

Troponin T Core Structure and the Regulatory NH₂-Terminal Variable Region[†]Brandon J. Biesiadecki,^{‡,||} Stephen M. Chong,[§] Thomas M. Nosek,[‡] and Jian-Ping Jin^{*,‡,§}

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ABSTRACT: The conserved central and COOH-terminal regions of troponin T (TnT) interact with troponin C, troponin I, and tropomyosin to regulate striated muscle contraction. Phylogenetic data show that the NH₂-terminal region has evolved as an addition to the conserved core structure of TnT. This NH₂-terminal region does not bind other thin filament proteins, and its sequence is hypervariable between fiber type and developmental isoforms. Previous studies have demonstrated that NH₂-terminal modifications alter the COOH-terminal conformation of TnT and thin filament Ca²⁺-activation, yet the functional core structure of TnT and the mechanism of NH₂-terminal modulation are not well understood. To define the TnT core structure and investigate the regulatory role of the NH₂-terminal variable region, we investigated two classes of model TnT molecules: (1) NH₂-terminal truncated cardiac TnT and (2) chimera proteins consisting of an acidic or basic skeletal muscle TnT NH₂-terminus spliced to the cardiac TnT core. Deletion of the TnT hypervariable NH₂-terminus preserved binding to troponin I and tropomyosin and sustained cardiac muscle contraction in the heart of transgenic mice. Further deletion of the conserved central region diminished binding to tropomyosin. The reintroduction of differently charged NH₂-terminal domains in the chimeric molecules produced long-range conformational changes in the central and COOH-terminal regions to alter troponin I and tropomyosin binding. Similar NH₂-terminal charge effects are demonstrated in naturally occurring cardiac TnT isoforms, indicating a physiological significance. These results suggest that the hypervariable NH₂-terminal region modulates the conformation and function of the TnT core structure to fine-tune muscle contractility.

The contraction of striated muscle is regulated by Ca²⁺ via the troponin (Tn¹) complex that is composed of three subunits: troponin C (TnC, the calcium binding subunit), troponin I (TnI, the inhibitory subunit), and troponin T (TnT, the tropomyosin (Tm)-binding subunit). During the activation of contraction, increased cytosolic Ca²⁺ binds to TnC to induce a series of conformational changes in Tn and Tm that “opens” the muscle thin filament to allow the interaction of myosin with actin and cross-bridge cycling (1). Through direct binding to TnC, TnI, Tm, and actin, TnT anchors the

Tn complex to the thin filament and transmits the Ca²⁺-induced conformational changes from Tn to Tm (2–4).

Three homologous TnT genes have evolved in higher vertebrates to encode fast skeletal, slow skeletal, and cardiac muscle TnT isoforms (5–8). These TnT isoforms have highly conserved amino acid sequences in the COOH-terminal region containing binding sites for TnC, TnI, and Tm (9–11). The central region of TnT is also highly conserved and contains a second Tm-binding site (2). The NH₂-terminal region of the TnT polypeptide chain does not contain any known protein binding sites, and phylogenetic data show that this region has evolved as an addition to the conserved core structure of TnT (12). In contrast to the central and COOH-terminal regions, the NH₂-terminal region of TnT is highly variable among the three muscle fiber type isoforms (13) and is alternatively spliced for additional length and charge variations (2, 14).

The crystal structures of the partial cardiac and skeletal muscle Tn complex have revealed the molecular interactions that occur in the globular domain of Tn (15, 16). In agreement with protein binding data (9–11), the high-resolution Tn structures demonstrate interactions between the TnT COOH-terminal domain and TnI and TnC. The partial crystal structure of Tn does not include the central region of TnT or the NH₂-terminal variable region. This is consistent with the previous observation that TnT is an elongated molecule with its central and NH₂-terminal regions forming the asymmetric tail portion of the Tn complex (17–19).

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¹ Abbreviations: Acidic TnT, chimeric TnT containing an acidic NH₂-terminal segment and the McTnT core; Basic TnT, chimeric TnT containing a basic NH₂-terminal segment and the McTnT core; BSA, bovine serum albumin; ELISA, enzyme-linked immunosorbent assay; McTnT, mouse cardiac TnT; ND₇₂, deletion of the NH₂-terminal 71 amino acids; ND₉₂, deletion of the NH₂-terminal 91 amino acids; PCR, polymerase chain reaction; pI, isoelectric point; SDS–PAGE, SDS–polyacrylamide gel electrophoresis; TBS, Tris-buffered saline; Tg, transgenic mouse; Emb, embryonic; Tm, tropomyosin; Tn, troponin; TnC, troponin C; TnI, troponin I; TnT, troponin T.

Phylogenetic lineage of TnT shows a short-to-long, simple-to-complex evolution of the NH₂-terminal variable region (12). Fast skeletal muscle TnT with the NH₂-terminal variable region removed retains activity in the calcium activation of actomyosin ATPase (20–22). Similarly, the exchange of NH₂-terminal truncated cardiac TnT into myofibrils retains calcium activated force development (23). However, removing the NH₂-terminal region from both skeletal and cardiac TnT decreased the maximal force (20, 23) and increased the binding affinity to Tm compared to that of the intact TnT molecule (20, 23, 24). Furthermore, suggesting a regulatory function of the TnT NH₂-terminal variation, alternative splicing within the NH₂-terminal region is responsible for the developmental isoform switches of both cardiac and fast skeletal muscle TnT (14). We have previously demonstrated that structural alterations within the TnT NH₂-terminal region alters the molecular conformation and flexibility of the central and COOH-terminal regions of TnT to affect the binding affinity for TnI, TnC, and Tm (19, 25, 26), Ca²⁺ activation of ATPase activity (27), shortening velocity of isolated cardiomyocytes (28), and Ca²⁺ sensitivity of force development (29). Its variable structure and the resulting functional effects suggest a hypothesis that the NH₂-terminal region of TnT is a modulatory addition to the TnT central and COOH-terminal core structure, which functions to fine-tune muscle contractility.

There are three variables in the TnT NH₂-terminal structure: amino acid sequence, length (mass), and charge. The NH₂-terminal sequence of TnT isoforms shows very little conservation and, therefore, provides little information for a functional correlation. Furthermore, the size of vertebrate TnT varies by as many as 70 amino acid residues, almost completely resulting from variations in the length of the NH₂-terminal region. A striking observation is that although the sizes of embryonic and adult fast skeletal TnT isoforms overlap, they fall into two distinct and non-overlapping acidic and basic charge classes, resulting from developmentally regulated alternative splicing of six exons that encode the NH₂-terminal region (14, 26). Therefore, charge variation may be a determinant for the TnT hypervariable NH₂-terminal region to modulate the conformation and function of the core structure.

In the present study, we investigated representative model proteins to define the TnT core structure and investigate the regulatory effects of NH₂-terminal charge. The results demonstrate preserved biochemical and physiological activity of the TnT core corresponding to the conserved central and COOH-terminal regions. The addition of differently charged exogenous NH₂-terminal segments alters the molecular conformation of the TnT core structure to affect the interactions with TnI and Tm. Similar NH₂-terminal effects are demonstrated in naturally occurring cardiac TnT isoforms, suggesting that the NH₂-terminal variable region plays a role in modulating the conformation and function of the TnT core to fine-tune muscle contractility.

EXPERIMENTAL PROCEDURES

Construction of the cDNAs Encoding NH₂-Terminal Truncated and Chimeric TnT Proteins. cDNAs encoding two cTnT proteins with different NH₂-terminal truncations were designed to remove either only the NH₂-terminal variable

region (McTnT-ND₇₂) or extend the deletion to include a portion of the conserved central region (McTnT-ND₉₂). The two cDNA coding templates were constructed using polymerase chain reaction (PCR) to place a methionine codon prior to leucine 72 (McTnT-ND₇₂) or aspartic acid 92 (McTnT-ND₉₂) of the adult mouse cardiac TnT-4 ((30); Figure 1). The truncated cDNA was amplified from an intact adult mouse cardiac TnT cDNA cloned in pBluescript SK(-) plasmid (31) using mutagenesis primers McTnT-ND-F₇₂ containing an ATG codon prior to the codon for Leu₇₂ and an *NdeI* restriction endonuclease cloning site (5'-AGCCCCATATGCTCTTCATGCCCAACTT-3') or McTnT-ND-F₉₂ containing an ATG codon prior to the codon for Asp₉₂ and an *NdeI* cloning site (5'-GACCATATGGACATCCACAGGAA-3') paired with a vector primer next to the cDNA 3' end. The resultant PCR fragments were modified by *NdeI* and *XhoI* at the ends and cloned into the pAED4 prokaryotic expression plasmid (32). Purified plasmid DNA was verified by dideoxy chain termination sequencing.

