Myung-Shin Kim

List of Publications by Year
in descending order

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272	3,616	218381	253896
papers	citations	26 h-index	43 g-index

291	291	291	6473
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hereditary renal hypouricemia with SLC22A12 mutation: A case report. Pediatrics and Neonatology, 2022, 63, 202-203.	0.3	0
2	Analytical and Clinical Assessment of Prostate Specific Antigen Using an HISCL-5000 Chemiluminescence Immunoassay. Laboratory Medicine Online, 2022, 12, 26-32.	0.0	0
3	Adrenocortical carcinoma and a sporadic MEN1 mutation in a 3-year-old girl: a case report. Annals of Pediatric Endocrinology and Metabolism, 2022, 27, 315-319.	0.8	4
4	A Novel HNPP Phenotype in Charcot-Marie-Tooth Type 2E With c.1319C>T Missense Mutation in the		

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19	The factors influencing clinical outcomes after leukapheresis in acute leukaemia. Scientific Reports, 2021, 11, 6426.	1.6	8
20	Marked thrombocytosis resulting in pseudohyperkalemia in a neonate with transient abnormal myelopoiesis. Pediatric Blood and Cancer, 2021, 68, e28986.	0.8	2
21	Generation of the human induced pluripotent stem cell lines (CMCi009-A) from a patient with Birt-Hogg-DubÃ© syndrome (BHD) with heterozygous frameshift deletion mutation c.1285delC of the FLCN gene. Stem Cell Research, 2021, 51, 102215.	0.3	1
22	Impact of Integrated Genetic Information on Diagnosis and Prognostication for Myeloproliferative Neoplasms in the Next-Generation Sequencing Era. Journal of Clinical Medicine, 2021, 10, 1033.	1.0	6
23	Targeted Next-Generation Sequencing of Plasma Cell-Free DNA in Korean Patients with Hepatocellular Carcinoma. Annals of Laboratory Medicine, 2021, 41, 198-206.	1.2	5
24	Optimizing the Diagnostic Strategy to Identify Genetic Abnormalities in Miscarriage. Molecular Diagnosis and Therapy, 2021, 25, 351-359.	1.6	5
25	Non-inferior long-term outcomes of adults with Philadelphia chromosome-like acute lymphoblastic leukemia. Bone Marrow Transplantation, 2021, 56, 1953-1963.	1.3	9
26	Diagnosis for Pheochromocytoma and Paraganglioma: A Joint Position Statement of the Korean Pheochromocytoma and Paraganglioma Task Force. Endocrinology and Metabolism, 2021, 36, 322-338.	1.3	11
27	Transient Neonatal Diabetes Mellitus in SHORT Syndrome: A Case Report. Frontiers in Pediatrics, 2021, 9, 650920.	0.9	0
28	Prognostic values of D816V KIT mutation and peri-transplant CBFB-MYH11 MRD monitoring on acute myeloid leukemia with CBFB-MYH11. Bone Marrow Transplantation, 2021, 56, 2682-2689.	1.3	3
29	Multilevel Airway Stenosis Being Bypassed by a Customized Tracheostomy Tube in an Infant with Myhre Syndrome. Pediatric, Allergy, Immunology, and Pulmonology, 2021, 34, 83-87.	0.3	1
30	Prognostic value of measurable residual disease monitoring by next-generation sequencing before and after allogeneic hematopoietic cell transplantation in acute myeloid leukemia. Blood Cancer Journal, 2021, 11, 109.	2.8	27
31	Analytical and Clinical Performance of Droplet Digital PCR in the Detection and Quantification of SARS-CoV-2. Molecular Diagnosis and Therapy, 2021, 25, 617-628.	1.6	12
32	Postoperative Circulating Tumor DNA Can Predict High Risk Patients with Colorectal Cancer Based on Next-Generation Sequencing. Cancers, 2021, 13, 4190.	1.7	3
33	Integrative Analysis of Gene Expression Data by RNA Sequencing for Differential Diagnosis of Acute Leukemia: Potential Application of Machine Learning. Frontiers in Oncology, 2021, 11, 717616.	1.3	6
