

# Jodie Ingles

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

146  
papers

4,745  
citations

35  
h-index

66  
g-index

182  
ext. papers

6,504  
ext. citations

5.5  
avg, IF

5.78  
L-index

#	Paper	IF	Citations
146	A prospective longitudinal study of health-related quality of life and psychological wellbeing after an implantable cardioverter-defibrillator in patients with genetic heart diseases.. <i>Heart Rhythm O2</i> , <b>2022</b> , 3, 143-151	1.5	0
145	Whole genome sequencing delineates regulatory, copy number, and cryptic splice variants in early onset cardiomyopathy.. <i>Npj Genomic Medicine</i> , <b>2022</b> , 7, 18	6.2	0
144	Associations Between Female Sex, Sarcomere Variants, and Clinical Outcomes in Hypertrophic Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , <b>2021</b> , 14, e003062	5.2	10
143	External validation of the HCM Risk-Kids model for predicting sudden cardiac death in childhood hypertrophic cardiomyopathy. <i>European Journal of Preventive Cardiology</i> , <b>2021</b> ,	3.9	5
142	Clinical characteristics and outcomes in childhood-onset hypertrophic cardiomyopathy. <i>European Heart Journal</i> , <b>2021</b> , 42, 1988-1996	9.5	20
141	Computational prediction of protein subdomain stability in MYBPC3 enables clinical risk stratification in hypertrophic cardiomyopathy and enhances variant interpretation. <i>Genetics in Medicine</i> , <b>2021</b> , 23, 1281-1287	8.1	3
140	Clinical Profile and Health Disparities in a Multiethnic Cohort of Patients With Hypertrophic Cardiomyopathy. <i>Circulation: Heart Failure</i> , <b>2021</b> , 14, e007537	7.6	2
139	Transcriptome Sequencing of Patients With Hypertrophic Cardiomyopathy Reveals Novel Splice-Altering Variants in. <i>Circulation Genomic and Precision Medicine</i> , <b>2021</b> , 14, e003202	5.2	5
138	Essentials of a new clinical practice guidance on familial hypercholesterolaemia for physicians. <i>Internal Medicine Journal</i> , <b>2021</b> , 51, 769-779	1.6	0
137	Investigation of current models of care for genetic heart disease in Australia: A national clinical audit. <i>International Journal of Cardiology</i> , <b>2021</b> , 330, 128-134	3.2	1
136	Harmonizing the Collection of Clinical Data on Genetic Testing Requisition Forms to Enhance Variant Interpretation in Hypertrophic Cardiomyopathy (HCM): A Study from the ClinGen Cardiomyopathy Variant Curation Expert Panel. <i>Journal of Molecular Diagnostics</i> , <b>2021</b> , 23, 589-598	5.1	1
135	Precision Medicine in Cardiovascular Disease: Genetics and Impact on Phenotypes: JACC Focus Seminar 1/5. <i>Journal of the American College of Cardiology</i> , <b>2021</b> , 77, 2517-2530	15.1	2
134	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. <i>Circulation</i> , <b>2021</b> , 144, 7-19	16.7	34
133	Synopsis of an integrated guidance for enhancing the care of familial hypercholesterolaemia: an Australian perspective. <i>American Journal of Preventive Cardiology</i> , <b>2021</b> , 6, 100151	1.9	0
132	Genetic variants associated with inherited cardiovascular disorders among 13,131 asymptomatic older adults of European descent. <i>Npj Genomic Medicine</i> , <b>2021</b> , 6, 51	6.2	3
131	The genetic architecture of Plakophilin 2 cardiomyopathy. <i>Genetics in Medicine</i> , <b>2021</b> , 23, 1961-1968	8.1	2
130	"Concealed cardiomyopathy" as a cause of previously unexplained sudden cardiac arrest. <i>International Journal of Cardiology</i> , <b>2021</b> , 324, 96-101	3.2	6

