Nelson Leung-Sang Tang

List of Publications by Year in descending order

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287 papers

15,127 citations

55 h-index 25983 112 g-index

295 all docs 295 docs citations

295 times ranked 23395 citing authors

#	Article	IF	CITATIONS
1	Nonalbuminuric Diabetic Kidney Disease and Risk of All-Cause Mortality and Cardiovascular and Kidney Outcomes in Type 2 Diabetes: Findings From the Hong Kong Diabetes Biobank. American Journal of Kidney Diseases, 2022, 80, 196-206.e1.	2.1	12
2	Clinical Predictors and Long-term Impact of Acute Kidney Injury on Progression of Diabetic Kidney Disease in Chinese Patients With Type 2 Diabetes. Diabetes, 2022, 71, 520-529.	0.3	6
3	Removal of the Prehospital Tourniquet in the Emergency Department. Journal of Emergency Medicine, 2021, 60, 98-102.	0.3	5
4	Development of genome-wide polygenic risk scores for lipid traits and clinical applications for dyslipidemia, subclinical atherosclerosis, and diabetes cardiovascular complications among East Asians. Genome Medicine, 2021, 13, 29.	3.6	18
5	Natural Selection of ATP2B1 Underlies Susceptibility to Essential Hypertension. Frontiers in Genetics, 2021, 12, 628516.	1.1	0
6	Promoter Methylation and Gene Expression of Pin1 Associated with the Risk of Alzheimer's Disease in Southern Chinese. Current Alzheimer Research, 2021, 17, 1232-1237.	0.7	4
7	Single-cell RNA sequencing identifies molecular targets associated with poor <i>in vitro</i> maturation performance of oocytes collected from ovarian stimulation. Human Reproduction, 2021, 36, 1907-1921.	0.4	10
8	Direct Measurement of B Lymphocyte Gene Expression Biomarkers in Peripheral Blood Transcriptomics Enables Early Prediction of Vaccine Seroconversion. Genes, 2021, 12, 971.	1.0	3
9	Peripheral Blood T Cell Gene Expression Responses to Exercise and HMB in Sarcopenia. Nutrients, 2021, 13, 2313.	1.7	9
10	A Decade in Review after Idiopathic Scoliosis Was First Called a Complex Trait—A Tribute to the Late Dr. Yves Cotrel for His Support in Studies of Etiology of Scoliosis. Genes, 2021, 12, 1033.	1.0	6
11	Differential Effects of Estrogen Receptor Alpha and Beta on Endogenous Ligands of Peroxisome Proliferator-Activated Receptor Gamma in Papillary Thyroid Cancer. Frontiers in Endocrinology, 2021, 12, 708248.	1.5	10
12	A Functional SNP in the Promoter of LBX1 Is Associated With the Development of Adolescent Idiopathic Scoliosis Through Involvement in the Myogenesis of Paraspinal Muscles. Frontiers in Cell and Developmental Biology, 2021, 9, 777890.	1.8	9
13	Biochemical Studies in Fibroblasts to Interpret Variants of Unknown Significance in the ABCD1 Gene. Genes, 2021, 12, 1930.	1.0	6
14	Dietary patterns and telomere length in community-dwelling Chinese older men and women: a cross-sectional analysis. European Journal of Nutrition, 2020, 59, 3303-3311.	1.8	8
15	Ambient Temperature is A Strong Selective Factor Influencing Human Development and Immunity. Genomics, Proteomics and Bioinformatics, 2020, 18, 489-500.	3.0	5
16	Genes Regulate Blood Pressure, but "Environments―Cause Hypertension. Frontiers in Genetics, 2020, 11, 580443.	1.1	10
17	Genome-Wide Search for SNP Interactions in GWAS Data: Algorithm, Feasibility, Replication Using Schizophrenia Datasets. Frontiers in Genetics, 2020, 11, 1003.	1.1	14
18	A randomized study of olanzapine-containing versus standard antiemetic regimens for the prevention of chemotherapy-induced nausea and vomiting in Chinese breast cancer patients. Breast, 2020, 50, 30-38.	0.9	19

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19	Reference intervals of spot urine copper excretion in preschool children and potential application in pre-symptomatic screening of Wilson disease. Pathology, 2020, 52, 439-446.	0.3	5
20	NEPA efficacy and tolerability during (neo)adjuvant breast cancer chemotherapy with cyclophosphamide and doxorubicin. BMJ Supportive and Palliative Care, 2020, , bmjspcare-2019-002037.	0.8	5
21	Status of inflammation in relation to health related quality of life in hepatocellular carcinoma patients. Quality of Life Research, 2019, 28, 2597-2607.	1.5	4
22	Effect of CYP2D6 and CYP3A4 Genotypes on the Efficacy of Cholinesterase Inhibitors in Southern Chinese Patients With Alzheimer's Disease. American Journal of Alzheimer's Disease and Other Dementias, 2019, 34, 302-307.	0.9	13
23	The association of liver function and quality of life of patients with liver cancer. BMC Gastroenterology, 2019, 19, 66.	0.8	21
24	GWAS of bone size yields twelve loci that also affect height, BMD, osteoarthritis or fractures. Nature Communications, 2019, 10, 2054.	5.8	74
25	Correlations of health-related quality of life with serum inflammatory indicators IL-8 and mIBI in patients with hepatocellular carcinoma. Cancer Management and Research, 2019, Volume 11, 2719-2727.	0.9	8
26	Revealing cellular and molecular transitions in neonatal germ cell differentiation using Single-cell RNA sequencing. Development (Cambridge), 2019, 146, .	1.2	20
27	Role of protein structure in variant annotation: structural insight of mutations causing 6-pyruvoyl-tetrahydropterin synthase deficiency. Pathology, 2019, 51, 274-280.	0.3	7
28	New Evidence Supporting the Role of FBN1 in the Development of Adolescent Idiopathic Scoliosis. Spine, 2019, 44, E225-E232.	1.0	10
29	Replication Study for the Association of GWAS-associated Loci With Adolescent Idiopathic Scoliosis Susceptibility and Curve Progression in a Chinese Population. Spine, 2019, 44, 464-471.	1.0	19
30	Genetic susceptibility to Tuberculosis: Interaction between HLA-DQA1 and age of onset. Infection, Genetics and Evolution, 2019, 68, 98-104.	1.0	7
31	Comparative analysis of single-cell parallel sequencing approaches in oocyte application. International Journal of Biochemistry and Cell Biology, 2019, 107, 1-5.	1.2	9
32	Progression of diabetic kidney disease and trajectory of kidney function decline in Chinese patients with Type 2 diabetes. Kidney International, 2019, 95, 178-187.	2.6	105
33	A Genetic Predictive Model Estimating the Risk of Developing Adolescent Idiopathic Scoliosis. Current Genomics, 2019, 20, 246-251.	0.7	14
34	A network approach to exploring the functional basis of gene–gene epistatic interactions in disease susceptibility. Bioinformatics, 2018, 34, 1741-1749.	1.8	11
35	Genetic Variant of PAX1 Gene Is Functionally Associated With Adolescent Idiopathic Scoliosis in the Chinese Population. Spine, 2018, 43, 492-496.	1.0	15
36	Transcriptome evidence reveals enhanced autophagy-lysosomal function in centenarians. Genome Research, 2018, 28, 1601-1610.	2.4	36

