

Human recombinant protein Ubiquitin-WT

Human Recombinant Ubiquitin-WT Catalog # PBV10630r

Specification

Human recombinant protein Ubiquitin-WT - Product info

Primary Accession P62979
Concentration 4

Calculated MW 8.564 kDa KDa

Human recombinant protein Ubiquitin-WT - Additional Info

Gene ID 6233
Gene Symbol RPS27A
Gene Source Human
Source E. Coli

Assay&Purity SDS-PAGE; ≥95%

Assay2&Purity2 N/A;
Recombinant Yes
Format

Format Liquid

Storage

-80°C; 4 mg/ml in PBS.

Human recombinant protein Ubiquitin-WT - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Human recombinant protein Ubiquitin-WT - Images

Human recombinant protein Ubiquitin-WT - Background

Ubiquitin is a small, evolutionarily conserved eukaryotic protein that can be attached to a wide variety of intracellular proteins including itself. Covalent attachment of ubiquitin to other proteins serves various functions, but its major role is to target cellular proteins for destruction. Cellular components that activate, transfer, remove, or simply recognize ubiquitin number in the hundreds, perhaps even in the thousands. In light of this complexity the ubiquitin pathway is ideal for a systems biology approach. Ubiquitin plays a very important role in regulated non-lysosomal ATP dependent protein degradation. The protein to be degraded is conjugated to ubiquitin and the





ubiquinated protein is then selectively degraded by a 26S complex, multi-catalytic cytosolic and nuclear protease termed proteasome. The Ub-proteasome proteolytic pathway, which is a complex process, is implicated to be of great importance for regulating numerous cellular processes.

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Pancre V., et al. Eur. J. Immunol. 21:2735-2741(1991). Adams S.M., et al. Br. J. Cancer 65:65-71(1992). Hillier L.W., et al. Nature 434:724-731(2005). Vladimirov S.N., et al. Eur. J. Biochem. 239:144-149(1996). Schlesinger D.H., et al. Nature 255:423-424(1975).