Sug Hyung Lee

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genomic Progression of Precancerous Actinic Keratosis to Squamous Cell Carcinoma. Journal of Investigative Dermatology, 2022, 142, 528-538.e8.	0.7	20
2	Dissection of molecular and histological subtypes of papillary thyroid cancer using alternative splicing profiles. Experimental and Molecular Medicine, 2022, 54, 263-272.	7.7	3
3	Different Molecular Features of Epithelioid and Giant Cells in Foreign Body Reaction Identified by Single-Cell RNA Sequencing. Journal of Investigative Dermatology, 2022, 142, 3232-3242.e16.	0.7	11
4	Cancer-related SRCAP and TPR mutations in colon cancers. Pathology Research and Practice, 2021, 217, 153292.	2.3	7
5	Mutation and expression alterations of histone methylation-related NSD2, KDM2B and SETMAR genes in colon cancers. Pathology Research and Practice, 2021, 219, 153354.	2.3	6
6	Clinical implications of copy number alteration detection using panel-based next-generation sequencing data in myelodysplastic syndrome. Leukemia Research, 2021, 103, 106540.	0.8	1
7	Somatic Mutation of NLRP Genes in Gastric and Colonic Cancers. Pathology and Oncology Research, 2021, 27, 607385.	1.9	2
8	Brief Research Report Regional Difference in TRAF2 and TRAF3 Gene Mutations in Colon Cancers. Pathology and Oncology Research, 2021, 27, 625438.	1.9	3
9	Expression and Mutation Alterations of ZMYM4 Gene in Gastric and Colonic Cancers. Applied Immunohistochemistry and Molecular Morphology, 2021, 29, 570-575.	1.2	3
10	Molecular genetic evidence supporting diverse histogenic origins of germ cell tumors. Journal of Pathology, 2021, , .	4.5	2
11	Mutational Alterations of DNA Methylation-related Genes CTCF, ZFP57 and ATF7IP Genes in Colon Cancers. Applied Immunohistochemistry and Molecular Morphology, 2021, Publish Ahead of Print, e16-e20.	1.2	0
12	Inactivating Frameshift Mutations of HACD4 and TCP10L Tumor Suppressor Genes in Colorectal and Gastric Cancers. Pathology and Oncology Research, 2020, 26, 583-584.	1.9	0
13	Intratumoral Heterogeneity for Inactivating Frameshift Mutation of CYB5R2 Gene in Colorectal Cancers. Pathology and Oncology Research, 2020, 26, 585-586.	1.9	3
14	Analysis of Promoter Mutation in Long Non-coding RNA NEAT1 in Acute Leukemias. Pathology and Oncology Research, 2020, 26, 1345-1346.	1.9	0
15	Intratumoral Heterogeneity of RPL22 Frameshift Mutation in Colorectal Cancers. Pathology and Oncology Research, 2020, 26, 587-588.	1.9	0
16	Promoter Mutation Analysis of Long-Non-coding RNA RMRP Gene in Solid Tumors. Pathology and Oncology Research, 2020, 26, 2809-2810.	1.9	3
17	Intratumoral heterogeneity of FLCN somatic mutations in gastric and colorectal cancers. Pathology and Oncology Research, 2020, 26, 2811-2812.	1.9	0
18	Inactivating mutations of tumor suppressor genes KLOTHO and DTWD1 in colorectal cancers. Pathology Research and Practice, 2020, 216, 152816.	2.3	4

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19	Somatic mutation and loss of expression of a candidate tumor suppressor gene TET3 in gastric and colorectal cancers. Pathology Research and Practice, 2020, 216, 152759.	2.3	11
20	Mutation and Expression of a Candidate Tumor Suppressor Gene EPB41L3 in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2020, 26, 2003-2005.	1.9	2
21	Somatic Mutations and Intratumoral Heterogeneity of Cancer-Related Genes NLK, YY1 and PA2G4 in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2020, 26, 2813-2815.	1.9	6
22	Frameshift Mutations and Loss of Expression of CLCA4 Gene are Frequent in Colorectal Cancers With Microsatellite Instability. Applied Immunohistochemistry and Molecular Morphology, 2020, 28, 489-494.	1.2	5
23	Somatic mutations of cancer-related genes PELP1 and BDP1 in colorectal cancers. Pathology Research and Practice, 2020, 216, 153107.	2.3	5
