Gil Atzmon

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/907005/publications.pdf

Version: 2024-02-01

117453 79541 12,297 78 34 73 citations h-index g-index papers 80 80 80 26918 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|----|--|--------------|-----------|
| 1 | Rare coding variants in 35 genes associate with circulating lipid levels—A multi-ancestry analysis of 170,000 exomes. American Journal of Human Genetics, 2022, 109, 81-96. | 2.6 | 24 |
| 2 | Greater effect of polygenic risk score for Alzheimer's disease among younger cases who are apolipoprotein E-ε4 carriers. Neurobiology of Aging, 2021, 99, 101.e1-101.e9. | 1.5 | 16 |
| 3 | The Hypothalamic-Pituitary-Testicular Axis in Exceptionally Old Men. Journal of the Endocrine Society, 2021, 5, A727-A727. | 0.1 | O |
| 4 | Novel ultra-rare exonic variants identified in a founder population implicate cadherins in schizophrenia. Neuron, 2021, 109, 1465-1478.e4. | 3.8 | 21 |
| 5 | Determinants of penetrance and variable expressivity in monogenic metabolic conditions across 77,184 exomes. Nature Communications, 2021, 12, 3505. | 5 . 8 | 49 |
| 6 | Genetic signature of human longevity in PKC and NFâ€ÎºB signaling. Aging Cell, 2021, 20, e13362. | 3.0 | 12 |
| 7 | Rare genetic coding variants associated with human longevity and protection against age-related diseases. Nature Aging, 2021, 1, 783-794. | 5 . 3 | 22 |
| 8 | Redox-mediated regulation of aging and healthspan by an evolutionarily conserved transcription factor HLH-2/Tcf3/E2A. Redox Biology, 2020, 32, 101448. | 3.9 | 10 |
| 9 | Prevalent skin cancer and conservative faith may be linked with cognitive impairment in Ashkenazi Jewish exceptionally longâ€lived individuals. Alzheimer's and Dementia, 2020, 16, e046002. | 0.4 | O |
| 10 | The mutational constraint spectrum quantified from variation in 141,456 humans. Nature, 2020, 581, 434-443. | 13.7 | 6,140 |
| 11 | Transcript expression-aware annotation improves rare variant interpretation. Nature, 2020, 581, 452-458. | 13.7 | 142 |
| 12 | Pregnancy as a model for aging. Ageing Research Reviews, 2020, 62, 101093. | 5.0 | 20 |
| 13 | Breaking the Glass Ceiling. Gerontology, 2020, 66, 309-314. | 1.4 | 1 |
| 14 | Exceptionally Long-Lived Individuals (ELLI) Demonstrate Slower Aging Rate Calculated by DNA Methylation Clocks as Possible Modulators for Healthy Longevity. International Journal of Molecular Sciences, 2020, 21, 615. | 1.8 | 18 |
| 15 | The effects of environmental stressors on candidate aging associated genes. Experimental Gerontology, 2020, 137, 110952. | 1.2 | 5 |
| 16 | The mitochondrial derived peptide humanin is a regulator of lifespan and healthspan. Aging, 2020, 12, 11185-11199. | 1.4 | 67 |
| 17 | Senescence and Longevity of Sea Urchins. Genes, 2020, 11, 573. | 1.0 | 7 |
| 18 | A meta-analysis of genome-wide association studies identifies multiple longevity genes. Nature Communications, 2019, 10, 3669. | 5.8 | 214 |

