## **Daniel Sinnett**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Screening for distress in pediatric cancer survivors: A systematic comparison of one-step and two-step strategies to minimize detection errors. Journal of Psychosocial Oncology, 2022, 40, 441-456.	0.6	2
2	Genetic factors contributing to late adverse musculoskeletal effects in childhood acute lymphoblastic leukemia survivors. Pharmacogenomics Journal, 2022, 22, 19-24.	0.9	2
3	Detection of doxorubicin-induced cardiotoxicity using myocardial T1 and T2 relaxation times in childhood acute lymphoblastic leukemia survivors. International Journal of Cardiovascular Imaging, 2022, 38, 873-882.	0.7	3
4	Magnetoencephalography resting-state correlates of executive and language components of verbal fluency. Scientific Reports, 2022, 12, 476.	1.6	3
5	Identification of new ETV6 modulators through a high throughput functional screening. IScience, 2022, 25, 103858.	1.9	3
6	Early Nutritional Intervention to Promote Healthy Eating Habits in Pediatric Oncology: A Feasibility Study. Nutrients, 2022, 14, 1024.	1.7	10
7	Whole-transcriptome analysis in acute lymphoblastic leukemia: a report from the DFCI ALL Consortium Protocol 16-001. Blood Advances, 2022, 6, 1329-1341.	2.5	30
8	Cardiometabolic Health After Pediatric Cancer Treatment: Adolescents Are More Affected than Children. Nutrition and Cancer, 2022, 74, 3236-3252.	0.9	3
9	Distinct transcriptomic profile of small arteries of hypertensive patients with chronic kidney disease identified miR-338-3p targeting GPX3 and PTPRS. Journal of Hypertension, 2022, 40, 1394-1405.	0.3	2
10	Contributing Factors of Unmet Needs Among Young Adult Survivors of Childhood Acute Lymphoblastic Leukemia with Comorbidities. Journal of Adolescent and Young Adult Oncology, 2021, 10, 462-475.	0.7	4
11	Developing and validating equations to predict V˙O <sub>2</sub> peak from the 6MWT in Childhood ALL Survivors. Disability and Rehabilitation, 2021, 43, 2937-2944.	0.9	5
12	Maximal cardiopulmonary exercise testing in childhood acute lymphoblastic leukemia survivors exposed to chemotherapy. Supportive Care in Cancer, 2021, 29, 987-996.	1.0	12
13	French-language adaptation of the 16D and 17D Quality of Life measures and score description in two Canadian pediatric samples. Health Psychology and Behavioral Medicine, 2021, 9, 619-635.	0.8	4
14	Elaboration and refinement of a motivational communication training program for healthcare professionals in pediatric oncology: a feasibility and acceptability study. Health Psychology and Behavioral Medicine, 2021, 9, 220-238.	0.8	2
15	Chromosome 2 Fragment Substitutions in Dahl Salt-Sensitive Rats and RNA Sequencing Identified Enpep and Hs2st1 as Vascular Inflammatory Modulators. Hypertension, 2021, 77, 178-189.	1.3	3
16	Frontline Ethico-Legal Issues in Childhood Cancer Genetics Research. , 2021, , 387-414.		1
17	Heart rate response and chronotropic incompetence during cardiopulmonary exercise testing in childhood acute lymphoblastic leukemia survivors. Pediatric Hematology and Oncology, 2021, 38, 564-580.	0.3	4
18	Repurposing proscillaridin A in combination with decitabine against embryonal rhabdomyosarcoma RD cells. Cancer Chemotherapy and Pharmacology, 2021, 88, 845-856.	1.1	2

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19	Genetic susceptibility to acute graft versus host disease in pediatric patients undergoing HSCT. Bone Marrow Transplantation, 2021, 56, 2697-2704.	1.3	2
20	Pre-implantation alcohol exposure induces lasting sex-specific DNA methylation programming errors in the developing forebrain. Clinical Epigenetics, 2021, 13, 164.	1.8	11
21	Genetic factors in treatment-related cardiovascular complications in survivors of childhood acute lymphoblastic leukemia. Pharmacogenomics, 2021, 22, 885-901.	0.6	1
22	Abstract MP60: Down-regulated Mir-338-3p In Subcutaneous Small Arteries Of Hypertensive Patients With Chronic Kidney Disease Targets Protein Tyrosine Phosphatase Receptor Type S And Glutathione Peroxidase 3. Hypertension, 2021, 78, .	1.3	0
23	Human Leucocyte Antigen alleles associated with asparaginase hypersensitivity in childhood Acute Lymphoblastic Leukemia patients treated with Pegylated asparaginase within Dana Farber Cancer Institute treatment protocols. Leukemia Research, 2021, 109, 106650.	0.4	0
24	"Taking back control together― Definition of a new intervention designed to support parents confronted with childhood cancer. Cogent Medicine, 2021, 8, 1944476.	0.7	3
25	Predictors of Vertebral Deformity in Long-Term Survivors of Childhood Acute Lymphoblastic Leukemia: The PETALE Study. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 512-525.	1.8	6
26	Performance of the McGill Interactive Pediatric OncoGenetic Guidelines for Identifying Cancer Predisposition Syndromes. JAMA Oncology, 2021, 7, 1806.	3.4	22
27	The VIE study: feasibility of a physical activity intervention in a multidisciplinary program in children with cancer. Supportive Care in Cancer, 2020, 28, 2627-2636.	1.0	8
28	Cryptic recurrent <i>ACIN1</i> â€ <i>NUTM1</i> fusions in nonâ€ <i>KMT2A</i> â€rearranged infant acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2020, 59, 125-130.	1.5	16
29	Genetic Susceptibility to Hepatic Sinusoidal Obstruction Syndrome in Pediatric Patients Undergoing Hematopoietic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2020, 26, 920-927.	2.0	11
30	Physical Activity and Sedentary Behaviors in Childhood Acute Lymphoblastic Leukemia Survivors. Journal of Pediatric Hematology/Oncology, 2020, 42, 53-60.	0.3	16
31	Role of rs10406069 in miR-5196 in hyperdiploid childhood acute lymphoblastic leukemia. Epigenomics, 2020, 12, 1949-1955.	1.0	2
32	Genetic factors in anthracycline-induced cardiotoxicity in patients treated for pediatric cancer. Expert Opinion on Drug Metabolism and Toxicology, 2020, 16, 865-883.	1.5	10
33	Diet Quality Is Associated with Cardiometabolic Outcomes in Survivors of Childhood Leukemia. Nutrients, 2020, 12, 2137.	1.7	16
34	Biomarkers of cardiometabolic complications in survivors of childhood acute lymphoblastic leukemia. Scientific Reports, 2020, 10, 21507.	1.6	15
35	Single-cell analysis of childhood leukemia reveals a link between developmental states and ribosomal protein expression as a source of intra-individual heterogeneity. Scientific Reports, 2020, 10, 8079.	1.6	37
36	Inconsistencies between measures of cognitive dysfunction in childhood acute lymphoblastic leukemia survivors: Description and understanding. Psycho-Oncology, 2020, 29, 1201-1208.	1.0	3

