## Winston A Hide

List of Publications by Year in descending order

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175 papers 25,224 citations

65 h-index 7518 151 g-index

267 all docs

267 docs citations

times ranked

267

44928 citing authors

#	Article	IF	CITATIONS
1	The Transcriptional Landscape of the Mammalian Genome. Science, 2005, 309, 1559-1563.	12.6	3,227
2	A promoter-level mammalian expression atlas. Nature, 2014, 507, 462-470.	27.8	1,838
3	Twelve type 2 diabetes susceptibility loci identified through large-scale association analysis. Nature Genetics, 2010, 42, 579-589.	21.4	1,631
4	Aberrant lipid metabolism disrupts calcium homeostasis causing liver endoplasmic reticulum stress in obesity. Nature, 2011, 473, 528-531.	27.8	864
5	A genome-wide approach accounting for body mass index identifies genetic variants influencing fasting glycemic traits and insulin resistance. Nature Genetics, 2012, 44, 659-669.	21.4	762
6	Integrating human sequence data sets provides a resource of benchmark SNP and indel genotype calls. Nature Biotechnology, 2014, 32, 246-251.	17.5	722
7	Gateways to the FANTOM5 promoter level mammalian expression atlas. Genome Biology, 2015, 16, 22.	8.8	687
8	An Atlas of Combinatorial Transcriptional Regulation in Mouse and Man. Cell, 2010, 140, 744-752.	28.9	667
9	The future of biocuration. Nature, 2008, 455, 47-50.	27.8	648
10	Meta-analysis of genome-wide association studies identifies eight new loci for type 2 diabetes in east Asians. Nature Genetics, 2012, 44, 67-72.	21.4	545
11	miR-24 Inhibits Cell Proliferation by Targeting E2F2, MYC, and Other Cell-Cycle Genes via Binding to "Seedless―3′UTR MicroRNA Recognition Elements. Molecular Cell, 2009, 35, 610-625.	9.7	544
12	COVID-19 tissue atlases reveal SARS-CoV-2 pathology and cellular targets. Nature, 2021, 595, 107-113.	27.8	537
13	Parental origin of sequence variants associated with complex diseases. Nature, 2009, 462, 868-874.	27.8	521
14	Genome-wide association study identifies loci influencing concentrations of liver enzymes in plasma. Nature Genetics, 2011, 43, 1131-1138.	21.4	501
15	Novel Loci for Adiponectin Levels and Their Influence on Type 2 Diabetes and Metabolic Traits: A Multi-Ethnic Meta-Analysis of 45,891 Individuals. PLoS Genetics, 2012, 8, e1002607.	3.5	419
16	The transcriptional network that controls growth arrest and differentiation in a human myeloid leukemia cell line. Nature Genetics, 2009, 41, 553-562.	21.4	408
17	Bayesian inference analyses of the polygenic architecture of rheumatoid arthritis. Nature Genetics, 2012, 44, 483-489.	21.4	402
18	Toward interoperable bioscience data. Nature Genetics, 2012, 44, 121-126.	21.4	362

