

# Prevalence of Hypertrophic Cardiomyopathy in a General

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Dyskinesia Associated with Chronic Antihistamine Use. New England Journal of Medicine, 1976, 294, 113-113.	27.0	13
2	TRIGGERS FOR SUDDEN CARDIAC DEATH IN THE ATHLETE. Cardiology Clinics, 1996, 14, 195-210.	2.2	46
4	The Cardiovascular Complications of Vigorous Physical Activity. Archives of Internal Medicine, 1996, 156, 2297.	3.8	65
5	Experience from clinical genetics in hypertrophic cardiomyopathy: proposal for new diagnostic criteria in adult members of affected families.. Heart, 1997, 77, 130-132.	2.9	252
6	Training Related Left Ventricular Hypertrophy in a Soldier. Journal of the Royal Army Medical Corps, 1997, 143, 118-121.	0.8	0
7	Sudden death from cardiovascular disease in young athletes: fact or fiction?. British Journal of Sports Medicine, 1997, 31, 269-276.	6.7	41
8	Epidemiology of Idiopathic Cardiomyopathies in Children and Adolescents: A Nationwide Study in Finland. American Journal of Epidemiology, 1997, 146, 385-393.	3.4	212
9	RISK PROFILES AND CARDIOVASCULAR PREPARTICIPATION SCREENING OF COMPETITIVE ATHLETES. Cardiology Clinics, 1997, 15, 473-483.	2.2	13
10	OUTER LIMITS OF THE ATHLETE'S HEART, THE EFFECT OF GENDER, AND RELEVANCE TO THE DIFFERENTIAL DIAGNOSIS WITH PRIMARY CARDIAC DISEASES. Cardiology Clinics, 1997, 15, 381-396.	2.2	76
11	The Management of Hypertrophic Cardiomyopathy. New England Journal of Medicine, 1997, 336, 775-785.	27.0	945
12	Hypertrophic cardiomyopathy. Lancet, The, 1997, 350, 127-133.	13.7	412
13	Title is missing!. Journal of Interventional Cardiac Electrophysiology, 1997, 1, 229-232.	1.0	1
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15	Heart disease and other causes of sudden death in young athletes. Current Problems in Cardiology, 1998, 23, 477-529.	2.4	20
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18	A high risk phenotype of hypertrophic cardiomyopathy associated with a compound genotype of two mutated $\beta$ -myosin heavy chain genes. Human Genetics, 1998, 102, 299-304.	3.8	68
20	Identification of two novel mutations in the ventricular regulatory myosin light chain gene ( MYL2 ) associated with familial and classical forms of hypertrophic cardiomyopathy. Journal of Molecular Medicine, 1998, 76, 208-214.	3.9	130

