Michael Morley

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11850248/publications.pdf

Version: 2024-02-01

26 5,164 papers citations

361413 552781 26
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26 26 all docs citations

26 times ranked 7862 citing authors

#	Article	IF	CITATIONS
1	Genetic analysis of genome-wide variation in human gene expression. Nature, 2004, 430, 743-747.	27.8	1,146
2	Making and reading microarrays. Nature Genetics, 1999, 21, 15-19.	21.4	606
3	Mapping determinants of human gene expression by regional and genome-wide association. Nature, 2005, 437, 1365-1369.	27.8	550
4	Natural variation in human gene expression assessed in lymphoblastoid cells. Nature Genetics, 2003, 33, 422-425.	21.4	533
5	Common genetic variants account for differences in gene expression among ethnic groups. Nature Genetics, 2007, 39, 226-231.	21.4	466
6	Differentiation of Human Pluripotent Stem Cells into Functional Lung Alveolar Epithelial Cells. Cell Stem Cell, 2017, 21, 472-488.e10.	11.1	406
7	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. Nature Genetics, 2014, 46, 826-836.	21.4	281
8	RNA-Seq identifies novel myocardial gene expression signatures of heart failure. Genomics, 2015, 105, 83-89.	2.9	220
9	Genetic analysis of radiation-induced changes in human gene expression. Nature, 2009, 459, 587-591.	27.8	192
10	Polymorphic Cis- and Trans-Regulation of Human Gene Expression. PLoS Biology, 2010, 8, e1000480.	5.6	142
11	RNA-sequence analysis of human B-cells. Genome Research, 2011, 21, 991-998.	5.5	126
12	Genomics-First Evaluation of Heart Disease Associated With Titin-Truncating Variants. Circulation, 2019, 140, 42-54.	1.6	97
13	Single-Cell Transcriptomic Profiling of Pluripotent Stem Cell-Derived SCGB3A2+ Airway Epithelium. Stem Cell Reports, 2018, 10, 1579-1595.	4.8	78
14	Monozygotic Twins Reveal Germline Contribution to Allelic Expression Differences. American Journal of Human Genetics, 2008, 82, 1357-1360.	6.2	55
15	Gene Expression Phenotype in Heterozygous Carriers of Ataxia Telangiectasia. American Journal of Human Genetics, 2002, 71, 791-800.	6.2	50
16	Truncated titin proteins in dilated cardiomyopathy. Science Translational Medicine, 2021, 13, eabd7287.	12.4	39
17	A common variant alters SCN5A–miR-24 interaction and associates with heart failure mortality. Journal of Clinical Investigation, 2018, 128, 1154-1163.	8.2	34
18	Discovery of Genetic Variation on Chromosome 5q22 Associated with Mortality in Heart Failure. PLoS Genetics, 2016, 12, e1006034.	3.5	34

#	Article	IF	CITATIONS
19	Genetic variation in radiation-induced cell death. Genome Research, 2012, 22, 332-339.	5.5	33
20	Method for manufacturing whole-genome microarrays by rolling circle amplification. Genes Chromosomes and Cancer, 2004, 40, 72-77.	2.8	20
21	Antisense regulation of atrial natriuretic peptide expression. JCI Insight, 2019, 4, .	5.0	14
22	A Resource of Mapped Human Bacterial Artificial Chromosome Clones. Genome Research, 1999, 9, 989-993.	5.5	12
23	Global analysis of histone modifications and long-range chromatin interactions revealed the differential cistrome changes and novel transcriptional players in human dilated cardiomyopathy. Journal of Molecular and Cellular Cardiology, 2020, 145, 30-42.	1.9	11
24	Whole-Transcriptome Profiling of Human Heart Tissues Reveals the Potential Novel Players and Regulatory Networks in Different Cardiomyopathy Subtypes of Heart Failure. Circulation Genomic and Precision Medicine, 2021, 14, e003142.	3.6	7
25	Direct IBD mapping: identical-by-descent mapping without genotyping. Genomics, 2004, 83, 335-345.	2.9	6
26	Differential expression of members of SOX family of transcription factors in failing human hearts. Translational Research, 2022, 242, 66-78.	5.0	6