Amnionless (F-7): sc-365734



The Power to Question

BACKGROUND

Megaloblastic anemia 1 (MGA1), also referred to as MGA1 Norwegian type or Imerslund-Gräsbeck syndrome (I-GS), is a hereditary, recessive disorder caused by defects in the AMN gene. Patients suffering from MGA1 have a selective malabsorption of vitamin B_{12} , causing impaired function of thymidine synthase which in turn interrupts DNA synthesis. Amnionless protein, encoded for by the AMN gene, is crucial for vitamin B_{12} absorption. It modulates a BMP (bone morphogenetic protein) signaling pathway and is therefore important for trunk mesoderm production during development. Amnionless is a membrane protein that interacts with cubulin and is primarily expressed in colon, kidney and small intestine. Shorter isoforms can also be detected in thymus, testis and peripheral blood leukocytes.

REFERENCES

- Tomihara-Newberger, C., et al. 1998. The AMN gene product is required in extraembryonic tissues for the generation of middle primitive streak derivatives. Dev. Biol. 204: 34-54.
- Kalantry, S., et al. 2001. The Amnionless gene, essential for mouse gastrulation, encodes a visceral-endoderm-specific protein with an extracellular cysteine-rich domain. Nat. Genet. 27: 412-416.
- Tanner, S.M., et al. 2003. Amnionless, essential for mouse gastrulation, is mutated in recessive hereditary megaloblastic anemia. Nat. Genet. 33: 426-429.
- Strope, S., et al. 2004. Mouse Amnionless, which is required for primitive streak assembly, mediates cell-surface localization and endocytic function of cubilin on visceral endoderm and kidney proximal tubules. Development 131: 4787-4795.
- He, Q., et al. 2005. Amnionless function is required for cubilin brush-border expression and intrinsic factor-cobalamin (vitamin B₁₂) absorption *in vivo*. Blood 106: 1447-1453.
- 6. Gräsbeck, R. 2006. Imerslund-Gräsbeck syndrome (selective vitamin B_{12} malabsorption with proteinuria). Orphanet J. Rare Dis. 1: 17.

CHROMOSOMAL LOCATION

Genetic locus: AMN (human) mapping to 14q32.32; Amn (mouse) mapping to 12 F1.

SOURCE

Amnionless (F-7) is a mouse monoclonal antibody raised against amino acids 20-300 mapping near the N-terminus of Amnionless of human origin.

PRODUCT

Each vial contains 200 $\mu g \; lg G_1$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

Amnionless (F-7) is recommended for detection of Amnionless of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for Amnionless siRNA (h): sc-60069, Amnionless siRNA (m): sc-60070, Amnionless shRNA Plasmid (h): sc-60069-SH, Amnionless shRNA Plasmid (m): sc-60070-SH, Amnionless shRNA (h) Lentiviral Particles: sc-60069-V and Amnionless shRNA (m) Lentiviral Particles: sc-60070-V.

Molecular Weight of extracellular Amnionless domain: 35 kDa.

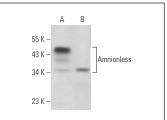
Molecular Weight of membrane-bound Amnionless: 45 kDa.

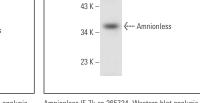
Positive Controls: Caki-1 cell lysate: sc-2224, KNRK whole cell lysate: sc-2214 or Hep G2 cell lysate: sc-2227.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA





Amnionless (F-7): sc-365734. Western blot analysis of Amnionless expression in Hep G2 (A) and KNRK (B) whole cell lysates.

Amnionless (F-7): sc-365734. Western blot analysis of Amnionless expression in Caki-1 whole cell lysate.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.