

## Product Data Sheet

### anti-human CEACAM5,6 monoclonal antibody

#### Product information

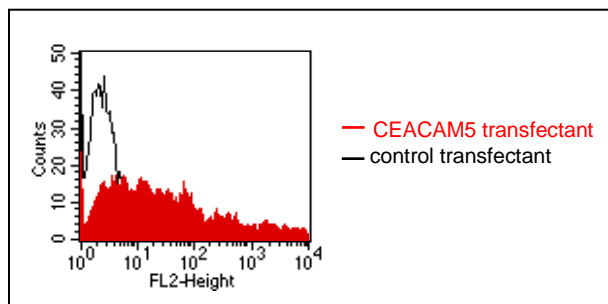
<b>Catalog Number:</b>	GM-0506
<b>Clone:</b>	MUS
<b>Description:</b>	purified monoclonal mouse antibody
<b>Specificity:</b>	anti-human CEACAM5,6 (CEA;NCA,CD66c)
<b>Isotype:</b>	IgG1
<b>Purification:</b>	Protein G
<b>Storage:</b>	short term: 2°C - 8°C; long term: -20°C (avoid repeated freezing and thawing)
<b>Buffer :</b>	phosphate buffered saline, pH 7.2
<b>Immunogen:</b>	immunisation with extracted protein of CEACAM5
<b>Selection:</b>	based on recognition of the complete <b>native protein</b> expressed on transfected mammalian cells

#### Working dilutions

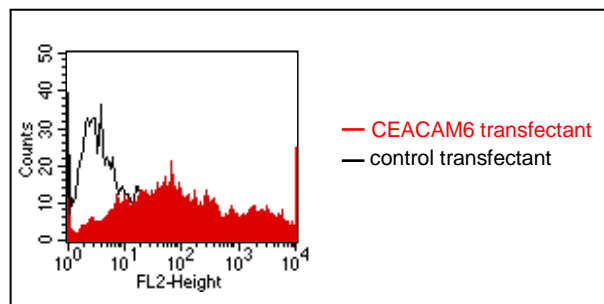
<b>Flow cytometry:</b>	1.2 µg/10 <sup>6</sup> cells
<b>ELISA:</b>	1:200 - 1:400
<b>CELISA:</b>	1:200
<b>Western blot:</b>	4µg/ml
<b>Immunohistology:</b>	1-2 µg/10 <sup>6</sup> cells (on cryosections)

For each application a titration should be performed to determine the optimal concentration.

#### Specificity testing by flow cytometry



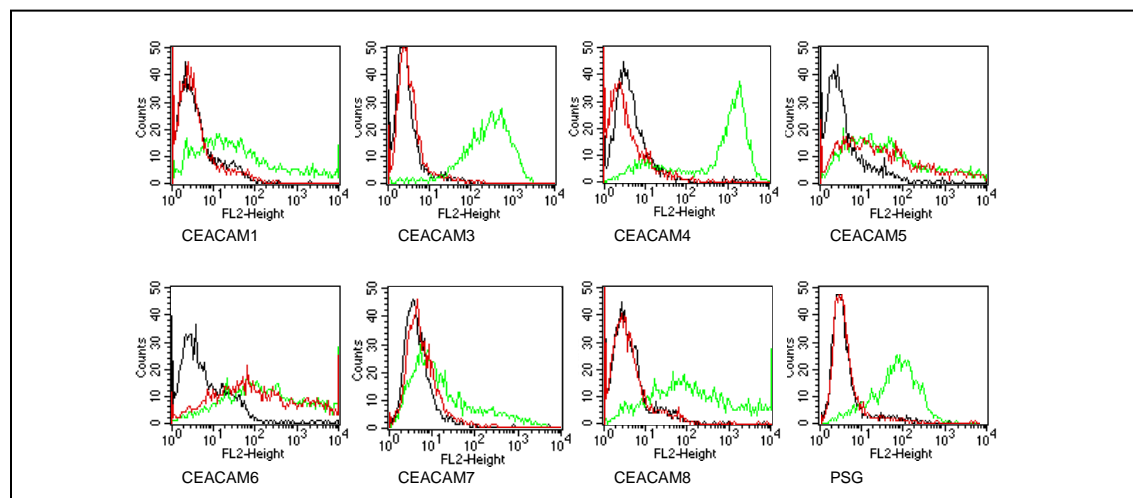
**Fig.1:** FACS analysis of BOSC23 cells using MUS Cat.# GM-0506. BOSC23 cells were transiently transfected with an expression vector encoding either CEACAM5 (red curve) or an irrelevant protein (control transfectant). Binding of MUS was detected with a PE-conjugated secondary antibody. A positive signal was obtained only with CEACAM5 transfected cells.



