

DESCRIPTION

Species Reactivity	Human/Mouse/Rat
Specificity	Detects human, mouse, and rat GULP1/CED-6 in direct ELISAs and Western blots.
Source	Polyclonal Sheep IgG
Purification	Antigen Affinity-purified
Immunogen	<i>E. coli</i> -derived recombinant human GULP1/CED-6 Met1-Asn92 Accession # Q9UBP9
Conjugate	Alexa Fluor 405 Excitation Wavelength: 405 nm Emission Wavelength: 421 nm
Formulation	Supplied 0.2mg/ml in 1X PBS with RDF1 and 0.09% Sodium Azide *Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. [General Protocols](#) are available in the Technical Information section on our website.

Western Blot Optimal dilution of this antibody should be experimentally determined.

PREPARATION AND STORAGE

Shipping	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Protect from light. Do not freeze. 12 months from date of receipt, 2 to 8 °C as supplied

BACKGROUND

GULP1 (also PTB domain-containing engulfment adaptor protein 1 and Cell death protein 6 homolog/CED6) is a 36-40 kDa, cytosolic member of the ced-6 family of proteins. It is expressed by multiple cell types, including macrophages, neurons and sinusoidal endothelial cells, and is believed to play a key role in phagocytosis, particularly involving apoptotic cells. Failure to remove apoptotic cells is suggested to contribute to chronic inflammatory conditions. GULP1 is involved with a MEGF10:GULF:stabilin-1:stabilin-2 internalization pathway. Possibly following endosome formation, it binds to transmembrane activated stabilins and/or LRP1 as a homodimer, and serves as an intermediate for signal transduction. GULP1 also binds to APP, facilitating its proteolysis and the generation of Aβ. Human GULP1 is 304 amino acids (aa) in length. It contains one PTB domain (aa 17-154) that interacts with transmembrane receptors, a Leu-zipper (aa 163-193) that mediates dimerization, and a Pro-rich region (aa 238-271) that is likely involved with signal transduction. There are three splice variants. One shows a deletion of aa 31-133, a second contains a 34 aa substitution for aa 134-304, and a third contains a 10 aa substitution for aa 282-304. Over aa 1-92, human GULP1 shares 99% aa sequence identity with mouse GULP1.

PRODUCT SPECIFIC NOTICES

This product is provided under an agreement between Life Technologies Corporation and R&D Systems, Inc, and the manufacture, use, sale or import of this product is subject to one or more US patents and corresponding non-US equivalents, owned by Life Technologies Corporation and its affiliates. The purchase of this product conveys to the buyer the non-transferable right to use the purchased amount of the product and components of the product only in research conducted by the buyer (whether the buyer is an academic or for-profit entity). The sale of this product is expressly conditioned on the buyer not using the product or its components (1) in manufacturing; (2) to provide a service, information, or data to an unaffiliated third party for payment; (3) for therapeutic, diagnostic or prophylactic purposes; (4) to resell, sell, or otherwise transfer this product or its components to any third party, or for any other commercial purpose. Life Technologies Corporation will not assert a claim against the buyer of the infringement of the above patents based on the manufacture, use or sale of a commercial product developed in research by the buyer in which this product or its components was employed, provided that neither this product nor any of its components was used in the manufacture of such product. For information on purchasing a license to this product for purposes other than research, contact Life Technologies Corporation, Cell Analysis Business Unit, Business Development, 29851 Willow Creek Road, Eugene, OR 97402, Tel: (541) 465-8300. Fax: (541) 335-0354.