

Patterns of inheritance in hypertrophic cardiomyopathy two-dimensional echocardiography

American Journal of Cardiology

53, 1087-1094

DOI: [10.1016/0002-9149\(84\)90643-x](https://doi.org/10.1016/0002-9149(84)90643-x)

Citation Report

#	ARTICLE	IF	CITATIONS
1	Hypertrophic cardiomyopathy compatible with successful completion of the marathon. American Journal of Cardiology, 1984, 53, 1470-1471.	1.6	13
2	Hypertrophic cardiomyopathy. The importance of the site and the extent of hypertrophy. A review. Progress in Cardiovascular Diseases, 1985, 28, 1-83.	3.1	751
3	Apical hypertrophic cardiomyopathy: evaluation by noninvasive and invasive techniques in 23 patients.. Circulation, 1985, 71, 45-56.	1.6	87
4	Task force III: Hypertrophic cardiomyopathy, other myopericardial diseases and mitral valve prolapse. Journal of the American College of Cardiology, 1985, 6, 1215-1217.	2.8	36
5	Unusual distribution of left ventricular hypertrophy in obstructive hypertrophic cardiomyopathy: Localized posterobasal free wall thickening in two patients. Journal of the American College of Cardiology, 1985, 5, 1474-1477.	2.8	15
6	Asymmetry in hypertrophic cardiomyopathy: The septal to free wall thickness ratio revisited. American Journal of Cardiology, 1985, 55, 835-838.	1.6	39
7	Hypertrophic cardiomyopathy with extreme increase in left ventricular wall thickness: Functional and morphologic features and clinical significance. Journal of the American College of Cardiology, 1986, 8, 57-65.	2.8	93
8	Structural features of the athlete heart as defined by echocardiography. Journal of the American College of Cardiology, 1986, 7, 190-203.	2.8	491
9	Causes of sudden death in competitive athletes. Journal of the American College of Cardiology, 1986, 7, 204-214.	2.8	528
10	Sudden death and the competitive athlete: Perspectives on preparticipation screening studies. Journal of the American College of Cardiology, 1986, 7, 220-230.	2.8	135
11	Human lymphocyte antigens in hypertrophic cardiomyopathy. International Journal of Cardiology, 1986, 12, 193-202.	1.7	5
12	The Genetics of Hypertrophic Cardiomyopathy. Annals of Internal Medicine, 1986, 105, 610.	3.9	37
13	Clinical significance and therapeutic implications of the left ventricular outflow tract pressure gradient in hypertrophic cardiomyopathy. American Journal of Cardiology, 1986, 58, 1093-1096.	1.6	64
14	Familial spontaneous complete heart block in hypertrophic cardiomyopathy.. Heart, 1986, 55, 469-474.	2.9	21
15	Development and Progression of Left Ventricular Hypertrophy in Children with Hypertrophic Cardiomyopathy. New England Journal of Medicine, 1986, 315, 610-614.	27.0	320
16	Hypertrophic cardiomyopathy in three generations of a large Norwegian family. A clinical, echocardiographic, and genetic study.. Heart, 1986, 55, 168-175.	2.9	9
17	Inheritance of hypertrophic cardiomyopathy: a cross sectional and M mode echocardiographic study of 50 families.. Heart, 1987, 58, 259-266.	2.9	76
18	Hypertrophic Cardiomyopathy. New England Journal of Medicine, 1987, 316, 780-789.	27.0	819

