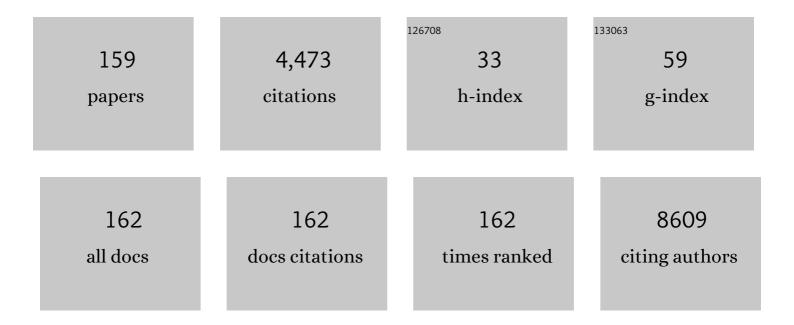
## Raffaele Palmirotta

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Correlation between targeted RNAseq signature of breast cancer CTCs and onset of bone-only metastases. British Journal of Cancer, 2022, 126, 419-429.	2.9	10
2	Comment on Kopańska et al. Disorders of the Cholinergic System in COVID-19 Era—A Review of the Latest Research. Int. J. Mol. Sci. 2022, 23, 672. International Journal of Molecular Sciences, 2022, 23, 2818.	1.8	2
3	SARS-CoV-2 infection after vaccination in Italian health care workers: a case report. The National Academy of Sciences, India, 2022, 45, 249-254.	0.8	2
4	Direct Oral Anticoagulants (DOAC): Are We Ready for a Pharmacogenetic Approach?. Journal of Personalized Medicine, 2022, 12, 17.	1.1	3
5	Association between Vitamin D Receptor Gene Polymorphisms and Periodontal Bacteria: A Clinical Pilot Study. Biomolecules, 2022, 12, 833.	1.8	3
6	Susceptibility to ischaemic heart disease: Focusing on genetic variants for ATP-sensitive potassium channel beyond traditional risk factors. European Journal of Preventive Cardiology, 2021, 28, 1495-1500.	0.8	22
7	Application of "omics―sciences to the prediction of bone metastases from breast cancer: State of the art. Journal of Bone Oncology, 2021, 26, 100337.	1.0	6
8	Targeted RNA-seq signature of breast cancer (BC) circulating tumor cells (CTCs) correlates with the onset of bone-only metastases. Bone Reports, 2021, 14, 100840.	0.2	0
9	Angiotensin System Polymorphisms' in SARS-CoV-2 Positive Patients: Assessment Between Symptomatic and Asymptomatic Patients: A Pilot Study. Pharmacogenomics and Personalized Medicine, 2021, Volume 14, 621-629.	0.4	37
10	The ATM Gene in Breast Cancer: Its Relevance in Clinical Practice. Genes, 2021, 12, 727.	1.0	29
11	Liquid Biopsy in Cervical Cancer: Hopes and Pitfalls. Cancers, 2021, 13, 3968.	1.7	9
12	Potential Role of eNOS Genetic Variants in Ischemic Heart Disease Susceptibility and Clinical Presentation. Journal of Cardiovascular Development and Disease, 2021, 8, 116.	0.8	18
13	Antioxidant and inflammatory polymorphism in patients with migraine with white matter hyperintensity (WMH). Journal of the Neurological Sciences, 2021, 429, 119300.	0.3	0
14	Could Small Neurotoxins-Peptides be Expressed during SARS-CoV-2 Infection?. Current Genomics, 2021, 22, 557-563.	0.7	4
15	239 Role of eNOS genetic variants on ischaemic heart disease susceptibility and acute coronary syndrome presentation. European Heart Journal Supplements, 2021, 23, .	0.0	0
16	Pharmacogenomics and Pharmacogenetics: In Silico Prediction of Drug Effects in Treatments for Novel Coronavirus SARS-CoV2 Disease. Pharmacogenomics and Personalized Medicine, 2020, Volume 13, 463-484.	0.4	8
17	DEAD-Box Helicase 4 (Ddx4)+ Stem Cells Sustain Tumor Progression in Non-Serous Ovarian Cancers. International Journal of Molecular Sciences, 2020, 21, 6096.	1.8	2
18	Pulmonary enteric adenocarcinoma: an overview. Expert Reviews in Molecular Medicine, 2020, 22, e1.	1.6	11

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19	Machine learning approach to predict medication overuse in migraine patients. Computational and Structural Biotechnology Journal, 2020, 18, 1487-1496.	1.9	23
20	Dual-procedural separation of CTCs in cutaneous melanoma provides useful information for both molecular diagnosis and prognosis. Therapeutic Advances in Medical Oncology, 2020, 12, 175883592090541.	1.4	10
21	ISCHEMIC HEART DISEASE AND GENETIC POLYMORPHISMS IN THE CROSS - TALK BETWEEN MYOCARDIAL METABOLISM AND CORONARY FLOW: AN UPDATE. Journal of the American College of Cardiology, 2019, 73, 151.	1.2	0