The most variable NH₂-terminal charge exists among alternative spliced fast skeletal muscle TnT isoforms (33, 34). The isoelectric point (pI) of embryonic versus adult mouse fast skeletal muscle TnT isoforms ranges from 5.04 to 9.19 (33). To build model molecules for testing the NH₂-terminal charge effect, two recombinant chimera cDNAs were constructed by joining the 5' variable region of embryonic or adult chicken fast skeletal muscle TnT cDNA (fsTnT-3 or fsTnT-4, respectively (35)) to cDNA encoding the conserved core structure of mouse cardiac TnT. As illustrated in Figure 1, the Acidic chimeric TnT consists of chicken fast skeletal TnT exons 2–8 spliced to mouse cardiac TnT exons 9–17. The Basic chimeric TnT consists of chicken fast skeletal muscle TnT exons 2, 3, 5, 6, and 7 spliced to mouse cardiac TnT exons 9–17.

The chimeric cDNA was constructed using a three-step recombinant PCR method. First, the chicken NH₂-terminal region containing exons 2–8 of TnT-3 or exons 2–3 and 5–7 of TnT-4 was amplified from intact cDNA templates cloned in M13mp18 phagemid (kindly provided by Dr. Larry Smillie, University of Alberta (35)). The PCR used primer CfTnTe7R (5'-GGGTTTGGTCTCCTCTGGAGG-3') with the 3' region complimentary to exon 7 of chicken fsTnT and the 5' region complimentary to exon 9 of mouse cardiac TnT paired with a vector primer M13R. In a separate PCR reaction, the exon 9–17 region of mouse cardiac TnT cDNA was amplified from intact TnT-4 cDNA cloned in pBluescript SK(-) (31) using the primer McTnTe7F (5'-AGAGGAGACCAAACCAAGCCCA-3') with the 3' region complimentary to exon 9 of mouse cardiac TnT cDNA and the 5' region complimentary to chicken fsTnT paired with a vector primer T7. The first PCR products were purified, and either the chicken fsTnT-3 or the chicken fsTnT-4 fragment was mixed with the mouse cardiac TnT fragment and hybrid cDNA templates produced by 5 cycles of denaturation-annealing and chain elongation under standard PCR conditions. Recombinant full length cDNAs were then amplified by PCR using the flanking M13R and T7 primers and cloned into the pBluescript KS(+) vector. Plasmid DNA was purified and sequenced to verify the construction of cDNA chimeras before subcloning the insert into the pAED4 expression vector as described above.

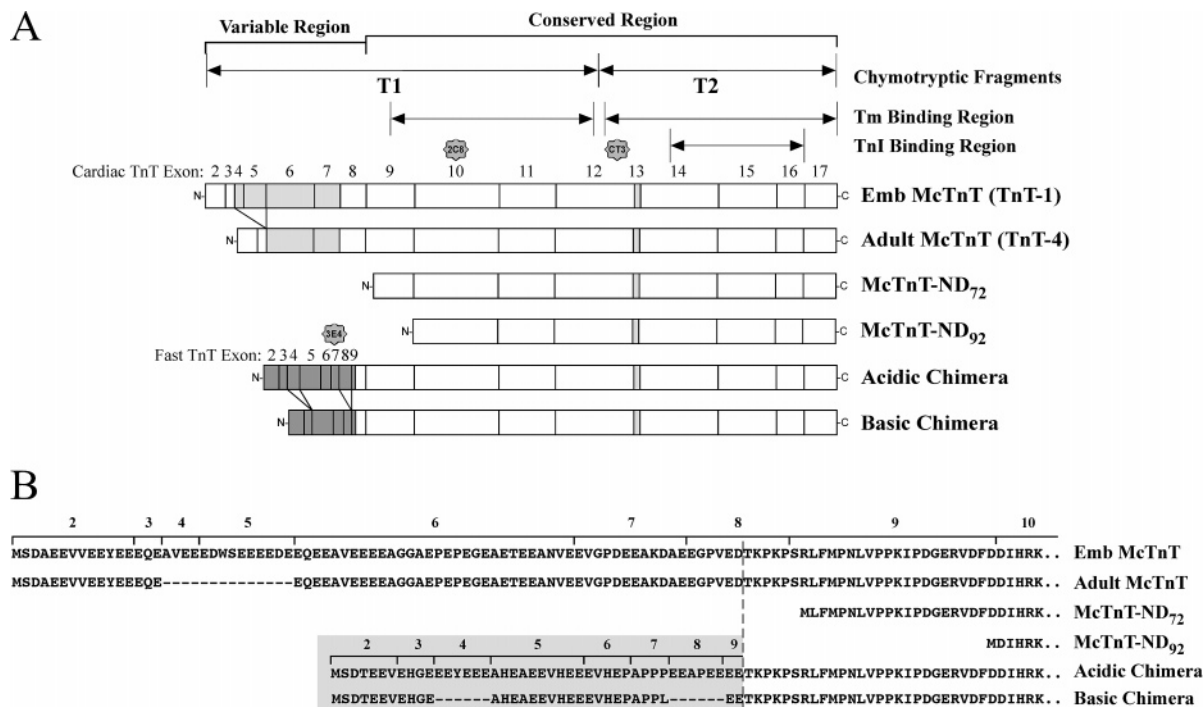


FIGURE 1: TnT molecules studied. (A) Linear maps show the two naturally occurring mouse cardiac TnT isoforms and the engineered model TnT molecules used in this study. The adult mouse cardiac TnT (TnT-4) differs from the embryonic cardiac TnT (TnT-1) by the exclusion of exons 4 and 5 from the NH₂-terminal variable region. The truncated mouse cardiac TnT molecules deleted only the NH₂-terminal variable region encoded by exons 2–8 (McTnT-ND₇₂) or further including a conserved segment encoded by exon 9 (McTnT-ND₉₂). The two chimeric TnT molecules are identical to the mouse cardiac TnT in their central and COOH-terminal regions (exons 9–17) but contain the NH₂-terminal domain of an acidic chicken fast skeletal muscle TnT (encoded by exons 2–8 from chicken fast TnT-3) (Acidic chimera) or a basic chicken fast skeletal muscle TnT (encoded by exons 2–3 and 5–7 from chicken fast TnT-4) (Basic chimera). The TnI binding region, tropomyosin binding regions, T1 and T2 chymotryptic fragments, and the mAb 3E4, CT3, and 2C8 epitopes are designated. The light gray exons indicate alternately spliced exons in cardiac TnT; dark gray exons indicate chicken fast skeletal muscle TnT exons. (B) The NH₂-terminal amino acid sequences of the six TnT molecules are aligned to demonstrate structural relationships. Residues that originated from chicken fast skeletal muscle TnT are highlighted in gray.

Two alternatively spliced cardiac TnT isoforms with the largest possible natural charge difference (the TnT-4 adult isoform and the TnT-1 embryonic isoform (31)) were chosen as a pair to study the naturally occurring NH₂-terminal charge effect. These previously cloned adult and embryonic mouse cardiac TnT cDNAs (31) were subcloned into the pAED4 vector as described above.

SDS–Polyacrylamide Gel Electrophoresis (SDS–PAGE) and Western Blotting. Protein samples were mixed with 3× SDS–PAGE sample buffer to a final concentration of 2% SDS, 0.3% bromophenol blue, 10% glycerol, and 50 mM Tris–HCl at pH 6.8, heated to 80 °C for 5 min and clarified by centrifugation. The separation of the McTnT-ND proteins was carried out by electrophoresis on a 14% gel with an acrylamide/bisacrylamide ratio of 120:1 in the Tris-tricine buffer system as previously described (32). All other protein bands were resolved on a 14% gel with an acrylamide/bisacrylamide ratio of 180:1 in the Laemmli buffer system. Resulting gels were stained with Coomassie Blue R250 to reveal the protein bands and duplicate gels transferred to a nitrocellulose membrane. The membrane was blocked with bovine serum albumin (BSA) and incubated with anti-cardiac TnT mAbs CT3 (25), 2C8 (36), or an anti-chicken fast skeletal TnT mAb 3E4 (26). Following washes with Tris-buffered saline (TBS, 136.9 mM NaCl, 2.7 mM KCl, and 25 mM Tris–HCl at pH 7.4) containing 0.5% Triton X-100 and 0.05% SDS, the membranes were incubated with alkaline phosphatase-labeled anti-mouse IgG second antibody (Sigma Chemical Co., St. Louis, MO), washed as above, and

developed with 5-bromo-4-chloro-3-indolylphosphate and nitro blue tetrazolium substrate solution as previously described (26).