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19	Enabling the genomic revolution in Africa. Science, 2014, 344, 1346-1348.	12.6	361
20	Quantitating the Multiplicity of Infection with Human Immunodeficiency Virus Type 1 Subtype C Reveals a Non-Poisson Distribution of Transmitted Variants. Journal of Virology, 2009, 83, 3556-3567.	3.4	354
21	Is the guinea-pig a rodent?. Nature, 1991, 351, 649-652.	27.8	318
22	The ESAT-6 gene cluster of Mycobacterium tuberculosis and other high G+C Gram-positive bacteria. Genome Biology, 2001, 2, research0044.1.	9.6	302
23	Genome-wide analysis of cancer/testis gene expression. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 20422-20427.	7.1	295
24	Integrative Annotation of 21,037 Human Genes Validated by Full-Length cDNA Clones. PLoS Biology, 2004, 2, e162.	5.6	290
25	Identification of an imprinted master trans regulator at the KLF14 locus related to multiple metabolic phenotypes. Nature Genetics, 2011, 43, 561-564.	21.4	289
26	Genome Sequence of the Tsetse Fly ( <i>Glossina morsitans</i> ): Vector of African Trypanosomiasis. Science, 2014, 344, 380-386.	12.6	254
27	ISA software suite: supporting standards-compliant experimental annotation and enabling curation at the community level. Bioinformatics, 2010, 26, 2354-2356.	4.1	247
28	Capture of MicroRNA–Bound mRNAs Identifies the Tumor Suppressor miR-34a as a Regulator of Growth Factor Signaling. PLoS Genetics, 2011, 7, e1002363.	3.5	222
29	Meta-Analysis of the Alzheimer's Disease Human Brain Transcriptome and Functional Dissection in Mouse Models. Cell Reports, 2020, 32, 107908.	6.4	199
30	A Genome-Wide Association Search for Type 2 Diabetes Genes in African Americans. PLoS ONE, 2012, 7, e29202.	2.5	197
31	Meta-Analysis of Genome-Wide Association Studies in African Americans Provides Insights into the Genetic Architecture of Type 2 Diabetes. PLoS Genetics, 2014, 10, e1004517.	3.5	191
32	Abdominal Aortic Aneurysm Is Associated with a Variant in Low-Density Lipoprotein Receptor-Related Protein 1. American Journal of Human Genetics, 2011, 89, 619-627.	6.2	185
33	Genome-Wide Association Study Identifies Novel Loci Associated with Circulating Phospho- and Sphingolipid Concentrations. PLoS Genetics, 2012, 8, e1002490.	3.5	181
34	d2_cluster: A Validated Method for Clustering EST and Full-Length cDNA Sequences. Genome Research, 1999, 9, 1135-1142.	5.5	176
35	A Comprehensive Approach to Clustering of Expressed Human Gene Sequence: The Sequence Tag Alignment and Consensus Knowledge Base. Genome Research, 1999, 9, 1143-1155.	5.5	175
36	Cell-specific translational profiling in acute kidney injury. Journal of Clinical Investigation, 2014, 124, 1242-1254.	8.2	172

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37	Integration of text- and data-mining using ontologies successfully selects disease gene candidates. Nucleic Acids Research, 2005, 33, 1544-1552.	14.5	167
38	A Pathway-Based View of Human Diseases and Disease Relationships. PLoS ONE, 2009, 4, e4346.	2.5	158
39	Transcriptome Analysis of Mouse Stem Cells and Early Embryos. PLoS Biology, 2003, 1, e74.	5.6	156
40	Human Proteinpedia enables sharing of human protein data. Nature Biotechnology, 2008, 26, 164-167.	17.5	155
41	Mutations in a novel retina-specific gene cause autosomal dominant retinitis pigmentosa. Nature Genetics, 1999, 22, 255-259.	21.4	154
42	A Genome-wide siRNA Screen Identifies Proteasome Addiction as a Vulnerability of Basal-like Triple-Negative Breast Cancer Cells. Cancer Cell, 2013, 24, 182-196.	16.8	147
43	eVOC: A Controlled Vocabulary for Unifying Gene Expression Data. Genome Research, 2003, 13, 1222-1230.	5.5	144
44	Computational disease gene identification: a concert of methods prioritizes type 2 diabetes and obesity candidate genes. Nucleic Acids Research, 2006, 34, 3067-3081.	14.5	134
45	Transmission of HIV-1 CTL Escape Variants Provides HLA-Mismatched Recipients with a Survival Advantage. PLoS Pathogens, 2008, 4, e1000033.	4.7	129
46	Project MinE: study design and pilot analyses of a large-scale whole-genome sequencing study in amyotrophic lateral sclerosis. European Journal of Human Genetics, 2018, 26, 1537-1546.	2.8	129
47	MicroRNA Expression Profile in Human Macrophages in Response to Leishmania major Infection. PLoS Neglected Tropical Diseases, 2013, 7, e2478.	3.0	125
48	Leveraging Cross-Species Transcription Factor Binding Site Patterns: From Diabetes Risk Loci to Disease Mechanisms. Cell, 2014, 156, 343-358.	28.9	113
49	STACK: Sequence Tag Alignment and Consensus Knowledgebase. Nucleic Acids Research, 2001, 29, 234-238.	14.5	112
50	Differential roles of epigenetic changes and Foxp3 expression in regulatory T cell-specific transcriptional regulation. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 5289-5294.	7.1	111
51	The generation and utilization of a cancer-oriented representation of the human transcriptome by using expressed sequence tags. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 13418-13423.	7.1	105
52	miR-200 promotes the mesenchymal to epithelial transition by suppressing multiple members of the Zeb2 and Snail1 transcriptional repressor complexes. Oncogene, 2016, 35, 158-172.	5.9	105
53	Transcriptomic Signature of Leishmania Infected Mice Macrophages: A Metabolic Point of View. PLoS Neglected Tropical Diseases, 2012, 6, e1763.	3.0	103
54	Alternative Gene Form Discovery and Candidate Gene Selection from Gene Indexing Projects. Genome Research, 1998, 8, 276-290.	5.5	102