24	Cancer-related gene mutations of ASPN in colon cancers. Pathology Research and Practice, 2020, 216, 153154.	2.3	1
25	Inactivating mutations of class II transactivator (CIITA) gene in gastric and colorectal cancers. Pathology Research and Practice, 2020, 216, 153110.	2.3	2
26	Distinct genomic profiles of gestational choriocarcinoma, a unique cancer of pregnant tissues. Experimental and Molecular Medicine, 2020, 52, 2046-2054.	7.7	12
27	Tight Junction-Related CLDN5 and CLDN6 Genes, and Gap Junction-Related GJB6 and GJB7 Genes Are Somatically Mutated in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2020, 26, 1983-1987.	1.9	12
28	Inactivating mutations of tumor suppressor genes ABCA1 and CAPN13 in colorectal cancers. Pathology Research and Practice, 2020, 216, 152870.	2.3	2
29	Mutational and expressional alterations of a candidate tumor suppressor HECA gene in gastric and colorectal cancers. Pathology Research and Practice, 2020, 216, 152896.	2.3	2
30	TGF-Î ² induced EMT and stemness characteristics are associated with epigenetic regulation in lung cancer. Scientific Reports, 2020, 10, 10597.	3.3	93
31	Downregulation of a putative tumor suppressor gene PHRF1 in gastric and colorectal cancers. Pathology Research and Practice, 2020, 216, 152984.	2.3	0
32	Differentially expressed genes in small intestine and colon adenocarcinomas identified by transcriptome sequencing. Pathology Research and Practice, 2020, 216, 152871.	2.3	0
33	Mutational alterations of TDRD 1, 4 and 9 genes in colorectal cancers. Pathology and Oncology Research, 2020, 26, 2007-2008.	1.9	5
34	Intratumoral heterogeneity of CSNK1G3 mutations, a casein kinase 1, in colon cancers. Pathology Research and Practice, 2020, 216, 152936.	2.3	1
35	Clinical Implications of Circulating Tumor DNA from Ascites and Serial Plasma in Ovarian Cancer. Cancer Research and Treatment, 2020, 52, 779-788.	3.0	24
36	Candidate Tumor Suppressor Gene EAF2 is Mutated in Colorectal and Gastric Cancers. Pathology and Oncology Research, 2019, 25, 823-824.	1.9	4

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37	Somatic frameshift mutations of cancer-related genes KIF3C and BARD1 in colorectal cancers. Pathology Research and Practice, 2019, 215, 152579.	2.3	4
38	Somatic mutations of candidate tumor suppressor genes folliculin-interacting proteins FNIP1 and FNIP2 in gastric and colon cancers. Pathology Research and Practice, 2019, 215, 152646.	2.3	2
39	Promoter mutation analysis of PMS2 gene in solid tumors and acute leukemias. Pathology Research and Practice, 2019, 215, 152583.	2.3	0
40	Analysis of driver somatic mutations in heterotopia of pancreas, spleen, liver and adrenal tissues. Pathology Research and Practice, 2019, 215, 152461.	2.3	0
41	Promoter Mutation Analysis of LEPROTL1 Gene in Acute Leukemias and Solid Tumors. Acta Haematologica, 2019, 141, 214-215.	1.4	0
42	Comparison of PANAMutyper and PNAClamp for Detecting KRAS Mutations from Patients With Malignant Pleural Effusion. In Vivo, 2019, 33, 945-954.	1.3	1
43	Integrative immunologic and genomic characterization of brain metastasis from ovarian/peritoneal cancer. Pathology Research and Practice, 2019, 215, 152404.	2.3	9
44	Hypoxia-induced cancer stemness acquisition is associated with CXCR4 activation by its aberrant promoter demethylation. BMC Cancer, 2019, 19, 148.	2.6	27
45	Comparison of PNA Clamping-assisted Fluorescence Melting Curve Analysis and PNA Clamping in Detecting <i>EGFR</i> Mutations in Matched Tumor Tissue, Cell Block, Pleural Effusion and Blood of Lung Cancer Patients With Malignant Pleural Effusion. In Vivo, 2019, 33, 595-603.	1.3	5
46	Somatic mutations in long-non-coding RNA RMRP in acute leukemias. Pathology Research and Practice, 2019, 215, 152647.	2.3	4
47	Promoter Mutation Analysis of ALDOA Gene in Solid Tumors and Acute Leukemias. Pathology and Oncology Research, 2019, 25, 825-826.	1.9	1