129	Characterization of clinically relevant copy-number variants from exomes of patients with inherited heart disease and unexplained sudden cardiac death. <i>Genetics in Medicine</i> , <b>2021</b> , 23, 86-93	8.1	4
128	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. <i>Heart Rhythm</i> , <b>2021</b> , 18, e1-e50	6.7	37
127	Disease-specific variant pathogenicity prediction significantly improves variant interpretation in inherited cardiac conditions. <i>Genetics in Medicine</i> , <b>2021</b> , 23, 69-79	8.1	4
126	The End Unexplained Cardiac Death (EndUCD) Registry for Young Australian Sudden Cardiac Arrest. <i>Heart Lung and Circulation</i> , <b>2021</b> , 30, 714-720	1.8	2
125	Integrated Guidance for Enhancing the Care of Familial Hypercholesterolaemia in Australia. <i>Heart Lung and Circulation</i> , <b>2021</b> , 30, 324-349	1.8	20
124	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. <i>Journal of Arrhythmia</i> , <b>2021</b> , 37, 481-534	1.5	3
123	Discordant clinical features of identical hypertrophic cardiomyopathy twins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2021</b> , 118,	11.5	5
122	Sex Disparities in Sudden Cardiac Death. <i>Circulation: Arrhythmia and Electrophysiology</i> , <b>2021</b> , 14, e009834	6.4	1
121	Worldwide differences in primary prevention implantable cardioverter defibrillator utilization and outcomes in hypertrophic cardiomyopathy. <i>European Heart Journal</i> , <b>2021</b> , 42, 3932-3944	9.5	6
120	Sex Differences in Hypertrophic Cardiomyopathy: Interaction With Genetics and Environment. <i>Current Heart Failure Reports</i> , <b>2021</b> , 18, 264-273	2.8	2
119	Arrhythmia in Cardiomyopathy: Sex and Gender Differences. <i>Current Heart Failure Reports</i> , <b>2021</b> , 18, 274-283	2.8	0
118	Global approaches to cardiogenetic evaluation after sudden cardiac death in the young: A survey among health care professionals. <i>Heart Rhythm</i> , <b>2021</b> , 18, 1637-1644	6.7	2
117	Inherited cardiomyopathies <b>2021</b> , 277-290		
116	Hypertrophic Cardiomyopathy: Genetic Testing and Risk Stratification. <i>Current Cardiology Reports</i> , <b>2021</b> , 23, 9	4.2	4
115	What do we do and how do we do it? Assessing genetic counselling in the modern era. <i>European Journal of Human Genetics</i> , <b>2020</b> , 28, 1137-1138	5.3	1
114	Psychological Issues in Managing Families with Inherited Cardiovascular Diseases. <i>Cold Spring Harbor Perspectives in Medicine</i> , <b>2020</b> , 10,	5.4	7
113	A Validated Model for Sudden Cardiac Death Risk Prediction in Pediatric Hypertrophic Cardiomyopathy. <i>Circulation</i> , <b>2020</b> , 142, 217-229	16.7	51
112	Epidemiology and clinical characteristics of atrial fibrillation in patients with inherited heart diseases. <i>Journal of Cardiovascular Electrophysiology</i> , <b>2020</b> , 31, 465-473	2.7	4