#	ARTICLE	IF	CITATIONS
19	Southwestern Internal Medicine Conference: Hypertrophic Cardiomyopathy: Current Views on Etiology, Pathophysiology, and Management. American Journal of the Medical Sciences, 1987, 294, 191-210.	1.1	11
20	Sudden cardiac death. Human Pathology, 1987, 18, 485-492.	2.0	38
21	Cardiomyopathies. Human Pathology, 1987, 18, 625-635.	2.0	33
22	HYPERTROPHIC CARDIOMYOPATHY ASSOCIATED WITH HEREDITARY SPHEROCYTOSIS IN THREE GENERATIONS OF ONE FAMILY. Lancet, The, 1987, 330, 853-854.	13.7	6
23	Bibliography of biomedical ultrasound 1984. Ultrasound in Medicine and Biology, 1987, 13, 803-942.	1.5	0
24	Results of screening a large group of intercollegiate competitive athletes for cardiovascular disease. Journal of the American College of Cardiology, 1987, 10, 1214-1221.	2.8	184
25	Apical Hypertrophic Cardiomyopathy: Clinical and Two-Dimensional Echocardiographic Assessment. Annals of Internal Medicine, 1987, 106, 663.	3.9	99
26	Dominantly inherited dilated cardiomyopathy. American Journal of Medical Genetics Part A, 1987, 27, 61-73.	2.4	18
27	Familial apical hypertrophic cardiomyopathy. American Journal of Cardiology, 1988, 62, 821-822.	1.6	15
28	Valvular aortic stenosis and asymmetric septal hypertrophy: diagnostic considerations and clinical and therapeutic implications. European Heart Journal, 1988, 9, 71-76.	2.2	44
29	Spontaneously occurring hypertrophic cardiomyopathy in the rat. II. Distribution of, and correlations between, various cardiac abnormalities in the WKY/NCrj and its related strains.. Japanese Circulation Journal, 1988, 52, 1156-1170.	1.0	9
31	Equivocal and Borderline Myocardial Hypertrophy in Relatives of Patients with Hypertrophic Cardiomyopathy: Possible Implications in Genetics of the Disease. Cardiology, 1988, 75, 348-356.	1.4	5
32	Mapping a Gene for Familial Hypertrophic Cardiomyopathy to Chromosome 14q1. New England Journal of Medicine, 1989, 321, 1372-1378.	27.0	511
33	Calcium-Antagonist Receptors in the Atrial Tissue of Patients with Hypertrophic Cardiomyopathy. New England Journal of Medicine, 1989, 320, 755-761.	27.0	99
34	Relation between extent of left ventricular hypertrophy and age in hypertrophic cardiomyopathy. Journal of the American College of Cardiology, 1989, 13, 820-823.	2.8	59
35	Long-term outcome of patients with hypertrophic cardiomyopathy successfully resuscitated after cardiac arrest. Journal of the American College of Cardiology, 1989, 13, 1283-1288.	2.8	127
36	Clinical Course and Prognosis of Hypertrophic Cardiomyopathy in an Outpatient Population. New England Journal of Medicine, 1989, 320, 749-755.	27.0	318
37	Morphological quantification and differentiation of left ventricular hypertrophy in hypertrophic cardiomyopathy and hypertensive heart disease A two dimensional echocardiographic stud. European Heart Journal, 1990, 11, 65-74.	2.2	12

#	ARTICLE	IF	CITATIONS
38	Genetic evidence of dissociation (generational skips) of electrical from morphologic forms of hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1990, 66, 627-631.	1.6	21
39	Diversity of patterns of hypertrophy in patients with systemic hypertension and marked left ventricular wall thickening. <i>American Journal of Cardiology</i> , 1990, 65, 874-881.	1.6	51
40	Preseason cardiovascular examination: A review. <i>Journal of Adolescent Health Care: Official Publication of the Society for Adolescent Medicine</i> , 1990, 11, 379-386.	0.3	1
41	Two-dimensional electrophoresis of heart muscle proteins in human cardiomyopathies. <i>Electrophoresis</i> , 1990, 11, 333-336.	2.4	13
42	Cosegregation of hypertrophic cardiomyopathy and a fragile site on chromosome 16 in a large Italian family.. <i>Journal of Medical Genetics</i> , 1990, 27, 363-366.	3.2	11
43	Hypertrophic cardiomyopathy: one disease or several?. <i>Heart</i> , 1990, 63, 263-264.	2.9	14
44	A molecular basis for familial hypertrophic cardiomyopathy: An β cardiac myosin heavy chain hybrid gene. <i>Cell</i> , 1990, 62, 991-998.	28.9	236
45	Apical hypertrophic cardiomyopathy: The continuing saga. <i>Journal of the American College of Cardiology</i> , 1990, 15, 91-93.	2.8	43
46	Hypertrophic cardiomyopathy characterized by marked hypertrophy of the posterior left ventricular free wall: Significance and clinical implications. <i>Journal of the American College of Cardiology</i> , 1991, 18, 421-428.	2.8	24
47	Simultaneous occurrence of mitral valve prolapse and systolic anterior motion in hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1991, 67, 404-410.	1.6	12
48	Prognosis of nonobstructive hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1991, 67, 215-217.	1.6	11
49	The genetics of hypertrophic cardiomyopathy.. <i>Heart</i> , 1991, 66, 193-195.	2.9	7
50	Hereditary Transmission of Tetralogy of Fallot, Cardiac Hypertrophy, and Anomalies of Great Vessels in WKY/NCrj Rats. <i>Pediatric Research</i> , 1991, 30, 227-230.	2.3	5
51	Anomalous insertion of papillary muscle directly into anterior mitral leaflet in hypertrophic cardiomyopathy. Significance in producing left ventricular outflow obstruction.. <i>Circulation</i> , 1991, 84, 1188-1197.	1.6	229
52	Progress in familial hypertrophic cardiomyopathy: molecular genetic analyses in the original family studied by Teare.. <i>Heart</i> , 1992, 67, 34-38.	2.9	24
53	Usefulness of Doppler echocardiographic assessment of diastolic filling in distinguishing "athlete's heart" from hypertrophic cardiomyopathy. <i>Heart</i> , 1992, 68, 296-300.	2.9	118
55	Diversity of structural mitral valve alterations in hypertrophic cardiomyopathy.. <i>Circulation</i> , 1992, 85, 1651-1660.	1.6	310
56	Evidence of genetic heterogeneity in five kindreds with familial hypertrophic cardiomyopathy.. <i>Circulation</i> , 1992, 85, 635-647.	1.6	59

