Edmond S K

List of Publications by Year
in descending order

157 papers	6,461 citations	125106 35 h-index	87275 74 g-index
157 all docs	157 docs citations	157 times ranked	12209 citing authors

#	Article	IF	Citations
1	Safety and Immunogenicity of a Booster Vaccination by CoronaVac or BNT162b2 in Previously Two-Dose Inactivated Virus Vaccinated Individuals with Negative Neutralizing Antibody. Vaccines, 2022, 10, 556.	2.1	9
2	A Case Report of Germline Compound Heterozygous Mutations in the BRCA1 Gene of an Ovarian and Breast Cancer Patient. International Journal of Molecular Sciences, 2021, 22, 889.	1.8	5
3	Rapid Breakpoint Mapping of a Novel Germline PALB2 Duplication by PCR-Free Long-Read Sequencing for Interpretation of Its Pathogenicity. JCO Precision Oncology, 2021, 5, 1044-1047.	1.5	1
4	Assessment of droplet digital polymerase chain reaction for measuring BCRâ€ABL1 in chronic myeloid leukaemia in an international interlaboratory study. British Journal of Haematology, 2021, 194, 53-60.	1.2	10
5	Characteristics and predictors of early hospital deaths in newly diagnosed APL: a 13-year population-wide study. Blood Advances, 2021, 5, 2829-2838.	2.5	18
6	Germline PALB2 Mutation in High-Risk Chinese Breast and/or Ovarian Cancer Patients. Cancers, 2021, 13, 4195.	1.7	7
7	Association of Genomic Domains in BRCA1 and BRCA2 with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638.	0.4	39
8	Sorafenib and omacetaxine mepesuccinate as a safe and effective treatment for acute myeloid leukemia carrying internal tandem duplication of Fmsâ€like tyrosine kinase 3. Cancer, 2020, 126, 344-353.	2.0	19
9	An economical Nanopore sequencing assay for human papillomavirus (HPV) genotyping. Diagnostic Pathology, 2020, 15, 45.	0.9	6
10	Clofarabine, cytarabine, and mitoxantrone in refractory/relapsed acute myeloid leukemia: High response rates and effective bridge to allogeneic hematopoietic stem cell transplantation. Cancer Medicine, 2020, 9, 3371-3382.	1.3	7
11	Germline Mutation in 1338 BRCA-Negative Chinese Hereditary Breast and/or Ovarian Cancer Patients. Journal of Molecular Diagnostics, 2020, 22, 544-554.	1.2	17
12	Potential utility of targeted Nanopore sequencing for improving etiologic diagnosis of bacterial and fungal respiratory infection. Diagnostic Pathology, 2020, 15, 41.	0.9	20
13	Rapid detection of chromosomal translocation and precise breakpoint characterization in acute myeloid leukemia by nanopore long-read sequencing. Cancer Genetics, 2019, 239, 22-25.	0.2	24
14	Distinct mutation spectrum, clinical outcome and therapeutic responses of typical complex/monosomy karyotype acute myeloid leukemia carrying TP53 mutations. American Journal of Hematology, 2019, 94, 650-657.	2.0	30
15	Potential utility of metagenomic sequencing for improving etiologic diagnosis of infective endocarditis. Future Cardiology, 2019, 15, 411-424.	0.5	15
16	Next-generation sequencing panel for diagnosis and management of chronic neutrophilic leukaemia: a case report. Hong Kong Medical Journal, 2019, 25, 248-250.	0.1	1
17	Prospective study on human fecal carriage of Enterobacteriaceae possessing mcr-1 and mcr-2 genes in a regional hospital in Hong Kong. BMC Infectious Diseases, 2018, 18, 81.	1.3	28
18	Rapid detection of BRCA1/2 recurrent mutations in Chinese breast and ovarian cancer patients with multiplex SNaPshot genotyping panels. Oncotarget, 2018, 9, 7832-7843.	0.8	9

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19	Draft Genome Sequence of Helicobacter cinaedi, Compiled by Direct Whole-Genome Sequencing of a Blood Culture-Positive Isolate in Hong Kong. Microbiology Resource Announcements, 2018, 7, .	0.3	1
20	Breast and ovarian cancer penetrance of BRCA1/2 mutations among Hong Kong women. Oncotarget, 2018, 9, 25025-25033.	0.8	8
