Kyung-A Lee

List of Publications by Year
in descending order

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147	1,716	361045	454577
papers	citations	20 h-index	30 g-index

157	157	157	3131
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Delta Neutrophil Index. Shock, 2012, 37, 242-246.	1.0	102
2	Novel interleukin 1ï¿½ polymorphism increased the risk of gastric cancer in a Korean population. Journal of Gastroenterology, 2004, 39, 429-433.	2.3	84
3	Haplotype Structure of the UDP-Glucuronosyltransferase 1A1 (UGT1A1) Gene and Its Relationship to Serum Total Bilirubin Concentration in a Male Korean Population. Clinical Chemistry, 2003, 49, 2078-2081.	1.5	55
4	Diagnostic performance of <scp>CA</scp> 125, <scp>HE</scp>4, and risk of Ovarian Malignancy Algorithm for ovarian cancer. Journal of Clinical Laboratory Analysis, 2019, 33, e22624.	0.9	49
5	Complete Sequencing of a Genetic Polymorphism in NAT2 in the Korean Population. Clinical Chemistry, 2002, 48, 775-777.	1.5	48
6	Prevalence of sexually transmitted infections among healthy Korean women: Implications of multiplex PCR pathogen detection on antibiotic therapy. Journal of Infection and Chemotherapy, 2014, 20, 74-76.	0.8	38
7	Assessment of real-time PCR method for detection of EGFR mutation using both supernatant and cell pellet of malignant pleural effusion samples from non-small-cell lung cancer patients. Clinical Chemistry and Laboratory Medicine, 2017, 55, 1962-1969.	1.4	36
8	Profiling cancer-associated genetic alterations and molecular classification of cancer in Korean gastric cancer patients. Oncotarget, 2017, 8, 69888-69905.	0.8	34
9	Three Cases of Manifesting Female Carriers in Patients with Duchenne Muscular Dystrophy. Yonsei Medical Journal, 2011, 52, 192.	0.9	32
10	A novel approach for tuberculosis diagnosis using exosomal DNA and droplet digital PCR. Clinical Microbiology and Infection, 2020, 26, 942.e1-942.e5.	2.8	29
11	Validation and optimization of the Ion Torrent S5 XL sequencer and Oncomine workflow for BRCA1 and BRCA2 genetic testing. Oncotarget, 2017, 8, 34858-34866.	0.8	29
12	Acute promyelocytic leukemia with insertion of PML exon 7a and partial deletion of exon 3 of RARA: a novel variant transcript related to aggressive course and not detected with real-time polymerase chain reaction analysis. Cancer Genetics and Cytogenetics, 2009, 188, 103-107.	1.0	25
13	Molecular characterization of alternative SET-NUP214 fusion transcripts in a case of acute undifferentiated leukemia. Cancer Genetics and Cytogenetics, 2010, 201, 73-80.	1.0	24
14	Spectrum of EGFR Gene Copy Number Changes and KRAS Gene Mutation Status in Korean Triple Negative Breast Cancer Patients. PLoS ONE, 2013, 8, e79014.	1.1	24
15	JAK2 V617F/C618R mutation in a patient with polycythemia vera:A case study and review of the literature. Cancer Genetics and Cytogenetics, 2009, 189, 43-47.	1.0	23
16	Clinical and Genetic Characterization of Female Dystrophinopathy. Journal of Clinical Neurology		

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19	Birt-Hogg-Dube syndrome prospectively detected by review of chest computed tomography scans. PLoS ONE, 2017, 12, e0170713.	1.1	22
20	TP53BP2 locus is associated with gastric cancer susceptibility. International Journal of Cancer, 2005, 117, 957-960.	2.3	21
21	Cytochrome P450 2C19 Polymorphism is Associated with Reduced Clopidogrel Response in Cerebrovascular Disease. Yonsei Medical Journal, 2011, 52, 734.	0.9	20
22	Delta neutrophil index discriminates true bacteremia from blood culture contamination. Clinica Chimica Acta, 2014, 427, 11-14.	0.5	20