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55	Integrated Genomic Analysis of Diverse Induced Pluripotent Stem Cells from the Progenitor Cell Biology Consortium. Stem Cell Reports, 2016, 7, 110-125.	4.8	101
56	Mining Biological Pathways Using WikiPathways Web Services. PLoS ONE, 2009, 4, e6447.	2.5	100
57	Sex-specific and blood meal-induced proteins of Anopheles gambiae midguts: analysis by two-dimensional gel electrophoresis. Malaria Journal, 2003, 2, 1.	2.3	96
58	Mice and Men: Their Promoter Properties. PLoS Genetics, 2006, 2, e54.	3 <b>.</b> 5	95
59	H3ABioNet, a sustainable pan-African bioinformatics network for human heredity and health in Africa. Genome Research, 2016, 26, 271-277.	5 <b>.</b> 5	94
60	Comparison of glioma stem cells to neural stem cells from the adult human brain identifies dysregulated Wnt- signaling and a fingerprint associated with clinical outcome. Experimental Cell Research, 2013, 319, 2230-2243.	2.6	92
61	ASTD: The Alternative Splicing and Transcript Diversity database. Genomics, 2009, 93, 213-220.	2.9	87
62	A network of epigenetic regulators guides developmental haematopoiesis in vivo. Nature Cell Biology, 2013, 15, 1516-1525.	10.3	81
63	Rapid evolution of cancer/testis genes on the X chromosome. BMC Genomics, 2007, 8, 129.	2.8	80
64	Novel Developmental Analyses Identify Longitudinal Patterns of Early Gut Microbiota that Affect Infant Growth. PLoS Computational Biology, 2013, 9, e1003042.	3.2	76
65	Association of TRIM22 with the Type 1 Interferon Response and Viral Control during Primary HIV-1 Infection. Journal of Virology, 2011, 85, 208-216.	3.4	66
66	Biological Evaluation of d <sup>2</sup> , an Algorithm for High-Performance Sequence Comparison. Journal of Computational Biology, 1994, 1, 199-215.	1.6	63
67	A data-driven approach links microglia to pathology and prognosis in amyotrophic lateral sclerosis. Acta Neuropathologica Communications, 2017, 5, 23.	5.2	63
68	The Molecular Taxonomy and Evolution of the Guinea Pig. Journal of Heredity, 1992, 83, 174-181.	2.4	61
69	A functional genomic screen reveals novel host genes that mediate interferon-alpha's effects against hepatitis C virus. Journal of Hepatology, 2012, 56, 326-333.	3.7	60
70	Transcriptome-Wide Assessment of Human Brain and Lymphocyte Senescence. PLoS ONE, 2008, 3, e3024.	2.5	60
71	The H-Invitational Database (H-InvDB), a comprehensive annotation resource for human genes and transcripts. Nucleic Acids Research, 2007, 36, D793-D799.	14.5	57
72	Prenatal Lead Levels, Plasma Amyloid $\hat{l}^2$ Levels, and Gene Expression in Young Adulthood. Environmental Health Perspectives, 2012, 120, 702-707.	6.0	57