22	Age-Dependent Levels of Protein Kinase Cs in Brain: Reduction of Endogenous Mechanisms of Neuroprotection. International Journal of Molecular Sciences, 2019, 20, 3544.	1.8	8
23	Cutaneous metastasis as a primary presentation of a pulmonary enteric adenocarcinoma. International Journal of Biological Markers, 2019, 34, 421-426.	0.7	8
24	Nutrition and Female Fertility: An Interdependent Correlation. Frontiers in Endocrinology, 2019, 10, 346.	1.5	95
25	Dopamine-beta-hydroxylase 19-bp insertion/deletion polymorphism affects medication overuse in patients with chronic migraine. Neurological Sciences, 2019, 40, 1717-1724.	0.9	9
26	Rare Dihydropyrimidine Dehydrogenase Variants and Toxicity by Floropyrimidines: A Case Report. Frontiers in Oncology, 2019, 9, 139.	1.3	10
27	Dissection of major cancer gene variants in subsets of circulating tumor cells in advanced breast cancer. Scientific Reports, 2019, 9, 17276.	1.6	16
28	DAXX mutations as potential genomic markers of malignant evolution in small nonfunctioning pancreatic neuroendocrine tumors. Scientific Reports, 2019, 9, 18614.	1.6	26
29	Bone Metastases from Solid Tumors. , 2019, , 141-163.		2
30	Gene Fusion in NSCLC. , 2019, , 443-464.		1
31	Molecular Assessment of Ovarian Cancer and Translation to Clinical Management. , 2019, , 501-519.		Ο
32	Genetic bases of the nutritional approach to migraine. Critical Reviews in Food Science and Nutrition, 2019, 59, 2308-2320.	5.4	4
33	Redox Mechanisms in Migraine: Novel Therapeutics and Dietary Interventions. Antioxidants and Redox Signaling, 2018, 28, 1144-1183.	2.5	27
34	SNPs in predicting clinical efficacy and toxicity of chemotherapy: walking through the quicksand. Oncotarget, 2018, 9, 25355-25382.	0.8	34
35	Liquid biopsy of cancer: a multimodal diagnostic tool in clinical oncology. Therapeutic Advances in Medical Oncology, 2018, 10, 175883591879463.	1.4	317
36	Animal-type melanoma: dog or wolf? A review of the literature and a case report. Expert Reviews in Molecular Medicine, 2018, 20, e5.	1.6	2

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37	Vitamin D in melanoma: Controversies and potential role in combination with immune check-point inhibitors. Cancer Treatment Reviews, 2018, 69, 21-28.	3.4	31
38	Double Heterozygosity for BRCA1 Pathogenic Variant and BRCA2 Polymorphic Stop Codon K3326X: A Case Report in a Southern Italian Family. International Journal of Molecular Sciences, 2018, 19, 285.	1.8	14
39	Diabetes Mellitus and Ischemic Heart Disease: The Role of Ion Channels. International Journal of Molecular Sciences, 2018, 19, 802.	1.8	48
40	Serotonin receptor targeted therapy for migraine treatment: an overview of drugs in phase I and II clinical development. Expert Opinion on Investigational Drugs, 2017, 26, 269-277.	1.9	14
41	<i>ALK</i> gene alterations in cancer: biological aspects and therapeutic implications. Pharmacogenomics, 2017, 18, 277-292.	0.6	8
42	Immune‑related adverse events during anticancer immunotherapy: Pathogenesis and management (Review). Oncology Letters, 2017, 14, 5671-5680.	0.8	54
43	Procoagulant imbalance in premenopausal women with chronic migraine. Neurology, 2017, 89, 1525-1527.	1.5	8
44	Ovarian cancer: Novel molecular aspects for clinical assessment. Critical Reviews in Oncology/Hematology, 2017, 117, 12-29.	2.0	25
45	Characterization of a Rare Nonpathogenic Methylenetetrahydrofolatereductase (MTHFR) Gene Mutation p.Lys215del in a Southern Italian family. Human Mutation, 2017, 38, 120-121.	1.1	2
46	Characterization of a Rare Nonpathogenic Sequence Variant (c.1905C>T) of the Dihydropyrimidine Dehydrogenase Gene (DPYD). International Journal of Biological Markers, 2017, 32, 357-360.	0.7	3
47	Sex-Genetic Interaction in the Risk for Cerebrovascular Disease. Current Medicinal Chemistry, 2017, 24, 2687-2699.	1.2	12
