

Product datasheet

Recombinant human BAFF-R protein ab157066

1 Image

Description

Product name	Recombinant human BAFF-R protein	
Biological activity	Inhibits rhsBAFF mediated splenocyte survival.	
Purity	> 95 % SDS-PAGE.	
Endotoxin level	< 0.100 Eu/μg	
Expression system	HEK 293 cells	
Accession	Q96RJ3	
Protein length	Protein fragment	
Animal free	No	
Nature	Recombinant	
Species	Human	
Sequence	RRGPRSLRGRDAPAPTPCVPAECFDLLVRHCVACGLLRT PRPKPAGASSP APRTALQPQESVGAGAGEAA	
Predicted molecular weight	40 kDa	
Amino acids	2 to 71	
Additional sequence information	The extracellular domain of human BAFF-R (aa 2-71) is fused at the C-terminus to the Fc portion of human IgG1.	

Specifications

Our [Abpromise guarantee](#) covers the use of **ab157066** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications	Functional Studies SDS-PAGE
Form	Lyophilized
Additional notes	Binds human and mouse BAFF. Detection of membrane-bound human and mouse BAFF in combination with PAb to human IgG1. Inhibits rhsBAFF mediated splenocyte survival.

Preparation and Storage

Stability and Storage

Shipped at 4°C. After reconstitution store at -20°C. Avoid freeze / thaw cycles.

Constituent: 99% PBS

This product is an active protein and may elicit a biological response in vivo, handle with caution.

Reconstitution

Reconstitute with 50µl sterile water. Further dilutions should be made with medium containing 5% fetal calf serum or a carrier protein. 1mg/ml (after reconstitution).

General Info

Function

B-cell receptor specific for TNFSF13B/TALL1/BAFF/BLyS. Promotes the survival of mature B-cells and the B-cell response.

Tissue specificity

Highly expressed in spleen and lymph node, and in resting B-cells. Detected at lower levels in activated B-cells, resting CD4+ T-cells, in thymus and peripheral blood leukocytes.

Involvement in disease

Defects in TNFRSF13C are the cause of immunodeficiency common variable type 4 (CVID4) [MIM:613494]; also called antibody deficiency due to BAFFR defect. CVID4 is a primary immunodeficiency characterized by antibody deficiency, hypogammaglobulinemia, recurrent bacterial infections and an inability to mount an antibody response to antigen. The defect results from a failure of B-cell differentiation and impaired secretion of immunoglobulins; the numbers of circulating B cells is usually in the normal range, but can be low.

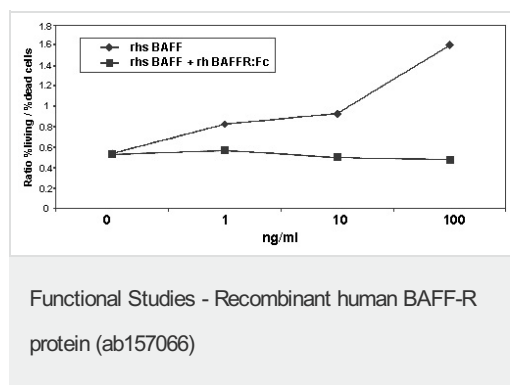
Sequence similarities

Contains 1 TNFR-Cys repeat.

Cellular localization

Membrane.

Images



ab157066 used to show rhBAFF-R:Fc inhibits rhSBAFF-mediated activation of freshly isolated mouse splenocytes.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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