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Review Series

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Genetic mutations and mechanisms in dilated cardiomyopathy

Elizabeth M. McNally, Jessica R. Golbus, and Megan J. Puckelwartz

Department of Human Genetics, University of Chicago, Chicago, Illinois, USA.

Genetic mutations account for a significant percentage of cardiomyopathies, which are a leading cause of congestive heart failure. In hypertrophic cardiomyopathy (HCM), cardiac output is limited by the thickened myocardium through impaired filling and outflow. Mutations in the genes encoding the thick filament components myosin heavy chain and myosin binding protein C (*MYH7* and *MYBPC3*) together explain 75% of inherited HCMs, leading to the observation that HCM is a disease of the sarcomere. Many mutations are “private” or rare variants, often unique to families. In contrast, dilated cardiomyopathy (DCM) is far more genetically heterogeneous, with mutations in genes encoding cytoskeletal, nucleoskeletal, mitochondrial, and calcium-handling proteins. DCM is characterized by enlarged ventricular dimensions and impaired systolic and diastolic function. Private mutations account for most DCMs, with few hotspots or recurring mutations. More than 50 single genes are linked to inherited DCM, including many genes that also link to HCM. Relatively few clinical clues guide the diagnosis of inherited DCM, but emerging evidence supports the use of genetic testing to identify those patients at risk for faster disease progression, congestive heart failure, and arrhythmia.

Cardiomyopathy: diagnosis and prevalence

Cardiomyopathy is defined by a pathologically abnormal myocardium. There are four major classifications of cardiomyopathy: dilated (DCM), hypertrophic (HCM), restrictive (RCM), and arrhythmogenic RV (ARVC). Cardiomyopathy is most commonly diagnosed through in vivo imaging, with either echocardiography or, increasingly, cardiac MRI. DCM refers to enlargement of the heart, which often affects all four chambers, especially late in the disease. Most commonly, DCM is associated with reduced LV function or systolic function, although early in the disease the LV may be dilated, with only minimally reduced function. In contrast, HCM is characterized by increased LV wall thickness, often targeting the septum that separates the LV from the RV (Figure 1). RCM is the most elusive, in part because the heart may appear morphologically close to normal, with only minor increased wall thickness or modestly decreased LV ejection fraction. The infiltrative process underlying RCM is often not readily detectable in vivo with even the most sensitive imaging technique. RCM is characterized by impaired filling of the heart, known as diastolic dysfunction, which reduces cardiac output. ARVC is characterized by reduced function and thinning of the RV with a fibrofatty infiltration that can be seen on MRI.

Myocardial ischemia remains a common cause of DCM, accounting for approximately half of DCM. Toxic, metabolic, and immunologic causes have each been linked to DCM, as well as hypertension and valvular disease. Genetic cardiomyopathy, which runs in families, is now more commonly diagnosed, largely because surveillance for and awareness of the condition have improved. It is also now increasingly understood that more than one etiology may contribute to DCM within a given individual. Unlike cardiomyopathy that results from some of the other causes, genetic cardiomyopathy is a progressive disease. The time course is highly dependent on the age of presentation, which relates to

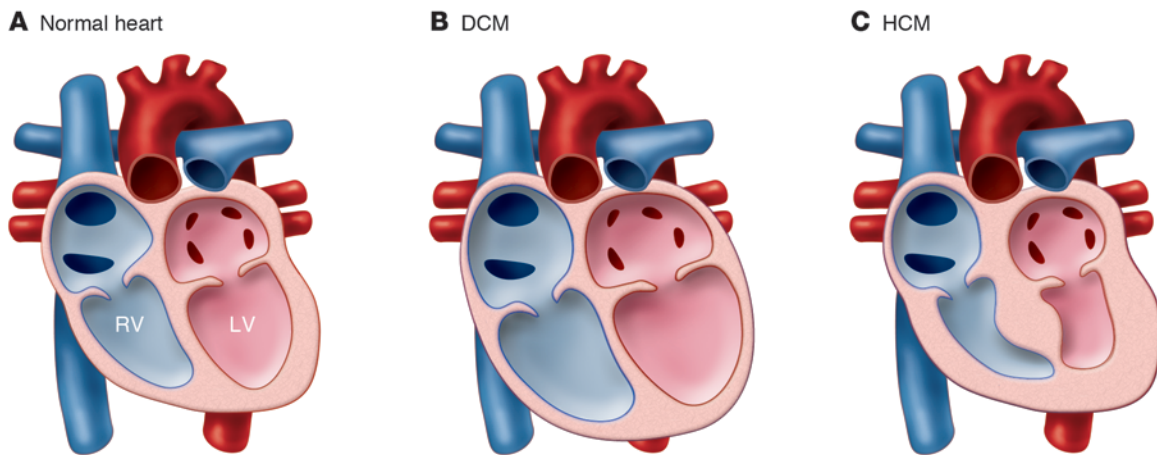
the specific genetic mutation as well as the presence of disease modifiers. Because it is now commonplace to screen relatives of individuals affected with cardiomyopathy, many individuals are diagnosed at much earlier stages of disease. This detection strategy is expected to change both the prevalence and treatment strategies for cardiomyopathy. Previous estimates of DCM prevalence were 1 in 2,500–3,000 (1, 2). However, these data reflect older diagnostic modalities and therefore likely significantly underestimate the disease prevalence. For HCM, previous estimates were similar to DCM, but newer modalities and heightened diagnostic awareness now estimate the prevalence of HCM to be 1 in 500 (3). The prevalence of DCM increases with age, and DCM is rare (1–2 in 100,000) in the pediatric population (4). DCM is also more common among males, and prevalence varies with ethnicity. Nonischemic DCM, defined as DCM not from myocardial ischemia or infarct, is familial in 25%–50% of cases, with estimates that vary based how family members were screened (5).