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37	HLA alleles associated with asparaginase hypersensitivity in childhood ALL: a report from the DFCI Consortium. Pharmacogenomics, 2020, 21, 541-547.	0.6	9
38	An optimized workflow to improve reliability of detection of KIAA1549:BRAF fusions from RNA sequencing data. Acta Neuropathologica, 2020, 140, 237-239.	3.9	5
39	The effect of cardiorespiratory fitness and physical activity levels on cognitive functions in survivors of childhood acute lymphoblastic leukemia. Pediatric Hematology and Oncology, 2020, 37, 582-598.	0.3	8
40	Children's physical activity behavior following a supervised physical activity program in pediatric oncology. Journal of Cancer Research and Clinical Oncology, 2020, 146, 3037-3048.	1.2	5
41	Circulating let-7g-5p and miR-191-5p Are Independent Predictors of Chronic Kidney Disease in Hypertensive Patients. American Journal of Hypertension, 2020, 33, 505-513.	1.0	18
42	Mining Heterogeneous Associations from Pediatric Cancer Data by Relational Concept Analysis. , 2020, , .		0
43	Inherited genetic susceptibility to acute lymphoblastic leukemia in Down syndrome. Blood, 2019, 134, 1227-1237.	0.6	37
44	Heritable variation at the chromosome 21 gene ERG is associated with acute lymphoblastic leukemia risk in children with and without Down syndrome. Leukemia, 2019, 33, 2746-2751.	3.3	18
45	Visual shortâ€ŧerm memory activation patterns in adult survivors of childhood acute lymphoblastic leukemia. Cancer, 2019, 125, 3639-3648.	2.0	3
46	Childhood Acute Lymphoblastic Leukemia Survivors Have a Substantially Lower Cardiorespiratory Fitness Level Than Healthy Canadians Despite a Clinically Equivalent Level of Physical Activity. Journal of Adolescent and Young Adult Oncology, 2019, 8, 674-683.	0.7	21
47	Impact of DARC, GSDMA and CXCL2 polymorphisms on induction toxicity in children with acute lymphoblastic leukemia: A complementary study. Leukemia Research, 2019, 86, 106228.	0.4	3
48	Genes identified through genome-wide association studies of osteonecrosis in childhood acute lymphoblastic leukemia patients. Pharmacogenomics, 2019, 20, 1189-1197.	0.6	7
49	<p>Identification of genetic variants associated with skeletal muscle function deficit in childhood acute lymphoblastic leukemia survivors</p> . Pharmacogenomics and Personalized Medicine, 2019, Volume 12, 33-45.	0.4	2
50	Influence of genetic factors on long-term treatment related neurocognitive complications, and on anxiety and depression in survivors of childhood acute lymphoblastic leukemia: The Petale study. PLoS ONE, 2019, 14, e0217314.	1.1	14
51	Heart failure drug proscillaridin A targets MYC overexpressing leukemia through global loss of lysine acetylation. Journal of Experimental and Clinical Cancer Research, 2019, 38, 251.	3.5	27
52	A Bayesian multivariate latent t-regression model for assessing the association between corticosteroid and cranial radiation exposures and cardiometabolic complications in survivors of childhood acute lymphoblastic leukemia: a PETALE study. BMC Medical Research Methodology, 2019, 19, 100.	1.4	3
53	Identification of genetic association between cardiorespiratory fitness and the trainability genes in childhood acute lymphoblastic leukemia survivors. BMC Cancer, 2019, 19, 443.	1.1	9
54	Identification of a single-nucleotide polymorphism within CDH2 gene associated with bone morbidity in childhood acute lymphoblastic leukemia survivors. Pharmacogenomics, 2019, 20, 409-420.	0.6	8

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55	Molecular Profiling of Hard-to-Treat Childhood and Adolescent Cancers. JAMA Network Open, 2019, 2, e192906.	2.8	36
56	Functional Analysis of Promoter Variants in Genes Involved in Sex Steroid Action, DNA Repair and Cell Cycle Control. Genes, 2019, 10, 186.	1.0	6
57	Definition and improvement of the concept and tools of a psychosocial intervention program for parents in pediatric oncology: a mixed-methods feasibility study conducted with parents and healthcare professionals. Pilot and Feasibility Studies, 2019, 5, 20.	0.5	13
58	Altered proteome of high-density lipoproteins from paediatric acute lymphoblastic leukemia survivors. Scientific Reports, 2019, 9, 4268.	1.6	11
59	miR-431-5p Knockdown Protects Against Angiotensin Il–Induced Hypertension and Vascular Injury. Hypertension, 2019, 73, 1007-1017.	1.3	21
60	ls there a relationship between vitamin D nutritional status and metabolic syndrome in childhood acute lymphoblastic leukemia survivors? A PETALE study. Clinical Nutrition ESPEN, 2019, 31, 28-32.	0.5	1
61	Doxorubicin treatments induce significant changes on the cardiac autonomic nervous system in childhood acute lymphoblastic leukemia long-term survivors. Clinical Research in Cardiology, 2019, 108, 1000-1008.	1.5	37
62	Nutrition education and cooking workshops for families of children with cancer: a feasibility study. BMC Nutrition, 2019, 5, 52.	0.6	9
63	Exercise Prescription Based on a Six Minute Walk Test in Childhood Acute Lymphoblastic Leukemia Survivors. Medicine and Science in Sports and Exercise, 2019, 51, 985-985.	0.2	0
64	Prevention of Long-term Adverse Health Outcomes With Cardiorespiratory Fitness and Physical Activity in Childhood Acute Lymphoblastic Leukemia Survivors. Journal of Pediatric Hematology/Oncology, 2019, 41, e450-e458.	0.3	33
65	Dietary Intakes Are Associated with HDL-Cholesterol in Survivors of Childhood Acute Lymphoblastic Leukaemia. Nutrients, 2019, 11, 2977.	1.7	11
66	Trainability Genes Provide Answers To The Cardiorespiratory Fitness Deficit In Childhood Acute Lymphoblastic Leukemia Survivors Medicine and Science in Sports and Exercise, 2019, 51, 157-157.	0.2	0
67	Preventive Action Of Cardiorespiratory Fitness On Health Outcomes In Childhood Acute Lymphoblastic Leukemia Survivors. Medicine and Science in Sports and Exercise, 2019, 51, 428-429.	0.2	0
68	Recurrent somatic BRAF insertion (p.V504_R506dup): a tumor marker and a potential therapeutic target in pilocytic astrocytoma. Oncogene, 2019, 38, 2994-3002.	2.6	13
69	Vitamin D nutritional status and bone turnover markers in childhood acute lymphoblastic leukemia survivors: A PETALE study. Clinical Nutrition, 2019, 38, 912-919.	2.3	17
70	Influence of BCL2L11 polymorphism on osteonecrosis during treatment of childhood acute lymphoblastic leukemia. Pharmacogenomics Journal, 2019, 19, 33-41.	0.9	16
71	Cardiometabolic Profile after Pediatric Cancer Treatment: Insight into HDL Composition and Nutritional Intake. FASEB Journal, 2019, 33, .	0.2	0
72	Development and relative validation of a food frequency questionnaire for French-Canadian adolescent and young adult survivors of acute lymphoblastic leukemia. Nutrition Journal, 2018, 17, 45.	1.5	13