48	Desmoid Tumors in Familial Adenomatous Polyposis. Anticancer Research, 2017, 37, 3357-3366.	0.5	62
49	Next-generation Sequencing (NGS) Analysis on Single Circulating Tumor Cells (CTCs) with No Need of Whole-genome Amplification (WGA). Cancer Genomics and Proteomics, 2017, 14, 173-179.	1.0	29
50	Sirtuins and Cancer: Role in the Epithelial-Mesenchymal Transition. Oxidative Medicine and Cellular Longevity, 2016, 2016, 1-9.	1.9	62
51	VEGF gene promoter polymorphisms and risk of VTE in chemotherapy-treated cancer patients. Thrombosis and Haemostasis, 2016, 115, 143-151.	1.8	13
52	The role of epsilon PKC in acute and chronic diseases: Possible pharmacological implications of its modulators. Pharmacological Research, 2016, 111, 659-667.	3.1	10
53	Parallelism of DOG1 expression with recurrence risk in gastrointestinal stromal tumors bearing KIT or PDGFRA mutations. BMC Cancer, 2016, 16, 87.	1.1	20
54	O039. Case-control genetic association studies in migraine: a 7-year experience at the Interinstitutional Multidisciplinary Biobank (BioBIM) of IRCCS San Raffaele Pisana. Journal of Headache and Pain, 2015, 16, A120.	2.5	0

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55	Look beyond Catechol-O-Methyltransferase genotype for cathecolamines derangement in migraine: the BioBIM rs4818 and rs4680 polymorphisms study. Journal of Headache and Pain, 2015, 16, 520.	2.5	16
56	Sample PREanalytical Code for labeling of biospecimens: an analysis of specimen labeling protocols. Journal of Biorepository Science for Applied Medicine, 2015, , 15.	0.2	0
57	DNA fingerprinting for sample authentication in biobanking: recent perspectives. Journal of Biorepository Science for Applied Medicine, 2015, , 35.	0.2	1
58	Type 2 Diabetes and Breast Cancer: The Interplay between Impaired Glucose Metabolism and Oxidant Stress. Oxidative Medicine and Cellular Longevity, 2015, 2015, 1-10.	1.9	63
59	Comparison of therapy with Ticagrelor, Prasugrel or high Clopidogrel dose in PCI patients with high on treatment platelet reactivity and genotype variation. TRIPLETE RESET trial. International Journal of Cardiology, 2015, 194, 60-62.	0.8	16
60	The DOG1 scoring system in GIST: a novel factor for measurement of the recurrence risk. Annals of Oncology, 2015, 26, vi29.	0.6	0
61	Is <i>SOD2</i> Ala16Val Polymorphism Associated with Migraine with Aura Phenotype?. Antioxidants and Redox Signaling, 2015, 22, 275-279.	2.5	22
62	Impact of <i>VEGF</i> gene polymorphisms in elderly cancer patients: clinical outcome and toxicity. Pharmacogenomics, 2015, 16, 61-78.	0.6	12
63	Progesterone Receptor Gene (PROGINS) Polymorphism Correlates with Late Onset of Migraine. DNA and Cell Biology, 2015, 34, 208-212.	0.9	13
64	Common variation in PHACTR1 is associated with susceptibility to cervical artery dissection. Nature Genetics, 2015, 47, 78-83.	9.4	195
65	Role of Serum and Glucocorticoid-Inducible Kinase (SGK)-1 in Senescence: A Novel Molecular Target Against Age-Related Diseases. Current Medicinal Chemistry, 2015, 22, 3765-3788.	1.2	17
66	Ensuring Sample Quality for Biomarker Discovery Studies - Use of ICT Tools to Trace Biosample Life-cycle. Cancer Genomics and Proteomics, 2015, 12, 291-9.	1.0	6
67	Obesity and colorectal cancer: Role of adipokines in tumor initiation and progression. World Journal of Gastroenterology, 2014, 20, 5177.	1.4	157
68	Pharmacogenomics and pharmacogenetics of thiazolidinediones: role in diabetes and cardiovascular risk factors. Pharmacogenomics, 2014, 15, 2063-2082.	0.6	37
69	EHMTI-0241. Association between migraine and sod1 and sod2 genes polymorphisms: the biobim study. Journal of Headache and Pain, 2014, 15, .	2.5	0