34	Poster: AML-282: Prognostic Values of D816V KIT Mutation and Peri-Transplant CBFB-MYH11 MRD Monitoring on Acute Myeloid Leukemia with CBFB-MYH11. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, S215.	0.2	0
35	AML-282: Prognostic Values of D816V KIT Mutation and Peri-Transplant CBFB-MYH11 MRD Monitoring on Acute Myeloid Leukemia with CBFB-MYH11. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, S299-S300.	0.2	0
36	Comparable Outcomes After Alternative and Matched Sibling Donor Hematopoietic Stem Cell Transplantation and the Role of Molecular Measurable Residual Disease for Acute Myeloid Leukemia in Elderly Patients. Transplantation and Cellular Therapy, 2021, 27, 774.e1-774.e12.	0.6	12

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37	Characteristics of RAS pathway mutations in juvenile myelomonocytic leukaemia: a singleâ€institution study from Korea. British Journal of Haematology, 2021, 195, 748-756.	1.2	3
38	Status of Next-Generation Sequencing-Based Genetic Diagnosis in Hematologic Malignancies in Korea (2017-2018). Laboratory Medicine Online, 2021, 11, 25-31.	0.0	1
39	Direct Detection of Low Abundance Genes of Single Point Mutation. Nano Letters, 2021, 21, 9061-9068.	4.5	11
40	Novel WTX nonsense mutation in a family diagnosed with osteopathia striata with cranial sclerosis. Medicine (United States), 2021, 100, e27346.	0.4	1
41	A Boy with X-Linked Inhibitor of Apoptosis Protein (XIAP) Deficiency as the Initial Presentation of Pure Red Cell Aplasia. Clinical Pediatric Hematology-Oncology, 2021, 28, 84-88.	0.0	0
42	Andersenâ€“Tawil Syndrome With Novel Mutation in KCNJ2: Case Report. Frontiers in Pediatrics, 2021, 9, 790075.	0.9	1
43	Evaluation of posaconazole plasma concentrations achieved with the delayedâ€release tablets in Korean highâ€risk patients with haematologic malignancy. Mycoses, 2020, 63, 131-138.	1.8	5
44	Clinical Validity of Next-Generation Sequencing Multi-Gene Panel Testing for Detecting Pathogenic Variants in Patients With Hereditary Breast-Ovarian Cancer Syndrome. Annals of Laboratory Medicine, 2020, 40, 148-154.	1.2	15
45	Prognostic Role of Postinduction Minimal Residual Disease and Myeloid Sarcoma Type Extramedullary Involvement in Pediatric RUNX1-RUNX1T1 (+) Acute Myeloid Leukemia. Journal of Pediatric Hematology/Oncology, 2020, 42, e132-e139.	0.3	7
46	De Novo Pure Trisomy 20p: Report of a Novel Case of a Marker Chromosome and Literature Review. Annals of Laboratory Medicine, 2020, 40, 277-280.	1.2	2
47	Generation of a human induced pluripotent stem cell line (CMCi001-A) from a patient with karyomegalic interstitial nephritis with homozygous frameshift deletion mutation c.1985_1994del10 of the FANCD2/FANCI-Associated Nuclease 1 gene. Stem Cell Research, 2020, 46, 101876.	0.3	1
48	Analytical and Potential Clinical Performance of Oncomine Myeloid Research Assay for Myeloid Neoplasms. Molecular Diagnosis and Therapy, 2020, 24, 579-592.	1.6	6
49	Gene expression signatures associated with sensitivity to azacitidine in myelodysplastic syndromes. Scientific Reports, 2020, 10, 19555.	1.6	7
50	Peroxidasin is essential for endothelial cell survival and growth signaling by sulfilimine crosslinkâ€dependent matrix assembly. FASEB Journal, 2020, 34, 10228-10241.	0.2	17
51	Common and different alterations of bone marrow mesenchymal stromal cells in myelodysplastic syndrome and multiple myeloma. Cell Proliferation, 2020, 53, e12819.	2.4	10
52	High incidence of RAS pathway mutations among sentinel genetic lesions of Korean pediatric BCRâ€ABL1 â€like acute lymphoblastic leukemia. Cancer Medicine, 2020, 9, 4632-4639.	1.3	7
