

Ray E Hershberger

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.

59 papers	5,404 citations	31 h-index	66 g-index
66 ext. papers	7,075 ext. citations	7.2 avg, IF	5.66 L-index

#	Paper	IF	Citations
59	Prevalence and Cumulative Risk of Familial Idiopathic Dilated Cardiomyopathy.. <i>JAMA - Journal of the American Medical Association</i> , 2022 , 327, 454-463	27.4	4
58	Validating an Idiopathic Dilated Cardiomyopathy Diagnosis Using Cardiovascular Magnetic Resonance: The Dilated Cardiomyopathy Precision Medicine Study.. <i>Circulation: Heart Failure</i> , 2022 , CIRCHEARTFAILURE1	7.6	0
57	Communal Coping as a Strategy to Enhance Family Engagement in Dilated Cardiomyopathy.. <i>Circulation Genomic and Precision Medicine</i> , 2022 , 101161CIRCGEN121003541	5.2	
56	Dilated Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2022 , 79, 2233-2235	15.1	0
55	Response to McGurk et al.. <i>Genetics in Medicine</i> , 2021 ,	8.1	
54	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2021 update: a policy statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021 , 23, 1391-1398	8.1	33
53	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> , 2021 , 23, 589-598	5.1	1
52	ACMG SF v3.0 list for reporting of secondary findings in clinical exome and genome sequencing: a policy statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021 , 23, 1381-1390	8.1	81
51	The Complex and Diverse Genetic Architecture of Dilated Cardiomyopathy. <i>Circulation Research</i> , 2021 , 128, 1514-1532	15.7	10
50	Evidence-Based Assessment of Genes in Dilated Cardiomyopathy. <i>Circulation</i> , 2021 , 144, 7-19	16.7	34
49	International Evidence Based Reappraisal of Genes Associated With Arrhythmogenic Right Ventricular Cardiomyopathy Using the Clinical Genome Resource Framework. <i>Circulation Genomic and Precision Medicine</i> , 2021 , 14, e003273	5.2	21
48	Considering complexity in the genetic evaluation of dilated cardiomyopathy. <i>Heart</i> , 2021 , 107, 106-112	5.1	2
47	Attitudes of Dilated Cardiomyopathy Patients and Investigators Toward Genomic Study Enrollment, Consent Process, and Return of Genetic Results. <i>Clinical and Translational Science</i> , 2021 , 14, 550-557	4.9	2
46	Evaluation of gene validity for CPVT and short QT syndrome in sudden arrhythmic death. <i>European Heart Journal</i> , 2021 ,	9.5	5
45	Effects of danicamtiv, a novel cardiac myosin activator, in heart failure with reduced ejection fraction: experimental data and clinical results from a phase 2a trial. <i>European Journal of Heart Failure</i> , 2020 , 22, 1649-1658	12.3	20
44	Variant Interpretation for Dilated Cardiomyopathy: Refinement of the American College of Medical Genetics and Genomics/ClinGen Guidelines for the DCM Precision Medicine Study. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e002480	5.2	27
43	Gain-of-Function Variants in Dilated Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e002892	5.2	5

42	Familial Dilated Cardiomyopathy. <i>Heart Lung and Circulation</i> , 2020 , 29, 566-574	1.8	11
41	Genetic Testing for Inherited Cardiovascular Diseases: A Scientific Statement From the American Heart Association. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e000067	5.2	59
40	Novel heterozygous truncating titin variants affecting the A-band are associated with cardiomyopathy and myopathy/muscular dystrophy. <i>Molecular Genetics & Genomic Medicine</i> , 2020 , 8, e1460	2.3	1
39	Evaluating the Clinical Validity of Hypertrophic Cardiomyopathy Genes. <i>Circulation Genomic and Precision Medicine</i> , 2019 , 12, e002460	5.2	132
38	Dilated cardiomyopathy. <i>Nature Reviews Disease Primers</i> , 2019 , 5, 32	51.1	143
37	Regional Variation in RBM20 Causes a Highly Penetrant Arrhythmogenic Cardiomyopathy. <i>Circulation: Heart Failure</i> , 2019 , 12, e005371	7.6	51
36	Hypertrophic cardiomyopathy genetic test reports: A qualitative study of patient understanding of uninformative genetic test results. <i>Journal of Genetic Counseling</i> , 2019 , 28, 1087-1097	2.5	4
35	Development of Clinical Domain Working Groups for the Clinical Genome Resource (ClinGen): lessons learned and plans for the future. <i>Genetics in Medicine</i> , 2019 , 21, 987-993	8.1	13
34	Genetic Evaluation of Cardiomyopathy-A Heart Failure Society of America Practice Guideline. <i>Journal of Cardiac Failure</i> , 2018 , 24, 281-302	3.3	160
33	Pilot Randomized Controlled Trial to Reduce Readmission for Heart Failure Using Novel Tablet and Nurse Practitioner Education. <i>American Journal of Medicine</i> , 2018 , 131, 974-978	2.4	5
32	Reappraisal of Reported Genes for Sudden Arrhythmic Death: Evidence-Based Evaluation of Gene Validity for Brugada Syndrome. <i>Circulation</i> , 2018 , 138, 1195-1205	16.7	158
31	Clinical Genetic Testing for Familial Hypercholesterolemia: JACC Scientific Expert Panel. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 662-680	15.1	215
30	Genetic evaluation of cardiomyopathy: a clinical practice resource of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2018 , 20, 899-909	8.1	96
29	Multigenic Disease and Bilineal Inheritance in Dilated Cardiomyopathy Is Illustrated in Nonsegregating LMNA Pedigrees. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002038	5.2	12
28	Toward Genetics-Driven Early Intervention in Dilated Cardiomyopathy: Design and Implementation of the DCM Precision Medicine Study. <i>Circulation: Cardiovascular Genetics</i> , 2017 , 10,		25
27	The Rationale and Timing of Molecular Genetic Testing for Dilated Cardiomyopathy. <i>Canadian Journal of Cardiology</i> , 2015 , 31, 1309-12	3.8	27
26	Novel familial dilated cardiomyopathy mutation in MYL2 affects the structure and function of myosin regulatory light chain. <i>FEBS Journal</i> , 2015 , 282, 2379-93	5.7	23
25	A novel human R25C-phospholamban mutation is associated with super-inhibition of calcium cycling and ventricular arrhythmia. <i>Cardiovascular Research</i> , 2015 , 107, 164-74	9.9	54