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73	Insight from mitochondrial functions and proteomics to understand cardiometabolic disorders in survivors of acute lymphoblastic leukemia. Metabolism: Clinical and Experimental, 2018, 85, 151-160.	1.5	12
74	Impact of genetic polymorphisms determining leukocyte/neutrophil count on chemotherapy toxicity. Pharmacogenomics Journal, 2018, 18, 270-274.	0.9	11
75	Could we use parent report as a valid proxy of child report on anxiety, depression, and distress? A systematic investigation of father–mother–child triads in children successfully treated for leukemia. Pediatric Blood and Cancer, 2018, 65, e26840.	0.8	27
76	A3964 Identification of chromosome 2 differentially expressed aortic genes linked to vascular inflammation using congenic rats fed a normal and high-salt diet. Journal of Hypertension, 2018, 36, e25.	0.3	0
77	A3952 Circulating miR-26a-5p and let-7g-5p are potential biomarkers of chronic kidney disease. Journal of Hypertension, 2018, 36, e138-e139.	0.3	0
78	A3978 miR-338–3p down-regulation was identified in small arteries of hypertensive patients with chronic kidney disease. Journal of Hypertension, 2018, 36, e139.	0.3	0
79	Very long intergenic non-coding RNA transcripts and expression profiles are associated to specific childhood acute lymphoblastic leukemia subtypes. PLoS ONE, 2018, 13, e0207250.	1.1	12
80	TRPV4 and KRAS and FGFR1 gain-of-function mutations drive giant cell lesions of the jaw. Nature Communications, 2018, 9, 4572.	5.8	58
81	Genome wide mapping of ETV6 binding sites in pre-B leukemic cells. Scientific Reports, 2018, 8, 15526.	1.6	9
82	Genetic risk factors for VIPN in childhood acute lymphoblastic leukemia patients identified using whole-exome sequencing. Pharmacogenomics, 2018, 19, 1181-1193.	0.6	27
83	Cancer-related effects on relationships, long-term psychological status and relationship satisfaction in couples whose child was treated for leukemia: A PETALE study. PLoS ONE, 2018, 13, e0203435.	1.1	10
84	Mutational dynamics of early and late relapsed childhood ALL: rapid clonal expansion and long-term dormancy. Blood Advances, 2018, 2, 177-188.	2.5	31
85	DIVERGT screening procedure predicts general cognitive functioning in adult longâ€ŧerm survivors of pediatric acute lymphoblastic leukemia: A PETALE study. Pediatric Blood and Cancer, 2018, 65, e27259.	0.8	14
86	Psychological risk in longâ€ŧerm survivors of childhood acute lymphoblastic leukemia and its association with functional health status: A PETALE cohort study. Pediatric Blood and Cancer, 2018, 65, e27356.	0.8	25
87	Trametinib for progressive pediatric low-grade gliomas. Journal of Neuro-Oncology, 2018, 140, 435-444.	1.4	75
88	Research- and Practice-Based Nutrition Education and Cooking Workshops in Pediatric Oncology: Protocol for Implementation and Development of Curriculum. JMIR Research Protocols, 2018, 7, e2.	0.5	7
89	Abstract 1381: Targeting histone acetyltransferases to reprogram high C-MYC expressing cancers. , 2018, , .		0
90	Abstract 222: Genome-wide association study of acute lymphoblastic leukemia in children with Down		0

syndrome. , 2018, , .

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91	A 2â€year dyadic longitudinal study of mothers' and fathers' marital adjustment when caring for a child with cancer. Psycho-Oncology, 2017, 26, 1660-1666.	1.0	16
92	Lipid and lipoprotein abnormalities in acute lymphoblastic leukemia survivors. Journal of Lipid Research, 2017, 58, 982-993.	2.0	49
93	KMT2E-ASNS: a novel relapse-specific fusion gene in early T-cell precursor acute lymphoblastic leukemia. Blood, 2017, 129, 1729-1732.	0.6	3
94	Nutriepigenomics and malnutrition. Epigenomics, 2017, 9, 893-917.	1.0	18
95	Novel therapy for childhood acute lymphoblastic leukemia. Expert Opinion on Pharmacotherapy, 2017, 18, 1081-1099.	0.9	24
96	The PETALE study: Late adverse effects and biomarkers in childhood acute lymphoblastic leukemia survivors. Pediatric Blood and Cancer, 2017, 64, e26361.	0.8	66
97	How to interpret high levels of distress when using the Distress Thermometer in the long-term follow-up clinic? A study with Acute Lymphoblastic Leukemia survivors. Pediatric Hematology and Oncology, 2017, 34, 131-135.	0.3	11
98	Tracking Silent Hypersensitivity Reactions to Asparaginase during Leukemia Therapy Using Single-Chip Indirect Plasmonic and Fluorescence Immunosensing. ACS Sensors, 2017, 2, 1761-1766.	4.0	2
99	A protective role of IL-37 in cancer: a new hope for cancer patients. Journal of Leukocyte Biology, 2017, 101, 395-406.	1.5	46
100	Cardiometabolic Risk Factors in Childhood, Adolescent and Young Adult Survivors of Acute Lymphoblastic Leukemia – A Petale Cohort. Scientific Reports, 2017, 7, 17684.	1.6	41
101	LncRNAs downregulated in childhood acute lymphoblastic leukemia modulate apoptosis, cell migration, and DNA damage response. Oncotarget, 2017, 8, 80645-80650.	0.8	28
102	Specific expression of novel long non-coding RNAs in high-hyperdiploid childhood acute lymphoblastic leukemia. PLoS ONE, 2017, 12, e0174124.	1.1	24
103	Genomic determinants of long-term cardiometabolic complications in childhood acute lymphoblastic leukemia survivors. BMC Cancer, 2017, 17, 751.	1.1	14
104	Spontaneous brain oscillations as neural fingerprints of working memory capacities: A resting-state MEG study. Cortex, 2017, 97, 109-124.	1.1	15
105	Visual short term memory related brain activity predicts mathematical abilities Neuropsychology, 2017, 31, 535-545.	1.0	2
106	Characterization of the microDNA through the response to chemotherapeutics in lymphoblastoid cell lines. PLoS ONE, 2017, 12, e0184365.	1.1	33
107	A childhood acute lymphoblastic leukemia-specific lncRNA implicated in prednisolone resistance, cell proliferation, and migration. Oncotarget, 2017, 8, 7477-7488.	0.8	36
108	Whole-exome sequencing identified genetic risk factors for asparaginase-related complications in childhood ALL patients. Oncotarget, 2017, 8, 43752-43767.	0.8	33

