

Product datasheet for BA627

Cardiac Troponin I Human Protein

Product data:

Product Type: Native Proteins

Description: Cardiac Troponin I human protein, 10 μg

Species: Human

Protein Source: Cardiac muscle

Concentration: lot specific

Purity: >95% by SDS-PAGE

Buffer: State: Liquid

Buffer System: 7M Urea, 5 mM EDTA, 20 mM Tris

Preservative: 15 mM Mercaptoethanol

Preparation: Liquid Applications: ELISA.

Protein Description: Purified native Human Troponin I from Human heart

Note: Caution: Blood samples of the tissue donor for this product was tested and found negative

for HBsAg, HCV, syphillis and HIV-I and HIV-II antibodies. Nevertheless, all products from

human sources should be handled as potentially infectious.

Storage: Upon receipt, store undiluted (in aliquots) at -20°C.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

RefSeq: NP 000354

Locus ID: 7137

Cytogenetics: 19q13.42

Synonyms: TNNI3, TNNC1



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Summary: Troponin I (TnI), along with troponin T (TnT) and troponin C (TnC), is one of 3 subunits that

form the troponin complex of the thin filaments of striated muscle. Tnl is the inhibitory subunit; blocking actin-myosin interactions and thereby mediating striated muscle relaxation. The Tnl subfamily contains three genes: Tnl-skeletal-fast-twitch, Tnl-skeletal-slow-twitch, and Tnl-cardiac. This gene encodes the Tnl-cardiac protein and is exclusively expressed in cardiac muscle tissues. Mutations in this gene cause familial hypertrophic cardiomyopathy type 7 (CMH7) and familial restrictive cardiomyopathy (RCM). Troponin I is useful in making a diagnosis of heart failure, and of ischemic heart disease. An elevated level of troponin is also now used as indicator of acute myocardial injury in patients hospitalized with moderate/severe Coronavirus Disease 2019 (COVID-19). Such elevation has also been associated with higher risk of mortality in cardiovascular disease patients hospitalized due to

Protein Families: ELISA.

Protein Pathways: Cardiac muscle contraction, Dilated cardiomyopathy, Hypertrophic cardiomyopathy (HCM)

COVID-19. [provided by RefSeq, Aug 2020]