#	ARTICLE	IF	CITATIONS
57	Idiopathic Hypertrophic Cardiomyopathy in Identical Twins. <i>Chest</i> , 1992, 102, 783-785.	0.8	21
58	Pathology of cardiomyopathies in childhood. <i>Progress in Pediatric Cardiology</i> , 1992, 1, 8-39.	0.4	6
60	Utility of continuous wave doppler echocardiography in the noninvasive assessment of left ventricular outflow tract pressure gradient in patients with hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 1992, 19, 91-99.	2.8	178
61	Novel missense mutation in cardiac β myosin heavy chain gene found in a japanese patient with hypertrophic cardiomyopathy. <i>Biochemical and Biophysical Research Communications</i> , 1992, 188, 379-387.	2.1	47
62	The genetic basis of hypertrophic cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 1992, 24, 1471-1477.	1.9	6
63	Outer Limits of Physiologic Hypertrophy and Relevance to the Diagnosis of Primary Cardiac Disease. <i>Cardiology Clinics</i> , 1992, 10, 267-279.	2.2	11
64	No evidence for linkage of familial hypertrophic cardiomyopathy and chromosome 14q1 locus D14S26 in a chinese family: evidence for genetic heterogeneity. <i>Human Genetics</i> , 1992, 89, 597-601.	3.8	15
65	Clinical course of middle-aged asymptomatic patients with hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1992, 69, 935-940.	1.6	21
66	Two brothers with unexplained cardiomegaly. <i>Trends in Cardiovascular Medicine</i> , 1992, 2, 2-5.	4.9	0
67	The prevalence of hypertrophic cardiomyopathy in men: an echocardiographic population screening study with a review of death records. <i>Journal of Internal Medicine</i> , 1992, 232, 499-506.	6.0	10
68	Mapping of a novel gene for familial hypertrophic cardiomyopathy to chromosome 11. <i>Nature Genetics</i> , 1993, 4, 311-313.	21.4	184
69	Hypertrophic cardiomyopathy. <i>Current Problems in Cardiology</i> , 1993, 18, 641-704.	2.4	53
70	Morphologic basis for obstruction to right ventricular outflow in hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1993, 71, 1089-1094.	1.6	48
71	Impact of patient selection biases on the perception of hypertrophic cardiomyopathy and its natural history. <i>American Journal of Cardiology</i> , 1993, 72, 970-972.	1.6	112
72	Genetic heterogeneity of familial hypertrophic cardiomyopathy. <i>Neuromuscular Disorders</i> , 1993, 3, 483-486.	0.6	9
73	Coexistence of sudden cardiac death and end-stage heart failure in familial hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 1993, 22, 489-497.	2.8	76
74	Identification of a mutation in the beta cardiac myosin heavy chain gene in a family with hypertrophic cardiomyopathy.. <i>Heart</i> , 1993, 69, 136-141.	2.9	24
75	Reduction in left ventricular wall thickness after deconditioning in highly trained Olympic athletes.. <i>Heart</i> , 1993, 69, 125-128.	2.9	155