111 Clinical Genetics **2020**, 27-50

110	Patients With Genetic Heart Disease and COVID-19: A Cardiac Society of Australia and New Zealand (CSANZ) Consensus Statement. <i>Heart Lung and Circulation</i> , <b>2020</b> , 29, e85-e87	1.8	2
109	Prevalence of Anderson-Fabry disease in a cohort with unexplained late gadolinium enhancement on cardiac MRI. <i>International Journal of Cardiology</i> , <b>2020</b> , 304, 122-124	3.2	2
108	How Patient Perceptions Shape Responses and Outcomes in Inherited Cardiac Conditions. <i>Heart Lung and Circulation</i> , <b>2020</b> , 29, 641-652	1.8	2
107	Genetic Testing in Inherited Heart Diseases. <i>Heart Lung and Circulation</i> , <b>2020</b> , 29, 505-511	1.8	14
106	Association of Race With Disease Expression and Clinical Outcomes Among Patients With Hypertrophic Cardiomyopathy. <i>JAMA Cardiology</i> , <b>2020</b> , 5, 83-91	16.2	23
105	Doctor-patient care relationship in genetic cardiomyopathies: An exploratory study on clinical consultations. <i>PLoS ONE</i> , <b>2020</b> , 15, e0236814	3.7	1
104	An International Multicenter Evaluation of Inheritance Patterns, Arrhythmic Risks, and Underlying Mechanisms of -Catecholaminergic Polymorphic Ventricular Tachycardia. <i>Circulation</i> , <b>2020</b> , 142, 932-947	16.7	12
103	Needs analysis of parents following sudden cardiac death in the young. <i>Open Heart</i> , <b>2020</b> , 7,	3	6
102	Genetic architecture of left ventricular noncompaction in adults. <i>Human Genome Variation</i> , <b>2020</b> , 7, 33	1.8	10
101	Spatial and Functional Distribution of Pathogenic Variants and Clinical Outcomes in Patients With Hypertrophic Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , <b>2020</b> , 13, 396-405	5.2	19
100	Hypertrophic Cardiomyopathy With Left Ventricular Systolic Dysfunction: Insights From the SHaRe Registry. <i>Circulation</i> , <b>2020</b> , 141, 1371-1383	16.7	43
99	Evaluating the Clinical Validity of Hypertrophic Cardiomyopathy Genes. <i>Circulation Genomic and Precision Medicine</i> , <b>2019</b> , 12, e002460	5.2	132
98	Implantable cardioverter-defibrillators in previously undiagnosed patients with catecholaminergic polymorphic ventricular tachycardia resuscitated from sudden cardiac arrest. <i>European Heart Journal</i> , <b>2019</b> , 40, 2953-2961	9.5	53
97	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy. <i>Heart Rhythm</i> , <b>2019</b> , 16, e301-e372	6.7	247
96	Perceptions of genetic variant reclassification in patients with inherited cardiac disease. <i>European Journal of Human Genetics</i> , <b>2019</b> , 27, 1134-1142	5.3	17
95	Revisiting Genome Sequencing Data in Light of Novel Disease Gene Associations. <i>Journal of the American College of Cardiology</i> , <b>2019</b> , 73, 1365-1366	15.1	5
94	Regional Variation in RBM20 Causes a Highly Penetrant Arrhythmogenic Cardiomyopathy. <i>Circulation: Heart Failure</i> , <b>2019</b> , 12, e005371	7.6	51

