Silke Szymczak

List of Publications by Year in descending order

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159585 91884 7,145 69 30 69 citations h-index g-index papers 71 71 71 15284 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Genomewide Association Analysis of Coronary Artery Disease. New England Journal of Medicine, 2007, 357, 443-453.	27.0	1,865
2	A catalog of genetic loci associated with kidney function from analyses of a million individuals. Nature Genetics, 2019, 51, 957-972.	21.4	549
3	Genetics and Beyond – The Transcriptome of Human Monocytes and Disease Susceptibility. PLoS ONE, 2010, 5, e10693.	2.5	539
4	Genome-wide association analysis identifies variation in vitamin D receptor and other host factors influencing the gut microbiota. Nature Genetics, 2016, 48, 1396-1406.	21.4	533
5	Evaluation of variable selection methods for random forests and omics data sets. Briefings in Bioinformatics, 2019, 20, 492-503.	6.5	342
6	Increased Tryptophan Metabolism Is Associated With Activity of Inflammatory Bowel Diseases. Gastroenterology, 2017, 153, 1504-1516.e2.	1.3	338
7	Trans-ancestry genome-wide association study identifies 12 genetic loci influencing blood pressure and implicates a role for DNA methylation. Nature Genetics, 2015, 47, 1282-1293.	21.4	294
8	Atopic Dermatitis Is an IL-13–Dominant Disease with Greater Molecular Heterogeneity Compared to Psoriasis. Journal of Investigative Dermatology, 2019, 139, 1480-1489.	0.7	283
9	A trans-acting locus regulates an anti-viral expression network and type 1 diabetes risk. Nature, 2010, 467, 460-464.	27.8	271
10	Metabolic Functions of Gut Microbes Associate With Efficacy ofÂTumor Necrosis Factor Antagonists in Patients With Inflammatory Bowel Diseases. Gastroenterology, 2019, 157, 1279-1292.e11.	1.3	180
11	Genome-wide association meta-analyses and fine-mapping elucidate pathways influencing albuminuria. Nature Communications, 2019, 10, 4130.	12.8	133
12	A Genome-Wide Association Study Identifies <i>LIPA</i> as a Susceptibility Gene for Coronary Artery Disease. Circulation: Cardiovascular Genetics, 2011, 4, 403-412.	5.1	130
13	Integrating Genome-Wide Genetic Variations and Monocyte Expression Data Reveals Trans-Regulated Gene Modules in Humans. PLoS Genetics, 2011, 7, e1002367.	3.5	126
14	A Hereditary Form of Small Intestinal Carcinoid Associated With a Germline Mutation in Inositol Polyphosphate Multikinase. Gastroenterology, 2015, 149, 67-78.	1.3	96
15	A fungal pathogen induces systemic susceptibility and systemic shifts in wheat metabolome and microbiome composition. Nature Communications, 2020, 11, 1910.	12.8	85
16	Construction of a Large Signature-Tagged Mini-Tn5 Transposon Library and Its Application to Mutagenesis of Sinorhizobium meliloti. Applied and Environmental Microbiology, 2006, 72, 4329-4337.	3.1	84
17	Individual Radiosensitivity Measured With Lymphocytes May Predict the Risk of Acute Reaction After Radiotherapy. International Journal of Radiation Oncology Biology Physics, 2008, 71, 256-264.	0.8	79
18	Whole Exome Sequencing of Distant Relatives in Multiplex Families Implicates Rare Variants in Candidate Genes for Oral Clefts. Genetics, 2014, 197, 1039-1044.	2.9	79