24	Design and implementation of a randomized controlled trial of genomic counseling for patients with chronic disease. <i>Journal of Personalized Medicine</i> , 2014 , 4, 1-19	3.6	17
23	Dilated cardiomyopathy: the complexity of a diverse genetic architecture. <i>Nature Reviews Cardiology</i> , 2013 , 10, 531-47	14.8	534
22	Exome sequencing and genome-wide linkage analysis in 17 families illustrate the complex contribution of TTN truncating variants to dilated cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2013 , 6, 144-53		81
21	Where genome meets phenotype: rationale for integrating genetic and protein biomarkers in the diagnosis and management of dilated cardiomyopathy and heart failure. <i>Journal of the American College of Cardiology</i> , 2012 , 60, 283-9	15.1	63
20	Reconsidering association testing methods using single-variant test statistics as alternatives to pooling tests for sequence data with rare variants. <i>PLoS ONE</i> , 2012 , 7, e30238	3.7	30
19	Update 2011: clinical and genetic issues in familial dilated cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2011 , 57, 1641-9	15.1	249
18	HRS/EHRA expert consensus statement on the state of genetic testing for the channelopathies and cardiomyopathies this document was developed as a partnership between the Heart Rhythm Society (HRS) and the European Heart Rhythm Association (EHRA). <i>Heart Rhythm</i> , 2011 , 8, 1308-39	6.7	737
17	Genome-wide studies of copy number variation and exome sequencing identify rare variants in BAG3 as a cause of dilated cardiomyopathy. <i>American Journal of Human Genetics</i> , 2011 , 88, 273-82	11	264
16	Rare variant mutations identified in pediatric patients with dilated cardiomyopathy. <i>Progress in Pediatric Cardiology</i> , 2011 , 31, 39-47	0.4	56
15	Coding sequence rare variants identified in MYBPC3, MYH6, TPM1, TNNC1, and TNNI3 from 312 patients with familial or idiopathic dilated cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2010 , 3, 155-61		173
14	Rare variant mutations in pregnancy-associated or peripartum cardiomyopathy. <i>Circulation</i> , 2010 , 121, 2176-82	16.7	143
13	Clinical and genetic issues in dilated cardiomyopathy: a review for genetics professionals. <i>Genetics in Medicine</i> , 2010 , 12, 655-67	8.1	179
12	A glimpse into multigene rare variant genetics: triple mutations in hypertrophic cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2010 , 55, 1454-5	15.1	11
11	Progress with genetic cardiomyopathies: screening, counseling, and testing in dilated, hypertrophic, and arrhythmogenic right ventricular dysplasia/cardiomyopathy. <i>Circulation: Heart Failure</i> , 2009 , 2, 253-61	7.6	158
10	Clinical and functional characterization of TNNT2 mutations identified in patients with dilated cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2009 , 2, 306-13		77
9	Genetic evaluation of cardiomyopathy—a Heart Failure Society of America practice guideline. <i>Journal of Cardiac Failure</i> , 2009 , 15, 83-97	3.3	388
8	Family history: an essential tool for cardiovascular genetic medicine. <i>Congestive Heart Failure</i> , 2008 , 14, 37-45		49
7	Coding sequence mutations identified in MYH7, TNNT2, SCN5A, CSRP3, LBD3, and TCAP from 313 patients with familial or idiopathic dilated cardiomyopathy. <i>Clinical and Translational Science</i> , 2008 , 1, 21-6	4.9	144

6	Lamin A/C mutation analysis in a cohort of 324 unrelated patients with idiopathic or familial dilated cardiomyopathy. <i>American Heart Journal</i> , 2008 , 156, 161-9	4.9	184
5	Cardiovascular genetic medicine: evolving concepts, rationale, and implementation. <i>Journal of Cardiovascular Translational Research</i> , 2008 , 1, 137-43	3.3	19
4	Clinical characteristics of 304 kindreds evaluated for familial dilated cardiomyopathy. <i>Journal of Cardiac Failure</i> , 2006 , 12, 422-9	3.3	36
3	Clinical and genetic issues in familial dilated cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2005 , 45, 969-81	15.1	279
2	A novel lamin A/C mutation in a family with dilated cardiomyopathy, prominent conduction system disease, and need for permanent pacemaker implantation. <i>American Heart Journal</i> , 2002 , 144, 1081-6	4.9	57
1	Genetic Counseling and Screening Issues in Familial Dilated Cardiomyopathy. <i>Journal of Genetic Counseling</i> , 2001 , 10, 397-415	2.5	22