

β-2-Microglobulin (2213): sc-80668

BACKGROUND

Major histocompatibility complex (MHC) class 1 molecules bind to antigens for presentation on the surface of cells. The proteasome is responsible for producing these antigens from the components of foreign pathogens. MHC class 1 molecules consist of an α heavy chain that contains three subdomains ($\alpha 1$, $\alpha 2$, $\alpha 3$), and a non-covalent associating light chain, known as β-2-Microglobulin. β-2-Microglobulin associates with the $\alpha 3$ subdomain of the α heavy chain and forms an immunoglobulin domain-like structure that mediates proper folding and expression of MHC class 1 molecules. The $\alpha 1$ and $\alpha 2$ domains of the α heavy chain form the peptide antigen-binding cleft. Mice that lack β-2-Microglobulin protein show a normal distribution of T cells, yet have no mature CD4-8+ T cells and are defective in CD4-8+ T cell-mediated cytotoxicity. Interferon- γ can stimulate production of β-2-Microglobulin transcripts. The human β-2-Microglobulin gene maps to chromosome 15q21.1 and encodes a 119 amino acid protein. Mutations in the β-2-Microglobulin gene can enhance the progression of malignant melanoma phenotypes.

REFERENCES

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CHROMOSOMAL LOCATION

Genetic locus: B2M (human) mapping to 15q21.1.

SOURCE

β-2-Microglobulin (2213) is a mouse monoclonal antibody raised against β-2-Microglobulin of human origin.

PRODUCT

Each vial contains 100 µg IgG₁ in 1.0 ml PBS with < 0.1% sodium azide, 1% glycerol, and 0.1% gelatin.

APPLICATIONS

β-2-Microglobulin (2213) is recommended for detection of β-2-Microglobulin of human origin by immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

Suitable for use as control antibody for β-2-Microglobulin siRNA (h): sc-29592, β-2-Microglobulin shRNA Plasmid (h): sc-29592-SH and β-2-Microglobulin shRNA (h) Lentiviral Particles: sc-29592-V.

Molecular Weight of β-2-Microglobulin: 12 kDa.

SELECT PRODUCT CITATIONS

- Tayman, C., et al. 2011. Mesenchymal stem cell therapy in necrotizing enterocolitis: a rat study. *Pediatr. Res.* 70: 489-494.
- Ryu, C.M., et al. 2018. Longitudinal intravital imaging of transplanted mesenchymal stem cells elucidates their functional integration and therapeutic potency in an animal model of interstitial cystitis/bladder pain syndrome. *Theranostics* 8: 5610-5624.
- Ryu, C.M., et al. 2019. N-acetylcysteine prevents bladder tissue fibrosis in a lipopolysaccharide-induced cystitis rat model. *Sci. Rep.* 9: 8134.
- Shin, J.H., et al. 2019. Synergistic effects of N-acetylcysteine and mesenchymal stem cell in a lipopolysaccharide-induced interstitial cystitis rat model. *Cells* 9 pii: E86.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.