93	Participation in thrill-seeking activities by patients with hypertrophic cardiomyopathy: Individual preferences, adverse events and physician attitude. <i>American Heart Journal</i> , <b>2019</b> , 214, 28-35	4.9	1
92	Response by Ingles and Semsarian to Letter Regarding Article, "Concealed Arrhythmogenic Right Ventricular Cardiomyopathy in Sudden Unexplained Cardiac Death Events". <i>Circulation Genomic and Precision Medicine</i> , <b>2019</b> , 12, e002475	5.2	2
91	Evaluating a custom-designed aid to improve communication of genetic results in families with hypertrophic cardiomyopathy: study protocol for a randomised controlled trial. <i>BMJ Open</i> , <b>2019</b> , 9, e026627	3.2	1
90	Genome sequencing as a first-line genetic test in familial dilated cardiomyopathy. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 650-662	8.1	36
89	2019 HRS expert consensus statement on evaluation, risk stratification, and management of arrhythmogenic cardiomyopathy: Executive summary. <i>Heart Rhythm</i> , <b>2019</b> , 16, e373-e407	6.7	73
88	A balanced translocation disrupting SCN5A in a family with Brugada syndrome and sudden cardiac death. <i>Heart Rhythm</i> , <b>2019</b> , 16, 231-238	6.7	8
87	Key Value of RNA Analysis of MYBPC3 Splice-Site Variants in Hypertrophic Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , <b>2019</b> , 12, e002368	5.2	18
86	Psychological adaptation to molecular autopsy findings following sudden cardiac death in the young. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 1452-1456	8.1	11
85	Genetics of HCM and Role of Genetic Testing <b>2019</b> , 83-91		0
84	Response to Brodehl et al. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 1248-1249	8.1	
83	Psychosocial Impact of a Positive Gene Result for Asymptomatic Relatives at Risk of Hypertrophic Cardiomyopathy. <i>Journal of Genetic Counseling</i> , <b>2018</b> , 27, 1040-1048	2.5	10
82	Daily Step Count as a Simple Marker of Disease Severity in Hypertrophic Cardiomyopathy. <i>Heart Lung and Circulation</i> , <b>2018</b> , 27, 752-755	1.8	2
81	Psychosocial Implications of Living with Catecholaminergic Polymorphic Ventricular Tachycardia in Adulthood. <i>Journal of Genetic Counseling</i> , <b>2018</b> , 27, 549-557	2.5	14
80	Adaptation and validation of the ACMG/AMP variant classification framework for MYH7-associated inherited cardiomyopathies: recommendations by ClinGen's Inherited Cardiomyopathy Expert Panel. <i>Genetics in Medicine</i> , <b>2018</b> , 20, 351-359	8.1	173
79	Rare desmin variant causing penetrant life-threatening arrhythmic cardiomyopathy. <i>HeartRhythm Case Reports</i> , <b>2018</b> , 4, 318-323	1	7
78	Genetic Testing for Cardiomyopathies in Clinical Practice. <i>Heart Failure Clinics</i> , <b>2018</b> , 14, 129-137	3.3	8
77	Genetic Testing <b>2018</b> , 668-673		
76	Whole Genome Sequencing Improves Outcomes of Genetic Testing in Patients With Hypertrophic Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , <b>2018</b> , 72, 419-429	15.1	92

75	A Control Theory-Based Pilot Intervention to Increase Physical Activity in Patients With Hypertrophic Cardiomyopathy. <i>American Journal of Cardiology</i> , <b>2018</b> , 122, 866-871	3	8
74	Communication of genetic information to families with inherited rhythm disorders. <i>Heart Rhythm</i> , <b>2018</b> , 15, 780-786	6.7	21
73	Implantable cardioverter-defibrillator therapy in Australia, 2002-2015. <i>Medical Journal of Australia</i> , <b>2018</b> , 209, 123-129	4	10
72	Concealed Arrhythmogenic Right Ventricular Cardiomyopathy in Sudden Unexplained Cardiac Death Events. <i>Circulation Genomic and Precision Medicine</i> , <b>2018</b> , 11, e002355	5.2	21
71	Lack of genotype-phenotype correlation in Brugada Syndrome and Sudden Arrhythmic Death Syndrome families with reported pathogenic SCN1B variants. <i>Heart Rhythm</i> , <b>2018</b> , 15, 1051-1057	6.7	10
70	Relations between right ventricular morphology and clinical, electrical and genetic parameters in Brugada Syndrome. <i>PLoS ONE</i> , <b>2018</b> , 13, e0195594	3.7	16
69	Temporal Trends in Sudden Cardiac Death From 1997 to 2010: A Data Linkage Study. <i>Heart Lung and Circulation</i> , <b>2017</b> , 26, 808-816	1.8	15
68	Cardiovascular Effects of Energy Drinks in Familial Long QT Syndrome: A Randomized Cross-Over Study. <i>International Journal of Cardiology</i> , <b>2017</b> , 231, 150-154	3.2	23
67	Nonfamilial Hypertrophic Cardiomyopathy: Prevalence, Natural History, and Clinical Implications. <i>Circulation: Cardiovascular Genetics</i> , <b>2017</b> , 10,		75
66	Attitudes, knowledge and consequences of uncertain genetic findings in hypertrophic cardiomyopathy. <i>European Journal of Human Genetics</i> , <b>2017</b> , 25, 809-815	5.3	19
65	Burden of Recurrent and Ancestral Mutations in Families With Hypertrophic Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , <b>2017</b> , 10,		11
64	Psychosocial care and cardiac genetic counseling following sudden cardiac death in the young. <i>Progress in Pediatric Cardiology</i> , <b>2017</b> , 45, 31-36	0.4	8
63	LAMP2 shines a light on cardiomyopathy in an athlete. <i>Heart Rhythm Case Reports</i> , <b>2017</b> , 3, 172-176	1	2
62	Exome sequencing-based molecular autopsy of formalin-fixed paraffin-embedded tissue after sudden death. <i>Genetics in Medicine</i> , <b>2017</b> , 19, 1127-1133	8.1	18
61	A clinical approach to genetic testing for non-specialists. <i>BMJ, The</i> , <b>2017</b> , 358, j4101	5.9	1
60	Development of a communication aid for explaining hypertrophic cardiomyopathy genetic test results. <i>Pilot and Feasibility Studies</i> , <b>2017</b> , 3, 53	1.9	6
59	Multiple Gene Variants in Hypertrophic Cardiomyopathy in the Era of Next-Generation Sequencing. <i>Circulation: Cardiovascular Genetics</i> , <b>2017</b> , 10,		45
58	Role of Genetic Testing in Inherited Cardiovascular Disease: A Review. <i>JAMA Cardiology</i> , <b>2017</b> , 2, 1153-1160	16.0	45