48	Frameshift mutation of candidate tumor suppressor genes QK1 and TMEFF2 in gastric and colorectal cancers. Cancer Biomarkers, 2019, 24, 1-6.	1.7	7
49	Absence of Promoter Mutation in TBC1D12 Gene in Solid and Hematologic Neoplasia. Pathology and Oncology Research, 2019, 25, 1675-1676.	1.9	0
50	Immune checkpoint blockade resistance-related B2M hotspot mutations in microsatellite-unstable colorectal carcinoma. Pathology Research and Practice, 2019, 215, 209-214.	2.3	28
51	Clinical Implications of Copy Number Variant Detection from Panel-Based Next-Generation Sequencing Data in Myelodysplastic Syndrome. Blood, 2019, 134, 4264-4264.	1.4	0
52	Expressional analysis of APLNR, an essential gene for cancer immunotherapy, in colon and prostate cancers. Pathology Research and Practice, 2018, 214, 599-600.	2.3	1
53	Absence of <i><scp>KRAS</scp></i> hotspot mutations in endometriosis of Korean patients. Histopathology, 2018, 73, 357-360.	2.9	3
54	Mutational signatures and chromosome alteration profiles of squamous cell carcinomas of the vulva. Experimental and Molecular Medicine, 2018, 50, e442-e442.	7.7	43

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55	Mutational intratumoral heterogeneity of a putative tumor suppressor gene RARRES3 in colorectal cancers. Pathology Research and Practice, 2018, 214, 601-602.	2.3	3
56	Rare frameshift mutations of putative tumor suppressor genes CSMD1 and SLX4 in colorectal cancers. Pathology Research and Practice, 2018, 214, 325-326.	2.3	6
57	Whole-exome sequencing identified mutational profiles of squamous cell carcinomas of anus. Human Pathology, 2018, 80, 1-10.	2.0	7
58	Whole-exome sequencing of chondroid hamartoma of lung identified no driver mutations. Pathology Research and Practice, 2018, 214, 459-462.	2.3	0
59	Low Frequent Mutation of ARHGAP35, a Candidate Tumor Suppressor Gene, in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2018, 24, 175-176.	1.9	4
60	TRIO Gene Encoding Trio Rho Guanine Nucleotide Exchange Factor Harbors Frameshift Mutations of in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2018, 24, 185-187.	1.9	0
61	Intratumoral Heterogeneity of Frameshift Mutations of GLI1 Encoding a Hedgehog Signaling Protein in Colorectal Cancers. Pathology and Oncology Research, 2018, 24, 477-481.	1.9	7
62	Intratumoral Heterogeneity of Somatic Mutations for NRIP1, DOK1, ULK1, ULK2, DLGAP3, PARD3 and PRKCI in Colon Cancers. Pathology and Oncology Research, 2018, 24, 827-832.	1.9	13
63	Frameshift Mutations in Repeat Sequences of ANK3, HACD4, TCP10L, TP53BP1, MFN1, LCMT2, RNMT, TRMT6, METTL8 and METTL16 Genes in Colon Cancers. Pathology and Oncology Research, 2018, 24, 617-622.	1.9	40
64	DAB2IP with tumor-inhibiting activities exhibits frameshift mutations in gastrointestinal cancers. Pathology Research and Practice, 2018, 214, 2075-2080.	2.3	2
65	TP53 mutation in allogeneic hematopoietic cell transplantation for de novo myelodysplastic syndrome. Leukemia Research, 2018, 74, 97-104.	0.8	9
66	Disparate genomic characteristics of concurrent endometrial adenocarcinoma and ovarian granulosa cell tumor, revealed by targeted next-generation sequencing. Pathology Research and Practice, 2018, 214, 1231-1233.	2.3	1
67	Clonal Structures of Regionally Synchronous Gastric Adenomas and Carcinomas. Clinical Cancer Research, 2018, 24, 4715-4725.	7.0	11
68	Targeted sequencing of burn scar-related squamous cell carcinomas identified PIK3CA amplification. Pathology, 2018, 50, 568-571.	0.6	1
69	Genomic structures of dysplastic nodule and concurrent hepatocellular carcinoma. Human Pathology, 2018, 81, 37-46.	2.0	6
70	Promoter mutation in long non-coding RNA NEAT1 is not common in common solid cancers. Pathology Research and Practice, 2018, 214, 1912-1913.	2.3	0
71	Intratumoral heterogeneity for frameshift mutations of TP53BP1 and MFN1 genes in colorectal cancers. Pathology Research and Practice, 2018, 214, 1514-1515.	2.3	0