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73	Ten Simple Rules for Organizing a Virtual Conference—Anywhere. PLoS Computational Biology, 2010, 6, e1000650.	3.2	54
74	Compound heterozygous variants in NBAS as a cause of atypical osteogenesis imperfecta. Bone, 2017, 94, 65-74.	2.9	54
75	Eukaryotic genes in Mycobacterium tuberculosis could have a role in pathogenesis and immunomodulation. Trends in Genetics, 2002, 18, 5-8.	6.7	53
76	edgeRun: an R package for sensitive, functionally relevant differential expression discovery using an unconditional exact test. Bioinformatics, 2015, 31, 2589-2590.	4.1	53
77	Circadian Gene Variants and Susceptibility to Type 2 Diabetes: A Pilot Study. PLoS ONE, 2012, 7, e32670.	2.5	52
78	Sequencing of Captive Target Transcripts Identifies the Network of Regulated Genes and Functions of Primate-Specific miR-522. Cell Reports, 2014, 8, 1225-1239.	6.4	50
79	Wholeâ€genome sequencing reveals new Alzheimer's disease–associated rare variants in loci related to synaptic function and neuronal development. Alzheimer's and Dementia, 2021, 17, 1509-1527.	0.8	50
80	Assembly, Verification, and Initial Annotation of the NIA Mouse 7.4K cDNA Clone Set. Genome Research, 2002, 12, 1999-2003.	5.5	49
81	Molecular, phenotypic, and sample-associated data to describe pluripotent stem cell lines and derivatives. Scientific Data, 2017, 4, 170030.	5.3	48
82	The Contribution of Exon-Skipping Events on Chromosome 22 to Protein Coding Diversity. Genome Research, 2001, 11, 1848-1853.	5.5	46
83	A direct fate exclusion mechanism by Sonic hedgehog-regulated transcriptional repressors. Development (Cambridge), 2015, 142, 3286-93.	2.5	42
84	Silencing of the Drosophila ortholog of SOX5 leads to abnormal neuronal development and behavioral impairment. Human Molecular Genetics, 2017, 26, 1472-1482.	2.9	42
85	The Pathway Coexpression Network: Revealing pathway relationships. PLoS Computational Biology, 2018, 14, e1006042.	3.2	41
86	Profiling the malaria genome: a gene survey of three species of malaria parasite with comparison to other apicomplexan species. Molecular and Biochemical Parasitology, 2001, 118, 201-210.	1.1	40
87	Immunotherapy for breast cancer using EpCAM aptamer tumor-targeted gene knockdown. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	38
88	New methods for finding disease-susceptibility genes: impact and potential. Genome Biology, 2003, 4, 119.	9.6	37
89	The LIFEdb database in 2006. Nucleic Acids Research, 2006, 34, D415-D418.	14.5	36
90	Computational selection and prioritization of candidate genes for Fetal Alcohol Syndrome. BMC Genomics, 2007, 8, 389.	2.8	36

