

## **Versican Polyclonal Antibody**

Catalog No: YT4874

**Reactivity:** Human; Rat; Mouse;

**Applications:** WB;IHC;IF;ELISA

Target: Versican

**Fields:** >>Cell adhesion molecules

P13611

Q62059

Gene Name: VCAN

Protein Name: Versican core protein

Human Gene Id: 1462

**Human Swiss Prot** 

Tullian Swiss F10

No:

**Mouse Swiss Prot** 

No:

**Immunogen:** The antiserum was produced against synthesized peptide derived from human

VCAN. AA range:532-581

**Specificity:** Versican Polyclonal Antibody detects endogenous levels of Versican protein.

**Formulation :** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution :** WB 1:500-2000 IHC 1:100 - 1:300. ELISA: 1:10000. IF 1:50-200

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

1/3



Molecularweight: 373kD

**Cell Pathway:** Cell adhesion molecules (CAMs);

**Background:** This gene is a member of the aggrecan/versican proteoglycan family. The

protein encoded is a large chondroitin sulfate proteoglycan and is a major component of the extracellular matrix. This protein is involved in cell adhesion, proliferation, proliferation, migration and angiogenesis and plays a central role in tissue morphogenesis and maintenance. Mutations in this gene are the cause of Wagner syndrome type 1. Multiple transcript variants encoding different isoforms

have been found for this gene. [provided by RefSeq, Aug 2009],

**Function:** alternative products:Additional isoforms seem to exist, developmental

stage:Disappears after the cartilage development., disease:Defects in VCAN are the cause of Wagner syndrome type 1 (WGN1) [MIM:143200]. WGN is a dominantly inherited vitreoretinopathy characterized by an optically empty vitreous cavity with fibrillary condensations and a preretinal avascular membrane.

Other optical features include progressive chorioretinal atrophy, perivascular sheating, subcapsular cataract and myopia. Systemic manifestations are absent in WGN.,function:May play a role in intercellular signaling and in connecting cells with the extracellular matrix. May take part in the regulation of cell motility, growth

and differentiation. Binds hyaluronic acid., online

information:Versican,similarity:Belongs to the aggrecan/versican proteoglycan

family., similarity: Contains 1 C-type lectin domain., similarity: Contains

Subcellular Location : Secreted, extracellular space, extracellular matrix. Cell projection, cilium, photoreceptor outer segment. Secreted, extracellular space, extracellular matrix,

interphotoreceptor matrix.

**Expression:** Expressed in the retina (at protein level) (PubMed:29777959). Cerebral white

matter and plasma (PubMed:2469524). Isoform V0: Expressed in normal brain, gliomas, medulloblastomas, schwannomas, neurofibromas, and meningiomas (PubMed:8627343). Isoform V1: Expressed in normal brain, gliomas, medulloblastomas, schwannomas, neurofibromas, and meningiomas

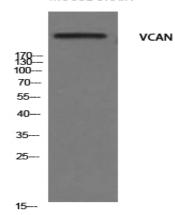
(PubMed:8627343). Isoform V2: Restricted to normal brain and gliomas (PubMed:8627343). Isoform V3: Found in all these tissues except

medulloblastomas (PubMed:8627343).

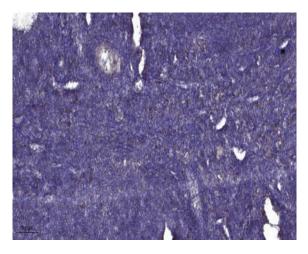
## **Products Images**



## MOUSE-BRAIN



Western Blot analysis of mouse-brain cells using Versican Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 30min).