

Silke Szymczak

List of Publications by Year in descending order

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Version: 2024-02-01

69
papers

7,145
citations

159585

30
h-index

91884

69
g-index

71
all docs

71
docs citations

71
times ranked

15284
citing authors

#	ARTICLE	IF	CITATIONS
1	Linkage analysis identifies novel genetic modifiers of microbiome traits in families with inflammatory bowel disease. <i>Gut Microbes</i> , 2022, 14, 2024415.	9.8	5
2	Blood transcriptome profiling identifies 2 candidate endotypes of atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 150, 385-395.	2.9	17
3	Genetic loci and prioritization of genes for kidney function decline derived from a meta-analysis of 62 longitudinal genome-wide association studies. <i>Kidney International</i> , 2022, 102, 624-639.	5.2	18
4	Early prediction of final infarct volume with material decomposition images of dual-energy CT after mechanical thrombectomy. <i>Neuroradiology</i> , 2021, 63, 695-704.	2.2	4
5	Meta-analysis uncovers genome-wide significant variants for rapid kidney function decline. <i>Kidney International</i> , 2021, 99, 926-939.	5.2	42
6	Genotype imputation in case-only studies of gene-environment interaction: validity and power. <i>Human Genetics</i> , 2021, 140, 1217-1228.	3.8	3
7	Host traits, lifestyle and environment are associated with human skin bacteria. <i>British Journal of Dermatology</i> , 2021, 185, 573-584.	1.5	14
8	The BIOMarkers in Atopic Dermatitis and Psoriasis (BIOMAP) glossary: developing a lingua franca to facilitate data harmonization and cross-cohort analyses. <i>British Journal of Dermatology</i> , 2021, 185, 1066-1069.	1.5	10
9	Integrating biological knowledge and gene expression data using pathway-guided random forests: a benchmarking study. <i>Bioinformatics</i> , 2020, 36, 4301-4308.	4.1	8
10	DNA methylation QTL analysis identifies new regulators of human longevity. <i>Human Molecular Genetics</i> , 2020, 29, 1154-1167.	2.9	15
11	Rare Variants in Specific Lysosomal Genes Are Associated With Parkinson's Disease. <i>Movement Disorders</i> , 2020, 35, 1245-1248.	3.9	37
12	A fungal pathogen induces systemic susceptibility and systemic shifts in wheat metabolome and microbiome composition. <i>Nature Communications</i> , 2020, 11, 1910.	12.8	85
13	Private variants in PRKN are associated with late-onset Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2020, 75, 24-26.	2.2	4
14	Genome-wide association meta-analyses and fine-mapping elucidate pathways influencing albuminuria. <i>Nature Communications</i> , 2019, 10, 4130.	12.8	133
15	Metabolic Functions of Gut Microbes Associate With Efficacy of Tumor Necrosis Factor Antagonists in Patients With Inflammatory Bowel Diseases. <i>Gastroenterology</i> , 2019, 157, 1279-1292.e11.	1.3	180
16	A catalog of genetic loci associated with kidney function from analyses of a million individuals. <i>Nature Genetics</i> , 2019, 51, 957-972.	21.4	549
17	Heterogeneous intracellular TRAIL-receptor distribution predicts poor outcome in breast cancer patients. <i>Journal of Molecular Medicine</i> , 2019, 97, 1155-1167.	3.9	9
18	Atopic Dermatitis Is an IL-13-Dominant Disease with Greater Molecular Heterogeneity Compared to Psoriasis. <i>Journal of Investigative Dermatology</i> , 2019, 139, 1480-1489.	0.7	283

