SunLong Biotech Co.,LTD
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[Marker Primary Antibody]

Rabbit Anti-CD19/AF594 Conjugated antibody

Catalog Number: SL0079R-AF594

FOR RESEARCH USE ONLY

[DATASHEET]

Host:	Rabbit
Isotype:	IgG
Clonality:	Polyclonal
Target:	Anti-CD19/AF594
Immunogen:	KLH conjugated synthetic peptide derived from human CD19 (485-526aa)
Purification:	affinity purified by Protein A
Concentration:	1mg/ml
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 1M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
Applications:	Flow-Cyt=1μg/Test IF=1:100-500 not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Reactivity:	Human,Mouse(predicted:Rat,Pig,Cow,Horse,GuineaPig)
Background:	background: CD19 is a transmembrane glycoprotein that is delectively expressed on the cell surface of B-lymphocytes,where it activates intracellular signaling cascades involving both Ras and phosphatidylinositol 3-kinase pathways.Lymphocytes proliferate and differentiate in response to various concentrations of different antigens. The ability of the B cell to respond in a specific, yet sensitive manner to the various antigens is achieved with the use of low-affinity antigen receptors. This gene encodes a cell surface molecule which assembles with the antigen receptor of B lymphocytes in order to decrease the threshold for antigen receptor-dependent stimulation. **Function:** Assembles with the antigen receptor of B-lymphocytes in order to decrease the threshold for antigen receptor-dependent stimulation. **Subunit:** Forms a complex with CD21, CD81 and CD225 in the membrane of mature B-cells. Interacts with VAV. Interacts with GRB2 and SOS when phosphorylated on Tyr-348 and/or Tyr-378. Interacts with PLCG2 when

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	phosphorylated on Tyr-409. Interacts with LYN. **Subcellular Location:** Membrane; Single-pass type I membrane protein. **Post-translational modifications:** Phosphorylated on serine and threonine upon DNA damage, probably by ATM or ATR. Phosphorylated on tyrosine following B-cell activation. Phosphorylated on tyrosine residues by LYN. DISEASE: Defects in CD19 are the cause of immunodeficiency common variable type 3 (CVID3) [MIM:613493]; also called antibody deficiency due to CD19 defect. CVID3 is a primary immunodeficiency characterized by antibody deficiency, hypogammaglobulinemia, recurrent bacterial infections and an inability to mount an antibody response to antigen. The defect results from a failure of B-cell differentiation and impaired secretion of immunoglobulins; the numbers of circulating B-cells is usually in the normal range, but can be low. **Similarity:** Contains 2 Ig-like C2-type (immunoglobulin-like) domains. **Database links:** **Entrez Gene:** 930 Human **Entrez Gene:** 12478 Mouse **Entrez Gene:** 365367 Rat Omim: 107265 Human **SwissProt:** P15391 Human **SwissProt:** P25918 Mouse **Unigene:** 652262 Human **Unigene:** 4360 Mouse Important **Note:** This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications. CD19 is a plasma membrane protein involved in signal transduction. It is expressed on the surface of pre-B cells and mature B The cell membrane and is associated with the regulation of B cell activation and developmental regulation; it is not expressed on T cells or normal granulocytes. This antibody can specifically recognize CD19 and is mainly used to label normal B cells and neoplastic B cells.