

# Rabbit Anti-MXRA8 antibody

## SL19118R

| <b>Product Name:</b>   | MXRA8   |
|------------------------|---|
| Chinese Name:          | 基质重塑相关蛋白8抗体   |
| Alias:                 | Limitrin; Matrix remodelling associated 8; Matrix-remodeling-associated protein8; Mxra8; MXRA8 HUMAN.   |
| Organism Species:      | Rabbit  |
| Clonality:             | Polyclonal  |
| React Species:         | Human, Mouse, Rat, Dog, Pig, Cow, Sheep,  |
| Applications:          | WB=1:500-2000ELISA=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:      | 47kDa   |
| Cellular localization: | The cell membrane   |
| Form:                  | Lyophilized or Liquid   |
| Concentration:         | 1mg/ml  |
| immunogen:             | KLH conjugated synthetic peptide derived from human MXRA8:181-280/442 <extracellular></extracellular>   |
| 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:                | <u>PubMed</u>   |
| Product Detail:        | MXRA8 is a 442 amino acid single-pass type I membrane protein that is implicated in the maintenance and maturation of the blood-brain barrier. Containing two Ig-like V-type (immunoglobulin-like) domains, MXRA8 exists as two alternatively spliced isoforms that are encoded by a gene that maps to human chromosome 1p36.33. Human chromosome 1 spans 260 million base pairs, contains over 3,000 genes and comprises |

nearly 8% of the human genome. Chromosome 1 houses a large number of disease-associated genes, including those that are involved in familial adenomatous polyposis, Stickler syndrome, Parkinson's disease, Gaucher disease, schizophrenia and Usher syndrome. Aberrations in chromosome 1 are found in a variety of cancers, including head and neck cancer, malignant melanoma and multiple myeloma

#### **Function:**

May play a role in the maturation and maintenance of blood-brain barrier.

### Subcellular Location:

Membrane.

#### Similarity:

Contains 2 Ig-like V-type (immunoglobulin-like) domains.

#### **SWISS:**

Q9BRK3

#### Gene ID:

54587

#### Database links:

Entrez Gene: 54587 Human

Entrez Gene: 313770 Rat

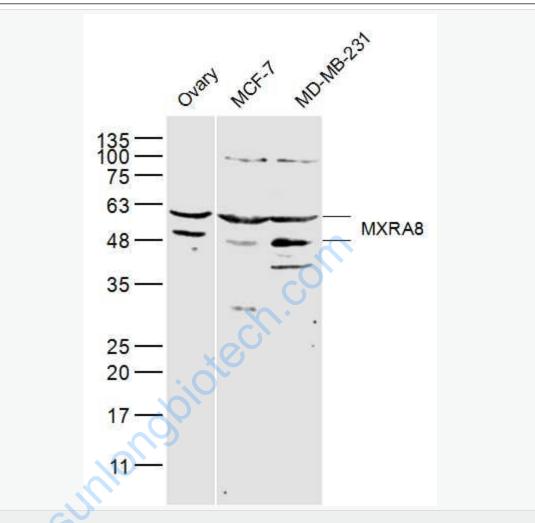
SwissProt: Q9BRK3 Human

SwissProt: Q5XI43 Rat

Unigene: 515687 Human

#### **Important Note:**

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



Picture:

Sample:

Ovary (Mouse) Lysate at 40 ug

MCF-7(Human) Cell Lysate at 30 ug

MD-MB-231(Human) Cell Lysate at 30 ug

Primary: Anti-MXRA8 (SL19118R) at 1/300 dilution

Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution

Predicted band size: 47 kD

Observed band size: 47 kD