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19	Surrogate minimal depth as an importance measure for variables in random forests. <i>Bioinformatics</i> , 2019, 35, 3663-3671.	4.1	26
20	The metabolic network coherence of human transcriptomes is associated with genetic variation at the cadherin 18 locus. <i>Human Genetics</i> , 2019, 138, 375-388.	3.8	6
21	Evaluation of variable selection methods for random forests and omics data sets. <i>Briefings in Bioinformatics</i> , 2019, 20, 492-503.	6.5	342
22	Heterogeneous intracellular TRAIL-receptor distribution predicts poor outcome in breast cancer patients. <i>Senologie - Zeitschrift für Mammadiagnostik Und -therapie</i> , 2019, 16, .	0.0	0
23	NGS-based methylation profiling differentiates TCF3-HLF and TCF3-PBX1 positive B-cell acute lymphoblastic leukemia. <i>Epigenomics</i> , 2018, 10, 133-147.	2.1	10
24	Histologic improvement of NAFLD in patients with obesity after bariatric surgery based on standardized NAS (NAFLD activity score). <i>Surgery for Obesity and Related Diseases</i> , 2018, 14, 1607-1616.	1.2	56
25	Evaluation of interleukin-6 and its soluble receptor components sIL-6R and sgp130 as markers of inflammation in inflammatory bowel diseases. <i>International Journal of Colorectal Disease</i> , 2018, 33, 927-936.	2.2	34
26	Topography of essential tremor. <i>Parkinsonism and Related Disorders</i> , 2017, 40, 58-63.	2.2	30
27	Increased Tryptophan Metabolism Is Associated With Activity of Inflammatory Bowel Diseases. <i>Gastroenterology</i> , 2017, 153, 1504-1516.e2.	1.3	338
28	Serologic Anti-GP2 Antibodies Are Associated with Genetic Polymorphisms, Fibrostenosis, and Need for Surgical Resection in Crohn's Disease. <i>Inflammatory Bowel Diseases</i> , 2016, 22, 2648-2657.	1.9	25
29	Genome-wide association study of serum coenzyme Q ₁₀ levels identifies susceptibility loci linked to neuronal diseases. <i>Human Molecular Genetics</i> , 2016, 25, ddw134.	2.9	15
30	Genome-wide association analysis identifies variation in vitamin D receptor and other host factors influencing the gut microbiota. <i>Nature Genetics</i> , 2016, 48, 1396-1406.	21.4	533
31	Paternal chronic colitis causes epigenetic inheritance of susceptibility to colitis. <i>Scientific Reports</i> , 2016, 6, 31640.	3.3	15
32	Comparison of parametric and machine methods for variable selection in simulated Genetic Analysis Workshop 19 data. <i>BMC Proceedings</i> , 2016, 10, 147-152.	1.6	1
33	r2VIM: A new variable selection method for random forests in genome-wide association studies. <i>BioData Mining</i> , 2016, 9, 7.	4.0	53
34	Association between SNPs in defined functional pathways and risk of early or late toxicity as well as individual radiosensitivity. <i>Strahlentherapie Und Onkologie</i> , 2015, 191, 59-66.	2.0	12
35	A Hereditary Form of Small Intestinal Carcinoid Associated With a Germline Mutation in Inositol Polyphosphate Multikinase. <i>Gastroenterology</i> , 2015, 149, 67-78.	1.3	96
36	Vy-PER: eliminating false positive detection of virus integration events in next generation sequencing data. <i>Scientific Reports</i> , 2015, 5, 11534.	3.3	42

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37	Trans-ancestry genome-wide association study identifies 12 genetic loci influencing blood pressure and implicates a role for DNA methylation. <i>Nature Genetics</i> , 2015, 47, 1282-1293.	21.4	294
38	Sparse Modeling Reveals miRNA Signatures for Diagnostics of Inflammatory Bowel Disease. <i>PLoS ONE</i> , 2015, 10, e0140155.	2.5	31
39	Increased cathepsin D protein expression is a biomarker for osteosarcomas, pulmonary metastases and other bone malignancies. <i>Oncotarget</i> , 2015, 6, 16517-16526.	1.8	44
40	Variable selection method for the identification of epistatic models. <i>Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing</i> , 2015, , 195-206.	0.7	6
41	Genomewide <i>RNAi</i> screen identifies protein kinase <i>C¹</i> and new members of mitogen-activated protein kinase pathway as regulators of melanoma cell growth and metastasis. <i>Pigment Cell and Melanoma Research</i> , 2014, 27, 418-430.	3.3	12
42	False-positive rates in two-point parametric linkage analysis. <i>BMC Proceedings</i> , 2014, 8, S110.	1.6	1
43	Whole Exome Sequencing of Distant Relatives in Multiplex Families Implicates Rare Variants in Candidate Genes for Oral Clefts. <i>Genetics</i> , 2014, 197, 1039-1044.	2.9	79
44	Risk estimation using probability machines. <i>BioData Mining</i> , 2014, 7, 2.	4.0	14
45	Adaptive linear rank tests for eQTL studies. <i>Statistics in Medicine</i> , 2013, 32, 524-537.	1.6	4
46	Chromosomal Aneuploidy Affects the Global Proteome Equilibrium of Colorectal Cancer Cells. <i>Analytical Cellular Pathology</i> , 2013, 36, 149-161.	1.4	17
47	Chromosomal aneuploidy affects the global proteome equilibrium of colorectal cancer cells. <i>Analytical Cellular Pathology</i> , 2013, 36, 149-61.	1.4	10
48	Association of single nucleotide polymorphisms in the genes ATM, GSTP1, SOD2, TGFB1, XPD and XRCC1 with risk of severe erythema after breast conserving radiotherapy. <i>Radiation Oncology</i> , 2012, 7, 65.	2.7	33
49	Deregulation of a distinct set of microRNAs is associated with transformation of gastritis into MALT lymphoma. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2012, 460, 371-377.	2.8	46
50	Protein profiling of genomic instability in endometrial cancer. <i>Cellular and Molecular Life Sciences</i> , 2012, 69, 325-333.	5.4	8
51	Influence of sex and genetic variability on expression of X-linked genes in human monocytes. <i>Genomics</i> , 2011, 98, 320-326.	2.9	23
52	A Genome-Wide Association Study Identifies <i>LIPA</i> as a Susceptibility Gene for Coronary Artery Disease. <i>Circulation: Cardiovascular Genetics</i> , 2011, 4, 403-412.	5.1	130
53	HDAC2 and TXNL1 distinguish aneuploid from diploid colorectal cancers. <i>Cellular and Molecular Life Sciences</i> , 2011, 68, 3261-3274.	5.4	17
54	Integrating Genome-Wide Genetic Variations and Monocyte Expression Data Reveals Trans-Regulated Gene Modules in Humans. <i>PLoS Genetics</i> , 2011, 7, e1002367.	3.5	126

