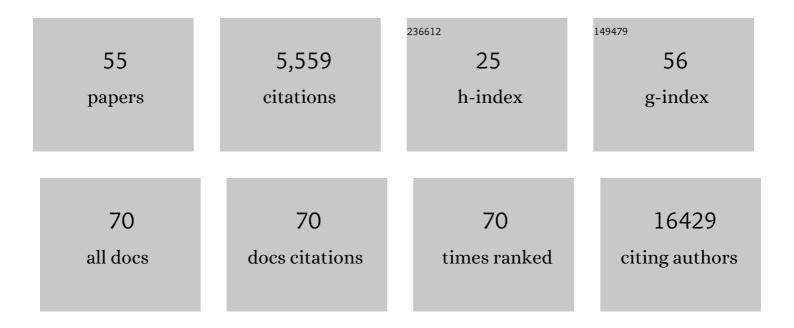
## Massimo Mezzavilla

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6702487/publications.pdf Version: 2024-02-01



| #  | Article   | IF   | CITATIONS |
|----|---|------|-----------|
| 1  | Runs of homozygosity are associated with staging of periodontitis in isolated populations. Human<br>Molecular Genetics, 2021, 30, 1154-1159.  | 1.4  | 3         |
| 2  | Natural human knockouts and Mendelian disorders: deep phenotyping in Italian isolates. European<br>Journal of Human Genetics, 2021, 29, 1272-1281.  | 1.4  | 6         |
| 3  | The landscape of autosomal-recessive pathogenic variants in European populations reveals phenotype-specific effects. American Journal of Human Genetics, 2021, 108, 608-619.                                      | 2.6  | 36        |
| 4  | Genetic insights into biological mechanisms governing human ovarian ageing. Nature, 2021, 596,<br>393-397.  | 13.7 | 183       |
| 5  | Poking COVID-19: Insights on Genomic Constraints among Immune-Related Genes between Qatari and Italian Populations. Genes, 2021, 12, 1842.  | 1.0  | 1         |
| 6  | Fine-scale population structure and demographic history of British Pakistanis. Nature<br>Communications, 2021, 12, 7189.  | 5.8  | 21        |
| 7  | A bird's-eye view of Italian genomic variation through whole-genome sequencing. European Journal of<br>Human Genetics, 2020, 28, 435-444.   | 1.4  | 29        |
| 8  | Glucagon-like peptide-1 receptor and sarcoglycan delta genetic variants can affect cardiovascular risk<br>in chronic kidney disease patients under hemodialysis. CKJ: Clinical Kidney Journal, 2020, 13, 666-673. | 1.4  | 3         |
| 9  | Deleterious variants in genes associated with bone mineral density are linked to susceptibility to periodontitis development. Meta Gene, 2020, 24, 100670.  | 0.3  | 2         |
| 10 | A population-based approach for gene prioritization in understanding complex traits. Human Genetics,<br>2020, 139, 647-655.   | 1.8  | 7         |
| 11 | Mutations in <i>PLS1</i> , encoding fimbrin, cause autosomal dominant nonsyndromic hearing loss.<br>Human Mutation, 2019, 40, 2286-2295.  | 1.1  | 23        |
| 12 | A transcriptomic atlas of mammalian olfactory mucosae reveals an evolutionary influence on food<br>odor detection in humans. Science Advances, 2019, 5, eaax0396.   | 4.7  | 59        |
| 13 | Next generation sequencing study in a cohort of Italian patients with syndromic hearing loss. Hearing Research, 2019, 381, 107769.  | 0.9  | 7         |
| 14 | Establishment and equilibrium levels of deleterious mutations in large populations. Scientific<br>Reports, 2019, 9, 10384.  | 1.6  | 2         |
| 15 | Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.   | 5.8  | 84        |
| 16 | Investigation of the link between PROP taste perception and vegetables consumption using FAOSTAT data. International Journal of Food Sciences and Nutrition, 2019, 70, 484-490.                                   | 1.3  | 6         |
| 17 | Response to Giem. American Journal of Human Genetics, 2018, 102, 331.   | 2.6  | 1         |
| 18 | MMAB, a novel candidate gene to be screened in the molecular diagnosis of Mevalonate Kinase<br>Deficiency. Rheumatology International, 2018, 38, 121-127.   | 1.5  | 1         |