72	Mutational analysis of H3F3B gene in acute leukaemias and solid tumours. Hematological Oncology, 2017, 35, 390-391.	1.7	0

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73	Inactivating frameshift mutation of putative tumor suppressor genes PLA2R1 and SRPK1 in gastric and colorectal cancers. Cancer Genetics, 2017, 210, 34-35.	0.4	6
74	WRN, the Werner Syndrome Gene, Exhibits Frameshift Mutations in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2017, 23, 451-452.	1.9	2
75	Molecular masquerading of rare EGFR L858M/L861R mutations as common L858R/L861Q mutations by PNA clamping assay. Pathology, 2017, 49, 453-455.	0.6	2
76	Mutational analysis of hypermutation-related POLE gene in acute leukemias and lymphomas. Experimental Hematology, 2017, 48, 39-40.	0.4	0
77	Frameshift Mutation of FXR1 Encoding a RNA-Binding Protein in Gastric and Colorectal Cancers with Microsatellite Instability. Pathology and Oncology Research, 2017, 23, 453-454.	1.9	2
78	USP9X, a Putative Tumor Suppressor Gene, Exhibits Frameshift Mutations in Colorectal Cancers. Pathology and Oncology Research, 2017, 23, 219-220.	1.9	4
79	Frameshift mutation and loss of expression of PLK2 , a serine/threonine kinase-encoding gene, in colorectal cancers. Pathology Research and Practice, 2017, 213, 1019-1020.	2.3	4
80	Histone Demethylase Gene PHF2 Is Mutated in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2017, 23, 471-476.	1.9	11
81	Intraindividual genomic heterogeneity of highâ€grade serous carcinoma of the ovary and clinical utility of ascitic cancer cells for mutation profiling. Journal of Pathology, 2017, 241, 57-66.	4.5	41
82	Candidate tumor suppressor gene MCPH1 is mutated in colorectal and gastric cancers. International Journal of Colorectal Disease, 2017, 32, 161-162.	2.2	5
83	Frameshift Mutations of SMG7 Essential for Nonsense-Mediated mRNA Decay in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2017, 23, 221-222.	1.9	4
84	Frameshift Mutations in the Mononucleotide Repeats of TAF1 and TAF1L Genes in Gastric and Colorectal Cancers with Regional Heterogeneity. Pathology and Oncology Research, 2017, 23, 125-130.	1.9	30
85	Intratumoral Heterogeneity of Frameshift Mutations in MECOM Gene is Frequent in Colorectal Cancers with High Microsatellite Instability. Pathology and Oncology Research, 2017, 23, 145-149.	1.9	17
86	Genomic profiles of a hepatoblastoma from a patient with Beckwith-Wiedemann syndrome with uniparental disomy on chromosome 11p15 and germline mutation of APC and PALB2. Oncotarget, 2017, 8, 91950-91957.	1.8	13
87	Intratumoral heterogeneity for inactivating frameshift mutation of CUX1 and SIRT1 genes in gastric and colorectal cancers. Polish Journal of Pathology, 2017, 68, 258-260.	0.3	3
88	Whole-exome sequencing identified mutational profiles of high-grade colon adenomas. Oncotarget, 2017, 8, 6579-6588.	1.8	27
89	Circulating Tumor DNA in a Breast Cancer Patient's Plasma Represents Driver Alterations in the Tumor Tissue. Genomics and Informatics, 2017, 15, 48.	0.8	7
90	<i>BPTF,</i> a chromatin remodelingâ€related gene, exhibits frameshift mutations in gastric and colorectal cancers. Apmis, 2016, 124, 425-427.	2.0	13

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91	ADNP encoding a transcription factor interacting with BAF complexes exhibits frameshift mutations in gastric and colorectal cancers. Scandinavian Journal of Gastroenterology, 2016, 51, 1269-1271.	1.5	2
92	Frameshift mutations of a tumor suppressor gene <i><scp>ZNF</scp>292</i> in gastric and colorectal cancers with high microsatellite instability. Apmis, 2016, 124, 556-560.	2.0	15
93	NSD1 encoding a histone methyltransferase exhibits frameshift mutations in colorectal cancers. Pathology, 2016, 48, 284-286.	0.6	6