Bacterial Expression and Purification of TnT. The six TnT cDNA constructs described above were expressed in BL21-(DE3)pLysS *E. coli*. Freshly transformed bacterial cells were cultured in liquid media (16 g/L Tryptone, 10 g/L yeast extract, 5 g/L NaCl, and 1.32 g/L Na₂HPO₄ at pH 7.3) containing 100 mg/L ampicillin and 25 mg/L chloramphenicol at 37 °C with vigorous shaking and induced with 0.4 mM isopropyl-1-thiol- β -D-glactoside at mid-log phase. After 3 additional hours of culture, the bacterial cells were harvested by centrifugation at 4 °C.

McTnT-ND₇₂ and McTnT-ND₉₂ proteins were purified by identical procedures. The bacterial pellet was suspended in 2.5 mM EDTA and 50 mM Tris–HCl at pH 8.0 and lysed by three passes through a French Press cell. The bacterial lysate was clarified by centrifugation and precipitated with ammonium sulfate to obtain the 0–35% saturation fraction. Following dialysis against 0.1 mM EDTA containing 6 mM β -mercaptoethanol, the fraction was brought to 6 M urea, 0.1 mM EDTA, 6 mM β -mercaptoethanol, and 20 mM sodium acetate at pH 6.0 and fractionated by chromatography on a CM52 cation-exchange column equilibrated in the same buffer. The column was eluted by a 0–500 mM linear KCl gradient and the protein peaks analyzed by SDS–PAGE. Fractions containing the NH₂-terminal truncated cardiac TnT were dialyzed against 0.1 mM EDTA containing 6 mM β -mercaptoethanol, concentrated by lyophilization, redis-

solved in 6 M urea, 0.5 M KCl, 0.1 mM EDTA, 6 mM β -mercaptoethanol, and 10 mM imidazole-HCl at pH 7.0 and fractionated by G75 gel filtration chromatography in the same buffer. Protein peaks were analyzed by SDS-PAGE, and the fractions containing pure McTnT-ND₇₂ or McTnT-ND₉₂ protein were dialyzed against 0.1% formic acid and lyophilized. All purification steps were carried out at 4 °C.

The Basic chimeric TnT was purified as described above for the NH₂-terminal truncated cardiac TnT with the following modifications. The bacterial lysate was fractionated by (NH₄)₂SO₄ to collect the 30–50% saturation precipitation. After dialysis, the proteins were fractionated by chromatography on a CM52 column at pH 4.8 (20 mM sodium acetate buffer). The column was eluted by a 0–500 mM linear KCl gradient and the protein peaks analyzed by SDS-PAGE. Fractions containing the Basic chimeric TnT were further purified by G75 chromatography.

The Acidic chimeric TnT was similarly purified except that the 30–50% saturation (NH₄)₂SO₄ fraction was fractionated by chromatography on a DE-52 anion-exchange column at pH 8.5 (20 mM Tris-HCl buffer) with a 0–300 mM linear KCl gradient. The eluted protein peaks were analyzed by SDS-PAGE, and fractions containing the Acidic chimeric TnT were further purified by G75 chromatography.

The embryonic and adult mouse cardiac TnTs were purified by identical procedures. The bacterial pellet was made into acetone powder and extracted by stirring with a buffer containing 6 M urea, 0.1 mM EDTA, 6 mM β -mercaptoethanol, and 20 mM Tris-HCl at pH 8.0 at 4 °C for 2 h. The protein extracts were fractionated by ammonium sulfate precipitation between 30 and 50% saturation. After dialysis, the proteins were fractionated on a DE52 column at pH 8.0 (20 mM Tris-HCl buffer) followed by G75 chromatography as described above.

Purification of Other Myofibrillar Proteins. Bovine cardiac TnI was purified from adult ventricular muscle as previously described (27). Rabbit α -tropomyosin was purified from adult cardiac muscle as previously described (37).

mAb Epitope Analysis. The binding affinity between an antibody and its antigenic epitope depends on a 3D structural fit. Enzyme-linked immunosorbent assay (ELISA) epitope analysis (26) was employed to examine conformational differences between the TnT molecules. mAb CT3 against an epitope in the central domain of cardiac TnT (25) and mAb 3E4 against an NH₂-terminal epitope on chicken fast skeletal TnT (26) (Figure 1) were used to monitor conformational changes that alter the antibody binding affinity. The Acidic or Basic chimeric TnT and mouse embryonic or adult cardiac TnT were dissolved in Buffer A (0.1 M KCl, 3 mM MgCl₂, and 10 mM PIPES at pH 7.0) and the protein concentration determined by UV absorbance according to the following formula: (E₂₈₀) (mm⁻¹ cm⁻¹) = 1.4 (# tyrosine) + 5.6 (# tryptophan). Five μ g/mL protein solutions were used at 100 μ L/well to coat microtitering plates by incubation at 4 °C overnight. After removing unbound TnT by washing once with Buffer A containing 0.05% Tween-20 (Buffer T) and blocking the plates with 150 μ L/well Buffer T containing 1% BSA at room temperature for 1.5 h, immobilized TnT was incubated with 100 μ L/well serial dilutions of CT3 or 3E4 mAb in Buffer T containing 0.1% BSA at room temperature for 2 h. Following three washes with Buffer T in a 10 min period to remove the unbound first antibody,

the plates were incubated with 100 μ L/well horseradish peroxidase-conjugated anti-mouse immunoglobulin second antibody (Sigma, St. Louis, MO) in Buffer T containing 0.1% BSA at room temperature for 1 h. Unbound second antibody was removed by three washes as described above. The binding of mAbs to a particular TnT construct was detected by H₂O₂/2,2'-azinobis-(3-ethylbenzthiazolinesulfonic acid) substrate reaction. The enzymatic reaction in each assay well was monitored at a series of time points by an automated microplate reader (BioRad Benchmark, Hercules, CA). A_{405nm} values in the linear course of the color development were used to plot the antibody titration curves for quantifying the binding affinity of the mAbs to the TnT epitopes. All experiments were done in triplicate.

Microtiter Plate Protein Binding Assays. An ELISA-based solid-phase protein binding assay (26) was used to investigate the interactions of the TnT variants with TnI and Tm. Similar to that described above for the epitope analysis, purified TnT or BSA control was dissolved in Buffer A at 5 μ g/mL and 100 μ L/well used to coat microtiter plates by incubation at 4 °C overnight. Unbound TnT was removed by a wash with Buffer T and the plate blocked with 150 μ L/well with Buffer T containing 1% BSA at room temperature for 1.5 h. The plates were then washed three times with Buffer T over a 10 min period and incubated with 100 μ L/well serial dilutions of bovine cardiac TnI or rabbit α -Tm in Buffer T containing 0.1% BSA. Following incubation at room temperature for 2 h, the plates were washed three times with Buffer T over a 10 min period. The bound TnI or Tm was quantified via an anti-TnI mAb TnI-1 (38) or an anti-Tm mAb CH1 (39) (a gift from Professor Jim J.-C. Lin, University of Iowa) followed by the standard ELISA procedure as described above. The A_{405nm} values in the linear course of color development were used to construct protein binding curves for the quantification of binding affinity between the TnT variants to Tm and TnI under various washing conditions. All experiments were done in triplicate.

Immunoaffinity Chromatographic Isolation of the Tn Complex from Mouse Cardiac Muscle. As described previously (40), McTnT-ND₇₂ transgenic mice were constructed on a C57BL/6 background using mouse cardiac α -myosin heavy chain (α -MHC) promoter ((41); generously provided by Dr. Jeffrey Robbins, University of Cincinnati) to direct a heart-specific expression of mouse cardiac TnT cDNA encoding cardiac TnT with the NH₂-terminal variable region deleted to investigate the functional preservation in the TnT core structure. The pronucleus injection was performed at the Transgenic Core Facility at Case Western University, and mouse line 14 was used in the present study. The expression of the exogenous cardiac TnT in the transgenic mouse hearts was verified by Western blots using CT3 mAb. Another transgenic mouse line with α -MHC promoter-directed heart-specific expression of the embryonic isoform of cardiac TnT in the adult cardiac muscle developed previously (28) was also used in the present study.