Genetic mutations in DCM

Familial DCM refers to DCM that is inherited as a single gene disorder in a Mendelian pattern. The primary mode of inheritance for familial DCM is autosomal dominant, with reduced penetrance and variability expressivity. Mitochondrial mutations also contribute to DCM with the expected matrilineal inheritance. Autosomal-recessive mutations are a less common cause of familial DCM, although X-linked recessive inheritance is associated with several genes such as dystrophin, tafazzin, and emerin. At least 50 single genes have been identified as linked to familial DCM, and the majority of these elicit disease as dominant mutations. With few exceptions, it is not possible to determine which gene is responsible for DCM on clinical grounds. That is, mutations in many single genes phenocopy DCM. Because mutations in many individual single genes lead to DCM, genetic testing commonly employs multi-gene panels, in which more than 50 genes can be tested simultaneously. With the use of multi-gene panels for clinical genetic testing, more than a single pathogenic variant can be identified within an individual, thus explaining some features of

Conflict of interest: An immediate family member of E.M. McNally receives patent royalties related to massively parallel sequencing.

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**Figure 1**

Morphological changes to the heart in cardiomyopathy. (A) Normal heart. (B) In DCM, the heart enlarges with increased diameter and reduced function. (C) In HCM, the myocardium — especially in the LV — becomes thickened, leading to impaired filling and emptying.

early onset in more severe forms of disease (6, 7). Some of the genes linked to DCM encode proteins of the sarcomere, costamere, Z band, and nuclear membrane, while others have functions distinct from these broad cell biological classifications (Table 1).

Sarcomere DCM genes. The sarcomere is the unit of contraction for striated muscle (Figure 2). Thin filaments emanate from the Z band with filamentous actin (gene name *ACTC1*) forming the core and tropomyosin (encoded by *TPM1*) and the troponin subunits (encoded by *TNNT2*, *TNNC1*, *TNNI3*) providing calcium regulation of the actomyosin interface. The thick filament core is formed by β -myosin heavy chain (β -MHC) (encoded by *MYH7*), with myosin binding protein C (MYBPC3) highly concentrated in the M band region. The giant protein titin (encoded by *TTN*) spans the length of half the sarcomere, where it acts as a stretch sensor, transmitting signals from its anchor at the Z band to its carboxyterminal kinase domain. Dominant mutations in the genes encoding these sarcomeric proteins cause DCM. Many sarcomeric gene mutations are nonsynonymous SNPs, small insertion/deletions that may be in-frame or, alternatively, protein-truncating mutations. An estimated 35%–40% of genetic DCMs may result from sarcomere gene mutations, with the majority attributed to protein-truncating mutations in *TTN* (8). Notably, arrhythmia risk may not correlate with morphological abnormality (9), and additional markers to predict arrhythmia risk are needed.

The mechanism of action for some sarcomeric DCM mutations is dominant negative. A gain of function by the protein produced from the mutant allele interferes with sarcomere formation and function. The experimental evidence to support expression of the mutant allele is limited by the availability of human hearts and often relies on demonstration of variant pathogenicity within an experimental model. The deletion of lysine at position 210 (del210K) in *TNNT2* has been linked to DCM (10). Although initially believed to be a highly potent mutation with early onset and lethal disease, subsequent reports revealed the *TNNT2* del210K mutation to be associated with a marked phenotypic expression range with both young and older affected individuals (11). The *TNNT2* del210K mutation was engineered as a knockin allele in the mouse, where it produced DCM and reduced survival

(12). Skinned cardiac muscle fibers from heterozygous del210K mice showed decreased calcium sensitivity for force generation (12). Homozygous *TNNT2* del210K mice showed an even more enhanced rightward shift of decreased calcium sensitivity, markedly enlarged hearts, and further reduced survival compared with heterozygous *TNNT2* del210K mice. These findings suggest that *TNNT2* del210K disrupts intermolecular interactions within the sarcomere, most likely with tropomyosin.

