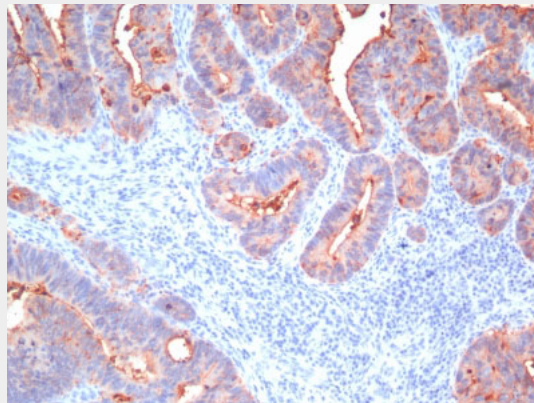
**Tissue Location**

Expressed in columnar epithelial and goblet cells of the colon (at protein level) (PubMed:10436421). Found in adenocarcinomas of endodermally derived digestive system epithelium and fetal colon.

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Images**

Formalin-fixed, paraffin-embedded human Colon Carcinoma stained with CEA, pan Monoclonal Antibody (C66/195+C66/261+ C66/1009+ C66/1030)

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - Background**

This antibody recognizes proteins of 80-200kDa, identified as different members of CEA family. CEA is synthesized during development in the fetal gut and is re-expressed in increased amounts in intestinal carcinomas and several other tumors. This MAb does not react with nonspecific cross-reacting antigen (NCA) and with human polymorphonuclear leucocytes. It shows no reaction with a variety of normal tissues and is suitable for staining of formalin/paraffin tissues. CEA is not found in benign glands, stroma, or malignant prostatic cells. Antibody to CEA is useful in detecting early foci of gastric carcinoma and in distinguishing pulmonary adenocarcinomas (60-70% are

CEA+) from pleural mesotheliomas (rarely or weakly CEA+). Anti-CEA positivity is seen in adenocarcinomas from the lung, colon, stomach, esophagus, pancreas, gallbladder, urachus, salivary gland, ovary, and endocervix.Ã

**Carcinoembryonic Antigen, pan (CEA) / CD66 Antibody - With BSA and Azide - References**

Muraro R, et. al. Cancer Research, 1985, 45:5769-80. | Siler K, et. al. Biotechnology Therapeutics, 1993, 4(3-4):163-81. | Robbins PF, et. al. International Journal of Cancer, 1993, 53(6):892-7. | Shi ZR, et. al. Journal of Histochemistry and Cytochemistry, 1994, 42(9):1215-9