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37	Discovering Genetic Factors for psoriasis through exhaustively searching for significant second order SNP-SNP interactions. Scientific Reports, 2018, 8, 15186.	1.6	16
38	Predicting Mouse Oocyte Methylome from Polar Body by Single ell Whole Genome Bisulfite Sequencing. FASEB Journal, 2018, 32, 818.12.	0.2	O
39	Prognostic values of EORTC QLQ-C30 and QLQ-HCC18 index-scores in patients with hepatocellular carcinoma – clinical application of health-related quality-of-life data. BMC Cancer, 2017, 17, 8.	1.1	38
40	The relationship between angiotensin-converting enzyme (ACE) insertion (I) / deletion (D) polymorphism, serum ACE activity and bone mineral density (BMD) in older Chinese. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2017, 18, 147032031668834.	1.0	1
41	VCNet: vector-based gene co-expression network construction and its application to RNA-seq data. Bioinformatics, 2017, 33, 2173-2181.	1.8	9
42	Genome-wide association study identifies novel susceptible loci and highlights Wnt/beta-catenin pathway in the development of adolescent idiopathic scoliosis. Human Molecular Genetics, 2017, 26, 1577-1583.	1.4	59
43	Genetic variant of BNC2 gene is functionally associated with adolescent idiopathic scoliosis in Chinese population. Molecular Genetics and Genomics, 2017, 292, 789-794.	1.0	12
44	Whole-genome sequencing identifies rare genotypes in COMP and CHADL associated with high risk of hip osteoarthritis. Nature Genetics, 2017, 49, 801-805.	9.4	75
45	Whole genome sequencing finds rare high-risk genotypes for hip osteoarthritis in the COMP and CHADL genes. Osteoarthritis and Cartilage, 2017, 25, S37-S38.	0.6	1
46	<scp>BRCA</scp> 1 and <scp>BRCA</scp> 2 tumor suppressors protect against endogenous acetaldehyde toxicity. EMBO Molecular Medicine, 2017, 9, 1398-1414.	3.3	57
47	Profiles of lipids, blood pressure and weight changes among premenopausal Chinese breast cancer patients after adjuvant chemotherapy. BMC Women's Health, 2017, 17, 55.	0.8	22
48	Telomeres and Physical Activity., 2017,, 103-116.		O
49	The mechanism of transactivation regulation due to polymorphic short tandem repeats (STRs) using IGF1 promoter as a model. Scientific Reports, 2016, 6, 38225.	1.6	23
50	CrossNorm: a novel normalization strategy for microarray data in cancers. Scientific Reports, 2016, 6, 18898.	1.6	46
51	Genome-wide association study in Chinese identifies new susceptibility loci associated with chronic kidney disease in type 2 diabetes. Diabetes Research and Clinical Practice, 2016, 120, S49-S50.	1.1	O
52	A network based covariance test for detecting multivariate eQTL in saccharomyces cerevisiae. BMC Systems Biology, 2016, 10, 8.	3.0	1
53	AGTR1 has undergone natural selection in Euro-Asian populations in relation to ambient temperature that predisposes Chinese populations to essential hypertension. International Journal of Cardiology, 2016, 209, 278-280.	0.8	7
54	Association of gene expression and methylation of UQCRC1 to the predisposition of Alzheimer's disease in a Chinese population. Journal of Psychiatric Research, 2016, 76, 143-147.	1.5	16

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55	Sequence variants in the PTCH1 gene associate with spine bone mineral density and osteoporotic fractures. Nature Communications, 2016, 7, 10129.	5.8	58
56	Gene-gene Interaction Analysis by IAC (Interaction Analysis by Chi-Square) - A Novel Biological Constraint-based Interaction Analysis Framework. , 2016, , .		3
57	Telomere length is not associated with frailty in older Chinese elderly: Cross-sectional and longitudinal analysis. Mechanisms of Ageing and Development, 2015, 152, 74-79.	2.2	32
58	Applicability of <scp>BALAD</scp> score in prognostication of hepatitis <scp>B</scp> â€related hepatocellular carcinoma. Journal of Gastroenterology and Hepatology (Australia), 2015, 30, 1529-1535.	1.4	14
59	Chemotherapy-Related Amenorrhea and Menopause in Young Chinese Breast Cancer Patients: Analysis on Incidence, Risk Factors and Serum Hormone Profiles. PLoS ONE, 2015, 10, e0140842.	1.1	25
60	EGFR: The Paradigm of an Oncogene-Driven Lung Cancer. Clinical Cancer Research, 2015, 21, 2221-2226.	3.2	72
61	Exome Sequencing Identifies a Rare <i>HSPG2</i> Variant Associated with Familial Idiopathic Scoliosis. G3: Genes, Genomes, Genetics, 2015, 5, 167-174.	0.8	49
62	MicroRNAs mediated targeting on the Yin-yang dynamics of DNA methylation in disease and development. International Journal of Biochemistry and Cell Biology, 2015, 67, 115-120.	1.2	20
63	Prehospital Emergency Care Training Practices Regarding Lesbian, Gay, Bisexual, and Transgender Patients in Maryland (USA). Prehospital and Disaster Medicine, 2015, 30, 163-166.	0.7	6
64	Genome-wide association study identifies new susceptibility loci for adolescent idiopathic scoliosis in Chinese girls. Nature Communications, 2015, 6, 8355.	5.8	104
65	Uniaxial cyclic stretch stimulates TRPV4 to induce realignment of human embryonic stem cell-derived cardiomyocytes. Journal of Molecular and Cellular Cardiology, 2015, 87, 65-73.	0.9	25
66	Dynamic changes of DNA epigenetic marks in mouse oocytes during natural and accelerated aging. International Journal of Biochemistry and Cell Biology, 2015, 67, 121-127.	1,2	26
67	The Role of Piezo1 in Regulating Cytosolic Calcium Level in Aortic Smooth Muscle Cells in Rats. FASEB Journal, 2015, 29, 943.3.	0.2	O
68	An Upregulation in the Expression of Vanilloid Transient Potential Channels 2 Enhances Hypotonicity-Induced Cytosolic Ca2+ Rise in Human Induced Pluripotent Stem Cell Model of Hutchinson Gillford Progeria. PLoS ONE, 2014, 9, e87273.	1.1	16
69	Gender difference of serum angiotensin-converting enzyme (ACE) activity in DD genotype of ACE insertion/deletion polymorphism in elderly Chinese. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2014, 15, 547-552.	1.0	15
70	The predisposition to thyrotoxic periodic paralysis (<scp>TPP</scp>) is due to a genetic variant in the inwardâ€rectifying potassium channel, <i><scp>KCNJ</scp>2</i> . Clinical Endocrinology, 2014, 80, 770-771.	1.2	2
71	Use of Tissue Adhesive as a Field Expedient Barrier Dressing for Hand Wounds in Disaster Responders. Prehospital and Disaster Medicine, 2014, 29, 107-109.	0.7	2
72	Use of a Hooked Cutting Device Compared With Scissors for the Emergency Exposure of Critically Ill and Injured Patients. Prehospital and Disaster Medicine, 2014, 29, 43-46.	0.7	3