One recent study of a *TNNC1* DCM mutation used mass spectrometry to document equimolar expression of the G159D allele and the normal *TNNC1* allele in an explanted human DCM heart (13). The calcium sensitivity of force generation was increased with this mutation, in opposition to what occurs with the *TNNT2* mutation described above. Tropomyosin is a thin filament protein that is also mutated in DCM. For example, the *TPM1* D230N mutation was described in two large, unrelated families with DCM (14). As with *TNNT2* del210K, the DCM clinical phenotype with *TPM1* D230N markedly varied. Some children presented with cardiomyopathy, while other family members with the identical gene mutation remained largely asymptomatic, with mildly increased LV diameter and mildly reduced systolic function until later life. The D230N *TPM1* DCM mutation was modeled in vitro, where a decreased calcium sensitivity of actin-activated myosin ATPase activity was seen (14). This result is in contrast to the increased calcium sensitivity seen with an HCM-inducing *TPM1* D175N mutation. Sarcomere mutations have been well studied using enzymatic assays of myosin function (e.g., assays of actin-activated myosin ATPase activity) and more complex biophysical assays of actin's sliding filament velocity, which can be further interrogated to monitor in vitro force production. Isolated muscles, usually the papillary muscle, are treated with detergents to strip away membrane components and leave intact, skinned fibers. These reductionist approaches (in vitro motility or isolation of fibers) simplify the sarcomere, often leading to useful results. However, the results should be cautiously interpreted, especially where there are inconsistencies, because these approaches do not replicate the full complexity of sarcomere function within a failing heart (for a more full discussion, see ref. 15).



Table 1
Cardiomyopathy genes

Gene	Gene name	Cardiomyopathy subtype(s)
<i>ABCC9</i>	ATP-binding cassette, sub-family C, member 9	DCM
<i>ACTC1</i>	Actin, α , cardiac muscle 1	DCM, HCM, LVNC
<i>ACTN2</i>	Actinin, α 2	DCM, HCM
<i>ANKRD1</i>	Ankyrin repeat domain 1 (cardiac muscle)	DCM, HCM
<i>BAG3</i>	BCL2-associated athanogene 3	DCM
<i>CASQ2</i>	Calsequestrin 2 (cardiac muscle)	LVNC, CPVT
<i>CAV3</i>	Caveolin 3	HCM
<i>COX15</i>	COX15 homolog, cytochrome c oxidase assembly protein	HCM
<i>CRYAB</i>	Crystallin α B	HCM
<i>CSRP3</i>	Cysteine and glycine-rich protein 3	DCM, HCM
<i>CTF1</i>	Cardiotrophin 1	DCM
<i>DES</i>	Desmin	DCM
<i>DMD</i>	Dystrophin	DCM
<i>DNAJC19</i>	DnaJ (Hsp40) homolog, subfamily C, member 19	DCM
<i>DSC2</i>	Desmocollin 2	DCM, ARVC
<i>DSG2</i>	Desmoglein 2	ARVC
<i>DSP</i>	Desmoplakin	DCM, ARVC
<i>DTNA</i>	Dystrobrevin, α	LVNC
<i>EMD</i>	Emerin	DCM
<i>EYA4</i>	Eyes absent homolog 4	DCM
<i>FHL2</i>	Four and a half LIM domains 2	DCM
<i>FKTN</i>	Fukutin	DCM
<i>FOXD4</i>	Forkhead box D4	DCM
<i>GLA</i>	Galactosidase, α	HCM
<i>JUP</i>	Junction plakoglobin	ARVC
<i>LAMA4</i>	Laminin, α 4	DCM
<i>LAMP2</i>	Lysosomal-associated membrane protein 2	DCM, HCM
<i>LDB3</i>	LIM domain binding 3	DCM, LVNC
<i>LMNA</i>	Lamin A/C	DCM, LVNC
<i>MYBPC3</i>	Myosin binding protein C, cardiac	DCM, HCM, LVNC
<i>MYH6</i>	Myosin, heavy chain 6, cardiac muscle, α	DCM, HCM
<i>MYH7</i>	Myosin, heavy chain 7, cardiac muscle, α	DCM, HCM, LVNC
<i>MYL2</i>	Myosin, light chain 2, regulatory, cardiac, slow	HCM
<i>MYL3</i>	Myosin, light chain 3, alkali; ventricular, skeletal, slow	HCM
<i>MYLK2</i>	Myosin light chain kinase 2	HCM
<i>MYOZ2</i>	Myozenin 2	HCM
<i>NEXN</i>	Nexilin (F actin binding protein)	DCM, HCM
<i>PKP2</i>	Plakophilin 2	ARVC
<i>PLN</i>	Phospholamban	DCM, HCM
<i>PRKAG2</i>	Protein kinase, AMP-activated, γ 2, non-catalytic subunit	HCM
<i>PSEN1</i>	Presenilin 1	DCM
<i>PSEN2</i>	Presenilin 2	DCM
<i>RBM20</i>	RNA binding motif protein 20	DCM
<i>RYR2</i>	Ryanodine receptor 2 (cardiac)	CPVT
<i>SCN5A</i>	Sodium channel, voltage-gated, type V, α subunit	DCM
<i>SDHA</i>	Succinate dehydrogenase complex, subunit A, flavoprotein	DCM
<i>SGCD</i>	Sarcoglycan, δ	DCM
<i>SYNE1</i>	Spectrin repeat containing, nuclear envelope 1	DCM
<i>SYNE2</i>	Spectrin repeat containing, nuclear envelope 2	DCM
<i>TAZ</i>	Tafazzin	DCM, LVNC
<i>TCAP</i>	Titin-cap (telethonin)	DCM
<i>TMEM43</i>	Transmembrane protein 43	ARVC
<i>TMPO</i>	Thymopoietin	DCM
<i>TNNC1</i>	Troponin C type 1 (slow)	DCM, HCM
<i>TNNI3</i>	Troponin I type 3 (cardiac)	DCM, HCM
<i>TNNT2</i>	Troponin T type 2 (cardiac)	DCM, HCM, LVNC
<i>TPM1</i>	Tropomyosin 1 (α)	DCM, HCM
<i>TTN</i>	Titin	DCM, HCM, ARVC
<i>TTR</i>	Transthyretin	HCM
<i>VCL</i>	Vinculin	DCM, HCM

