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## Cardiac and Neuromuscular Features of Patients With LMNA-Related Cardiomyopathy

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#	Paper	IF	Citations
26	X-Linked Emery-Dreifuss Muscular Dystrophy: Study Of X-Chromosome Inactivation and Its Relation with Clinical Phenotypes in Female Carriers. <i>Genes</i> , <b>2019</b> , 10,	4.2	4
25	Limb Girdle Muscular Dystrophies. <i>Neurologic Clinics</i> , <b>2020</b> , 38, 493-504	4.5	1
24	Genotype-phenotype correlation of LMNA variants involving the Arg541 residue: a case report with multimodality imaging and literature review. <i>ESC Heart Failure</i> , <b>2020</b> , 7, 3169-3173	3.7	1
23	The proof is in the appendage: A case report of a fenestrated membrane overlying the left atrial appendage. <i>Echocardiography</i> , <b>2020</b> , 37, 1864-1868	1.5	
22	Cardiovascular Involvement in Pediatric Laminopathies. Report of Six Patients and Literature Revision. <i>Frontiers in Pediatrics</i> , <b>2020</b> , 8, 374	3.4	2
21	The Broad Spectrum of Cardiac Diseases: From Molecular Mechanisms to Clinical Phenotype. <i>Frontiers in Physiology</i> , <b>2020</b> , 11, 761	4.6	18
20	Late gadolinium enhancement role in arrhythmic risk stratification of patients with LMNA cardiomyopathy: results from a long-term follow-up multicentre study. <i>Europace</i> , <b>2020</b> , 22, 1864-1872	3.9	7
19	Inflammation as a Predictor of Recurrent Ventricular Tachycardia After Ablation in Patients With Myocarditis. <i>Journal of the American College of Cardiology</i> , <b>2020</b> , 76, 1644-1656	15.1	11
18	Cytokine Profile in Striated Muscle Laminopathies: New Promising Biomarkers for Disease Prediction. <i>Cells</i> , <b>2020</b> , 9,	7.9	2
17	Genetic basis and molecular biology of cardiac arrhythmias in cardiomyopathies. <i>Cardiovascular Research</i> , <b>2020</b> , 116, 1600-1619	9.9	15
16	Lamin cardiomyopathy risk stratification: Authorsbreply. <i>Europace</i> , <b>2021</b> , 23, 487-488	3.9	0
15	Epidemiology of the inherited cardiomyopathies. <i>Nature Reviews Cardiology</i> , <b>2021</b> , 18, 22-36	14.8	32
14	Echocardiographic Features of Cardiomyopathy in Emery-Dreifuss Muscular Dystrophy. <i>Cardiology Research and Practice</i> , <b>2021</b> , 2021, 8812044	1.9	2
13	Cardiac Arrhythmias in Muscular Dystrophies Associated with Emerinopathy and Laminopathy: A Cohort Study. <i>Journal of Clinical Medicine</i> , <b>2021</b> , 10,	5.1	1
12	Genetics of dilated cardiomyopathy. <i>Current Opinion in Cardiology</i> , <b>2021</b> , 36, 288-294	2.1	4
11	Skeletal and Cardiac Muscle Disorders Caused by Mutations in Genes Encoding Intermediate Filament Proteins. <i>International Journal of Molecular Sciences</i> , <b>2021</b> , 22,	6.3	4
10	Biallelic mutations in RNF220 cause laminopathies featuring leukodystrophy, ataxia and deafness. <i>Brain</i> , <b>2021</b> , 144, 3020-3035	11.2	1

9	Quantitative disease risk scores from EHR with applications to clinical risk stratification and genetic studies. <i>Npj Digital Medicine</i> , <b>2021</b> , 4, 116	15.7	1
8	Importance of clinical suspicion and multidisciplinary management for early diagnosis of a cardiac laminopathy patient: A case report. <i>World Journal of Clinical Cases</i> , <b>2021</b> , 9, 7472-7477	1.6	
7	Genetic cause of heterogeneous inherited myopathies in a cohort of Greek patients. <i>Molecular Genetics and Metabolism Reports</i> , <b>2020</b> , 25, 100682	1.8	2
6	Continuous Electrical Monitoring in Patients with Arrhythmic Myocarditis: Insights from a Referral Center. <i>Journal of Clinical Medicine</i> , <b>2021</b> , 10,	5.1	1
5	Most myopathic lamin variants aggregate: a functional genomics approach for assessing variants of uncertain significance. <i>Npj Genomic Medicine</i> , <b>2021</b> , 6, 103	6.2	1
4	Clinical Profile, Arrhythmias, and Adverse Cardiac Outcomes in Emery-Dreifuss Muscular Dystrophies: A Systematic Review of the Literature.. <i>Biology</i> , <b>2022</b> , 11,	4.9	0
3	Arrhythmogenic Cardiomyopathy: One, None and a Hundred Thousand Diseases. <b>2022</b> , 12, 1256		1
2	Characterization of cardiac involvement in children with LMNA-related muscular dystrophy. 11,		0
1	Myocardial Inflammation as a Manifestation of Genetic Cardiomyopathies: From Bedside to the Bench. <b>2023</b> , 13, 646		0