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|----|---|-----|-----------|
| 19 | Genetic Landscape of Slovenians: Past Admixture and Natural Selection Pattern. Frontiers in Genetics, 2018, 9, 551.   | 1.1 | 8         |
| 20 | Demographic History and Genetic Adaptation in the Himalayan Region Inferred from Genome-Wide SNP<br>Genotypes of 49 Populations. Molecular Biology and Evolution, 2018, 35, 1916-1933.                | 3.5 | 36        |
| 21 | Mutations in L-type amino acid transporter-2 support SLC7A8 as a novel gene involved in age-related hearing loss. ELife, 2018, 7, .   | 2.8 | 38        |
| 22 | Investigation of genetic variation and lifestyle determinants in vitamin D levels in Arab individuals.<br>Journal of Translational Medicine, 2018, 16, 20.  | 1.8 | 9         |
| 23 | FineMAV: prioritizing candidate genetic variants driving local adaptations in human populations.<br>Genome Biology, 2018, 19, 5.  | 3.8 | 20        |
| 24 | Whole-genome sequencing reveals new insights into age-related hearing loss: cumulative effects, pleiotropy and the role of selection. European Journal of Human Genetics, 2018, 26, 1167-1179.        | 1.4 | 22        |
| 25 | Continuity and Admixture in the Last Five Millennia of Levantine History from Ancient Canaanite and<br>Present-Day Lebanese Genome Sequences. American Journal of Human Genetics, 2017, 101, 274-282. | 2.6 | 102       |
| 26 | "Like sugar in milk― reconstructing the genetic history of the Parsi population. Genome Biology, 2017,<br>18, 110.  | 3.8 | 29        |
| 27 | Enrichment of low-frequency functional variants revealed by whole-genome sequencing of multiple isolated European populations. Nature Communications, 2017, 8, 15927.                                 | 5.8 | 64        |
| 28 | A reference panel of 64,976 haplotypes for genotype imputation. Nature Genetics, 2016, 48, 1279-1283.   | 9.4 | 2,421     |
| 29 | Global diversity in the TAS2R38 bitter taste receptor: revisiting a classic evolutionary PROPosal.<br>Scientific Reports, 2016, 6, 25506.   | 1.6 | 69        |
| 30 | Chad Genetic Diversity Reveals an African History Marked by Multiple Holocene Eurasian Migrations.<br>American Journal of Human Genetics, 2016, 99, 1316-1324.  | 2.6 | 37        |
| 31 | Wide distribution and altitude correlation of an archaic high-altitude-adaptive EPAS1 haplotype in the<br>Himalayas. Human Genetics, 2016, 135, 393-402.  | 1.8 | 41        |
| 32 | Genetic evidence for an origin of the Armenians from Bronze Age mixing of multiple populations.<br>European Journal of Human Genetics, 2016, 24, 931-936.   | 1.4 | 44        |
| 33 | Response to Hellenthal etÂal.:. American Journal of Human Genetics, 2016, 98, 398.  | 2.6 | 1         |
| 34 | Understanding the role of personality and alexithymia in food preferences and PROP taste perception.<br>Physiology and Behavior, 2016, 157, 72-78.  | 1.0 | 37        |
| 35 | Tracking of the origin of recurrent mutations of the BRCA1 and BRCA2 genes in the North-East of Italy and improved mutation analysis strategy. BMC Medical Genetics, 2016, 17, 11.                    | 2.1 | 16        |
| 36 | Ancient DNA and the rewriting of human history: be sparing with Occam's razor. Genome Biology,<br>2016, 17, 1.  | 3.8 | 1,335     |