53	Genotypic and phenotypic characteristics of Korean children with childhood-onset Leberâ€™s hereditary optic neuropathy. Graefe's Archive for Clinical and Experimental Ophthalmology, 2020, 258, 2283-2290.	1.0	7
54	Evaluation of Two EGFR Mutation Tests on Tumor and Plasma from Patients with Non-Small Cell Lung Cancer. Cancers, 2020, 12, 785.	1.7	5

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55	Seroprevalence of Neutralizing Antibodies against Japanese Encephalitis Virus among Adolescents and Adults in Korea: A Prospective Multicenter Study. Vaccines, 2020, 8, 328.	2.1	7
56	Sex Chromosomes Are Severely Disrupted in Gastric Cancer Cell Lines. International Journal of Molecular Sciences, 2020, 21, 4598.	1.8	3
57	Detection of BRCA1/2 large genomic rearrangement including BRCA1 promoter-region deletions using next-generation sequencing. Clinica Chimica Acta, 2020, 505, 49-54.	0.5	13
58	Characteristics ofDNMT3Amutations in acute myeloid leukemia. Blood Research, 2020, 55, 17-26.	0.5	44
59	Revision of Laboratory Testing Guidelines for Initial Diagnosis of Hematologic Neoplasms. Laboratory Medicine Online, 2020, 10, 10.	0.0	1
60	Influence of plasma methotrexate level and MTHFR genotype in Korean paediatric patients with acute lymphoblastic leukaemia. Journal of Chemotherapy, 2020, 32, 251-259.	0.7	7
61	A newborn with developmental delay diagnosed with 4q35 deletion and 10p duplication. Journal of Genetic Medicine, 2020, 17, 102-107.	0.1	0
62	Prenatal Genetic Diagnosis of Congenital Nephrotic Syndrome. Perinatology, 2020, 31, 99.	0.0	0
63	KRAS, NRAS, and BRAF mutations in plasma cell myeloma at a single Korean institute. Blood Research, 2020, 55, 159-168.	0.5	0
64	Prognostic Impacts of Next-Generation Sequencing-Based Measurable Residual Disease Monitoring before and after Allogeneic Hematopoietic Cell Transplantation in AML. Blood, 2020, 136, 1-2.	0.6	0
65	Analytical and Clinical Evaluation of Chemiluminescent Carcinoembryonic Antigen (CEA) by HISCL-5000 Immunoanalyzer. Annals of Clinical and Laboratory Science, 2020, 50, 417-422.	0.2	3
66	A Mutation in ZNF143 as a Novel Candidate Gene for Endothelial Corneal Dysplasia. Journal of Clinical Medicine, 2019, 8, 1174.	1.0	3
67	Procalcitonin as a prognostic marker for sepsis based on SEPSISâ€3. Journal of Clinical Laboratory Analysis, 2019, 33, e22996.	0.9	53
68	Exon sequencing of the alpha-2-globin gene for the differential diagnosis of central cyanosis in newborns: a case report. BMC Pediatrics, 2019, 19, 221.	0.7	1
69	A case of early diagnosis of pulmonary capillary hemangiomatosis in a worker with exposure to silica. BMC Pulmonary Medicine, 2019, 19, 133.	0.8	2
70	BRCA1/2 somatic mutation detection in formalin-fixed paraffin embedded tissue by next-generation sequencing in Korean ovarian cancer patients. Pathology Research and Practice, 2019, 215, 152595.	1.0	4
71	Diagnosis and Prognosis of Sepsis Based on Use of Cytokines, Chemokines, and Growth Factors. Disease Markers, 2019, 2019, 1-11.	0.6	25
72	Analytical evaluation and clinical application of insulin and C-peptide by a whole blood, lateral flow, point of care (POC) assay system. Scandinavian Journal of Clinical and Laboratory Investigation, 2019, 79, 347-353.	0.6	4

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73	Clonal Cell Proliferation in Paroxysmal Nocturnal Hemoglobinuria: Evaluation of PIGA Mutations and T-cell Receptor Clonality. Annals of Laboratory Medicine, 2019, 39, 438-446.	1.2	6
