

Andrew J Pakstis

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

2,656
citations

394286

19
h-index

345118

36
g-index

40
all docs

40
docs citations

40
times ranked

2331
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | The world-wide distribution of allele frequencies at the human dopamine D4 receptor locus. <i>Human Genetics</i> , 1996, 98, 91-101. | 1.8 | 429 |
| 2 | Evidence against linkage of schizophrenia to markers on chromosome 5 in a northern Swedish pedigree. <i>Nature</i> , 1988, 336, 167-170. | 13.7 | 405 |
| 3 | Progress toward an efficient panel of SNPs for ancestry inference. <i>Forensic Science International: Genetics</i> , 2014, 10, 23-32. | 1.6 | 211 |
| 4 | A global survey of haplotype frequencies and linkage disequilibrium at the DRD2 locus. <i>Human Genetics</i> , 1998, 103, 211-227. | 1.8 | 197 |
| 5 | SNPs for a universal individual identification panel. <i>Human Genetics</i> , 2010, 127, 315-324. | 1.8 | 194 |
| 6 | Current sequencing technology makes microhaplotypes a powerful new type of genetic marker for forensics. <i>Forensic Science International: Genetics</i> , 2014, 12, 215-224. | 1.6 | 182 |
| 7 | Evaluating 130 microhaplotypes across a global set of 83 populations. <i>Forensic Science International: Genetics</i> , 2017, 29, 29-37. | 1.6 | 117 |
| 8 | Copy Number Variation in Obsessive-Compulsive Disorder and Tourette Syndrome: A Cross-Disorder Study. <i>Journal of the American Academy of Child and Adolescent Psychiatry</i> , 2014, 53, 910-919. | 0.3 | 111 |
| 9 | Candidate SNPs for a universal individual identification panel. <i>Human Genetics</i> , 2007, 121, 305-317. | 1.8 | 96 |
| 10 | Genome scan for linkage to Gilles de la Tourette syndrome. , 1999, 88, 437-445. | | 82 |
| 11 | DRD2 Haplotypes Containing the TaqI A1 Allele: Implications for Alcoholism Research. <i>Alcoholism: Clinical and Experimental Research</i> , 1996, 20, 697-705. | 1.4 | 63 |
| 12 | A panel of 74 AISNPs: Improved ancestry inference within Eastern Asia. <i>Forensic Science International: Genetics</i> , 2016, 23, 101-110. | 1.6 | 63 |
| 13 | Ancestry inference of 96 population samples using microhaplotypes. <i>International Journal of Legal Medicine</i> , 2018, 132, 703-711. | 1.2 | 48 |
| 14 | Mini-haplotypes as lineage informative SNPs and ancestry inference SNPs. <i>European Journal of Human Genetics</i> , 2012, 20, 1148-1154. | 1.4 | 45 |
| 15 | Validation of novel forensic DNA markers using multiplex microhaplotype sequencing. <i>Forensic Science International: Genetics</i> , 2020, 47, 102275. | 1.6 | 42 |
| 16 | 52 additional reference population samples for the 55 AISNP panel. <i>Forensic Science International: Genetics</i> , 2015, 19, 269-271. | 1.6 | 41 |
| 17 | Improving ancestry distinctions among Southwest Asian populations. <i>Forensic Science International: Genetics</i> , 2018, 35, 14-20. | 1.6 | 40 |
| 18 | Selecting microhaplotypes optimized for different purposes. <i>Electrophoresis</i> , 2018, 39, 2815-2823. | 1.3 | 39 |