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|----|--|-----|-----------|
| 37 | Early modern human dispersal from Africa: genomic evidence for multiple waves of migration.<br>Investigative Genetics, 2015, 6, 13.  | 3.3 | 34        |
| 38 | Neon: An R Package to Estimate Human Effective Population Size and Divergence Time from Patterns of Linkage Disequilibrium between SNPS. Journal of Computer Science and Systems Biology, 2015, 8, . | 0.0 | 28        |
| 39 | Increased Rate of Deleterious Variants in Long Runs of Homozygosity of an Inbred Population from<br>Qatar. Human Heredity, 2015, 79, 14-19.  | 0.4 | 28        |
| 40 | Analysis of functional variants reveals new candidate genes associated with alexithymia. Psychiatry Research, 2015, 227, 363-365.  | 1.7 | 12        |
| 41 | The Kalash Genetic Isolate: Ancient Divergence, Drift, and Selection. American Journal of Human<br>Genetics, 2015, 96, 775-783.  | 2.6 | 46        |
| 42 | Population genetic differentiation of height and body mass index across Europe. Nature Genetics, 2015, 47, 1357-1362.  | 9.4 | 227       |
| 43 | Insights into the origin of rare haplogroup C3* Y chromosomes in South America from high-density autosomal SNP genotyping. Forensic Science International: Genetics, 2015, 15, 115-120.              | 1.6 | 10        |
| 44 | Assessment of the Olfactory Function in Italian Patients with Type 3 von Willebrand Disease Caused by a Homozygous 253 Kb Deletion Involving VWF and TMEM16B/ANO2. PLoS ONE, 2015, 10, e0116483.     | 1.1 | 7         |
| 45 | A Population-Based Approach to Study the Impact of PROP Perception on Food Liking in Populations along the Silk Road. PLoS ONE, 2014, 9, e91716.   | 1.1 | 34        |
| 46 | Genetic characterization of Greek population isolates reveals strong genetic drift at missense and trait-associated variants. Nature Communications, 2014, 5, 5345.                                  | 5.8 | 60        |
| 47 | Consanguinity and Hereditary Hearing Loss in Qatar. Human Heredity, 2014, 77, 175-182.   | 0.4 | 15        |
| 48 | Genetic landscape of populations along the Silk Road: admixture and migration patterns. BMC Genetics, 2014, 15, 131.   | 2.7 | 24        |
| 49 | Insight into genetic determinants of resting heart rate. Gene, 2014, 545, 170-174.   | 1.0 | 7         |
| 50 | Evidence for past and present hybridization in three <scp>A</scp> ntarctic icefish species provides new perspectives on an evolutionary radiation. Molecular Ecology, 2013, 22, 5148-5161.           | 2.0 | 29        |
| 51 | Genetic characterization of northeastern Italian population isolates in the context of broader<br>European genetic diversity. European Journal of Human Genetics, 2013, 21, 659-665.                 | 1.4 | 64        |
| 52 | High variance in reproductive success generates a false signature of a genetic bottleneck in populations of constant size: a simulation study. BMC Bioinformatics, 2013, 14, 309.                    | 1.2 | 29        |
| 53 | Genetics of eye colours in different rural populations on the Silk Road. European Journal of Human<br>Genetics, 2013, 21, 1320-1323.   | 1.4 | 11        |
| 54 | Population genetic structure and gene flow patterns between populations of the Antarctic icefish<br><i>Chionodraco rastrospinosus</i> . Journal of Biogeography, 2012, 39, 1361-1372.                | 1.4 | 27        |

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| 55 | Gone with the currents: lack of genetic differentiation at the circum-continental scale in the Antarctic krill Euphausia superba. BMC Genetics, 2011, 12, 32. | 2.7 | 51        |