21	Frequent functional activation of RAS signalling not explained by RAS/RAF mutations in relapsed/refractory multiple myeloma. Scientific Reports, 2018, 8, 13522.	1.6	11
22	BAMClipper: removing primers from alignments to minimize false-negative mutations in amplicon next-generation sequencing. Scientific Reports, 2017, 7, 1567.	1.6	27
23	Recurrent Cytogenetic Abnormalities in Non-Hodgkinâ€™s Lymphoma and Chronic Lymphocytic Leukemia. Methods in Molecular Biology, 2017, 1541, 279-293.	0.4	12
24	Next-generation sequencing and molecular cytogenetic characterization of ETV6-LYN fusion due to chromosomes 1, 8 and 12 rearrangement in acute myeloid leukemia. Cancer Genetics, 2017, 218-219, 15-19.	0.2	14
25	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	13.7	1,099
26	INDELseek: detection of complex insertions and deletions from next-generation sequencing data. BMC Genomics, 2017, 18, 16.	1.2	19
27	Next-generation sequencing with a myeloid gene panel in core-binding factor AML showed KIT activation loop and TET2 mutations predictive of outcome. Blood Cancer Journal, 2016, 6, e442-e442.	2.8	29
28	Clinical evaluation of panel testing by next-generation sequencing (NGS) for gene mutations in myeloid neoplasms. Diagnostic Pathology, 2016, 11, 11.	0.9	77
29	Germline RECQL mutations in high risk Chinese breast cancer patients. Breast Cancer Research and Treatment, 2016, 157, 211-215.	1.1	23
30	Detection of Germline Mutation in Hereditary Breast and/or Ovarian Cancers by Next-Generation Sequencing on a Four-Gene Panel. Journal of Molecular Diagnostics, 2016, 18, 580-594.	1.2	38
31	Target fluorescence in-situ hybridization (Target FISH) for plasma cell enrichment in myeloma. Molecular Cytogenetics, 2016, 9, 63.	0.4	7
32	Homoharringtonine (omacetaxine mepesuccinate) as an adjunct for FLT3 - ITD acute myeloid leukemia. Science Translational Medicine, 2016, 8, 359ra129.	5.8	53
33	Comprehensive spectrum ofBRCA1andBRCA2deleterious mutations in breast cancer in Asian countries. Journal of Medical Genetics, 2016, 53, 15-23.	1.5	82
34	Evaluation of the new red cell parameters on Beckman Coulter DxH800 in distinguishing iron deficiency anaemia from thalassaemia trait. International Journal of Laboratory Hematology, 2015, 37, 199-207.	0.7	20
35	Emergency internal carotid artery stenting with platelet glycoprotein IIbIIIa antagonist coverage in a patient with essential thrombocytosis. Annals of Hematology, 2015, 94, 497-499.	0.8	1
36	Alpha thalassemia trait masquerading as hemoglobin H disease due to co-existing primary myelofibrosis. Annals of Hematology, 2015, 94, 875-877.	0.8	1

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37	Complete response of acquired FV inhibitor to rituximab. International Journal of Hematology, 2015, 101, 421-422.	0.7	5
38	Plasma DNA tissue mapping by genome-wide methylation sequencing for noninvasive prenatal, cancer, and transplantation assessments. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E5503-12.	3.3	579
39	The importance of analysis of long-range rearrangement of BRCA1 and BRCA2 in genetic diagnosis of familial breast cancer. Cancer Genetics, 2015, 208, 448-454.	0.2	43
40	Elevation of methylated DNA in KILLIN/PTEN in the plasma of patients with thyroid and/or breast cancer. OncoTargets and Therapy, 2014, 7, 2085.	1.0	8
41	Asymptomatic bulky Hodgkin lymphoma over 8 years with elevated alpha fetoprotein. Leukemia and Lymphoma, 2014, 55, 207-208.	0.6	1
42	Intrauterine therapy in a fetus with congenital dyserythropoietic anaemia type I. Journal of Obstetrics and Gynaecology, 2014, 34, 352-353.	0.4	4
43	MicroRNA-143 is downregulated in breast cancer and regulates DNA methyltransferases 3A in breast cancer cells. Tumor Biology, 2014, 35, 2591-2598.	0.8	79
