

The Importance of Genetic Counseling, DNA Diagnostic in Left Ventricular Noncompaction Cardiomyopathy

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Left ventricular noncompaction and myocardial fibrosis: a case report. International Archive of Medicine, 2010, 3, 20.	1.2	13
2	Imaging Phenotype vs Genotype in Nonhypertrophic Heritable Cardiomyopathies. Circulation: Cardiovascular Imaging, 2010, 3, 753-765.	1.3	10
3	Mutations in the Sarcomere Gene <i>MYH7</i> in Ebstein Anomaly. Circulation: Cardiovascular Genetics, 2011, 4, 43-50.	5.1	153
4	HRS/EHRA Expert Consensus Statement on the State of Genetic Testing for the Channelopathies and Cardiomyopathies. Heart Rhythm, 2011, 8, 1308-1339.	0.3	995
5	Systematic review of pregnancy in women with inherited cardiomyopathies. European Journal of Heart Failure, 2011, 13, 584-594.	2.9	73
6	A de novo mutation in NKX2.5 associated with atrial septal defects, ventricular noncompaction, syncope and sudden death. Clinica Chimica Acta, 2011, 412, 170-175.	0.5	75
7	HRS/EHRA Expert Consensus Statement on the State of Genetic Testing for the Channelopathies and Cardiomyopathies: This document was developed as a partnership between the Heart Rhythm Society (HRS) and the European Heart Rhythm Association (EHRA). Europace, 2011, 13, 1077-1109.	0.7	699
8	Recent progress in the genetics of cardiomyopathy and its role in the clinical evaluation of patients with cardiomyopathy. Current Opinion in Cardiology, 2011, 26, 155-164.	0.8	25
10	Cardiac Magnetic Resonance of Left Ventricular Trabeculation. Circulation: Cardiovascular Imaging, 2011, 4, 84-86.	1.3	4
12	Sarcomere Gene Mutations in Isolated Left Ventricular Noncompaction Cardiomyopathy Do Not Predict Clinical Phenotype. Circulation: Cardiovascular Genetics, 2011, 4, 367-374.	5.1	167
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18	Left Ventricular Noncompaction. Circulation Journal, 2012, 76, 1556-1562.	0.7	72
19	Congenital Cardiomyopathies. , 2012, , 459-472.		0
21	Almanac 2011: Cardiomyopathies. The national society journals present selected research that has driven recent advances in clinical cardiology. Revista Portuguesa De Cardiologia (English Edition), 2012, 31, 255-261.	0.2	0
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23	Left Ventricular Noncompaction: A 25-Year Odyssey. <i>Journal of the American Society of Echocardiography</i> , 2012, 25, 363-375.	1.2	97
24	Left ventricular noncompaction in patients with β^0 -thalassemia: Uncovering a previously unrecognized abnormality. <i>American Journal of Hematology</i> , 2012, 87, 1079-1083.	2.0	23
25	No relationship between left ventricular radial wall motion and longitudinal velocity and the extent and severity of noncompaction cardiomyopathy. <i>Cardiovascular Ultrasound</i> , 2012, 10, 9.	0.5	6
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43	Isolated left ventricular non-compaction: A single-center experience. <i>Revista Portuguesa De Cardiologia (English Edition)</i> , 2013, 32, 229-238.	0.2	1
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57	Novel Mutation in Exon 14 of the Sarcomere Gene <i><sc>MYH7</sc></i> in Familial Left Ventricular Noncompaction With Bicuspid Aortic Valve. <i>Circulation: Heart Failure</i> , 2014, 7, 1059-1062.	1.6	27
58	Loss of consciousness and convulsion induced by a ventricular tachycardia mimicking epilepsy in a patient with noncompaction cardiomyopathy: a case report. <i>Netherlands Heart Journal</i> , 2014, 22, 301-303.	0.3	3
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82	A low prevalence of sarcomeric gene variants in a Chinese cohort with left ventricular non-compaction. <i>Heart and Vessels</i> , 2015, 30, 258-264.	0.5	21
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91	Human Genetics of Cardiomyopathies. , 2016, , 675-686.		1
92	Clinical Applications for Next Generation Sequencing in Cardiology. , 2016, , 189-215.		0
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94	Left ventricular noncompaction cardiomyopathy: cardiac, neuromuscular, and genetic factors. <i>Nature Reviews Cardiology</i> , 2017, 14, 224-237.	6.1	166
95	Next Generation Sequencing Based Clinical Molecular Diagnosis of Human Genetic Disorders. , 2017, , .		2
96	Application of NGS in the Diagnosis of Cardiovascular Genetic Diseases. , 2017, , 243-286.		0
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136	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , 2019, 16, e301-e372.	0.3	494

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142	Cardiac Phenotypes, Genetics, and Risks in Familial Noncompaction Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2019, 73, 1601-1611.	1.2	65
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147	<i>RBM20</i> mutations in left ventricular non-compaction cardiomyopathy. <i>Pediatric Investigation</i> , 2020, 4, 61-63.	0.6	4
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150	Left ventricular non-compaction cardiomyopathy: how many needles in the haystack?. <i>Heart</i> , 2021, 107, 1344-1352.	1.2	20
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160	Ventricular non-compaction review. Heart Failure Reviews, 2022, 27, 1063-1076.	1.7	22
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164	The Genetic Landscape of Cardiomyopathies. Cardiac and Vascular Biology, 2019, , 45-91.	0.2	20
165	Human Genetics of Ebstein Anomaly. , 2016, , 613-620.		2
167	Is Left Ventricular Noncompaction a Trait, Phenotype, or Disease?. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	15
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169	4-Dimensional light-sheet microscopy to elucidate shear stress modulation of cardiac trabeculation. Journal of Clinical Investigation, 2016, 126, 1679-1690.	3.9	100
171	Importance of genetic evaluation and testing in pediatric cardiomyopathy. World Journal of Cardiology, 2014, 6, 1156.	0.5	40
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