74	Enumeration of CD34-positive Stem Cells Using the ADAMII Image-based Fluorescence Cell Counter. Annals of Laboratory Medicine, 2019, 39, 388-395.	1.2	6
75	Recent progress in laboratory diagnosis of thalassemia and hemoglobinopathy: a study by the Korean Red Blood Cell Disorder Working Party of the Korean Society of Hematology. Blood Research, 2019, 54, 17-22.	0.5	25
76	Hereditary spherocytosis caused by copy number variation in SPTB gene identified through targeted next-generation sequencing. International Journal of Hematology, 2019, 110, 250-254.	0.7	4
77	Complex interaction networks of cytokines after transarterial chemotherapy in patients with hepatocellular carcinoma. PLoS ONE, 2019, 14, e0224318.	1.1	20
78	Simultaneous Monitoring of Mutation and Chimerism Using Next-Generation Sequencing in Myelodysplastic Syndrome. Journal of Clinical Medicine, 2019, 8, 2077.	1.0	23
79	Germline CEBPA mutations in Korean patients with acute myeloid leukemia. Leukemia Research, 2019, 76, 84-86.	0.4	14
80	Considerations for monitoring minimal residual disease using immunoglobulin clonality in patients with precursor B-cell lymphoblastic leukemia. Clinica Chimica Acta, 2019, 488, 81-89.	0.5	7
81	Diagnostic approach with genetic tests for global developmental delay and/or intellectual disability: Single tertiary center experience. Annals of Human Genetics, 2019, 83, 115-123.	0.3	14
82	Development of immunocompatible pluripotent stem cells via CRISPR-based human leukocyte antigen engineering. Experimental and Molecular Medicine, 2019, 51, 1-11.	3.2	28
83	CDKN2B downregulation and other genetic characteristics in T-acute lymphoblastic leukemia. Experimental and Molecular Medicine, 2019, 51, 1-15.	3.2	29
84	Chromosomal Microarray Analysis as a First-Tier Clinical Diagnostic Test in Patients With Developmental Delay/Intellectual Disability, Autism Spectrum Disorders, and Multiple Congenital Anomalies: A Prospective Multicenter Study in Korea. Annals of Laboratory Medicine, 2019, 39, 299-310.	1.2	44
85	Reclassification of Acute Myeloid Leukemia According to the 2016 WHO Classification. Annals of Laboratory Medicine, 2019, 39, 311-316.	1.2	18
86	Effects of Copy Number Variations on Developmental Aspects of Children With Delayed Development. Annals of Rehabilitation Medicine, 2019, 43, 215-223.	0.6	6
87	Bone marrow histology in CALR mutated thrombocythemia and myelofibrosis: results from two cross sectional studies in 70 newly diagnosed JAK2/MPL wild type thrombocythemia patients. , 2019, 2, 064-078.		0
88	Analysis of a 6-year pilot external quality assurance survey of free light chain using Sigma metrics. Journal of Laboratory Medicine, 2019, 43, 235-242.	1.1	1
89	A Boy with Chronic Active EBV Infection Presented as Mosquito Bite Hypersensitivity Progressed to Fatal Hemophagocytic Lymphohistiocytosis due to NK Cell Neoplasm. Clinical Pediatric Hematology-Oncology, 2019, 26, 95-98.	0.0	1
90	Targeted next-generation sequencing identifies a novel nonsense mutation in SPTB for hereditary spherocytosis. Medicine (United States), 2018, 97, e9677.	0.4	7

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91	Measurement of Teicoplanin Concentration With Liquid Chromatography-Tandem Mass Spectrometry Method Demonstrates the Usefulness of Therapeutic Drug Monitoring in Hematologic Patient Populations. Therapeutic Drug Monitoring, 2018, 40, 330-336.	1.0	11
92	Hereditary dehydrated stomatocytosis with splicing site mutation of PIEZO1 mimicking myelodysplastic syndrome diagnosed by targeted nextâ€generation sequencing. Pediatric Blood and Cancer, 2018, 65, e27053.	0.8	9
