

Sarah Sandmann

List of Publications by Year in descending order

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Version: 2024-02-01

27
papers

636
citations

933447

10
h-index

642732

23
g-index

30
all docs

30
docs citations

30
times ranked

1370
citing authors

#	ARTICLE	IF	CITATIONS
1	Exploring Current Challenges and Perspectives for Automatic Reconstruction of Clonal Evolution. <i>Cancer Genomics and Proteomics</i> , 2022, 19, 194-204.	2.0	1
2	Design of a targeted next-generation DNA sequencing panel for pediatric T-cell lymphoblastic lymphoma to unravel biology and optimize treatment. <i>Genes Chromosomes and Cancer</i> , 2022, 61, 459-470.	2.8	2
3	Machine learning in the detection and management of atrial fibrillation. <i>Clinical Research in Cardiology</i> , 2022, 111, 1010-1017.	3.3	23
4	MEDB-67. Subgroup specific analysis of cellular metabolism in medulloblastoma. <i>Neuro-Oncology</i> , 2022, 24, i122-i122.	1.2	1
5	Clinical relevance of molecular characteristics in Burkitt lymphoma differs according to age. <i>Nature Communications</i> , 2022, 13, .	12.8	28
6	Reconstructing clonal evolution in relapsed and non-relapsed Burkitt lymphoma. <i>Leukemia</i> , 2021, 35, 639-643.	7.2	16
7	Integrative genomic analysis of pediatric T-cell lymphoblastic lymphoma reveals candidates of clinical significance. <i>Blood</i> , 2021, 137, 2347-2359.	1.4	31
8	Mutation patterns in recurrent and/or metastatic oropharyngeal squamous cell carcinomas in relation to human papillomavirus status. <i>Cancer Medicine</i> , 2021, 10, 1347-1356.	2.8	9
9	Comparison of Open-access Databases for Clinical Variant Interpretation in Cancer: A Case Study of MDS/AML. <i>Cancer Genomics and Proteomics</i> , 2021, 18, 157-166.	2.0	6
10	Multicenter Next-Generation Sequencing Studies between Theory and Practice. <i>Journal of Molecular Diagnostics</i> , 2021, 23, 347-357.	2.8	1
11	Persistent symptoms and lab abnormalities in patients who recovered from COVID-19. <i>Scientific Reports</i> , 2021, 11, 12775.	3.3	36
12	SimFFPE and FilterFFPE: improving structural variant calling in FFPE samples. <i>GigaScience</i> , 2021, 10, .	6.4	4
13	Calcitonin receptor-like (CALCRL) is a marker of stemness and an independent predictor of outcome in pediatric AML. <i>Blood Advances</i> , 2021, 5, 4413-4421.	5.2	9
14	Clonal Evolution at First Sight: A Combined Visualization of Diverse Diagnostic Methods Improves Understanding of Leukemia Progression. <i>Blood</i> , 2021, 138, 1293-1293.	1.4	0
15	Divergent Effects of EZH1 and EZH2 Protein Expression on the Prognosis of Patients with T-Cell Lymphomas. <i>Biomedicines</i> , 2021, 9, 1842.	3.2	6
16	CopyDetective: Detection threshold-aware copy number variant calling in whole-exome sequencing data. <i>GigaScience</i> , 2020, 9, .	6.4	5
17	Mutation Profiles Identify Distinct Clusters of Lower Risk Myelodysplastic Syndromes with Unique Clinical and Biological Features and Clinical Endpoints. <i>Blood</i> , 2020, 136, 29-29.	1.4	2
18	Genetic alterations in human papillomavirus-associated oropharyngeal squamous cell carcinoma of patients with treatment failure. <i>Oral Oncology</i> , 2019, 93, 59-65.	1.5	10

#	ARTICLE	IF	CITATIONS
19	Effects of computerized decision support system implementations on patient outcomes in inpatient care: a systematic review. <i>Journal of the American Medical Informatics Association: JAMIA</i> , 2018, 25, 593-602.	4.4	98
20	appreci8: a pipeline for precise variant calling integrating 8 tools. <i>Bioinformatics</i> , 2018, 34, 4205-4212.	4.1	26
21	Online Information Infrastructure Increases Inter-Rater Reliability of Medical Coders: A Quasi-Experimental Study (Preprint). <i>Journal of Medical Internet Research</i> , 2018, 20, e274.	4.3	5
22	Evaluating Variant Calling Tools for Non-Matched Next-Generation Sequencing Data. <i>Scientific Reports</i> , 2017, 7, 43169.	3.3	185
23	Clonal evolution in myelodysplastic syndromes. <i>Nature Communications</i> , 2017, 8, 15099.	12.8	118
24	BBCAnalyzer: a visual approach to facilitate variant calling. <i>BMC Bioinformatics</i> , 2017, 18, 133.	2.6	0
25	GLM-based optimization of NGS data analysis: A case study of Roche 454, Ion Torrent PGM and Illumina NextSeq sequencing data. <i>PLoS ONE</i> , 2017, 12, e0171983.	2.5	7
26	Medical Effect of Venous Thromboembolism Prophylaxis Systems and Common Input Categories: Preliminary Findings from a Systematic Review. <i>Studies in Health Technology and Informatics</i> , 2017, 245, 1175-1179.	0.3	1
27	Clonal Evolution at First Sight: A Combined Visualization of Diverse Diagnostic Methods Improves Understanding of Leukemic Progression. <i>Frontiers in Oncology</i> , 0, 12, .	2.8	2