CPVT, catecholeminergic polymorphic ventricular tachycardia; LVNC, LV noncompaction.

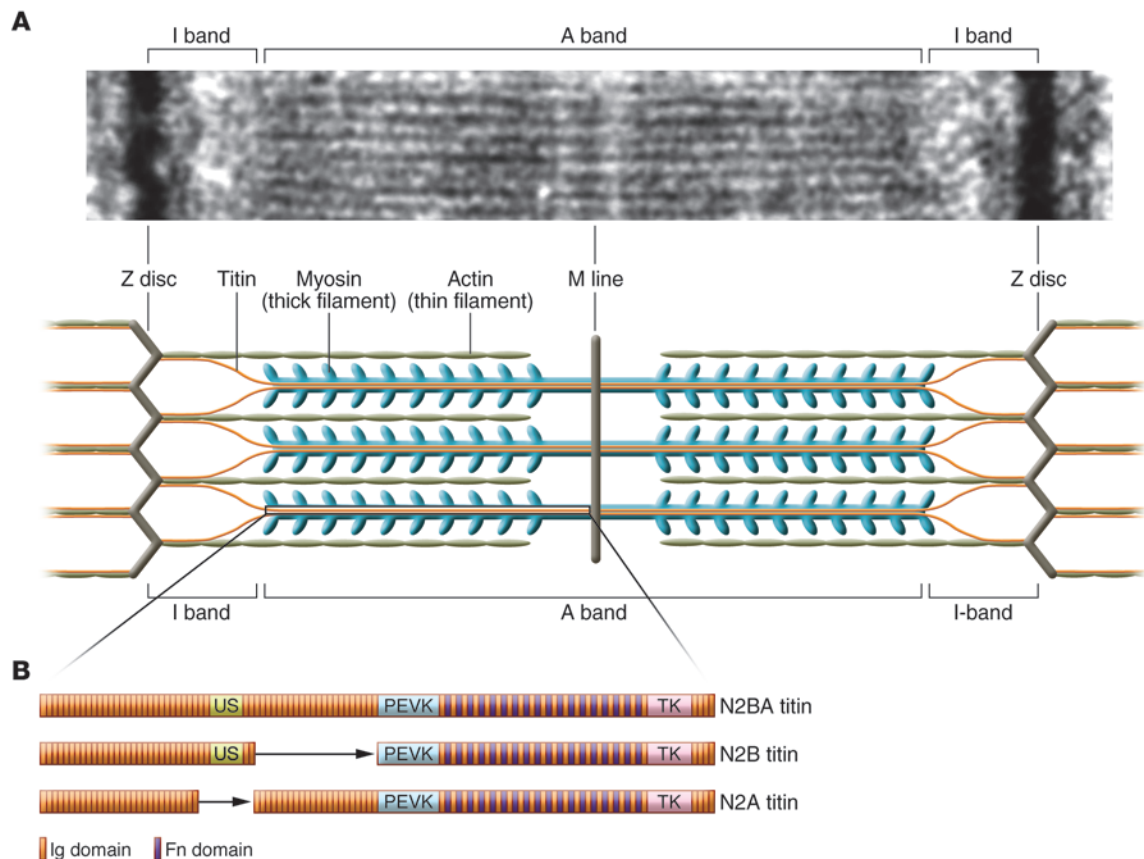


Figure 2

The sarcomere is a target for cardiomyopathy mutations. **(A)** The sarcomere from Z disc to Z disc. Electron micrograph of a sarcomere from a human heart. The *TTN* gene encoding the giant protein titin is mutated in DCM. Titin's amino terminus anchors in the Z band, and its carboxy terminus ends in the M band. The titin kinase (TK) domain is found at the carboxy terminus and, when mutated, results in impaired stretch sensing and signaling. Titin interacts with both the thin and thick filaments. The thick-filament proteins (grey) are encoded by *MYH7* and *MYBPC3*, two genes commonly linked to HCM. **(B)** Isoforms of titin. Alternative splicing in the region of titin that encodes the I band gives rise to isoforms with varying spring properties. The N2B isoform is found exclusively in cardiac muscle and the N2A isoform in skeletal muscle. The N2BA isoform is also found in cardiac muscle and contains features found in both N2B and N2A titin. N2BA titin has a longer extensible I band region than N2B titin, making it more compliant. US, unique sequence; Fn, fibronectin domains; PEVK, repeating units of amino acids (proline, glutamic acid, valine, and lysine).

DCM mutations also occur in the sarcomere's thick filament (16). *MYH7* and *MYBPC3* mutations are the most common causes of HCM, and HCM may progress to DCM. However, there are thick filament mutations that have no HCM phase. Two *MYH7* mutations, S532P and F764L, each linked to familial DCM, were independently modeled in mice by modifying the endogenous α -MHC (encoded by *MYH6*), which is the major MHC of the small rodent heart. Each of these mutations maps within the S1 head of myosin that harbors the actin-binding, ATPase, and force-producing domains. The S532P mutation occurs within the actin-binding domain, and the F764L mutation occurs within the converter domain. Actin-activated ATPase is reduced, as is in vitro sliding motility of actin. Force production is not decreased, but rather a reduction in myosin step size is observed. Since ATP hydrolysis produces less displacement of myosin, this is a less energetically efficient sarcomere. Mutations in *MYH7* have also been described in patients with LV noncompaction cardiomyopathy (LVNC), a pathological variant with deep trabecular invaginations in the LV. Truncating and

point mutations in *MYH7* have been identified in LVNC families (17). In addition, *TPM1* and *MYBPC3* mutations have also been found in LVNC patients (17, 18).

Most recently, mutations in the giant protein titin were estimated to be responsible for approximately 25% of DCM (8). The titin protein includes more than 35,000 amino acids, containing many repeating fibronectin-like and Ig-like domains. Titin is considered a molecular ruler that regulates and transmits information about sarcomere length to the sarcomere and the cardiomyocyte (19). Consistent with this role, the carboxy terminus titin kinase domain is stimulated by activity and stretch (20). The *TTN* gene at chromosome 2 was previously linked to familial DCM (21), and those findings have been reinforced by studies in animal models (22). To evaluate N2BA, one of the numerous isoforms encoded by the *TTN* gene, an exon-capture strategy was employed. The N2BA form contains additional domains compared to the N2B form, including the Ig and PEVK domains that lend additional recoil (23). In heart failure, the ratio of N2BA to N2B is altered, which may be both a cause and effect of cardiac ventricular dilation (24).

