## Alexander A Kanapin

List of Publications by Year in descending order

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59 papers

15,537 citations

126708 33 h-index 57 g-index

63 all docs 63
docs citations

63 times ranked

23973 citing authors

#	Article	IF	Citations
1	Systematic functional analysis of the Caenorhabditis elegans genome using RNAi. Nature, 2003, 421, 231-237.	13.7	3,343
2	The Transcriptional Landscape of the Mammalian Genome. Science, 2005, 309, 1559-1563.	6.0	3,227
3	Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs. Nature, 2002, 420, 563-573.	13.7	1,548
4	The InterPro database, an integrated documentation resource for protein families, domains and functional sites. Nucleic Acids Research, 2001, 29, 37-40.	6.5	928
5	Germline mutations affecting the proofreading domains of POLE and POLD1 predispose to colorectal adenomas and carcinomas. Nature Genetics, 2013, 45, 136-144.	9.4	851
6	Reactome knowledgebase of human biological pathways and processes. Nucleic Acids Research, 2009, 37, D619-D622.	6.5	760
7	The InterPro Database, 2003 brings increased coverage and new features. Nucleic Acids Research, 2003, 31, 315-318.	<b>6.</b> 5	640
8	InterPro, progress and status in 2005. Nucleic Acids Research, 2004, 33, D201-D205.	<b>6.</b> 5	478
9	New developments in the InterPro database. Nucleic Acids Research, 2007, 35, D224-D228.	6.5	444
10	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. Nature Genetics, 2015, 47, 717-726.	9.4	310
11	InterProan integrated documentation resource for protein families, domains and functional sites. Bioinformatics, 2000, 16, 1145-1150.	1.8	301
12	Integrative Annotation of 21,037 Human Genes Validated by Full-Length cDNA Clones. PLoS Biology, 2004, 2, e162.	2.6	290
13	Clinical whole-genome sequencing in severe early-onset epilepsy reveals new genes and improves molecular diagnosis. Human Molecular Genetics, 2014, 23, 3200-3211.	1.4	222
14	Mutations in TCF12, encoding a basic helix-loop-helix partner of TWIST1, are a frequent cause of coronal craniosynostosis. Nature Genetics, 2013, 45, 304-307.	9.4	181
15	Disruption of SF3B1 results in deregulated expression and splicing of key genes and pathways in myelodysplastic syndrome hematopoietic stem and progenitor cells. Leukemia, 2015, 29, 1092-1103.	3.3	161
16	Choice of transcripts and software has a large effect on variant annotation. Genome Medicine, 2014, 6, 26.	3.6	158
17	InterPro: An integrated documentation resource for protein families, domains and functional sites. Briefings in Bioinformatics, 2002, 3, 225-235.	3.2	155
18	Integr8 and Genome Reviews: integrated views of complete genomes and proteomes. Nucleic Acids Research, 2004, 33, D297-D302.	6.5	125

#	Article	IF	Citations
19	Clinical whole-genome sequencing from routine formalin-fixed, paraffin-embedded specimens: pilot study for the 100,000 Genomes Project. Genetics in Medicine, 2018, 20, 1196-1205.	1.1	125
20	Unfixed Endogenous Retroviral Insertions in the Human Population. Journal of Virology, 2014, 88, 9529-9537.	1.5	118
21	Recessive Mutations in SPTBN2 Implicate $\hat{l}^2$ -III Spectrin in Both Cognitive and Motor Development. PLoS Genetics, 2012, 8, e1003074.	1.5	94
22	Mouse Proteome Analysis. Genome Research, 2003, 13, 1335-1344.	2.4	91
23	Multivalent Histone and DNA Engagement by a PHD/BRD/PWWP Triple Reader Cassette Recruits ZMYND8 to K14ac-Rich Chromatin. Cell Reports, 2016, 17, 2724-2737.	2.9	86
24	Proteome Analysis Database: online application of InterPro and CluSTr for the functional classification of proteins in whole genomes. Nucleic Acids Research, 2001, 29, 44-48.	6.5	84
25	Whole-genome sequencing of bladder cancers reveals somatic CDKN1A mutations and clinicopathological associations with mutation burden. Nature Communications, 2014, 5, 3756.	5.8	81
26	RASSF1A uncouples Wnt from Hippo signalling and promotes YAP mediated differentiation via p73. Nature Communications, 2018, 9, 424.	5.8	72
27	NOX1 loss-of-function genetic variants in patients with inflammatory bowel disease. Mucosal Immunology, 2018, 11, 562-574.	2.7	71
28	The Proteome Analysis database: a tool for the in silico analysis of whole proteomes. Nucleic Acids Research, 2003, 31, 414-417.	6.5	64
29	Human Accelerated Regions and Other Human-Specific Sequence Variations in the Context of Evolution and Their Relevance for Brain Development. Genome Biology and Evolution, 2018, 10, 166-188.	1.1	61
30	Applications of InterPro in protein annotation and genome analysis. Briefings in Bioinformatics, 2002, 3, 285-295.	3.2	54
31	The pss4 gene from Rhizobium leguminosarum bv viciae VF39: cloning, sequence and the possible role in polysaccharide production and nodule formation. Gene, 1994, 150, 111-116.	1.0	45
32	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.	0.5	40
33	Erythrocytosis associated with a novel missense mutation in the BPGM gene. Haematologica, 2014, 99, e201-e204.	1.7	35
34	A point mutation in the ion conduction pore of AMPA receptor GRIA3 causes dramatically perturbed sleep patterns as well as intellectual disability. Human Molecular Genetics, 2017, 26, 3869-3882.	1.4	35
35	Development and Evaluation of an Automated Annotation Pipeline and cDNA Annotation System. Genome Research, 2003, 13, 1542-1551.	2.4	34
36	PRMT5 promotes cancer cell migration and invasion through the E2F pathway. Cell Death and Disease, 2020, 11, 572.	2.7	20