The troponin complex was isolated from transgenic mouse hearts by immunoaffinity chromatography in manner similar to that previously described (42). The troponin complex was isolated using mAb TnI-1 against an epitope at the extreme COOH-terminus of TnI (38), which is exposed in the Tn complex, providing a specific handle for affinity purification. TnI-1 mAb (IgG1) was purified from hybridoma ascites fluid

using a Protein G column (Amersham-GE Healthcare Bio-Sciences Corp., Piscataway, NJ) and coupled to CNBr-activated Sepharose 4B (Amersham-GE Healthcare Bio-Sciences Corp., Piscataway, NJ) according to the manufacturer's protocols. The immunoaffinity isolation of the Tn complex was then carried out using Sepharose 4B-TnI-1 mAb affinity column chromatography. Transgenic mouse ventricular muscle was minced into 1–2 mm³ pieces and extracted by 20 volumes (w/v) of Guba–Straub solution containing 300 mM KCl, 100 mM K₂HPO₄, 50 mM KH₂PO₄, 2.5 mM MgCl₂, 1 mM EGTA, and 0.1 mM PMSF at pH 6.5 on ice for 15 min. After centrifugation at 16,000g at 4 °C for 20 min, the supernatant containing mainly myosin and cytosolic proteins was removed. The pellet was extracted next in 20 volumes (w/v) of 1 M KCl, 10 mM Tris-HCl at pH 8.0, and 0.1 mM PMSF by stirring on ice for 30 min. After centrifugation as described above, the extract was diluted 5-fold in TBS and loaded onto a 0.5 mL TnI-1 mAb affinity column equilibrated in TBS. The column was washed with TBS and then eluted with 50 mM glycine-HCl at pH 2.7. Then, 0.5 mL fractions were collected into tubes containing 0.1 mL of neutralizing buffer (1 M Tris-HCl, 1.5 M NaCl, and 1 mM EDTA at pH 8.0) and analyzed by SDS–PAGE and Western blotting to identify the Tn peak as described above.

Incorporation of the NH₂-terminal truncated cardiac TnT in the transgenic mouse cardiac Tn complex was compared with that of the intact adult and embryonic cardiac TnT by densitometric quantification of the SDS gels and Western blots as described previously (28).

Data Analysis. The DNA and protein sequence analyses were done using DNASTar computer programs. Statistical analysis of the ELISA antibody epitope analysis, protein binding assays, and gel densitometry was done by Student's *t* test. All values are presented as the mean \pm SD.

RESULTS

Preserved Activity of the Conserved Core Structure of TnT. To identify the TnT core structure required to maintain protein-binding activity, we constructed two truncated cardiac TnT molecules with progressive NH₂-terminal deletions. The deletion of the first 71 residues in the McTnT-ND₇₂ molecule preserves the central and COOH-terminal conserved regions, whereas the deletion of the first 91 residues in the McTnT-ND₉₂ molecule results in a partial removal of the central conserved region (Figure 1). The physical properties of the NH₂-terminal truncated cardiac TnTs and that of the intact adult mouse cardiac TnT (TnT-4) are summarized in Table 1. The large-scale expression of the McTnT-ND₇₂, McTnT-ND₉₂, and adult mouse cardiac TnT in *E. coli* and the effective purification methods provided proteins for structural and functional characterizations (Figure 2A). The authenticity of bacterial expressed McTnT-ND proteins was confirmed by Western blotting.

TnT contains two Tm binding sites, one is located in the COOH-terminal region and the other in the central region. To investigate the binding of the NH₂-terminal truncated TnT constructs to Tm, we carried out ELISA solid-phase protein binding experiments. The results in Figure 3 demonstrate the binding affinity of McTnT-ND₇₂ to Tm was not significantly affected compared to that of the intact mouse cardiac TnT, but McTnT-ND₉₂ showed a decreased Tm binding

Table 1: Physical Properties of the TnT Proteins Studied^a

| TnT | number of amino acids | molecular weight | pI | NH ₂ -terminal charge at pH 7.0 |
|------------------------|-----------------------|------------------|-------|--|
| McTnT-ND ₇₂ | 221 | 26,859 | 9.87 | N/A |
| McTnT-ND ₉₂ | 201 | 24,578 | 10.00 | N/A |
| Emb McTnT (TnT-1) | 305 | 36,253 | 4.87 | –42 |
| adult McTnT (TnT-4) | 291 | 34,546 | 5.19 | –32 |
| Acidic model TnT | 271 | 32,634 | 6.33 | –19 |
| Basic model TnT | 259 | 31,157 | 8.99 | –10 |
| chicken fast TnT3 | 263 | 31,141 | 6.75 | –19 |
| chicken fast TnT4 | 251 | 29,647 | 9.20 | –10 |

^a The physical properties of the NH₂-terminal deleted mouse cardiac TnTs (McTnT-ND₇₂ and McTnT-ND₉₂), embryonic (Emb McTnT, TnT-1), and adult (Adult McTnT, TnT-4) mouse cardiac TnTs (31), Acidic and Basic model TnT chimeras, and chicken fast skeletal muscle TnT3 and TnT4 (accession number J04198) were calculated from amino acid sequences.

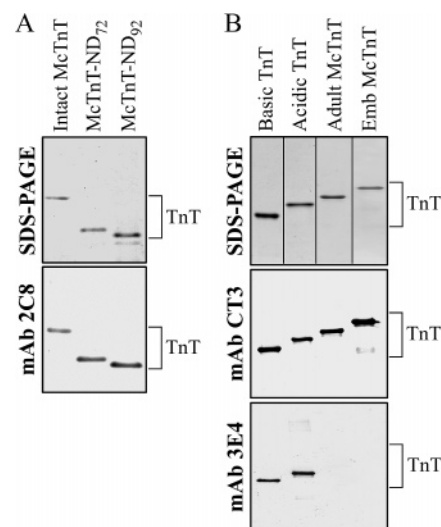


FIGURE 2: TnT Protein Preparations. Recombinant TnT proteins were expressed in *E. coli* and purified as shown by SDS–PAGE gels. (A) NH₂-terminal truncated TnT (McTnT-ND₇₂ and McTnT-ND₉₂) were analyzed along with the intact adult mouse cardiac TnT-4 (Intact McTnT). The Western blot demonstrates that the truncated TnT proteins are positive to mAb 2C8, which recognizes an epitope in the exon 10 region (Figure 1). (B) The Basic and Acidic chimeric TnTs were analyzed along with the adult mouse cardiac TnT-4 (Adult McTnT) and embryonic mouse cardiac TnT-1 (Emb McTnT). The Western blot demonstrates that the chimeric TnTs, similar to the native cardiac TnTs, are positive to mAb CT3, recognizing a central cardiac TnT epitope. The chimeric TnTs are also positive to mAb 3E4, which recognizes an epitope within exon 7 of the chicken fast skeletal muscle TnT. The results verified the construction of the NH₂-terminal deleted and recombinant chimeric TnT molecules.

affinity (the Tm concentration required for 50% maximum binding was as follows: intact mouse cardiac = 19.4 ± 2.2 nM, McTnT-ND₇₂ = 20.7 ± 3.1 nM, and McTnT-ND₉₂ = 39.4 ± 8.3 nM, $P < 0.001$). At concentrations of Tm that render saturated binding to comparable amounts of TnT coated on the plate, the maximal binding of Tm to McTnT-ND₇₂ was slightly increased (0.423 ± 0.001 , $P < 0.05$), whereas that to McTnT-ND₉₂ decreased (0.383 ± 0.008 , $P < 0.01$) when compared to that of the intact McTnT control (0.416 ± 0.003) (Figure 3, inset). The results showed that deletion of the entire variable region in McTnT-ND₇₂ retained both Tm binding sites intact, whereas further deletion of 20 residues from the central conserved region in McTnT-ND₉₂

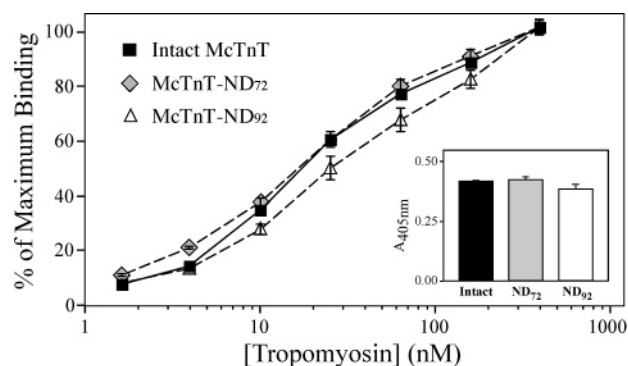


FIGURE 3: Selective deletion of the NH₂-terminal variable region from TnT does not abolish binding to Tm. Solid-phase protein binding curves of intact mouse cardiac TnT, McTnT-ND₇₂, and McTnT-ND₉₂ to Tm demonstrate the effect of TnT NH₂-terminal truncations. The binding affinity of McTnT-ND₇₂ to Tm was similar to that of the intact McTnT, whereas the Tm binding affinity of McTnT-ND₉₂ was decreased ($P < 0.001$). The maximal binding of McTnT-ND₇₂ to Tm was slightly increased, whereas that of McTnT-ND₉₂ to Tm was decreased (inset). In contrast to the preserved activity of McTnT-ND₇₂ with only the NH₂-terminal variable region deleted, the decreased binding of McTnT-ND₉₂ to Tm indicates that deletion into the conserved region damages the TnT core structure by disrupting the central Tm binding site (Figure 1).

resulted in decreased Tm binding, indicating a destruction of the TnT core structure.

