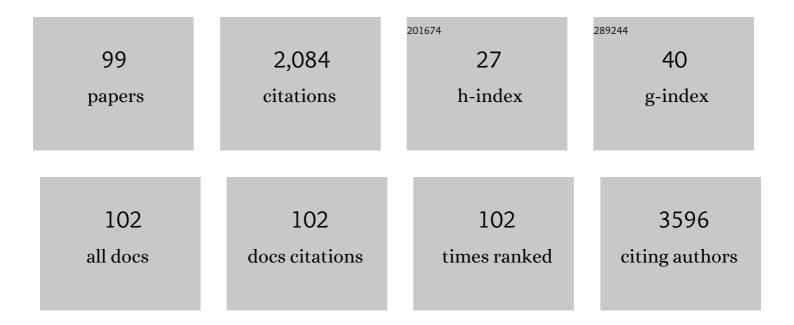
Jan-Gowth Chang

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genetic contributions to female gout and hyperuricaemia using genome-wide association study and polygenic risk score analyses. Rheumatology, 2023, 62, 638-646.	1.9	3
2	Application of Whole Exome Sequencing and Functional Annotations to Identify Genetic Variants Associated with Marfan Syndrome. Journal of Personalized Medicine, 2022, 12, 198.	2.5	2
3	Clinical application of liquid biopsy in cancer patients. BMC Cancer, 2022, 22, 413.	2.6	3
4	Mutation Analysis of Second Primary Tumors in Oral Cancer in Taiwanese Patients through Next-Generation Sequencing. Diagnostics, 2022, 12, 951.	2.6	2
5	A nine-gene signature identification and prognostic risk prediction for patients with lung adenocarcinoma using novel machine learning approach. Computers in Biology and Medicine, 2022, 145, 105493.	7.0	5
6	Construction and Validation of a Prognostic Gene-Based Model for Overall Survival Prediction in Hepatocellular Carcinoma Using an Integrated Statistical and Bioinformatic Approach. International Journal of Molecular Sciences, 2021, 22, 1632.	4.1	8
7	The Fusion Gene Landscape in Taiwanese Patients with Non-Small Cell Lung Cancer. Cancers, 2021, 13, 1343.	3.7	4
8	Immunohistochemical Expression of Five Protein Combinations Revealed as Prognostic Markers in Asian Oral Cancer. Frontiers in Genetics, 2021, 12, 643461.	2.3	6
9	A novel miRNA-based classification model of risks and stages for clear cell renal cell carcinoma patients. BMC Bioinformatics, 2021, 22, 270.	2.6	12
10	Oxidative Stress-Induced Unscheduled CDK1–Cyclin B1 Activity Impairs ER–Mitochondria-Mediated Bioenergetic Metabolism. Cells, 2021, 10, 1280.	4.1	5
11	Long Noncoding RNA NTT Context-Dependently Regulates MYB by Interacting With Activated Complex in Hepatocellular Carcinoma Cells. Frontiers in Oncology, 2021, 11, 592045.	2.8	2
12	Metatranscriptomic Analysis of Human Lung Metagenomes from Patients with Lung Cancer. Genes, 2021, 12, 1458.	2.4	17
13	Mutation profile of non-small cell lung cancer revealed by next generation sequencing. Respiratory Research, 2021, 22, 3.	3.6	35
14	Mitochondrial DNA haplogroups affect physical performances in Han older adults: an 8â€year followâ€up prospective cohort study. Geriatrics and Gerontology International, 2021, 21, 166-171.	1.5	1
15	Genome-Wide Analysis of Prognostic Alternative Splicing Signature and Splicing Factors in Lung Adenocarcinoma. Genes, 2020, 11, 1300.	2.4	13
16	Congenital dyserythropoiesis anemia type Ia with a novel <i>CDAN1</i> mutation diagnosed by whole exome sequencing. Molecular Genetics & amp; Genomic Medicine, 2020, 8, e1220.	1.2	1
17	Fasting glucose-to-HbA1c ratio is a good indicator of G6PD deficiency, but not thalassemia, in patients with type 2 diabetes mellitus. Clinica Chimica Acta, 2020, 506, 9-15.	1.1	3
18	Metabolic Imaging Phenotype Using Radiomics of [18F]FDG PET/CT Associated with Genetic Alterations of Colorectal Cancer. Molecular Imaging and Biology, 2019, 21, 183-190.	2.6	35

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19	Male-Specific Long Noncoding RNA TTTY15 Inhibits Non-Small Cell Lung Cancer Proliferation and Metastasis via TBX4. International Journal of Molecular Sciences, 2019, 20, 3473.	4.1	29