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73	Inherited metabolic diseases in the Southern Chinese population: spectrum of diseases and estimated incidence from recurrent mutations. Pathology, 2014, 46, 375-382.	0.3	21
74	Gender and telomere length: Systematic review and meta-analysis. Experimental Gerontology, 2014, 51, 15-27.	1.2	394
75	Rare variants in FBN1 and FBN2 are associated with severe adolescent idiopathic scoliosis. Human Molecular Genetics, 2014, 23, 5271-5282.	1.4	111
76	Deciphering global signal features of high-throughput array data from cancers. Molecular BioSystems, 2014, 10, 1549-1556.	2.9	7
77	Molecular evolution in the CREB1 signal pathway and a rare haplotype in CREB1 with genetic predisposition to schizophrenia. Journal of Psychiatric Research, 2014, 57, 84-89.	1.5	18
78	Telomere length is associated with decline in grip strength in older persons aged 65Âyears and over. Age, 2014, 36, 9711.	3.0	37
79	Estrogen Receptor α Polymorphisms and the Risk of Cognitive Decline: A 2-Year Follow-Up Study. American Journal of Geriatric Psychiatry, 2014, 22, 489-498.	0.6	10
80	A meta-analysis identifies adolescent idiopathic scoliosis association with <i>LBX1</i> locus in multiple ethnic groups. Journal of Medical Genetics, 2014, 51, 401-406.	1.5	79
81	Effect of whole soy and purified isoflavone daidzein on renal function—a 6-month randomized controlled trial in equol-producing postmenopausal women with prehypertension. Clinical Biochemistry, 2014, 47, 1250-1256.	0.8	18
82	Familial Young-Onset Diabetes, Pre-Diabetes and Cardiovascular Disease Are Associated with Genetic Variants of DACH1 in Chinese. PLoS ONE, 2014, 9, e84770.	1.1	16
83	Urinary Sodium Excretion and Dietary Sources of Sodium Intake in Chinese Postmenopausal Women with Prehypertension. PLoS ONE, 2014, 9, e104018.	1.1	20
84	Medical support for law enforcement-extended operations incidents. American Journal of Disaster Medicine, 2014, 9, 127-135.	0.1	0
85	A single nucleotide polymorphism in microRNAâ€146a is associated with the risk for nasopharyngeal carcinoma. Molecular Carcinogenesis, 2013, 52, 28-38.	1.3	38
86	Association of polymorphisms in the Chr18q11.2 locus with tuberculosis in Chinese population. Human Genetics, 2013, 132, 691-695.	1.8	21
87	Genome-wide association study in a Chinese population identifies a susceptibility locus for type 2 diabetes at 7q32 near PAX4. Diabetologia, 2013, 56, 1291-1305.	2.9	94
88	Functional Interaction Between SNPs and Microsatellite in the Transcriptional Regulation of Insulin-Like Growth Factor 1. Human Mutation, 2013, 34, 1289-1297.	1.1	18
89	Carnitine Uptake Defect (Primary Carnitine Deficiency): Risk in Genotype-Phenotype Correlation. Human Mutation, 2013, 34, 655-655.	1.1	13
90	Lack of association between polymorphisms from genome-wide association studies and tuberculosis in the Chinese population. Scandinavian Journal of Infectious Diseases, 2013, 45, 310-314.	1.5	15

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91	Role of pharmacogenetics on adjuvant chemotherapy-induced neutropenia in Chinese breast cancer patients. Journal of Cancer Research and Clinical Oncology, 2013, 139, 419-427.	1.2	35
92	Mitochondrial DNA haplogroup Y is associated to Leigh syndrome in Chinese population. Gene, 2013, 512, 460-463.	1.0	11
93	The role of the Ala746Thr variant in the ATP13A2 gene among Chinese patients with Parkinson's disease. Journal of Clinical Neuroscience, 2013, 20, 761-762.	0.8	7
94	Isolated persistent elevation of alanine transaminase for early diagnosis of pre-symptomatic Wilson's disease in Chinese children. World Journal of Pediatrics, 2013, 9, 361-364.	0.8	8
95	Telomere length and cognitive function in southern Chinese community-dwelling male elders. Age and Ageing, 2013, 42, 450-455.	0.7	37
96	Correlation of telomere length shortening with TP53 somatic mutations, polymorphisms and allelic loss in breast tumors and esophageal cancer. Oncology Reports, 2013, 29, 226-236.	1.2	17
97	Serotonin receptor 2C gene polymorphism associated with post-stroke depression in Chinese patients. Genetics and Molecular Research, 2013, 12, 1546-1553.	0.3	17
98	Genetic Associations of Type 2 Diabetes with Islet Amyloid Polypeptide Processing and Degrading Pathways in Asian Populations. PLoS ONE, 2013, 8, e62378.	1.1	7
99	GP-Pi: Using Genetic Programming with Penalization and Initialization on Genome-Wide Association Study. Lecture Notes in Computer Science, 2013, , 330-341.	1.0	O
100	Chinese Tea and Telomere Length in Elderly Chinese Men. , 2013, , 1117-1127.		0
101	NEDD9 Gene Polymorphism Influences the Risk of Alzheimer Disease and Cognitive Function in Chinese Older Persons. Alzheimer Disease and Associated Disorders, 2012, 26, 88-90.	0.6	4
102	MxA Polymorphisms Are Associated with Risk and Age-at-Onset in Alzheimer Disease and Accelerated Cognitive Decline in Chinese Elders. Rejuvenation Research, 2012, 15, 516-522.	0.9	8
103	A novel ALDH5A1 mutation in a patient with succinic semialdehyde dehydrogenase deficiency. Pathology, 2012, 44, 280-282.	0.3	6
104	Genome-wide meta-analysis identifies 56 bone mineral density loci and reveals 14 loci associated with risk of fracture. Nature Genetics, 2012, 44, 491-501.	9.4	1,100
105	Comparative analysis of protein-coding genes and long non-coding RNAs of prostate cancer between Caucasian and Chinese populations. , 2012, , .		1
106	A PIN1 polymorphism that prevents its suppression by AP4 associates with delayed onset of Alzheimer's disease. Neurobiology of Aging, 2012, 33, 804-813.	1.5	68
107	Genetic Adaptation of the Hypoxia-Inducible Factor Pathway to Oxygen Pressure among Eurasian Human Populations. Molecular Biology and Evolution, 2012, 29, 3359-3370.	3.5	29
108	The first reported HLCS gene mutation causing holocarboxylase synthetase deficiency in a Vietnamese patient. World Journal of Pediatrics, 2012, 8, 278-280.	0.8	4