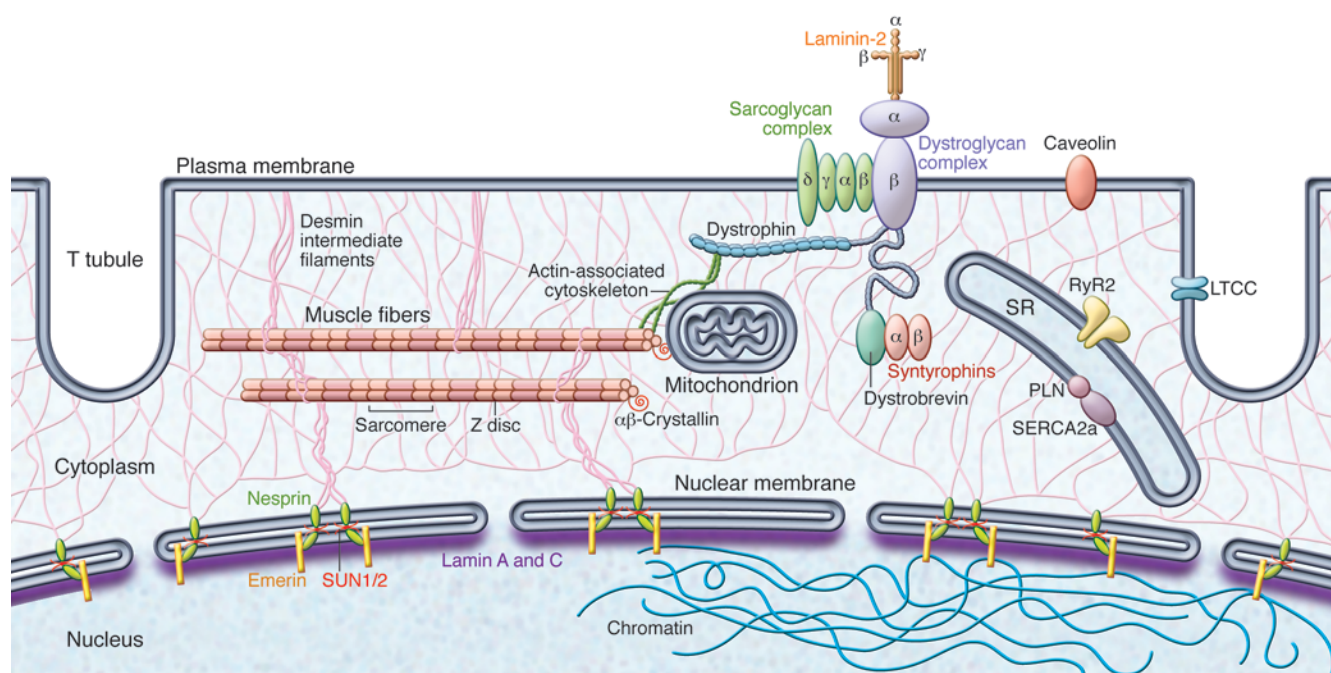


Figure 3

A view of the cardiomyocyte. The cytoplasm of the cardiomyocyte contains sarcomeres, which contain thin and thick filaments. Mutations in sarcomere genes lead to HCM and DCM. Plasma membrane-associated proteins such as dystrophin and its associated proteins, the sarcoglycans, are mutated in inherited DCM associated with skeletal muscle disease. Mutations of genes encoding nuclear membrane proteins such as lamins A and C, emerin, and nesprins lead to inherited DCM due to an inappropriate transcriptional response to mechanical stress. Many of these nuclear membrane genes also induce cardiac conduction system disease.

Many of these same mutations are also expected to alter skeletal muscle titin, which is encoded by the related isoform N2-A titin, but whether these variants contribute to subclinical muscle disease remains to be documented.

A dilated heart may be even more dependent on signaling through titin, and frameshifting mutations that disrupt this intramolecular signaling pathway would be expected to adversely affect the heart, as it regularly contends with repetitive and dynamic load. In the study by Herman et al., 54 of 312 DCM patients were found to have truncating *TTN* mutations, which the authors defined as frameshifts, nonsense, or predicted splice site mutations (8). In comparison, only 3 of 231 HCM cases and, intriguingly, 7 of 249 control subjects had protein-disrupting *TTN* mutations. The protein-disrupting *TTN* variants were nonrandomly distributed along the length of titin, with a preference for the A band region of titin. Most *TTN* variants mapped in regions common to both the N2B and N2BA forms, and nearly all *TTN* mutations were heterozygous. A subset of subjects had documented familial disease, thus providing strong support for *TTN* as a dominant DCM gene. In addition to these protein-truncating *TTN* mutations, a large number of *TTN* coding