Physical Properties of the Acidic and Basic Chimeric TnT Model Molecules. To evaluate the effect of NH₂-terminal charge on the molecular conformation and protein binding activity of TnT, we constructed two chimeric model TnT molecules consisting of the NH₂-terminal variable region of naturally occurring acidic (Acidic TnT) or basic (Basic TnT) chicken fast skeletal muscle TnT and the conserved central and COOH-terminal core regions of mouse cardiac TnT (Figure 1). The anticipated sequence combinations of the chimeric TnT were confirmed by Western blotting using mAb 3E4, specific for an epitope in the NH₂-terminal domain of chicken fast TnT, and mAb CT3, against an epitope in the central region of cardiac TnT (Figure 2B). The physical properties of the chimeric model TnT molecules along with that of the chicken fast skeletal muscle TnT-3, TnT-4, and adult mouse cardiac TnT-4 are summarized in Table 1. In the presence of an identical central and COOH-terminal core structure, the NH₂-terminal charge difference of 9 between the Acidic and Basic chimera produced an overall pI difference of 2.66, reproducing the NH₂-terminal charge effect seen in the original chicken fast TnT isoforms (Table 1). These designs are consistent with the proposed universal role of the NH₂-terminal variable region in regulating the overall charge of TnT.

NH₂-Terminal Charge Alters the Core Conformation in TnT Chimeras. The results of the ELISA epitope analysis demonstrated the effect of NH₂-terminal charge variation in Basic and Acidic chimeric TnT on molecular conformation. Figure 4A shows that the Basic TnT exhibited a binding affinity to mAb 3E4 higher than that observed for the Acidic TnT chimera. The mAb 3E4 recognizes an epitope encoded by exon 7 of chicken skeletal muscle TnT (Figure 1A). The different binding affinities of mAb 3E4 to Basic and Acidic chimeric TnTs demonstrate that the inclusion or exclusion of exon 4 and 8 segments affects the local molecular

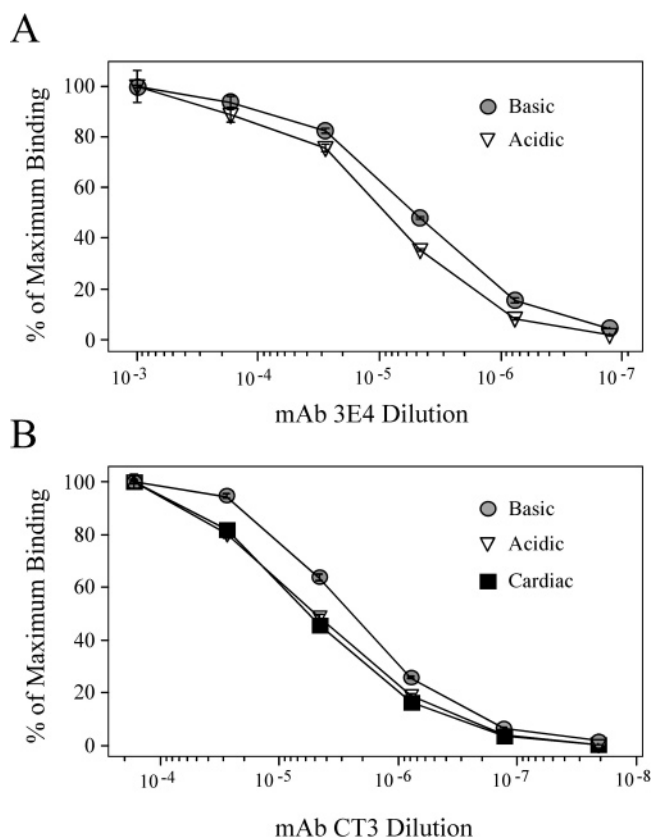


FIGURE 4: Alteration of NH₂-terminal charge affects the nearby and distal conformation in chimeric TnT model molecules. (A) The ELISA epitope affinity titration curves of mAb 3E4 demonstrate that the Acidic and Basic chimeric TnTs exhibit a conformational difference at the 3E4 epitope near the NH₂-terminal variation site, as defined by the difference in antibody concentrations required for 50% maximal binding. (B) The titration curves of mAb CT3 demonstrate that the Acidic and Basic chimeric TnTs and the native adult mouse cardiac TnT exhibit different conformations at the CT3 epitope in the central region (Figure 1A), indicating a remote conformational effect of the NH₂-terminal variation.

conformation in the NH₂-terminal domain. The Acidic chimeric TnT further exhibited a lower binding affinity to mAb CT3 when compared to that of the Basic chimeric TnT (Figure 4B). The CT3 mAb recognizes an epitope in the central region of the mouse cardiac TnT core removed from the altered NH₂-terminal region (Figure 1A). This result indicates a long-range conformational effect resulting from NH₂-terminal TnT charge variation. The adult mouse cardiac TnT, with an NH₂-terminus slightly more negatively charged than that of the Acidic chimeric TnT, confirmed the trend of lower binding affinity to mAb CT3 (Figure 4B). These results demonstrate that changes in the NH₂-terminal charge not only affect the local conformation but also have long-range conformational effects on the core structure of TnT.

NH₂-Terminal Charge Variation in TnT Model Molecules Affects Binding to TnI and Tm. To investigate the functional significance of the long-range conformational effects of NH₂-terminal charge variation on the TnT core structure, ELISA protein binding experiments showed that the Basic chimera exhibits a higher binding affinity to TnI compared to that of the Acidic chimera (Figure 5A), the TnI concentrations required for 50% maximum binding were 7.0 ± 0.9 nM and 13.7 ± 0.8 nM, respectively; $P < 0.05$). Consistent with this trend, the binding affinity of the more acidic adult mouse cardiac TnT to TnI was even lower than that of Acidic

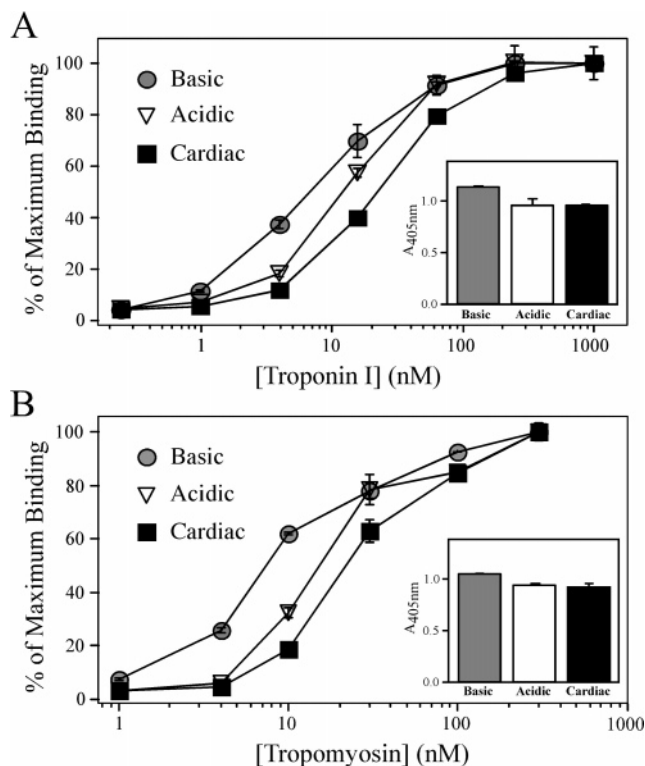


FIGURE 5: NH₂-terminal charge affects the binding of chimeric TnTs to TnI and Tm. (A) Solid-phase protein binding curves demonstrate that the Basic TnT bound with higher affinity to TnI than to Acidic TnT ($P < 0.05$). Both Basic and Acidic chimeric TnTs exhibited higher binding affinities to TnI than to the more acidic mouse cardiac TnT ($P < 0.05$). The A_{405nm} values in the inset show that the Basic chimeric TnT also had a higher level of maximal binding to TnI compared to those of the Acidic chimeric TnT and mouse cardiac TnT ($P < 0.05$). (B) Solid-phase protein binding curves demonstrate that the Basic chimeric TnT exhibited a higher binding affinity to Tm than to Acidic chimeric TnT ($P < 0.05$), whereas the more acidic mouse cardiac TnT had the lowest binding affinity as defined by the concentration of Tm required for 50% maximum binding ($P < 0.05$). The A_{405nm} values in the inset show that the Basic TnT exhibits a higher level of maximal binding to Tm compared to the bindings of Acidic TnT and mouse cardiac TnT ($P < 0.05$).

chimeric TnT. (The TnI concentration required for 50% maximum binding was 23.3 ± 1.4 nM; $P < 0.05$.) At TnI concentrations that render saturated binding to comparable amounts of TnT coated on the microtiter plate, the maximal binding of TnI to Basic chimeric TnT was also significantly greater than that of the Acidic chimera and the adult mouse cardiac TnT (Figure 5A, inset; A_{405nm} values are 1.13 ± 0.014 , 0.96 ± 0.060 , and 0.955 ± 0.010 , respectively; $P < 0.05$), indicating a stronger coupling strength that may affect the function of the Tn complex. The results demonstrate a trend that the more basic the TnT NH₂-terminus, the stronger the binding of the TnT molecule to TnI.