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91	Protein kinase C $\hat{\Gamma}$ is essential for optimal macrophage-mediated phagosomal containment of <i>i&gt;Listeria</i> monocytogenes <i>i&gt;.</i> Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 16251-16256.	7.1	35
92	Deep phenotyping of peripheral tissue facilitates mechanistic disease stratification in sporadic Parkinson's disease. Progress in Neurobiology, 2020, 187, 101772.	5.7	35
93	An overview of the wcd EST clustering tool. Bioinformatics, 2008, 24, 1542-1546.	4.1	34
94	A Model of Directional Selection Applied to the Evolution of Drug Resistance in HIV-1. Molecular Biology and Evolution, 2007, 24, 1025-1031.	8.9	33
95	The Constrained Maximal Expression Level Owing to Haploidy Shapes Gene Content on the Mammalian X Chromosome. PLoS Biology, 2015, 13, e1002315.	5.6	32
96	A case for a Glossina genome project. Trends in Parasitology, 2005, 21, 107-111.	3.3	31
97	Identification of Novel Alzheimer's Disease Loci Using Sex-Specific Family-Based Association Analysis of Whole-Genome Sequence Data. Scientific Reports, 2020, 10, 5029.	3.3	31
98	The Stem Cell Discovery Engine: an integrated repository and analysis system for cancer stem cell comparisons. Nucleic Acids Research, 2012, 40, D984-D991.	14.5	29
99	Chronic Endotoxin Exposure Produces Airflow Obstruction and Lung Dendritic Cell Expansion. American Journal of Respiratory Cell and Molecular Biology, 2012, 47, 209-217.	2.9	27
100	Comparison of Illumina and 454 Deep Sequencing in Participants Failing Raltegravir-Based Antiretroviral Therapy. PLoS ONE, 2014, 9, e90485.	2.5	27
101	Relatively frequent switching of transcription start sites during cerebellar development. BMC Genomics, 2017, 18, 461.	2.8	27
102	Adaptive changes in HIV-1 subtype C proteins during early infection are driven by changes in HLA-associated immune pressure. Virology, 2010, 396, 213-225.	2.4	26
103	The biochemical phylogeny of guinea-pigs and gundis, and the paraphyly of the order Rodentia. Comparative Biochemistry and Physiology Part B: Comparative Biochemistry, 1992, 101, 495-498.	0.2	25
104	Molecular evolution of Mycobacterium tuberculosis: phylogenetic reconstruction of clonal expansion. Tuberculosis, 2001, 81, 291-302.	1.9	25
105	An in-depth comparison of the male pediatric and adult urinary proteomes. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2014, 1844, 1044-1050.	2.3	25
106	Origin of rodents and guinea-pigs. Nature, 1992, 359, 277-278.	27.8	24
107	Comparative analysis of resistant and susceptible macrophage gene expression response to Leishmania major parasite. BMC Genomics, 2013, 14, 723.	2.8	22
108	Inferring an Evolutionary Tree of Uveal Melanoma From Genomic Copy Number Aberrations. , 2015, 56, 6801.		22

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109	Exome sequencing reveals recurrent germ line variants in patients with familial WaldenstrĶm macroglobulinemia. Blood, 2016, 127, 2598-2606.	1.4	22
110	Aβ-accelerated neurodegeneration caused by Alzheimer's-associated <i>ACE</i> variant R1279Q is rescued by angiotensin system inhibition in mice. Science Translational Medicine, 2020, 12, .	12.4	22
111	Vertical HIV transmission in South Africa: translating research into policy and practice. Lancet, The, 2002, 359, 992-993.	13.7	21
112	Conserved Domains of Subtype C Nef from South African HIV Type 1-Infected Individuals Include Cytotoxic T Lymphocyte Epitope-Rich Regions. AIDS Research and Human Retroviruses, 2001, 17, 1681-1687.	1.1	20
113	Transcript-Specific Expression Profiles Derived from Sequence-Based Analysis of Standard Microarrays. PLoS ONE, 2009, 4, e4702.	2.5	20
114	The S-star trial bioinformatics course: An on-line learning success. Biochemistry and Molecular Biology Education, 2003, 31, 20-23.	1.2	18
115	The Human Anatomic Gene Expression Library (H-ANGEL), the H-Inv integrative display of human gene expression across disparate technologies and platforms. Nucleic Acids Research, 2004, 33, D567-D572.	14.5	16
116	CLU: A new algorithm for EST clustering. BMC Bioinformatics, 2005, 6, S3.	2.6	16
117	A missense (Asp250Asn) mutation in the lipoprotein lipase gene in two unrelated families with familial lipoprotein lipase deficiency. Journal of Lipid Research, 1992, 33, 745-54.	4.2	16
118	Assessment of the parallelization approach ofd2_cluster for high-performance sequence clustering. Journal of Computational Chemistry, 2002, 23, 755-757.	3.3	15
119	An Assessment of the Role of DNA Adenine Methyltransferase on Gene Expression Regulation in E coli. PLoS ONE, 2007, 2, e273.	2.5	15
120	Iterative sorting reveals CD133+ and CD133- melanoma cells as phenotypically distinct populations. BMC Cancer, 2016, 16, 726.	2.6	15
121	Physical and transcriptional map of the critical region for keratolytic winter erythema (KWE) on chromosome 8p22-p23 between D8S550 and D8S1759. European Journal of Human Genetics, 2002, 10, 17-25.	2.8	14
122	The relative resistance of children to sepsis mortality: from pathways to drug candidates. Molecular Systems Biology, 2018, 14, e7998.	7.2	14
123	Pathprinting: An integrative approach to understand the functional basis of disease. Genome Medicine, 2013, 5, 68.	8.2	13
124	Mycoplasma Infection Alters Cancer Stem Cell Properties in Vitro. Stem Cell Reviews and Reports, 2016, 12, 156-161.	5.6	13
125	Most Pathways Can Be Related to the Pathogenesis of Alzheimer's Disease. Frontiers in Aging Neuroscience, 0, 14, .	3.4	13
126	Simplified ontologies allowing comparison of developmental mammalian gene expression. Genome Biology, 2007, 8, R229.	9.6	12