23	Automated Detection of Malaria-Associated Pseudoeosinophilia and Abnormal WBC Scattergram by the Sysmex XE-2100 Hematology Analyzer: A Clinical Study with 1,801 Patients and Real-Time Quantitative PCR Analysis in Vivax Malaria-Endemic Area. American Journal of Tropical Medicine and Hygiene, 2010, 82, 412-414.	0.6	19
24	PRSS1, SPINK1, CFTR, and CTRC Pathogenic Variants in Korean Patients With Idiopathic Pancreatitis. Annals of Laboratory Medicine, 2016, 36, 555-560.	1.2	19
25	A Comparative Study for Detection of EGFR Mutations in Plasma Cell-Free DNA in Korean Clinical Diagnostic Laboratories. BioMed Research International, 2018, 2018, 1-11.	0.9	19
26	Ethnic differences in gastric cancer genetic susceptibility: Allele flips of interleukin gene. World Journal of Gastroenterology, 2014, 20, 4558.	1.4	19
27	Rare translocations involving chromosome band 8p11 in myeloid neoplasms. Cancer Genetics and Cytogenetics, 2008, 186, 127-129.	1.0	18
28	Exosome-based detection of EGFR T790M in plasma and pleural fluid of prospectively enrolled non-small cell lung cancer patients after first-line tyrosine kinase inhibitor therapy. Cancer Cell International, 2021, 21, 50.	1.8	18
29	Detection of FUSâ€“ERG chimeric transcript in two cases of acute myeloid leukemia with t(16;21)(p11.2;q22) with unusual characteristics. Cancer Genetics and Cytogenetics, 2009, 194, 111-118.	1.0	17
30	Detection of Immunoglobulin Heavy Chain Gene Clonality by Next-Generation Sequencing for Minimal Residual Disease Monitoring in B-Lymphoblastic Leukemia. Annals of Laboratory Medicine, 2017, 37, 331-335.	1.2	17
31	Selecting short length nucleic acids localized in exosomes improves plasma EGFR mutation detection in NSCLC patients. Cancer Cell International, 2019, 19, 251.	1.8	17
32	Interaction of polymorphisms in the Interleukin 1B-31 and general transcription factor 2A1 genes on the susceptibility to gastric cancer. Cytokine, 2007, 38, 96-100.	1.4	16
33	Detection of a novel CBFB/MYH11 variant fusion transcript (K-type) showing partial insertion of exon 6 of CBFB gene using two commercially available multiplex RT-PCR kits. Cancer Genetics and Cytogenetics, 2009, 189, 87-92.	1.0	16
34	Routine Chromosomal Microarray Analysis is Necessary in Korean Patients With Unexplained Developmental Delay/Mental Retardation/Autism Spectrum Disorder. Annals of Laboratory Medicine, 2015, 35, 510-518.	1.2	16
35	Multiplex Ligation-Dependent Probe Amplification in X-linked Recessive Muscular Dystrophy in Korean Subjects. Yonsei Medical Journal, 2017, 58, 613.	0.9	16
36	Novel in-frame deletion mutation in FLCN gene in a Korean family with recurrent primary spontaneous pneumothorax. Gene, 2012, 499, 339-342.	1.0	15

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37	Low PR in ER(+)/HER2(â˜) breast cancer: high rates of TP53 mutation and high SUV. Endocrine-Related Cancer, 2019, 26, 177-185.	1.6	15
38	Clinical Practice Guidelines for Pre-Analytical Procedures of Plasma Epidermal Growth Factor Receptor Variant Testing. Annals of Laboratory Medicine, 2022, 42, 141-149.	1.2	15
39	Efficacy of imatinib mesylate (STI571) in chronic neutrophilic leukemia with t(15;19): Case report. American Journal of Hematology, 2004, 77, 366-369.	2.0	14
40	Electrocardiography based prediction of hypertrophy pattern and fibrosis amount in hypertrophic cardiomyopathy: comparative study with cardiac magnetic resonance imaging. International Journal of Cardiovascular Imaging, 2018, 34, 1619-1628.	0.7	14
41	Acute promyelocytic leukemia in early pregnancy with translocation t(15;17) and variant PML/RARA fusion transcripts. Cancer Genetics and Cytogenetics, 2009, 188, 48-51.	1.0	13
42	Three-way translocation involving MLL, MLLT1, and a novel third partner, NRXN1, in a patient with acute lymphoblastic leukemia and t(2;19;11) (p12;p13.3;q23). Cancer Genetics and Cytogenetics, 2010, 197, 32-38.	1.0	13