93	Ubiquitin C decrement plays a pivotal role in replicative senescence of bone marrow mesenchymal stromal cells. Cell Death and Disease, 2018, 9, 139.	2.7	14
94	Heavy/light chain assay as a biomarker for diagnosis and follow-up of multiple myeloma. Clinica Chimica Acta, 2018, 479, 7-13.	0.5	7
95	Molecular drug resistance profiles of Mycobacterium tuberculosis from sputum specimens using ion semiconductor sequencing. Journal of Microbiological Methods, 2018, 145, 1-6.	0.7	9
96	Cytokine and molecular networks in sepsis cases: a network biology approach. European Cytokine Network, 2018, 29, 103-111.	1.1	25
97	PCM1-JAK2 Fusion in a Patient With Acute Myeloid Leukemia. Annals of Laboratory Medicine, 2018, 38, 492-494.	1.2	12
98	Diagnosis of Liver Fibrosis With Wisteria floribunda Agglutinin-Positive Mac-2 Binding Protein (WFA-M2BP) Among Chronic Hepatitis B Patients. Annals of Laboratory Medicine, 2018, 38, 348-354.	1.2	29
99	Clinical Presentation with High Penetrance in a Korean Family with Pulmonary Arterial Hypertension Associated with a BMPR2 Intron 3 Splice Site Pathogenic Variant. Laboratory Medicine Online, 2018, 8, 119.	0.0	0
100	Two Korean Cases of Hereditary Spherocytosis Caused by Mutations inSLC4A1. Laboratory Medicine Online, 2018, 8, 114.	0.0	0
101	Genetic Characteristics and Long-Term Outcomes of Korean Adult Patients with Ph-like Acute Lymphoblastic Leukemia Versus Non-Ph-like Acute Lymphoblastic Leukemia. Blood, 2018, 132, 4087-4087.	0.6	0
102	Report on the External Quality Assessment Scheme for Cytogenetics and Molecular Cytogenetics in Korea (2016â€“2017). Journal of Laboratory Medicine and Quality Assurance, 2018, 40, 188-198.	0.1	0
103	Nonautoimmune congenital hyperthyroidism due to p.Asp633Glu mutation in the TSHR gene. Annals of Pediatric Endocrinology and Metabolism, 2018, 23, 235-239.	0.8	4
104	Practical informativeness of short tandem repeat loci for chimerism analysis in hematopoietic stem cell transplantation. Clinica Chimica Acta, 2017, 468, 51-59.	0.5	9
105	Identification of large genomic rearrangement of BRCA1/2 in high risk patients in Korea. BMC Medical Genetics, 2017, 18, 38.	2.1	10
106	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.	0.5	7
107	Peptide nucleic acid probe-based fluorescence melting curve analysis for rapid screening of common JAK2, MPL, and CALR mutations. Clinica Chimica Acta, 2017, 465, 82-90.	0.5	6
108	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.	0.7	15

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109	Systemic Epstein-Barr Virus-positive T-Cell Lymphoproliferative Disease of Childhood With Good Response to Steroid Therapy. Journal of Pediatric Hematology/Oncology, 2017, 39, e497-e500.	0.3	3
110	Diagnostic exome sequencing identifies a heterozygous MBD5 frameshift mutation in a family with intellectual disability and epilepsy. European Journal of Medical Genetics, 2017, 60, 559-564.	0.7	8
111	Passage-dependent accumulation of somatic mutations in mesenchymal stromal cells during in vitro culture revealed by whole genome sequencing. Scientific Reports, 2017, 7, 14508.	1.6	50
112	Distribution of somatic mutations of cancer-related genes according to microsatellite instability status in Korean gastric cancer. Medicine (United States), 2017, 96, e7224.	0.4	11
113	Considerations when using next-generation sequencing for genetic diagnosis of long-QT syndrome in the clinical testing laboratory. Clinica Chimica Acta, 2017, 464, 128-135.	0.5	7
