

Genotype, phenotype: upstairs, downstairs in the family of cardiomyopathies

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Erratum

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Table 1
Molecular defects linked to human cardiomyopathies

Genomic defects	Human defects		
	HCM	DCM	RCM
Sarcomere			
Myosin heavy chain	Missense (17–19)	Missense (20)	
Myosin essential light chain	Missense (21)		
Myosin regulatory light chain	Missense (21)		
Cardiac actin	Missense (22)	Missense (3)	
Troponin-T	Missense/deletion (19, 23)	Deletion (20)	
Troponin-I	Missense (7)		Missense (6)
α-Tropomyosin	Missense (19, 23)	Missense (24)	
Myosin binding protein-C	Missense/deletion (19, 25)		
Titin/titin-related protein			
Titin	Missense (26)	Missense/deletion (27, 28)	
Telethonin (T-cap)		Missense (14)	
Z-disk-associated proteins			
MLP		Missense (14)	
Sarcolemma cytoskeleton			
Dystrophin		Deletion (29–31)	
β-Sarcoglycan		Deletion/duplication (32)	
δ-Sarcoglycan		Missense (33)	
α-Dystrobrevin		Missense (34)	
Metavinculin		Deletion (35, 36)	
Intermediate filaments			
Desmin		Missense (37, 38)	
Lamin A/C		Missense (39)	