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109	Abstract 4885: Identification of actionable targets for refractory/relapsed childhood cancer leading to personalized targeted therapy (TRICEPS Study). , 2017, , .		0
110	Abstract 1375: Loss of C-MYC and chromatin acetylation induce epigenetic reprogramming in acute lymphoblastic leukemia. , 2017, , .		0
111	Abstract P159: MicroRNA Profiling in Small Resistance Arteries of Hypertensive Patients With or Without Chronic Kidney Disease. Hypertension, 2017, 70, .	1.3	Ο
112	Abstract P162: <i>In vivo</i> miR-431 Inhibition Protects Against Vascular Damage and Hypertension. Hypertension, 2017, 70, .	1.3	0
113	Abstract P158: MicroRNA Profiling in Peripheral Blood Mononuclear Cells From Hypertensive Patients With or Without Chronic Kidney Disease. Hypertension, 2017, 70, .	1.3	Ο
114	CLIC5: a novel ETV6 target gene in childhood acute lymphoblastic leukemia. Haematologica, 2016, 101, 1534-1543.	1.7	27
115	SNooPer: a machine learning-based method for somatic variant identification from low-pass next-generation sequencing. BMC Genomics, 2016, 17, 912.	1.2	50
116	Sa1985 Microbiota-Related Acute Phase Proteins Are Predictors of Cardiometabolic Complications in Survivors of Pediatric Leukemia. Gastroenterology, 2016, 150, S424.	0.6	0
117	Adverse neuropsychological effects associated with cumulative doses of corticosteroids to treat childhood acute lymphoblastic leukemia: A literature review. Critical Reviews in Oncology/Hematology, 2016, 107, 138-148.	2.0	5
118	Genome-wide repression of eRNA and target gene loci by the ETV6-RUNX1 fusion in acute leukemia. Genome Research, 2016, 26, 1468-1477.	2.4	31
119	DNA methylome analysis of acute lymphoblastic leukemia cells reveals stochastic <i>de novo</i> DNA methylation in CpG islands. Epigenomics, 2016, 8, 1367-1387.	1.0	19
120	MPS 01-01 A CONSERVED microRNA CLUSTER AS A POTENTIAL MASTER GENE EXPRESSION REGULATOR IN ANGIOTENSIN II-INDUCED VASCULAR DAMAGE. Journal of Hypertension, 2016, 34, e78.	0.3	0
121	Polymorphisms of ABCC5 and NOS3 genes influence doxorubicin cardiotoxicity in survivors of childhood acute lymphoblastic leukemia. Pharmacogenomics Journal, 2016, 16, 530-535.	0.9	81
122	Genomic characterization of pediatric T-cell acute lymphoblastic leukemia reveals novel recurrent driver mutations. Oncotarget, 2016, 7, 65485-65503.	0.8	54
123	Abstract A41: TRICEPS: A feasibility study of personalized targeted therapy in relapsed/refractory childhood cancers. , 2016, , .		Ο
124	Abstract 107: Mir-431 as a Potential Master Regulator in Angiotensin li-induced Vascular Injury. Hypertension, 2016, 68, .	1.3	0
125	A novel somatic mutation in ACD induces telomere lengthening and apoptosis resistance in leukemia cells. BMC Cancer, 2015, 15, 621.	1.1	13
126	Hepatocyte Nuclear Factor 4 Alpha Polymorphisms and the Metabolic Syndrome in French-Canadian Youth. PLoS ONE, 2015, 10, e0117238.	1.1	19

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127	Association between CEBPE Variant and Childhood Acute Leukemia Risk: Evidence from a Meta-Analysis of 22 Studies. PLoS ONE, 2015, 10, e0125657.	1.1	7
128	Statement of principles on the return of research results and incidental findings in paediatric research: a multi-site consultative process. Genome, 2015, 58, 541-548.	0.9	25
129	Polymorphisms of Asparaginase Pathway and Asparaginase-Related Complications in Children with Acute Lymphoblastic Leukemia. Clinical Cancer Research, 2015, 21, 329-334.	3.2	52
130	To disclose, or not to disclose? Context matters. European Journal of Human Genetics, 2015, 23, 279-284.	1.4	15
131	Whole-exome sequencing of a rare case of familial childhood acute lymphoblastic leukemia reveals putative predisposing mutations in Fanconi anemia genes. BMC Cancer, 2015, 15, 539.	1.1	30
132	Abstract 1629: Prednisone versus dexamethasone acute toxicity and cumulative doses variations in childhood acute lymphoblastic leukemia. , 2015, , .		1
133	Abstract P625: Non-coding Rna Regulation Of Gene Expression In Angiotensin li-induced Vascular Damage. Hypertension, 2015, 66, .	1.3	0
134	Contribution of Polymorphisms in IKZF1 Gene to Childhood Acute Leukemia: A Meta-Analysis of 33 Case-Control Studies. PLoS ONE, 2014, 9, e113748.	1.1	18
135	Polymorphisms of the vincristine pathway and response to treatment in children with childhood acute lymphoblastic leukemia. Pharmacogenomics, 2014, 15, 1105-1116.	0.6	75
136	Hoxa9 collaborates with E2Aâ€₽BX1 in mouse B cell leukemia in association with Flt3 activation and decrease of B cell gene expression. Developmental Dynamics, 2014, 243, 145-158.	0.8	12
137	Tissue Distribution and Regulation of the Small Sar1b GTPase in Mice. Cellular Physiology and Biochemistry, 2014, 33, 1815-1826.	1.1	9
138	Pharmacogenetic considerations for acute lymphoblastic leukemia therapies. Expert Opinion on Drug Metabolism and Toxicology, 2014, 10, 699-719.	1.5	14
139	Impact of promoter polymorphisms in key regulators of the intrinsic apoptosis pathway on the outcome of childhood acute lymphoblastic leukemia. Haematologica, 2014, 99, 314-321.	1.7	10
140	Physical Activity and Fitness in Long Term Leukemia Survivors. Medicine and Science in Sports and Exercise, 2014, 46, 661.	0.2	0
141	Joint genotype inference with germline and somatic mutations. BMC Bioinformatics, 2013, 14, S3.	1.2	4
142	Integration of High-Resolution Methylome and Transcriptome Analyses to Dissect Epigenomic Changes in Childhood Acute Lymphoblastic Leukemia. Cancer Research, 2013, 73, 4323-4336.	0.4	44
143	Genome-wide signatures of differential DNA methylation in pediatric acute lymphoblastic leukemia. Genome Biology, 2013, 14, r105.	13.9	314
144	The Childhood Leukemia International Consortium. Cancer Epidemiology, 2013, 37, 336-347.	0.8	89