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19	Association of single nucleotide polymorphisms in ATM, GSTP1, SOD2, TGFB1, XPD and XRCC1 with clinical and cellular radiosensitivity. Radiotherapy and Oncology, 2010, 97, 26-32.	0.6	69
20	Identification of Genes Relevant to Symbiosis and Competitiveness in <i>Sinorhizobium meliloti</i> Using Signature-Tagged Mutants. Molecular Plant-Microbe Interactions, 2008, 21, 219-231.	2.6	63
21	Histologic improvement of NAFLD in patients with obesity after bariatric surgery based on standardized NAS (NAFLD activity score). Surgery for Obesity and Related Diseases, 2018, 14, 1607-1616.	1.2	56
22	r2VIM: A new variable selection method for random forests in genome-wide association studies. BioData Mining, 2016, 9, 7.	4.0	53
23	Deregulation of a distinct set of microRNAs is associated with transformation of gastritis into MALT lymphoma. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2012, 460, 371-377.	2.8	46
24	Increased cathepsin D protein expression is a biomarker for osteosarcomas, pulmonary metastases and other bone malignancies. Oncotarget, 2015, 6, 16517-16526.	1.8	44
25	Vy-PER: eliminating false positive detection of virus integration events in next generation sequencing data. Scientific Reports, 2015, 5, 11534.	3.3	42
26	Meta-analysis uncovers genome-wide significant variants for rapid kidney function decline. Kidney International, 2021, 99, 926-939.	5.2	42
27	Rare Variants in Specific Lysosomal Genes Are Associated With Parkinson's Disease. Movement Disorders, 2020, 35, 1245-1248.	3.9	37
28	Evaluation of interleukin-6 and its soluble receptor components sIL-6R and sgp130 as markers of inflammation in inflammatory bowel diseases. International Journal of Colorectal Disease, 2018, 33, 927-936.	2.2	34
29	Association of single nucleotide polymorphisms in the genes ATM, GSTP1, SOD2, TGFB1, XPD and XRCC1 with risk of severe erythema after breast conserving radiotherapy. Radiation Oncology, 2012, 7, 65.	2.7	33
30	Sparse Modeling Reveals miRNA Signatures for Diagnostics of Inflammatory Bowel Disease. PLoS ONE, 2015, 10, e0140155.	2.5	31
31	Topography of essential tremor. Parkinsonism and Related Disorders, 2017, 40, 58-63.	2.2	30
32	Picking single-nucleotide polymorphisms in forests. BMC Proceedings, 2007, 1, S59.	1.6	28
33	Surrogate minimal depth as an importance measure for variables in random forests. Bioinformatics, 2019, 35, 3663-3671.	4.1	26
34	Serologic Anti-GP2 Antibodies Are Associated with Genetic Polymorphisms, Fibrostenosis, and Need for Surgical Resection in Crohn $\hat{E}\frac{1}{4}$ s Disease. Inflammatory Bowel Diseases, 2016, 22, 2648-2657.	1.9	25
35	Influence of sex and genetic variability on expression of X-linked genes in human monocytes. Genomics, 2011, 98, 320-326.	2.9	23
36	Stress sensitivity is increased in transgenic rats with low brain angiotensinogen. Journal of Endocrinology, 2010, 204, 85-92.	2.6	18

