

Rabbit Anti-MOBKL2A antibody

SL17699R

| Product Name: | MOBKL2A |
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| Chinese Name: | MOBKL2A蛋白抗体 |
| Alias: | MOB-LAK; Mob1 homolog 2A; MOBKL2A; MOL2A_HUMAN; Mps one binder |
| | kinase activator-like 2A; Protein Mob3A。 |
| Organism Species: | Rabbit |
| Clonality: | Polyclonal |
| React Species: | Human,Mouse,Rat,Dog, |
| | ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100- |
| Applications: | 500 (Paraffin sections need antigen repair) |
| Applications. | not yet tested in other applications. |
| | optimal dilutions/concentrations should be determined by the end user. |
| Molecular weight: | 25kDa |
| Form: | Lyophilized or Liquid |
| Concentration: | 1mg/ml |
| immunogen: | KLH conjugated synthetic peptide derived from human MOBKL2A:61-160/217 |
| 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: | MOBKL2A is a 217 amino acid protein that regulates kinase activity. A member of the MOB1/phocein family, MOBKL2A is encoded by a gene that maps to human chromosome 19. Consisting of around 63 million bases with over 1,400 genes, chromosome 19 makes up over 2% of human genomic DNA. Chromosome 19 includes a diversity of interesting genes and is recognized for having the greatest gene density of the human chromosomes. It is the genetic home for a number of immunoglobulin superfamily members including the killer cell and leukocyte Ig-like receptors, a number |

| of ICAMs, the CEACAM and PSG family, and Fc?receptors. Key genes for eye color and hair color also map to chromosome 19. Peutz-Jeghers syndrome, spinocerebellar ataxia type 6, the stroke disorder CADASIL, hypercholesterolemia and insulin- dependent diabetes have been linked to chromosome 19. |
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| Function: |
| May regulate the activity of kinases. |
| Similarity: |
| Belongs to the MOB1/phocein family. |
| SWISS: |
| Q96BX8 |
| Gene ID: |
| 126308 |
| Q96BX8 Gene ID: 126308 Database links: Entrez Gene: 126308 Human Entrez Gene: 208228 Mouse |
| Entrez Gene: 126308 Human |
| Entrez Gene: 208228 Mouse |
| Entrez Gene: 362833 Rat |
| SwissProt: Q96BX8 Human |
| SwissProt: Q8BSU7 Mouse |
| Unigene: 86912 Human |
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| Important Note: |
| This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications. |
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