20	Chromosomal microarray and wholeâ€exome sequence analysis in Taiwanese patients with autism spectrum disorder. Molecular Genetics & Genomic Medicine, 2019, 7, e996.	1.2	11
21	Molecular characterization of colorectal cancer using wholeâ€exome sequencing in a Taiwanese population. Cancer Medicine, 2019, 8, 3738-3747.	2.8	18
22	Alternative splicing in human cancer cells is modulated by the amiloride derivative 3,5â€diaminoâ€6â€chloroâ€Nâ€(Nâ€(2,6â€dichlorobenzoyl)carbamimidoyl)pyrazineâ€2â€carboxide. Molecular (2019, 13, 1744-1762.	D raco logy,	9
23	Antrodia cinnamomea, a Treasured Medicinal Mushroom, Induces Growth Arrest in Breast Cancer Cells, T47D Cells: New Mechanisms Emerge. International Journal of Molecular Sciences, 2019, 20, 833.	4.1	11
24	Genome-wide analysis of lncRNAs in 3'-untranslated regions: CR933609 acts as a decoy to protect the INO80D gene. International Journal of Oncology, 2018, 53, 417-433.	3.3	5
25	Mutation Analysis of Second Primary Tumors in the Head and Neck Cancer by Next Generation Sequencing. , 2018, , .		0
26	lncRNA NTT/PBOV1 Axis Promotes Monocyte Differentiation and Is Elevated in Rheumatoid Arthritis. International Journal of Molecular Sciences, 2018, 19, 2806.	4.1	51
27	Whole-exome sequencing for the genetic diagnosis of congenital red blood cell membrane disorders in Taiwan. Clinica Chimica Acta, 2018, 487, 311-317.	1.1	13
28	Correlation of genomic alterations between tumor tissue and circulating tumor DNA by next-generation sequencing. Journal of Cancer Research and Clinical Oncology, 2018, 144, 2167-2175.	2.5	9
29	Whole exome sequencing in Dandy-Walker variant with intellectual disability reveals an activating CIP2A mutation as novel genetic cause. Neurogenetics, 2018, 19, 157-163.	1.4	6
30	G6PD as a predictive marker for glioma risk, prognosis and chemosensitivity. Journal of Neuro-Oncology, 2018, 139, 661-670.	2.9	22
31	Prognostic Value of RNASEH2A-, CDK1-, and CD151-Related Pathway Gene Profiling for Kidney Cancers. International Journal of Molecular Sciences, 2018, 19, 1586.	4.1	8
32	Determination of the mutational landscape in Taiwanese patients with papillary thyroid cancer by whole-exome sequencing. Human Pathology, 2018, 78, 151-158.	2.0	9
33	Detection of Molecular Alterations in Taiwanese Patients with Medullary Thyroid Cancer Using Whole-Exome Sequencing. Endocrine Pathology, 2018, 29, 324-331.	9.0	23
34	Securinine enhances SMN2 exon 7 inclusion in spinal muscular atrophy cells. Biomedicine and Pharmacotherapy, 2017, 88, 708-714.	5.6	9
35	Evaluation of whole exome sequencing by targeted gene sequencing and Sanger sequencing. Clinica Chimica Acta, 2017, 471, 222-232.	1.1	10
36	Muscle developmental defects in heterogeneous nuclear Ribonucleoprotein A1 knockout mice. Open Biology, 2017, 7, 160303.	3.6	36

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37	DNA-Sensing and Nuclease Gene Expressions as Markers for Colorectal Cancer Progression. Oncology, 2017, 92, 115-124.	1.9	41
38	4β-Hydroxywithanolide E Modulates Alternative Splicing of Apoptotic Genes in Human Hepatocellular Carcinoma Huh-7 Cells. Scientific Reports, 2017, 7, 7290.	3.3	10
39	Identification of novel mutations in endometrial cancer patients by whole-exome sequencing. International Journal of Oncology, 2017, 50, 1778-1784.	3.3	43
