

# Bovine Serum Albumin Polyclonal Antibody

## Product Details

Size	500 µL
Species Reactivity	Bovine
Published Species	Rabbit, Baboon, Non-human primate, Bovine, Human, Mouse
Host/Isotype	Rabbit / IgG
Class	Polyclonal
Type	Antibody
Conjugate	Unconjugated
Immunogen	Bovine Serum Albumin
Form	Liquid
Concentration	2 mg/mL
Purification	IgG fraction
Storage buffer	PBS, pH 7.2
Contains	5mM sodium azide
Storage conditions	4° C
RRID	AB_2534139

Applications	Tested Dilution	Publications
Western Blot (WB)	1:1,000	10 Publications
Immunohistochemistry (Paraffin) (IHC (P))	Assay-dependent	-
Immunocytochemistry (ICC/IF)	Assay-dependent	3 Publications
Flow Cytometry (Flow)	-	1 Publication
ELISA (ELISA)	Assay-dependent	1 Publication
Immunoprecipitation (IP)	Assay-dependent	1 Publication
Functional Assay (FN)	-	1 Publication
Miscellaneous PubMed (Misc)	-	2 Publications

## Product Specific Information

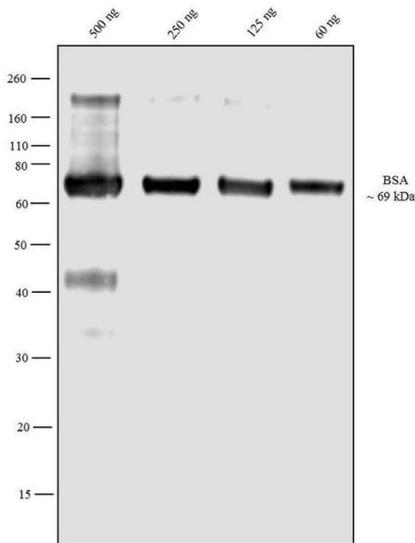
Anti-bovine serum albumin (anti-BSA) antibody is a highly purified rabbit IgG fraction that can be used for the detection of BSA in tracer studies or in protease assays. This antibody demonstrates high specificity for BSA and little or no crossreactivity to other serum proteins. Cat. No. A11133 can be used in enzyme and fluorescence immunoassays and blots, cell or tissue immunostaining, and as a precipitating antibody or as an immuno-adsorbant when attached to solid supports. For detection of this rabbit antibody, we offer a variety of second-step reagents labeled with biotin or with a wide range of fluorophores.

Because staining protocols vary with application, the appropriate dilution of antibody should be determined empirically. We suggest using serial dilutions from about 1:100 to 1:1,000 to determine the optimal conditions for your experiments.

## Product Images For Bovine Serum Albumin Polyclonal Antibody

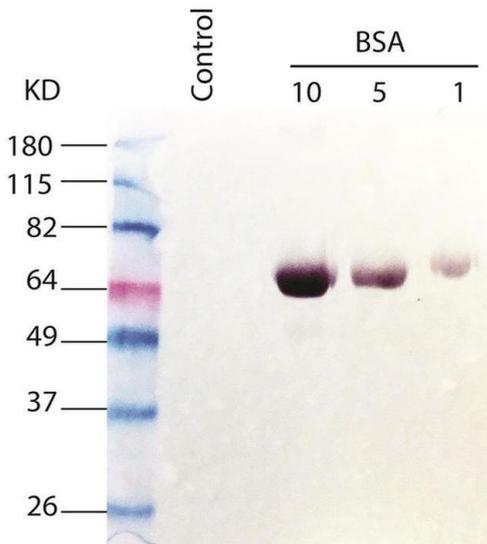
### Bovine Serum Albumin Antibody (A11133) in WB

Western blot analysis was performed by loading 500 ng (Lane 1), 250 ng (Lane 2), 125 ng (Lane 3) and 60 ng (Lane 4) of BSA. The blot was probed with Anti-Bovine Serum Albumin Rabbit Polyclonal Antibody (Product # A11133, 1:1000 dilution) and detected by chemiluminescence using Goat anti-Rabbit IgG (Heavy Chain) Superclonal™ Secondary Antibody, HRP conjugate (Product # A27036, 0.25 µg/mL, 1:4000 dilution). A 69 kDa band corresponding to Bovine Serum Albumin was observed. Known quantity of protein samples were electrophoresed using Novex® NuPAGE® 4-12 % Bis-Tris gel (Product # NP0322BOX), XCell SureLock™ Electrophoresis System (Product # EI0002) and Novex® Sharp Pre-Stained Protein Standard (Product # LC5800). Resolved proteins were then transferred onto a nitrocellulose membrane with iBlot® 2 Dry Blotting System (Product # IB21001). The membrane was probed with the relevant primary and secondary Antibody following blocking with 5 % skimmed milk. Chemiluminescent detection was performed using Pierce™ ECL Western Blotting Substrate (Product # 32106).