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145	Rare allelic forms of <i>PRDM9</i> associated with childhood leukemogenesis. Genome Research, 2013, 23, 419-430.	2.4	45
146	Whole-Exome Sequencing Reveals a Rapid Change in the Frequency of Rare Functional Variants in a Founding Population of Humans. PLoS Genetics, 2013, 9, e1003815.	1.5	70
147	<i>Bim</i> Polymorphisms: Influence on Function and Response to Treatment in Children with Acute Lymphoblastic Leukemia. Clinical Cancer Research, 2013, 19, 5240-5249.	3.2	21
148	Role of NOS3 DNA Variants in Externalizing Behavioral Problems Observed in Childhood Leukemia Survivors. Journal of Pediatric Hematology/Oncology, 2013, 35, e157-e162.	0.3	9
149	Frequency of Chromosomally-Integrated Human Herpesvirus 6 in Children with Acute Lymphoblastic Leukemia. PLoS ONE, 2013, 8, e84322.	1.1	21
150	Polymorphism in multidrug resistance-associated protein gene 3 is associated with outcomes in childhood acute lymphoblastic leukemia. Pharmacogenomics Journal, 2012, 12, 386-394.	0.9	26
151	CD133 expression is associated with poor outcome in neuroblastoma via chemoresistance mediated by the AKT pathway. Histopathology, 2012, 60, 1144-1155.	1.6	52
152	Promoter polymorphisms in CHI3L1 are associated with asthma. Journal of Allergy and Clinical Immunology, 2012, 130, 533-535.	1.5	10
153	Identification of functional DNA variants in the constitutive promoter region of MDM2. Human Genomics, 2012, 6, 15.	1.4	15
154	Association between genetic variants in the HNF4A gene and childhood-onset Crohn's disease. Genes and Immunity, 2012, 13, 556-565.	2.2	44
155	Functional analysis of promoter variants in <i>KU70</i> and their role in cancer susceptibility. Genes Chromosomes and Cancer, 2012, 51, 1007-1013.	1.5	8
156	Interaction between genetic and epigenetic variation defines gene expression patterns at the asthma-associated locus 17q12-q21 in lymphoblastoid cell lines. Human Genetics, 2012, 131, 1161-1171.	1.8	55
157	Abstract 4335: The genomic landscape of childhood pre-B acute lymphoblastic leukemia. , 2012, , .		0
158	Abstract 2484: Whole-exome sequencing of a rare case of familial childhood acute lymphoblastic leukemia. , 2012, , .		0
159	Plants growing on contaminated and brownfield sites appropriate for use in Organisation for Economic Coâ€operation and Development terrestrial plant growth test. Environmental Toxicology and Chemistry, 2011, 30, 124-131.	2.2	6
160	Glypican-3 (GPC3). Atlas of Genetics and Cytogenetics in Oncology and Haematology, 2011, , .	0.1	0
161	ALG: Automated Genotype Calling of Luminex Assays. PLoS ONE, 2011, 6, e19368.	1.1	8
162	Novel associations between activating killer-cell immunoglobulin-like receptor genes and childhood leukemia. Blood, 2011, 118, 1323-1328.	0.6	63

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163	ATF5 polymorphisms influence ATF function and response to treatment in children with childhood acute lymphoblastic leukemia. Blood, 2011, 118, 5883-5890.	0.6	46
164	Rationale for an international consortium to study inherited genetic susceptibility to childhood acute lymphoblastic leukemia. Haematologica, 2011, 96, 1049-1054.	1.7	36
165	Pediatric Research and the Return of Individual Research Results. Journal of Law, Medicine and Ethics, 2011, 39, 593-604.	0.4	35
166	Genomic and genealogical investigation of the French Canadian founder population structure. Human Genetics, 2011, 129, 521-531.	1.8	69
167	Genome-wide detection and characterization of mating asymmetry in human populations. Genetic Epidemiology, 2011, 35, n/a-n/a.	0.6	0
168	Expression of Sar1b Enhances Chylomicron Assembly and Key Components of the Coat Protein Complex II System Driving Vesicle Budding. Arteriosclerosis, Thrombosis, and Vascular Biology, 2011, 31, 2692-2699.	1.1	45
169	Common variants of the BRCA1 wild-type allele modify the risk of breast cancer in BRCA1 mutation carriers. Human Molecular Genetics, 2011, 20, 4732-4747.	1.4	32
170	Replication analysis confirms the association of ARID5B with childhood B-cell acute lymphoblastic leukemia. Haematologica, 2010, 95, 1608-1611.	1.7	71
171	Food-chain transfer of zinc from contaminated Urtica dioica and Acer pseudoplatanus L. to the aphids Microlophium carnosum and Drepanosiphum platanoidis Schrank. Environmental Pollution, 2010, 158, 267-271.	3.7	4
172	Polymorphisms in glucocorticoid receptor gene and the outcome of childhood acute lymphoblastic leukemia (ALL). Leukemia Research, 2010, 34, 492-497.	0.4	27
173	Variation in CDKN2A at 9p21.3 influences childhood acute lymphoblastic leukemia risk. Nature Genetics, 2010, 42, 492-494.	9.4	248
174	The Peptidomimetic CXCR4 Antagonist TC14012 Recruits β-Arrestin to CXCR7. Journal of Biological Chemistry, 2010, 285, 37939-37943.	1.6	77
175	Detection of Fetomaternal Genotype Associations in Early-Onset Disorders: Evaluation of Different Methods and Their Application to Childhood Leukemia. Journal of Biomedicine and Biotechnology, 2010, 2010, 1-13.	3.0	8
176	Modification in Oxidative Stress, Inflammation, and Lipoprotein Assembly in Response to Hepatocyte Nuclear Factor 41± Knockdown in Intestinal Epithelial Cells. Journal of Biological Chemistry, 2010, 285, 40448-40460.	1.6	52
177	HNF4 alpha: A New Susceptibility Gene for Crohn's Disease?: 2010 Presidential Poster. American Journal of Gastroenterology, 2010, 105, S464.	0.2	0
178	No evidence for association between TGFB1 promoter SNPs and the risk of childhood pre-B acute lymphoblastic leukemia among French Canadians. Haematologica, 2009, 94, 1034-1035.	1.7	1
179	DNA Variants in Region for Noncoding Interfering Transcript of Dihydrofolate Reductase Gene and Outcome in Childhood Acute Lymphoblastic Leukemia. Clinical Cancer Research, 2009, 15, 6931-6938.	3.2	34
180	Population genomics in a disease targeted primary cell model. Genome Research, 2009, 19, 1942-1952.	2.4	89

#	Article	IF	CITATIONS
181	Genetic heterogeneity in regional populations of Quebec—Parental lineages in the Gaspe Peninsula. American Journal of Physical Anthropology, 2009, 139, 512-522.	2.1	20
182	Functional impact of sequence variation in the promoter region of <i>TGFB1</i> . International Journal of Cancer, 2009, 125, 1483-1489.	2.3	19
183	Localization, function and regulation of the two intestinal fatty acid-binding protein types. Histochemistry and Cell Biology, 2009, 132, 351-367.	0.8	67
184	Clobal patterns of cis variation in human cells revealed by high-density allelic expression analysis. Nature Genetics, 2009, 41, 1216-1222.	9.4	206
185	The TP53 Arg72Pro and MDM2 309G>T polymorphisms are not associated with breast cancer risk in BRCA1 and BRCA2 mutation carriers. British Journal of Cancer, 2009, 101, 1456-1460.	2.9	19
186	FOOD-CHAIN TRANSFER OF CADMIUM AND ZINC FROM CONTAMINATED URTICA DIOICA TO HELIX ASPERSA AND LUMBRICUS TERRESTRIS. Environmental Toxicology and Chemistry, 2009, 28, 1756.	2.2	6
187	Allele-Specific Chromatin Remodeling in the ZPBP2/CSDMB/ORMDL3 Locus Associated with the Risk of Asthma and Autoimmune Disease. American Journal of Human Genetics, 2009, 85, 377-393.	2.6	262
188	Comparative expression analysis reveals differences in the regulation of intestinal paraoxonase family members. International Journal of Biochemistry and Cell Biology, 2009, 41, 1628-1637.	1.2	35
189	Assessment of Health and Ecological Impact of Policy on a Legacy Contamination Case Study. Epidemiology, 2009, 20, S230.	1.2	0
190	Development of a Software Based Decision Support Platform for Assessing the Impacts of Urban Pollutants. Epidemiology, 2009, 20, S230-S231.	1.2	0
191	Response: MRP4 gene polymorphisms and treatment response in adult ALL. Blood, 2009, 114, 5401-5402.	0.6	6
192	Polymorphisms in multidrug resistance-associated protein gene 4 is associated with outcome in childhood acute lymphoblastic leukemia. Blood, 2009, 114, 1383-1386.	0.6	83
193	The protective role of HNF4alpha in an intestinal epithelial cell model and the exploration of its genetic polymorphisms in inflammatory bowel diseases. Inflammatory Bowel Diseases, 2009, 15, S52.	0.9	0
194	IL-10 and TNF-α promoter haplotypes are associated with childhood Crohn's disease location. World Journal of Gastroenterology, 2009, 15, 3776.	1.4	48
195	Anderson or chylomicron retention disease: Molecular impact of five mutations in the SAR1B gene on the structure and the functionality of Sar1b protein. Molecular Genetics and Metabolism, 2008, 93, 74-84.	0.5	77
196	Differential Allelic Expression in the Human Genome: A Robust Approach To Identify Genetic and Epigenetic Cis-Acting Mechanisms Regulating Gene Expression. PLoS Genetics, 2008, 4, e1000006.	1.5	199
197	Parental Effect of DNA (Cytosine-5) Methyltransferase 1 on Grandparental-Origin-Dependent Transmission Ratio Distortion in Mouse Crosses and Human Families. Genetics, 2008, 178, 35-45.	1.2	12
198	DNA variants in the dihydrofolate reductase gene and outcome in childhood ALL. Blood, 2008, 111, 3692-3700.	0.6	104

