



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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## Product Information

**Description:**

This anti-BCMA antibody is a purified, biotinylated, recombinant human monoclonal antibody which recognizes the human BCMA protein. BCMA is also known as tumor necrosis factor receptor superfamily member 17 (TNFRSF17) and is expressed on the surface of plasma cells. BCMA is important for plasma cell differentiation and B cell proliferation. BCMA is a validated therapeutic target in multiple myeloma. The first anti-BCMA CAR-T cell therapy (idecabt agene vicleucel) was approved by the FDA in 2021.

This antibody has been tested for specific binding to purified human BCMA protein (BPS Bioscience #79467) in an ELISA binding assay with colorimetric detection.

**Concentration:**

1.0 mg/ml

**Species:**

Human

**Isotype:**

IgG1

**Formulated In:**

8 mM phosphate, pH 7.4, 110 mM NaCl, 2.2 mM KCl, 20% glycerol

**Expression System:**

HEK293

**Clonality:**

Monoclonal

**Purification:**

Protein A affinity chromatography from HEK293 supernatants

**Cross Reactivity:**

This antibody recognizes human BCMA. It has not been tested with other species.

**Format:**

Aqueous buffer solution

**Stability:**

At least 12 months at -80°C. Avoid freeze/thaw cycles.

**Storage:**

-80°C

**MW:**

~146 kDa (Anti-BCMA HC: 50 kDa; Anti-BCMA LC: 23 kDa)

**Purity:**

≥90%

**Assay Conditions:**

Experimental design and assay protocol for measuring anti-BCMA specific binding to human BCMA in an ELISA assay:

1. Purified anti-BCMA (cat#101219) was thawed on ice and coated onto a clear 96-well plate overnight at 4°C (1 µg/ml in PBS, 50 µl per well).  
*\*\* "No Coat" controls and "Blank" wells were included by coating PBS only to determine background levels\*\**
2. The coated wells were washed three times with 150 µl of BPS Immuno Buffer 1.
3. Wells were blocked with 100 µl of Blocking Buffer 2 (BPS Bioscience, #79728) per well for 1 hour at room temperature with slow shaking.
4. Serial dilutions of purified human biotin-labeled BCMA protein (BPS Bioscience, #79467) were prepared in Blocking Buffer 2 (Titration from 0 nM to 100 nM, using 50 µl per well)
5. At the end of the blocking step, 50 µl of biotinylated-BCMA dilutions was added to appropriate wells and incubated for 1 hour at room temperature with slow shaking  
*\*\*Wells designated as "BLANK" were skipped\*\**
6. Wells were then washed as in step 2
7. An additional blocking step was performed with 100 µl of Blocking Buffer 2 at room temperature on the benchtop. The wells were then decanted and tapped to dry.

8. Streptavidin-HRP was prepared in 1x Blocking Buffer 2 and 50  $\mu$ l was added to each well immediately after the blocking buffer was decanted, and incubated for 1 hour at room temperature with slow shaking.
9. Wells were then washed as in step 2
10. 100  $\mu$ l of Colorimetric HRP Substrate (BPS Bioscience #79651) was added to all wells. Samples were allowed to develop color until positive control wells became blue (typically between 30 seconds to 5 minutes).
11. The reaction was quickly quenched with an equal volume (100  $\mu$ l) of 1N HCl.
12. Absorbance was read at 450 nm. The "blank" value was subtracted from all other measurements.

**Applications:**

This product is for research use only. It is not suitable for human, diagnostic, or therapeutic use.

## Quality Control Data

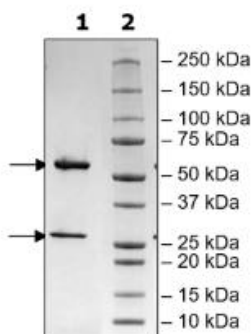
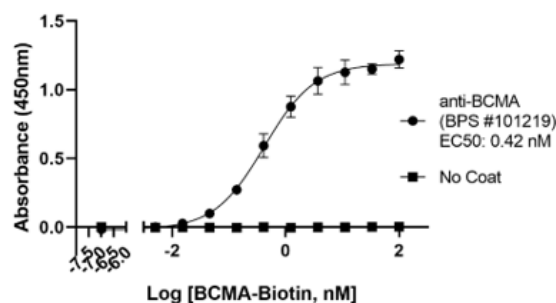
## 4-20% SDS-Page Coomassie Staining

**4-20% SDS-PAGE  
Coomassie staining**

**Lane 1:**  
5  $\mu$ g Anti-BCMA

**Lane 2:**  
Protein Marker

**MW:**  
LC: 23kDa  
HC: 50kDa  
**Purity:**  $\geq 90\%$

**anti-hBCMA vs. hBCMA-biotin Binding Assay**

This binding assay was performed following the assay conditions detailed above. Anti-BCMA was coated at 1  $\mu$ g/ml overnight. A "No Coat" control was included to determine the background levels and was coated with PBS only