57	Impact of the implantable cardioverter defibrillator on confidence to undertake physical activity in inherited heart disease: A cross-sectional study. <i>European Journal of Cardiovascular Nursing</i> , <b>2017</b> , 16, 742-752	3.3	5
56	Minding the Genes: a Multidisciplinary Approach towards Genetic Assessment of Cardiovascular Disease. <i>Journal of Genetic Counseling</i> , <b>2017</b> , 26, 224-231	2.5	14
55	Routinely collected health data to study inherited heart disease: a systematic review (2000-2016). <i>Open Heart</i> , <b>2017</b> , 4, e000686	3	2
54	A Prospective Study of Sudden Cardiac Death among Children and Young Adults. <i>New England Journal of Medicine</i> , <b>2016</b> , 374, 2441-52	59.2	396
53	Clinical and genetic features of Australian families with long QT syndrome: A registry-based study. <i>Journal of Arrhythmia</i> , <b>2016</b> , 32, 456-461	1.5	7
52	Interdisciplinary psychosocial care for families with inherited cardiovascular diseases. <i>Trends in Cardiovascular Medicine</i> , <b>2016</b> , 26, 647-53	6.9	38
51	Posttraumatic Stress and Prolonged Grief After the Sudden Cardiac Death of a Young Relative. <i>JAMA Internal Medicine</i> , <b>2016</b> , 176, 402-5	11.5	33
50	Clinical Genetics <b>2016</b> , 25-47		
49	Factors influencing uptake of familial long QT syndrome genetic testing. <i>American Journal of Medical Genetics, Part A</i> , <b>2016</b> , 170A, 418-425	2.5	31
48	Physical activity in hypertrophic cardiomyopathy: prevalence of inactivity and perceived barriers. <i>Open Heart</i> , <b>2016</b> , 3, e000484	3	26
47	Sudden deaths during the largest community running event in Australia: A 25-year review. <i>International Journal of Cardiology</i> , <b>2016</b> , 203, 1029-31	3.2	4
46	Molecular autopsy in victims of inherited arrhythmias. <i>Journal of Arrhythmia</i> , <b>2016</b> , 32, 359-365	1.5	26
45	A novel heterozygous mutation in cardiac caldesmon causes autosomal dominant catecholaminergic polymorphic ventricular tachycardia. <i>Heart Rhythm</i> , <b>2016</b> , 13, 1652-60	6.7	43
44	Exome sequencing identifies a novel mutation in the gene in a family with early-onset sinus node dysfunction, ventricular arrhythmias, and cardiac arrest. <i>HeartRhythm Case Reports</i> , <b>2015</b> , 1, 141-145	1	7
43	Letter by Semsarian and Ingles regarding article, "A randomized trial of social media from Circulation". <i>Circulation</i> , <b>2015</b> , 131, e394	16.7	1
42	Social determinants of health in the setting of hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , <b>2015</b> , 184, 743-749	3.2	14
41	New perspectives on the prevalence of hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , <b>2015</b> , 65, 1249-1254	15.1	557
40	Genetics of HCM and Role of Genetic Testing <b>2015</b> , 77-84		1