The binding of chimera TnT to Tm also exhibits a similar charge-dependent trend. Figure 5B demonstrates that the Tm concentrations required for 50% maximum binding to Basic and Acidic chimeric TnT were 7.7 ± 0.1 nM and 14.2 ± 1.8 nM, respectively ($P < 0.05$). Consistently, the binding affinity of the more acidic adult mouse cardiac TnT to Tm was even lower than that of the Acidic chimeric TnT. (The Tm concentration required for 50% maximum binding was 23.5 ± 1.5 nM; $P < 0.05$.) The maximum binding level of Basic chimeric TnT to Tm was greater than that of Acidic

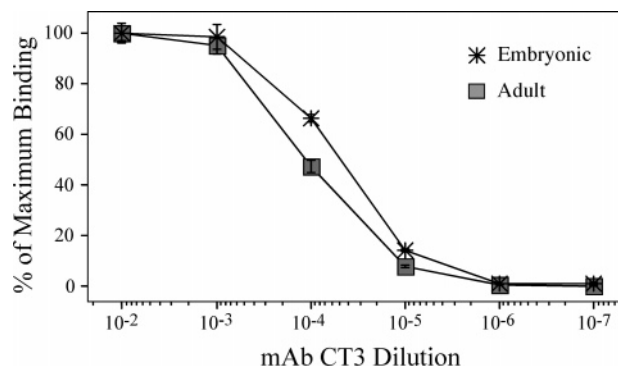


FIGURE 6: Alternative splicing-generated developmental isoforms of cardiac TnT differing in NH₂-terminal charge show differences in molecular conformation. The ELISA epitope affinity titration curves of mAb CT3 demonstrate that exclusion or inclusion of the exon 4 and 5 encoded segments from the adult and embryonic mouse cardiac TnTs has long-range conformational effects on the CT3 epitope located in the central region.

chimeric TnT and adult mouse cardiac TnT (Figure 5B, inset; the A_{405nm} values are 1.04 ± 0.005 , 0.93 ± 0.018 , and 0.92 ± 0.031 , respectively; $P < 0.05$), also indicating a higher coupling strength, which may affect the Tn-thin filament anchoring.

Native Embryonic and Adult Mouse Cardiac TnT Isoforms Exhibit Conformational and Functional Differences. To evaluate whether the long-range conformational effects and altered protein interactions observed for the chimeric model TnT molecules are representative of the natural NH₂-terminal charge variations occurring among TnT isoforms, we investigated embryonic and adult mouse cardiac TnT isoforms. The NH₂-terminal charge of the embryonic and adult TnT isoforms differ as a result of the developmentally regulated alternative mRNA splicing of the exon 5 segment (43) and the non-developmental alternative splicing of exon 4 (31) (TnT-1 and TnT-4, Figure 1). The additive NH₂-terminal charge effect of the segments of exons 4 and 5 imparts these two isoforms with the largest NH₂-terminal charge difference (-10 in TnT-1 compared with that in TnT-4; Table 1) among the four cardiac TnT isoforms (44). In contrast to the chimeric TnT molecules in which nine negative charges in the NH₂-terminal region produced a pI difference of 2.66, in the two naturally occurring mouse cardiac TnT molecules, the NH₂-terminal charge difference of 10 only accounts for a pI difference of 0.32 because of the fact that the NH₂-terminal region of cardiac TnT is already highly negatively charged (Figure 1; Table 1). An ELISA epitope analysis using mAb CT3 against a central epitope shared by the two isoforms (Figure 1) showed a binding affinity to the embryonic cardiac TnT higher than that to the adult isoform (Figure 6). The response of CT3 epitope affinity to NH₂-terminal variation here did not follow the charge trend seen in the chimeric TnTs (Figure 4) and may reflect other modulatory factors (see discussions below). Nonetheless, this result demonstrates that the naturally occurring NH₂-terminal splicing alterations in TnT isoforms also have long-range conformational effects on the core structure, validating the observation from the chimeric model TnT molecules.

Corresponding to the conformational difference, the naturally occurring cardiac TnT isoforms also demonstrate altered interactions to TnI and Tm in solid-phase protein binding experiments (Figure 7). The more acidic embryonic

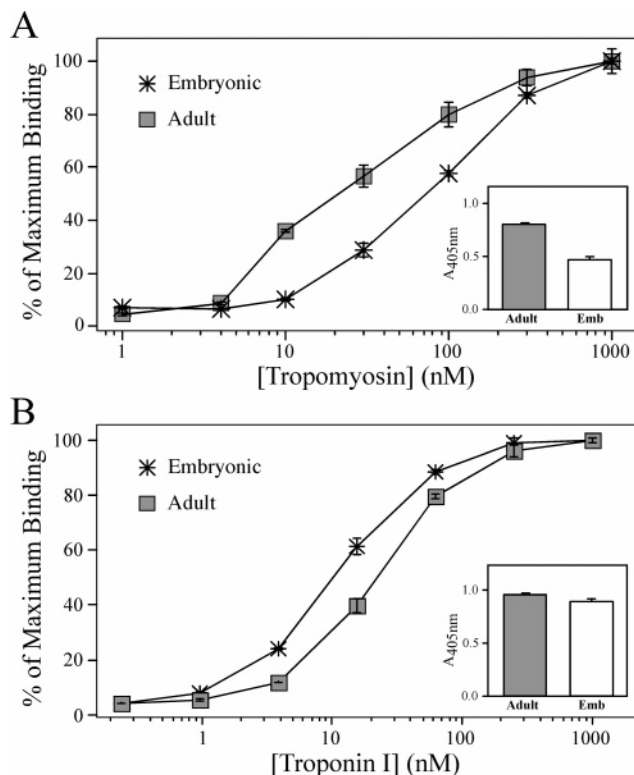


FIGURE 7: Alternative splicing-generated NH₂-terminal charge difference in cardiac TnT affects binding affinity to Tm and TnI. (A) Solid-phase protein binding curves demonstrate that the less acidic adult mouse cardiac TnT immobilized to microtiter plate bound with a higher affinity to Tm than the more acidic embryonic cardiac TnT, as determined by the concentration of Tm required for 50% maximum binding ($P < 0.05$). The adult cardiac TnT also exhibited a higher level of maximal binding to Tm than the embryonic cardiac TnT (shown by the A_{405nm} values in the inset). (B) However, the adult cardiac TnT bound to TnI with a lower affinity compared to that of the embryonic cardiac TnT ($P < 0.05$), whereas the A_{405nm} values in the inset show that the adult cardiac TnT demonstrated a moderate but statistically significant higher maximal binding to TnI compared to that of the embryonic cardiac TnT ($P < 0.05$).

cardiac TnT exhibits a lower binding affinity to Tm compared to that of the less acidic adult cardiac TnT (Figure 7A, the Tm concentration required for 50% maximum binding was 89.9 ± 14.1 nM and 22.3 ± 3.7 nM, respectively; $P < 0.05$). The embryonic cardiac TnT also exhibited a lower level of maximal binding to Tm compared to that of the adult isoform (Figure 7A inset, $A_{405nm} = 0.51 \pm 0.027$ and 0.81 ± 0.041 , respectively; $P < 0.05$). These results were similar to those observed for chimeric TnT, indicating that the more negative NH₂-terminal charge produces a lower binding affinity and weaker coupling strength to Tm.