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55	A trans-acting locus regulates an anti-viral expression network and type 1 diabetes risk. <i>Nature</i> , 2010, 467, 460-464.	27.8	271
56	Genetics and Beyond – The Transcriptome of Human Monocytes and Disease Susceptibility. <i>PLoS ONE</i> , 2010, 5, e10693.	2.5	539
57	Stress sensitivity is increased in transgenic rats with low brain angiotensinogen. <i>Journal of Endocrinology</i> , 2010, 204, 85-92.	2.6	18
58	Association of single nucleotide polymorphisms in ATM, GSTP1, SOD2, TGFB1, XPD and XRCC1 with clinical and cellular radiosensitivity. <i>Radiotherapy and Oncology</i> , 2010, 97, 26-32.	0.6	69
59	The potential role of G2- but not of G0-radiosensitivity for predisposition of prostate cancer. <i>Radiotherapy and Oncology</i> , 2010, 96, 19-24.	0.6	15
60	ACPA: automated cluster plot analysis of genotype data. <i>BMC Proceedings</i> , 2009, 3, S58.	1.6	9
61	Genetic Analysis Workshop 16: Strategies for genome-wide association study analyses. <i>BMC Proceedings</i> , 2009, 3, S1.	1.6	8
62	Evaluation of single-nucleotide polymorphism imputation using random forests. <i>BMC Proceedings</i> , 2009, 3, S65.	1.6	7
63	Detecting SNP-expression associations: A comparison of mutual information and median test with standard statistical approaches. <i>Statistics in Medicine</i> , 2009, 28, 3581-3596.	1.6	17
64	Individual Radiosensitivity Measured With Lymphocytes May Predict the Risk of Acute Reaction After Radiotherapy. <i>International Journal of Radiation Oncology Biology Physics</i> , 2008, 71, 256-264.	0.8	79
65	Identification of Genes Relevant to Symbiosis and Competitiveness in <i>Sinorhizobium meliloti</i> Using Signature-Tagged Mutants. <i>Molecular Plant-Microbe Interactions</i> , 2008, 21, 219-231.	2.6	63
66	Genomewide Association Analysis of Coronary Artery Disease. <i>New England Journal of Medicine</i> , 2007, 357, 443-453.	27.0	1,865
67	Genetic association studies for gene expressions: permutation-based mutual information in a comparison with standard ANOVA and as a novel approach for feature selection. <i>BMC Proceedings</i> , 2007, 1, S9.	1.6	6
68	Picking single-nucleotide polymorphisms in forests. <i>BMC Proceedings</i> , 2007, 1, S59.	1.6	28
69	Construction of a Large Signature-Tagged Mini-Tn5 Transposon Library and Its Application to Mutagenesis of <i>Sinorhizobium meliloti</i> . <i>Applied and Environmental Microbiology</i> , 2006, 72, 4329-4337.	3.1	84