40	Impact of Enterobius vermicularis infection and mebendazole treatment on intestinal microbiota and host immune response. PLoS Neglected Tropical Diseases, 2017, 11, e0005963.	3.0	25
41	Modulation the alternative splicing of GLA (IVS4+919G>A) in Fabry disease. PLoS ONE, 2017, 12, e0175929.	2.5	15
42	Long noncoding RNA MIAT promotes non-small cell lung cancer proliferation and metastasis through MMP9 activation. Oncotarget, 2017, 8, 98148-98162.	1.8	50
43	Mutation analysis of 13 driver genes of colorectal cancer-related pathways in Taiwanese patients. World Journal of Gastroenterology, 2016, 22, 2314-2325.	3.3	28
44	Validating the Sensitivity of Highâ€Resolution Melting Analysis for JAK2 V617F Mutation in the Clinical Setting. Journal of Clinical Laboratory Analysis, 2016, 30, 838-844.	2.1	5
45	High-resolution Melting Analysis for Gene Scanning of Adenomatous Polyposis Coli (APC) Gene With Oral Squamous Cell Carcinoma Samples. Applied Immunohistochemistry and Molecular Morphology, 2016, 24, 97-104.	1.2	5
46	Long noncoding RNA TUG1 is downregulated in non-small cell lung cancer and can regulate CELF1 on binding to PRC2. BMC Cancer, 2016, 16, 583.	2.6	94
47	Analysing the mutational status of adenomatous polyposis coli (APC) gene in breast cancer. Cancer Cell International, 2016, 16, 23.	4.1	11
48	Development of a high-resolution melting method for the screening of TNFAIP3 gene mutations. Oncology Reports, 2016, 35, 2936-2942.	2.6	4
49	Somatic Mutations and Genetic Variants of NOTCH1 in Head and Neck Squamous Cell Carcinoma Occurrence and Development. Scientific Reports, 2016, 6, 24014.	3.3	33
50	TCH1036, a indeno[1,2-c]quinoline derivative, potentially inhibited the growth of human brain malignant glioma (GBM) 8401 cells via suppression of the expression of Suv39h1 and PARP. Biomedicine and Pharmacotherapy, 2016, 82, 649-659.	5.6	2
51	Genetic alterations in endometrial cancer by targeted next-generation sequencing. Experimental and Molecular Pathology, 2016, 100, 8-12.	2.1	24
52	Uncovering synthetic lethal interactions for therapeutic targets and predictive markers in lung adenocarcinoma. Oncotarget, 2016, 7, 73664-73680.	1.8	14
53	An XIST-related small RNA regulates KRAS G-quadruplex formation beyond X-inactivation. Oncotarget, 2016, 7, 86713-86729.	1.8	4
54	Genetic Alterations in Colorectal Cancer Have Different Patterns on 18F-FDG PET/CT. Clinical Nuclear Medicine, 2015, 40, 621-626.	1.3	24

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55	Rapid Identification of FGFR2 Gene Mutations in Taiwanese Patients With Endometrial Cancer Using High-resolution Melting Analysis. Applied Immunohistochemistry and Molecular Morphology, 2015, 23, 532-537.	1.2	2
56	BubR1 Acts as a Promoter in Cellular Motility of Human Oral Squamous Cancer Cells through Regulating MMP-2 and MMP-9. International Journal of Molecular Sciences, 2015, 16, 15104-15117.	4.1	9
57	High Expression Level of Tra2-β1 Is Responsible for Increased SMN2 Exon 7 Inclusion in the Testis of SMA Mice. PLoS ONE, 2015, 10, e0120721.	2.5	12
58	Mutation Analysis of KCNQ1, KCNH2 and SCN5A Genes in Taiwanese Long QT Syndrome Patients. International Heart Journal, 2015, 56, 450-453.	1.0	11
59	The Overexpression of FEN1 and RAD54B May Act as Independent Prognostic Factors of Lung Adenocarcinoma. PLoS ONE, 2015, 10, e0139435.	2.5	28