43	Interleukin 10 polymorphisms differentially influence the risk of gastric cancer in East Asians and Caucasians. Cytokine, 2010, 51, 73-77.	1.4	13
44	Patterns and Biologic Features of p53 Mutation Types in Korean Breast Cancer Patients. Journal of Breast Cancer, 2014, 17, 1.	0.8	13
45	Development and Comparison of Warfarin Dosing Algorithms in Stroke Patients. Yonsei Medical Journal, 2016, 57, 635.	0.9	13
46	Cerebellar vermis hypoplasia in CHARGE syndrome: clinical and molecular characterization of 18 unrelated Korean patients. Journal of Human Genetics, 2016, 61, 235-239.	1.1	13
47	Analytical validation of the droplet digital PCR assay for diagnosis of spinal muscular atrophy. Clinica Chimica Acta, 2020, 510, 787-789.	0.5	13
48	Therapy-related acute lymphoblastic leukemia with t(9;22)(q34;q11.2):a case study and review of the literature. Cancer Genetics and Cytogenetics, 2009, 191, 51-54.	1.0	12
49	Determination of SMN1 and SMN2 Copy Numbers in a Korean Population using Multiplex Ligation-dependent Probe Amplification. Annals of Laboratory Medicine, 2010, 30, 93-96.	1.2	12
50	LEOPARD Syndrome with PTPN11 Gene Mutation Showing Six Cardinal Symptoms of LEOPARD. Annals of Dermatology, 2011, 23, 232.	0.3	12
51	CD5-negative Blastoid Variant Mantle Cell Lymphoma with Complex CCND1/IGH and MYC Aberrations. Annals of Laboratory Medicine, 2012, 32, 95-98.	1.2	12
52	Populationâ€specific spectrum of the F11 mutations in Koreans: evidence for a founder effect. Clinical Genetics, 2012, 82, 180-186.	1.0	12
53	A novel synonymous mutation causing complete skipping of exon 16 in the SLC26A4 gene in a Korean family with hearing loss. Biochemical and Biophysical Research Communications, 2013, 430, 1147-1150.	1.0	12
54	Cataloging Coding Sequence Variations in Human Genome Databases. PLoS ONE, 2008, 3, e3575.	1.1	12

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55	Heterozygosities of 735 microsatellite markers and background linkage disequilibrium in the Korean population. Experimental and Molecular Medicine, 2006, 38, 662-667.	3.2	11
56	Intrafamilial phenotypic variability in families with biallelic SLC26A4 mutations. Laryngoscope, 2014, 124, E194-202.	1.1	11
57	Effects of Triflusal and Clopidogrel on the Secondary Prevention of Stroke Based on Cytochrome P450 2C19 Genotyping. Journal of Stroke, 2017, 19, 356-364.	1.4	11
58	DeviCNV: detection and visualization of exon-level copy number variants in targeted next-generation sequencing data. BMC Bioinformatics, 2018, 19, 381.	1.2	11
59	A novel missense mutation (I344K) in the SPG4 gene in a Korean family with autosomal-dominant hereditary spastic paraplegia. Journal of Human Genetics, 2002, 47, 473-477.	1.1	10
60	BCR/ABL rearrangement with b3a3 fusion transcript in a case of childhood acute lymphoblastic leukemia. Cancer Genetics and Cytogenetics, 2009, 189, 132-137.	1.0	10
61	CASP8AP2 is a novel partner gene of MLL rearrangement with t(6;11)(q15;q23) in acute myeloid leukemia. Cancer Genetics and Cytogenetics, 2009, 195, 94-95.	1.0	10
62	HomozygousSMN2Deletion is a Major Risk Factor among Twenty-Five Korean Sporadic Amyotrophic Lateral Sclerosis Patients. Yonsei Medical Journal, 2012, 53, 53.	0.9	10
63	Copy number variation and gene rearrangements inCYP2D6genotyping using multiplex ligation-dependent probe amplification in Koreans. Pharmacogenomics, 2012, 13, 963-973.	0.6	10
64	Evaluation of Three Automated Nucleic Acid Extraction Systems for Identification of Respiratory Viruses in Clinical Specimens by Multiplex Real-Time PCR. BioMed Research International, 2014, 2014, 1-8.	0.9	10
