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Product Datasheet

APRIL Antibody [Aprily-8], IgG1, Unconjugated, Mouse, Monoclonal PRS-XA-1020

Artikelname	APRIL Antibody [Aprily-8], IgG1, Unconjugated, Mouse, Monoclonal
Artikelnummer	PRS-XA-1020
Hersteller Artikelnummer	XA-1020
Alternativnummer	PRS-XA-1020-0.1
Hersteller	ProSci
Wirt	Mouse
Kategorie	Antikörper
Applikation	FC, ICC, WB
Spezies Reaktivität	Human
Immunogen	APRIL (monoclonal Aprily-8) antibody was raised against recombinant human APRIL (aa. 93-233) expressed in bacteria.
Konjugation	Unconjugated
Klonalität	Monoclonal
Konzentration	batch dependent
Klon-Bezeichnung	[Aprily-8]
Isotyp	IgG1
NCBI	8741
UniProt	O75888

Puffer	PBS containing 0.02% sodium azide
Formulierung	Liquid
Application Verdünnung	Optimal dilutions for each application to be determined by the researcher.
Anwendungsbeschreibung	<p>APRIL antibody can be used to recognize human APRIL by Flow Cytometry, immunocytochemistry and Western Blot. Method: 3 days after transfection of cells with the indicated constructs, cells were fixed with 4% formaldehyde 5 min at RT. After a wash in PBS, samples were dehydrated by washes in 60%, 80%, 90%, 100% EtOH and xylol. Cells were then dried and embedded in paraffin. Sections were cut, mounted on slides and dried overnight at 50°C. Slides were then successively washed 2x 10min in xylol, 2x 10min in 100% EtOH, and then treated 10 min in 100% MeOH/ 0.6% H2O2 to inhibit endogenous peroxidase. Samples were rehydrated by washes in 90%, 80%, 60% EtOH and PBS. After micro-wave treatment, slides were washed 3x in PBS, blocked with IgG, and incubated for 1 hour with 5 µg/mL Aprily-8 or control mouse IgG (isotype control) in 1%BSA / 1x PBS for 1 hour. After PBS washes, samples were incubated with the secondary Ab for 1 hour, washed in PBS and revealed with StreptABComplex/HRP (Vector) and AEC. Method: HEK 293 cells were mock transfected or transfected with an expression plasmid coding for human APRIL. Cells (5x10⁵) were incubated on ice for 30 min in 50 µl FACS buffer (PBS, 5% fetal calf serum, 0.02% azide) containing 10 µg/mL of Aprily-8 antibody. After washing in FACS buffer, FITC-conjugated antibody to mouse IgG was added. Cells were incubated on ice for 30 min, washed and analyzed by flow cytometry. Method: 10 µg of protein was applied to the gel, and revealed with Aprily-8 (1 µg/mL) and HRP-coupled anti-mouse secondary antibody (1:4,000). Note: The hAPRIL construct used in this experiment is an uncleavable fusion protein between hBAFF (aa 1-132) and hAPRIL (aa 93-233). This product is for research use only.</p>