

Ana Morales

List of Publications by Year in descending order

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31
papers

3,096
citations

394421

19
h-index

434195

31
g-index

32
all docs

32
docs citations

32
times ranked

4406
citing authors

#	ARTICLE	IF	CITATIONS
1	Dilated cardiomyopathy: the complexity of a diverse genetic architecture. <i>Nature Reviews Cardiology</i> , 2013, 10, 531-547.	13.7	763
2	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> , 2018, 20, 351-359.	2.4	283
3	Genetic Evaluation of Cardiomyopathy—A Heart Failure Society of America Practice Guideline. <i>Journal of Cardiac Failure</i> , 2018, 24, 281-302.	1.7	280
4	Clinical and genetic issues in dilated cardiomyopathy: A review for genetics professionals. <i>Genetics in Medicine</i> , 2010, 12, 655-667.	2.4	223
5	Coding Sequence Rare Variants Identified in <i>MYBPC3</i> , <i>MYH6</i> , <i>TPM1</i> , <i>TNNC1</i> , and <i>TNNI3</i> From 312 Patients With Familial or Idiopathic Dilated Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2010, 3, 155-161.	5.1	218
6	Progress With Genetic Cardiomyopathies. <i>Circulation: Heart Failure</i> , 2009, 2, 253-261.	3.9	191
7	Rare Variant Mutations in Pregnancy-Associated or Peripartum Cardiomyopathy. <i>Circulation</i> , 2010, 121, 2176-2182.	1.6	179
8	Identification of Novel Mutations in <i>RBM20</i> in Patients with Dilated Cardiomyopathy. <i>Clinical and Translational Science</i> , 2010, 3, 90-97.	3.1	159
9	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.	3.6	112
10	Clinical and Functional Characterization of <i>TNNT2</i> Mutations Identified in Patients With Dilated Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2009, 2, 306-313.	5.1	95
11	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, 144-153.	5.1	95
12	A novel human R25C-phospholamban mutation is associated with super-inhibition of calcium cycling and ventricular arrhythmia. <i>Cardiovascular Research</i> , 2015, 107, 164-174.	3.8	72
13	Variant Interpretation for Dilated Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e002480.	3.6	70
14	Family History: An Essential Tool for Cardiovascular Genetic Medicine. <i>Congestive Heart Failure</i> , 2008, 14, 37-45.	2.0	56
15	Genetic Evaluation of Dilated Cardiomyopathy. <i>Current Cardiology Reports</i> , 2013, 15, 375.	2.9	42
16	Novel familial dilated cardiomyopathy mutation in <i>MYL2</i> affects the structure and function of myosin regulatory light chain. <i>FEBS Journal</i> , 2015, 282, 2379-2393.	4.7	42
17	Toward Genetics-Driven Early Intervention in Dilated Cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2017, 10, .	5.1	41
18	Genetic Testing in Inherited Heart Diseases. <i>Heart Lung and Circulation</i> , 2020, 29, 505-511.	0.4	34

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19	The Rationale and Timing of Molecular Genetic Testing for Dilated Cardiomyopathy. <i>Canadian Journal of Cardiology</i> , 2015, 31, 1309-1312.	1.7	31
20	Variants of Uncertain Significance. <i>Circulation Genomic and Precision Medicine</i> , 2018, 11, e002169.	3.6	24
21	Is Left Ventricular Noncompaction a Trait, Phenotype, or Disease?. <i>Circulation: Cardiovascular Genetics</i> , 2017, 10, .	5.1	15
22	Impact of variant reclassification in the clinical setting of cardiovascular genetics. <i>Journal of Genetic Counseling</i> , 2021, 30, 503-512.	1.6	13
23	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.	1.2	10
24	Late Onset Sporadic Dilated Cardiomyopathy Caused by a Cardiac Troponin T Mutation. <i>Clinical and Translational Science</i> , 2010, 3, 219-226.	3.1	9
25	Communication pitfalls in interpreted genetic counseling sessions. <i>Journal of Genetic Counseling</i> , 2019, 28, 897-907.	1.6	8
26	Expanded newborn screening in Puerto Rico and the US Virgin Islands: education and barriers assessment. <i>Genetics in Medicine</i> , 2009, 11, 169-175.	2.4	6
27	Clinical Application of Genetic Testing in Heart Failure. <i>Current Heart Failure Reports</i> , 2017, 14, 543-553.	3.3	6
28	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.	1.6	6
29	At the Heart of the Pregnancy: What Prenatal and Cardiovascular Genetic Counselors Need to Know about Maternal Heart Disease. <i>Journal of Genetic Counseling</i> , 2017, 26, 669-688.	1.6	5
30	Harmonizing the Collection of Clinical Data on Genetic Testing Requisition Forms to Enhance Variant Interpretation in Hypertrophic Cardiomyopathy (HCM). <i>Journal of Molecular Diagnostics</i> , 2021, 23, 589-598.	2.8	5
31	The Family History as a Tool to Identify Patients at Risk for Dilated Cardiomyopathy. <i>Progress in Cardiovascular Nursing</i> , 2008, 23, 41-44.	0.4	3