65	Breakpoint mapping by whole genome sequencing identifiesPTH2Rgene disruption in a patient with midline craniosynostosis and a de novo balanced chromosomal rearrangement. Journal of Medical Genetics, 2015, 52, 706-709.	1.5	10
66	Performance evaluation of cobas HBV real-time PCR assay on Roche cobas 4800 System in comparison with COBAS AmpliPrep/COBAS TaqMan HBV Test. Clinical Chemistry and Laboratory Medicine, 2018, 56, 1133-1139.	1.4	10
67	Significant therapeutic effects of adult human multipotent neural cells on spinal cord injury. Stem Cell Research, 2018, 31, 71-78.	0.3	10
68	Acute erythroleukemia with der(1;7)(q10;p10) as a sole acquired abnormality after treatment with azathioprine. Cancer Genetics and Cytogenetics, 2008, 186, 58-60.	1.0	9
69	A tandem triplication, trp(1)(q21q32), in a patient with follicular lymphoma: a case study and review of the literature. Cancer Genetics and Cytogenetics, 2009, 189, 127-131.	1.0	9
70	Hereditary protein S deficiency from a novel large deletion mutation of the PROS1 gene detected by multiplex ligation-dependent probe amplification (MLPA). Thrombosis Research, 2009, 123, 793-795.	0.8	9
71	Rapid Identification of Thrombocytopenia-Associated Multiple Organ Failure Using Red Blood Cell Parameters and a Volume/Hemoglobin Concentration Cytogram. Yonsei Medical Journal, 2011, 52, 845.	0.9	9
72	Non-syndromic hearing loss caused by the dominant cis mutation R75Q with the recessive mutation V37I of the GJB2 (Connexin 26) gene. Experimental and Molecular Medicine, 2015, 47, e169-e169.	3.2	9

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73	A New Integrated Newborn Screening Workflow Can Provide a Shortcut to Differential Diagnosis and Confirmation of Inherited Metabolic Diseases. Yonsei Medical Journal, 2018, 59, 652.	0.9	9
74	Contribution of sarcomere gene mutations to left atrial function in patients with hypertrophic cardiomyopathy. Cardiovascular Ultrasound, 2021, 19, 4.	0.5	9
75	Primary endocrine resistance of ER+ breast cancer with ESR1 mutations interrogated by droplet digital PCR. Npj Breast Cancer, 2022, 8, 58.	2.3	9
76	ALK-positive anaplastic large cell lymphoma with TPM3-ALK translocation. Leukemia Research, 2012, 36, e143-e145.	0.4	8
77	A Case of Late-Onset Li-Fraumeniâ€“like Syndrome with Unilateral Breast Cancer. Annals of Laboratory Medicine, 2013, 33, 212-216.	1.2	8
78	Differential contributions of sarcomere and mitochondria-related multigene variants to the endophenotype of hypertrophic cardiomyopathy. Mitochondrion, 2020, 53, 48-56.	1.6	8
79	Application of CRISPR/Cas9-based mutant enrichment technique to improve the clinical sensitivity of plasma EGFR testing in patients with non-small cell lung cancer. Cancer Cell International, 2022, 22, 82.	1.8	8
80	Genetic relevance and determinants of mitral leaflet size in hypertrophic cardiomyopathy. Cardiovascular Ultrasound, 2019, 17, 21.	0.5	7
81	De Novo Cancer Incidence after Kidney Transplantation in South Korea from 2002 to 2017. Journal of Clinical Medicine, 2021, 10, 3530.	1.0	7
82	Identification of cell morphology parameters from automatic hematology analyzers to predict the peripheral blood CD34-positive cell count after mobilization. PLoS ONE, 2017, 12, e0174286.	1.1	7
83	A novel de novo mutation in the serine-threonine kinase STK11 gene in a Korean patient with Peutz-Jeghers syndrome. BMC Medical Genetics, 2008, 9, 44.	2.1	6
84	Tetrasomy 8 in a Patient with Acute Monoblastic Leukemia. Annals of Laboratory Medicine, 2008, 28, 262-266.	1.2	6
85	The First Korean Case of Camurati-Engelmann Disease (Progressive Diaphyseal Dysplasia) Confirmed by TGFB1 Gene Mutation Analysis. Journal of Korean Medical Science, 2009, 24, 737.	1.1	6
