## MAGEA11 antibody

Catalog No: #22350



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

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| Product Name          | MAGEA11 antibody  |  |
|-----------------------|---|--|
| Host Species          | Rabbit  |  |
| Clonality             | Polyclonal  |  |
| Purification          | Purified by antigen-affinity chromatography.  |  |
| Applications          | WB IHC IF   |  |
| Species Reactivity    | Hu  |  |
| Immunogen Type        | Peptide   |  |
| Immunogen Description | Synthetic peptide contain a sequence corresponding to a region within amino acids 78 and 143 of MAGEA11 |  |
| Target Name           | MAGEA11   |  |
| Accession No.         | Swiss-Prot:P43364Gene ID:4110   |  |
| Uniprot               | P43364  |  |
| GeneID                | 4110;   |  |
| Concentration         | 1mg/ml  |  |
| Formulation           | Supplied in 0.1M Tris-buffered saline with 10% Glycerol (pH7.0). 0.01% Thimerosal was added as a        |  |
|                       | preservative.   |  |
| Storage               | Store at -20°C for long term preservation (recommended). Store at 4°C for short term use.               |  |
|                       |   |  |

## Application Details

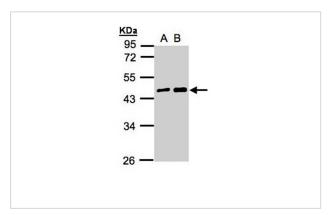
Predicted MW: 48kd

Western blotting: 1:500-1:3000

Immunohistochemistry: 1:100-1:500

Immunofluorescence: 1:100-1:200

## **Images**



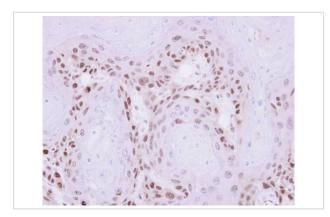
Sample(30 ug whole cell lysate)

A: A431

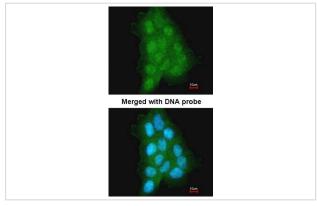
B: H1299

10% SDS PAGE

Primary antibody diluted at 1: 5000



Immunohistochemical analysis of paraffin-embedded Ca922 xenograft, using MAGEA11 antibody at 1: 500 dilution.



Immunofluorescence analysis of paraformaldehyde-fixed A431, using MAGEA11 antibody at 1: 500 dilution.

## Background

This gene is a member of the MAGEA gene family. The members of this family encode proteins with 50 to 80% sequence identity to each other. The promoters and first exons of the MAGEA genes show considerable variability, suggesting that the existence of this gene family enables the same function to be expressed under different transcriptional controls. The MAGEA genes are clustered at chromosomal location Xq28. They have been implicated in some hereditary disorders, such as dyskeratosis congenita. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq]

Note: This product is for in vitro research use only