# abcam

### Product datasheet

## Human Versican peptide ab39784

| Description  |   |  |
|--|---|--|
| Product name   | Human Versican peptide  |  |
| Purity   | > 90 % SDS-PAGE.  |  |
|  | This peptide is greater than 70% pure.  |  |
|  |   |  |
| Animal free  | No  |  |
| Nature   | Synthetic   |  |
| Amino Acid Sequence 1  |   |  |
| Species  | Human   |  |
| Sequence   | CGG-DPEAAE  |  |
| Amino acids  | 436 to 441  |  |
| Amino Acid Sequence 2  |   |  |
| Species  | Human   |  |
| Sequence   | CGG-DPEAAE  |  |
| Amino acids  | 436 to 441  |  |
|  |   |  |
| Specifications   |   |  |
| Our Abpromise guarantee covers the use of ab39784 in the following tested applications.  |   |  |
| The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user. |   |  |
| Applications   | Blocking - Blocking peptide for Anti-Versican antibody (ab19345)                      |  |
|  | Neutralising  |  |
| Form   | Lyophilized   |  |
|  |   |  |
| Preparation and Storage  |   |  |
| Stability and Storage  | Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles. |  |
| ·····, ·····, ······   | Double distilled water or equivalent after reconstitution.                            |  |
| Reconstitution   | Reconstitute with 0.1 ml of distilled water.  |  |
|  |   |  |

| 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.   |
|----------------------------------|--|
| Tissue specificity               | Cerebral white matter and plasma. Isoform V0 and isoform V1 are expressed in normal brain, gliomas, medulloblastomas, schwannomas, neurofibromas, and meningiomas. Isoform V2 is restricted to normal brain and gliomas. Isoform V3 is found in all these tissues except medulloblastomas.   |
| Involvement in 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. |
| Sequence similarities            | Belongs to the aggrecan/versican proteoglycan family.<br>Contains 1 C-type lectin domain.<br>Contains 2 EGF-like domains.<br>Contains 1 Ig-like V-type (immunoglobulin-like) domain.<br>Contains 2 Link domains.<br>Contains 1 Sushi (CCP/SCR) domain.   |
| Developmental stage              | Disappears after the cartilage development.  |
| Post-translational modifications | Phosphorylation sites are present in the extracelllular medium.  |
| Cellular localization            | Secreted > extracellular space > extracellular matrix.   |

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit <u>https://www.abcam.com/abpromise</u> or contact our technical team.

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