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37	Arginine methylation expands the regulatory mechanisms and extends the genomic landscape under E2F control. Science Advances, 2019, 5, eaaw4640.	4.7	19
38	Neanderthal and Denisovan retroviruses in modern humans. Current Biology, 2013, 23, R994-R995.	1.8	17
39	Current challenges and possible future developments in personalized psychiatry with an emphasis on psychotic disorders. Heliyon, 2020, 6, e03990.	1.4	15
40	Interactive InterPro-based comparisons of proteins in whole genomes. Bioinformatics, 2002, 18, 374-375.	1.8	14
41	The Genome Sequence of Five Highly Pathogenic Isolates of Fusarium oxysporum f. sp. lini. Molecular Plant-Microbe Interactions, 2020, 33, 1112-1115.	1.4	14
42	Characterising a human endogenous retrovirus(HERV)-derived tumour-associated antigen: enriched RNA-Seq analysis of HERV-K(HML-2) in mantle cell lymphoma cell lines. Mobile DNA, 2020, 11, 9.	1.3	13
43	The Genetic Landscape of Fiber Flax. Frontiers in Plant Science, 2021, 12, 764612.	1.7	11
44	Interplay between RNA interference and heat shock response systems in Drosophila melanogaster. Open Biology, 2016, 6, 160224.	1.5	9
45	Proteome Complexity Measures Based on Counting of Domain-to-Protein Links for Replicative and Non-Replicative Domains., 2006,, 329-341.		9
46	A Genomic Blueprint of Flax Fungal Parasite Fusarium oxysporum f. sp. lini. International Journal of Molecular Sciences, 2021, 22, 2665.	1.8	6
47	Application of InterPro for the functional classification of the proteins of fish origin in SWISS-PROT and TrEMBL. Journal of Biosciences, 2001, 26, 277-284.	0.5	5
48	Mutations of TCF12, encoding a basic-helix-loop-helix partner of TWIST1, are a frequent cause of coronal craniosynostosis. Lancet, The, 2013, 381, S114.	6.3	5
49	A comprehensive dataset of flax (Linum uitatissimum L.) phenotypes. Data in Brief, 2021, 37, 107224.	0.5	5
50	A genome-wide association study identifies a gene network associated with paranoid schizophrenia and antipsychotics-induced tardive dyskinesia. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2021, 105, 110134.	2.5	4
51	Projection of gene-protein networks to the functional space of the proteome and its application to analysis of organism complexity. BMC Genomics, 2010, 11, S4.	1.2	3
52	Antisence oligodeoxyribonucleotides for fragments of the reverse transcriptase gene of the LINE-1 element of rats disturb the formation of long-term memory. Doklady Biochemistry and Biophysics, 2002, 383, 93-95.	0.3	2
53	Stochastic Effects in Retrotransposon Dynamics Revealed by Modeling under Competition for Cellular Resources. Life, 2021, 11, 1209.	1.1	2
54	An account of Fusarium wilt resistance in flax Linum usitatissimum: The disease severity data. Data in Brief, 2022, 41, 107869.	0.5	2

#	Article	IF	CITATIONS
55	Regions Of 3D Similarity in Potential ORF1 Prducts of Mobile Genetics Classes. Protein Engineering, Design and Selection, 0, , .	1.0	O
56	Supervised Learning-Aided Optimization of Expert-Driven Functional Protein Sequence Annotation. Lecture Notes in Computer Science, 2004, , 159-169.	1.0	0
57	InterPro as a new tool for complete genome analysis: An example of comparative analysis. Biophysics (Russian Federation), 2006, 51, 587-591.	0.2	O
58	Reactome - a knowledgebase of human biological pathways. Nature Precedings, 2009, , .	0.1	0
59	Aberration Of SF3B1 Results In Deregulated Splicing Of Key Genes and Pathways In Myelodysplastic Syndromes. Blood, 2013, 122, 2747-2747.	0.6	0