

Jinyoung Hong

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

23
citations

2258059

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2053705

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docs citations

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#	ARTICLE	IF	CITATIONS
1	RNA Sequencing for Elucidating an Intronic Variant of Uncertain Significance (<i>c.5074+3A>C</i>) of <i>BRCA1</i> by RNA Sequencing and TOPO Cloning. <i>Genes</i> , 2021, 12, 810.	2.5	4
2	RNA Sequencing Provides Evidence for Pathogenicity of a Novel <i>CHEK2</i> Splice Variant (C.1009-7T>G). <i>Annals of Laboratory Medicine</i> , 2022, 42, 380-383.	2.5	2
3	Commutability Assessment of Frozen Human Serum Pools for External Quality Assessment of Tumor Markers. <i>Journal of Laboratory Medicine and Quality Assurance</i> , 2022, 44, 111-120.		1
4	Practical Considerations for Clinical Laboratories in Top-down Approach for Assessing the Measurement Uncertainty of Clinical Chemistry Analytes. <i>Annals of Laboratory Medicine</i> , 2022, 42, 630-637.	2.5	0
5	Development of a Method for Manufacturing External Quality Assessment Material for Genetic Testing of Solid Tumors Using Mutant and Wild-Type Cell Lines. <i>Journal of Laboratory Medicine and Quality Assurance</i> , 2021, 43, 31-36.	0.4	0
6	Prevalence and endoscopic treatment outcomes of upper gastrointestinal neoplasms in familial adenomatous polyposis. <i>Surgical Endoscopy and Other Interventional Techniques</i> , 2021, , 1.	2.4	3
7	Identification of a Splice Variant (<i>c.5074+3A>C</i>) of <i>BRCA1</i> by RNA Sequencing and TOPO Cloning. <i>Genes</i> , 2021, 12, 810.	2.4	0
8	Performance evaluation of an amplicon-based next-generation sequencing panel for <i>BRCA1</i> and <i>BRCA2</i> variant detection. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23524.	2.1	4
9	Application and optimization of reference change values for Delta Checks in clinical laboratory. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23550.	2.1	7
10	A Case of Anti-LebH Antibody Identified in a Patient with Ulcerative Colitis. <i>Laboratory Medicine Online</i> , 2019, 9, 254.	0.2	0
11	A Case of Lynch Syndrome with the Deletion of Multiple Exons of the <i>MLH1</i> Gene, Detected by Next-Generation Sequencing. <i>Journal of Laboratory Medicine and Quality Assurance</i> , 2019, 41, 220-224.	0.4	2