

Datasheet

TGM1 MaxPab rabbit polyclonal antibody (D01)

Catalog Number: H00007051-D01

Regulation Status: For research use only (RUO)

Product Description: Rabbit polyclonal antibody raised against a full-length human TGM1 protein.

Immunogen: TGM1 (NP_000350.1, 1 a.a. ~ 817 a.a) full-length human protein.

Sequence:

MMDGPRSDVGRWGGNPLQPPTTPSPEPEPEPDGRS
RRGGGRSFWARCCGCCSCRNAADDDWGPEPSDSR
GRGSSSGTRRPGSRGSDSRPVSRLSGVNAAGDGTI
REGMLVVNGVDLLSSRSDQNRREHHTDEYEYDELIVR
RGQPFHMLLLSRTYESSDRITLLELLIGNNPEVGKGT
VIIPVGKGGSGGWKAQVVKASGQNLNLRVHTSPNAIIG
KFQFTVRTQSDAGEFQLPFDPRNEIYILFNPWCPEDIV
YVDHEDWRQEYVLNESGRIYYGTEAQIGERTWNYGQ
FDHGVLDACLYILDRRGMPYGGRGDPVNVSRVISAMV
NSLDDNGVLIGNWSGDYSRGTNPVSAWVGSVEILLSYL
RTGYSVPYGGCWFVAGVTTTTLRCLGLATRTVTNFNS
AHDTDTSLTMDIYFDENMKPLEHLNHDSVWNFHVWN
DCWMKRPDLPSGFDGWQVVDATPQETSSGIFCCGPC
SVESIKNGLVYMKYDTPFIFAENVSDKVYVWQRQDDGS
FKIVYVEEKAIGTLIVTKAISSNMREDITYLYKHPEGSDA
ERKAVETAAAHGSKPNVYANRGAEDVAMQVEAQDA
VMGQDLMVSVMLINHSSSRRTVKLHLYLSVTFTYTGVS
GTIFKETKKEVELAPGASDRVTMPVAYKEYRPHLVDQ
GAMLLNVSGHVKESGQVLAKQHTFRLRTPDLSLTLLG
AAVVGQECEVQIVFKNPLPVTLTNVVFRLEGSGLQRP
KILNVGDIGGNETVTLRQSFVPRPGPRQLIASLDSPQ
LSQVHGVVIQVDVAPAPGDGGFFSDAGGDSHLGETIPM
ASRGA

Host: Rabbit

Reactivity: Human, Mouse

Applications: IP, WB-Ti, WB-Tr

(See our web site product page for detailed applications information)

Protocols: See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Storage Buffer: No additive

Storage Instruction: Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 7051

Gene Symbol: TGM1

Gene Alias: ICR2, KTG, LI, LI1, TGASE, TGK

Gene Summary: The protein encoded by this gene is a membrane protein that catalyzes the addition of an alkyl group from an alkylamine to a glutamine residue of a protein, forming an alkylglutamine in the protein. This protein alkylation leads to crosslinking of proteins and catenation of polyamines to proteins. This gene contains either one or two copies of a 22 nt repeat unit in its 3' UTR. Mutations in this gene have been associated with autosomal recessive lamellar ichthyosis (LI) and nonbullous congenital ichthyosiform erythroderma (NCIE). [provided by RefSeq]