114	Maternal serum placental growth factor combined with second trimester aneuploidy screening to predict small-for-gestation neonates without preeclampsia. Taiwanese Journal of Obstetrics and Gynecology, 2017, 56, 801-805.	0.5	4
115	Two Cases of Pre-descemet Corneal Dystrophy Associated with X-linked Ichthyosis: A Case Report by Genetic Analysis. Journal of Korean Ophthalmological Society, 2017, 58, 993.	0.0	1
116	Diagnostic approaches for inherited hemolytic anemia in the genetic era. Blood Research, 2017, 52, 84.	0.5	40
117	Hemoglobin Kansas: First Korean Family and Literature Review. Annals of Laboratory Medicine, 2017, 37, 352-354.	1.2	3
118	Association of FLG single nucleotide variations with clinical phenotypes of atopic dermatitis. PLoS ONE, 2017, 12, e0190077.	1.1	10
119	Clinical Characteristics and Prognosis of Fuchs Dystrophy According to COL8A2 Gene Mutation Status. Journal of Korean Ophthalmological Society, 2017, 58, 380.	0.0	0
120	A Case of Pseudohypoparathyroidism Type Ib Caused by Aberrant Methylation in theGNASComplex Locus. Laboratory Medicine Online, 2017, 7, 83.	0.0	0
121	Phenotype of a Patient With a 1p36.11-p35.3 Interstitial Deletion Encompassing the AHDC1. Annals of Laboratory Medicine, 2017, 37, 563-565.	1.2	13
122	Genetic Profiles of Korean Patients With Glucose-6-Phosphate Dehydrogenase Deficiency. Annals of Laboratory Medicine, 2017, 37, 108-116.	1.2	15
123	Better transplant outcome with pre-transplant marrow response after hypomethylating treatment in higher-risk MDS with excess blasts. Oncotarget, 2017, 8, 12342-12354.	0.8	27
124	Outcome and Prognostic Factors for ETV6/RUNX1 Positive Pediatric Acute Lymphoblastic Leukemia Treated at a Single Institution in Korea. Cancer Research and Treatment, 2017, 49, 446-453.	1.3	12
125	Birt-Hogg-DubÃ© Syndrome Manifesting as Spontaneous Pneumothorax: A Novel Mutation of the Folliculin Gene. Korean Journal of Thoracic and Cardiovascular Surgery, 2017, 50, 386-390.	0.6	3
126	Patient With Delayed Development Resulting From De Novo Duplication of 7q36.1-q36.3 and Deletion of 9p24.3. Annals of Rehabilitation Medicine, 2017, 41, 881.	0.6	1

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127	Comparison of Targeted Next-Generation and Sanger Sequencing for the BRCA1 and BRCA2 Mutation Screening. Annals of Laboratory Medicine, 2016, 36, 197-201.	1.2	7
128	Extranasal natural killer/T-cell lymphoma initially presenting as myelofibrosis. Blood Research, 2016, 51, 7.	0.5	0
129	Molecular Features of Three Children Diagnosed With Early T-Cell Precursor Acute Lymphoblastic Leukemia. Annals of Laboratory Medicine, 2016, 36, 384-386.	1.2	1
130	A Novel Syntaxin 11 Gene (STX11) Mutation c.650T>C, p.Leu217Pro, in a Korean Child With Familial Hemophagocytic Lymphohistiocytosis. Annals of Laboratory Medicine, 2016, 36, 170-173.	1.2	2
131	Budding Yeast Cells in Peripheral Blood Smear: Clue to Candidemia. Infection and Chemotherapy, 2016, 48, 342.	1.0	1
132	Potential Risk Factors Associated With Vascular Diseases in Patients Receiving Treatment for Hypertension. Annals of Laboratory Medicine, 2016, 36, 215-222.	1.2	0
133	Analytical Performance Evaluation of Infopia Elementâ„¢ Autoâ€coding Blood Glucose Monitoring System for Selfâ€Monitoring of Blood Glucose. Journal of Clinical Laboratory Analysis, 2016, 30, 849-858.	0.9	3
134	A twin sibling with Prader-Willi syndrome caused by type 2 microdeletion following assisted reproductive technology: A case report. Biomedical Reports, 2016, 5, 18-22.	0.9	11