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127	Divergent LIN28-mRNA associations result in translational suppression upon the initiation of differentiation. Nucleic Acids Research, 2014, 42, 7997-8007.	14.5	12
128	A prioritization analysis of disease association by data-mining of functional annotation of human genes. Genomics, 2012, 99, 1-9.	2.9	11
129	Discovery of widespread transcription initiation at microsatellites predictable by sequence-based deep neural network. Nature Communications, 2021, 12, 3297.	12.8	11
130	A comprehensive promoter landscape identifies a novel promoter for CD133 in restricted tissues, cancers, and stem cells. Frontiers in Genetics, 2013, 4, 209.	2.3	10
131	GCH1 Deficiency Activates Brain Innate Immune Response and Impairs Tyrosine Hydroxylase Homeostasis. Journal of Neuroscience, 2022, 42, 702-716.	3.6	10
132	Region-based analysis of rare genomic variants in whole-genome sequencing datasets reveal two novel Alzheimer's disease-associated genes: DTNB and DLG2. Molecular Psychiatry, 2022, 27, 1963-1969.	7.9	9
133	Application of eVOC: controlled vocabularies for unifying gene expression data. Comptes Rendus - Biologies, 2003, 326, 1089-1096.	0.2	8
134	Integrative analysis of intraerythrocytic differentially expressed transcripts yields novel insights into the biology of Plasmodium falciparum. Malaria Journal, 2003, 2, 38.	2.3	8
135	Prioritizing genes of potential relevance to diseases affected by sex hormones: an example of Myasthenia Gravis. BMC Genomics, 2008, 9, 481.	2.8	8
136	Computational Analysis of Constraints on Noncoding Regions, Coding Regions and Gene Expression in Relation to Plasmodium Phenotypic Diversity. PLoS ONE, 2008, 3, e3122.	2.5	8
137	Population Differences in Transcript-Regulator Expression Quantitative Trait Loci. PLoS ONE, 2012, 7, e34286.	2.5	8
138	Positive Selection Scanning Reveals Decoupling of Enzymatic Activities of Carbamoyl Phosphate Synthetase in Helicobacter pylori. Journal of Molecular Evolution, 2002, 54, 458-464.	1.8	7
139	CAGExploreR: an R package for the analysis and visualization of promoter dynamics across multiple experiments. Bioinformatics, 2014, 30, 1183-1184.	4.1	5
140	A Network Of Epigenetic Regulators Guide Developmental Hematopoiesis In Vivo. Blood, 2013, 122, 1174-1174.	1.4	5
141	An Atlas of Combinatorial Transcriptional Regulation in Mouse and Man. Cell, 2010, 141, 369.	28.9	4
142	FRAGS: estimation of coding sequence substitution rates from fragmentary data. BMC Bioinformatics, 2004, 5, 8.	2.6	3
143	Quantitating the Multiplicity of Infection with Human Immunodeficiency Virus Type 1 Subtype C Reveals a Non-Poisson Distribution of Transmitted Variants. Journal of Virology, 2009, 83, 6974-6974.	3.4	3
144	Message from the ISCB: ISCB Ebola award for important future research on the computational biology of Ebola virus. Bioinformatics, 2015, 31, 616-617.	4.1	3