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21	Genes and disease expression in hypertrophic cardiomyopathy. Lancet, The, 1998, 352, 1162-1163.	13.7	28
22	Hypertrophic cardiomyopathy. Evidence-based Cardiovascular Medicine, 1998, 2, 89-91.	0.0	1
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24	Prevention of Cardiac Hypertrophy in Mice by Calcineurin Inhibition. Science, 1998, 281, 1690-1693.	12.6	421
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50	Unusual Myocardial Hypertrophy in a Young Woman. Echocardiography, 2000, 17, 451-451.	0.9	0
51	Pearls for Practice. Recognizing Young People at Risk For Sudden Cardiac Death in Preparticipation Sports Physicals. Journal of the American Academy of Nurse Practitioners, 2000, 12, 11-14.	1.4	1
52	Molecular genetics of hypertrophic cardiomyopathy. Current Cardiology Reports, 2000, 2, 134-140.	2.9	44
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55	Athlete's heart and hypertrophic cardiomyopathy. Current Cardiology Reports, 2000, 2, 166-171.	2.9	23
56	Ventricular arrhythmias, sudden death, and prevention in patients with hypertrophic cardiomyopathy. Current Cardiology Reports, 2000, 2, 522-528.	2.9	11
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62	Reversal of Cardiac Hypertrophy in Transgenic Disease Models by Calcineurin Inhibition. Journal of Molecular and Cellular Cardiology, 2000, 32, 697-709.	1.9	69
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67	Nonsurgical septal reduction therapy for hypertrophic obstructive cardiomyopathy: one-year follow-up. Journal of the American College of Cardiology, 2000, 36, 852-855.	2.8	135
68	Spectrum of clinical phenotypes and gene variants in cardiac myosin-binding protein C mutation carriers with hypertrophic cardiomyopathy. Journal of the American College of Cardiology, 2001, 38, 322-330.	2.8	125
69	Beta-myosin Heavy Chain Gene Mutations and Hypertrophic Cardiomyopathy in Austrian Children. Journal of Molecular and Cellular Cardiology, 2001, 33, 141-148.	1.9	18
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80	Molecular Pathophysiology of Cardiomyopathies. , 2001, , 1045-1063.		2
81	Use of implantable pacemakers and implantable defibrillators in hypertrophic cardiomyopathy. Current Opinion in Cardiology, 2001, 16, 58-65.	1.8	9
82	Arrhythmia substrate and management in hypertrophic cardiomyopathy: from molecules to implantable card ioverter-defibrillators. European Heart Journal Supplements, 2001, 3, L15-L20.	0.1	4
83	Percutaneous transluminal septal myocardial ablation: A new catheter-based therapy for patients with hypertrophic obstructive cardiomyopathy. Heart Lung and Circulation, 2001, 10, 63-67.	0.4	0
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87	Lack of Tacrolimus-Induced Cardiomyopathy. Annals of Pharmacotherapy, 2001, 35, 985-989.	1.9	21
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95	The need for European Registries in inherited cardiomyopathies. European Heart Journal, 2002, 23, 1972-1974.	2.2	3
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98	Molecular diagnosis of myocardial disease. <i>Expert Review of Molecular Diagnostics</i> , 2002, 2, 587-602.	3.1	18
99	Hypertrophic cardiomyopathy: management, risk stratification, and prevention of sudden death. <i>British Heart Journal</i> , 2002, 87, 169-176.	2.1	183
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105	Two mutations in troponin I that cause hypertrophic cardiomyopathy have contrasting effects on cardiac muscle contractility. <i>Biochemical Journal</i> , 2002, 362, 443.	3.7	27
106	Epidemiologic and Clinical Characteristics of Cardiomyopathies in Japan. Results From Nationwide Surveys.. <i>Circulation Journal</i> , 2002, 66, 323-336.	1.6	76
107	Clinical profile of stroke in 900 patients with hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2002, 39, 301-307.	2.8	329
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111	The prevention of sudden death in hypertrophic cardiomyopathy. <i>Expert Opinion on Pharmacotherapy</i> , 2002, 3, 499-504.	1.8	4
112	Genetic Causes of Inherited Cardiac Hypertrophy: Robert L. Frye Lecture. <i>Mayo Clinic Proceedings</i> , 2002, 77, 1315-1319.	3.0	16
113	Microarray gene expression profiles in dilated and hypertrophic cardiomyopathic end-stage heart failure. <i>Physiological Genomics</i> , 2002, 10, 31-44.	2.3	220
114	A redução do gradiente na via de saída do ventrículo esquerdo pelo marcapasso DDD em pacientes com miocardiopatia hipertrófica obstrutiva. <i>Brazilian Journal of Cardiovascular Surgery</i> , 2002, 17, 248.	0.6	0
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129	Fabry disease: a functional and anatomical study of cardiac manifestations in 20 hemizygous male patients. Clinical Genetics, 2003, 63, 46-52.	2.0	46
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131	Molecular Genetics of Arrhythmias and Cardiovascular Conditions Associated with Arrhythmias. PACE - Pacing and Clinical Electrophysiology, 2003, 26, 2194-2208.	1.2	15
132	Hypertrophic cardiomyopathy: from gene defect to clinical disease. Cell Research, 2003, 13, 9-20.	12.0	66
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137	Cardiovascular Disease. New England Journal of Medicine, 2003, 349, 60-72.	27.0	382
138	On predictors of sudden cardiac death in hypertrophic cardiomyopathy**Editorials published in the Journal of the American College of Cardiologyreflect the views of the authors and do not necessarily represent the views of JACC or the American College of Cardiology.. Journal of the American College of Cardiology, 2003, 41, 994-996.	2.8	27
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146	Comparison of fluorescent SSCP and denaturing HPLC analysis with direct sequencing for mutation screening in hypertrophic cardiomyopathy. Journal of Medical Genetics, 2003, 40, 59e-59.	3.2	40
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154	Familial Hypertrophic Cardiomyopathy-linked Alterations in Ca <sup>2+</sup> Binding of Human Cardiac Myosin Regulatory Light Chain Affect Cardiac Muscle Contraction. Journal of Biological Chemistry, 2004, 279, 3535-3542.	3.4	65