44	Novel Point Mutation of the Î±2-Globin Gene (HBA2) and a Rare 2.4â€‰kb Deletion of the Î±1-Globin Gene (HBA1), Identified in Two Chinese Patients with Hb H Disease. Hemoglobin, 2014, 38, 213-215.	0.4	5
45	Aberrant large tumor suppressor 2 (LATS2) gene expression correlates with EGFR mutation and survival in lung adenocarcinomas. Lung Cancer, 2014, 85, 282-292.	0.9	25
46	Sudden blindness due to bilateral central retinal artery occlusion in a patient on eltrombopag. Annals of Hematology, 2014, 93, 881-882.	0.8	5
47	Large granular lymphocytic leukemia of gammaâ€“delta T cells: cytogenetics and fluorescence in situ hybridization study. Annals of Hematology, 2014, 93, 1247-1250.	0.8	1
48	JAK2 V617F mutation positive primary myelofibrosis with concomitant t(9;11;22)(q34;p15;q11.2) but no BCR/ABL fusion. International Journal of Hematology, 2013, 97, 435-437.	0.7	1
49	Survival of >20 years in a myeloma patient with an unusual combination of t(14;16) and hyperdiploidy: A case report. Oncology Letters, 2013, 6, 1663-1664.	0.8	2
50	Evaluation of 2 Real-Time PCR Assays for In Vitro Diagnostic Use in the Rapid and Multiplex Detection of EGFR Gene Mutations in NSCLC. Diagnostic Molecular Pathology, 2013, 22, 138-143.	2.1	14
51	Circulating microRNAs as Specific Biomarkers for Breast Cancer Detection. PLoS ONE, 2013, 8, e53141.	1.1	212
52	Association of Exon 19 and 21 EGFR Mutation Patterns with Treatment Outcome after First-Line Tyrosine Kinase Inhibitor in Metastatic Nonâ€“Small-Cell Lung Cancer. Journal of Thoracic Oncology, 2013, 8, 1148-1155.	0.5	97
53	A spuriously â€˜normalâ€™ haemoglobin A1c result. Annals of Clinical Biochemistry, 2012, 49, 408-411.	0.8	2
54	Identification of BRCA1/2 Founder Mutations in Southern Chinese Breast Cancer Patients Using Gene Sequencing and High Resolution DNA Melting Analysis. PLoS ONE, 2012, 7, e43994.	1.1	93

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55	EGFR Gene Mutation Study in Cytology Specimens. Acta Cytologica, 2012, 56, 661-668.	0.7	17
56	Association Between <emph type="ital">BRCA1</emph> and <emph type="ital">BRCA2</emph> Mutations and Survival in Women With Invasive Epithelial Ovarian Cancer. JAMA - Journal of the American Medical Association, 2012, 307, 382.	3.8	546
57	Small cell lung cancer with an epidermal growth factor receptor mutation in primary gefitinib-resistant adenocarcinoma of the lung. Acta OncolÃ³gica, 2012, 51, 557-559.	0.8	13
58	Treatment outcome and prognostic factor analysis in transplant-eligible Chinese myeloma patients receiving bortezomib-based induction regimens including the staged approach, PAD or VTD. Journal of Hematology and Oncology, 2012, 5, 28.	6.9	13
59	Is HbA2 level a reliable diagnostic measurement for Î²-thalassemia trait in people with iron deficiency?. American Journal of Hematology, 2012, 87, 114-116.	2.0	26
60	A 3-bp deletion in the HBS1L-MYB intergenic region on chromosome 6q23 is associated with HbF expression. Blood, 2011, 117, 4935-4945.	0.6	116
61	A novel de novo BRCA1 mutation in a Chinese woman with early onset breast cancer. Familial Cancer, 2011, 10, 233-237.	0.9	21
62	Quantitative Analysis and Diagnostic Significance of Methylated SLC19A3 DNA in the Plasma of Breast and Gastric Cancer Patients. PLoS ONE, 2011, 6, e22233.	1.1	53
63	Nearâ€tetraploid acute myeloid leukaemia. British Journal of Haematology, 2011, 155, 285-285.	1.2	4
64	High-resolution melting analysis for rapid screening of BRCA2 founder mutations in Southern Chinese breast cancer patients. Breast Cancer Research and Treatment, 2010, 122, 605-607.	1.1	5
65	Synchronous Primary Lung Cancer and Epidermal Growth Factor Receptor Mutation. Annals of Thoracic Surgery, 2010, 90, e38-e39.	0.7	6