70	Association between migraine and <i>ACE</i> gene (insertion/deletion) polymorphism: the BioBIM study. Pharmacogenomics, 2014, 15, 147-155.	0.6	10
71	Diagnostic Procedures for Paraffin-Embedded Tissues Analysis in Pharmacogenomic Studies. Methods in Molecular Biology, 2014, 1175, 45-65.	0.4	3
72	Plasma plasminogen activator inhibitor-1 (PAI-1) levels in breast cancer - relationship with clinical outcome. Anticancer Research, 2014, 34, 1153-61.	0.5	36

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73	Establishment of a biorepository for migraine research: the experience of Interinstitutional Multidisciplinary BioBank (BioBIM). Neurological Sciences, 2013, 34, 1659-1663.	0.9	14
74	An International Ki67 Reproducibility Study. Journal of the National Cancer Institute, 2013, 105, 1897-1906.	3.0	498
75	Peripheral CD45RO, PD-1, and TLR4 expression in metastatic colorectal cancer patients treated with bevacizumab, fluorouracil, and irinotecan (FOLFIRI-B). Medical Oncology, 2013, 30, 743.	1.2	20
76	Role of genetic polymorphisms of ion channels in the pathophysiology of coronary microvascular dysfunction and ischemic heart disease. Basic Research in Cardiology, 2013, 108, 387.	2.5	63
77	Heterogeneity of Triple-Negative Breast Cancer: Histologic Subtyping to Inform the Outcome. Clinical Breast Cancer, 2013, 13, 31-39.	1.1	128
78	Research Highlights: Highlights from the latest articles in stroke pharmacogenomics. Pharmacogenomics, 2013, 14, 13-16.	0.6	1
79	TNF-α gene promoter polymorphisms and risk of venous thromboembolism in gastrointestinal cancer patients undergoing chemotherapy. Annals of Oncology, 2013, 24, 2571-2575.	0.6	16
80	<scp>P</scp> rion Protein Gene <scp>M129V</scp> Polymorphism and Variability in Age at Migraine Onset. Headache, 2013, 53, 540-545.	1.8	7
81	Impact of chemotherapy on activated protein Câ€dependent thrombin generation—Association with VTE occurrence. International Journal of Cancer, 2013, 133, 1253-1258.	2.3	26
82	An importance of identification of double variant methylenetetrahydrofolate reductase gene C677T and A1298C in cis configuration for pharmacogenetic studies. Blood Coagulation and Fibrinolysis, 2013, 24, 784-786.	0.5	4
83	Age-related reduction of cerebral ischemic preconditioning: myth or reality?. Clinical Interventions in Aging, 2013, 8, 1055.	1.3	19
84	A Reliable and Reproducible Technique for DNA Fingerprinting in Biorepositories: A Pilot Study from BioBIM. International Journal of Biological Markers, 2013, 28, 398-404.	0.7	5
85	MDR1 and LPL genetic variants are associated with unfavorable outcome in stroke patients under atorvastatin treatment. Pharmacogenomics, 2013, 14, 14-5.	0.6	Ο
86	Polymorphisms on IL1B and VWF genes are associated with response to fibrinolysis in Hispanics with ischemic stroke. Pharmacogenomics, 2013, 14, 15-6.	0.6	0
87	Mutational analysis of gastrointestinal stromal tumors (GISTs): procedural approach for diagnostic purposes. Cancer Genomics and Proteomics, 2013, 10, 115-23.	1.0	5
88	Pharmacodynamic Effect of Switching Therapy in Patients With High On-Treatment Platelet Reactivity and Genotype Variation With High Clopidogrel Dose Versus Prasugrel. Circulation: Cardiovascular Interventions, 2012, 5, 698-704.	1.4	44
89	Brachyury, a Driver of the Epithelial–Mesenchymal Transition, Is Overexpressed in Human Lung Tumors: An Opportunity for Novel Interventions against Lung Cancer. Clinical Cancer Research, 2012, 18, 3868-3879.	3.2	112
90	Factor Seven Activating Protease Activity Levels in Women With Recurrent Pregnancy Loss. Reproductive Sciences, 2012, 19, 317-321.	1.1	2

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91	Angiogenesis and Hypertension: The Dual Role of Anti-Hypertensive and Anti-Angiogenic Therapies. Current Vascular Pharmacology, 2012, 10, 479-493.	0.8	26