#	Article	IF	CITATIONS
199	Effect of Oxidative Stress on the Status of Adhesion Molecules, Nuclear Receptors and Cholesterol Flux in Endothelial Cells: Priming of Monocytes. Clinical Medicine Cardiology, 2008, 2, CMC.S708.	0.1	0
200	Connections between ETV6-Modulated Genes: Identification of Shared Features. Cancer Informatics, 2008, 6, CIN.S556.	0.9	2
201	Biological role, protein expression, subcellular localization, and oxidative stress response of paraoxonase 2 in the intestine of humans and rats. American Journal of Physiology - Renal Physiology, 2007, 293, G1252-G1261.	1.6	64
202	Promoter SNPs in G1/S checkpoint regulators and their impact on the susceptibility to childhood leukemia. Blood, 2007, 109, 683-692.	0.6	62
203	Intestinal cholesterol transport proteins: an update and beyond. Current Opinion in Lipidology, 2007, 18, 310-318.	1.2	114
204	Patterns of variation in DNA segments upstream of transcription start sites. Human Mutation, 2007, 28, 441-450.	1.1	4
205	Asymmetrical regulation of scavenger receptor class B type I by apical and basolateral stimuli using Caco-2 cells. Journal of Cellular Biochemistry, 2007, 100, 421-433.	1.2	11
206	Subcellular proteomics of cell differentiation: Quantitative analysis of the plasma membrane proteome of Caco-2 cells. Proteomics, 2007, 7, 2201-2215.	1.3	53
207	Identification of transcripts modulated by ETV6 expression. British Journal of Haematology, 2007, 136, 48-62.	1.2	16
208	In vivo footprinting analysis of the Glypican 3 (GPC3) promoter region in neuroblastoma cells. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 2007, 1769, 182-193.	2.4	6
209	Detection and characterization of DNA variants in the promoter regions of hundreds of human disease candidate genes. Genomics, 2006, 87, 704-710.	1.3	26
210	Mapping cis-acting regulatory variation in recombinant congenic strains. Physiological Genomics, 2006, 25, 294-302.	1.0	20
211	Challenges Identifying Genetic Determinants of Pediatric Cancers – the Childhood Leukemia Experience. Familial Cancer, 2006, 5, 35-47.	0.9	25
212	Gene expression profiles of normal proliferating and differentiating human intestinal epithelial cells: A comparison with the Caco-2 cell model. Journal of Cellular Biochemistry, 2006, 99, 1175-1186.	1.2	65
213	Localization and role of NPC1L1 in cholesterol absorption in human intestine. Journal of Lipid Research, 2006, 47, 2112-2120.	2.0	141
214	Paraoxonase 1, 2 and 3 DNA variants and susceptibility to childhood inflammatory bowel disease. Gut, 2006, 55, 1820-1821.	6.1	8
215	Evaluation of BRCA1 and BRCA2 mutation prevalence, risk prediction models and a multistep testing approach in French-Canadian families with high risk of breast and ovarian cancer. Journal of Medical Genetics, 2006, 44, 107-121.	1.5	72
216	Iron-ascorbic acid-induced oxidant stress and its quenching by paraoxonase 1 in HDL and the liver: Comparison between humans and rats. Journal of Cellular Biochemistry, 2005, 96, 404-411.	1.2	22

#	Article	IF	CITATIONS
217	Macrophage Scavenger Receptor 1 999C>T (R293X) Mutation and Risk of Prostate Cancer. Cancer Epidemiology Biomarkers and Prevention, 2005, 14, 397-402.	1.1	21
218	Functional promoter SNPs in cell cycle checkpoint genes. Human Molecular Genetics, 2005, 14, 2641-2648.	1.4	28
219	Role of MTHFR genetic polymorphisms in the susceptibility to childhood acute lymphoblastic leukemia. Blood, 2004, 103, 252-257.	0.6	193
220	Genetic Diversity Patterns in the SR-BI/II Locus Can Be Explained by a Recent Selective Sweep. Molecular Biology and Evolution, 2004, 21, 760-769.	3.5	13
221	Mutational and expression analysis of the chromosome 12p candidate tumor suppressor genes in pre-B acute lymphoblastic leukemia. Leukemia, 2004, 18, 1499-1504.	3.3	35
222	Methylation analysis of the glypican 3 gene in embryonal tumours. British Journal of Cancer, 2004, 90, 1606-1611.	2.9	19
223	Association of metabolic gene polymorphisms with tobacco consumption in healthy controls. International Journal of Cancer, 2004, 110, 266-270.	2.3	21
224	Polymorphisms in Genes Involved in the Corticosteroid Response and the Outcome of Childhood Acute Lymphoblastic Leukemia. Molecular Diagnosis and Therapy, 2004, 4, 331-341.	3.3	34
225	Association of Vitamin D Receptor Genetic Variants with Susceptibility to Asthma and Atopy. American Journal of Respiratory and Critical Care Medicine, 2004, 170, 967-973.	2.5	217
226	A survey of genetic and epigenetic variation affecting human gene expression. Physiological Genomics, 2004, 16, 184-193.	1.0	228
227	Variable continental distribution of polymorphisms in the coding regions of DNA-repair genes. Journal of Human Genetics, 2003, 48, 659-664.	1.1	6
228	Analysis of the conservation of synteny between Fugu and human chromosome 12. BMC Genomics, 2003, 4, 30.	1.2	6
229	Role of DNA mismatch repair genetic polymorphisms in the risk of childhood acute lymphoblastic leukaemia. British Journal of Haematology, 2003, 123, 45-48.	1.2	47
230	Isolation of Cosmid and BAC DNA from E. coli. , 2003, 235, 99-102.		6
231	Characterization of the Bcll Polymorphism in the Glucocorticoid Receptor Gene. Clinical Chemistry, 2003, 49, 1528-1531.	1.5	49
232	Prostate cancer susceptibility genes: lessons learned and challenges posed Endocrine-Related Cancer, 2003, 10, 225-259.	1.6	81
233	Cellular Aspects of Intestinal Lipoprotein Assembly in Psammomys Obesus: A Model of Insulin Resistance and Type 2 Diabetes. Diabetes, 2003, 52, 2539-2545.	0.3	73
234	Inflammatory reaction without endogenous antioxidant response in Caco-2 cells exposed to iron/ascorbate-mediated lipid peroxidation. American Journal of Physiology - Renal Physiology, 2003, 285, G898-G906.	1.6	48

