

Max Köller

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

5,057
citations

687363

13
h-index

713466

21
g-index

21
all docs

21
docs citations

21
times ranked

10499
citing authors

#	ARTICLE	IF	CITATIONS
1	MultiQC: summarize analysis results for multiple tools and samples in a single report. <i>Bioinformatics</i> , 2016, 32, 3047-3048.	4.1	4,633
2	T-cell receptorâ€“HLA-DRB1 associations suggest specific antigens in pulmonary sarcoidosis. <i>European Respiratory Journal</i> , 2016, 47, 898-909.	6.7	65
3	Whole-Genome Sequencing of Cytogenetically Balanced Chromosome Translocations Identifies Potentially Pathological Gene Disruptions and Highlights the Importance of Microhomology in the Mechanism of Formation. <i>Human Mutation</i> , 2017, 38, 180-192.	2.5	58
4	Comprehensive structural variation genome map of individuals carrying complex chromosomal rearrangements. <i>PLoS Genetics</i> , 2019, 15, e1007858.	3.5	36
5	Replicative and non-replicative mechanisms in the formation of clustered CNVs are indicated by whole genome characterization. <i>PLoS Genetics</i> , 2018, 14, e1007780.	3.5	28
6	De novo assembly of <i>Dekkera bruxellensis</i> : a multi technology approach using short and long-read sequencing and optical mapping. <i>GigaScience</i> , 2015, 4, 56.	6.4	26
7	Transcriptomics and methylomics of CD4-positive T cells in arsenic-exposed women. <i>Archives of Toxicology</i> , 2017, 91, 2067-2078.	4.2	26
8	Arrayed identification of DNA signatures. <i>Expert Review of Molecular Diagnostics</i> , 2007, 7, 65-76.	3.1	25
9	Stationary and portable sequencing-based approaches for tracing wastewater contamination in urban stormwater systems. <i>Scientific Reports</i> , 2018, 8, 11907.	3.3	24
10	Sarek: A portable workflow for whole-genome sequencing analysis of germline and somatic variants. <i>F1000Research</i> , 2020, 9, 63.	1.6	21
11	Cluster Flow: A user-friendly bioinformatics workflow tool. <i>F1000Research</i> , 2016, 5, 2824.	1.6	18
12	Single base resolution analysis of 5-hydroxymethylcytosine in 188 human genes: implications for hepatic gene expression. <i>Nucleic Acids Research</i> , 2016, 44, 6756-6769.	14.5	15
13	Microarray-based AMASE as a novel approach for mutation detection. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2004, 554, 77-88.	1.0	13
14	High throughput barcoding method for genome-scale phasing. <i>Scientific Reports</i> , 2019, 9, 18116.	3.3	13
15	Droplet Barcode Sequencing for targeted linked-read haplotyping of single DNA molecules. <i>Nucleic Acids Research</i> , 2017, 45, e125-e125.	14.5	11
16	¹⁸ F-Fluorodeoxyglucose-Positron Emission Tomography Imaging Detects Response to Therapeutic Intervention and Plaque Vulnerability in a Murine Model of Advanced Atherosclerotic Diseaseâ€“Brief Report. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2020, 40, 2821-2828.	2.4	10
17	Assessment of Whole Genome Amplification for Sequence Capture and Massively Parallel Sequencing. <i>PLoS ONE</i> , 2014, 9, e84785.	2.5	10
18	Chromosomal genome assembly of the ethanol production strain CBS 11270 indicates a highly dynamic genome structure in the yeast species <i>Brettanomyces bruxellensis</i> . <i>PLoS ONE</i> , 2019, 14, e0215077.	2.5	8

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19	Tag-array based HPV genotyping by competitive hybridization and extension. Journal of Virological Methods, 2005, 129, 102-112.	2.1	6
20	Detection of MC1R Polymorphisms with Protease-Mediated Allele-Specific Extension as an Alternative to Direct Sequencing. Clinical Chemistry, 2005, 51, 2388-2391.	3.2	6
21	Comparison of PrASE and Pyrosequencing for SNP Genotyping. BMC Genomics, 2006, 7, 291.	2.8	5