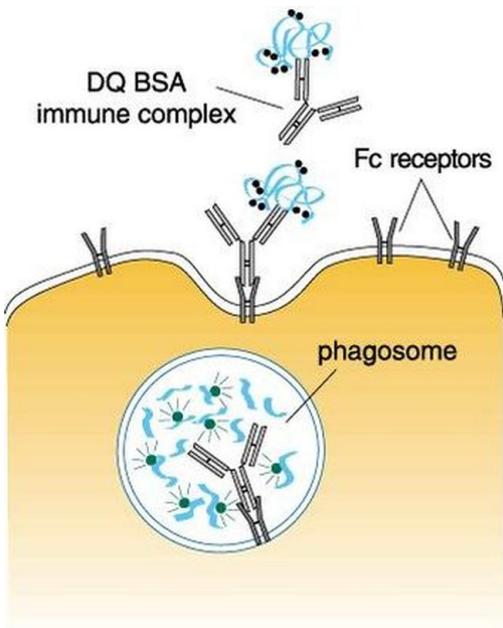
### Bovine Serum Albumin Antibody (A11133) in WB

Western blot analysis of BSA was performed by loading an unrelated protein (lane 1), 1 µg (lane 2), 5 µg (lane 3) and 10 µg of BSA (lane 4) in reducing sample buffer and protein ladder onto an SDS-PAGE gel. Proteins were transferred to nitrocellulose membrane. Membrane was blocked in 2% skim milk in PBS. BSA was detected at approximately 69 kDa using a BSA polyclonal antibody (Product # A11133) at a dilution of 1:1000 in 1% skim milk in PBST overnight at 4 degrees, followed by a 1:2000 dilution of HRP-conjugated goat anti-rabbit IgG antibody at room temperature for one hour. HRP color development substrate was used. Data courtesy of Antibody Data Exchange Program.



### Bovine Serum Albumin Antibody (A11133)

Immune complex of DQ™ BSA conjugate (Product # D12050, D12051) with an anti-bovine serum albumin (BSA) antibody (Product # A11133) for the fluorescent detection of the Fc receptor-mediated phagocytosis pathway. The DQ™ BSA is a derivative of BSA that is labeled to such a high degree with either the green-fluorescent BODIPY® FL or red-fluorescent BODIPY® TR-X dye that the fluorescence is self-quenched. Upon binding to an Fc receptor, the nonfluorescent immune complex is internalized and the protein is subsequently hydrolyzed to fluorescent peptides within the phagosome.



[View more figures on thermofisher.com](#)

## 19 References

### Western Blot (10)

Frontiers in molecular neuroscience

#### SUMOylation of the Kv4.2 Ternary Complex Increases Surface Expression and Current Amplitude by Reducing Internalization in HEK 293 Cells.

"A11133 was used in Western Blot to show that SUMOylation of Kv4.2 at K579 regulates TC internalization most likely by promoting channel recycling."

Authors: Welch MA, Jansen LR, Baro DJ

Year  
2022

Species  
Human

Dilution  
1:20000

Scientific reports

#### Nanostructure of bioactive glass affects bone cell attachment via protein restructuring upon adsorption.

"A11133 was used in Western Blot to provide new insight into the interaction of cells with the scaffolds' interfacial layer, which is vital for the continued development of engineered tissue scaffolds."

Authors: Thamma U, Kowal TJ, Falk MM, Jain H

Year  
2021

Species  
Mouse

Dilution  
1:5000

[View more WB references on thermofisher.com](#)

### Immunocytochemistry (3)

Journal of tissue engineering and regenerative medicine

#### Human dental pulp stem cells derived from cryopreserved dental pulp tissues of vital extracted teeth with disease demonstrate hepatic-like differentiation.

"Published figure using Bovine Serum Albumin polyclonal antibody (Product # A11133) in Immunofluorescence"

Authors: Chen YK, Huang AH, Chan AW, Lin LM

Year  
2016

The Journal of biological chemistry

#### CR1-mediated ATP release by human red blood cells promotes CR1 clustering and modulates the immune transfer process.

"Published figure using Bovine Serum Albumin polyclonal antibody (Product # A11133) in Immunofluorescence"

Authors: Melhorn MI, Brodsky AS, Estanislau J, Khoory JA, Illigens B, Hamachi I, Kurishita Y, Fraser AD, Nicholson-Weller A, Dolmatova E, Duffy HS, Ghiran IC

Year  
2013

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