variants of unknown significance were found. This type of protein-altering variation may contribute to DCM complexity, as some variants are predicted to be pathogenic. With this frequency of protein-disrupting variation, the *TTN* gene should be investigated in DCM, although the challenges associated with evaluating the large and repetitive *TTN* gene are enormous. Given this frequency, *TTN* mutations in combination with other forms of heart disease would be expected to enhance the severity of cardiomyopathy expression. For example,

TTN variants in combination with valve defects, hypertension, or myocardial infarction would be expected to result in more significant cardiomyopathy and congestive heart failure. A larger heart, such as that dilated from regurgitant valve disease, may be more susceptible to impaired elastic recoil or a reduction in titin signaling from mutations ablating the titin kinase domain. It also follows that *TTN* mutations in combination with other pathogenic genetic variants would enhance the severity of DCM. It is notable that HCM subjects had a strikingly lower percentage of *TTN* frameshifting mutations in the Herman study, possibly supporting the idea that HCM sarcomere mutations along with *TTN*

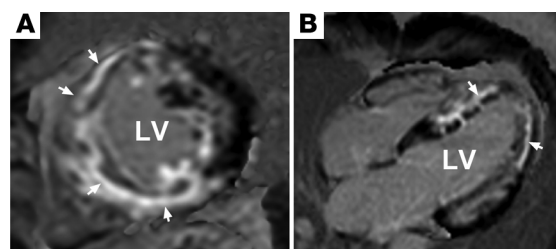


Figure 4

Fibrosis occurs commonly in DCM. Cardiac MRI of a patient with an *MYH7* mutation. (A) Short-axis view sectioning through the LV shows late gadolinium enhancement, which indicates fibrosis. Late gadolinium enhancement is visualized as white regions (arrows) and is never seen in a normal heart. (B) Four-chamber view shows late gadolinium enhancement (arrows) in the same *MYH7* mutant individual.



frameshifting mutations predispose to DCM. More comprehensive genetic testing is required to explore this possibility, and these observations strongly favor the inclusion of *TTN* in genetic screening panels. *TTN* variants have also been described in ARVC (25).

