Soyoung Shin

List of Publications by Year
in descending order

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Version: 2024-02-01

26	251	1163117	996975
papers	citations	8 h-index	15 g-index

27	27	27	545
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The impact of the Xpert MTB/RIF screening among hospitalized patients with pneumonia on timely isolation of patients with pulmonary tuberculosis. Scientific Reports, 2021, 11, 1694.	3.3	3
2	Measles seroprevalence among healthcare workers in South Korea during the post-elimination period. Human Vaccines and Immunotherapeutics, 2021, 17, 2517-2521.	3.3	8
3	Development and Stability Evaluation of In- House Prepared External Quality Controls for Autoimmune Disease Tests. Journal of Laboratory Medicine and Quality Assurance, 2021, 43, 72-79.	0.4	0
4	Analyzing Genetic Differences Between Sporadic Primary and Secondary/Tertiary Hyperparathyroidism by Targeted Next-Generation Panel Sequencing. Endocrine Pathology, 2021, 32, 501-512.	9.0	3
5	790. Evaluation of an Enhanced CPE Screening Program in an Acute Care Hospital in South Korea. Open Forum Infectious Diseases, 2021, 8, S491-S492.	0.9	0
6	Seizure duration may increase thyroid-stimulating hormone levels in children experiencing a seizure. Journal of International Medical Research, 2020, 48, 030006051988840.	1.0	5
7	Genetically confirmed limb-girdle muscular dystrophy type 2B with DYSF mutation using gene panel sequencing. Medicine (United States), 2020, 99, e20810.	1.0	2
8	A novel EPB41 p.Trp704* mutation in a Korean patient with hereditary elliptocytosis: a case report. Hematology, 2020, 25, 321-326.	1.5	2
9	Coexistence of digenic mutations in the collagen VI genes (COL6A1 and COL6A3) leads to Bethlem myopathy. Clinica Chimica Acta, 2020, 508, 28-32.	1.1	4
10	A novel SYNE2 mutation identified by whole exome sequencing in a Korean family with Emery-Dreifuss muscular dystrophy. Clinica Chimica Acta, 2020, 506, 50-54.	1.1	2
11	Genetic profiling of somatic alterations by Oncomine Focus Assay in Korean patients with advanced gastric cancer. Oncology Letters, 2020, 20, 1-1.	1.8	7
12	Genetic Characterization of Molecular Targets in Korean Patients with Gastrointestinal Stromal Tumors. Journal of Gastric Cancer, 2020, 20, 29.	2.5	6
13	Complement 4 levels of a 4-year-old girl with angioedema. Clinical and Experimental Pediatrics, 2020, 63, 30-31.	2.2	1
14	Comparison of Six Commercial Diagnostic Tests for the Detection of Dengue Virus Non-Structural-1 Antigen and IgM/IgG Antibodies. Annals of Laboratory Medicine, 2019, 39, 566-571.	2.5	36
15	Targeted next-generation sequencing identifies a novel nonsense mutation in SPTB for hereditary spherocytosis. Medicine (United States), 2018, 97, e9677.	1.0	7
16	Molecular drug resistance profiles of Mycobacterium tuberculosis from sputum specimens using ion semiconductor sequencing. Journal of Microbiological Methods, 2018, 145, 1-6.	1.6	9
17	Expression profile of microRNAs following bone marrowâ€derived mesenchymal stem cell treatment in lipopolysaccharideâ€induced acute lung injury. Experimental and Therapeutic Medicine, 2018, 15, 5495-5502.	1.8	20
18	Incidental Identification of Plasmodium vivax During Routine Complete Blood Count Analysis Using the UniCel DxH 800. Annals of Laboratory Medicine, 2018, 38, 165-168.	2.5	3

#	Article	IF	Citations
19	Determining Genotypic Drug Resistance by Ion Semiconductor Sequencing With the Ion AmpliSeqâ„¢ TB Panel in Multidrug-Resistant Mycobacterium tuberculosis Isolates. Annals of Laboratory Medicine, 2018, 38, 316-323.	2.5	15
20	Identification of a novel de novo nonsense mutation of the NSD1 gene in monozygotic twins discordant for Sotos syndrome. Clinica Chimica Acta, 2017, 470, 31-35.	1.1	7
21	The first patient with sporadic X-linked intellectual disability with de novo ZDHHC9 mutation identified by targeted next-generation sequencing. European Journal of Medical Genetics, 2017, 60, 499-503.	1.3	15
22	Comparison between Mononucleotide and Dinucleotide Marker Panels in Gastric Cancer with Loss of hMLH1 or hMSH2 Expression. International Journal of Biological Markers, 2017, 32, 352-356.	1.8	5
23	A Novel Inherited Mutation of in a Korean Family with Benign Familial Infantile Epilepsy Using Diagnostic Exome Sequencing. Annals of Clinical and Laboratory Science, 2017, 47, 747-753.	0.2	11
24	Expression profiling of microRNAs in lipopolysaccharide-induced acute lung injury after hypothermia treatment. Molecular and Cellular Toxicology, 2016, 12, 243-253.	1.7	9
25	Analysis of Immunoglobulin and T Cell Receptor Gene Rearrangement in the Bone Marrow of Lymphoid Neoplasia Using BIOMED-2 Multiplex Polymerase Chain Reaction. International Journal of Medical Sciences, 2013, 10, 1510-1517.	2.5	12
26	The Therapeutic Effect of Human Adult Stem Cells Derived from Adipose Tissue in Endotoxemic Rat Model. International Journal of Medical Sciences, 2013, 10, 8-18.	2.5	57