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156	The pathology of hypertrophic cardiomyopathy. Histopathology, 2004, 44, 412-427.	2.9	236
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159	Cellular and molecular aspects of familial hypertrophic cardiomyopathy caused by mutations in the cardiac troponin I gene. Molecular and Cellular Biochemistry, 2004, 263, 99-114.	3.1	60
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161	Decreased interactions of mutant muscle LIM protein (MLP) with N-RAP and $\beta$ -actinin and their implication for hypertrophic cardiomyopathy. Cell and Tissue Research, 2004, 317, 129-136.	2.9	56
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164	Cardiovascular disease in athletes. Clinics in Sports Medicine, 2004, 23, 455-471.	1.8	5
165	Molecular genetic basis of sudden cardiac death. Pediatric Clinics of North America, 2004, 51, 1229-1255.	1.8	26
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168	Risk of sudden cardiac death in young athletes: which screening strategies are appropriate?. Pediatric Clinics of North America, 2004, 51, 1421-1441.	1.8	20
169	Hypertrophic Cardiomyopathy: From "Heart Tumour" to a Complex Molecular Genetic Disorder. Heart Lung and Circulation, 2004, 13, 15-25.	0.4	17
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171	Comprehensive Analysis of the Beta-Myosin Heavy Chain Gene in 389 Unrelated Patients With Hypertrophic Cardiomyopathy. Journal of the American College of Cardiology, 2004, 44, 602-610.	2.8	178
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176	Prevalence of idiopathic hypertrophic cardiomyopathy in China: a population-based echocardiographic analysis of 8080 adults. American Journal of Medicine, 2004, 116, 14-18.	1.5	225
177	Hypertrophic cardiomyopathy: an important global disease. American Journal of Medicine, 2004, 116, 63-65.	1.5	157
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179	Subaortic myectomy for patients with hypertrophic obstructive cardiomyopathy. Operative Techniques in Thoracic and Cardiovascular Surgery, 2004, 9, 254-260.	0.3	2
180	Hypertrophic cardiomyopathy. Lancet, The, 2004, 363, 1881-1891.	13.7	558
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184	Parasympathetic dysfunction in hypertrophic cardiomyopathy assessed by heart rate variability: comparison between short-term and 24-h measurements. Clinical Physiology and Functional Imaging, 2005, 25, 90-99.	1.2	21
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188	Cardiomyopathie hypertrophique. EMC - Cardiologie-Angéiologie, 2005, 2, 103-119.	0.8	3
189	Sudden Cardiac Death in Young Athletes. Clinical Pediatric Emergency Medicine, 2005, 6, 194-199.	0.4	12
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195	Weaning failure from mechanical ventilation due to hypertrophic obstructive cardiomyopathy. <i>Intensive Care Medicine</i> , 2005, 31, 734-737.	8.2	24
196	Elevated serum markers for collagen synthesis in patients with hypertrophic cardiomyopathy and diastolic dysfunction. <i>Clinical Research in Cardiology</i> , 2005, 94, 328-335.	1.1	31
197	Array lessons from the heart: focus on the genome and transcriptome of cardiomyopathies. <i>Physiological Genomics</i> , 2005, 21, 131-143.	2.3	34
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899	Population Burden of Sudden Death Associated With Hypertrophic Cardiomyopathy. <i>Circulation</i> , 2017, 136, 1665-1667.	1.6	29
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901	Relationship of cardiac troponin to systolic global longitudinal strain in hypertrophic cardiomyopathy. <i>Echocardiography</i> , 2017, 34, 1470-1477.	0.9	11
902	Hypertrophic Cardiomyopathy. <i>Circulation Research</i> , 2017, 121, 749-770.	4.5	790
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1079	A Contraction Stress Model of Hypertrophic Cardiomyopathy due to Sarcomere Mutations. <i>Stem Cell Reports</i> , 2019, 12, 71-83.	4.8	82
1080	Fibrotic Signaling in Cardiomyopathies. <i>Molecular and Translational Medicine</i> , 2019, , 273-317.	0.4	1
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1082	Catecholamine response to exercise in patients with non-obstructive hypertrophic cardiomyopathy. <i>Journal of Physiology</i> , 2019, 597, 1337-1346.	2.9	18
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1088	Surgical myectomy versus alcohol septal ablation for obstructive hypertrophic cardiomyopathy: A propensity score-matched cohort. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2019, 157, 306-315.e3.	0.8	77
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1090	QT prolongation and sudden cardiac death risk in hypertrophic cardiomyopathy. <i>Acta Cardiologica</i> , 2019, 74, 53-58.	0.9	27
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1161	Risk of arrhythmic events after alcohol septal ablation for hypertrophic cardiomyopathy using continuous implantable cardiac monitoring. <i>Heart Rhythm</i> , 2021, 18, 50-56.	0.7	7
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1169	Prevalence and Clinical Correlates of Aortic Dilation in Hypertrophic Cardiomyopathy. <i>Journal of the American Society of Echocardiography</i> , 2021, 34, 279-285.	2.8	6
1170	Cardiovascular genetics: the role of genetic testing in diagnosis and management of patients with hypertrophic cardiomyopathy. <i>Heart</i> , 2021, 107, 183-189.	2.9	12
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1172	Implantable cardiac monitors in low risk hypertrophic cardiomyopathy: To protect and serve, or observe and report?. <i>Journal of Cardiovascular Electrophysiology</i> , 2021, 32, 136-137.	1.7	0
1173	Thromboembolism in Patients with Hypertrophic Cardiomyopathy. <i>International Journal of Medical Sciences</i> , 2021, 18, 727-735.	2.5	5
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1218	Molecular Genetic Basis of Hypertrophic Cardiomyopathy. <i>Circulation Research</i> , 2021, 128, 1533-1553.	4.5	88
1219	Transgenic rabbit models for cardiac disease research. <i>British Journal of Pharmacology</i> , 2022, 179, 938-957.	5.4	13
1221	Mavacamten, a Novel Therapeutic Strategy for Obstructive Hypertrophic Cardiomyopathy. <i>Current Cardiology Reports</i> , 2021, 23, 79.	2.9	17
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1235	The function of LncRNA-H19 in cardiac hypertrophy. <i>Cell and Bioscience</i> , 2021, 11, 153.	4.8	17

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