Ana Morales

List of Publications by Year
in descending order

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	Impact of variant reclassification in the clinical setting of cardiovascular genetics. Journal of Genetic Counseling, 2021, 30, 503-512.	1.6	13
2	Harmonizing the Collection of Clinical Data on Genetic Testing Requisition Forms to Enhance Variant Interpretation in Hypertrophic Cardiomyopathy (HCM). Journal of Molecular Diagnostics, 2021, 23, 589-598.	2.8	5
3	International Evidence Based Reappraisal of Genes Associated With Arrhythmogenic Right Ventricular Cardiomyopathy Using the Clinical Genome Resource Framework. Circulation Genomic and Precision Medicine, 2021, 14, e003273.	3.6	112
4	Genetic Testing in Inherited Heart Diseases. Heart Lung and Circulation, 2020, 29, 505-511.	0.4	34
5	Novel heterozygous truncating titin variants affecting the Aâ€band are associated with cardiomyopathy and myopathy/muscular dystrophy. Molecular Genetics & Genomic Medicine, 2020, 8, e1460.	1.2	10
6	Variant Interpretation for Dilated Cardiomyopathy. Circulation Genomic and Precision Medicine, 2020, 13, e002480.	3.6	70
7	Hypertrophic cardiomyopathy genetic test reports: A qualitative study of patient understanding of uninformative genetic test results. Journal of Genetic Counseling, 2019, 28, 1087-1097.	1.6	6
8	Communication pitfalls in interpreted genetic counseling sessions. Journal of Genetic Counseling, 2019, 28, 897-907.	1.6	8
9	Adaptation and validation of the ACMG/AMP variant classification framework for MYH7-associated inherited cardiomyopathies: recommendations by ClinGenâ€™s Inherited Cardiomyopathy Expert Panel. Genetics in Medicine, 2018, 20, 351-359.	2.4	283
10	Genetic Evaluation of Cardiomyopathyâ€”A Heart Failure Society of America Practice Guideline. Journal of Cardiac Failure, 2018, 24, 281-302.	1.7	280
11	Variants of Uncertain Significance. Circulation Genomic and Precision Medicine, 2018, 11, e002169.	3.6	24
12	At the Heart of the Pregnancy: What Prenatal and Cardiovascular Genetic Counselors Need to Know about Maternal Heart Disease. Journal of Genetic Counseling, 2017, 26, 669-688.	1.6	5
13	Toward Genetics-Driven Early Intervention in Dilated Cardiomyopathy. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	41
14	Clinical Application of Genetic Testing in Heart Failure. Current Heart Failure Reports, 2017, 14, 543-553.	3.3	6
15	Is Left Ventricular Noncompaction a Trait, Phenotype, or Disease?. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	15
16	A novel human R25C-phospholamban mutation is associated with super-inhibition of calcium cycling and ventricular arrhythmia. Cardiovascular Research, 2015, 107, 164-174.	3.8	72
17	The Rationale and Timing of Molecular Genetic Testing forÂDilated Cardiomyopathy. Canadian Journal of Cardiology, 2015, 31, 1309-1312.	1.7	31
18	Novel familial dilated cardiomyopathy mutation in <scp>MYL</scp>2 affects the structure and function of myosin regulatory light chain. FEBS Journal, 2015, 282, 2379-2393.	4.7	42

#	Article	IF	Citations
19	Dilated cardiomyopathy: the complexity of a diverse genetic architecture. Nature Reviews Cardiology, 2013, 10, 531-547.	13.7	763
20	Exome Sequencing and Genome-Wide Linkage Analysis in 17 Families Illustrate the Complex Contribution of TTN Truncating Variants to Dilated Cardiomyopathy. Circulation: Cardiovascular Genetics, 2013, 6, 144-153.	5.1	95
21	Genetic Evaluation of Dilated Cardiomyopathy. Current Cardiology Reports, 2013, 15, 375.	2.9	42
22	Identification of Novel Mutations in RBM20 in Patients with Dilated Cardiomyopathy. Clinical and Translational Science, 2010, 3, 90-97.	3.1	159
23	Late Onset Sporadic Dilated Cardiomyopathy Caused by a Cardiac Troponin T Mutation. Clinical and Translational Science, 2010, 3, 219-226.	3.1	9
24	Coding Sequence Rare Variants Identified in MYBPC3 , MYH6 , TPM1 , TNNC1 , and TNNI3 From 312 Patients With Familial or Idiopathic Dilated Cardiomyopathy. Circulation: Cardiovascular Genetics, 2010, 3, 155-161.	5.1	218
25	Rare Variant Mutations in Pregnancy-Associated or Peripartum Cardiomyopathy. Circulation, 2010, 121, 2176-2182.	1.6	179
26	Clinical and genetic issues in dilated cardiomyopathy: A review for genetics professionals. Genetics in Medicine, 2010, 12, 655-667.	2.4	223
27	Expanded newborn screening in Puerto Rico and the US Virgin Islands: education and barriers assessment. Genetics in Medicine, 2009, 11, 169-175.	2.4	6
28	Progress With Genetic Cardiomyopathies. Circulation: Heart Failure, 2009, 2, 253-261.	3.9	191
29	Clinical and Functional Characterization of TNNT2 Mutations Identified in Patients With Dilated Cardiomyopathy. Circulation: Cardiovascular Genetics, 2009, 2, 306-313.	5.1	95
30	The Family History as a Tool to Identify Patients at Risk for Dilated Cardiomyopathy. Progress in Cardiovascular Nursing, 2008, 23, 41-44.	0.4	3
31	Family History: An Essential Tool for Cardiovascular Genetic Medicine. Congestive Heart Failure, 2008, 14, 37-45.	2.0	56