#	ARTICLE	IF	CITATIONS
76	Inherited Cardiomyopathies. <i>New England Journal of Medicine</i> , 1994, 330, 913-919.	27.0	295
77	Multiple disease genes cause hypertrophic cardiomyopathy. <i>Heart</i> , 1994, 72, S4-S9.	2.9	34
78	Risk factors and stratification for sudden cardiac death in patients with hypertrophic cardiomyopathy. <i>Heart</i> , 1994, 72, S13-S18.	2.9	49
79	Natural history of hypertrophic cardiomyopathy. <i>Heart</i> , 1994, 72, S10-S12.	2.9	43
80	Surviving competitive athletics with hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , 1994, 73, 1098-1104.	1.6	55
81	Clinical and morphologic expression of hypertrophic cardiomyopathy in patients \geq 65 years of age. <i>American Journal of Cardiology</i> , 1994, 73, 1105-1111.	1.6	54
82	Hypertrophic cardiomyopathy. <i>Progress in Cardiovascular Diseases</i> , 1994, 36, 275-308.	3.1	81
83	Task force 3: Hypertrophic cardiomyopathy, myocarditis and other myopericardial diseases and mitral valve prolapse. <i>Journal of the American College of Cardiology</i> , 1994, 24, 880-885.	2.8	113
85	Chronic congestive heart failure. <i>European Heart Journal</i> , 1994, 15, 328-334.	2.2	3
86	The Genetic Basis of Paediatric Heart Disease. <i>Annals of Medicine</i> , 1995, 27, 289-300.	3.8	23
87	Long-term evaluation of patients with apical hypertrophic cardiomyopathy. <i>European Heart Journal</i> , 1995, 16, 210-217.	2.2	43
88	Sudden cardiac death in patients with hypertrophic cardiomyopathy: From bench to bedside with an emphasis on genetic markers. <i>Clinical Cardiology</i> , 1995, 18, 189-198.	1.8	33
89	Analysis of randomness of atrial and ventricular rhythm in atrial fibrillation. <i>European Heart Journal</i> , 1995, 16, 971-976.	2.2	10
90	The COX8 gene is not the disease gene of the CMH4 locus in familial hypertrophic cardiomyopathy.. <i>Journal of Medical Genetics</i> , 1995, 32, 670-671.	3.2	1
91	Doppler Echocardiography in Familial Hypertrophic Cardiomyopathy. <i>Echocardiography</i> , 1995, 12, 235-241.	0.9	14
92	Hypertrophic cardiomyopathy in tuscany: Clinical course and outcome in an unselected regional population. <i>Journal of the American College of Cardiology</i> , 1995, 26, 1529-1536.	2.8	265
93	Phenotypic spectrum and patterns of left ventricular hypertrophy in hypertrophic cardiomyopathy: Morphologic observations and significance as assessed by two-dimensional echocardiography in 600 patients. <i>Journal of the American College of Cardiology</i> , 1995, 26, 1699-1708.	2.8	594
94	Syncope in Athletes. <i>Sports Medicine</i> , 1995, 19, 223-234.	6.5	21

#	ARTICLE	IF	CITATIONS
95	GÃ©nÃ©tique molÃ©culaire des cardiomyopathies. Annales De L'Institut Pasteur / ActualitÃ©s, 1996, 7, 199-2030.1		0
96	Mapping the locus for familial hypertrophic cardiomyopathy to chromosome 11 in a family with a case of apical hypertrophic cardiomyopathy of the Japanese type. Human Genetics, 1996, 97, 457-461.	3.8	2
97	Malignant familial hypertrophic cardiomyopathy in a family with a 453Argâ†’Cys mutation in the Î²-myosin heavy chain gene: Coexistence of sudden death and end-stage heart failure. Human Genetics, 1996, 97, 585-590.	3.8	19
98	Familial hypertrophic cardiomyopathy: diagnostic and therapeutic implications of recent genetic studies. Trends in Molecular Medicine, 1996, 2, 387-393.	2.6	9
99	TRIGGERS FOR SUDDEN CARDIAC DEATH IN THE ATHLETE. Cardiology Clinics, 1996, 14, 195-210.	2.2	46
100	HLA-DR2 Antigen Linkage in Patients with Apical Hypertrophic Cardiomyopathy in Japan. Cardiology, 1996, 87, 488-491.	1.4	5
101	Molecular pathology of dilated cardiomyopathies. Current Problems in Cardiology, 1996, 21, 99-144.	2.4	10
102	Myocardial beta adrenoceptors and left ventricular function in hypertrophic cardiomyopathy.. Heart, 1996, 75, 50-54.	2.9	50
103	Clinical profile and prognosis of hypertrophic cardiomyopathy when first diagnosed in infancy as opposed to childhood. Cardiology in the Young, 1997, 7, 410-416.	0.8	3
104	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
105	THE PREPARTICIPATION SPORTS EXAMINATION FOR HIGH SCHOOL AND COLLEGE ATHLETES. Clinics in Sports Medicine, 1997, 16, 569-591.	1.8	21
106	SYNCOPE IN THE PEDIATRIC PATIENT. Cardiology Clinics, 1997, 15, 277-294.	2.2	33
107	ACC/AHA Guidelines for the Clinical Application of Echocardiography: Executive Summary. Journal of the American College of Cardiology, 1997, 29, 862-879.	2.8	402
108	Differences in Myocardial Velocity Gradient Measured Throughout the Cardiac Cycle in Patients With Hypertrophic Cardiomyopathy, Athletes and Patients With Left Ventricular Hypertrophy Due to Hypertension. Journal of the American College of Cardiology, 1997, 30, 760-768.	2.8	204
109	The natural history of left ventricular wall thickening in hypertrophic cardiomyopathy. Australian and New Zealand Journal of Medicine, 1997, 27, 51-58.	0.5	25
110	Molecular mechanisms regulating the myofilament response to Ca ²⁺ : Implications of mutations causal for familial hypertrophic cardiomyopathy. Basic Research in Cardiology, 1997, 92, 63-74.	5.9	40
111	Editorial. American Journal of Cardiology, 1998, 81, 1339-1344.	1.6	159
112	Heart disease and other causes of sudden death in young athletes. Current Problems in Cardiology, 1998, 23, 477-529.	2.4	20