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109	Familial aggregation and heritability of insomnia in a community-based study. Sleep Medicine, 2012, 13, 985-990.	0.8	60
110	The Origin and Evolution of Variable Number Tandem Repeat of CLEC4M Gene in the Global Human Population. PLoS ONE, 2012, 7, e30268.	1.1	6
111	The VNTR Polymorphism of the DC-SIGNR Gene and Susceptibility to HIV-1 Infection: A Meta-Analysis. PLoS ONE, 2012, 7, e42972.	1.1	8
112	Genetic epidemiology and heritability of AIS: A study of 415 Chinese female patients. Journal of Orthopaedic Research, 2012, 30, 1464-1469.	1,2	42
113	The ALU polymorphism of angiotensin I converting enzyme (ACE) and atherosclerosis, incident chronic diseases and mortality in an elderly Chinese population. Journal of Nutrition, Health and Aging, 2012, 16, 262-268.	1.5	4
114	Insomnia, sleep quality, pain, and somatic symptoms: Sex differences and shared genetic components. Pain, 2012, 153, 666-673.	2.0	87
115	Novel mutations in PHKA2 gene in glycogen storage disease type IX patients from Hong Kong, China. Molecular Genetics and Metabolism, 2011, 102, 222-225.	0.5	13
116	Familial aggregation of narcolepsy. Sleep Medicine, 2011, 12, 947-951.	0.8	15
117	Haplotype effect in the IGF1 promoter accounts for the association between microsatellite and serum IGF1 concentration. Clinical Endocrinology, 2011, 74, 520-527.	1.2	13
118	Top Theories for the Etiopathogenesis of Adolescent Idiopathic Scoliosis. Journal of Pediatric Orthopaedics, 2011, 31, S14-S27.	0.6	134
119	CBS gene mutations found in a Chinese pyridoxine-responsive homocystinuria patient. Pathology, 2011, 43, 81-83.	0.3	4
120	A hidden two-locus disease association pattern in genome-wide association studies. BMC Bioinformatics, 2011, 12, 156.	1,2	2
121	Shorter Telomere Length is Associated With Greater Decrease in Ankle-Brachial Index in Elderly Chinese Women but not Men. Angiology, 2011, 62, 87-91.	0.8	3
122	An HIV-1 Resistance Polymorphism in TRIM5α Gene Among Chinese Intravenous Drug Users. Journal of Acquired Immune Deficiency Syndromes (1999), 2011, 56, 306-311.	0.9	28
123	Chinese tea consumption is associated with longer telomere length in elderly Chinese men. British Journal of Nutrition, 2010, 103, 107-113.	1.2	56
124	Lack of Association Between the Promoter Polymorphisms of MMP-3 and IL-6 Genes and Adolescent Idiopathic Scoliosis. Spine, 2010, 35, 1701-1705.	1.0	27
125	BOOST: A Fast Approach to Detecting Gene-Gene Interactions in Genome-wide Case-Control Studies. American Journal of Human Genetics, 2010, 87, 325-340.	2.6	452
126	The effect of telomere length, a marker of biological aging, on bone mineral density in elderly population. Osteoporosis International, 2010, 21, 89-97.	1.3	28

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127	<i>PHF11</i> is not a major candidate gene for asthma or eczema in Chinese children. Pediatric Pulmonology, 2010, 45, 890-897.	1.0	1
128	Association of disease-predisposition polymorphisms of the melatonin receptors and sunshine duration in the global human populations. Journal of Pineal Research, 2010, 48, 133-141.	3.4	18
129	Common variants near CAV1 and CAV2 are associated with primary open-angle glaucoma. Nature Genetics, 2010, 42, 906-909.	9.4	357
130	SCALY RASH. Journal of Paediatrics and Child Health, 2010, 46, 441-441.	0.4	1
131	Polarized Secretion of Interleukin (IL)-6 and IL-8 by Human Airway Epithelia 16HBE140- Cells in Response to Cationic Polypeptide Challenge. PLoS ONE, 2010, 5, e12091.	1.1	61
132	CNGA2 Contributes to ATP-Induced Noncapacitative Ca ²⁺ Influx in Vascular Endothelial Cells. Journal of Vascular Research, 2010, 47, 148-156.	0.6	17
133	Association of Plasma Soluble CTLA-4 with Lung Function and GenePolymorphism in Chinese Asthmatic Children. International Archives of Allergy and Immunology, 2010, 152, 113-121.	0.9	3
134	Telomeres and the ageing process. Reviews in Clinical Gerontology, 2010, 20, 1-9.	0.5	3
135	Association of SRD5A2 Variants and Serum Androstane-3α,17β-Diol Glucuronide Concentration in Chinese Elderly Men. Clinical Chemistry, 2010, 56, 1742-1749.	1.5	12
136	Predictive rule inference for epistatic interaction detection in genome-wide association studies. Bioinformatics, 2010, 26, 30-37.	1.8	156
137	Detecting two-locus associations allowing for interactions in genome-wide association studies. Bioinformatics, 2010, 26, 2517-2525.	1.8	23
138	421 The expression of CYP1B1 predicts neutropenia after adjuvant chemotherapy in breast cancer patients. European Journal of Cancer, Supplement, 2010, 8, 179.	2.2	0
139	Fine-scale stratification analysis of Hong Kong Chinese population. , 2010, , .		1
140	Sex-specific effect of Pirin gene on bone mineral density in a cohort of 4000 Chinese. Bone, 2010, 46, 543-550.	1.4	15
141	Association of genetic variations in aromatase gene with serum estrogen and estrogen/testosterone ratio in Chinese elderly men. Clinica Chimica Acta, 2010, 411, 53-58.	0.5	23
142	High-level expression of early growth response-1 and association of polymorphism with total IgE and atopy in allergic rhinitis adults. Clinica Chimica Acta, 2010, 411, 67-71.	0.5	5
143	ACE inhibitor use was associated with lower serum dehydroepiandrosterone concentrations in older men. Clinica Chimica Acta, 2010, 411, 1122-1125.	0.5	12
144	Diagnoses of newborns and mothers with carnitine uptake defects through newborn screening. Molecular Genetics and Metabolism, 2010, 100, 46-50.	0.5	86

