

PERFORMANCE DATA SHEET

2216

Monoclonal anti-human GARP (LRRC32)/R-PE *

mAb name/Clone: ANC8C9

Isotype: Mouse IgG2ak

Immunogen: Recombinant human GARP

CATALOG#: 277-050

QUANTITY: 120 tests

VOLUME IN VIAL: 0.2 ml

WORKING DILUTION: 1:50 (or use 1.6µl of concentrated stock per 5 x 10⁵-cell test)

INFORMATION: Human GARP (Glycoprotein A Repetitions Predominant/ Leucine Rich Repeat Containing 32) is a roughly 80 kd transmembrane cell surface molecule expressed on a variety of tissue types, including a subset of activated T regulatory cells. GARP covalently binds latent TGF-beta 1 serving as a regulatory reservoir(1). Antibody ANC8C9 binds to recombinant GARP in EIA and to cell surface GARP expressed on a subset of FOXP3 positive cells.

References: 1) [R Wang, T A Springer, et al. \(2012\) Mol Biol Cell 23\(6\): 1129-1139.](#)

STORAGE CONDITIONS: Store at 2 - 5°C. Do not freeze! Protect from light.

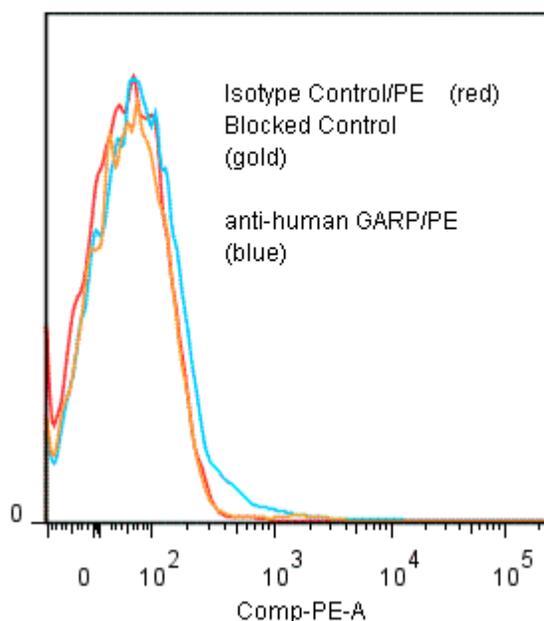
PRODUCT STABILITY: Product should retain activity for at least 12 months after shipping date when stored as recommended. Ship Date: _____

BUFFER: 50 mM Sodium Phosphate pH 7.5, 500 mM Potassium Chloride, 150mM NaCl, 15% Glycerol, 0.2% BSA, 0.04% NaN₃ (as a preservative).

PRODUCTION: Protein A purified antibody from tissue culture supernatant was conjugated to R-Phycoerythrin through a sulfo-ester linkage. Unconjugated antibody was removed using size exclusion chromatography. Antibody conjugate is at 0.5 mg/ml with an A₅₆₅/A₂₈₀ ratio of 2.63.

PERFORMANCE: (FACS analysis) Ficoll prepared human PBMC were stimulated in culture overnight with immobilized anti-human CD3 antibody (cat #144-020), anti-human CD28 mAb (cat #177-020), and human IL-2. Cells were harvested and 5 x 10⁵ cells per tube were washed and pre incubated ~5 minutes with 20 µl human IgG at 250 µg/ml (to block non specific binding) after which they were incubated 45 minutes on ice with 80 µl of anti-CD4/APC conjugate (cat #147-060), and anti-human GARP/PE at dilution factor of 1:50 (10 µg/ml). Cells were washed twice, fixed and analyzed by FACS, gating on CD4 positive lymphoid cells. A net 3.9% sub population of gated cells stained positive with a mean shift of 1.58 log₁₀ fluorescent units when compared to a Mouse IgG2a/PE isotype control (Catalog # 281-050). This binding was blocked when cells were pre incubated with 50 µg/ml unlabeled anti-human GARP mAb.

Blinding of anti-GARP/PE to CD4 positive stimulated human PBL



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