

IGF2BP1/IMP1 Rabbit mAb

Catalog No.: A25715 **Recombinant**

Basic Information

Observed MW

64kDa

Calculated MW

49kDa/63kDa

Category

Primary antibody

Applications

ELISA, WB, IHC-P, IF/ICC, IP

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC66817

Background

This gene encodes a member of the insulin-like growth factor 2 mRNA-binding protein family. The protein encoded by this gene contains four K homology domains and two RNA recognition motifs. It functions by binding to the mRNAs of certain genes, including insulin-like growth factor 2, beta-actin and beta-transducin repeat-containing protein, and regulating their translation. Two transcript variants encoding different isoforms have been found for this gene.

Recommended Dilutions

WB	1:500 - 1:1000
IHC-P	1:50 - 1:200
IF/ICC	1:50 - 1:200
IP	0.5µg-4µg antibody for 400µg-600µg extracts of whole cells

Immunogen Information

Gene ID

10642

Swiss Prot

Q9NZI8

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 101-200 of human IGF2BP1/IMP1(NP_006537.3).

Synonyms

IMP1; ZBP1; CRDBP; IMP-1; CRD-BP; VICKZ1

Contact

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Product Information

Source

Rabbit

Isotype

IgG

Purification

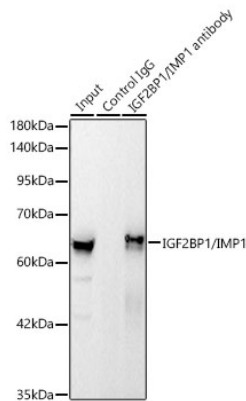
Affinity purification

Storage

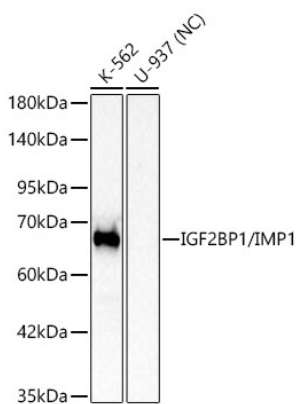
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300, 0.05% BSA, 50% glycerol, pH7.3.

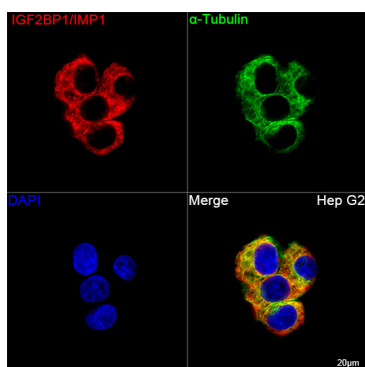
Validation Data



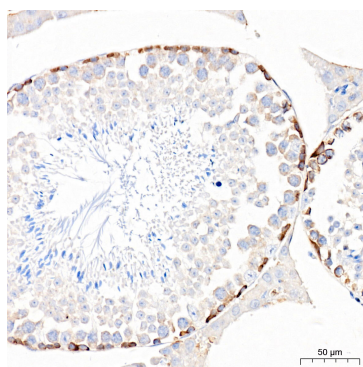
Immunoprecipitation of IGF2BP1/IMP1 from 500 µg extracts of 293F cells was performed using 2 µg of IGF2BP1/IMP1 Rabbit mAb (A25715). Rabbit IgG isotype control (AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X non-reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using IGF2BP1/IMP1 Rabbit mAb (A25715) at a dilution of 1:1000.



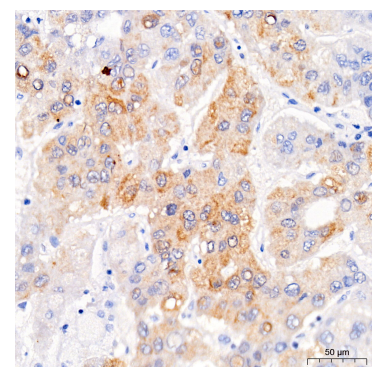
Western blot analysis of various lysates using IGF2BP1/IMP1 Rabbit mAb (A25715) at 1:1000 dilution incubated overnight at 4°C.
 Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
 Lysates/proteins: 25 µg per lane.
 Blocking buffer: 3% nonfat dry milk in TBST.
 Detection: ECL Basic Kit (RM00020)
 Negative control (NC): U-937
 Exposure time: 1 s.



Confocal immunofluorescence analysis of Hep G2 cells using IGF2BP1/IMP1 Rabbit mAb (A25715, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with Alpha-tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

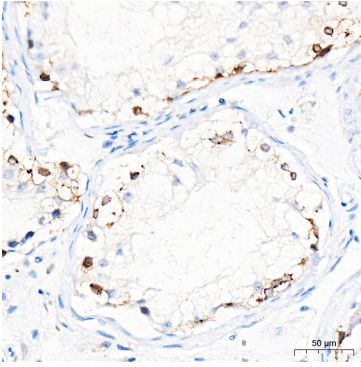


Immunohistochemistry analysis of IGF2BP1/IMP1 in paraffin-embedded Mouse testis tissue using IGF2BP1/IMP1 Rabbit mAb (A25715) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.

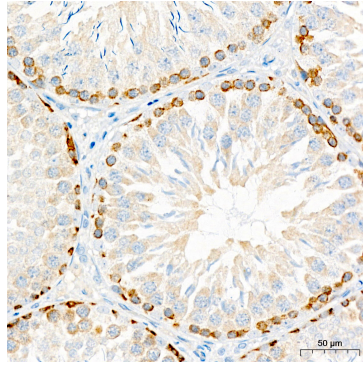


Immunohistochemistry analysis of IGF2BP1/IMP1 in paraffin-embedded Human liver cancer tissue using IGF2BP1/IMP1 Rabbit mAb (A25715) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.

Validation Data



Immunohistochemistry analysis of IGF2BP1/IMP1 in paraffin-embedded Human testis tissue using IGF2BP1/IMP1 Rabbit mAb (A25715) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of IGF2BP1/IMP1 in paraffin-embedded Rat testis tissue using IGF2BP1/IMP1 Rabbit mAb (A25715) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.