

# PRODUCT DATA SHEET

## ANTI-HUMAN CEACAM1, 5, 6, 8 MONOCLONAL ANTIBODY

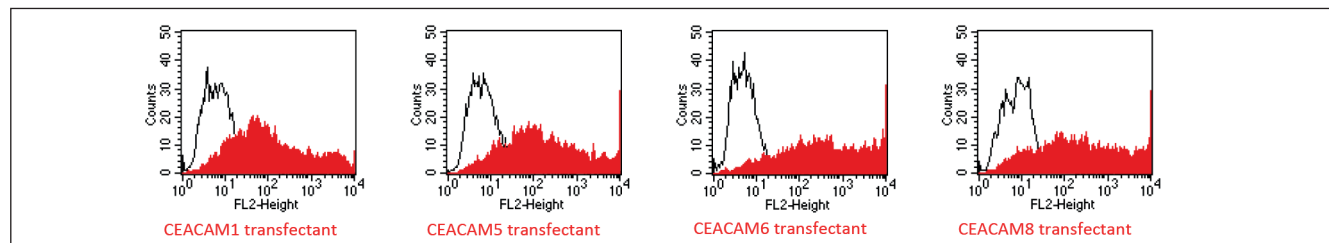
### PRODUCT INFORMATION

<b>Catalog Number:</b>	GM-0504	<b>Clone:</b>	TET2
<b>Description:</b>	purified monoclonal mouse antibody	<b>Specificity:</b>	anti-human CEACAM1, 5, 6, 8 (CD66a,b,c,e)
<b>Isotype:</b>	IgG2b/kappa	<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>	immunization with extracted protein of CEACAM5	<b>Selection:</b>	based on recognition of the complete native protein expressed on transfected mammalian cells

### WORKING DILUTIONS

<b>Flow cytometry:</b>	1 µg/10 <sup>6</sup> cells	<b>CELISA:</b>	1:200
<b>ELISA:</b>	1:200 – 1:400	<b>Immunohistology:</b>	1-2 µg/10 <sup>6</sup> cells (on cryosections)
<b>Western Blot:</b>	4µg/ml	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 TET2 Cat.# GM-0504. BOSC23 cells were transiently transfected with an expression vector encoding either CEACAM1,5,6,8 (red curves) or an irrelevant protein (control transfectant). Binding of TET2 was detected with a PE-conjugated secondary antibody. A positive signal was obtained only with CEACAM1, CEACAM5, CEACAM6 and CEACAM8 ex-pressing cells.

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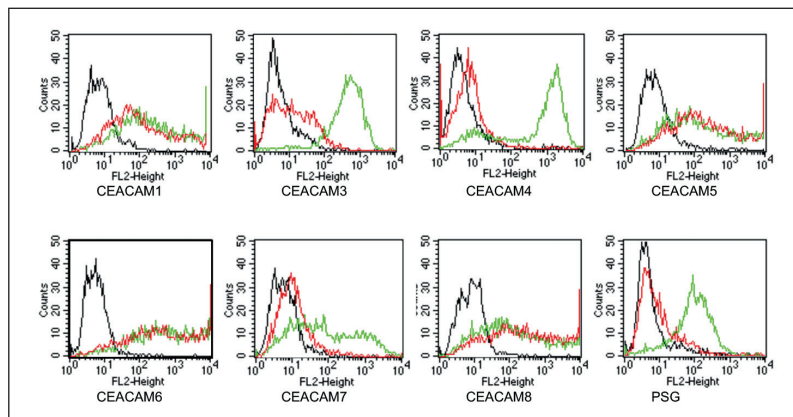
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## ANTIBODY CROSS-REACTIVITY WITH MEMBERS OF THE CEA FAMILY



**Fig. 2:** Specificity testing of TET2. 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; CEA-CAM7: CAC2; CEACAM8: 80H3; PSG1: BAP1; green curves). An irrelevant monoclonal antibody served as a negative control (black curves). For specificity testing, protein G purified TET2 was tested on all CEACAM transfectants. A positive signal was obtained with CEACAM1, CEACAM5, CEACAM6 and CEACAM8 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 immuno-globulin (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). The function of CEACAM 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. **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.
4. **Grunert F, Stocks SC, Nagel G, Zimmermann W, Thompson JA, Jantschke P and Kromer B (1996).** CD66 family Workshop: Binding of myeloid blind panel antibodies and CD66 Subsection antibodies to HeLa transfectants expressing individual CD66 molecules. In *Leukocyte Typing VI: White cell Differentiation Antigens* (T. Kishimoto et al., eds.), Garland Publishing Inc., New York and London, pp. 1012-1025.

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