

## Anti-Human TIE-2/tek Clone 16, Monoclonal Antibody

**Catalog No.** CMT204 **Quantity:** 100 μg

Clone: 16

**Synonyms:** Angiopoietin 1 receptor precursor antibody, CD202B antibody, CD202b antigen antibody,

P140 TEK antibody, TIE2 antibody, TIE-2 antibody, Tunica interna endothelial cell kinase

antibody, Tyrosine-protein kinase receptor TEK antibody, Tyrosine-protein kinase

receptor TIE-2 antibody, VMCM antibody, VMCM1 antibody

**Description:** Monoclonals were produced with the help of BALB/c mice using recombinant human

soluble extracellular TIE-2 as the immunizing antigen. Mouse IgG<sub>1</sub> antibody (#tek16)

from hybridomas was purified from cell culture supernatant by Protein G

chromatography.

Host Species: Mouse

Antigen: Recombinant human soluble TIE-2 protein

**Purification:** Protein G chromatography.

Stabilizer: none

**Buffer:** PBS pH 7.4 w/o preservative.

Formulation: Lyophilized.

**Reconstitution:** When reconstituted in sterile water to a concentration of 1.0 mg/ml the antibody is stable

for at least six weeks at 2-4°C.

**Stability:** The lyophilized antibody, though stable at room temperature, is best stored desiccated

below 0°C.

Reconstituted anti-TIE-2/tek is stable at 4°C for >one month or can be stored in working

aliquots at 20°C for more than six months.

Specificity: The monoclonal antibody will detect native human TIE-2/tek in ELISA experiments and

on the surface of different human cell types. The antibody can be used for ELISA

experiments, Western blotting, FACS and cell sorting.

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**Applications: ELISA**: Use at 1-15 μg/ml.

Western blotting: Use at 1-2 µg/ml

FACS analysis and cell sorting: Use at 2-5  $\mu g/ml$  together with the appropriate

secondary reagents.

Optimal dilutions should be determined by each laboratory for each application.

E-mail: info@cellsciences.com

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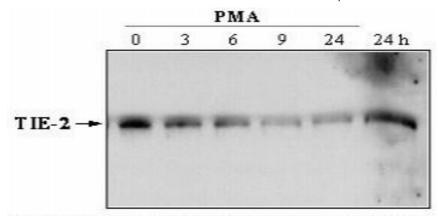
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**References:** Search <u>PubMed</u> (MEDLINE) for references to this product.

## Please note: always centrifuge vials before opening.

Effects of PMA treatment on TIE-2 mRNA and protein. HUVECs (passage 1) were stimulated for the indicated periods of time with PMA at 25 ng/ml or left untreated. Western blot analysis for the presence of TIE-2 protein by immunoprecipitation using antibodies directed against the extracellular domain of human TIE-2)



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