## Omer Gokcumen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9274696/publications.pdf

Version: 2024-02-01

59 papers 4,632 citations

218677 26 h-index 58 g-index

72 all docs 72 docs citations

times ranked

72

9091 citing authors

#	Article	IF	Citations
1	Similarity-Based Analysis of Allele Frequency Distribution among Multiple Populations Identifies Adaptive Genomic Structural Variants. Molecular Biology and Evolution, 2022, 39, .	8.9	6
2	Sex-specific phenotypic effects and evolutionary history of an ancient polymorphic deletion of the human growth hormone receptor. Science Advances, 2021, 7, eabi4476.	10.3	11
3	Protein acylation by saturated very long chain fatty acids and endocytosis are involved in necroptosis. Cell Chemical Biology, 2021, 28, 1298-1309.e7.	5.2	21
4	Evolutionary context of psoriatic immune skin response. Evolution, Medicine and Public Health, 2021, 9, 474-486.	2.5	6
5	An Evolutionary Perspective on the Impact of Genomic Copy Number Variation on Human Health. Journal of Molecular Evolution, 2020, 88, 104-119.	1.8	27
6	Human and Nonhuman Primate Lineage-Specific Footprints in the Salivary Proteome. Molecular Biology and Evolution, 2020, 37, 395-405.	8.9	19
7	Archaic hominin introgression into modern human genomes. American Journal of Physical Anthropology, 2020, 171, 60-73.	2.1	33
8	The Impact of Ancient Genome Studies in Archaeology. Annual Review of Anthropology, 2020, 49, 277-298.	1.5	13
9	Functional Specialization of Human Salivary Glands and Origins of Proteins Intrinsic to Human Saliva. Cell Reports, 2020, 33, 108402.	6.4	54
10	Analysis of Haplotypic Variation and Deletion Polymorphisms Point to Multiple Archaic Introgression Events, Including from Altai Neanderthal Lineage. Genetics, 2020, 215, 497-509.	2.9	17
11	Genome-Wide Selection Scan in an Arabian Peninsula Population Identifies a TNKS Haplotype Linked to Metabolic Traits and Hypertension. Genome Biology and Evolution, 2020, 12, 77-87.	2.5	11
12	Loss-of-function tolerance of enhancers in the human genome. PLoS Genetics, 2020, 16, e1008663.	3.5	12
13	Resolving the Insertion Sites of Polymorphic Duplications Reveals a HERC2 Haplotype under Selection. Genome Biology and Evolution, 2019, 11, 1679-1690.	2.5	6
14	Filamentation Regulatory Pathways Control Adhesion-Dependent Surface Responses in Yeast. Genetics, 2019, 212, 667-690.	2.9	20
15	Fine-Scale Characterization of Genomic Structural Variation in the Human Genome Reveals Adaptive and Biomedically Relevant Hotspots. Genome Biology and Evolution, 2019, 11, 1136-1151.	2.5	41
16	Shades of complexity: New perspectives on the evolution and genetic architecture of human skin. American Journal of Physical Anthropology, 2019, 168, 4-26.	2.1	45
17	Independent amylase gene copy number bursts correlate with dietary preferences in mammals. ELife, 2019, 8, .	6.0	78
18	Complex Haplotypes of <i>GSTM1</i> Gene Deletions Harbor Signatures of a Selective Sweep in East Asian Populations. G3: Genes, Genomes, Genetics, 2018, 8, 2953-2966.	1.8	8

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19	The Year In Genetic Anthropology: New Lands, New Technologies, New Questions. American Anthropologist, 2018, 120, 266-277.	1.4	5
20	An evolutionary transcriptomics approach links CD36 to membrane remodeling in replicative senescence. Molecular Omics, 2018, 14, 237-246.	2.8	12
21	Structural Variants in Ancient Genomes. Population Genomics, 2018, , 375-391.	0.5	1
22	Complex evolution of the GSTM gene family involves sharing of GSTM1 deletion polymorphism in humans and chimpanzees. BMC Genomics, 2018, 19, 293.	2.8	12
23	Regulation of lipids is central to replicative senescence. Molecular BioSystems, 2017, 13, 498-509.	2.9	69
24	Variation and Functional Impact of Neanderthal Ancestry in Western Asia. Genome Biology and Evolution, 2017, 9, 3516-3524.	2.5	14
25	Archaic Hominin Introgression in Africa Contributes to Functional Salivary MUC7 Genetic Variation. Molecular Biology and Evolution, 2017, 34, 2704-2715.	8.9	57
26	Segmental duplications and evolutionary acquisition of UV damage response in the SPATA31 gene family of primates and humans. BMC Genomics, 2017, 18, 222.	2.8	8
27	VCFtoTree: a user-friendly tool to construct locus-specific alignments and phylogenies from thousands of anthropologically relevant genome sequences. BMC Bioinformatics, 2017, 18, 426.	2.6	14
28	The Multiple Histories of Western Asia: Perspectives from Ancient and Modern Genomes. Human Biology, 2017, 89, 107.	0.2	6
29	Evolution, Function, and Deconstructing Histories: A New Generation of Anthropological Genetics. Human Biology, 2017, 89, 5.	0.2	2
30	Atopic Dermatitis Susceptibility Variants in Filaggrin (i) Hitchhike (i) Hornerin Selective Sweep. Genome Biology and Evolution, 2016, 8, 3240-3255.	2.5	35
31	Recent evolution of the salivary mucin MUC7. Scientific Reports, 2016, 6, 31791.	3.3	30
32	The psoriasis-associated deletion of late cornified envelope genes LCE3B and LCE3C has been maintained under balancing selection since Human Denisovan divergence. BMC Evolutionary Biology, 2016, 16, 265.	3.2	35
33	Next-generation sequencing-based detection of germline L1-mediated transductions. BMC Genomics, 2016, 17, 342.	2.8	7
34	The Evolution and Functional Impact of Human Deletion Variants Shared with Archaic Hominin Genomes. Molecular Biology and Evolution, 2015, 32, 1008-1019.	8.9	45
35	Global Survey of Variation in a Human Olfactory Receptor Gene Reveals Signatures of Non-Neutral Evolution. Chemical Senses, 2015, 40, 481-488.	2.0	31
36	Whole genome sequencing of Turkish genomes reveals functional private alleles and impact of genetic interactions with Europe, Asia and Africa. BMC Genomics, 2014, 15, 963.	2.8	46