66	Establishment of the first World Health Organization International Genetic Reference Panel for quantitation of BCR-ABL mRNA. Blood, 2010, 116, e111-e117.	0.6	141
67	MicroRNAs as New Players for Diagnosis, Prognosis, and Therapeutic Targets in Breast Cancer. Journal of Oncology, 2009, 2009, 1-6.	0.6	29
68	A BRCA2 founder mutation and seven novel deleterious BRCA mutations in southern Chinese women with breast and ovarian cancer. Breast Cancer Research and Treatment, 2009, 117, 683-686.	1.1	40
69	Clinical and pathological characteristics of Chinese patients with BRCA related breast cancer. The HUGO Journal, 2009, 3, 63-76.	4.1	30
70	Application of tri-colour, dual fusion fluorescence in situ hybridization (FISH) system for the characterization of BCR-ABL1 fusion in chronic myelogenous leukaemia (CML) and residual disease monitoring. BMC Hematology, 2009, 9, 4.	2.6	1
71	Burkitt lymphoma presenting as posterior reversible encephalopathy syndrome secondary to hypercalcaemia. British Journal of Haematology, 2009, 146, 584-584.	1.2	11
72	Detection and characterisation of Â-globin gene cluster deletions in Chinese using multiplex ligation-dependent probe amplification. Journal of Clinical Pathology, 2009, 62, 1107-1111.	1.0	24

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73	Detection of KRAS mutations in colorectal cancer by high-resolution melting analysis. Journal of Clinical Pathology, 2009, 62, 886-891.	1.0	46
74	Characterization of the pathogenic mechanism of a novel BRCA2 variant in a Chinese family. Familial Cancer, 2008, 7, 125-133.	0.9	15
75	Inherited thrombophilic factors do not increase central venous catheter blockage in children with malignancy. Pediatric Blood and Cancer, 2008, 51, 509-512.	0.8	10
76	Variation and heritability of Hb F and Fâ€cells among Î²â€thalassemia heterozygotes in Hong Kong. American Journal of Hematology, 2008, 83, 458-464.	2.0	30
77	Clinicopathological features of unbalanced translocation Der(1;7)(q10;p10) in myeloid neoplasms. Leukemia Research, 2008, 32, 1000-1001.	0.4	5
78	The HBS1L-MYB intergenic region on chromosome 6q23 is a quantitative trait locus controlling fetal haemoglobin level in carriers of Â-thalassaemia. Journal of Medical Genetics, 2008, 45, 745-751.	1.5	42
79	BCL11A is a major HbF quantitative trait locus in three different populations with Î²-hemoglobinopathies. Blood Cells, Molecules, and Diseases, 2008, 41, 255-258.	0.6	158
80	DOUBLE HETEROZYGOSITY FOR Hb NEW YORK [Î²113 GTGâ†'GAG; VALâ†'GLU] AND Î²Â°-THALASSEMIA MUTATIONS MANIFESTS AS A THALASSEMIA TRAIT. Pediatric Hematology and Oncology, 2008, 25, 227-231.	0.3	5
81	An unbalanced translocation, der(17)t(1;17)(p13;p11.2), leads to heterozygous loss of TP53 and is associated with clinical evolution in myelodysplastic syndrome. British Journal of Biomedical Science, 2008, 65, 36-38.	1.2	2
82	HLA associations, microsatellite instability and epigenetic changes in thyroid lymphoma in Chinese. Leukemia and Lymphoma, 2007, 48, 531-534.	0.6	8
83	A Case of IgM Paraproteinemia in Which Serum Free Light Chain Values Were Within Reference Intervals. Clinical Chemistry, 2007, 53, 362-363.	1.5	2
84	Cough mixture abuse as a novel cause of folate deficiency: a prospective, community-based, controlled study. Haematologica, 2007, 92, 562-563.	1.7	7
85	Diagnostic utility of dual fusion PML/RARÎ± translocation DNA probe (D-FISH) in acute promyelocytic leukemia. Oncology Reports, 2007, , .	1.2	8
86	The study of sequence configuration and functional impact of the (AC)n(AT)xTy motif in human Î²-globin gene promoter. American Journal of Hematology, 2007, 82, 342-348.	2.0	5
87	Amplification, mutation and loss of heterozygosity of the EGFR gene in metastatic lung cancer. International Journal of Cancer, 2007, 120, 1828-1831.	2.3	9