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|----|---|-----|-----------|
| 19 | Increasing the reference populations for the 55 AISNP panel: the need and benefits. <i>International Journal of Legal Medicine</i> , 2017, 131, 913-917. | 1.2 | 38 |
| 20 | Genetic variation in Tunisia in the context of human diversity worldwide. <i>American Journal of Physical Anthropology</i> , 2016, 161, 62-71. | 2.1 | 29 |
| 21 | Genetic relationships of European, Mediterranean, and SW Asian populations using a panel of 55 AISNPs. <i>European Journal of Human Genetics</i> , 2019, 27, 1885-1893. | 1.4 | 22 |
| 22 | The redesigned Forensic Research/Reference on Genetics-knowledge base, FROG-kb. <i>Forensic Science International: Genetics</i> , 2018, 33, 33-37. | 1.6 | 21 |
| 23 | Development of a map of chromosome 11p. <i>Genetic Epidemiology</i> , 1986, 3, 153-158. | 0.6 | 17 |
| 24 | Detection of a large CTG/CAG trinucleotide repeat expansion in a Danish schizophrenia kindred. , 1997, 74, 546-548. | | 15 |
| 25 | Population relationships based on 170 ancestry SNPs from the combined Kidd and Seldin panels. <i>Scientific Reports</i> , 2019, 9, 18874. | 1.6 | 15 |
| 26 | The population genetics characteristics of a 90 locus panel of microhaplotypes. <i>Human Genetics</i> , 2021, 140, 1753-1773. | 1.8 | 15 |
| 27 | Mongolians in the Genetic Landscape of Central Asia: Exploring the Genetic Relations among Mongolians and Other World Populations. <i>Human Biology</i> , 2015, 87, 73. | 0.4 | 14 |
| 28 | ALFRED: An Allele Frequency Database for Microevolutionary Studies. <i>Evolutionary Bioinformatics</i> , 2005, 1, 117693430500100. | 0.6 | 12 |
| 29 | ALFRED: A WEB-ACCESSIBLE ALLELE FREQUENCY DATABASE. , 1999, , 639-50. | | 12 |
| 30 | The distinctive geographic patterns of common pigmentation variants at the OCA2 gene. <i>Scientific Reports</i> , 2020, 10, 15433. | 1.6 | 8 |
| 31 | FrogAncestryCalc: A standalone batch likelihood computation tool for ancestry inference panels catalogued in FROG-kb. <i>Forensic Science International: Genetics</i> , 2020, 46, 102237. | 1.6 | 8 |
| 32 | Usefulness of COMT gene polymorphisms in North African populations. <i>Gene</i> , 2019, 696, 186-196. | 1.0 | 7 |
| 33 | Genetic relationships of Southwest Asian and Mediterranean populations. <i>Forensic Science International: Genetics</i> , 2021, 53, 102528. | 1.6 | 7 |
| 34 | North Asian population relationships in a global context. <i>Scientific Reports</i> , 2022, 12, 7214. | 1.6 | 3 |
| 35 | A more powerful method to evaluate p-values in GENEHUNTER. <i>Genetic Epidemiology</i> , 1999, 17, S415-S420. | 0.6 | 2 |
| 36 | Genetic diversity of the North African population revealed by the typing of SNPs in the DRD2/ANKK1 genomic region. <i>Gene</i> , 2021, 777, 145466. | 1.0 | 2 |

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|----|---|-----|-----------|
| 37 | New Insight into the human genetic diversity in North African populations by genotyping of <sc>SNPs</sc> in <sc><i>DRD3</i></sc>, <sc><i>CSMD1</i></sc> and <sc><i>NRG1</i></sc> genes. Molecular Genetics & Genomic Medicine, 2022, 10, e1871. | 0.6 | 2 |
| 38 | Dinucleotide polymorphism at the DXS1178 locus is tightly linked to PGK1 at Xq13. Human Genetics, 1995, 95, 467-8. | 1.8 | 1 |
| 39 | STAT3 polymorphisms in North Africa and its implication in breast cancer. Molecular Genetics & Genomic Medicine, 2021, 9, e1744. | 0.6 | 1 |
| 40 | Genetic analysis workshop III: Sampling considerations and assumptions in gene mapping. Genetic Epidemiology, 1985, 2, 219-220. | 0.6 | 0 |