39	Rarity and phenotypic heterogeneity provide challenges in the diagnosis of Andersen-Tawil syndrome: Two cases presenting with ECGs mimicking catecholaminergic polymorphic ventricular tachycardia (CPVT). <i>International Journal of Cardiology</i> , <b>2015</b> , 201, 473-5	3.2	6
38	Reply: What Is the True Prevalence of Hypertrophic Cardiomyopathy?. <i>Journal of the American College of Cardiology</i> , <b>2015</b> , 66, 1846-1847	15.1	2
37	NOS1AP Polymorphisms Modify QTc Interval Duration But Not Cardiac Arrest Risk in Hypertrophic Cardiomyopathy. <i>Journal of Cardiovascular Electrophysiology</i> , <b>2015</b> , 26, 1346-51	2.7	3
36	Sudden cardiac death in the young: the molecular autopsy and a practical approach to surviving relatives. <i>European Heart Journal</i> , <b>2015</b> , 36, 1290-6	9.5	144
35	Sudden cardiac death rates in an Australian population: a data linkage study. <i>Australian Health Review</i> , <b>2015</b> , 39, 561-567	1.8	4
34	Application of Genetic Testing in Hypertrophic Cardiomyopathy for Preclinical Disease Detection. <i>Circulation: Cardiovascular Genetics</i> , <b>2015</b> , 8, 852-9		35
33	Challenges of exercise recommendations and sports participation in genetic heart disease patients. <i>Circulation: Cardiovascular Genetics</i> , <b>2015</b> , 8, 178-86		14
32	Determining pathogenicity of genetic variants in hypertrophic cardiomyopathy: importance of periodic reassessment. <i>Genetics in Medicine</i> , <b>2014</b> , 16, 286-93	8.1	60
31	Delay to diagnosis amongst patients with catecholaminergic polymorphic ventricular tachycardia. <i>International Journal of Cardiology</i> , <b>2014</b> , 176, 1402-4	3.2	18
30	The value of cardiac genetic testing. <i>Trends in Cardiovascular Medicine</i> , <b>2014</b> , 24, 217-24	6.9	29
29	Conveying a probabilistic genetic test result to families with an inherited heart disease. <i>Heart Rhythm</i> , <b>2014</b> , 11, 1073-8	6.7	41
28	Genetic Testing <b>2014</b> , 685-690		
27	Clinical predictors of genetic testing outcomes in hypertrophic cardiomyopathy. <i>Genetics in Medicine</i> , <b>2013</b> , 15, 972-7	8.1	79
26	Prolongation of the QTc interval predicts appropriate implantable cardioverter-defibrillator therapies in hypertrophic cardiomyopathy. <i>JACC: Heart Failure</i> , <b>2013</b> , 1, 149-55	7.9	34
25	Psychological wellbeing and posttraumatic stress associated with implantable cardioverter defibrillator therapy in young adults with genetic heart disease. <i>International Journal of Cardiology</i> , <b>2013</b> , 168, 3779-84	3.2	65
24	The Australian Genetic Heart Disease Registry. <i>International Journal of Cardiology</i> , <b>2013</b> , 168, e127-8	3.2	17
23	Tweeting our way to cardiovascular health. <i>Journal of the American College of Cardiology</i> , <b>2013</b> , 61, 1657-8	15.1	24
22	Homozygous mutation in the cardiac troponin I gene: clinical heterogeneity in hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , <b>2013</b> , 168, 1530-1	3.2	9



