

Milos Pjanic

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

1,377
citations

471509

17
h-index

713466

21
g-index

29
all docs

29
docs citations

29
times ranked

2270
citing authors

#	ARTICLE	IF	CITATIONS
1	Atheroprotective roles of smooth muscle cell phenotypic modulation and the TCF21 disease gene as revealed by single-cell analysis. <i>Nature Medicine</i> , 2019, 25, 1280-1289.	30.7	494
2	Integrative functional genomics identifies regulatory mechanisms at coronary artery disease loci. <i>Nature Communications</i> , 2016, 7, 12092.	12.8	123
3	Coronary Artery Disease Associated Transcription Factor TCF21 Regulates Smooth Muscle Precursor Cells That Contribute to the Fibrous Cap. <i>PLoS Genetics</i> , 2015, 11, e1005155.	3.5	86
4	Genetic Regulatory Mechanisms of Smooth Muscle Cells Map to Coronary Artery Disease Risk Loci. <i>American Journal of Human Genetics</i> , 2018, 103, 377-388.	6.2	76
5	Coronary Disease-Associated Gene <i>TCF21</i> Inhibits Smooth Muscle Cell Differentiation by Blocking the Myocardin-Serum Response Factor Pathway. <i>Circulation Research</i> , 2020, 126, 517-529.	4.5	67
6	Environment-Sensing Aryl Hydrocarbon Receptor Inhibits the Chondrogenic Fate of Modulated Smooth Muscle Cells in Atherosclerotic Lesions. <i>Circulation</i> , 2020, 142, 575-590.	1.6	57
7	TCF21 and the environmental sensor aryl-hydrocarbon receptor cooperate to activate a pro-inflammatory gene expression program in coronary artery smooth muscle cells. <i>PLoS Genetics</i> , 2017, 13, e1006750.	3.5	52
8	Nuclear factor I revealed as family of promoter binding transcription activators. <i>BMC Genomics</i> , 2011, 12, 181.	2.8	48
9	TCF21 and AP-1 interact through epigenetic modifications to regulate coronary artery disease gene expression. <i>Genome Medicine</i> , 2019, 11, 23.	8.2	43
10	Characterization of TCF21 Downstream Target Regions Identifies a Transcriptional Network Linking Multiple Independent Coronary Artery Disease Loci. <i>PLoS Genetics</i> , 2015, 11, e1005202.	3.5	41
11	Coronary artery disease genes SMAD3 and TCF21 promote opposing interactive genetic programs that regulate smooth muscle cell differentiation and disease risk. <i>PLoS Genetics</i> , 2018, 14, e1007681.	3.5	41
12	The role of polycarbonate monomer bisphenol-A in insulin resistance. <i>PeerJ</i> , 2017, 5, e3809.	2.0	41
13	Genomic profiling of human vascular cells identifies TWIST1 as a causal gene for common vascular diseases. <i>PLoS Genetics</i> , 2020, 16, e1008538.	3.5	40
14	Advances in Transcriptomics. <i>Circulation Research</i> , 2018, 122, 1200-1220.	4.5	38
15	Genetics and Genomics of Coronary Artery Disease. <i>Current Cardiology Reports</i> , 2016, 18, 102.	2.9	31
16	Functional regulatory mechanism of smooth muscle cell-restricted LMOD1 coronary artery disease locus. <i>PLoS Genetics</i> , 2018, 14, e1007755.	3.5	30
17	Nuclear Factor I genomic binding associates with chromatin boundaries. <i>BMC Genomics</i> , 2013, 14, 99.	2.8	24
18	Molecular mechanisms of coronary disease revealed using quantitative trait loci for TCF21 binding, chromatin accessibility, and chromosomal looping. <i>Genome Biology</i> , 2020, 21, 135.	8.8	16

#	ARTICLE	IF	CITATIONS
19	From Locus Association to Mechanism of Gene Causality. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, 2079-2080.	2.4	12
20	CRISPR-Cas9-mediated knockout of <i>SPRY2</i> in human hepatocytes leads to increased glucose uptake and lipid droplet accumulation. <i>BMC Endocrine Disorders</i> , 2019, 19, 115.	2.2	6
21	Detailed Functional Characterization of a Waist-Hip Ratio Locus in 7p15.2 Defines an Enhancer Controlling Adipocyte Differentiation. <i>iScience</i> , 2019, 20, 42-59.	4.1	6
22	Abstract 21021: Functional Regulatory Mechanism of Smooth Muscle Cell-Restricted <i>LMOD1</i> Coronary Artery Disease Locus. <i>Circulation</i> , 2017, 136, .	1.6	1
23	Abstract 62: Molecular Basis of Regulatory Variation at Coronary Heart Disease-Associated Loci. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, .	2.4	0
24	Transcription factor regulation as a mechanism of confounding effects between distinct human traits. <i>F1000Research</i> , 0, 4, 1349.	1.6	0