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37	Geographic Distribution and Adaptive Significance of Genomic Structural Variants: An Anthropological Genetics Perspective. Human Biology, 2014, 86, 260.	0.2	11
38	Impact of constitutional copy number variants on biological pathway evolution. BMC Evolutionary Biology, 2013, 13, 19.	3.2	25
39	Primate genome architecture influences structural variation mechanisms and functional consequences. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 15764-15769.	7.1	80
40	Analysis of variable retroduplications in human populations suggests coupling of retrotransposition to cell division. Genome Research, 2013, 23, 2042-2052.	5.5	52
41	Balancing Selection on a Regulatory Region Exhibiting Ancient Variation That Predates Human–Neandertal Divergence. PLoS Genetics, 2013, 9, e1003404.	3.5	26
42	Vif Proteins from Diverse Primate Lentiviral Lineages Use the Same Binding Site in APOBEC3G. Journal of Virology, 2013, 87, 11861-11871.	3.4	36
43	Extensive genetic diversity and substructuring among zebrafish strains revealed through copy number variant analysis. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 529-534.	7.1	102
44	Regulatory element copy number differences shape primate expression profiles. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 12656-12661.	7.1	37
45	Landscape of Somatic Retrotransposition in Human Cancers. Science, 2012, 337, 967-971.	12.6	631
46	Mitochondrial DNA and Y Chromosome Variation Provides Evidence for a Recent Common Ancestry between Native Americans and Indigenous Altaians. American Journal of Human Genetics, 2012, 90, 229-246.	6.2	146
47	Mitochondrial DNA and Y Chromosome Variation Provides Evidence for a Recent Common Ancestry between Native Americans and Indigenous Altaians. American Journal of Human Genetics, 2012, 90, 573.	6.2	6
48	Exploring the role of copy number variants in human adaptation. Trends in Genetics, 2012, 28, 245-257.	6.7	126
49	Extensive genomic and transcriptional diversity identified through massively parallel DNA and RNA sequencing of eighteen Korean individuals. Nature Genetics, 2011, 43, 745-752.	21.4	121
50	Refinement of primate copy number variationhotspots identifies candidate genomic regions evolving under positive selection. Genome Biology, 2011, 12, R52.	8.8	58
51	Biological Ancestries, Kinship Connections, and Projected Identities in Four Central Anatolian Settlements: Insights from Culturally Contextualized Genetic Anthropology. American Anthropologist, 2011, 113, 116-131.	1.4	22
52	Origins and functional impact of copy number variation in the human genome. Nature, 2010, 464, 704-712.	27.8	1,721
53	Discovery of common Asian copy number variants using integrated high-resolution array CGH and massively parallel DNA sequencing. Nature Genetics, 2010, 42, 400-405.	21.4	179
54	Y-chromosome and autosomal STR diversity in four proximate settlements in Central Anatolia. Forensic Science International: Genetics, 2010, 4, e135-e137.	3.1	13

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55	A highly annotated whole-genome sequence of a Korean individual. Nature, 2009, 460, 1011-1015.	27.8	295
56	Copy number variants (CNVs) in primate species using array-based comparative genomic hybridization. Methods, 2009, 49, 18-25.	3.8	22
57	Genetic variation in the enigmatic Altaian Kazakhs of Southâ€Central Russia: Insights into Turkic population history. American Journal of Physical Anthropology, 2008, 136, 278-293.	2.1	46
58	Russian Old Believers: Genetic Consequences of Their Persecution and Exile, as Shown by Mitochondrial DNA Evidence. Human Biology, 2008, 80, 203-237.	0.2	10
59	GENETİK VE KAMUSAL ALAN. Ankara Üniversitesi Dil Ve Tarih-CoÄŸrafya FakÃ⅓ltesi Dergisi, 0, , 019-031.	0.8	1