88	JAK2 V617F due to a novel TG â†' CT mutation at nucleotides 1848â€"1849: diagnostic implication. Leukemia, 2007, 21, 1344-1346.	3.3	12
89	Diagnostic clues to megaloblastic anaemia without macrocytosis. International Journal of Laboratory Hematology, 2007, 29, 163-171.	0.7	27
90	A novel beta-delta globin gene fusion, anti-Lepore Hong Kong, leads to overexpression of delta globin chain and a mild thalassaemia intermedia phenotype when co-inherited with ?0-thalassaemia. British Journal of Haematology, 2007, 136, 158-162.	1.2	24

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91	Successful treatment of thrombotic microangiopathy after haematopoietic stem cell transplantation with rituximab. British Journal of Haematology, 2007, 137, 475-478.	1.2	91
92	Detection of FIP1L1-PDGFRA fusion by FISH. British Journal of Haematology, 2007, 138, 279-279.	1.2	3
93	Non-secretory plasma cell myeloma of the true non-producer type. British Journal of Haematology, 2007, 138, 561-561.	1.2	7
94	Combined factors V and VIII deficiency (F5F8D) in a Chinese family due to compound heterozygosity for nonsense mutations of the LMAN1 gene. British Journal of Haematology, 2007, 139, 509-511.	1.2	7
95	Molecular Diagnosis of a Case of Hb Phnom Penh [Î±117(GH5)Phe-I1e-Î±118(H1)Thr (Î±1)]. Hemoglobin, 2006, 30, 397-399.	0.4	7
96	Concomitant post-transplantation lymphoproliferative disease and therapy-related myelodysplastic syndrome after lung transplantation. Journal of Heart and Lung Transplantation, 2006, 25, 259-260.	0.3	1
97	CD4+/CD56+ hematologic malignancy with rearranged MLL gene. Human Pathology, 2006, 37, 247-249.	1.1	10
98	Preclinical evaluation of pharmacokinetics and safety of melatonin in propylene glycol for intravenous administration. Journal of Pineal Research, 2006, 41, 337-343.	3.4	54
99	Ethnic Differences in Coagulation Factor Abnormalities After the Fontan Procedure. Pediatric Cardiology, 2006, 27, 96-101.	0.6	12
100	Therapy-related lymphomas in patients with autoimmune diseases after treatment with disease-modifying anti-rheumatic drugs. American Journal of Hematology, 2006, 81, 5-11.	2.0	40
101	Glucose-6-Phosphate Dehydrogenase Deficiency in Female Octogenarians, Nanogenarians, and Centenarians. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2006, 61, 1086-1089.	1.7	31
102	Pernicious Anemia in Chinese. Medicine (United States), 2006, 85, 129-138.	0.4	34
103	Lymphomatous proptosis as a novel feature of mantle cell lymphoma. Leukemia and Lymphoma, 2006, 47, 71-75.	0.6	13
104	Diagnostic Pitfall in PCR-Based Î±-Thalassemia Genotyping Resulting from a (Gâ†'C) Polymorphism at Nucleotide 71 3â€² to the Î±2-Globin Gene Termination Codon. Clinical Chemistry, 2006, 52, 536-537.	1.5	3
105	Ascaris-induced eosinophilic pneumonitis in an HIV-infected patient. Journal of Clinical Pathology, 2006, 60, 202-203.	1.0	8
106	A laboratory strategy for genotyping haemoglobin H disease in the Chinese. Journal of Clinical Pathology, 2006, 60, 931-934.	1.0	24
107	Dutcher-Fahey intranuclear inclusions in multiple myeloma. British Journal of Haematology, 2005, 129, 164-164.	1.2	2
108	Disseminated Penicillium marneffei infection. British Journal of Haematology, 2005, 130, 2-2.	1.2	4

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109	Serial studies of methylation of CDKN2B and CDKN2A in relapsed acute promyelocytic leukaemia treated with arsenic trioxide. British Journal of Haematology, 2005, 131, 632-635.	1.2	6
110	Systemic Oxygen Saturation and Coagulation Factor Abnormalities Before and After the Fontan Procedure. American Journal of Cardiology, 2005, 96, 1571-1575.	0.7	26
111	Plasma Epstein-Barr virus (EBV) DNA: Role as a screening test for nasopharyngeal carcinoma (NPC)?. International Journal of Cancer, 2005, 117, 515-516.	2.3	14
