

Recombinant Tobacco Etch Virus Protease, His Tag

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| Catalog No. | CSI20139A CSI20139B CSI20139C | Quantity: | 300 IU 1,000 IU 10,000 IU |
| Alternate Names: | P1 protease, TEV protease | | |
| Description: | TEV protease encoded by the tobacco etch virus is a catalytic domain of the Nuclear Inclusion a (NIa) protein. It consists of 241 a.a. amino acids with the molecular weight of 27kDa. TEV recognizes the amino acid sequence of the general form E-X-X-Y-X-Q (or S)/X', and cleaves between Q (or S)/X'. In this form X and X' stand for any of the amino acid residues, except that X' cannot be P. The optimal cleavage site is ENLYFQ/G. As having the absolute specificity and widely using conditions like broad pH range and ionic strength, the TEV protease became more versatile than EK, thrombin and other protease used in biochemical applications, especially recombinant protein production. The optimal temperature for cleavage is 30°C; however, the enzyme can be used at temperatures as low as 4°C. Following digestion, TEV Protease can be removed from the reaction via the His tag sequence by Ni ²⁺ -chelate affinity chromatography. | | |
| UniProt ID: | Q0GDU8 | | |
| Source: | <i>E. coli</i> | | |
| Concentration: | ~10 IU/μl, lot specific | | |
| Molecular Weight: | 27 kDa (241 aa) | | |
| Formulation: | Sterile-filtered 25 mM Tris-HCl, pH 8.0, 75 mM NaCl, 5 mM EDTA, 10 mM GSH, containing 50 % Glycerol | | |
| Purity: | > 90 % by SDS-PAGE analysis | | |
| Biological Activity: | One unit is defined as the amount of enzyme needed to cleave 3 μg of fusion protein in 1 hour to 85 % completion at 30°C in a buffer containing 50 mM Tris-HCl, pH 8.0, 0.5 mM EDTA, and 1 mM DTT. | | |
| Storage & Stability: | Store as supplied at -20°C to -80°C for up to 6 months. Once opened under sterile conditions, store at -20°C to -80°C for up to 3 months. Avoid repeated freeze-thaw cycles. | | |
| Application Notes: | A number of variables can be changed to optimize the cleavage of any specific protein. The amount of rTEV, the temperature of the incubation, and the time needed for cleavage may be examined. If the protein of interest is heat-labile, then 4 °C incubations are recommended. Reactions at 4 °C will require longer incubation times and/or more rTEV. | | |

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