94	Inactivating Frameshift Mutation of INPP4B Encoding a PI3K Pathway Phosphatase in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2016, 22, 653-654.	1.9	4
95	Frameshift Mutations of HSPA4 and MED13 in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2016, 22, 769-772.	1.9	11
96	Frameshift Mutation of ASPM Gene in Colorectal Cancers with Regional Heterogeneity. Pathology and Oncology Research, 2016, 22, 877-879.	1.9	5
97	Mutational and expressional alterations of ZMPSTE24, DNA damage response-related gene, in gastric and colorectal cancers. Pathology Research and Practice, 2016, 212, 1113-1118.	2.3	2
98	Frameshift Mutation of MED25, a Transcription Regulator, and its Mutational Heterogeneity in Colorectal Cancers. Pathology and Oncology Research, 2016, 22, 875-876.	1.9	0
99	Somatic mutation of a candidate tumour suppressor MGA gene and its mutational heterogeneity in colorectal cancers. Pathology, 2016, 48, 525-527.	0.6	17
100	Whole-exome sequencing identifies recurrent <i>AKT1</i> mutations in sclerosing hemangioma of lung. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 10672-10677.	7.1	42
101	Putative Tumor Suppressor Genes <i>EGR1 </i> and <i> BRSK1</i> Are Mutated in Gastric and Colorectal Cancers. Oncology, 2016, 91, 289-294.	1.9	15
102	Absence of PRKD1 Mutation, a Salivary Tumor-Specific Mutation, in Solid Tumors and Leukemias. Pathology and Oncology Research, 2016, 22, 231-232.	1.9	1
103	Whole-exome sequencing identified the genetic origin of a mucinous neoplasm in a mature cystic teratoma. Pathology, 2016, 48, 372-376.	0.6	8
104	Frameshift Mutations of AKAP9 Gene in Gastric and Colorectal Cancers with High Microsatellite Instability. Pathology and Oncology Research, 2016, 22, 587-592.	1.9	20
105	Inactivating frameshift mutation of <i>PBRM1</i> , a putative tumour suppressor gene, in colorectal cancers. Scandinavian Journal of Gastroenterology, 2016, 51, 639-640.	1.5	1
106	Elevated Coexpression of KITENIN and the ErbB4 CYT-2 Isoform Promotes the Transition from Colon Adenoma to Carcinoma Following <i>APC</i> loss. Clinical Cancer Research, 2016, 22, 1284-1294.	7.0	23
107	Frameshift mutation of WISP3 gene and its regional heterogeneity in gastric and colorectal cancers. Human Pathology, 2016, 50, 146-152.	2.0	9
108	Frameshift Mutations of CAB39L, an Activator of LKB1 Tumor Suppressor, in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2016, 22, 225-226.	1.9	2

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109	Leukemia Relapse-Associated Mutation of NT5C2 Gene is Rare in de Novo Acute Leukemias and Solid Tumors. Pathology and Oncology Research, 2016, 22, 223-224.	1.9	3
110	Genetic Progression of High Grade Prostatic Intraepithelial Neoplasia to Prostate Cancer. European Urology, 2016, 69, 823-830.	1.9	39
111	Absence of KNSTRN Mutation, a Cutaneous Squamous Carcinoma-Specific Mutation, in Other Solid Tumors and Leukemias Pathology and Oncology Research, 2016, 22, 227-228.	1.9	7
112	Somatic mutations predict outcomes of hypomethylating therapy in patients with myelodysplastic syndrome. Oncotarget, 2016, 7, 55264-55275.	1.8	62
113	Mutational burdens and evolutionary ages of thyroid follicular adenoma are comparable to those of follicular carcinoma. Oncotarget, 2016, 7, 69638-69648.	1.8	70
114	Predictive microRNAs for lymph node metastasis in endoscopically resectable submucosal colorectal cancer. Oncotarget, 2016, 7, 32902-32915.	1.8	20
115	Preferential occurrence of RHOA mutation in gastric and colorectal cancers. Pathology, 2015, 47, 598-599.	0.6	5
116	Genomic landscape of endometrial stromal sarcoma of uterus. Oncotarget, 2015, 6, 33319-33328.	1.8	26
117	Subclonal Genomic Architectures of Primary and Metastatic Colorectal Cancer Based on Intratumoral Genetic Heterogeneity. Clinical Cancer Research, 2015, 21, 4461-4472.	7.0	157