#	ARTICLE	IF	CITATIONS
113	Hamartoma of mature cardiac myocytes*1, *2. <i>Human Pathology</i> , 1998, 29, 904-909.	2.0	68
114	Familial Hypertrophic Cardiomyopathy. <i>Circulation Research</i> , 1998, 83, 580-593.	4.5	354
115	Coexistence of mitochondrial DNA and beta Âmyosin heavy chain mutations in hypertrophic cardiomyopathy with late congestive heart failure. <i>Heart</i> , 1998, 80, 548-558.	2.9	65
116	Expression of Proto-oncogenes and Gene Mutation of Sarcomeric Proteins in Patients With Hypertrophic Cardiomyopathy. <i>Circulation Research</i> , 1998, 83, 594-601.	4.5	87
117	Impact of Laboratory Molecular Diagnosis on Contemporary Diagnostic Criteria for Genetically Transmitted Cardiovascular Diseases: Hypertrophic Cardiomyopathy, Long-QT Syndrome, and Marfan Syndrome. <i>Circulation</i> , 1998, 98, 1460-1471.	1.6	128
118	Echocardiographic pitfalls in the diagnosis of hypertrophic cardiomyopathy. <i>Heart</i> , 1999, 82, 8iii-15.	2.9	37
119	Resolution of Neonatal Hypertrophic Cardiomyopathy in an Infant with an Affected Mother. <i>Pediatric Cardiology</i> , 1999, 20, 208-211.	1.3	7
120	The Inheritance of Hypertrophic Cardiomyopathy. <i>Pediatric Cardiology</i> , 1999, 20, 313-316.	1.3	24
121	Echocardiographic Diagnosis of Congenital Heart Disease: An Embryologic and Anatomic Approach. <i>Pediatric Cardiology</i> , 1999, 20, 316-316.	1.3	1
122	Persistent ST segment elevation: A new ECG finding in hypertrophic cardiomyopathy. <i>American Journal of Emergency Medicine</i> , 1999, 17, 296-299.	1.6	16
123	PEDIATRIC MYOCARDIAL DISEASE. <i>Pediatric Clinics of North America</i> , 1999, 46, 289-312.	1.8	69
124	Prognostic value of systemic blood pressure response during exercise in a community-based patient population with hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 1999, 33, 2044-2051.	2.8	230
125	Genetic aspects of heart failure. <i>European Journal of Heart Failure</i> , 1999, 1, 121-126.	7.1	17
126	Hypertrophic cardiomyopathy in a litter of five mixed-breed cats. <i>Journal of the American Animal Hospital Association</i> , 1999, 35, 293-296.	1.1	27
127	Situs inversus with hypertrophic cardiomyopathy in identical twins. , 2000, 91, 327-330.		14
128	Diagnosis of hypertrophic cardiomyopathy and screening for the phenotype suggestive of gene carriage in familial disease: a simple echocardiographic procedure. <i>Journal of Medical Screening</i> , 2000, 7, 82-90.	2.3	10
129	Deletion in the Cardiac Troponin I Gene in a Family From Northern Sweden with Hypertrophic Cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 2000, 32, 521-525.	1.9	35
130	Inherited and de novo Mutations in the Cardiac Actin Gene Cause Hypertrophic Cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 2000, 32, 1687-1694.	1.9	200