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145	<i>CISH</i> and Susceptibility to Infectious Diseases. New England Journal of Medicine, 2010, 362, 2092-2101.	13.9	129
146	Evidence for Positive Selection on the Osteogenin (BMP3) Gene in Human Populations. PLoS ONE, 2010, 5, e10959.	1.1	9
147	European Bone Mineral Density Loci Are Also Associated with BMD in East-Asian Populations. PLoS ONE, 2010, 5, e13217.	1.1	81
148	Older men with higher self-rated socioeconomic status have shorter telomeres. Age and Ageing, 2009, 38, 553-558.	0.7	40
149	Interindividual and Interethnic Variation in Genomewide Gene Expression: Insights into the Biological Variation of Gene Expression and Clinical Implications. Clinical Chemistry, 2009, 55, 774-785.	1.5	20
150	Energy intake and expenditure profile in chronic peritoneal dialysis patients complicated with circulatory congestion. American Journal of Clinical Nutrition, 2009, 90, 1179-1184.	2.2	19
151	MegaSNPHunter: a learning approach to detect disease predisposition SNPs and high level interactions in genome wide association study. BMC Bioinformatics, 2009, 10, 13.	1.2	86
152	0150 The severity of chemotherapy-induced neutropenia in association with PTGS-2 gene polymorphism in breast cancer patients. Breast, 2009, 18, S57.	0.9	0
153	Cholesterol 24â€hydroxylase (CYP46A1) polymorphisms are associated with faster cognitive deterioration in Chinese older persons: a twoâ€year follow up study. International Journal of Geriatric Psychiatry, 2009, 24, 921-926.	1.3	26
154	Choice of study phenotype in osteoporosis genetic research. Journal of Bone and Mineral Metabolism, 2009, 27, 121-126.	1.3	17
155	The neck-region polymorphism of DC-SIGNR in peri-centenarian from Han Chinese Population. BMC Medical Genetics, 2009, 10, 134.	2.1	5
156	Promoter polymorphism of matrilin-1 gene predisposes to adolescent idiopathic scoliosis in a Chinese population. European Journal of Human Genetics, 2009, 17, 525-532.	1.4	89
157	Asthma and atopy are associated with chromosome 17q21 markers in Chinese children. Allergy: European Journal of Allergy and Clinical Immunology, 2009, 64, 621-628.	2.7	102
158	Association of <i>early growth responseâ€4</i> gene polymorphisms with total IgE and atopy in asthmatic children. Pediatric Allergy and Immunology, 2009, 20, 142-150.	1.1	11
159	Delayed Onset of Brown-Sequard Syndrome Involving Upper Extremity Pain. Pain Practice, 2009, 9, 150-151.	0.9	0
160	Genome-wide Association and Replication Studies Identified TRHR as an Important Gene for Lean Body Mass. American Journal of Human Genetics, 2009, 84, 418-423.	2.6	103
161	Persistence of osteopenia in adolescence — A longitudinal study of 147 girls till maturity with dual energy X-ray absorptiometry (DXA). Bone, 2009, 45, S63.	1.4	O
162	Novel mutations in ETFDH gene in Chinese patients with riboflavin-responsive multiple acyl-CoA dehydrogenase deficiency. Clinica Chimica Acta, 2009, 404, 95-99.	0.5	44

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163	Genetic association between a chemokine gene CXCL-10 (IP-10, interferon gamma inducible protein 10) and susceptibility to tuberculosis. Clinica Chimica Acta, 2009, 406, 98-102.	0.5	51
164	A community-based study of insomnia in Hong Kong Chinese children: Prevalence, risk factors and familial aggregation. Sleep Medicine, 2009, 10, 1040-1046.	0.8	84
165	Dietary intake, blood pressure and osteoporosis. Journal of Human Hypertension, 2009, 23, 451-455.	1.0	44
166	Polymorphisms of the estrogen receptor \hat{l}_{\pm} (ESR1) gene and the risk of Alzheimer's disease in a southern Chinese community. International Psychogeriatrics, 2009, 21, 977.	0.6	36
167	Green space, psychological restoration, and telomere length. Lancet, The, 2009, 373, 299-300.	6.3	65
168	Glycogen synthase kinaseâ€3β and tau genes interact in Alzheimer's disease. Annals of Neurology, 2008, 64, 446-454.	2.8	65
169	Dopamine Receptor D4 Gene â^'521C/T Polymorphism Is Associated with Opioid Dependence through Coldâ€Pain Responses. Annals of the New York Academy of Sciences, 2008, 1139, 20-26.	1.8	23
170	The BOADICEA model of genetic susceptibility to breast and ovarian cancers: updates and extensions. British Journal of Cancer, 2008, 98, 1457-1466.	2.9	461
171	Study of gene–gene interactions for endophenotypic quantitative traits in Chinese asthmatic children. Allergy: European Journal of Allergy and Clinical Immunology, 2008, 63, 1031-1039.	2.7	39
172	Telomeres and frailty. Mechanisms of Ageing and Development, 2008, 129, 642-648.	2.2	91
173	Association of prostaglandin-endoperoxide synthase 2 (PTGS2) polymorphisms and Alzheimer's disease in Chinese. Neurobiology of Aging, 2008, 29, 856-860.	1.5	29
174	BRE is an antiapoptotic protein in vivo and overexpressed in human hepatocellular carcinoma. Oncogene, 2008, 27, 1208-1217.	2.6	42
175	Narcolepsy in Southern Chinese patients: clinical characteristics, HLA typing and seasonality of birth. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 1262-1267.	0.9	27
176	Association between HLA-A Alleles and Alzheimer's Disease in a Southern Chinese Community. Dementia and Geriatric Cognitive Disorders, 2008, 26, 391-397.	0.7	24
177	Polymorphisms in the C-type lectin genes cluster in chromosome 19 and predisposition to severe acute respiratory syndrome coronavirus (SARS-CoV) infection. Journal of Medical Genetics, 2008, 45, 752-758.	1.5	22
178	No Association Between Physical Activity and Telomere Length in an Elderly Chinese Population 65 Years and Older. Archives of Internal Medicine, 2008, 168, 2163.	4.3	39
179	Association study between adolescent idiopathic scoliosis and the DPP9 gene which is located in the candidate region identified by linkage analysis. Postgraduate Medical Journal, 2008, 84, 498-501.	0.9	11
180	Lack of Association Between the Promoter Polymorphism of the MTNR1A Gene and Adolescent Idiopathic Scoliosis. Spine, 2008, 33, 2204-2207.	1.0	24