118	Frameshift Mutations of TAF7L Gene, a Core Component for Transcription by RNA Polymerase II, in Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 849-850.	1.9	5
119	Frameshift Mutations of MUC15 Gene in Gastric and its Regional Heterogeneity in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 713-718.	1.9	24
120	Mutation of HELLS, a Chromatin Remodeling Gene, Gastric and Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 851-852.	1.9	7
121	Regional Bias of Intratumoral Genetic Heterogeneity of Apoptosis-Related Genes BAX, APAF1, and FLASH in Colon Cancers with High Microsatellite Instability. Digestive Diseases and Sciences, 2015, 60, 1674-1679.	2.3	10
122	Frameshift mutations in mammalian target of rapamycin pathway genes and their regional heterogeneity in sporadic colorectal cancers. Human Pathology, 2015, 46, 753-760.	2.0	6
123	Mutational analysis of oncogenic CSF3R p.T618I in acute leukemias and common solid cancers. Annals of Hematology, 2015, 94, 889-890.	1.8	0
124	Frameshift Mutation of an Angiogenesis Factor VEGFB and its Mutational Heterogeneity in Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 853-855.	1.9	1
125	Mutation and Expression of a Methyl-Binding Protein 6 (MBD6) in Gastric and Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 857-858.	1.9	4
126	GNAS Mutation Affecting Codon 201 is Rare in Most Human Tumors. Pathology and Oncology Research, 2015, 21, 859-860.	1.9	6

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127	Frameshift mutations of TAF1C gene, a core component for transcription by RNA polymerase I, and its regional heterogeneity in gastric and colorectal cancers. Pathology, 2015, 47, 101-104.	0.6	10
128	Inactivating frameshift mutation of <i>AKT1S1,</i> an mTOR inhibitory gene, in colorectal cancers. Scandinavian Journal of Gastroenterology, 2015, 50, 503-504.	1.5	3
129	Novel oncogenic <scp><i>PTPN11</i></scp> mutations in myelodysplastic syndrome in Korean patients. Hematological Oncology, 2015, 33, 166-167.	1.7	2
130	Absence ofEGFR, ERBB2andERBB4mutation homologous to the oncogenicERBB3Val-104 mutation in colorectal cancers. Apmis, 2015, 123, 87-88.	2.0	0
131	Laminin gene <i><scp>LAMB</scp>4</i> is somatically mutated and expressionally altered in gastric and colorectal cancers. Apmis, 2015, 123, 65-71.	2.0	31
132	<scp>TEAD</scp> 2, a Hippo pathway gene, is somatically mutated in gastric and colorectal cancers with high microsatellite instability. Apmis, 2015, 123, 359-360.	2.0	3
133	Mutational Heterogeneity of MED23 Gene in Colorectal Cancers. Pathology and Oncology Research, 2015, 21, 1281-1282.	1.9	5
134	Frequent frameshift mutations in 2 mononucleotide repeats of RNF43 gene and its regional heterogeneity in gastric and colorectal cancers. Human Pathology, 2015, 46, 1640-1646.	2.0	24
135	Oncogenic PTPN11 Mutations are Rare in Solid Tumors. Pathology and Oncology Research, 2015, 21, 225-227.	1.9	2
136	Frameshift Mutations of Cadherin Genes DCHS2, CDH10 and CDH24 Genes in Gastric and Colorectal Cancers with High Microsatellite Instability. Pathology and Oncology Research, 2015, 21, 181-185.	1.9	31
137	HMCN1, a cell polarity-related gene, is somatically mutated in gastric and colorectal cancers. Pathology and Oncology Research, 2015, 21, 847-848.	1.9	22
138	Progression of naive intraepithelial neoplasia genome to aggressive squamous cell carcinoma genome of uterine cervix. Oncotarget, 2015, 6, 4385-4393.	1.8	16
139	Genomic differences between pure ductal carcinoma <i>in situ</i> and synchronous ductal carcinoma <i>in situ</i> with invasive breast cancer. Oncotarget, 2015, 6, 7597-7607.	1.8	67
140	Clonal origins and parallel evolution of regionally synchronous colorectal adenoma and carcinoma. Oncotarget, 2015, 6, 27725-27735.	1.8	31
141	Mutational and expressional analysis of ERBB3 gene in common solid cancers. Apmis, 2014, 122, 1207-1212.	2.0	11
