

# **PRODUCT DATA SHEET**

## Product: BAFF, (human recombinant soluble)

### Cat. No.: BC-321 (10 µg)

#### Synonyms:

BlyS; TALL-1; THANK; zTNF4; TNFSF 13B/20; CD257

#### Source/Host:

Produced in *E. coli.* The extracellular domain of human BAFF (aa 83-285) is fused at the N-terminus to a linker peptide (6 aa) and a  $FLAG^{\textcircled{B}}$  tag.

#### Molecular Weight:

~28 kDa as determined by SDS-PAGE

#### Format:

Lyophilized. Contains PBS. Reconstitute with 100  $\mu$ L of sterile water for 0.1 mg/mL solution. Further dilutions should be made with medium containing 5% fetal calf serum.

#### **Purity:**

>95% as determined by SDS-PAGE.

Endotoxin: <0.1 EU/ $\mu$ g of purified protein (LAL test).

#### **Biological Activity and Application:**

Mediates splenocyte survival. Detection of receptors of BAFF on target cells. Binds to human and mouse(weak) BCMA, TACI and BAFF-R.

#### Storage:

Store at -20°C. After reconstitution, prepare aliquots and store at -20°C.

#### Limitations:

For *in vitro* research use only. Not for use in diagnostics or in humans.

#### Warranty:

No warranties, expressed or implied, are made regarding the use of this product. **KAMIYA BIOMEDICAL COMPANY** is not liable for any damage, personal injury, or economic loss caused by this product.

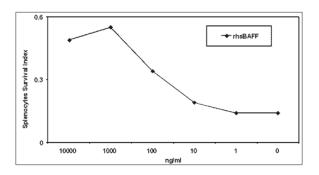
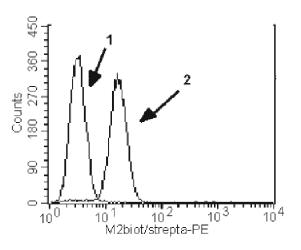


Figure: BAFF (BC-321) mediates survival of freshly isolated splenocytes.

Method: On day 0 splenocytes were isolated from a freshly collected C57Bl6 spleen. An aliquot of the splenocytes was analyzed on FACS and gated on the SSC-FSC panel. FACS settings were saved. The rest of the cells was put in culture with media alone or with increasing concentrations of BAFF (BC-321) as indicated. After three days in culture, cells were harvested and analyzed on FACS with the saved setting. Splenocytes Survival Index (ratio % living/% dead cells) was calculated and plotted



rhsBAFF can be used as a tool to detect its receptors on target cells.

Figure: Flow cytometric profile of BJAB cells incubated with (2) or without (1) rhsBAFF.

Method: BJAB cells (5 x  $10^5$ ) were incubated on ice for 30 min in 50 µL FACS buffer (PBS, 5% fetal calf serum, 0.02% sodium azide) containing 2 µg/mL of BAFF (BC-321). After washing in FACS buffer, biotinylated antibody to Flag was added at 2.5 µg/mL for 30 min. After washing in FACS buffer, streptavidin-PE (Jackson Lab) at 2 µg/mL was added, incubated for 30 min, washed, and finally cells were analyzed by flow cytometry.

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