

C7orf64 Rabbit pAb

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Catalog # AP55916

Product Information

| | |
|--------------------------------|--|
| Application | WB |
| Primary Accession | Q5RL73 |
| Reactivity | Human |
| Predicted | Mouse, Rat, Chicken, Dog, Horse, Rabbit, Sheep |
| Host | Rabbit |
| Clonality | Polyclonal |
| Calculated MW | 41808 |
| Physical State | Liquid |
| Immunogen | KLH conjugated synthetic peptide derived from human C7orf64 |
| Epitope Specificity | 1-100/367 |
| Isotype | IgG |
| Purity | affinity purified by Protein A |
| Buffer | 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. |
| SIMILARITY | Belongs to the RBM48 family. Contains 1 RRM (RNA recognition motif) domain. |
| Important Note | This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications. |
| Background Descriptions | Chromosome 7 has been linked to Osteogenesis imperfecta, Pendred syndrome, Lissencephaly, Citrullinemia and Shwachman-Diamond syndrome. The deletion of a portion of the q arm of chromosome 7 is associated with Williams-Beuren syndrome, a condition characterized by mild mental retardation, an unusual comfort and friendliness with strangers and an elfin appearance. Deletions of portions of the q arm of chromosome 7 are also seen in a number of myeloid disorders including cases of acute myelogenous leukemia and myelodysplasia. The C7orf64 gene product has been provisionally designated C7orf64 pending further characterization. |

Additional Information

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|--------------------|---|
| Gene ID | 84060 |
| Other Names | RNA-binding protein 48, RBM48, C7orf64 |
| Dilution | WB=1:500-2000 |
| Storage | Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. 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. |

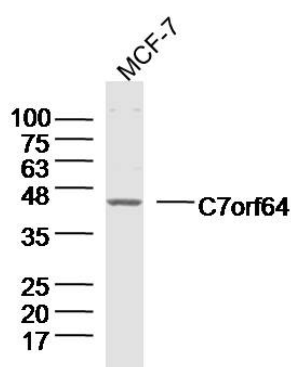
Protein Information

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|-----------------|---|
| Name | RBM48 |
| Synonyms | C7orf64 |
| Function | As a component of the minor spliceosome, involved in the splicing of U12-type introns in pre-mRNAs. |

Background

Chromosome 7 has been linked to Osteogenesis imperfecta, Pendred syndrome, Lissencephaly, Citrullinemia and Shwachman-Diamond syndrome. The deletion of a portion of the q arm of chromosome 7 is associated with Williams-Beuren syndrome, a condition characterized by mild mental retardation, an unusual comfort and friendliness with strangers and an elfin appearance. Deletions of portions of the q arm of chromosome 7 are also seen in a number of myeloid disorders including cases of acute myelogenous leukemia and myelodysplasia. The C7orf64 gene product has been provisionally designated C7orf64 pending further characterization.

Images



Sample: MCF-7 Cell (Human) Lysate at 40 ug
 Primary: Anti-C7orf64(AP55916) at 1/300 dilution
 Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution
 Predicted band size: 42kD
 Observed band size: 42kD

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.