Kun Sun

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7125169/publications.pdf

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		172457]	168389
55	4,399	29		53
papers	citations	h-index		g-index
58	58	58		6667
all docs	docs citations	times ranked		citing authors

#	Article	IF	CITATIONS
1	Plasma DNA tissue mapping by genome-wide methylation sequencing for noninvasive prenatal, cancer, and transplantation assessments. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E5503-12.	7.1	579
2	BayesHammer: Bayesian clustering for error correction in single-cell sequencing. BMC Genomics, 2013, 14, S7.	2.8	429
3	Noninvasive detection of cancer-associated genome-wide hypomethylation and copy number aberrations by plasma DNA bisulfite sequencing. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 18761-18768.	7.1	363
4	LncRNA Dum interacts with Dnmts to regulate Dppa2 expression during myogenic differentiation and muscle regeneration. Cell Research, 2015, 25, 335-350.	12.0	259
5	Linc-YY1 promotes myogenic differentiation and muscle regeneration through an interaction with the transcription factor YY1. Nature Communications, 2015, 6, 10026.	12.8	168
6	Orientation-aware plasma cell-free DNA fragmentation analysis in open chromatin regions informs tissue of origin. Genome Research, 2019, 29, 418-427.	5 . 5	159
7	Plasma DNA End-Motif Profiling as a Fragmentomic Marker in Cancer, Pregnancy, and Transplantation. Cancer Discovery, 2020, 10, 664-673.	9.4	152
8	Second generation noninvasive fetal genome analysis reveals de novo mutations, single-base parental inheritance, and preferred DNA ends. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E8159-E8168.	7.1	142
9	iSeeRNA: identification of long intergenic non-coding RNA transcripts from transcriptome sequencing data. BMC Genomics, 2013, 14, S7.	2.8	141
10	Preferred end coordinates and somatic variants as signatures of circulating tumor DNA associated with hepatocellular carcinoma. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E10925-E10933.	7.1	140
11	Genome-wide survey by ChIP-seq reveals YY1 regulation of lincRNAs in skeletal myogenesis. EMBO Journal, 2013, 32, 2575-2588.	7.8	138
12	<i>Dnase1 3</i> deletion causes aberrations in length and end-motif frequencies in plasma DNA. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 641-649.	7.1	134
13	Noninvasive Prenatal Methylomic Analysis by Genomewide Bisulfite Sequencing of Maternal Plasma DNA. Clinical Chemistry, 2013, 59, 1583-1594.	3.2	131
14	Combination of inflammation-related cytokines promotes long-term muscle stem cell expansion. Cell Research, 2015, 25, 655-673.	12.0	123
15	A Novel Wnt Regulatory Axis in Endometrioid Endometrial Cancer. Cancer Research, 2014, 74, 5103-5117.	0.9	114
16	Size-tagged preferred ends in maternal plasma DNA shed light on the production mechanism and show utility in noninvasive prenatal testing. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E5106-E5114.	7.1	107
17	Epigenetic Biomarkers in Cell-Free DNA and Applications in Liquid Biopsy. Genes, 2019, 10, 32.	2.4	96
18	A Novel YY1-miR-1 Regulatory Circuit in Skeletal Myogenesis Revealed by Genome-Wide Prediction of YY1-miRNA Network. PLoS ONE, 2012, 7, e27596.	2.5	88

