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List of Publications by Year in descending order

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36
papers

2,490
citations

471509

17
h-index

395702

33
g-index

39
all docs

39
docs citations

39
times ranked

4676
citing authors

#	ARTICLE	IF	CITATIONS
1	Towards accurate and reliable resolution of structural variants for clinical diagnosis. <i>Genome Biology</i> , 2022, 23, 68.	8.8	34
2	Ultra-deep sequencing data from a liquid biopsy proficiency study demonstrating analytic validity. <i>Scientific Data</i> , 2022, 9, 170.	5.3	4
3	Ultra-deep multi-oncopanel sequencing of benchmarking samples with a wide range of variant allele frequencies. <i>Scientific Data</i> , 2022, 9, .	5.3	0
4	Deep oncopanel sequencing reveals within block position-dependent quality degradation in FFPE processed samples. <i>Genome Biology</i> , 2022, 23, .	8.8	8
5	Evaluating the analytical validity of circulating tumor DNA sequencing assays for precision oncology. <i>Nature Biotechnology</i> , 2021, 39, 1115-1128.	17.5	126
6	Cross-oncopanel study reveals high sensitivity and accuracy with overall analytical performance depending on genomic regions. <i>Genome Biology</i> , 2021, 22, 109.	8.8	20
7	Optimized imaging methods for species-level identification of food-contaminating beetles. <i>Scientific Reports</i> , 2021, 11, 7957.	3.3	1
8	A verified genomic reference sample for assessing performance of cancer panels detecting small variants of low allele frequency. <i>Genome Biology</i> , 2021, 22, 111.	8.8	29
9	The Sequencing Quality Control 2 study: establishing community standards for sequencing in precision medicine. <i>Genome Biology</i> , 2021, 22, 306.	8.8	7
10	Advancing NGS quality control to enable measurement of actionable mutations in circulating tumor DNA. <i>Cell Reports Methods</i> , 2021, 1, 100106.	2.9	9
11	FDA-led consortium studies advance quality control of targeted next generation sequencing assays for precision oncology. <i>Precision Cancer Medicine</i> , 2021, 4, 32-32.	1.8	4
12	FDALabel for drug repurposing studies and beyond. <i>Nature Biotechnology</i> , 2020, 38, 1378-1379.	17.5	8
13	Study of pharmacogenomic information in FDA-approved drug labeling to facilitate application of precision medicine. <i>Drug Discovery Today</i> , 2020, 25, 813-820.	6.4	29
14	DLI-IT: a deep learning approach to drug label identification through image and text embedding. <i>BMC Medical Informatics and Decision Making</i> , 2020, 20, 68.	3.0	10
15	HetEnc: a deep learning predictive model for multi-type biological dataset. <i>BMC Genomics</i> , 2019, 20, 638.	2.8	4
16	A deep learning model to recognize food contaminating beetle species based on elytra fragments. <i>Computers and Electronics in Agriculture</i> , 2019, 166, 105002.	7.7	28
17	Estimating Bacterial Concentrations in Fibrous Substrates Through a Combination of Scanning Electron Microscopy and ImageJ. <i>Analytical Chemistry</i> , 2019, 91, 4405-4412.	6.5	7
18	Technical advance in targeted NGS analysis enables identification of lung cancer risk-associated low frequency TP53, PIK3CA, and BRAF mutations in airway epithelial cells. <i>BMC Cancer</i> , 2019, 19, 1081.	2.6	12

#	ARTICLE	IF	CITATIONS
19	Advancing Genomics for Drug Development and Safety Evaluation. <i>International Journal of Genomics</i> , 2018, 2018, 1-2.	1.6	0
20	Comparing SVM and ANN based Machine Learning Methods for Species Identification of Food Contaminating Beetles. <i>Scientific Reports</i> , 2018, 8, 6532.	3.3	72
21	Comprehensive Assessments of RNA-seq by the SEQC Consortium: FDA-Led Efforts Advance Precision Medicine. <i>Pharmaceutics</i> , 2016, 8, 8.	4.5	53
22	FDA drug labeling: rich resources to facilitate precision medicine, drug safety, and regulatory science. <i>Drug Discovery Today</i> , 2016, 21, 1566-1570.	6.4	38
23	Species Identification of Food Contaminating Beetles by Recognizing Patterns in Microscopic Images of Elytra Fragments. <i>PLoS ONE</i> , 2016, 11, e0157940.	2.5	11
24	NETBAGs: a network-based clustering approach with gene signatures for cancer subtyping analysis. <i>Biomarkers in Medicine</i> , 2015, 9, 1053-1065.	1.4	9
25	Comparison of RNA-seq and microarray-based models for clinical endpoint prediction. <i>Genome Biology</i> , 2015, 16, 133.	8.8	325
26	An investigation of biomarkers derived from legacy microarray data for their utility in the RNA-seq era. <i>Genome Biology</i> , 2014, 15, 523.	8.8	147
27	Transcriptomic profiling of rat liver samples in a comprehensive study design by RNA-Seq. <i>Scientific Data</i> , 2014, 1, 140021.	5.3	30
28	ATP-Binding Cassette Genes Genotype and Expression: A Potential Association with Pancreatic Cancer Development and Chemoresistance?. <i>Gastroenterology Research and Practice</i> , 2014, 2014, 1-9.	1.5	15
29	Assessing technical performance in differential gene expression experiments with external spike-in RNA control ratio mixtures. <i>Nature Communications</i> , 2014, 5, 5125.	12.8	122
30	The concordance between RNA-seq and microarray data depends on chemical treatment and transcript abundance. <i>Nature Biotechnology</i> , 2014, 32, 926-932.	17.5	420
31	Cross-platform ultradeep transcriptomic profiling of human reference RNA samples by RNA-Seq. <i>Scientific Data</i> , 2014, 1, 140020.	5.3	21
32	SNPTrack TM : an integrated bioinformatics system for genetic association studies. <i>Human Genomics</i> , 2012, 6, 5.	2.9	2
33	ArrayTrack: a free FDA bioinformatics tool to support emerging biomedical research – an update. <i>Human Genomics</i> , 2010, 4, 428-34.	2.9	19
34	An FDA bioinformatics tool for microbial genomics research on molecular characterization of bacterial foodborne pathogens using microarrays. <i>BMC Bioinformatics</i> , 2010, 11, S4.	2.6	53
35	Two new ArrayTrack libraries for personalized biomedical research. <i>BMC Bioinformatics</i> , 2010, 11, S6.	2.6	12
36	The MicroArray Quality Control (MAQC)-II study of common practices for the development and validation of microarray-based predictive models. <i>Nature Biotechnology</i> , 2010, 28, 827-838.	17.5	795