86	First Korean Case of SATB2-Associated 2q32-q33 Microdeletion Syndrome. Annals of Laboratory Medicine, 2015, 35, 275-278.	1.2	6
87	Effect of sarcomere and mitochondria-related mutations on myocardial fibrosis in patients with hypertrophic cardiomyopathy. Journal of Cardiovascular Magnetic Resonance, 2021, 23, 18.	1.6	6
88	A der(1;15)(q10;q10) is a rare nonrandom whole-arm translocation in patients with acute lymphoblastic leukemia. Cancer Genetics and Cytogenetics, 2007, 179, 132-135.	1.0	5
89	MLL rearrangement with t(6;11)(q15;q23) as a sole abnormality in a patient with de novo acute myeloid leukemia: conventional cytogenetics, FISH, and multicolor FISH analyses for detection of rare MLL-related chromosome abnormalities. Cancer Genetics and Cytogenetics, 2008, 187, 50-53.	1.0	5
90	Two case reports of 1q triplication in myeloproliferative neoplasms. Cancer Genetics and Cytogenetics, 2009, 191, 111-112.	1.0	5

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91	Ambras syndrome in a Korean patient with balanced pericentric inversion (8)(p11.2q24.2). Journal of Dermatological Science, 2010, 59, 204-206.	1.0	5
92	Cytogenetic features of 5q deletion and 5qâ^' syndrome in myelodysplastic syndrome in Korea; marker chromosomes proved to be chromosome 5 with interstitial deletion by fluorescence in situ hybridization. Cancer Genetics and Cytogenetics, 2010, 203, 193-202.	1.0	5
93	A Novel PTEN Mutation in a Korean Patient with Cowden Syndrome and Vascular Anomalies. Acta Dermato-Venereologica, 2011, 91, 88-90.	0.6	5
94	A novel association between relaxin receptor polymorphism and hematopoietic stem cell yield after mobilization. PLoS ONE, 2017, 12, e0179986.	1.1	5
95	Concomitant AID Expression and BCL7A Loss Associates With Accelerated Phase Progression and Imatinib Resistance in Chronic Myeloid Leukemia. Annals of Laboratory Medicine, 2017, 37, 177-179.	1.2	5
96	Clinical Features of Multiple Acyl-CoA Dehydrogenase Deficiency With ETFDH Variants in the First Korean Cases. Annals of Laboratory Medicine, 2018, 38, 616-618.	1.2	5
97	Novel and Recurrent ACADS Mutations and Clinical Manifestations Observed in Korean Patients with Short-chain Acyl-coenzyme a Dehydrogenase Deficiency. Annals of Clinical and Laboratory Science, 2016, 46, 360-6.	0.2	5
98	t(5;12)(q13;p13) in acute myeloid leukemia with preceding granulocytic sarcoma. Cancer Genetics and Cytogenetics, 2007, 177, 158-160.	1.0	4
99	Paracentric inversionâ€“associated t(8;21) variant in de novo acute myelogenous leukemia: characteristic patterns of conventional cytogenetics, FISH, and multicolor banding analysis. Cancer Genetics and Cytogenetics, 2008, 183, 72-76.	1.0	4
100	Differential association of RANTES-403 and IL-1B-1464 polymorphisms on histological subtypes in male Korean patients with gastric cancer. Tumor Biology, 2014, 35, 3765-3770.	0.8	4
101	Frequency and Clinical Characteristics of Unselected Korean Gastric Cancer Patients with a Germline CDH1 V832M Mutation. Journal of Cancer, 2020, 11, 208-212.	1.2	4
102	Genetic Spectrum ofUGT1A1in Korean Patients with Unconjugated Hyperbilirubinemia. Annals of Laboratory Medicine, 2020, 40, 281-283.	1.2	4
103	Detection of EGFA-SEPT14 fusion in cell-free DNA of a patient with advanced gastric cancer: A case report. World Journal of Clinical Cases, 2021, 9, 2884-2889.	0.3	4
104	A Population-Based Analysis of BRCA1/2 Genes and Associated Breast and Ovarian Cancer Risk in Korean Patients: A Multicenter Cohort Study. Cancers, 2021, 13, 2192.	1.7	4
105	Clinical Pharmacogenetic Testing and Application: Laboratory Medicine Clinical Practice Guidelines Part 2. Laboratory Medicine Online, 2016, 6, 193.	0.0	4