#	ARTICLE	IF	CITATIONS
131	A malignant phenotype of hypertrophic cardiomyopathy caused by Arg719Gln cardiac beta-myosin heavy-chain mutation in a Chinese family. <i>Clinica Chimica Acta</i> , 2001, 310, 131-139.	1.1	10
132	Cardiomyopathies: from genetics to the prospect of treatment. <i>Lancet, The</i> , 2001, 358, 1627-1637.	13.7	159
133	Clinical and Echocardiographic Features of Hypertrophic Cardiomyopathy in the Elderly. <i>The American Journal of Geriatric Cardiology</i> , 2001, 10, 11-19.	0.6	1
134	Genes and their polymorphisms in mono- and multifactorial cardiomyopathies. <i>Pharmacogenomics</i> , 2002, 3, 367-378.	1.3	12
135	Hypertrophic Cardiomyopathy. <i>JAMA - Journal of the American Medical Association</i> , 2002, 287, 1308-20.	7.4	1,981
136	Clinical features of hypertrophic cardiomyopathy in the young. <i>Cardiology in the Young</i> , 2002, 12, 147-152.	0.8	10
137	Hypertrophic Cardiomyopathy with Shared Morphology in Identical Twins: A Case Report. <i>Scottish Medical Journal</i> , 2002, 47, 64-65.	1.3	4
138	Sarcomere Protein Gene Mutations in Hypertrophic Cardiomyopathy of the Elderly. <i>Circulation</i> , 2002, 105, 446-451.	1.6	311
139	Hypertension, left ventricular hypertrophy, and sudden death. <i>Current Cardiology Reports</i> , 2002, 4, 449-457.	2.9	48
140	Hypertrophic cardiomyopathy: state-of-the-art review, with focus on the management of outflow obstruction. <i>Internal Medicine Journal</i> , 2003, 33, 521-529.	0.8	18
141	Identification of the genotypes causing hypertrophic cardiomyopathy in northern Sweden. <i>Journal of Molecular and Cellular Cardiology</i> , 2003, 35, 841-849.	1.9	96
142	Hypertrophic Cardiomyopathy: Low Frequency of Mutations in the β -Myosin Heavy Chain (MYH7) and Cardiac Troponin T (TNNT2) Genes among Spanish Patients. <i>Clinical Chemistry</i> , 2003, 49, 1279-1285.	3.2	62
143	Sudden Cardiac Death in Athletes. <i>Cardiology</i> , 2003, 100, 186-195.	1.4	11
144	Familial aggregation of genetically heterogeneous hypertrophic cardiomyopathy: A boy with LEOPARD syndrome due to PTPN11 mutation and his nonsyndromic father lacking PTPN11 mutations. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2004, 70, 95-98.	1.6	14
145	Proposal for contemporary screening strategies in families with hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2004, 44, 2125-2132.	2.8	176
146	The Usual Causes of Left Ventricular Outflow Tract Obstruction Below the Aortic Valve in Normal Ventriculoarterial Connection: Review of the Physiopathology and Surgical Implications. <i>Acta Chirurgica Belgica</i> , 2005, 105, 475-481.	0.4	6
147	Amyloid heart disease mimicking hypertrophic cardiomyopathy*. <i>Journal of Internal Medicine</i> , 2005, 258, 225-230.	6.0	47
148	Cardiomyopathie hypertrophique. <i>EMC - Cardiologie-Angeiologie</i> , 2005, 2, 103-119.	0.8	3

#	ARTICLE	IF	CITATIONS
149	Cardiac Function Assessment in Patients with Family History of Nonhypertrophic Cardiomyopathy: A Prenatal and Postnatal Study. <i>Pediatric Cardiology</i> , 2005, 26, 543-552.	1.3	16
150	Array lessons from the heart: focus on the genome and transcriptome of cardiomyopathies. <i>Physiological Genomics</i> , 2005, 21, 131-143.	2.3	34
151	Adult Echocardiography and Doppler. <i>Journal of Diagnostic Medical Sonography</i> , 2005, 21, 91-110.	0.3	1
152	Recent advances in genetics and treatment of hypertrophic cardiomyopathy. <i>Future Cardiology</i> , 2005, 1, 341-353.	1.2	4
153	Benign outcome in a long-term follow-up of patients with hypertrophic cardiomyopathy in Brazil. <i>American Heart Journal</i> , 2005, 149, 1099-1105.	2.7	31
155	Large-scale mutation screening in patients with dilated or hypertrophic cardiomyopathy: a pilot study using DGGE. <i>Journal of Molecular Medicine</i> , 2006, 84, 682-691.	3.9	31
156	Molecular genetics in hypertrophic cardiomyopathy: towards individualized management of the disease. <i>Expert Review of Molecular Diagnostics</i> , 2006, 6, 65-78.	3.1	24
158	Barry Joel Maron, MD: A Conversation With the Editor. This series of interviews was underwritten by an unrestricted grant from Bristol-Myers Squibb.. <i>American Journal of Cardiology</i> , 2007, 99, 1334-1349.	1.6	1
159	The genetics of cardiomyopathies: What clinicians should know. <i>Current Heart Failure Reports</i> , 2007, 4, 229-235.	3.3	2
160	Value of Real Time Three-Dimensional Echocardiography in Patients with Hypertrophic Cardiomyopathy: Comparison with Two-Dimensional Echocardiography and Magnetic Resonance Imaging. <i>Echocardiography</i> , 2008, 25, 717-726.	0.9	62
161	The ubiquitin-proteasome system in cardiac dysfunction. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2008, 1782, 749-763.	3.8	129
162	Chapter 18 Clinical genetic issues in stroke. <i>Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn</i> , 2008, 92, 355-372.	1.8	0
163	Imaging studies in patients with heart failure: Current and evolving technologies. <i>Critical Care Medicine</i> , 2008, 36, S28-S39.	0.9	2
164	Hypertrophic cardiomyopathy family with double-heterozygous mutations; does disease severity suggest doubleheterozygosity?. <i>Netherlands Heart Journal</i> , 2009, 17, 458-463.	0.8	16
165	Hypertrophic cardiomyopathy family with double-heterozygous mutations; does disease severity suggest double-heterozygosity?. <i>Netherlands Heart Journal</i> , 2010, , 1.	0.8	0
166	Hypertrophic cardiomyopathy: from genetics to treatment. <i>European Journal of Clinical Investigation</i> , 2010, 40, 360-369.	3.4	99
167	Clinical Approach to Sudden Cardiac Death Syndromes. , 2010, , .		5
168	Increased left ventricular torsion in hypertrophic cardiomyopathy mutation carriers with normal wall thickness. <i>Journal of Cardiovascular Magnetic Resonance</i> , 2011, 13, 3.	3.3	60