142	Somatic mutations of amino acid metabolism-related genes in gastric and colorectal cancers and their regional heterogeneity - a short report. Cellular Oncology (Dordrecht), 2014, 37, 455-461.	4.4	14
143	Mutational analysis of <i>H3F3A</i> , a chromatin remodeling gene in common solid tumors. Apmis, 2014, 122, 81-82.	2.0	1
144	Mutations in exon 2 of <i>TBX3</i> are rare in common human tumors. Apmis, 2014, 122, 161-163.	2.0	0

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145	Mutational and expressional analysis ofSMC2gene in gastric and colorectal cancers with microsatellite instability. Apmis, 2014, 122, 499-504.	2.0	13
146	Somatic mutation ofSPOPtumor suppressor gene is rare in breast, lung, liver cancers, and acute leukemias. Apmis, 2014, 122, 164-166.	2.0	9
147	The mutational burdens and evolutionary ages of early gastric cancers are comparable to those of advanced gastric cancers. Journal of Pathology, 2014, 234, 365-374.	4.5	33
148	Frameshift mutation of a histone methylation-related gene SETD1B and its regional heterogeneity in gastric and colorectal cancers with high microsatellite instability. Human Pathology, 2014, 45, 1674-1681.	2.0	37
149	Expressional and Mutational Analysis of CREBBP Gene in Gastric and Colorectal Cancers with Microsatellite Instability. Pathology and Oncology Research, 2014, 20, 221-222.	1.9	2
150	<i>PAX5</i> somatic mutation is rare in multiple myelomas and nonâ€Hodgkin lymphomas of Korean patients. Hematological Oncology, 2014, 32, 110-111.	1.7	1
151	Down-regulation of ROBO2 Expression in Prostate Cancers. Pathology and Oncology Research, 2014, 20, 517-519.	1.9	14
152	OncogenicERBB3mutations altering p.Val104 is rare in acute leukemias and non-Hodgkin lymphomas. European Journal of Haematology, 2014, 92, 177-178.	2.2	2
153	Regional Bias of Intratumoral Genetic Heterogeneity of Nucleotide Repeats in Colon Cancers with Microsatellite Instability. Pathology and Oncology Research, 2014, 20, 965-971.	1.9	13
154	Nutlin-3 induces BCL2A1 expression by activating ELK1 through the mitochondrial p53-ROS-ERK1/2 pathway. International Journal of Oncology, 2014, 45, 675-682.	3.3	16
155	Frameshift mutations of PRKAG1 gene encoding an AMPK gamma subunit in colorectal cancers. Journal of Gastrointestinal and Liver Diseases, 2014, 23, 343-5.	0.9	1
156	Mutational analysis of <scp> <i>DNMT3A</i> </scp> gene in acute leukemias and common solid cancers. Apmis, 2013, 121, 85-94.	2.0	47
157	Somatic Mutation of PARK2 Tumor Suppressor Gene is not Common in Common Solid Cancers. Pathology and Oncology Research, 2013, 19, 393-395.	1.9	10
158	Frameshift mutation of a tumor suppressor gene <i>PALB2</i> in gastric and colorectal cancers with microsatellite instability. Apmis, 2013, 121, 1015-1016.	2.0	2
159	STAT3exon 21 mutation is rare in common human cancers. Acta OncolÃ ³ gica, 2013, 52, 1221-1222.	1.8	4
160	NIPBL, a Cohesion Loading Factor, Is Somatically Mutated in Gastric and Colorectal Cancers with High Microsatellite Instability. Digestive Diseases and Sciences, 2013, 58, 3376-3378.	2.3	10
161	Somatic mutation of IL7R exon 6 in acute leukemias and solid cancers. Human Pathology, 2013, 44, 551-555.	2.0	13
162	Mutational analysis of splicing machinery genes <scp><i>SF3B1, U2AF1</i></scp> and <scp><i>SRSF2</i></scp> in myelodysplasia and other common tumors. International Journal of Cancer, 2013, 133, 260-265.	5.1	64

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163	Frameshift mutations of a chromatinâ€remodeling gene <scp> <i>SMARCC2 </i> </scp> in gastric and colorectal cancers with microsatellite instability. Apmis, 2013, 121, 168-169.	2.0	17
164	Frameshift mutations of tumor suppressor gene EP300 in gastric and colorectal cancers with high microsatellite instability. Human Pathology, 2013, 44, 2064-2070.	2.0	23
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