

Technical Data Sheet

RecombiMAb anti-mouse FAP



Attention: Use of this product constitutes an agreement to Bio X Cell's Terms and Conditions which are included with this product in print and can also be found at <https://bioxccl.com/terms-and-conditions>.

Lot Specific Information

Lot Number: Lot Specific*
Volume: Lot Specific*
Concentration: Lot Specific* (generally 4 to 11 mg/ml) *
Total Protein: Lot Specific*

*This information will be noted on the certificate of analysis that ships with this product.

Product Information

Catalog Number: CP057
Clone: 73.3-CP057
Isotype: Mouse IgG2a, κ
Recommended Isotype Control(s): RecombiMAb mouse IgG2a isotype control, unknown specificity
Recommended Dilution Buffer: InVivoPure pH 7.0 Dilution Buffer
Immunogen: Immunization of FAP-null mice with FAP-expressing 3T3 cells
Reported Applications: Immunoprecipitation
Flow cytometry
Western blot
Chimeric Antigen Receptor construction
*Reported for the original 73.3 antibody.
For information on *in vivo* applications, please contact technicalservice@bioxccl.com
Formulation: PBS, pH 7.0
Contains no stabilizers or preservatives
Endotoxin: ≤ 0.5 EU/mg (≤ 0.0005 EU/ μ g)
Determined by LAL gel clotting assay
Purity: $\geq 95\%$
Determined by SDS-PAGE
Sterility: 0.2 μ m filtration
Production: Purified from HEK293 cell supernatant in an animal-free facility
Purification: Protein G
Aggregation: $< 5\%$
Determined by SEC
RRID:
Molecular Weight: 150 kDa

Murine Pathogen Test Results

Mouse Norovirus: Negative, Mouse Parvovirus: Negative, Mouse Minute Virus: Negative, Mouse Hepatitis Virus: Negative, Reovirus Screen: Negative, Lymphocytic Choriomeningitis virus: Negative, Lactate Dehydrogenase-Elevating Virus: Negative, Mouse Rotavirus: Negative, Theiler's Murine Encephalomyelitis: Negative, Ectromelia/Mousepox Virus: Negative, Hantavirus: Negative, Polyoma Virus: Negative, Mouse Adenovirus: Negative, Sendai Virus: Negative, Mycoplasma Pulmonis: Negative, Pneumonia Virus of Mice: Negative, Mouse Cytomegalovirus: Negative, K Virus: Negative

Description

The CP057 monoclonal antibody is a recombinant, chimeric version of the original 73.3 antibody. The variable domain sequences are identical but the constant region sequences have been switched from Mouse IgG1, κ to Mouse IgG2a, κ . The highly controlled sequence and lack of genetic drift in recombinant antibodies provide more reliable and reproducible results

over hybridoma derived antibodies. The 73.3 monoclonal antibody reacts with mouse fibroblast activation protein (FAP), a cell-surface serine protease that acts on various hormones and extracellular matrix components. FAP is expressed during embryonic development, in tissues of healing wounds, and in chronic inflammatory and fibrotic conditions. FAP protein is highly upregulated in cancer-associated fibroblasts (CAFs) in epithelial tumors. CAF overexpression of FAP promotes tumor development and metastasis by influencing extracellular matrix remodeling, intracellular signaling, angiogenesis, epithelial-to-mesenchymal transition, and immunosuppression. The scFv of the 73.3 antibody has been used to construct 73.3-FAP-CAR T cells specific for FAP. Adoptively transferred 73.3-FAP-CAR T cells have been shown to inhibit the growth of multiple syngeneic mouse tumor models.

Storage

Store at the stock concentration at 4°C . **Do not freeze.**

It is not uncommon for a floccule or precipitate to appear during storage. The floccule is typically buffer salts precipitating out of solution or a small bit of protein aggregation. For information on how to remove floccules or precipitates see our FAQ's at <https://bioxcell.com/faqs>.

Protocol Information

Since applications vary, each investigator should use the application references as a guide to help estimate the appropriate dose or concentration. The dose or concentration can be further optimized experimentally in a dose response or titration experiment.

Application References

For a complete list of references, visit https://bioxcell.com/cp057?bxcs=9k1b3a#tab_references or scan the QR code below.



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