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|----|---|------|-----------|
| 19 | Telomeres and Longevity: A Cause or an Effect?. International Journal of Molecular Sciences, 2019, 20, 3233. | 1.8 | 28 |
| 20 | Exome sequencing of 20,791Âcases of type 2 diabetes and 24,440Âcontrols. Nature, 2019, 570, 71-76. | 13.7 | 248 |
| 21 | Empirical design of a variant quality control pipeline for whole genome sequencing data using replicate discordance. Scientific Reports, 2019, 9, 16156. | 1.6 | 14 |
| 22 | Screening Human Embryos for Polygenic Traits Has Limited Utility. Cell, 2019, 179, 1424-1435.e8. | 13.5 | 78 |
| 23 | PopCluster: an algorithm to identify genetic variants with ethnicity-dependent effects. Bioinformatics, 2019, 35, 3046-3054. | 1.8 | 3 |
| 24 | Evaluating the contribution of rare variants to type 2 diabetes and related traits using pedigrees. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 379-384. | 3.3 | 28 |
| 25 | Functional variants in the <i>LRRK2</i> gene confer shared effects on risk for Crohn's disease and Parkinson's disease. Science Translational Medicine, 2018, 10, . | 5.8 | 273 |
| 26 | High-depth whole genome sequencing of an Ashkenazi Jewish reference panel: enhancing sensitivity, accuracy, and imputation. Human Genetics, 2018, 137, 343-355. | 1.8 | 24 |
| 27 | Effects of FOXO3 Polymorphisms on Survival to Extreme Longevity in Four Centenarian Studies. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2018, 73, 1439-1447. | 1.7 | 32 |
| 28 | <i>>FOXP3</i> mutations causing earlyâ€onset insulinâ€requiring diabetes but without other features of immune dysregulation, polyendocrinopathy, enteropathy, Xâ€linked syndrome. Pediatric Diabetes, 2018, 19, 388-392. | 1.2 | 25 |
| 29 | Body mass index is negatively associated with telomere length: a collaborative cross-sectional meta-analysis of 87 observational studies. American Journal of Clinical Nutrition, 2018, 108, 453-475. | 2.2 | 137 |
| 30 | Genomic Instabilities, Cellular Senescence, and Aging: In Vitro, In Vivo and Aging-Like Human Syndromes. Frontiers in Medicine, 2018, 5, 104. | 1.2 | 60 |
| 31 | Genetic Insights Into Frailty: Association of 9p21-23 Locus With Frailty. Frontiers in Medicine, 2018, 5, 105. | 1.2 | 19 |
| 32 | Insights into the genetic epidemiology of Crohn's and rare diseases in the Ashkenazi Jewish population. PLoS Genetics, 2018, 14, e1007329. | 1.5 | 66 |
| 33 | Activation-Induced Autophagy Is Preserved in CD4+ T-Cells in Familial Longevity. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2017, 72, 1201-1206. | 1.7 | 35 |
| 34 | The GH receptor exon 3 deletion is a marker of male-specific exceptional longevity associated with increased GH sensitivity and taller stature. Science Advances, 2017, 3, e1602025. | 4.7 | 47 |
| 35 | A Low-Frequency Inactivating <i>AKT2</i> Variant Enriched in the Finnish Population Is Associated With Fasting Insulin Levels and Type 2 Diabetes Risk. Diabetes, 2017, 66, 2019-2032. | 0.3 | 47 |
| 36 | Effect of Exceptional Parental Longevity and Lifestyle Factors on Prevalence of Cardiovascular Disease in Offspring. American Journal of Cardiology, 2017, 120, 2170-2175. | 0.7 | 27 |

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|----|---|------|-----------|
| 37 | Association of anti-inflammatory cytokine IL10 polymorphisms with motoric cognitive risk syndrome in an Ashkenazi Jewish population. Neurobiology of Aging, 2017, 58, 238.e1-238.e8. | 1.5 | 22 |
| 38 | Sequence data and association statistics from 12,940 type 2 diabetes cases and controls. Scientific Data, 2017, 4, 170179. | 2.4 | 31 |
| 39 | [P3–118]: INCREASED BURDEN OF RARE LOSSâ€OFâ€FUNCTION VARIANTS IN ALZHEIMER's DISEASE PATIENTS COMPARED TO CENTENARIANS. Alzheimer's and Dementia, 2017, 13, P980. | 0.4 | O |
| 40 | The Genetics of Bene Israel from India Reveals Both Substantial Jewish and Indian Ancestry. PLoS ONE, 2016, 11, e0152056. | 1.1 | 17 |
| 41 | The genetic architecture of type 2 diabetes. Nature, 2016, 536, 41-47. | 13.7 | 952 |
| 42 | A Frameshift in CSF2RB Predominant Among Ashkenazi Jews Increases Risk for Crohn's Disease and Reduces Monocyte Signaling via GM-CSF. Gastroenterology, 2016, 151, 710-723.e2. | 0.6 | 51 |
| 43 | Genetic landscape of APOE in human longevity revealed by high-throughput sequencing. Mechanisms of Ageing and Development, 2016, 155, 7-9. | 2.2 | 35 |
| 44 | New Locus for Skin Intrinsic Fluorescence in Type 1 Diabetes Also Associated With Blood and Skin Glycated Proteins. Diabetes, 2016, 65, 2060-2071. | 0.3 | 10 |
| 45 | <scp>GWAS</scp> analysis of handgrip and lower body strength in older adults in the <scp>CHARGE</scp> consortium. Aging Cell, 2016, 15, 792-800. | 3.0 | 51 |
| 46 | The genetic history of Cochin Jews from India. Human Genetics, 2016, 135, 1127-1143. | 1.8 | 12 |
| 47 | Genome-wide analysis identifies 12 loci influencing human reproductive behavior. Nature Genetics, 2016, 48, 1462-1472. | 9.4 | 284 |
| 48 | Differential burden of rare protein truncating variants in Alzheimer's disease patients compared to centenarians. Human Molecular Genetics, 2016, 25, ddw150. | 1.4 | 10 |
| 49 | Transancestral fine-mapping of four type 2 diabetes susceptibility loci highlights potential causal regulatory mechanisms. Human Molecular Genetics, 2016, 25, 2070-2081. | 1.4 | 21 |
| 50 | Expanded genetic screening panel for the Ashkenazi Jewish population. Genetics in Medicine, 2016, 18, 522-528. | 1.1 | 33 |
| 51 | Positive attitude toward life, emotional expression, self-rated health, and depressive symptoms among centenarians and near-centenarians. Aging and Mental Health, 2016, 20, 930-939. | 1.5 | 41 |
| 52 | Genome-Wide Scan Informed by Age-Related Disease Identifies Loci for Exceptional Human Longevity. PLoS Genetics, 2015, 11, e1005728. | 1.5 | 128 |
| 53 | Genetic variation in Sirtuin 1 (SIRT1) is associated with lipid profiles but not with longevity in Ashkenazi Jews. Translational Research, 2015, 165, 480-481. | 2.2 | 9 |
| 54 | Genetics, lifestyle and longevity: Lessons from centenarians. Applied & Translational Genomics, 2015, 4, 23-32. | 2.1 | 90 |