Z band proteins and the costamere. The Z disc or Z band is an electron-dense structure into which the thin filaments and titin anchor. Gene mutations that have been identified in DCM patients that affect multiple Z band proteins include muscle LIM protein (MLP), cardiac ankyrin repeat protein (CARP), myopallidin, α -actinin 2, TCAP, and nexilin (26–32). TCAP and MLP modulate the elastic properties of titin. Phosphorylation of Z band proteins, or even nuclear translocation of Z band elements, are just two mechanisms by which Z band elements signal within the cardiomyocyte.

At the plasma membrane, with a periodicity that matches Z discs, is the costamere, a rib-like structure of cytoplasmic and transmembrane proteins that links the cytoskeleton to the plasma membrane and the extracellular matrix (Figure 3). Dystrophin and its associated proteins, sarcoglycans and dystroglycan, enrich at the costamere. Linked to the cytoplasm alongside talin, vinculin, and integrin, the dystrophin complex protects against contraction-induced injury in both cardiac and skeletal muscle (33). The integrity of the dystrophin complex is critical for mechanotransduction because disruption of this complex from loss-of-function mutations renders cardiomyocytes susceptible to damage elicited from contraction (34). Mutations in dystrophin and the sarcoglycan produce skeletal muscle disease and cardiomyopathy, and so heart failure in these patients is further compromised by hypoventilation from respiratory muscle weakness.

Because of the narrow window in which DCM develops in patients with Duchenne muscular dystrophy (DMD), this population has been ideal for testing whether medications for cardiomyopathy can prevent or slow progression of DCM. DCM prevention was tested by giving the angiotensin converting enzyme perindopril to boys with DMD who began treatment beginning at age 9.5 years, before the development of reduced LV function. This early treatment was associated with reduced progression of cardiomyopathy after five years and also showed a benefit in 10-year mortality (35, 36). The addition of β -adrenergic blockade may be expected to further slow progression of disease. Although animals have been used to test whether early treatment slows progression of DCM, the perindopril study in a human population with DMD provides strong evidence to support what was seen using animal models.

Nuclear membrane defects in genetic DCM. Lamins A and C, two alternatively spliced products of the *LMNA* gene, are expressed in all somatic cells, but mutations in *LMNA* strike in a very tissue-specific manner, with the majority causing cardiomyopathy with variable skeletal muscle involvement. More than 200 different *LMNA* mutations are associated with inherited cardiomyopathy, primarily DCM that may be associated with conduction system disease prior to the evidence of ventricular dilation. *LMNA* mutations that cause DCM are primarily inherited in an autosomal-dominant manner, although autosomal-recessive cases have been reported. Cardiac conduction system disease caused by *LMNA* mutations includes sinus atrial node disease, atrial dysrhythmias, atrioventricular heart block, and ventricular tachyarrhythmias (37). In a study of patients with familial DCM and prominent conduction block, 33% had *LMNA* mutations (38). A recent report found that frameshifting *LMNA* mutations have the highest risk for ventricular arrhythmias (39).

Lamins A and C are intermediate filament proteins that are present at the nuclear membrane and in the nucleoplasm. Lamins A and C, along with the B-type lamins and other lamina-associated proteins, comprise the nuclear lamina, a fibrous structure that provides support to the nuclear membrane. The lamina proteins are part of the LINC complex that Links the Nucleus to the Cytoplasm. In addition to lamins A and C, mutations in other members of the LINC complex cause cardiomyopathy, including emerin and nesprins-1 and -2 (40–42). Stress signals in the cytoplasm are hypothesized to act on the LINC complex to affect gene expression in the nucleus. Both *LMNA* and emerin-null fibroblasts have altered expression of mechanosensitive genes in response to mechanical stress (43, 44). *LMNA* mutations variably affect skeletal muscle, and skeletal muscle from both *LMNA* mouse models and human patients with *LMNA* mutations reveal aberrant expression of electrical activity-dependent genes and epigenetic chromatin modifications, indicating that components of the LINC complex are crucial for an appropriate transcriptional response to mechanical stress (45).

