

## Rabbit Anti-MS4A15 antibody

SL13691R

Product Name:	MS4A15
Chinese Name:	MS4A15蛋白抗体
Alias:	FLJ34527; Hypothetical protein MGC35295; M4A15_HUMAN; Membrane spanning 4 domains subfamily A member 15; Membrane-spanning 4-domains subfamily A member 15; MGC35295; MS4A15.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Dog, Pig, Cow, Horse, Rabbit, Sheep,
Applications:	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100- 500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	25kDa
<b>Cellular localization:</b>	The cell membrane
Form:	Lyophilized or Liquid
<b>Concentration:</b>	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human MS4A15:31-130/240
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
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 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
Product Detail:	MS4A (membrane-spanning 4-domain family, subfamily A) is a large family of proteins that includes at least 26 members in mouse and humans. Flanked by amino- and carboxyl- cytoplasmic regions, MS4A family members contain four highly conserved transmembrane domains. CD20, the most well-known MS4A family member, is a B- cell-specific molecule that functions as a calcium-permeable cation channel and is

known to accelerate the G0 to G1 progression induced by IGF-1. MS4A15 (membranespanning 4-domains, subfamily A, member 15) is a 240 amino acid multi-pass membrane protein that exists as two alternatively spliced isoforms. The gene encoding MS4A15 maps to human chromosome 11, which houses over 1,400 genes and comprises nearly 4% of the human genome. Jervell and Lange-Nielsen syndrome, Jacobsen syndrome, Niemann-Pick disease, hereditary angioedema and Smith-Lemli-Opitz syndrome are associated with defects in genes that maps to chromosome 11.

## Function:

May be involved in signal transduction as a component of a multimeric receptor complex.

Subcellular Location: Membrane.

joiotech.col Similarity: Belongs to the MS4A family.

SWISS: O8N5U1

Gene ID: 219995

Database links:

Entrez Gene: 219995 Human

Entrez Gene: 545279 Mouse

SwissProt: Q8N5U1 Human

SwissProt: Q3UPL6 Mouse

Unigene: 207465 Human

Unigene: 329729 Mouse

## **Important Note:**

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.