**Fig.2:** FACS analysis of BOSC23 cells using MUS Cat.# GM-0506. BOSC23 cells were transiently transfected with an expression vector encoding either CEACAM6 (red curve) or an irrelevant protein (control transfectant). Binding of MUS was detected with a PE conjugated secondary antibody. A positive signal was obtained only with CEACAM6 transfected cells.

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## Antibody cross-reactivity with members of the CEA family



**Fig. 3: Specificity testing of MUS.** BOSC cells were transiently transfected with expression vectors containing either the cDNA of CEACAM1, 3, 5, 6, 7, 8 or a recombinant transmembrane-anchored PSG1 fusion protein. Recognition of CEACAM4 was tested on CHO cells stably transfected with a CEACAM4 expression vector. Expression of the constructs was confirmed with monoclonal antibodies known to recognise the corresponding proteins (CEACAM1, 3, 4, 5 and 6: D14HD11; CEACAM7: CAC2; CEACAM8: 80H3; PSG: BAP1; green curves). An irrelevant monoclonal antibody served as a negative control (black curves). For specificity testing, protein G purified MUS was tested on all CEACAM transfectants. A positive signal was obtained with CEACAM5 and CEACAM6 expressing cells (red curves).

### Background

CEA-related cell adhesion molecules (CEACAM) belong to the carcinoembryonic antigen (CEA) family (1). The CEA family proteins belong to the immunoglobulin (Ig) superfamily and are composed of one Ig variable-like (IgV) and a varying number (0-6) of Ig constant-like (IgC) domains. CEACAM molecules are membrane-bound either via a transmembrane domain or a glycosyl phosphatidyl inositol (GPI) anchor. CEACAM molecules are differentially expressed in epithelial cells or in leucocytes. Over-expression of CEA/CEACAM5 in tumors of epithelial origin is the basis of its wide-spread use as a tumor marker (2). CEACAM6 expression is strongly up-regulated already during early stages of adenocarcinoma formation (3). The function of CEA family members varies widely: they function as cell adhesion molecules, tumor suppressors, regulators of lymphocyte and dendritic cell activation, receptors of *Neisseria* species and other bacteria (1).

### References

1. **Zimmermann W (2002).** Carcinoembryonic antigen. In *Wiley Encyclopedia of Molecular Medicine* (T. Creighton, ed.), John Wiley & Sons Inc., New York, USA, pp. 459-462.
2. **Hammarström S (1999).** The carcinoembryonic antigen (CEA) family: structures, suggested functions and expression in normal and malignant tissues. *Semin Cancer Biol.* 9, 67-81.
3. **Schölzel S, Zimmermann W, Schwarzkopf G, Grunert F, Rogaczewski B, Thompson J (2000).** Carcinoembryonic antigen family members CEACAM6 and CEACAM7 are differentially expressed in normal tissues and oppositely deregulated in hyperplastic colorectal polyps and early adenomas. *Am J Pathol* 156, 595-605.
4. **Grunert F, AbuHarfeil N, Schwarz K and von Kleist S (1985).** Two CEA and three NCA species, although distinguishable by monoclonal antibodies, have nearly identical peptide patterns. *Int J Cancer* 36, 357-362.

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