

Product datasheet for TP723443

OriGene Technologies, Inc.

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Thrombomodulin (THBD) (NM_000361) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Human thrombomodulin (THBD).

Species: Human Expression Host: HEK293

Expression cDNA Clone

or AA Sequence:

APAEPQPGGS QCVEHDCFAL YPGPATFLNA SQICDGLRGH LMTVRSSVAA DVISLLLNGD GGVGRRRLWI GLQLPPGCGD PKRLGPLRGF QWVTGDNNTS YSRWARLDLN GAPLCGPLCV

AVSAAEATVP SEPIWEEQQC EVKADGFLCE FHFPATCRPL AVEPGAAAAA VSITYGTPFA

ARGADFQALP VGSSAAVAPL GLQLMCTAPP GAVQGHWARE APGAWDCSVE NGGCEHACNA IPGAPRCQCP AGAALQADGR SCTASATQSC NDLCEHFCVP NPDQPGSYSC MCETGYRLAA DQHRCEDVDD CILEPSPCPQ RCVNTQGGFE CHCYPNYDLV DGECVEPVDP CFRANCEYQC QPLNQTSYLC VCAEGFAPIP HEPHRCQMFC NQTACPADCD PNTQASCECP EGYILDDGFI CTDIDECENG GFCSGVCHNL PGTFECICGP DSALARHIGT DCDSGKVDGG DSGSGEPPPS

PTPGSTLTPP A

Tag: Tag Free
Predicted MW: 51 kDa
Concentration: N/A

Purity: >98% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Lyophilized from a 0.2 μM filtered solution of 10 mM phosphate buffer, pH 7.5

Bioactivity: Measured by its ability to activate protein C induced cleavage of the chromogenic substrate,

BOC-Asp-Pro Arg-AMC in the presence of thrombin. The specific activity is greater than 500

pmoles/min/ug.

Endotoxin: Endotoxin level is < 0.1 ng/μg of protein (< 1 EU/μg)

Reconstitution Method: Restore in sterile water to a concentration of 0.1-1.0 mg/ml.

Storage: Store at -80°C.

Stability: Stable for at least 6 months from date of receipt under proper storage and handling

conditions.

RefSeq: NP 000352

Locus ID: 7056





Thrombomodulin (THBD) (NM_000361) Human Recombinant Protein - TP723443

UniProt ID: P07204

RefSeq Size: 4048

Cytogenetics: 20p11.21 RefSeq ORF: 1725

Synonyms: AHUS6; BDCA-3; BDCA3; CD141; THPH12; THRM; TM

Summary: The protein encoded by this intronless gene is an endothelial-specific type I membrane

receptor that binds thrombin. This binding results in the activation of protein C, which degrades clotting factors Va and VIIIa and reduces the amount of thrombin generated. Mutations in this gene are a cause of thromboembolic disease, also known as inherited

thrombophilia. [provided by RefSeq, Jul 2008]

Protein Families: Druggable Genome, Transmembrane
Protein Pathways: Complement and coagulation cascades