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|----|--|-----|-----------|
| 55 | Genotyping of geographically diverse Druze trios reveals substructure and a recent bottleneck. European Journal of Human Genetics, 2015, 23, 1093-1099. | 1.4 | 10 |
| 56 | The influence of gender on inheritance of exceptional longevity. Aging, 2015, 7, 412-418. | 1.4 | 12 |
| 57 | Identification of Genes Promoting Skin Youthfulness by Genome-Wide Association Study. Journal of Investigative Dermatology, 2014, 134, 651-657. | 0.3 | 30 |
| 58 | Novel candidate genes putatively involved in stress fracture predisposition detected by whole-exome sequencing. Genetical Research, 2014, 96, e004. | 0.3 | 14 |
| 59 | Genome-wide mapping of IBD segments in an Ashkenazi PD cohort identifies associated haplotypes. Human Molecular Genetics, 2014, 23, 4693-4702. | 1.4 | 49 |
| 60 | Sequencing an Ashkenazi reference panel supports population-targeted personal genomics and illuminates Jewish and European origins. Nature Communications, 2014, 5, 4835. | 5.8 | 156 |
| 61 | Disrupting Mitochondrial–Nuclear Coevolution Affects OXPHOS Complex I Integrity and Impacts Human Health. Genome Biology and Evolution, 2014, 6, 2665-2680. | 1.1 | 68 |
| 62 | Loss-of-function mutations in SLC30A8 protect against type 2 diabetes. Nature Genetics, 2014, 46, 357-363. | 9.4 | 428 |
| 63 | Clonal Hematopoiesis with Somatic Mutations Is a Common, Age-Related Condition Associated with Adverse Outcomes. Blood, 2014, 124, 840-840. | 0.6 | 1 |
| 64 | Association of telomere length (TL) with clinical outcomes in patients with colorectal carcinoma (CRC) Journal of Clinical Oncology, 2013, 31, 418-418. | 0.8 | 0 |
| 65 | Association of telomere length with clinical outcomes in patients with colorectal carcinoma Journal of Clinical Oncology, 2013, 31, e14540-e14540. | 0.8 | 0 |
| 66 | A Genome-Wide Scan of Ashkenazi Jewish Crohn's Disease Suggests Novel Susceptibility Loci. PLoS Genetics, 2012, 8, e1002559. | 1.5 | 144 |
| 67 | A genome-wide association study of aging. Neurobiology of Aging, 2011, 32, 2109.e15-2109.e28. | 1.5 | 127 |
| 68 | Obesity/diabetesâ€associated gene screening in rhesus monkeys. FASEB Journal, 2011, 25, 859.4. | 0.2 | 0 |
| 69 | Abraham's Children in the Genome Era: Major Jewish Diaspora Populations Comprise Distinct Genetic Clusters with Shared Middle Eastern Ancestry. American Journal of Human Genetics, 2010, 86, 850-859. | 2.6 | 217 |
| 70 | Genetic variation in human telomerase is associated with telomere length in Ashkenazi centenarians. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 1710-1717. | 3.3 | 203 |
| 71 | Genetic Predisposition to Elevated Serum Thyrotropin Is Associated with Exceptional Longevity. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 4768-4775. | 1.8 | 132 |
| 72 | Extreme Longevity Is Associated with Increased Serum Thyrotropin. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 1251-1254. | 1.8 | 223 |

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|----|--|-----|----------|
| 73 | Adiponectin Levels and Genotype: A Potential Regulator of Life Span in Humans. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2008, 63, 447-453. | 1.7 | 121 |
| 74 | Lipoprotein Genotype and Conserved Pathway for Exceptional Longevity in Humans. PLoS Biology, 2006, 4, e113. | 2.6 | 197 |
| 75 | Biological evidence for inheritance of exceptional longevity. Mechanisms of Ageing and Development, 2005, 126, 341-345. | 2.2 | 100 |
| 76 | Buffering Mechanisms in Aging: A systems approach towards uncovering the genetic component of aging. PLoS Computational Biology, 2005, preprint, e170. | 1.5 | 2 |
| 77 | Clinical Phenotype of Families with Longevity. Journal of the American Geriatrics Society, 2004, 52, 274-277. | 1.3 | 174 |
| 78 | Plasma HDL Levels Highly Correlate With Cognitive Function in Exceptional Longevity. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2002, 57, M712-M715. | 1.7 | 130 |