To further understand how the LINC complex affects gene expression, recent studies have examined the role the nuclear lamina plays in chromatin organization. The organization of chromatin in the nucleus is not random and is thought to play a role in transcriptional regulation. Whole chromosomes or portions of chromosomes reorganize in the nucleus, coincident with changes in gene expression. The nuclear lamina participates in the scaffolding of the chromatin through interactions between chromatin and lamins A and C. Fibroblasts from patients with *LMNA*-induced cardiomyopathy have changes in the three-dimensional location of chromatin compared with normal dermal fibroblasts. This chromatin mislocalization occurs concomitant with changes in gene expression, suggesting that the spatial organization of chromatin may be important for disease pathogenesis (46, 47). Together, these data indicate that a functional LINC complex is critical for the heart to properly respond to constant mechanical stress, potentially by regulating gene expression or other nuclear functions.

Genetic assessment and modifiers for cardiomyopathy

As with other genetic disorders, the study of DCM is undergoing a revolution, with implementation of newer, massively parallel next-generation sequencing. Herman et al. developed an exon-capture approach to sequence the more than 350 exons that comprise *TTN* (8). Whole exome and whole genome approaches have now become feasible and are in the clinical domain (48). While it may not be necessary to determine the sequence of all exons or the whole genome to arrive at a genetic diagnosis, it is rapidly becoming cost effective to determine the whole exome or whole genome and then selectively investigate sequence variation in specific gene(s). This analysis expands on the current strategy, which simultaneously evaluates the sequence of many DCM genes. This expansive screening is often undertaken in a single affected family member. With identification, the index mutation is then evaluated in other family members. With expansive testing, more than one pathogenic variant or variants of unknown significance may be uncovered. Determination of pathogenicity has typically relied on segregation in family members and experimental techniques performed in research laboratories. Not all families are large enough for segregation analysis. Laboratory-based experiments, while experimentally satisfying, are not performed with any type of regulatory oversight. More importantly, with the number of



variants currently being identified through expansive testing, such experimental validation is not possible. Future approaches to validate the pathogenicity of mutations will likely rely more heavily on computational prediction algorithms coupled tightly to better estimates of population frequency (49).

The same genetic mutation, even within families, can manifest with distinct cardiac morphology (HCM or DCM), varying age of onset, and range of arrhythmia risk. Modifiers for cardiomyopathy can include both environmental and genetic influences. Animal models are particularly useful for identifying genetic modifiers because of the availability of inbred strains, more uniform housing conditions, and the ability to make an unbiased genomewide search. Le Corvoisier and colleagues mapped modifiers of heart failure using a calsequestrin-expressing transgenic model of cardiomyopathy (50). With this approach, seven different loci were identified and one was identified as *TNNI3K*, a gene encoding cardiac troponin I interacting kinase (51). This gene was implicated in several traits linked to cardiac function, such as fractional shortening and ventricular diameter. These data fit well with the observation that *TNNI3K* is a gene expressed in cardiomyocytes.

Human studies for heart failure modifiers have utilized candidate gene and whole genome approaches. The β_1 -, β_2 -, and the α_2 -adrenergic receptors have common DNA variants that are associated with outcome in heart failure, including survival and exercise capacity (52–55). A genomewide association study identified a region that included the *HSPB7* gene and the *CLCNKA* gene (56, 57). Further evaluation revealed G83R, a specific SNP with functional consequences in *CLCNKA* (rs10927887), which was overrepresented in three cohorts of patients with heart failure, including both nonischemic and ischemic heart failure (58).

Fibrosis is an important biological modifier for cardiomyopathy and cardiopulmonary function (Figure 4). A recent study showed fibrosis in young individuals who had HCM mutations but had not yet developed hypertrophy (59). This indicates that fibrosis

is an early step in HCM pathology. The TGF- β pathway mediates fibrosis in many different disease states including the heart (60). *LTBP4* was identified as a modifier using a genomewide scan in a mouse model of cardiomyopathy and muscular dystrophy (61), and additional loci were identified that differentially modified heart and muscle disease (62). Additional modifiers for cardiomyopathy and heart failure will emerge from genomewide surveys currently ongoing in many laboratories. Hopefully these modifiers will not only yield diagnostic information that may predict rate of progression and arrhythmia risk, but also indicate pathways to therapeutic intervention.

Conclusions

The genetic complexity of cardiomyopathy, especially DCM, is being increasingly understood with the application of massively parallel sequencing technologies. Genetic diagnosis is now readily accessible for DCM and provides useful information about prognosis and associated risks. The pathways uncovered from studies on genetic cardiomyopathy highlight the importance of the sarcomere, including the Z band, the connections to the plasma membrane, and the nuclear membrane. Modifier genes for DCM include secondary rare variants, common variants in essential pathways for heart function, and each of these affords opportunities for diagnosis and prognosis as well as therapeutic intervention.

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Address correspondence to: Elizabeth M. McNally, University of Chicago, 5841 S. Maryland MC6088, Chicago, Illinois 60637, USA. Phone: 773.702.2672; Fax: 773.702.2681; E-mail: emcnally@uchicago.edu.

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