60	Developing and Evaluating the HRM Technique for Identifying Cytochrome P450 2D6 Polymorphisms. Journal of Clinical Laboratory Analysis, 2015, 29, 220-225.	2.1	6
61	MALAT1 long non-coding RNA is overexpressed in multiple myeloma and may serve as a marker to predict disease progression. BMC Cancer, 2014, 14, 809.	2.6	134
62	Combined mutational analysis of RAS, BRAF, PIK3CA, and TP53 genes in Taiwanese patients with oral squamous cell carcinoma. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2014, 118, 110-116.e1.	0.4	16
63	High-resolution melting analyses for genetic variants in ARID5B and IKZF1 with childhood acute lymphoblastic leukemia susceptibility loci in Taiwan. Blood Cells, Molecules, and Diseases, 2014, 52, 140-145.	1.4	31
64	lncRNAMap: A map of putative regulatory functions in the long non-coding transcriptome. Computational Biology and Chemistry, 2014, 50, 41-49.	2.3	36
65	Direct assessment of cytochrome P450 2D6 genotypes by high-resolution melting analysis and DNA sequencing. Environmental Toxicology and Pharmacology, 2014, 38, 821-828.	4.0	1
66	CSNK1E/CTNNB1 Are Synthetic Lethal To TP53 in Colorectal Cancer and Are Markers for Prognosis. Neoplasia, 2014, 16, 441-450.	5.3	23
67	Synthesis and antitumor activity evaluation of anilinoquinoline derivatives by the effect on the expression of polo-like kinase. Medicinal Chemistry Research, 2014, 23, 1437-1446.	2.4	2
68	Detection of KRAS codon 12 and 13 mutations by mutant-enriched PCR assay. Clinica Chimica Acta, 2014, 436, 169-175.	1.1	11
69	Demethylation within the proximal promoter region of human estrogen receptor alpha gene correlates with its enhanced expression: Implications for female bias in lupus. Molecular Immunology, 2014, 61, 28-37.	2.2	39
70	Pseudogene-Derived Endogenous siRNAs and Their Function. Methods in Molecular Biology, 2014, 1167, 227-239.	0.9	21
71	Histone-modifying genes as biomarkers in hepatocellular carcinoma. International Journal of Clinical and Experimental Pathology, 2014, 7, 2496-507.	0.5	31
72	Rapid detection of K-, N-, H-RAS, and BRAF hotspot mutations in thyroid cancer using the multiplex primer extension. Clinical Biochemistry, 2013, 46, 1572-1577.	1.9	6

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73	pseudoMap: an innovative and comprehensive resource for identification of siRNA-mediated mechanisms in human transcribed pseudogenes. Database: the Journal of Biological Databases and Curation, 2013, 2013, bat001-bat001.	3.0	7
74	Transcribed pseudogene Ï`PPM1K generates endogenous siRNA to suppress oncogenic cell growth in hepatocellular carcinoma. Nucleic Acids Research, 2013, 41, 3734-3747.	14.5	57
75	<i>RAS, BRAF,</i> and <i>TP53</i> Gene Mutations in Taiwanese Colorectal Cancer Patients. Oncology Research and Treatment, 2013, 36, 719-724.	1.2	17
76	High-resolution melting: Applications in genetic disorders. Clinica Chimica Acta, 2012, 414, 197-201.	1.1	77
77	Rapid identification of CYP2C8 polymorphisms by high resolution melting analysis. Clinica Chimica Acta, 2012, 413, 298-302.	1.1	8
78	The use of high resolution melting analysis to detect Fabry mutations in heterozygous females via dry bloodspots. Clinica Chimica Acta, 2012, 413, 422-427.	1.1	21
79	High-resolution melting (HRM) analysis for the detection of single nucleotide polymorphisms in microRNA target sites. Clinica Chimica Acta, 2012, 413, 1092-1097.	1.1	9