112	Therapy-related acute myeloid leukemia after single-agent treatment with fludarabine for chronic lymphocytic leukemia. American Journal of Hematology, 2005, 79, 288-290.	2.0	35
113	Peripartum pulmonary embolism in a patient with concomitant hemoglobin newcastle, hemoglobin constant spring, and hereditary spherocytosis. American Journal of Hematology, 2005, 80, 166-166.	2.0	3
114	Concomitant zoster myelitis and cerebral leukemia relapse after stem cell transplantation. Annals of Hematology, 2005, 84, 59-60.	0.8	6
115	Life-threatening cryoglobulinemia in HCV-negative Southern Chinese and a novel association with structural aortic abnormalities. Annals of Hematology, 2005, 84, 95-98.	0.8	13
116	A patient with anophthalmia, hemifacial microsomia, and hemoglobin anti-Lepore. Annals of Hematology, 2005, 84, 623-624.	0.8	2
117	Use of midazolam and ketamine as sedation for children undergoing minor operative procedures. Supportive Care in Cancer, 2005, 13, 1001-1009.	1.0	27
118	Proliferating cell nuclear antigen (PCNA) as a proliferative marker during embryonic and adult zebrafish hematopoiesis. Histochemistry and Cell Biology, 2005, 124, 105-111.	0.8	63
119	Kocuria kristinae infection associated with acute cholecystitis. BMC Infectious Diseases, 2005, 5, 60.	1.3	67
120	High-grade lymphoma after azathioprine treatment for Vogt-Kaganayi-Harada syndrome. Leukemia and Lymphoma, 2005, 46, 289-292.	0.6	4
121	CNS Manifestations of Malignancies. Journal of Clinical Oncology, 2005, 23, 4229-4230.	0.8	3
122	Side Effects Related to Cancer Treatment. Journal of Clinical Oncology, 2005, 23, 8535-8536.	0.8	5
123	Plasma Cell Problems. Journal of Clinical Oncology, 2005, 23, 3140-3143.	0.8	6
124	Unusual Rearrangement of the Î±-Globin Gene Cluster Containing Both the â˜Î±3.7 and Î±Î±Î±anti-4.2 Crossover Junctions: Clinical Diagnostic Implications and Possible Mechanisms. Clinical Chemistry, 2005, 51, 2167-2170.	1.5	30
125	Increased Alpha 7 Nicotinic Acetylcholine Receptor Protein Levels in Alzheimerâ€™s Disease Patients. Dementia and Geriatric Cognitive Disorders, 2005, 19, 106-112.	0.7	32
126	Diagnostic cues for natural killer cell lymphoma: primary nodal presentation and the role of in situ hybridisation for Epstein-Barr virus encoded early small RNA in detecting occult bone marrow involvement. Journal of Clinical Pathology, 2005, 58, 443-445.	1.0	31

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127	Clinical phenotype of haemoglobin Q-H disease. Journal of Clinical Pathology, 2004, 57, 81-82.	1.0	19
128	Unusual Sites of Metastatic Malignancy. Journal of Clinical Oncology, 2004, 22, 5015-5016.	0.8	1
129	FLT-3 aberrations in acute promyelocytic leukaemia: clinicopathological associations and prognostic impact. British Journal of Haematology, 2004, 125, 463-469.	1.2	56
130	Prolonged disturbances of in vitro cytokine production in patients with severe acute respiratory syndrome (SARS) treated with ribavirin and steroids. Clinical and Experimental Immunology, 2004, 135, 467-473.	1.1	70
131	Late-onset pure red blood cell aplasia owing to delayed lymphoid engraftment complicating ABO-mismatched hematopoietic stem cell transplantation. Transfusion, 2004, 44, 946-947.	0.8	3
132	Low frequency of FLT3 gene internal tandem duplication and activating loop mutation in therapy-related acute myelocyticleukemia and myelodysplastic syndrome. Cancer Genetics and Cytogenetics, 2004, 149, 169-172.	1.0	11
133	Detection of increased middle cerebral artery peak systolic velocity in fetuses affected by hemoglobin H Quong Sze disease. Ultrasound in Obstetrics and Gynecology, 2004, 23, 525-526.	0.9	5