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145	The Stem Cell Commons: an exemplar for data integration in the biomedical domain driven by the ISA framework. AMIA Summits on Translational Science Proceedings, 2013, 2013, 70.	0.4	3
146	Continuing tsetse and Trypanosoma genome sequencing projects. Trends in Parasitology, 2004, 20, 308-309.	3.3	2
147	ISCB Ebola Award for Important Future Research on the Computational Biology of Ebola Virus. PLoS Computational Biology, 2015, 11, e1004087.	3.2	2
148	Cell-specific translational profiling in acute kidney injury. Journal of Clinical Investigation, 2014, 124, 2288-2288.	8.2	2
149	Positive Selection Scanning of Parasite DNA Sequences. , 2004, 270, 127-150.		1
150	ISCB Ebola Award for Important Future Research on the Computational Biology of Ebola Virus. F1000Research, 2015, 4, 12.	1.6	1
151	Clonal-Heterogeneity and Propensity for Bone Metastasis in Multiple Myeloma. Blood, 2014, 124, 3370-3370.	1.4	1
152	Integrating Murine Gene Expression Studies to Understand Obstructive Lung Disease Due to Chronic Inhaled Endotoxin. PLoS ONE, 2013, 8, e62910.	2.5	1
153	The South African EMBnet Node: AGM 2010 report. EMBnet Journal, 2011, 16, 25.	0.6	1
154	Clinical allergy-state of the art. The British Journal of Clinical Practice, 1990, 44, 85-7.	0.2	1
155	A platform for genomics in South Africa. South African Medical Journal, 2001, 91, 1006-7.	0.6	1
156	USE OF A VISUAL COMPARATIVE METHOD TO RESOLVE CONSERVED SEQUENCE MOTIFS IN PROTEINS. , 1993, , .		0
157	ExScript: AN 'EX'-CENTRIC APPROACH TO THE DESCRIPTION OF TRANSCRIPT DIVERSITY. Bioinformatics, 2001, 17, 485-486.	4.1	0
158	Response to Hertz-Fowler and Berriman: Continuing tsetse and Trypanosoma genome sequencing projects. Trends in Parasitology, 2004, 20, 309-310.	3.3	0
159	EST clustering: a short tutorial. , 2005, , .		0
160	IDENTIFICATION OF NOVEL URINARY BIOMARKERS OF RENAL OBSTRUCTION USING TEMPORAL QUANTITATIVE PROTEOMICS. Journal of Urology, 2009, 181, 251-252.	0.4	O
161	Gene-Expression Ontologies and Tag-Based Expression Profiling. , 2009, , 169-178.		О
162	The Data Sharing Challenge. Learning Community From Consortia. Nature Precedings, 2010, , .	0.1	0

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163	Airway Reactivity And Persistent Inflammation Is Associated With Antigen Presenting Cell Population Shifts In A Murine Model Of Chronic Inhalational Endotoxin Exposure. , $2011, , .$		0
164	732 AN IN-DEPTH ANALYSIS OF THE PEDIATRIC URINARY PROTEOME. Journal of Urology, 2012, 187, .	0.4	0
165	In Vivo Profiling of Leukemic Stem Cell Fitness Identifies Therapeutically Actionable Determinants of Growth. Experimental Hematology, 2018, 64, S86.	0.4	0
166	Abstract SY43-03: Screening for triple negative breast cancer vulnerabilities. , 2014, , .		0
167	Identification of a Gene Expression Signature Characterizing Clonal Fitness and Dominance in Vivo in a Murine Model of MLL-AF9 Leukemia. Blood, 2014, 124, 2383-2383.	1.4	0
168	Reply to Neupane etÂal.: Replication study of ADâ€associated rare variants. Alzheimer's and Dementia, 2022, , .	0.8	0
169	Title is missing!. , 2013, 9, e1003042.		0
170	Title is missing!. , 2013, 9, e1003042.		0
171	Population Differences in Transcript-Regulator Expression Quantitative Trait Loci., 2012, 7, e34286.		0
172	Population Differences in Transcript-Regulator Expression Quantitative Trait Loci., 2012, 7, e34286.		0
173	Population Differences in Transcript-Regulator Expression Quantitative Trait Loci., 2012, 7, e34286.		0
174	Title is missing!. , 2008, 4, e1000033.		0
175	Title is missing!. , 2008, 4, e1000033.		0