

Shared genetic pathways contribute to risk of hypertrophy with opposite directions of effect

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Common genetic variants and modifiable risk factors underpin hypertrophic cardiomyopathy susceptibility and expressivity. <i>Nature Genetics</i> , 2021, 53, 135-142.	9.4	165
2	Diagnosis and Risk Prediction of Dilated Cardiomyopathy in the Era of Big Data and Genomics. <i>Journal of Clinical Medicine</i> , 2021, 10, 921.	1.0	16
3	New insights into the genetics of cardiomyopathies. <i>Nature Reviews Cardiology</i> , 2021, 18, 229-229.	6.1	1
4	Genomic Context Differs Between Human Dilated Cardiomyopathy and Hypertrophic Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2021, 10, e019944.	1.6	9
5	Genome-wide association for heart failure: from discovery to clinical use. <i>European Heart Journal</i> , 2021, 42, 2012-2014.	1.0	2
6	Editorial commentary: Genetic testing for congenital heart disease: The future is now. <i>Trends in Cardiovascular Medicine</i> , 2022, 32, 320-321.	2.3	1
7	Improving risk prediction in hypertrophic cardiomyopathy: the key role of Dutch founder variants. <i>Netherlands Heart Journal</i> , 2021, 29, 299-300.	0.3	0
8	The TRIB3 R84 variant is associated with increased left ventricular mass in a sample of 2426 White individuals. <i>Cardiovascular Diabetology</i> , 2021, 20, 115.	2.7	1
9	The Complex and Diverse Genetic Architecture of Dilated Cardiomyopathy. <i>Circulation Research</i> , 2021, 128, 1514-1532.	2.0	49
10	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. <i>Circulation</i> , 2021, 144, 7-19.	1.6	213
11	Time to Think Differently About Sarcomere-Negative Hypertrophic Cardiomyopathy. <i>Circulation</i> , 2021, 143, 2415-2417.	1.6	32
12	Understanding the genetics of adult-onset dilated cardiomyopathy: what a clinician needs to know. <i>European Heart Journal</i> , 2021, 42, 2384-2396.	1.0	28
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16	Alpha-protein kinase 3 (<i>ALPK3</i>) truncating variants are a cause of autosomal dominant hypertrophic cardiomyopathy. <i>European Heart Journal</i> , 2021, 42, 3063-3073.	1.0	51
17	Childhood Hypertrophic Cardiomyopathy: A Disease of the Cardiac Sarcomere. <i>Frontiers in Pediatrics</i> , 2021, 9, 708679.	0.9	10
18	Phenotypic variability and modifier variants in children with hereditary heart diseases. <i>Rossiyskiy Vestnik Perinatologii i Pediatrii</i> , 2021, 66, 12-19.	0.1	1

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19	ALPK3: a full spectrum cardiomyopathy gene?. <i>European Heart Journal</i> , 2021, 42, 3074-3077.	1.0	4
21	Pathogenic Mechanisms of Hypertrophic Cardiomyopathy beyond Sarcomere Dysfunction. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8933.	1.8	20
23	Phenotypic Expression and Outcomes in Individuals With Rare Genetic Variants of Hypertrophic Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2021, 78, 1097-1110.	1.2	55
24	Genetic Testing in Patients with Hypertrophic Cardiomyopathy. <i>International Journal of Molecular Sciences</i> , 2021, 22, 10401.	1.8	21
25	Minor hypertrophic cardiomyopathy genes, major insights into the genetics of cardiomyopathies. <i>Nature Reviews Cardiology</i> , 2022, 19, 151-167.	6.1	50
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31	Recent Findings Related to Cardiomyopathy and Genetics. <i>International Journal of Molecular Sciences</i> , 2021, 22, 12522.	1.8	21
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36	Predicting Sudden Cardiac Death in Genetic Heart Disease. <i>Canadian Journal of Cardiology</i> , 2022, 38, 479-490.	0.8	3
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85	Phenotypic and Genetic Factors Associated with Absence of Cardiomyopathy Symptoms in PLN:c.40_42delAGA Carriers. <i>Journal of Cardiovascular Translational Research</i> , 0, , .	1.1	2
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