#	ARTICLE	IF	CITATIONS
169	Cellular mechanisms of cardiomyopathy. <i>Journal of Cell Biology</i> , 2011, 194, 355-365.	5.2	308
170	Assessing the knowledge of sudden unexpected death in the young among Canadian medical students and recent graduates: a cross-sectional study. <i>BMJ Open</i> , 2012, 2, e001798.	1.9	5
171	Transcriptional Regulation of Cardiac Genes Balance Pro- and Anti-Hypertrophic Mechanisms in Hypertrophic Cardiomyopathy. <i>Neurology International</i> , 2012, 2, e5.	0.5	1
172	Mendelian Forms of Structural Cardiovascular Disease. <i>Current Cardiology Reports</i> , 2013, 15, 399.	2.9	4
173	Familial Hypertrophic Cardiomyopathy: New Insight on Mode of Inheritance among Egyptian Children. <i>Journal of Clinical & Experimental Cardiology</i> , 2014, 05, .	0.0	0
174	Advances in medical treatment of hypertrophic cardiomyopathy. <i>Journal of Cardiology</i> , 2014, 64, 1-10.	1.9	31
175	Mechanistic Heterogeneity in Contractile Properties of β -Tropomyosin (TPM1) Mutants Associated with Inherited Cardiomyopathies. <i>Journal of Biological Chemistry</i> , 2015, 290, 7003-7015.	3.4	41
176	The Remarkable 50 Years of Imaging in HCM and How It Has Changed Diagnosis and Management. <i>JACC: Cardiovascular Imaging</i> , 2016, 9, 858-872.	5.3	43
177	Midterm postoperative follow-up after surgical correction of hypertrophic cardiomyopathy in infancy and childhood. <i>Journal of the Egyptian Society of Cardio-Thoracic Surgery</i> , 2017, 25, 133-141.	0.2	0
178	Septal alcoholization in hypertrophic cardiomyopathy: about 11 cases. <i>Pan African Medical Journal</i> , 2017, 27, 196.	0.8	2
179	Hypertrophic Cardiomyopathy in South Western Nigeria. <i>SA Heart Journal</i> , 2017, 6, .	0.0	1
181	Pathology and Pathophysiology. , 2019, , 23-39.		1
182	Thromboembolism in Patients with Hypertrophic Cardiomyopathy. <i>International Journal of Medical Sciences</i> , 2021, 18, 727-735.	2.5	5
183	The Cardiomyopathies. , 1994, , 196-222.		5
185	Hypertrophische Kardiomyopathie (HCM). <i>Spezielle Pathologische Anatomie</i> , 2000, , 1055-1140.	0.0	1
186	The Molecular Genetics of Familial Hypertrophic Cardiomyopathy. , 1993, , 289-305.		1
187	Evolution of Left Ventricular Hypertrophy in Patients with Hypertrophic Cardiomyopathy. , 1990, , 7-24.		3
188	The Radiologic Evaluation of Chest Pain in the Athlete. <i>Clinics in Sports Medicine</i> , 1987, 6, 845-870.	1.8	2

