

TEV-cut site

Cat.No. 265 003; Polyclonal rabbit antibody, 50 µg specific antibody (lyophilized)

Data Sheet

Reconstitution/ Storage	50 µg specific antibody, lyophilized. Affinity purified with the immunogen. Rabbit serum albumin was added for stabilization. For reconstitution add 50 µl H ₂ O to get a 1mg/ml solution in PBS. Then aliquot and store at -20°C until use.
Applications	WB: 1 : 1000 (AP staining) IP: not tested yet ICC: 1 : 500 IHC: not tested yet IHC-P/FFPE: not tested yet
Immunogen	Synthetic peptide corresponding to AA 1 to 6 from TEV-cut site
Specificity	TEV-cut protease site is recognized with very strong preference.

TO BE USED IN VITRO / FOR RESEARCH ONLY
NOT TOXIC, NOT HAZARDOUS, NOT INFECTIOUS, NOT CONTAGIOUS

TEV is a highly specific cysteine protease derived from Tobacco Etch Virus (TEV). The optimum recognition site for this enzyme is the sequence (ENLYFQ(G/S)) and cleavage occurs between the Q and G/S residues leaving a free carboxy-terminus.

Selected General References

Tobacco etch virus protease retains its activity in various buffers and in the presence of diverse additives.

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Exploring the activity of tobacco etch virus protease in detergent solutions.

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Structural basis for the substrate specificity of tobacco etch virus protease.

Phan J, Zdanov A, Evdokimov AG, Tropea JE, Peters HK, Kapust RB, Li M, Wlodawer A, Waugh DS
The Journal of biological chemistry (2002) 277(52): 50564-72.

The P1' specificity of tobacco etch virus protease.

Kapust RB, Tózsér J, Copeland TD, Waugh DS

Biochemical and biophysical research communications (2002) 294(5): 949-55.

Tobacco etch virus protease: mechanism of autolysis and rational design of stable mutants with wild-type catalytic proficiency.

Kapust RB, Tózsér J, Fox JD, Anderson DE, Cherry S, Copeland TD, Waugh DS
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