Different from the trend of charge dependence observed in the chimeric TnTs, the more acidic embryonic mouse cardiac TnT bound to TnI with a greater affinity than that of the adult cardiac TnT (Figure 7B; the TnI concentration required for 50% maximum binding was 11.4 ± 1.0 nM and 23.3 ± 1.4 nM, respectively; $P < 0.05$). It is worth noting that the conformational difference between mouse adult and embryonic TnTs at the CT3 epitope adjacent to the TnI-binding site (Figure 1) also did not follow the charge trend (Figure 6). In contrast, the maximum binding of the embryonic cardiac TnT to TnI was not higher but lower than

that of the adult isoform (Figure 7B inset, A_{405nm} values were 0.89 ± 0.019 and 0.95 ± 0.010 , respectively; $P < 0.05$), similar to the general trend of charge dependence. Although the data are consistent with a role for the long-range effect of NH₂-terminal charge on the COOH-terminal functional domain, the TnI binding behavior of the embryonic versus adult cardiac TnT isoforms suggests that the negative charge effect may have been maximized in the already highly acidic adult cardiac TnT and other factors such as primary structure features may play a role in the modulatory function of the TnT NH₂-terminal domain. The binding of TnT to Tm involves an additional central site in contrast to the single COOH-terminal site binding to TnI (2). The closer proximity of the central Tm site to the NH₂-terminal domain may confer a higher sensitivity of TnT-Tm interaction to TnT NH₂-terminal charge variation (Figure 7A). Nonetheless, the consistent charge effect in the cardiac TnT isoforms on the coupling strength to TnI (Figure 7B inset) is more closely related to the function in the assembled Tn complex than it is to binding kinetics. Together with Tm coupling strength (Figure 7A), these results support the hypothesis that charge variation within the TnT NH₂-terminus plays a role in fine-tuning the function of the muscle thin filament.

Cardiac TnT Core Structure Forms a Functional Tn Complex. To verify that the proposed core structure of TnT indeed retains physiological function *in vivo*, we developed transgenic mouse lines that overexpress the cardiac TnT core structure in the heart (Tg McTnT-ND₇₂). In Figure 8A, SDS-PAGE and Western blots using the anti-cardiac TnT mAb CT3 demonstrate a high level of McTnT-ND₇₂ in the cardiac muscle of transgenic mice. As shown in Figure 8B, densitometry of the Western blot demonstrates the cardiac TnT core was expressed as $49.5 \pm 1.8\%$ of the total cardiac TnT in the adult transgenic mouse ventricular muscle. All McTnT-ND₇₂ transgenic mouse lines exhibit normal baseline life activity without apparent cardiac abnormalities. These transgenic mice expressing the cardiac TnT core in the heart muscle provide an integrated physiological system to evaluate the function of the TnT core structure *in vivo*.

To investigate the function of the TnT core structure and the effect of NH₂-terminal variation in forming the Tn complex *in vivo*, we examined the Tn complex isolated from cardiac muscle thin filaments of Tg McTnT-ND₇₂ mice, transgenic mice expressing the embryonic TnT-1 in the heart (Tg McTnT-Emb), and wild-type mice by TnI-1 mAb affinity chromatography (42). As shown in Figure 8B, the Tn complex isolated from Tg McTnT-ND₇₂ hearts contained the McTnT-ND₇₂ core as $42.2 \pm 4.7\%$ of the total TnT recovered. The Tn complex isolated from Tg McTnT-Emb hearts contained embryonic TnT equal to $68.3 \pm 0.4\%$ of the total TnT recovered. These amounts of exogenous TnT incorporated in the isolated Tn complex were not statistically different from the exogenous TnT detected in the whole heart homogenates (Figure 8B, $49.5 \pm 1.8\%$ for Tg McTnT-ND₇₂ and $68.8 \pm 10.1\%$ for Tg McTnT-Emb). These results demonstrate that the engineered TnT core structure and the exogenous embryonic cardiac TnT effectively incorporate into the adult cardiac Tn complex *in vivo*.

DISCUSSION

Defining the Functional Core Structure of TnT. The structure-function relationship of TnT has been extensively

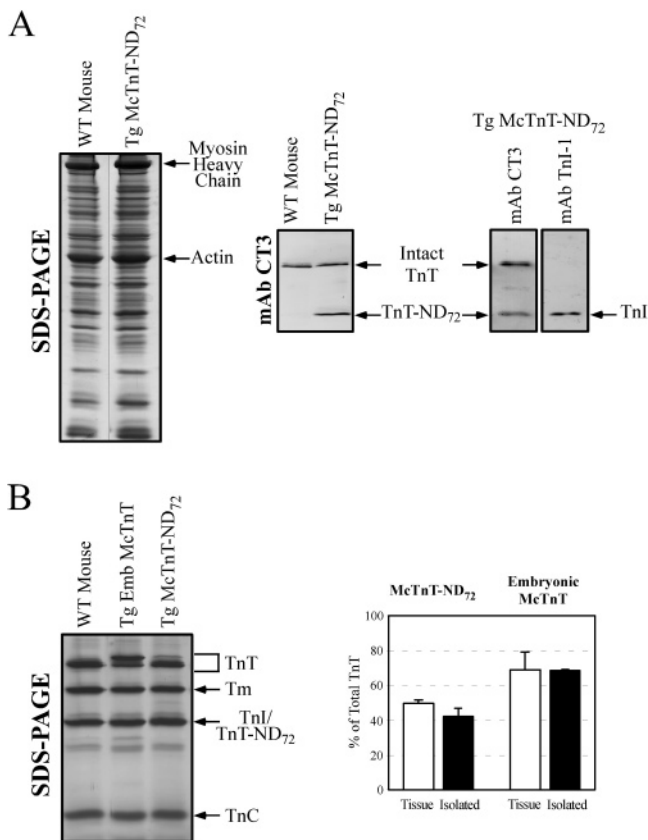


FIGURE 8: Preserved myofilament incorporation and sustained *in vivo* function of the TnT core structure. (A) SDS-PAGE and Western blot using the CT3 mAb demonstrates the high level expression of the cardiac TnT core in the heart of adult McTnT-ND₇₂ transgenic mice. The mAb TnI-1 Western blot shows that McTnT-ND₇₂ comigrates with the endogenous cardiac TnI. (B) Coomassie Blue staining of SDS-PAGE gels demonstrates the stoichiometric recovery of the tertiary Tn complex from wild-type and Tg McTnT-ND₇₂ hearts using TnI-1 mAb immunoaffinity chromatography. As described above, the comigration of TnT-ND₇₂ with TnI formed a fused band in the Tn sample from Tg McTnT-ND₇₂ hearts. In addition to the stoichiometric recovery of Tn subunits, the TnI affinity column also recovered Tm, indicating the native biological activity of the Tn complex. High percentages of the exogenous cardiac TnT in the total cardiac tissue of the transgenic mouse heart (open bars) were recovered in the isolated Tn complex (solid bars), indicating that McTnT-ND₇₂ and embryonic TnT both effectively incorporate into the myofilament *in vivo*.

studied by characterization of the chymotryptic NH₂-terminal fragment T1 (residues 1–158) and COOH-terminal fragment T2 (residues 159–259) defined two decades ago using rabbit fast skeletal muscle TnT (46, 47). The T1 region is often referred to as the tail domain of TnT, rendering the Tn complex asymmetric. Primary structures of the three muscle type-specific TnT isoforms from many species have demonstrated that the T1 fragment consists of the highly divergent NH₂-terminal region and a highly conserved portion that interacts with the Tm head-to-tail overlap (48–50). The present study investigated the hypothesis that the conserved portion of T1 (i.e., the central region of the TnT polypeptide chain) is a part of the TnT core structure, whereas the hypervariable NH₂-terminal region of T1 (i.e., the CB3 fragment corresponding to residues 1–49 in rabbit fast skeletal muscle TnT) that does not bind to any known proteins of the muscle thin filament is an auxiliary modulating structure. Investigating the TnT structure–function

relationship by dissecting the molecule according to its intrinsic features rather than the exogenous protease-generated fragments will help to better understand the two-site binding between TnT and Tm as well as the functional mechanism of the NH₂-terminal variable region.

By progressive NH₂-terminal deletions of cardiac TnT, we investigated the proposed core structure that confers the basal protein binding activities of TnT. A selective deletion of the entire hypervariable region corresponding to the NH₂-terminal 71 amino acids in mouse cardiac TnT was designed to represent the core structure of TnT (Figure 1). This construct, that is, McTnT-ND₇₂, contains the whole conserved region and demonstrates preserved binding to TnI and Tm. Like the intact TnT molecule, McTnT-ND₇₂ fully retains both Tm binding sites (Figure 1). The slightly increased maximal binding of McTnT-ND₇₂ to Tm demonstrates that the deletion of the entire NH₂-terminal variable region did not destroy the TnT core structure, although having a conformational effect on the remaining core structure's binding to Tm.