#	Article	IF	CITATIONS
181	Genetic Association Study of Growth Hormone Receptor and Idiopathic Scoliosis. Clinical Orthopaedics and Related Research, 2007, 462, 53-58.	0.7	31
182	Genetic Association of Complex Traits. Clinical Orthopaedics and Related Research, 2007, 462, 38-44.	0.7	64
183	Melatonin Receptor 1B (MTNR1B) Gene Polymorphism Is Associated With the Occurrence of Adolescent Idiopathic Scoliosis. Spine, 2007, 32, 1748-1753.	1.0	122
184	Gaucher disease among Chinese patients: Review on genotype/phenotype correlation from 29 patients and identification of novel and rare alleles. Blood Cells, Molecules, and Diseases, 2007, 38, 287-293.	0.6	55
185	Novel missense mutations in the first Chinese patient with very-long-chain acyl-CoA dehydrogenase deficiency. Clinica Chimica Acta, 2007, 375, 173-174.	0.5	2
186	A novel functional assay for simultaneous determination of total fatty acid \hat{l}^2 -oxidation flux and acylcarnitine profiling in human skin fibroblasts using 2H31-palmitate by isotope ratio mass spectrometry and electrospray tandem mass spectrometry. Clinica Chimica Acta, 2007, 382, 25-30.	0.5	19
187	The Familial Risk and HLA Susceptibility among Narcolepsy Patients in Hong Kong Chinese. Sleep, 2007, 30, 851-858.	0.6	15
188	Association analysis of GABA receptor subunit genes on 5q33 with heroin dependence in a Chinese male population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 439-443.	1.1	26
189	Lack of support for an association between CLEC4M homozygosity and protection against SARS coronavirus infection. Nature Genetics, 2007, 39, 691-692.	9.4	20
190	Lack of support for an association between CLEC4M homozygosity and protection against SARS coronavirus infection. Nature Genetics, 2007, 39, 692-693.	9.4	19
191	Association of prostaglandinâ€endoperoxide synthase 2 gene polymorphisms with asthma and atopy in Chinese children. Allergy: European Journal of Allergy and Clinical Immunology, 2007, 62, 802-809.	2.7	17
192	Association between candidate genes and lung function growth in Chinese asthmatic children. Clinical and Experimental Allergy, 2007, 37, 070806205546004-???.	1.4	21
193	The ?1Ssubunit of the L-type calcium channel is not a predisposition gene for thyrotoxic periodic paralysis. Clinical Endocrinology, 2007, 66, 229-234.	1.2	7
194	A study on whey protein supplement on physical performance and quality of life among elderly patients with chronic obstructive pulmonary disease. Australasian Journal on Ageing, 2007, 26, 168-172.	0.4	1
195	Abnormal expressions of the subunits of the UDP-N-acetylglucosamine: lysosomal enzyme, N-acetylglucosamine-1-phosphotransferase, result in the formation of cytoplasmic vacuoles resembling those of the I-cells. Journal of Molecular Medicine, 2007, 85, 351-360.	1.7	4
196	Deficiency of the carnitine transporter (OCTN2) with partial N-acetylglutamate synthase (NAGS) deficiency. Journal of Inherited Metabolic Disease, 2007, 30, 816-816.	1.7	11
197	Complete recovery from acute encephalopathy of lateâ€onset ornithine transcarbamylase deficiency in a 3â€yearâ€old boy. Journal of Inherited Metabolic Disease, 2007, 30, 981-981.	1.7	11
198	Genetic Association Study Between Asthma and Plasma IgE and Human Beta-Defensin-1 Gene in Chinese Children. Journal of Allergy and Clinical Immunology, 2006, 117, S324.	1.5	2

#	Article	IF	CITATIONS
199	Gene-gene interactions for asthma and plasma total IgE concentration in Chinese children. Journal of Allergy and Clinical Immunology, 2006, 117, 127-133.	1.5	74
200	A Relook Into the Association of the Estrogen Receptor α Gene (PvuII, XbaI) and Adolescent Idiopathic Scoliosis. Spine, 2006, 31, 2463-2468.	1.0	56
201	Polymorphisms of the cholesterol 24-hydroxylase (CYP46A1) gene and the risk of Alzheimer's disease in a Chinese population. International Psychogeriatrics, 2006, 18, 37-45.	0.6	24
202	How the SARS coronavirus causes disease: host or organism?. Journal of Pathology, 2006, 208, 142-151.	2.1	46
203	Genetic association study between mbl2 and asthma phenotypes in Chinese children. Pediatric Allergy and Immunology, 2006, 17, 501-507.	1.1	21
204	Topical 5-fluorouracil has no additional benefit in treating common warts with cryotherapy: a single-centre, double-blind, randomized, placebo-controlled trial. Clinical and Experimental Dermatology, 2006, 31, 394-397.	0.6	20
205	Asthma and atopy are associated with DEFB1 polymorphisms in Chinese children. Genes and Immunity, 2006, 7, 59-64.	2.2	44
206	Evaluation of an algorithm of tagging SNPs selection by linkage disequilibrium. Clinical Biochemistry, 2006, 39, 240-243.	0.8	11
207	Hepatitis C virus genotype distribution among intravenous drug user and the general population in Hong Kong. Journal of Medical Virology, 2006, 78, 574-581.	2.5	42
208	Apolipoprotein epsilon-4 allele and the two-year progression of cognitive function in Chinese subjects with late-onset Alzheimer's disease. American Journal of Alzheimer's Disease and Other Dementias, 2006, 21, 92-99.	0.9	14
209	Apolipoprotein E $\hat{l}\mu$ 4 Allele Is Associated with Vascular Dementia. Dementia and Geriatric Cognitive Disorders, 2006, 22, 301-305.	0.7	52
210	Very long-chain acyl-CoA dehydrogenase deficiency presenting as acute hypercapnic respiratory failure. European Respiratory Journal, 2006, 28, 447-450.	3.1	19
211	Nitric oxide synthase polymorphisms and asthma phenotypes in Chinese children. Clinical and Experimental Allergy, 2005, 35, 1288-1294.	1.4	53
212	Novel mutations in type 2 Gaucher disease in Chinese and their functional characterization by heterologous expression. Human Mutation, 2005, 26, 59-60.	1.1	13
213	Absence of association between angiotensin converting enzyme polymorphism and development of adult respiratory distress syndrome in patients with severe acute respiratory syndrome: a case control study. BMC Infectious Diseases, 2005, 5, 26.	1.3	41
214	Type I Gaucher disease with exophthalmos and pulmonary arteriovenous malformation. BMC Medical Genetics, 2005, 6, 25.	2.1	3
215	CD14 and Toll-Like Receptors: Potential Contribution of Genetic Factors and Mechanisms to Inflammation and Allergy. Inflammation and Allergy: Drug Targets, 2005, 4, 169-175.	3.1	41
216	Breast and ovarian cancer risks to carriers of the BRCA1 5382insC and 185delAG and BRCA2 6174delT mutations: a combined analysis of 22 population based studies. Journal of Medical Genetics, 2005, 42, 602-603.	1.5	121