92	SPRECware: Software Tools for Standard PREanalytical Code (SPREC) Labeling – Effective Exchange and Search of Stored Biospecimens. International Journal of Biological Markers, 2012, 27, 272-279.	0.7	19
93	Impact of Preanalytical Handling and Timing for Peripheral Blood Mononuclear Cells Isolation and RNA Studies: The Experience of the Interinstitutional Multidisciplinary BioBank (BioBIM). International Journal of Biological Markers, 2012, 27, 90-98.	0.7	21
94	730 Analysis of EMSY in Italian Male Breast Cancer Patients. European Journal of Cancer, 2012, 48, S173.	1.3	0
95	Impact of Statins on the Coagulation Status of Type 2 Diabetes Patients Evaluated by a Novel Thrombin-Generation Assay. Cardiovascular Drugs and Therapy, 2012, 26, 301-309.	1.3	12
96	Pharmacogenetics and pharmacogenomics: role of mutational analysis in anti-cancer targeted therapy. Pharmacogenomics Journal, 2012, 12, 277-286.	0.9	32
97	Association between increased tumor necrosis factor alpha levels and acquired activated protein C resistance in patients with metastatic colorectal cancer. International Journal of Colorectal Disease, 2012, 27, 1561-1567.	1.0	18
98	Genetics and genomics of ischemic tolerance: focus on cardiac and cerebral ischemic preconditioning. Pharmacogenomics, 2012, 13, 1741-1757.	0.6	34
99	Genetics of ischemic stroke, stroke-related risk factors, stroke precursors and treatments. Pharmacogenomics, 2012, 13, 595-613.	0.6	115
100	An AT-rich region in the APC gene may cause misinterpretation of familial adenomatous polyposis molecular screening. Human Mutation, 2012, 33, 895-898.	1.1	2
101	Concurrent mutation in exons 1 and 2 of the K-ras oncogene in colorectal cancer. Folia Histochemica Et Cytobiologica, 2012, 49, 729-733.	0.6	7
102	A comprehensive procedural approach to genotyping KRAS and BRAF from paraffin embedded tissues for diagnostic purposes. In Vivo, 2012, 26, 537-47.	0.6	3
103	PAI-1 4G/5G repeat is a target in gastric carcinomas with microsatellite instability. Digestive and Liver Disease, 2011, 43, 454-458.	0.4	0
104	RFID as a New ICT Tool to Monitor Specimen Life Cycle and Quality Control in a Biobank. International Journal of Biological Markers, 2011, 26, 129-135.	0.7	22
105	PRKCSH GAG trinucleotide repeat is a mutational target in gastric carcinomas with high-level microsatellite instability. Clinical Genetics, 2011, 79, 397-398.	1.0	3
106	Platinum-Based Compounds and Risk for Cardiovascular Toxicity in the Elderly: Role of the Antioxidants in Chemoprevention. Rejuvenation Research, 2011, 14, 293-308.	0.9	47
107	Predictive value of VEGF gene polymorphisms for metastatic colorectal cancer patients receiving first-line treatment including fluorouracil, irinotecan, and bevacizumab. International Journal of Colorectal Disease, 2011, 26, 143-151.	1.0	70
108	Importance of haplotype analysis in association studies considering VEGF promoter polymorphisms. Clinical Biochemistry, 2011, 44, 748.	0.8	0

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109	VEGF gene polymorphisms may be associated with an increased risk of fluorouracil-induced diarrhea. Annals of Oncology, 2011, 22, 1928-1929.	0.6	1
110	Preanalytical Procedures for DNA Studies: The Experience of the Interinstitutional Multidisciplinary BioBank (BioBIM). Biopreservation and Biobanking, 2011, 9, 35-45.	0.5	44
111	Serum sEâ€selectin levels and carcinoembryonic antigen mRNAâ€expressing cells in peripheral blood as prognostic factors in colorectal cancer patients. Cancer, 2010, 116, 2913-2921.	2.0	32