135	Circulating immune cell phenotype can predict the outcome of lenalidomide plus low-dose dexamethasone treatment in patients with refractory/relapsed multiple myeloma. Cancer Immunology, Immunotherapy, 2016, 65, 983-994.	2.0	21
136	Molecular analysis of myocilin and optineurin genes in Korean primary glaucoma patients. Molecular Medicine Reports, 2016, 14, 2439-2448.	1.1	9
137	Geneticâ€“pathologic characterization of myeloproliferative neoplasms. Experimental and Molecular Medicine, 2016, 48, e247-e247.	3.2	14
138	Tissue-specific Differentiation Potency of Mesenchymal Stromal Cells from Perinatal Tissues. Scientific Reports, 2016, 6, 23544.	1.6	92
139	Characterization of leukemias with ETV6-ABL1 fusion. Haematologica, 2016, 101, 1082-1093.	1.7	66
140	Identification of small marker chromosomes using microarray comparative genomic hybridization and multicolor fluorescent in situ hybridization. Molecular Cytogenetics, 2016, 9, 61.	0.4	20
141	Mutational spectrum of Korean patients with corneal dystrophy. Clinical Genetics, 2016, 89, 678-689.	1.0	14
142	Mutational characteristics of <scp>ANK1</scp> and <scp>SPTB</scp> genes in hereditary spherocytosis. Clinical Genetics, 2016, 90, 69-78.	1.0	69
143	Significance of KIT exon 17 mutation depends on mutant level rather than positivity in core-binding factor acute myeloid leukemia. Blood Cancer Journal, 2016, 6, e387-e387.	2.8	14
144	Evaluation of the Performance of Two Point-of-Care Analyzers for Total Cholesterol, Triglyceride, and High-Density Lipoprotein Cholesterol Analysis. Clinical Laboratory, 2016, 62, 1201-1208.	0.2	1

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145	Novel <scp>FLG</scp> null mutations in Korean patients with atopic dermatitis and comparison of the mutational spectra in Asian populations. Journal of Dermatology, 2015, 42, 867-873.	0.6	39
146	Novel 5.712 kb mitochondrial DNA deletion in a patient with Pearson syndrome: A case report. Molecular Medicine Reports, 2015, 11, 3741-3745.	1.1	4
147	Copy number variations could predict the outcome of bortezomib plus melphalan and prednisone for initial treatment of multiple myeloma. Genes Chromosomes and Cancer, 2015, 54, 20-27.	1.5	9
148	Fragmented Red Cell as a Possible Favorable Prognostic Marker of Hematopoietic Stem Cell Transplantation Associated Thrombotic Microangiopathy. Journal of Clinical Laboratory Analysis, 2015, 29, 444-450.	0.9	2
149	Two Cases of Shwachman-Diamond Syndrome in Adolescents Confirmed by Genetic Analysis. Annals of Laboratory Medicine, 2015, 35, 269-271.	1.2	3
150	Identification of Compound Heterozygous Mutations in the BBS7 Gene in a Korean Family with Bardet-Biedl Syndrome. Annals of Laboratory Medicine, 2015, 35, 181-184.	1.2	11
151	Flow Cytometric White Blood Cell Differential Using CytoDiff is Excellent for Counting Blasts. Annals of Laboratory Medicine, 2015, 35, 28-34.	1.2	13
152	T618I-Mutated Colony Stimulating Factor 3 Receptor in Chronic Neutrophilic Leukemia and Chronic Myelomonocytic Leukemia Patients who Underwent Allogeneic Stem Cell Transplantation. Annals of Laboratory Medicine, 2015, 35, 376-378.	1.2	13
153	3D printing technology to control BMP-2 and VEGF delivery spatially and temporally to promote large-volume bone regeneration. Journal of Materials Chemistry B, 2015, 3, 5415-5425.	2.9	151
154	Diagnosis and evaluation of severity of sepsis via the use of biomarkers and profiles of 13 cytokines: a multiplex analysis. Clinical Chemistry and Laboratory Medicine, 2015, 53, 575-81.	1.4	58