#	ARTICLE	IF	CITATIONS
189	Aortic/Mitral Obstruction and Coarctation of the Aorta. <i>Cardiology Clinics</i> , 1993, 11, 617-642.	2.2	11
190	Molecular Basis of Familial Cardiomyopathies. <i>Circulation</i> , 1995, 91, 532-540.	1.6	140
191	Cardiac Disease in Young Trained Athletes. <i>Circulation</i> , 1995, 91, 1596-1601.	1.6	332
192	Prevalence of Hypertrophic Cardiomyopathy and Limitations of Screening Methods. <i>Circulation</i> , 1995, 92, 700-704.	1.6	42
193	Prevalence of Hypertrophic Cardiomyopathy in a General Population of Young Adults. <i>Circulation</i> , 1995, 92, 785-789.	1.6	1,753
194	Recent Advances in the Molecular Genetics of Hypertrophic Cardiomyopathy. <i>Circulation</i> , 1995, 92, 1336-1347.	1.6	162
195	Clinical Approach to Genetic Cardiomyopathy in Children. <i>Circulation</i> , 1996, 94, 2021-2038.	1.6	138
196	ACC/AHA Guidelines for the Clinical Application of Echocardiography. <i>Circulation</i> , 1997, 95, 1686-1744.	1.6	513
197	Diagnostic Value of Electrocardiography and Echocardiography for Familial Hypertrophic Cardiomyopathy in a Genotyped Adult Population. <i>Circulation</i> , 1997, 96, 214-219.	1.6	143
198	Sporadic hypertrophic cardiomyopathy due to de novo myosin mutations.. <i>Journal of Clinical Investigation</i> , 1992, 90, 1666-1671.	8.2	120
199	Familial hypertrophic cardiomyopathy. Microsatellite haplotyping and identification of a hot spot for mutations in the beta-myosin heavy chain gene.. <i>Journal of Clinical Investigation</i> , 1993, 92, 2807-2813.	8.2	78
202	Medical Imaging. , 0, , 634-712.		2
205	Miocardiopatía hipertrófica: a propósito de un caso. <i>Medifam - Revista De Medicina Familiar Y Comunitaria</i> , 2003, 13, .	0.0	0
206	Familial Hypertrophic Cardiomyopathy With Triphasic Transmitral Flow Velocity. <i>Journal of Echocardiography</i> , 2006, 4, 37-42.	0.8	0
207	Evaluation of Myocardial Disease in the Cardiac Catheterization Laboratory. , 2007, , 1349-1357.		0
208	Echocardiography in the Evaluation of the Cardiomyopathies. , 2007, , 1359-1378.		0
209	A rare presentation of hypertrophic cardiomyopathy in a neonate. <i>Sri Lanka Journal of Child Health</i> , 2007, 36, 114.	0.1	0
210	Hypertrophic Cardiomyopathy In A Patient With Craniofacial Syndrom: New Cardiocranial Syndrome?. <i>The Internet Journal of Pediatrics and Neonatology</i> , 2011, 13, .	0.0	0

#	ARTICLE	IF	CITATIONS
211	Hypertrophische obstruktive Kardiomyopathie (HOCM). , 1989, , 70-109.		3
212	Preventive cardiology. , 1989, , 345-353.		0
213	Hypertrophische nicht obstruktive Kardiomyopathie (HNCM). , 1989, , 110-129.		0
214	Latente Kardiomyopathie (LCM). , 1989, , 130-155.		1
215	Cytogenetic Studies in Familial Hypertrophic Cardiomyopathy: Identification of a Fragile Site on Human Chromosome 16. , 1990, , 97-102.		0
216	Left Ventricular Systolic and Diastolic Function in Hypertrophic Cardiomyopathy. , 1990, , 32-37.		0
218	Clinical Investigation: Current Approaches. , 1993, , 94-107.		0
220	Methods for Distinguishing Athlete's Heart from Structural Heart Disease, with Emphasis on Hypertrophic Cardiomyopathy. , 1997, , 108-114.		0
221	Mitochondrial DNA Mutations and Heart Disease. , 1998, , 239-263.		1
222	Molecular mechanisms regulating the myofilament response to Ca ²⁺ : Implications of mutations causal for familial hypertrophic cardiomyopathy. , 1998, , 105-121.		0
224	Pathology and Pathophysiology. , 2015, , 23-38.		0
226	Ventricular dysfunction in hypertrophic obstructive cardiomyopathy. Texas Heart Institute Journal, 1991, 18, 165-9.	0.3	0
227	Molecular basis of hypertrophic and dilated cardiomyopathy. Texas Heart Institute Journal, 1994, 21, 6-15.	0.3	17
228	Myocardial diseases of animals. American Journal of Pathology, 1986, 124, 98-178.	3.8	106
229	Contemporary treatment of hypertrophic cardiomyopathy. Texas Heart Institute Journal, 2009, 36, 194-204.	0.3	37
232	Malignant familial hypertrophic cardiomyopathy in a family with a 453Arg→Cys mutation in the β -myosin heavy chain gene: coexistence of sudden death and end-stage heart failure. Human Genetics, 1996, 97, 585-590.	3.8	0
233	Novel MYBPC3 Mutations in Indian Population with Cardiomyopathies. Pharmacogenomics and Personalized Medicine, 0, Volume 16, 883-893.	0.7	0