106	Association between TP53 mutation and high 21-gene recurrence score in estrogen receptor-positive/HER2-negative breast cancer. Npj Breast Cancer, 2022, 8, 19.	2.3	4
107	Comparison of ILâ€6 measurement methods with a special emphasis on COVIDâ€19 patients according to equipment and sample type. Journal of Clinical Laboratory Analysis, 2022, 36, e24182.	0.9	4
108	Complex t(8;19;21)(q22;p13;q22) as a sole abnormality in a patient with de novo acute myeloid leukemia. Cancer Genetics and Cytogenetics, 2008, 185, 109-112.	1.0	3

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109	Trisomy 8 in an elderly patient with acute lymphoblastic leukemia as a sole abnormality. Cancer Genetics and Cytogenetics, 2008, 187, 57-58.	1.0	3
110	Concomitant t(3;3)(q21;q26), trisomy 19, and E255V mutation associated with imatinib mesylate resistance in chronic myelogenous leukemia. Cancer Genetics and Cytogenetics, 2009, 190, 46-48.	1.0	3
111	A novel F11 mutation in a Korean pediatric patient with recurrent epistaxis. Blood Coagulation and Fibrinolysis, 2013, 24, 433-435.	0.5	3
112	Partial Gene Deletions ofPMP22Causing Hereditary Neuropathy with Liability to Pressure Palsies. Case Reports in Genetics, 2014, 2014, 1-3.	0.1	3
113	Clinical Pharmacogenetic Testing and Application: Laboratory Medicine Clinical Practice Guidelines Part 1. Laboratory Medicine Online, 2016, 6, 119.	0.0	3
114	Burden of premature ventricular contractions beyond nonsustained ventricular tachycardia is related to the myocardial extracellular space expansion in patients with hypertrophicâ€cardiomyopathy. Clinical Cardiology, 2020, 43, 1317-1325.	0.7	3
115	Identification of a novel <scp>HLAâ€C</scp>*03:04 allele, <scp>HLAâ€C</scp>*03:04:84, in a Korean individual. Hla, 2021, 97, 156-158.	0.4	3
116	Comparison of antinuclear antibody profiles obtained using line immunoassay and fluorescence enzyme immunoassay. Journal of International Medical Research, 2021, 49, 030006052110143.	0.4	3
117	An optimizedBRCA1/2next-generation sequencing for different clinical sample types. Journal of Gynecologic Oncology, 2020, 31, e9.	1.0	3
118	Establishment of Reference Intervals for Serum Insulin-Like Growth Factor I in Korean Adult Population. Endocrinology and Metabolism, 2020, 35, 960-964.	1.3	3
119	A novel missense MSH2 gene mutation in a patient of a Korean family with hereditary nonpolyposis colorectal cancer. Cancer Genetics and Cytogenetics, 2008, 182, 136-139.	1.0	2
120	Clinical Significance of von Willebrand Factor-Cleaving Protease (ADAMTS13) Deficiency in Patients with Sepsis-Induced Disseminated Intravascular Coagulation. Infection and Chemotherapy, 2009, 41, 78.	1.0	2
121	Three new nonsense mutations of MLH1 and MSH2 genes in Korean families with hereditary nonpolyposis colorectal cancer. Cancer Genetics and Cytogenetics, 2009, 188, 61-64.	1.0	2
122	Biphenotypic acute leukemia with b2a2 fusion transcript and trisomy 21. Cancer Genetics and Cytogenetics, 2009, 188, 129-131.	1.0	2
123	Acute Promyelocytic Leukemia with Trisomy 8 and del(9)(q22) after Treatment of Cervical Cancer with Concurrent Chemoradiotherapy: A Case Report. Onkologie, 2011, 34, 388-390.	1.1	2
124	Functional Haplotype Frequencies of the Interleukin-1B Promoter in the Korean Population. Genomics and Informatics, 2008, 6, 29-31.	0.4	2
125	Detection of Anti-Extractable Nuclear Antigens in Patients with Systemic Rheumatic Disease via Fluorescence Enzyme Immunoassay and Its Clinical Utility. Yonsei Medical Journal, 2020, 61, 73.	0.9	2
126	Effectiveness of in silico tagSNP selection methods: virtual analysis of the genotypes of pharmacogenetic genes. Pharmacogenomics, 2007, 8, 1347-1357.	0.6	1