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19	Malat1 regulates myogenic differentiation and muscle regeneration through modulating MyoD transcriptional activity. Cell Discovery, 2017, 3, 17002.	6.7	86
20	Pax3/7BP Is a Pax7- and Pax3-Binding Protein that Regulates the Proliferation of Muscle Precursor Cells by an Epigenetic Mechanism. Cell Stem Cell, 2012, 11, 231-241.	11.1	84
21	MyoD induced enhancer RNA interacts with hnRNPL to activate target gene transcription during myogenic differentiation. Nature Communications, 2019, 10, 5787.	12.8	70
22	DNA of Erythroid Origin Is Present in Human Plasma and Informs the Types of Anemia. Clinical Chemistry, 2017, 63, 1614-1623.	3.2	63
23	Liver- and Colon-Specific DNA Methylation Markers in Plasma for Investigation of Colorectal Cancers with or without Liver Metastases. Clinical Chemistry, 2018, 64, 1239-1249.	3.2	60
24	Atlas of ACE2 gene expression reveals novel insights into transmission of SARS-CoV-2. Heliyon, 2021, 7, e05850.	3.2	59
25	Cell-free DNA in maternal plasma and serum: A comparison of quantity, quality and tissue origin using genomic and epigenomic approaches. Clinical Biochemistry, 2016, 49, 1379-1386.	1.9	58
26	Methy-Pipe: An Integrated Bioinformatics Pipeline for Whole Genome Bisulfite Sequencing Data Analysis. PLoS ONE, 2014, 9, e100360.	2.5	54
27	Characterization of the nasopharyngeal carcinoma methylome identifies aberrant disruption of key signaling pathways and methylated tumor suppressor genes. Epigenomics, 2015, 7, 155-173.	2.1	52
28	A Molecular Switch Regulating Cell Fate Choice between Muscle Progenitor Cells and Brown Adipocytes. Developmental Cell, 2017, 41, 382-391.e5.	7.0	48
29	Noninvasive Prenatal Testing by Nanopore Sequencing of Maternal Plasma DNA: Feasibility Assessment. Clinical Chemistry, 2015, 61, 1305-1306.	3.2	44
30	FetalQuantSD: accurate quantification of fetal DNA fraction by shallow-depth sequencing of maternal plasma DNA. Npj Genomic Medicine, 2016, 1, 16013.	3.8	31
31	Ktrim: an extra-fast and accurate adapter- and quality-trimmer for sequencing data. Bioinformatics, 2020, 36, 3561-3562.	4.1	26
32	mTFkb: a knowledgebase for fundamental annotation of mouse transcription factors. Scientific Reports, 2017, 7, 3022.	3.3	21
33	Sebnif: An Integrated Bioinformatics Pipeline for the Identification of Novel Large Intergenic Noncoding RNAs (lincRNAs) - Application in Human Skeletal Muscle Cells. PLoS ONE, 2014, 9, e84500.	2.5	21
34	COFFEE: controlâ€free noninvasive fetal chromosomal examination using maternal plasma DNA. Prenatal Diagnosis, 2017, 37, 336-340.	2.3	17
35	Noninvasive reconstruction of placental methylome from maternal plasma DNA: Potential for prenatal testing and monitoring. Prenatal Diagnosis, 2018, 38, 196-203.	2.3	16
36	Gestational Age Assessment by Methylation and Size Profiling of Maternal Plasma DNA: A Feasibility Study. Clinical Chemistry, 2017, 63, 606-608.	3.2	14

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37	GeneCT: a generalizable cancerous status and tissue origin classifier for pan-cancer biopsies. Bioinformatics, 2018, 34, 4129-4130.	4.1	13
38	The impact of digital DNA counting technologies on noninvasive prenatal testing. Expert Review of Molecular Diagnostics, 2015, 15, 1261-1268.	3.1	9
39	YY1TargetDB: an integral information resource for Yin Yang 1 target loci. Database: the Journal of Biological Databases and Curation, 2013, 2013, bat007.	3.0	8
40	Genome-wide RNA-seq and ChIP-seq reveal Linc-YY1 function in regulating YY1/PRC2 activity during skeletal myogenesis. Genomics Data, 2016, 7, 247-249.	1.3	8
41	BSviewer: a genotype-preserving, nucleotide-level visualizer for bisulfite sequencing data. Bioinformatics, 2017, 33, 3495-3496.	4.1	5
42	Msuite: A High-Performance and Versatile DNA Methylation Data-Analysis Toolkit. Patterns, 2020, 1, 100127.	5.9	5
43	Clonal hematopoiesis: background player in plasma cell-free DNA variants. Annals of Translational Medicine, 2019, 7, S384-S384.	1.7	4
44	Transcriptional Start Site Coverage Analysis in Plasma Cell-Free DNA Reveals Disease Severity and Tissue Specificity of COVID-19 Patients. Frontiers in Genetics, 2021, 12, 663098.	2.3	4
45	Recent advances in blood-based and artificial intelligence-enhanced approaches for gastrointestinal cancer diagnosis. World Journal of Gastroenterology, 2021, 27, 5666-5681.	3.3	4
46	Genome-wide profiling of YY1 binding sites during skeletal myogenesis. Genomics Data, 2014, 2, 89-91.	1.3	3
47	Bioinformatics for Novel Long Intergenic Noncoding RNA (lincRNA) Identification in Skeletal Muscle Cells. Methods in Molecular Biology, 2017, 1556, 355-362.	0.9	3
48	Msuite2: All-in-one DNA methylation data analysis toolkit with enhanced usability and performance. Computational and Structural Biotechnology Journal, 2022, 20, 1271-1276.	4.1	3
49	The Online Diagnosis System for sanger sequencing based genetic testing. , 2014, , .		2
50	NAMS: Noncoding Assessment of long RNAs in Magnoliophyta Species. Methods in Molecular Biology, 2019, 1933, 257-264.	0.9	2
51	Rapid preliminary purity evaluation of tumor biopsies using deep learning approach. Computational and Structural Biotechnology Journal, 2020, 18, 1746-1753.	4.1	2
52	Methy-Pipe: An integrated bioinformatics data analysis pipeline for whole genome methylome analysis. , 2010, , .		1
53	Online Diagnosis System: A webserver for analysis of Sanger sequencing-based genetic testing data. Methods, 2014, 69, 230-236.	3.8	1
54	NAMS webserver: coding potential assessment and functional annotation of plant transcripts. Briefings in Bioinformatics, 2021, 22, .	6.5	1

#	Article	IF	CITATIONS
55	Cover Image, Volume 37, Issue 4. Prenatal Diagnosis, 2017, 37, i.	2.3	O