

HAS2 Rabbit pAb

HAS2 Rabbit pAb Catalog # AP50809

#### **Product Information**

Application WB Primary Accession Q92819

**Reactivity** Rat, Pig, Rabbit, Chicken, Horse

Host Rabbit
Clonality Polyclonal
Calculated MW 63566
Physical State Liquid

**Immunogen** KLH conjugated synthetic peptide derived from human HAS2/Hyaluronan

synthase 2

**Epitope Specificity** 401-500/552

**Isotype** IgG

**Purity** affinity purified by Protein A

**Buffer** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

**SUBCELLULAR LOCATION** Membrane.

**SIMILARITY** Belongs to the nodC/HAS family.

**Post-translational**Autophosphorylated on several Ser and Thr residues. Autophosphorylation of Thr-451 is dependent on Thr-446 and is stimulated by dsRNA binding and

dimerization. Autophosphorylation apparently leads to the activation of the

kinase.

**DISEASE**Note=A chromosomal aberration involving HAS2 may be a cause of

lipoblastomas, which are benign tumors resulting from transformation of adipocytes, usually diagnosed in children. 8q12.1 to 8q24.1 intrachromosomal

rearrangement with PLAG1.

**Important Note** This product as supplied is intended for research use only, not for use in

human, therapeutic or diagnostic applications.

**Background Descriptions** Hyaluronan or hyaluronic acid (HA) is a high molecular weight unbranched

polysaccharide synthesized by a wide variety of organisms from bacteria to mammals, and is a constituent of the extracellular matrix. It consists of alternating glucuronic acid and N-acetylglucosamine residues that are linked

by beta-1-3 and beta-1-4 glycosidic bonds. HA is synthesized by

membrane-bound synthase at the inner surface of the plasma membrane,

and the chains are extruded through pore-like structures into the

extracellular space. It serves a variety of functions, including space filling, lubrication of joints, and provision of a matrix through which cells can migrate. HA is actively produced during wound healing and tissue repair to provide a framework for ingrowth of blood vessels and fibroblasts. Changes in

the serum concentration of HA are associated with inflammatory and degenerative arthropathies such as rheumatoid arthritis. In addition, the interaction of HA with the leukocyte receptor CD44 is important in

tissue-specific homing by leukocytes, and overexpression of HA receptors has been correlated with tumor metastasis. HAS2 is a member of the newly identified vertebrate gene family encoding putative hyaluronan synthases,

and its amino acid sequence shows significant homology to

glycosaminoglycan synthetase (DG42) from Xenopus laevis, and human and murine hyaluronan synthase 1. [provided by RefSeq, Jul 2008]

#### **Additional Information**

Gene ID 3037

Other Names Hyaluronan synthase 2, 2.4.1.212, Hyaluronate synthase 2, Hyaluronic acid

synthase 2, HA synthase 2, HAS2 (HGNC:4819)

**Target/Specificity** Expressed in fibroblasts.

**Dilution** WB=1:500-2000,Flow-Cyt=1ug/Test

**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**

Name HAS2 ( <u>HGNC:4819</u>)

**Function** Catalyzes the addition of GlcNAc or GlcUA monosaccharides to the nascent

hyaluronan polymer (Probable) (PubMed:<u>20507985</u>, PubMed:<u>21228273</u>,

PubMed: 23303191, PubMed: 32993960). Therefore, it is essential to hyaluronan synthesis a major component of most extracellular matrices that has a structural role in tissues architectures and regulates cell adhesion, migration and differentiation (PubMed: 20507985, PubMed: 21228273, PubMed: 8798477). This is one of three isoenzymes responsible for cellular hyaluronan synthesis and it is particularly responsible for the synthesis of

high molecular mass hyaluronan (By similarity).

**Cellular Location** Cell membrane; Multi-pass membrane protein Endoplasmic reticulum

membrane; Multi- pass membrane protein. Vesicle. Golgi apparatus membrane; Multi-pass membrane protein. Lysosome Note=Travels from endoplasmic reticulum (ER), Golgi to plasma membrane and either back to

endosomes and lysosomes, or out into extracellular vesicles

(PubMed:30394292). Post-translational modifications control HAS2 trafficking

(PubMed:30394292).

**Tissue Location** Expressed in fibroblasts.

# Background

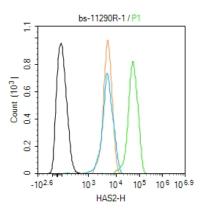
Hyaluronan or hyaluronic acid (HA) is a high molecular weight unbranched polysaccharide synthesized by a wide variety of organisms from bacteria to mammals, and is a constituent of the extracellular matrix. It consists of alternating glucuronic acid and N-acetylglucosamine residues that are linked by beta-1-3 and beta-1-4 glycosidic bonds. HA is synthesized by membrane-bound synthase at the inner surface of the plasma membrane, and the chains are extruded through pore-like structures into the extracellular space. It serves a variety of functions, including space filling, lubrication of joints, and provision of a matrix through which cells can migrate. HA is actively produced during wound healing and tissue repair to provide a framework for ingrowth of blood vessels and fibroblasts. Changes in the serum concentration of HA are associated with inflammatory and degenerative arthropathies such as rheumatoid arthritis. In addition, the interaction of HA with the leukocyte receptor CD44 is important in tissue-specific homing by leukocytes, and overexpression of HA receptors has been correlated with tumor metastasis. HAS2 is a member of the newly

identified vertebrate gene family encoding putative hyaluronan synthases, and its amino acid sequence shows significant homology to glycosaminoglycan synthetase (DG42) from Xenopus laevis, and human and murine hyaluronan synthase 1. [provided by RefSeq, Jul 2008]

## References

Watanabe K., et al.J. Biol. Chem. 271:22945-22948(1996). Morerio C., et al. Cancer Genet. Cytogenet. 156:183-184(2005).

## **Images**



Blank control(black line):Hela.

Primary Antibody (green line): Rabbit Anti-HAS2 antibody

(AP50809)

Dilution:1ug/Test;

Secondary Antibody(white blue line): Goat anti-rabbit

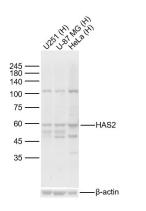
IgG-AF488

Dilution: 0.5ug/Test.

Isotype control(orange line): Normal Rabbit IgG

Protocol

The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C, The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.



#### Sample:

Lane 1: Human U251 cell Lysates Lane 2: Human U-87 MG cell Lysates Lane 3: Human Hela cell Lysates

Primary: Anti-HAS2 (AP50809) at 1/1000 dilution

Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000

dilution

Predicted band size: 64kDa Observed band size: 60kDa

#### **Citations**

• Role of Hyaluronan and Glucose on 4-Methylumbelliferone-inhibited Cell Proliferation in Breast Carcinoma Cells.

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