#	Article	IF	CITATIONS
217	Early Enhanced Expression of Interferon-Inducible Protein-10 (CXCL-10) and Other Chemokines Predicts Adverse Outcome in Severe Acute Respiratory Syndrome. Clinical Chemistry, 2005, 51, 2333-2340.	1.5	184
218	Seasonal pattern of tuberculosis in Hong Kong. International Journal of Epidemiology, 2005, 34, 924-930.	0.9	64
219	The association between promoter polymorphism of the interleukin-10 gene and Alzheimer's disease. Neurobiology of Aging, 2005, 26, 1005-1010.	1.5	90
220	RANTES G-401A polymorphism is associated with allergen sensitization and FEV1 in Chinese children. Respiratory Medicine, 2005, 99, 216-219.	1.3	17
221	ACE2 Gene Polymorphisms Do Not Affect Outcome of Severe Acute Respiratory Syndrome. Clinical Chemistry, 2004, 50, 1683-1686.	1.5	76
222	Resting Energy Expenditure and Subsequent Mortality Risk in Peritoneal Dialysis Patients. Journal of the American Society of Nephrology: JASN, 2004, 15, 3134-3143.	3.0	112
223	No mutation in the KCNE3 potassium channel gene in Chinese thyrotoxic hypokalaemic periodic paralysis patients. Clinical Endocrinology, 2004, 61, 109-112.	1.2	26
224	Isolate diffuse thickening of glomerular capillary basement membrane: a renal lesion in prediabetes?. Modern Pathology, 2004, 17, 1506-1512.	2.9	50
225	5-HT2A T102C receptor polymorphism and neuropsychiatric symptoms in Alzheimer's disease. International Journal of Geriatric Psychiatry, 2004, 19, 523-526.	1.3	55
226	Association between tumor necrosis factor-α promoter polymorphism and Alzheimer's disease. Neurology, 2004, 62, 307-309.	1.5	34
227	STAT6 in 17 SNP1 and in 18 SNP1 polymorphisms are not associated with asthma or atopy in Chinese children*1. Journal of Allergy and Clinical Immunology, 2004, 113, S205.	1.5	2
228	Association between TARC C-431T and atopy and asthma in children. Journal of Allergy and Clinical Immunology, 2004, 114, 199-202.	1.5	29
229	Case Report: 3-Methyglutaconic Aciduria in a Chinese Patient with Glycogen Storage Disease lb. Journal of Inherited Metabolic Disease, 2003, 26, 705-709.	1.7	11
230	Frequent hypermethylation of promoter region of RASSF1A in tumor tissues and voided urine of urinary bladder cancer patients. International Journal of Cancer, 2003, 104, 611-616.	2.3	97
231	A genomic approach to mutation analysis of holocarboxylase synthetase gene in three Chinese patients with late-onset holocarboxylase synthetase deficiency. Clinical Biochemistry, 2003, 36, 145-149.	0.8	13
232	The Câ^'159T polymorphism in the CD14 promoter is associated with serum total IgE concentration in atopic Chinese children. Pediatric Allergy and Immunology, 2003, 14, 255-260.	1.1	90
233	A genetic association study between asthma and CD14 promoter polymorphism in Chinese children. Journal of Allergy and Clinical Immunology, 2003, 111, S314.	1.5	0
234	Average Risks of Breast and Ovarian Cancer Associated with BRCA1 or BRCA2 Mutations Detected in Case Series Unselected for Family History: A Combined Analysis of 22 Studies. American Journal of Human Genetics, 2003, 72, 1117-1130.	2.6	3,105

#	Article	IF	CITATIONS
235	A novel mutation (G233D) in the glycogen phosphorylase gene in a patient with hepatic glycogen storage disease and residual enzyme activity. Molecular Genetics and Metabolism, 2003, 79, 142-145.	0.5	25
236	Carnitine level in Chinese epileptic patients taking sodium valproate. Pediatric Neurology, 2003, 28, 24-27.	1.0	19
237	A Clinical Guide to Inherited Metabolic Diseases, 2nd edition. Joe T.R. Clarke. Cambridge: Cambridge University Press, 2002, 306 pp., \$40.00, softcover. ISBN 0-521-89076-4 Clinical Chemistry, 2003, 49, 1232-1233.	1.5	0
238	Re: Hormonal Markers and Hepatitis B Virus-Related Hepatocellular Carcinoma Risk: A Nested Case-Control Study Among Men. Journal of the National Cancer Institute, 2003, 95, 559-560.	3.0	7
239	Lack of Association of the Interleukin-1β Gene Polymorphism with Alzheimer's Disease in a Chinese Population. Dementia and Geriatric Cognitive Disorders, 2003, 16, 265-268.	0.7	23
240	Renal Outcome in Type 2 Diabetic Patients With or Without Coexisting Nondiabetic Nephropathies. Diabetes Care, 2002, 25, 900-905.	4.3	118
241	Determinants of Cervical Human Papillomavirus Infection: Differences between High―and Lowâ€Oncogenic Risk Types. Journal of Infectious Diseases, 2002, 185, 28-35.	1.9	63
242	Total daily energy expenditure in wasted chronic obstructive pulmonary disease patients. European Journal of Clinical Nutrition, 2002, 56, 282-287.	1.3	32
243	Impact of Experimental Acute Hyponatremia on Severe Traumatic Brain Injury in Rats: Influences on Injuries, Permeability of Blood–Brain Barrier, Ultrastructural Features, and Aquaporin-4 Expression. Experimental Neurology, 2002, 178, 194-206.	2.0	46
244	The-401 polymorphism in RANTES gene promoter is associated with sensitization to inhalant allergens but not asthma in Chinese children. Journal of Allergy and Clinical Immunology, 2002, 109, S223-S223.	1.5	0
245	Primary IgA nephropathy with low histologic grade and disease progression: Is there a "point of no return�. American Journal of Kidney Diseases, 2002, 39, 401-406.	2.1	20
246	Inhalant Allergens as Risk Factors for the Development and Severity of Mild-to-Moderate Asthma in Hong Kong Chinese Children. Journal of Asthma, 2002, 39, 323-330.	0.9	25
247	Sensitization to Common Food Allergens Is a Risk Factor for Asthma in Young Chinese Children in Hong Kong. Journal of Asthma, 2002, 39, 523-529.	0.9	31
248	Low-density lipoprotein receptor-related protein 8 (apolipoprotein E receptor 2) gene polymorphisms in Alzheimer's disease. Neuroscience Letters, 2002, 332, 216-218.	1.0	32
249	Genotype spectrum of cervical human papillomavirus infection among sexually transmitted disease clinic patients in Hong Kong. Journal of Medical Virology, 2002, 68, 273-277.	2.5	24
250	A founder mutation (R254X) of SLC22A5 (OCTN2) in Chinese primary carnitine deficiency patients. Human Mutation, 2002, 20, 232-232.	1.1	43
251	Mutational and haplotype analysis of AGL in patients with glycogen storage disease type III. Journal of Human Genetics, 2002, 47, 55-59.	1.1	21
252	Thromboxane A2 receptor gene polymorphism is associated with the serum concentration of cat-specific immunoglobulin E as well as the development and severity of asthma in Chinese children. Pediatric Allergy and Immunology, 2002, 13, 10-17.	1.1	29

