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anti-FYN antibody (C-Term)

Images



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Target:

Quantity:	100 μL
Target:	FYN
Binding Specificity:	C-Term
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This FYN antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human Fyn, corresponding to a region within C-terminal amino acids.
Isotype:	IgG
Specificity:	Fyn Antibody detects endogenous levels of total Fyn.
Predicted Reactivity:	Pig,Bovine,Sheep,Rabbit,Dog,Chicken,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	

FYN

Alternative Name:	FYN (FYN Products)
Background:	Description: Non-receptor tyrosine-protein kinase that plays a role in many biological processe
	including regulation of cell growth and survival, cell adhesion, integrin-mediated signaling,
	cytoskeletal remodeling, cell motility, immune response and axon guidance. Inactive FYN is
	phosphorylated on its C-terminal tail within the catalytic domain. Following activation by PKA,
	the protein subsequently associates with PTK2/FAK1, allowing PTK2/FAK1 phosphorylation,
	activation and targeting to focal adhesions. Involved in the regulation of cell adhesion and
	motility through phosphorylation of CTNNB1 (beta-catenin) and CTNND1 (delta-catenin).
	Regulates cytoskeletal remodeling by phosphorylating several proteins including the actin
	regulator WAS and the microtubule-associated proteins MAP2 and MAPT. Promotes cell
	survival by phosphorylating AGAP2/PIKE-A and preventing its apoptotic cleavage. Participates
	in signal transduction pathways that regulate the integrity of the glomerular slit diaphragm (an
	essential part of the glomerular filter of the kidney) by phosphorylating several slit diaphragm
	components including NPHS1, KIRREL1 and TRPC6. Plays a role in neural processes by
	phosphorylating DPYSL2, a multifunctional adapter protein within the central nervous system,
	ARHGAP32, a regulator for Rho family GTPases implicated in various neural functions, and
	SNCA, a small pre-synaptic protein. Participates in the downstream signaling pathways that
	lead to T-cell differentiation and proliferation following T-cell receptor (TCR) stimulation. Also
	participates in negative feedback regulation of TCR signaling through phosphorylation of PAG
	thereby promoting interaction between PAG1 and CSK and recruitment of CSK to lipid rafts.
	CSK maintains LCK and FYN in an inactive form. Promotes CD28-induced phosphorylation of
	VAV1.
	Gene: FYN
Molecular Weight:	60kDa
Gene ID:	2534
UniProt:	P06241
Pathways:	JAK-STAT Signaling, TCR Signaling, Fc-epsilon Receptor Signaling Pathway, EGFR Signaling
	Pathway, Neurotrophin Signaling Pathway, Feeding Behaviour, CXCR4-mediated Signaling
	Events, Signaling Events mediated by VEGFR1 and VEGFR2, Activated T Cell Proliferation,
	Thromboxane A2 Receptor Signaling
Application Details	
Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000

Application Details

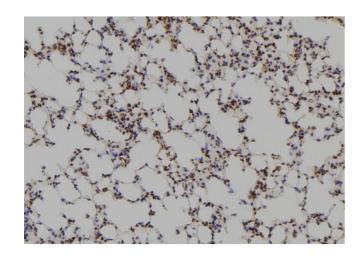
ed saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 %
azide: a POISONOUS AND HAZARDOUS SUBSTANCE which raff only.
6

Store at -20 °C. Stable for 12 months from date of receipt.

Images

Expiry Date:

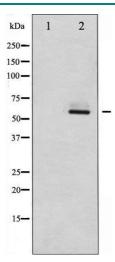
Storage Comment:



12 months

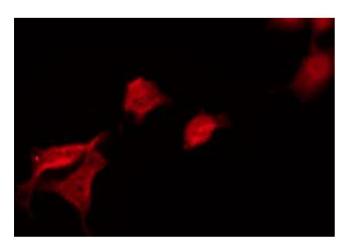
Immunohistochemistry

Image 1. ABIN6269100 at 1/100 staining Mouse lung tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.



Western Blotting

Image 2. Western blot analysis of Fyn expression in 293 whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



Immunofluorescence (fixed cells)

Image 3. ABIN6269100 staining 293 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.