155	Quantitative fragment analysis of FLT3-ITD efficiently identifying poor prognostic group with high mutant allele burden or long ITD length. Blood Cancer Journal, 2015, 5, e336-e336.	2.8	59
156	Genetic and epigenetic alterations of bone marrow stromal cells in myelodysplastic syndrome and acute myeloid leukemia patients. Stem Cell Research, 2015, 14, 177-184.	0.3	40
157	Wilms Tumor Gene 1 Expression as a Predictive Marker for Relapse and Survival after Hematopoietic Stem Cell Transplantation for Myelodysplastic Syndromes. Biology of Blood and Marrow Transplantation, 2015, 21, 460-467.	2.0	16
158	FLT3 expression and IL10 promoter polymorphism in acute myeloid leukemia with RUNX1-RUNX1T1. Molecular Biology Reports, 2015, 42, 451-456.	1.0	5
159	Autistic and Rettâ€like features associated with 2q33.3â€“q34 interstitial deletion. American Journal of Medical Genetics, Part A, 2015, 167, 2213-2218.	0.7	24
160	Mutational analysis of oncogenic CSF3R p.T618I in acute leukemias and common solid cancers. Annals of Hematology, 2015, 94, 889-890.	0.8	0
161	Microenvironmental Remodeling as a Parameter and Prognostic Factor of Heterogeneous Leukemogenesis in Acute Myelogenous Leukemia. Cancer Research, 2015, 75, 2222-2231.	0.4	124
162	Novel oncogenic <scp>PTPN11</scp> mutations in myelodysplastic syndrome in Korean patients. Hematological Oncology, 2015, 33, 166-167.	0.8	2

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163	Founder Haplotype Analysis of Fanconi Anemia in the Korean Population Finds Common Ancestral Haplotypes for a FANCG Variant. Annals of Human Genetics, 2015, 79, 153-161.	0.3	4
164	Determination of posaconazole concentration with LCâ€“MS/MS in adult patients with hematologic malignancy. Clinica Chimica Acta, 2015, 450, 220-226.	0.5	15
165	The effect of fibroblast growth factor on distinct differentiation potential of cord bloodâ€“derived unrestricted somatic stem cells and Wharton's jellyâ€“derived mesenchymal stem/stromal cells. Cytotherapy, 2015, 17, 1723-1731.	0.3	9
166	Clinical outcomes of venous thromboembolism with dalteparin therapy in multiple myeloma patients. Thrombosis Research, 2015, 136, 974-979.	0.8	11
167	Neutrophil Gelatinase-Associated Lipocalin as a Biomarker of Renal Impairment in Patients With Multiple Myeloma. Clinical Lymphoma, Myeloma and Leukemia, 2015, 15, 35-40.	0.2	8
168	Two cases of concurrent development of essential thrombocythemia with chronic lymphocytic leukemia, one related to clonal B-cell lymphocytosis, tested by array comparative genomic hybridization. International Journal of Hematology, 2015, 101, 612-619.	0.7	5
169	Characterization of Leukemias with ETV6-ABL1 Fusion. Blood, 2015, 126, 84-84.	0.6	1
170	Molecular analysis of the CHST6 gene in Korean patients with macular corneal dystrophy: Identification of three novel mutations. Molecular Vision, 2015, 21, 1201-9.	1.1	8
171	White blood cell differential counts in severely leukopenic samples: a comparative analysis of different solutions available in modern laboratory hematology. Blood Research, 2014, 49, 120.	0.5	21
172	Effects of ECM Protein Mimetics on Adhesion and Proliferation of Chorion Derived Mesenchymal Stem Cells. International Journal of Medical Sciences, 2014, 11, 298-308.	1.1	27
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174	Novel Markers of Early Neutrophilic and Monocytic Engraftment after Hematopoietic Stem Cell Transplantation. Annals of Laboratory Medicine, 2014, 34, 92-97.	1.2	9
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