80	Characteristics and prevalence of KRAS, BRAF, and PIK3CA mutations in colorectal cancer by high-resolution melting analysis in Taiwanese population. Clinica Chimica Acta, 2012, 413, 1605-1611.	1.1	56
81	High-resolution melting (HRM) analysis as a feasible method for detecting spinal muscular atrophy via dried blood spots. Clinica Chimica Acta, 2012, 413, 1781-1785.	1.1	11
82	Increased expression of PRL-1 protein correlates with shortened patient survival in human hepatocellular carcinoma. Clinical and Translational Oncology, 2012, 14, 287-293.	2.4	14
83	High-resolution melting curve (HRM) analysis to establish CYP21A2 mutations converted from the CYP21A1P in congenital adrenal hyperplasia. Clinica Chimica Acta, 2011, 412, 1918-1923.	1.1	14
84	Small Molecule Amiloride Modulates Oncogenic RNA Alternative Splicing to Devitalize Human Cancer Cells. PLoS ONE, 2011, 6, e18643.	2.5	53
85	Detection of N-, H-, and KRAS codons 12, 13, and 61 mutations with universal RAS primer multiplex PCR and N-, H-, and KRAS-specific primer extension. Clinical Biochemistry, 2010, 43, 296-301.	1.9	28
86	Development of a high-resolution melting method for the detection of hemoglobin alpha variants. Clinical Biochemistry, 2010, 43, 671-676.	1.9	15
87	High resolution melting analysis facilitates mutation screening of ETFDH gene: Applications in riboflavin-responsive multiple acyl-CoA dehydrogenase deficiency. Clinica Chimica Acta, 2010, 411, 690-699.	1.1	34
88	Development of a high-resolution melting method for the screening of Wilson disease-related ATP7B gene mutations. Clinica Chimica Acta, 2010, 411, 1223-1231.	1.1	25
89	Expression of BUBR1 in human oral potentially malignant disorders and squamous cell carcinoma. Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics, 2010, 109, 257-267.	1.4	16
90	Comparison of two different screening methods for the KRAS mutation in colorectal cancer. Clinical Laboratory, 2010, 56, 175-86.	0.5	15

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#	Article	IF	CITATIONS
91	Fast simultaneous detection of K-RASmutations in colorectal cancer. BMC Cancer, 2009, 9, 179.	2.6	38
92	Rapid identification of HBB gene mutations by high-resolution melting analysis. Clinical Biochemistry, 2009, 42, 1667-1676.	1.9	41
93	Detection of the JAK2 V617F missense mutation by high resolution melting analysis and its validation. Clinica Chimica Acta, 2009, 408, 39-44.	1.1	28
94	5â€(<i>N</i> â€ethylâ€Nâ€isopropyl)â€amiloride enhances <i>SMN2</i> exon 7 inclusion and protein expression spinal muscular atrophy cells. Annals of Neurology, 2008, 63, 26-34.	in 5.3	35
95	Subcellular and Functional Proteomic Analysis of the Cellular Responses Induced by Helicobacter pylori. Molecular and Cellular Proteomics, 2006, 5, 702-713.	3.8	27
96	Epigenetic alteration of the SOCS1 gene in chronic myeloid leukaemia. British Journal of Haematology, 2003, 123, 654-661.	2.5	75
97	The correlation between CpG methylation on promoter and protein expression of E-cadherin in oral squamous cell carcinoma. Anticancer Research, 2002, 22, 3971-5.	1.1	34
98	Prenatal diagnosis of thalassemia in the Chinese. American Journal of Hematology, 1997, 55, 65-68.	4.1	8
99	Mutation Analysis of K- <i>ras</i> Oncogenes in Gastroenterologic Cancers by the Amplified Created Restriction Sites Method. American Journal of Clinical Pathology, 1993, 100, 686-689.	0.7	31