#	Article	IF	CITATIONS
253	Diagnosing Wilson's disease in a 5-year-old child. Journal of Paediatrics and Child Health, 2002, 38, 412-413.	0.4	2
254	Distribution in allele frequencies of predisposition-to-atopy genotypes in Chinese children. Pediatric Pulmonology, 2002, 34, 419-424.	1.0	20
255	Hypermethylation of multiple genes in tumor tissues and voided urine in urinary bladder cancer patients. Clinical Cancer Research, 2002, 8, 464-70.	3. 2	221
256	Somatic \hat{l}^2 -catenin mutation in gastric carcinoma $\hat{a} \in \hat{l}^4$ an infrequent event that is not specific for microsatellite instability. Cancer Letters, 2001, 163, 125-130.	3.2	23
257	Reversible Renal Failure in Paroxysmal Nocturnal Hemoglobinuria. American Journal of Kidney Diseases, 2001, 37, e17.1-e17.6.	2.1	44
258	Prevalence of breast cancer predisposition gene mutations in Chinese women and guidelines for genetic testing. Clinica Chimica Acta, 2001, 313, 179-185.	0.5	20
259	Overview of common inherited metabolic diseases in a Southern Chinese population of Hong Kong. Clinica Chimica Acta, 2001, 313, 195-201.	0.5	12
260	Association between mu opioid receptor gene polymorphisms and Chinese heroin addicts. NeuroReport, 2001, 12, 1103-1106.	0.6	138
261	A polymorphism in the coding region of interleukin-13 gene is associated with atopy but not asthma in Chinese children. Clinical and Experimental Allergy, 2001, 31, 1515-1521.	1.4	75
262	Prevalence and genotype distribution of TT virus in various specimen types from thalassaemic patients. Journal of Viral Hepatitis, 2001, 8, 304-309.	1.0	19
263	Pathology of fatal human infection associated with avian influenza A H5N1 virus. Journal of Medical Virology, 2001, 63, 242-246.	2.5	405
264	Genetic imbalances in pT2 breast cancers of southern Chinese women. Cancer Genetics and Cytogenetics, 2001, 124, 56-61.	1.0	16
265	Vitamin B12 deficiency—need for a new guideline. Nutrition, 2001, 17, 917-920.	1.1	34
266	Assessment of total energy expenditure in a Chinese population by a physical activity questionnaire: examination of validity. International Journal of Food Sciences and Nutrition, 2001, 52, 269-282.	1.3	69
267	Congenital Hypertrophy of the Retinal Pigment Epithelium and <i>APC</i> Mutations in Chinese with Familial Adenomatous Polyposis. Ophthalmologica, 2001, 215, 408-411.	1.0	11
268	Recurrent and novel mutations of GCDH gene in Chinese glutaric acidemia type I families. Human Mutation, 2000, 16, 446-446.	1.1	26
269	Application of Urine Magnesium/Creatinine Ratio as an Indicator for Insufficient Magnesium Intake. Clinical Biochemistry, 2000, 33, 675-678.	0.8	18
270	Validation of prediction equations for basal metabolic rate in Chinese subjects. European Journal of Clinical Nutrition, 2000, 54, 551-554.	1.3	47

#	Article	IF	CITATIONS
271	A New Recommendation for Maternal Weight Gain in Chinese Women. Journal of the American Dietetic Association, 2000, 100, 791-796.	1.3	64
272	Congenital hypertrophy of the retinal pigment epithelium and APC mutations in two Chinese families with familial adenomatous polyposis. Eye, 2000, 14, 18-22.	1.1	16
273	New Features of Renal Lesion Induced by Stroma Free Hemoglobin. Toxicologic Pathology, 2000, 28, 635-642.	0.9	28
274	Outcome of IgA nephropathy in adults graded by chronic histological lesions. American Journal of Kidney Diseases, 2000, 35, 392-400.	2.1	64
275	Characterization of early IgA nephropathy. American Journal of Kidney Diseases, 2000, 36, 703-708.	2.1	34
276	The Impact of Acute Hyponatraemia on Severe Traumatic Brain Injury in Rats., 2000, 76, 405-408.		4
277	Prevalence of Mutations in the BRCA1 Gene Among Chinese Patients With Breast Cancer. Journal of the National Cancer Institute, 1999, 91, 882-885.	3.0	59
278	Mutations of OCTN2, an Organic Cation/Carnitine Transporter, Lead to Deficient Cellular Carnitine Uptake in Primary Carnitine Deficiency. Human Molecular Genetics, 1999, 8, 655-660.	1.4	148
279	Classical galactosaemia in Chinese: A case report and review of disease incidence. Journal of Paediatrics and Child Health, 1999, 35, 399-400.	0.4	21
280	Severe hypokalemic myopathy in Gitelman's syndrome. , 1999, 22, 545-547.		8
281	Avoiding cardiopulmonary bypass in multivessel CABG reduces cytokine response and myocardial injury. Annals of Thoracic Surgery, 1999, 68, 52-56.	0.7	254
282	Primary plasmalemmal carnitine transporter defect manifested with dicarboxylic aciduria and impaired fatty acid oxidation. Journal of Inherited Metabolic Disease, 1998, 21, 423-425.	1.7	9
283	Determination of the risk for familial disease in RET mutationâ€negative patients with medullary thyroid cancer. Journal of Internal Medicine, 1998, 244, 185-187.	2.7	1
284	Acute Renal Failure and Proximal Tubule Lesions after Trichosanthin Injection in Rats. Experimental and Molecular Pathology, 1997, 64, 78-89.	0.9	8
285	Histological changes of parathyroid adenoma after percutaneous injection of ethanol. Histopathology, 1997, 30, 87-89.	1.6	8
286	Polar spongioblastoma with cerebrospinal fluid metastases. World Neurosurgery, 1994, 41, 137-142.	1.3	6
287	The otological status of patients with nasopharyngeal carcinoma after megavoltage radiotherapy. Journal of Laryngology and Otology, 1992, 106, 1055-1058.	0.4	18