#	Article	IF	CITATIONS
235	Pharmacogenetics of Childhood Acute Lymphoblastic Leukemia. Current Pharmacogenomics and Personalized Medicine: the International Journal for Expert Reviews in Pharmacogenomics, 2003, 1, 87-100.	0.3	2
236	Childhood Acute Lymphoblastic Leukemia Associated with Parental Alcohol Consumption and Polymorphisms of Carcinogen-Metabolizing Genes. Epidemiology, 2002, 13, 277-281.	1.2	73
237	Glutathione S-transferase P1 genetic polymorphisms and susceptibility to childhood acute lymphoblastic leukaemia. Pharmacogenetics and Genomics, 2002, 12, 655-658.	5.7	65
238	A detailed transcriptional map of the chromosome 12p12 tumour suppressor locus. European Journal of Human Genetics, 2002, 10, 62-71.	1.4	38
239	Parental Genotypes in the Risk of a Complex Disease. American Journal of Human Genetics, 2002, 71, 193-197.	2.6	28
240	Reply to Comments by Kraft and Wilson and by Weinberg and Mitchell on "Parental Genotypes in the Risk of a Complex Disease― American Journal of Human Genetics, 2002, 71, 1240-1242.	2.6	1
241	Analyses of bulky DNA adduct levels in human breast tissue and genetic polymorphisms of cytochromes P450 (CYPs), myeloperoxidase (MPO), quinone oxidoreductase (NQO1), and glutathione S-transferases (GSTs). Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2002, 516, 41-47.	0.9	41
242	GSTT1 and CYP2E1 polymorphisms and trihalomethanes in drinking water: effect on childhood leukemia Environmental Health Perspectives, 2002, 110, 591-593.	2.8	50
243	Role ofNQO1,MPO andCYP2E1 genetic polymorphisms in the susceptibility to childhood acute lymphoblastic leukemia. International Journal of Cancer, 2002, 97, 230-236.	2.3	137
244	Polymorphisms in genes encoding drugs and xenobiotic metabolizing enzymes, DNA repair enzymes, and response to treatment of childhood acute lymphoblastic leukemia. Clinical Cancer Research, 2002, 8, 802-10.	3.2	106
245	Genetic susceptibility to breast cancer in French-Canadians: Role of carcinogen-metabolizing enzymes and gene-environment interactions. International Journal of Cancer, 2001, 92, 220-225.	2.3	111
246	Comparative analysis of the ETV6 gene in vertebrate genomes from pufferfish to human. Oncogene, 2001, 20, 3437-3442.	2.6	10
247	Childhood Acute Lymphoblastic Leukemia: Genetic Determinants of Susceptibility and Disease Outcome. Reviews on Environmental Health, 2001, 16, 263-79.	1.1	30
248	Expression of glypican 3(GPC3) in embryonal tumors. International Journal of Cancer, 2000, 89, 418-422.	2.3	94
249	Parental smoking, CYP1A1 genetic polymorphisms and childhood leukemia (Québec, Canada). Cancer Causes and Control, 2000, 11, 547-553.	0.8	75
250	Risk of childhood leukemia associated with diagnostic irradiation and polymorphisms in DNA repair genes Environmental Health Perspectives, 2000, 108, 495-498.	2.8	111
251	Genetic Susceptibility to Childhood Acute Lymphoblastic Leukemia. Leukemia and Lymphoma, 2000, 38, 447-462.	0.6	97
252	Pharmacogenomics and metabolite measurement for 6-mercaptopurine therapy in inflammatory bowel disease. Gastroenterology, 2000, 118, 705-713.	0.6	989

#	Article	IF	CITATIONS
253	Expression of glypican 3 (GPC3) in embryonal tumors. , 2000, 89, 418.		2
254	Expression of glypican 3 GPC3 in embryonal tumors. International Journal of Cancer, 2000, 89, 418-422.	2.3	4
255	Genetic polymorphisms of N-acetyltransferases 1 and 2 and gene-gene interaction in the susceptibility to childhood acute lymphoblastic leukemia. Cancer Epidemiology Biomarkers and Prevention, 2000, 9, 557-62.	1.1	28
256	Expression of glypican 3 (GPC3) in embryonal tumors. International Journal of Cancer, 2000, 89, 418-22.	2.3	37
257	Susceptibility to Childhood Acute Lymphoblastic Leukemia: Influence of CYP1A1, CYP2D6, GSTM1, and GSTT1 Genetic Polymorphisms. Blood, 1999, 93, 1496-1501.	0.6	211
258	Childhood Acute Lymphoblastic Leukemia: Is There a Tumor Suppressor Gene in Chromosome 12p 12.3 ?. Leukemia and Lymphoma, 1999, 34, 231-239.	0.6	22
259	Frequent loss of heterozygosity at the DNA mismatch-repair loci hMLH1 and hMSH3 in sporadic breast cancer. British Journal of Cancer, 1999, 79, 1012-1017.	2.9	31
260	Physical mapping of the G-protein coupled receptor 19 (GPR19) in the chromosome 12p12.3 region frequently rearranged in cancer cells. Human Genetics, 1999, 105, 162-164.	1.8	1
261	Physical mapping of the G-protein coupled receptor 19 (GPR19) in the chromosome 12p12.3 region frequently rearranged in cancer cells. Human Genetics, 1999, 105, 162-164.	1.8	7
262	Rapid Detection of CYP1A1, CYP2D6, and NAT Variants by Multiplex Polymerase Chain Reaction and Allele-Specific Oligonucleotide Assay. Analytical Biochemistry, 1999, 275, 84-92.	1.1	68
263	Fine physical and transcript mapping of a 1.8 Mb region spanning the locus for childhood acute lymphoblastic leukemia on chromosome 12p12.3. Gene, 1999, 240, 297-305.	1.0	9
264	Preconceptional paternal exposure to pesticides and increased risk of childhood leukaemia. Lancet, The, 1999, 354, 1819.	6.3	41
265	DNA methylation of retinoic acid receptor β in breast cancer and possible therapeutic role of 5-aza-2'-deoxycytidine. Anti-Cancer Drugs, 1999, 10, 471-476.	0.7	64
266	Risk of Childhood Leukemia Associated with Exposure to Pesticides and with Gene Polymorphisms. Epidemiology, 1999, 10, 481-487.	1.2	187
267	Susceptibility to Childhood Acute Lymphoblastic Leukemia: Influence of CYP1A1, CYP2D6, GSTM1, and GSTT1 Genetic Polymorphisms. Blood, 1999, 93, 1496-1501.	0.6	8
268	Susceptibility to childhood acute lymphoblastic leukemia: influence of CYP1A1, CYP2D6, GSTM1, and GSTT1 genetic polymorphisms. Blood, 1999, 93, 1496-501.	0.6	64
269	Risk of childhood leukemia associated with exposure to pesticides and with gene polymorphisms. Epidemiology, 1999, 10, 481-7.	1.2	53
270	Genomic loci susceptible to replication errors in cancer cells. British Journal of Cancer, 1998, 78, 981-985.	2.9	15