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127	Distinct Linkage Disequilibrium (LD) Runs of Single Nucleotide Polymorphisms and Microsatellite Markers; Implications for Use of Mixed Marker Haplotypes in LD-based Mapping. Journal of Korean Medical Science, 2007, 22, 425.	1.1	1
128	Three-way Philadelphia variant t(9;22;14)(q34;q11.2;p11) in chronic myeloid leukemia. Cancer Genetics and Cytogenetics, 2009, 191, 55-56.	1.0	1
129	Cys482Trp Missense Mutation in the Coagulation Factor XI Gene (F11) in a Korean Patient with Factor XI Deficiency. Annals of Laboratory Medicine, 2014, 34, 332-335.	1.2	1
130	Bone marrow hypoplasia, isochromosome 8q and deletion of chromosome 6q preceding B-cell lymphoma. Blood Research, 2014, 49, 200.	0.5	1
131	Deletion of 20p13 and Duplication of 20p13p12.3 in a Patient with Delayed Speech and Development. Annals of Laboratory Medicine, 2018, 38, 77-79.	1.2	1
132	Korean Monozygotic Twins with Lethal Acantholytic Epidermolysis Bullosa Caused by Two Novel Mutations. Annals of Clinical and Laboratory Science, 2017, 47, 213-216.	0.2	1
133	Comparison of VERSANT Hepatitis B Virus DNA 3.0 Assay with Digene Hybrid Capture II Hepatitis B Virus DNA Test in Relation to Clinical Status of Hepatitis B Virus. Annals of Laboratory Medicine, 2007, 27, 451-457.	1.2	0
134	Refractory anemia with ring sideroblasts associated with marked thrombocytosis harboring cytogenetic abnormality dup(2)(p15p22) treated with decitabine. Leukemia and Lymphoma, 2012, 53, 2287-2289.	0.6	0
135	A novel three-way variant t(4;17;5)(p16;q23;q31) in a case of secondary plasma cell leukemia. Leukemia Research, 2012, 36, e101-e102.	0.4	0
136	Novel Non-contiguous Duplications in theDMDGene in Five Patients with Duchenne Muscular Dystrophy. Laboratory Medicine Online, 2015, 5, 121.	0.0	0
137	Diagnosis of Smith-Magenis Syndrome in a Patient with Mental Retardation and Sleep Disturbance Confirmed by Multiplex Ligation-dependent Probe Amplification. Laboratory Medicine Online, 2018, 8, 71.	0.0	0
138	Hereditary cancer syndrome-associated pathogenic variants are common in patients with hematologic malignancies subsequent to primary solid cancer. Journal of Cancer, 2021, 12, 4288-4294.	1.2	0
139	Lineage switch of Bâ€lymphoblastic leukemia into acute myeloid leukemia with residual lymphoblasts in a patient with previous breast cancer. International Journal of Laboratory Hematology, 2021, 43, O197-O199.	0.7	0
140	Performance Evaluation of Aptima HBV and HCV Quant Assays in the Panther System. Laboratory Medicine Online, 2021, 11, 177-182.	0.0	0
141	Identification of a UGT1A1*37 Allele in a Korean Patient with Pancreatic Cancer. Laboratory Medicine Online, 2021, 11, 199-202.	0.0	0
142	Evaluation of a hybridization capture-based hereditary cancer panel for the ion semiconductor-based next-generation sequencing system. Clinica Chimica Acta, 2021, 521, 223-228.	0.5	0
143	CYP2C19 Polymorphisms and Smoking Status Affects Responsiveness to the Platelet P2Y12 Receptor Antagonist Clopidogrel. Cardiovascular Prevention and Pharmacotherapy, 2019, 1, 63.	0.0	0
144	Diagnostic Challenge: Primary Bone Marrow Diffuse Large B-cell Lymphoma Mimicking Systemic Autoimmune Diseases. Laboratory Medicine Online, 2019, 9, 242.	0.0	0

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145	Rare Korean Cases of Very-long-chain Acyl-CoA Dehydrogenase Deficiency with a Novel Recurrent Mutation. Annals of Clinical and Laboratory Science, 2016, 46, 97-101.	0.2	0
146	Performance Evaluation of the KRYPTOR Compact PLUS Analyzer-Based B.R.A.H.M.S. CgA â…¡ KRYPTOR Assay for Chromogranin A Measurement. Diagnostics, 2021, 11, 2400.	1.3	0
147	Applying Functional Assay Evidence to Interpret Sequence Variants Identified in Hereditary Cancer Genes. Laboratory Medicine Online, 2022, 12, 145-158.	0.0	0