We previously demonstrated that McTnT-ND₇₂ exhibits preserved binding to TnI (45) through the COOH-terminal region. The present study showed that McTnT-ND₇₂ also preserved the binding affinity to Tm. In comparison to McTnT-ND₇₂, McTnT-ND₉₂ lacks a portion of the conserved region and shows lower binding affinity to Tm. This difference suggests that the central Tm-binding site was disrupted by the deletion of 20 amino acids from the conserved region in McTnT-ND₉₂, which diminishes the two-site high affinity binding between TnT and Tm. Altogether, these data suggest a TnT core structure consisting of the conserved central and COOH-terminal regions from amino acids 72 to 291 in adult mouse cardiac TnT. The functional importance of amino acids 72–91 in the TnT core structure is supported by a previous study that a cardiac TnT fragment equivalent to McTnT-ND₉₂ produced by caspase proteolytic truncation in myocardial ischemia reperfusion injury had impaired function (51).

TnT Core Structure Conveys Physiological Function *in Vivo*. The preserved core activity of McTnT-ND₇₂ shown by ELISA binding experiments using *E. coli*-expressed recombinant proteins was confirmed *in vivo*. Transgenic mice overexpressing the cardiac TnT core lacking the entire NH₂-terminal variable region in the heart exhibited a normal basal life activity without apparent disruption of cardiac muscle function. Previously, we demonstrated that a COOH-terminal truncated slow skeletal muscle TnT was unable to accumulate in the muscle cells because it did not incorporate into the myofibrils (52). The large amount of McTnT-ND₇₂ recovered in the Tn complex from the cardiac muscle thin filaments (the Guba–Straub solution extracted fraction) of the transgenic mice demonstrates that McTnT-ND₇₂ effectively incorporates into the Tn complex and renders a functional thin filament. The McTnT-ND₇₂ transgenic mice did not exhibit any cardiomyopathy phenotype in the absence of stress stimulation, indicating the ability to sustain basal cardiac function. Although full phenotypic characterization of these mice is underway, this initial observation supports the idea that deletion of the entire NH₂-terminal variable region preserves the core structure and function of TnT. This is in agreement with previous findings that a restricted deletion of the NH₂-terminal variable region occurs *in vivo* (45), and its removal does not abolish the Ca²⁺ activated regulation

of muscle activation (20–23). These findings demonstrate that the TnT NH₂-terminal variable region is nonessential to the basic regulatory function of Tn *in vivo*, although its deletion does alter the interaction of TnT with Tm (20, 24) and the maximal activation of the thin filament (20, 23), consistent with modulatory function.

Regulatory Function of the TnT NH₂-Terminal Variable Region. Because the NH₂-terminal variable region of TnT does not directly bind to other thin filament proteins, its functional significance is a standing question in understanding muscle thin filament regulation. Deletion of the 71 NH₂-terminal residues from TnT did not destroy the function of the McTnT-ND₇₂ molecule but represents a default function in the absence of NH₂-terminal charge modification. The finding of decreased coupling strength of TnT to TnI and Tm resulting from the increased NH₂-terminal negative charge demonstrates that the NH₂-terminal variable region is modulatory to TnT function. Previous findings also indicate a functional significance of the TnT NH₂-terminal variable region: its structure is regulated by alternative RNA splicing during heart and skeletal muscle development (14); the developmentally regulated alternative splicing of the NH₂-terminal region alters the production of force in skinned fibers (29, 44, 53); and its aberrant splicing is related to dilated cardiomyopathy, correlating to changes in actomyosin ATPase activation (27, 28), cardiomyocyte contractility (27, 28), and heart failure (54).

The thin filament regulatory system of muscle, including the Tn tail domain that interacts with the Tm head-to-tail overlap, is an allosteric structure in which long-range conformational effects are essential to the function. Our previous studies suggest a model that the NH₂-terminal variable region of TnT regulates the overall conformation and function of TnT in the Tn complex (14, 19, 25, 26). Despite the extended conformation of TnT (17–19), epitope conformational assays demonstrated that the effect of NH₂-terminal variation is transmitted along the TnT structure to affect the conformation of the central and COOH-terminal regions. Similarly, the Acidic and Basic chimeric TnT models result in conformational differences at both the local NH₂-terminal mAb 3E4 epitope (Figure 4A) and the mAb CT3 epitope located in the central region (Figures 4B). These findings demonstrate that the long-range structural effects of the NH₂-terminal variable region can be caused by variation within the NH₂-terminus, which was also true in naturally occurring cardiac TnT isoforms (Figure 6).

The effects of the NH₂-terminal conformational modulation on TnI and Tm binding are also present in model TnT molecules. The differential charge effects of the TnT NH₂-terminus to regulate the binding affinity and coupling strength of TnT to TnI and Tm are worth investigating. Altered binding affinity may effect the assembling of the Tn complex; however, NH₂-terminal truncated (Figure 8), aberrantly spliced cardiac TnT (28) and chicken fast skeletal muscle TnT (40) all effectively incorporate into the thin filament when overexpressed in the transgenic mouse heart. Therefore, it seems that the NH₂-terminal regulation of the coupling strength between TnT and other thin filament proteins in the assembled myofibril is more critical to contractile function. Consistent with the propagation of the NH₂-terminal conformational modulation along the extended TnT molecule,

the TnT NH₂-terminal charge variation would most effectively alter the more proximal central Tm-binding site as opposed to the distal COOH-terminal TnI binding site. This is significant to muscle regulation in that the central TnT-Tm binding site is critical to the propagation of Ca²⁺ activation along the thin filament (2). Stronger versus weaker coupling among the regulatory proteins may correspond to a more rigid or more flexible molecular switch to alter the activation dynamics of the myofibril. This idea is supported by findings that skinned muscle fibers containing an acidic TnT isoform exhibited decreased Ca²⁺ sensitivity and increased cooperativity of activation, whereas Ca²⁺ sensitivity was increased and cooperativity decreased in muscle fibers containing a basic TnT isoform (40). The role of TnT NH₂-terminal splicing as a mechanism to regulate the dynamics of thin filament activation is interesting and deserves further investigation.

Role of NH₂-Terminal Charge. The hypervariable nature of the TnT NH₂-terminal region imposes difficulties in understanding its role in the Ca²⁺ regulation of muscle contraction. Among the physical features of the NH₂-terminal variable region, the charge variation demonstrates a consistent acidic-to-basic trend among the developmentally regulated isoforms (14, 33). Focusing on this feature, we sought to dissect the effects of the TnT NH₂-terminal charge on the interaction of TnT with TnI and Tm using model TnT molecules and naturally occurring NH₂-terminal splice variants. Consistent with a role for the TnT NH₂-terminal charge in Ca²⁺ regulation, all TnT molecules investigated exhibit a strong quantity-dependent effect of NH₂-terminal charge on Tm binding such that the more basic TnT coupled more strongly to Tm than the more acidic TnT (Figures 5B and 7A).

Unlike the two-site (central and COOH-terminal) binding to Tm, TnT binding to TnI occurs at a single COOH-terminal site (2). In the chimeric TnT molecules, the difference in pI is large (Table 1; Basic TnT = 8.99; Acidic TnT = 6.33; adult cardiac TnT = 5.19). These chimera demonstrate significant quantity-dependent effects of the NH₂-terminal charge on the binding to TnI such that the more basic TnT bound more strongly (Figure 5A). However, the total NH₂-terminal negative charge is strong in the cardiac TnT isoforms, diminishing the overall effect of NH₂-terminal charge alterations on the pI (Table 1; adult 5.19 vs embryonic 4.87) and TnI-binding affinity (Figure 7B) of the isoforms. This could result from the possibility that the NH₂-terminal charge difference between TnT isoforms is buffered by the already large amount of negative charge in the cardiac TnT NH₂-terminus to maintain the NH₂-terminal charge modulation effective within the range of physiological pH.

The results reported here suggest that although the core structure of TnT confers the basal function in sustaining muscle contraction, the evolutionarily additive hypervariable NH₂-terminal domain regulates TnT's interactions within the thin filament regulatory system through charge variation-based long-range conformational effects. Although other mechanisms that convey the physiological function of the TnT NH₂-terminal variable region remain to be investigated, the high tolerance of the TnT NH₂-terminal domain to structural variation or removal allows this mechanism to effectively extend the functional capacity of muscle during development and functional adaptation.

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