134	Cost-effectiveness of prenatal screening for thalassaemia in Hong Kong. Prenatal Diagnosis, 2004, 24, 899-907.	1.1	36
135	Magnetic resonance imaging of leptomeningeal lymphoma. British Journal of Haematology, 2004, 124, 566-566.	1.2	0
136	Thalassemia Intermedia Due to Coâ€inheritance of Î²0/Î²+â€Thalassemia and (â€“ â€“SEA)Î±â€Thalassemia/Hb Westmead [Î±122(H5)Hisâ€‰›â€‰Gln (Î±2)] in a Chinese Family. Hemoglobin, 2004, 28, 151-156.	0.4	9
137	Molecular Characterization of Hb Val de Marne [Î±133(H16)Serâ†'Arg; AGCâ†'AGA; (Î±2)] in a Chinese Family. Hemoglobin, 2004, 28, 213-216.	0.4	3
138	MS analysis of single-nucleotide differences in circulating nucleic acids: Application to noninvasive prenatal diagnosis. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 10762-10767.	3.3	193
139	Desmopressin does not decrease blood loss and transfusion requirements in patients undergoing hepatectomy. Canadian Journal of Anaesthesia, 2003, 50, 14-20.	0.7	73
140	Deletion 9q as the sole karyotypic abnormality in myelocytic disorders: a new case of myelodysplastic syndrome and its prognostic implications in acute myelocytic leukemia. Cancer Genetics and Cytogenetics, 2003, 145, 184-186.	1.0	7
141	Chromosomal aberrations of primary lung adenocarcinomas in nonsmokers. Cancer, 2003, 97, 1263-1270.	2.0	83
142	The spectrum of acute lymphoblastic leukemia with mature B-cell phenotype. Leukemia Research, 2003, 27, 231-234.	0.4	17
143	Prevalence and specificity of clinically significant red cell alloantibodies in Chinese women during pregnancy - a review of cases from 1997 to 2001. Transfusion Medicine, 2003, 13, 227-231.	0.5	58
144	Amyloid deposits in the bone marrow. British Journal of Haematology, 2003, 121, 679-679.	1.2	2

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145	Plasmablastic transformation of multiple myeloma. Human Pathology, 2003, 34, 710-714.	1.1	17
146	Single-Tube Multiplex-PCR Screen for Anti-3.7 and Anti-4.2 Î±-Globin Gene Triplications. Clinical Chemistry, 2003, 49, 1679-1682.	1.5	90
147	Multiple Minisequencing Screen for Seven Southeast Asian Nondeletional Î±-Thalassemia Mutations. Clinical Chemistry, 2003, 49, 800-803.	1.5	9
148	Unusual Sites of Involvement in Non-Hodgkinâ€™s Lymphoma. Journal of Clinical Oncology, 2002, 20, 4394-4395.	0.8	14
149	Establishment, characterization, karyotyping, and comparative genomic hybridization analysis of HKESC-2 and HKESC-3. Cancer Genetics and Cytogenetics, 2002, 135, 120-127.	1.0	54
150	Therapy-related myelodysplastic syndrome after eradication of acute promyelocytic leukemia: Cytogenetic and molecular features. Human Pathology, 2001, 32, 126-129.	1.1	28
151	Establishment and characterization of a new xenograft-derived human esophageal squamous cell carcinoma cell line SLMT-1 of Chinese origin. Cancer Genetics and Cytogenetics, 2001, 124, 36-41.	1.0	49
152	EVANS??? SYNDROME COMPLICATING CHRONIC GRAFT VERSUS HOST DISEASE AFTER CADAVERIC LIVER TRANSPLANTATION1. Transplantation, 2001, 72, 527-528.	0.5	33
153	Primitive small round cell tumour of the adrenal gland presenting with fever of unknown origin and t(12;22)(q13;q12) cytogenetic finding. Journal of Clinical Pathology, 2001, 54, 966-969.	1.0	3
154	Establishment and Characterization of HKESC-1, a New Cancer Cell Line from Human Esophageal Squamous Cell Carcinoma. Cancer Genetics and Cytogenetics, 2000, 118, 112-120.	1.0	62
155	Immunorestitution Disease Involving the Innate and Adaptive Response. Clinical Infectious Diseases, 2000, 30, 882-892.	2.9	221
156	CLARITHROMYCIN ATTENUATES THE INFLAMMATORY RESPONSE INDUCED BY SURGICAL TRAUMA IN A GUINEA PIG MODEL. Pharmacological Research, 1999, 39, 49-54.	3.1	18
157	Sibling HLA-Matched Cord Blood Transplant for p-Thalassemia. Journal of Pediatric Hematology/Oncology, 1998, 20, 477-481.	0.3	19