

DESCRIPTION

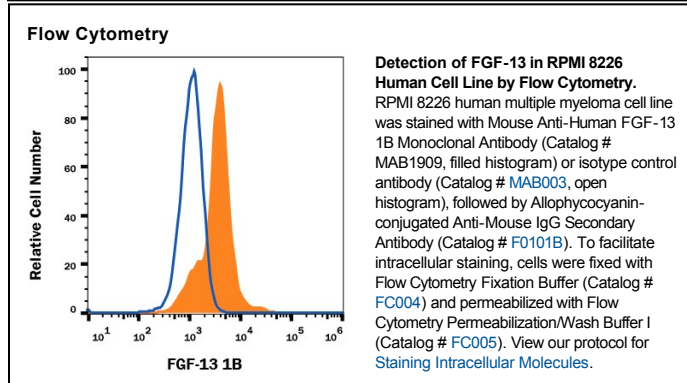
Species Reactivity	Human
Specificity	Detects human FGF-13 1B in direct ELISAs and Western blots. In direct ELISAs and Western blots, no cross-reactivity with recombinant human (rh) FGF-3, -4, -5, -6, -7, -9, -10, -11, -12, -16, -17, -19, -20, -21, rhFGF basic, recombinant mouse FGF-8b, or -8c is observed.
Source	Monoclonal Mouse IgG _{2A} Clone # 298923
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> -derived recombinant human FGF-13 isoform 1B Ala2-Thr192 Accession # NP_378668
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
Western Blot	1 µg/mL	Recombinant Human FGF-13 1B
Flow Cytometry	0.25 µg/10 ⁶ cells	See Below
CyTOF-ready	Ready to be labeled using established conjugation methods. No BSA or other carrier proteins that could interfere with conjugation.	

DATA



PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> ● 12 months from date of receipt, -20 to -70 °C as supplied. ● 1 month, 2 to 8 °C under sterile conditions after reconstitution. ● 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

FGF-13 (Fibroblast Growth Factor-13), also known as FGF homologous factor 2 (FHF-2), is a member of the FGF family. It is expressed in brain and skeletal muscle and has been implicated in nervous system development and function. The gene for FGF-13 has been localized in human chromosome X, in a region associated with Borjeson-Forssman-Lehmann syndrome, a syndromal X-linked mental retardation. Two alternately spliced isoforms that differ in their amino-terminal residues have been described.