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37	Genetic loci and prioritization of genes for kidney function decline derived from a meta-analysis of 62 longitudinal genome-wide association studies. Kidney International, 2022, 102, 624-639.	5.2	18
38	Detecting SNPâ€expression associations: A comparison of mutual information and median test with standard statistical approaches. Statistics in Medicine, 2009, 28, 3581-3596.	1.6	17
39	HDAC2 and TXNL1 distinguish aneuploid from diploid colorectal cancers. Cellular and Molecular Life Sciences, 2011, 68, 3261-3274.	5.4	17
40	Chromosomal Aneuploidy Affects the Global Proteome Equilibrium of Colorectal Cancer Cells. Analytical Cellular Pathology, 2013, 36, 149-161.	1.4	17
41	Blood transcriptome profiling identifies 2 candidate endotypes of atopic dermatitis. Journal of Allergy and Clinical Immunology, 2022, 150, 385-395.	2.9	17
42	The potential role of G2- but not of G0-radiosensitivity for predisposition of prostate cancer. Radiotherapy and Oncology, 2010, 96, 19-24.	0.6	15
43	Genome-wide association study of serum coenzyme Q ₁₀ levels identifies susceptibility loci linked to neuronal diseases. Human Molecular Genetics, 2016, 25, ddw134.	2.9	15
44	Paternal chronic colitis causes epigenetic inheritance of susceptibility to colitis. Scientific Reports, 2016, 6, 31640.	3.3	15
45	DNA methylation QTL analysis identifies new regulators of human longevity. Human Molecular Genetics, 2020, 29, 1154-1167.	2.9	15
46	Risk estimation using probability machines. BioData Mining, 2014, 7, 2.	4.0	14
47	Host traits, lifestyle and environment are associated with human skin bacteria. British Journal of Dermatology, 2021, 185, 573-584.	1.5	14
48	Genomewide <scp>RNA</scp> i screen identifies protein kinase <scp>C</scp> β and new members of mitogenâ€activated protein kinase pathway as regulators of melanoma cell growth and metastasis. Pigment Cell and Melanoma Research, 2014, 27, 418-430.	3.3	12
49	Association between SNPs in defined functional pathways and risk of early or late toxicity as well as individual radiosensitivity. Strahlentherapie Und Onkologie, 2015, 191, 59-66.	2.0	12
50	NGS-based methylation profiling differentiates TCF3-HLF and TCF3-PBX1 positive B-cell acute lymphoblastic leukemia. Epigenomics, 2018, 10, 133-147.	2.1	10
51	The BIOMarkers in Atopic Dermatitis and Psoriasis (BIOMAP) glossary: developing a lingua franca to facilitate data harmonization and crossâ€cohort analyses. British Journal of Dermatology, 2021, 185, 1066-1069.	1.5	10
52	Chromosomal aneuploidy affects the global proteome equilibrium of colorectal cancer cells. Analytical Cellular Pathology, 2013, 36, 149-61.	1.4	10
53	ACPA: automated cluster plot analysis of genotype data. BMC Proceedings, 2009, 3, S58.	1.6	9
54	Heterogeneous intracellular TRAIL-receptor distribution predicts poor outcome in breast cancer patients. Journal of Molecular Medicine, 2019, 97, 1155-1167.	3.9	9

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55	Genetic Analysis Workshop 16: Strategies for genome-wide association study analyses. BMC Proceedings, 2009, 3, S1.	1.6	8
56	Protein profiling of genomic instability in endometrial cancer. Cellular and Molecular Life Sciences, 2012, 69, 325-333.	5.4	8
57	Integrating biological knowledge and gene expression data using pathway-guided random forests: a benchmarking study. Bioinformatics, 2020, 36, 4301-4308.	4.1	8
58	Evaluation of single-nucleotide polymorphism imputation using random forests. BMC Proceedings, 2009, 3, S65.	1.6	7
59	Genetic association studies for gene expressions: permutation-based mutual information in a comparison with standard ANOVA and as a novel approach for feature selection. BMC Proceedings, 2007, 1, S9.	1.6	6
60	The metabolic network coherence of human transcriptomes is associated with genetic variation at the cadherin 18 locus. Human Genetics, 2019, 138, 375-388.	3.8	6
61	Variable selection method for the identification of epistatic models. Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing, 2015, , 195-206.	0.7	6
62	Linkage analysis identifies novel genetic modifiers of microbiome traits in families with inflammatory bowel disease. Gut Microbes, 2022, 14, 2024415.	9.8	5
63	Adaptive linear rank tests for eQTL studies. Statistics in Medicine, 2013, 32, 524-537.	1.6	4
64	Early prediction of final infarct volume with material decomposition images of dual-energy CT after mechanical thrombectomy. Neuroradiology, 2021, 63, 695-704.	2.2	4
65	Private variants in PRKN are associated with late-onset Parkinson's disease. Parkinsonism and Related Disorders, 2020, 75, 24-26.	2.2	4
66	Genotype imputation in case-only studies of gene-environment interaction: validity and power. Human Genetics, 2021, 140, 1217-1228.	3.8	3
67	False-positive rates in two-point parametric linkage analysis. BMC Proceedings, 2014, 8, S110.	1.6	1
68	Comparison of parametric and machine methods for variable selection in simulated Genetic Analysis Workshop 19 data. BMC Proceedings, 2016, 10, 147-152.	1.6	1
69	Heterogeneous intracellular TRAIL-receptor distribution predicts poor outcome in breast cancer patients. Senologie - Zeitschrift Für Mammadiagnostik Und -therapie, 2019, 16, .	0.0	O