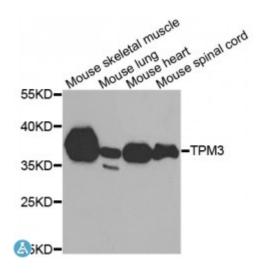


Anti-TPM3 Antibody



Description This gene encodes a member of the tropomyosin family of actin-binding

proteins. Tropomyosins are dimers of coiled-coil proteins that provide stability to actin filaments and regulate access of other actin-binding proteins. Mutations in this gene result in autosomal dominant nemaline myopathy and other muscle disorders. This locus is involved in translocations with other loci, including anaplastic lymphoma receptor tyrosine kinase (ALK) and neurotrophic tyrosine kinase receptor type 1 (NTRK1), which result in the formation of fusion proteins that act as oncogenes. There are numerous pseudogenes for this gene on different chromosomes. Alternative splicing results in multiple transcript variants.

Model STJ115518

Host Rabbit **Reactivity** Mouse

Applications IHC, WB

Immunogen Recombinant fusion protein containing a sequence corresponding to amino

acids 1-284 of human TPM3 (NP_689476.2).

Gene ID 7170

Gene Symbol TPM3

Dilution range WB 1:500 - 1:2000

IHC 1:50 - 1:200

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name Tropomyosin alpha-3 chain Gamma-tropomyosin Tropomyosin-3

Tropomyosin-5 hTM5

Molecular Weight 32.95 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Storage Instruction Store at -20C. Avoid freeze / thaw cycles.

Database Links HGNC:120120MIM:164970Reactome:R-HSA-390522

Alternative Names Tropomyosin alpha-3 chain Gamma-tropomyosin Tropomyosin-3

Tropomyosin-5 hTM5

Function Binds to actin filaments in muscle and non-muscle cells, Plays a central role,

in association with the troponin complex, in the calcium dependent regulation of vertebrate striated muscle contraction, Smooth muscle contraction is regulated by interaction with caldesmon, In non-muscle cells is implicated in

stabilizing cytoskeleton actin filaments,

Cellular Localization Cytoplasm, cytoskeleton

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