#	Article	IF	CITATIONS
271	High resolution deletion mapping reveals frequent allelic losses at the DNA mismatch repair locihMLH1 andhMSH3 in non-small cell lung cancer. , 1998, 77, 173-180.		46
272	Monophyletic Origin of Alu Elements in Primates. Journal of Molecular Evolution, 1998, 47, 172-182.	0.8	36
273	Chromosomal assignment of loci susceptible to replication errors by radiation hybrid mapping. Mutation Research - Mutation Research Genomics, 1998, 382, 81-83.	1.2	2
274	Allelic losses and DNA methylation at DNA mismatch repair loci in sporadic colorectal cancer. Carcinogenesis, 1998, 19, 1925-1929.	1.3	16
275	Demethylation by 5-aza-2??-deoxycytidine of specific 5-methylcytosine sites in the promoter region of the retinoic acid receptor ?? gene in human colon carcinoma cells. Anti-Cancer Drugs, 1998, 9, 743-750.	0.7	89
276	Isolation of Stable Bacterial Artificial Chromosome DNA Using a Modified Alkaline Lysis Method. BioTechniques, 1998, 24, 752-754.	0.8	34
277	High resolution deletion mapping reveals frequent allelic losses at the DNA mismatch repair loci hMLH1 and hMSH3 in non-small cell lung cancer. , 1998, 77, 173.		2
278	Human γ-Aminobutyric Acid-Type A Receptor α5 Subunit Gene (GABRA5): Characterization and Structural Organization of the 5′ Flanking Region. Genomics, 1997, 42, 378-387.	1.3	24
279	Microsatellite instability in childhood T cell acute lymphoblastic leukemia. Leukemia, 1997, 11, 797-802.	3.3	26
280	Allelic loss in childhood acute lymphoblastic leukemia. Leukemia Research, 1997, 21, 817-823.	0.4	44
281	Frequent deletion of chromosome 12p12.3 in children with acute lymphoblastic leukaemia. British Journal of Haematology, 1997, 99, 107-114.	1.2	41
282	Identification of a putative DNA replication origin in the λ-aminobutyric acid receptor subunit β3 and α5 gene cluster on human chromosome 15q11-q13, a region associated with parental imprinting and allele-specific replication timing. Gene, 1996, 173, 171-177.	1.0	24
283	Insulin modulation of newly synthesized apolipoproteins B-100 and B-48 in human fetal intestine: Gene expression and mRNA editing are not involved. FEBS Letters, 1996, 393, 253-258.	1.3	60
284	Overall Informativity,OI, in DNA Polymorphisms Revealed by inter-AluPCR: Detection of Genomic Rearrangements. Genomics, 1996, 36, 388-398.	1.3	13
285	Detection of a mutator phenotype in cancer cells by inter-Alu polymerase chain reaction. Cancer Research, 1996, 56, 2733-7.	0.4	18
286	Expression of 2 variant forms of fibroblast growth factor receptor 1 in human breast. International Journal of Cancer, 1995, 64, 274-279.	2.3	41
287	The Human γ-Aminobutyric Acid Receptor Subunit β3 and α5 Gene Cluster in Chromosome 15q11-q13 Is Rich in Highly Polymorphic (CA)n Repeats. Genomics, 1994, 19, 157-160.	1.3	45
288	FISH ordering of reference markers and of the gene for the α5 subunit of the γ-aminobutyric acid receptor (GABRA5) within the Angelman and Prader–Willi syndrome chromosomal regions. Human Molecular Genetics, 1993, 2, 183-189.	1.4	76

#	Article	IF	CITATIONS
289	Determination of DNA replication kinetics in synchronized human cells using a PCR-based assay. Nucleic Acids Research, 1993, 21, 3227-3232.	6.5	13
290	High-resolution mapping of the gamma-aminobutyric acid receptor subunit beta 3 and alpha 5 gene cluster on chromosome 15q11-q13, and localization of breakpoints in two Angelman syndrome patients. American Journal of Human Genetics, 1993, 52, 1216-29.	2.6	44
291	Mapping of the Angelman and Prader-Willi syndromes. Progress in Clinical and Biological Research, 1993, 384, 225-34.	0.2	0
292	Dinucleotide repeat polymorphism at the GABAA receptor α5 (GABRA5) locus at chromosome 15q11-q13. Human Molecular Genetics, 1992, 1, 348-348.	1.4	32
293	Linkage mapping by simultaneous screening of multiple polymorphic loci using Alu oligonucleotide-directed PCR Proceedings of the National Academy of Sciences of the United States of America, 1992, 89, 8448-8451.	3.3	47
294	Alu RNA transcripts in human embryonal carcinoma cells. Journal of Molecular Biology, 1992, 226, 689-706.	2.0	96
295	Single-strand conformational polymorphisms (SSCP): detection of useful polymorphisms at the dystrophin locus. Human Genetics, 1992, 89, 453-6.	1.8	13
296	G proteins in normal rat pituitaries and in prolactin-secreting rat pituitary tumors. Molecular and Cellular Endocrinology, 1991, 78, 33-44.	1.6	22
297	Evolution of mouse B1 repeats: 7SL RNA folding pattern conserved. Journal of Molecular Evolution, 1991, 32, 405-414.	0.8	71
298	Alu RNA secondary structure consists of two independent 7 SL RNA-like folding units. Journal of Biological Chemistry, 1991, 266, 8675-8678.	1.6	104
299	Alu RNA secondary structure consists of two independent 7 SL RNA-like folding units. Journal of Biological Chemistry, 1991, 266, 8675-8.	1.6	91
300	Reverse transcriptase activity from human embryonal carcinoma cells NTera2D1 EMBO Journal, 1990, 9, 3363-3368.	3.5	85
301	Use of Î <sup>3</sup> irradiation to eliminate DNA contamination for PCR. Nucleic Acids Research, 1990, 18, 6149-6149.	6.5	41
302	Alumorphs—Human DNA polymorphisms detected by polymerase chain reaction using Alu-specific primers. Genomics, 1990, 7, 331-334.	1.3	60
303	Reverse transcriptase activity from human embryonal carcinoma cells NTera2D1. EMBO Journal, 1990, 9, 3363-8.	3.5	49
304	The gene for incontinentia pigmenti is assigned to Xq28. Genomics, 1989, 4, 427-429.	1.3	107
305	Linkage studies do not confirm the cytogenetic location of incontinentia pigmenti on Xp11. Human Genetics, 1988, 80, 282-286.	1.8	33
306	Lesch-Nyhan syndrome: molecular investigation of three French Canadian families using a hypoxanthine-guanine phosphoribosyltransferase cDNA probe. Human Genetics, 1988, 81, 4-8.	1.8	16

#	Article	IF	CITATIONS
307	PRENATAL DIAGNOSIS USING DNA PROBES IN TWINS AT RISK FOR DUCHENNE MUSCULAR DYSTROPHY. Lancet, The, 1986, 328, 216-217.	6.3	4