21	Health status of cardiac genetic disease patients and their at-risk relatives. <i>International Journal of Cardiology</i> , <b>2013</b> , 165, 448-53	3.2	35
20	Poor psychological wellbeing particularly in mothers following sudden cardiac death in the young. <i>European Journal of Cardiovascular Nursing</i> , <b>2013</b> , 12, 484-91	3.3	22
19	Genetic testing for inherited heart diseases: longitudinal impact on health-related quality of life. <i>Genetics in Medicine</i> , <b>2012</b> , 14, 749-752	8.1	19
18	A cost-effectiveness model of genetic testing for the evaluation of families with hypertrophic cardiomyopathy. <i>Heart</i> , <b>2012</b> , 98, 625-30	5.1	83
17	Guidelines for genetic testing of inherited cardiac disorders. <i>Heart Lung and Circulation</i> , <b>2011</b> , 20, 681-7	1.8	56
16	Natural history of genotype positive-phenotype negative patients with hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , <b>2011</b> , 152, 258-9	3.2	51
15	The emerging role of the cardiac genetic counselor. <i>Heart Rhythm</i> , <b>2011</b> , 8, 1958-62	6.7	82
14	Molecular diagnostics of cardiomyopathies: the future is here. <i>Circulation: Cardiovascular Genetics</i> , <b>2011</b> , 4, 103-4		13
13	Mutations in alpha-actinin-2 cause hypertrophic cardiomyopathy: a genome-wide analysis. <i>Journal of the American College of Cardiology</i> , <b>2010</b> , 55, 1127-35	15.1	137
12	Establishment of an Australian National Genetic Heart Disease Registry. <i>Heart Lung and Circulation</i> , <b>2008</b> , 17, 463-7	1.8	10
11	Postmortem molecular analysis of KCNQ1 and SCN5A genes in sudden unexplained death in young Australians. <i>International Journal of Cardiology</i> , <b>2008</b> , 127, 138-41	3.2	35
10	Sex hormone receptor gene variation associated with phenotype in male hypertrophic cardiomyopathy patients. <i>Journal of Molecular and Cellular Cardiology</i> , <b>2008</b> , 45, 217-22	5.8	39
9	Psychosocial impact of specialized cardiac genetic clinics for hypertrophic cardiomyopathy. <i>Genetics in Medicine</i> , <b>2008</b> , 10, 117-20	8.1	79
8	Sudden cardiac death in the young: a clinical genetic approach. <i>Internal Medicine Journal</i> , <b>2007</b> , 37, 32-7	1.6	66
7	Genetic screening of calcium regulation genes in familial hypertrophic cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , <b>2007</b> , 43, 337-43	5.8	61
6	Mutation analysis of the natriuretic peptide precursor B (NPPB) gene in patients with hypertrophic cardiomyopathy. <i>DNA Sequence</i> , <b>2006</b> , 17, 392-5		1
5	Cardiac troponin I mutations in Australian families with hypertrophic cardiomyopathy: clinical, genetic and functional consequences. <i>Journal of Molecular and Cellular Cardiology</i> , <b>2005</b> , 38, 387-93	5.8	37
4	Long-term follow-up of patients with obstructive hypertrophic cardiomyopathy treated with dual-chamber pacing. <i>American Journal of Cardiology</i> , <b>2005</b> , 95, 991-3	3	34

3	Compound and double mutations in patients with hypertrophic cardiomyopathy: implications for genetic testing and counselling. <i>Journal of Medical Genetics</i> , <b>2005</b> , 42, e59	5.8	270
2	Long-term follow-up of implantable cardioverter defibrillator therapy for hypertrophic cardiomyopathy. <i>American Journal of Cardiology</i> , <b>2004</b> , 93, 1192-4	3	63
1	Progression of left ventricular hypertrophy and the angiotensin-converting enzyme gene polymorphism in hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , <b>2004</b> , 96, 157-63	3.2	33