112	VEGF-A gene promoter polymorphisms and microvascular complications in patients with essential hypertension. Clinical Biochemistry, 2010, 43, 1090-1095.	0.8	17
113	TNFA Gene Promoter Polymorphisms and Susceptibility to Recurrent Pregnancy Loss in Italian Women. Reproductive Sciences, 2010, 17, 659-666.	1.1	21
114	Predictive value of thrombopath determination in women with infertility and pregnancy complications. Clinica Chimica Acta, 2010, 411, 37-42.	0.5	5
115	Association between Birt Hogg Dube syndrome and cancer predisposition. Anticancer Research, 2010, 30, 751-7.	0.5	18
116	Laparoscopic resection of sporadic synchronous gastric and jejunal gastrointestinal stromal tumors: Report of a case. Surgery Today, 2009, 39, 335-339.	0.7	11
117	Prognostic value of pre-surgical plasma PAI-1 (plasminogen activator inhibitor-1) levels in breast cancer. Thrombosis Research, 2009, 124, 403-408.	0.8	30
118	Chemoprevention of Colonic Cancer Is There a Foreseeable Future?. , 2009, , 77-93.		0
119	Prognostic significance of serum adipokine levels in colorectal cancer patients. Anticancer Research, 2009, 29, 3321-7.	0.5	56
120	A novel K-ras mutation in colorectal cancer. A case report and literature review. Anticancer Research, 2009, 29, 3369-74.	0.5	12
121	Survival of hereditary non-polyposis colorectal cancer patients compared with sporadic colorectal cancer patients. Journal of Experimental and Clinical Cancer Research, 2008, 27, 39.	3.5	37
122	Mismatch of neurophysiological findings in partial recovery of consciousness: a case report. Brain Injury, 2008, 22, 633-637.	0.6	8
123	Birt-Hogg-Dubé (BHD) syndrome: report of two novel germline mutations in the folliculin (FLCN) gene. European Journal of Dermatology, 2008, 18, 382-6.	0.3	18
124	Prognostic significance of interleukin-6 measurement in the diagnosis of acute myocardial infarction in emergency department. Clinica Chimica Acta, 2007, 381, 151-156.	0.5	11
125	PO-32 Plasminogen activator inhibitor-1 4G/5G polymorphism in breast cancer. Thrombosis Research, 2007, 120, S155.	0.8	0
126	543 POSTER Soluble E-selectin levels and CEA expressing blood-borne cells in colorectal cancer patients. A causal relationship?. European Journal of Cancer, Supplement, 2007, 5, 102.	2.2	0

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127	Prognostic significance of adiponectin levels in non-metastatic colorectal cancer. Anticancer Research, 2007, 27, 483-9.	0.5	41
128	Review. TNF/VEGF cross-talk in chronic inflammation-related cancer initiation and progression: an early target in anticancer therapeutic strategy. In Vivo, 2007, 21, 147-61.	0.6	47
129	Non-steroidal anti-inflammatory drugs in cancer prevention and therapy. Anticancer Research, 2007, 27, 3147-62.	0.5	31
130	Variation of the insulin receptor substrate gene (IRS-1) in African Pygmies and Bantus. Diabetes Research and Clinical Practice, 2006, 72, 108-109.	1.1	1
131	Prognostic Value of Carcinoembryonic Antigen and Vascular Endothelial Growth Factor Tumor Tissue Content in Colorectal Cancer. Oncology, 2006, 71, 176-184.	0.9	27
132	Mitochondrial DNA from Prehistoric Canids Highlights Relationships Between Dogs and South-East European Wolves. Molecular Biology and Evolution, 2005, 22, 2541-2551.	3.5	68
133	Soluble CD40 Ligand Plasma Levels in Lung Cancer. Clinical Cancer Research, 2004, 10, 610-614.	3.2	62
134	Prognostic value of soluble P-selectin levels in colorectal cancer. International Journal of Cancer, 2004, 111, 404-408.	2.3	37
135	Prognostic value of serum and tumor tissue CA 72-4 content in gastric cancer. International Journal of Biological Markers, 2003, 18, 21-27.	0.7	25
136	Correlations between Phenotype and Microsatellite Instability in HNPCC: Implications for Genetic Testing. Familial Cancer, 2002, 3, 117-121.	0.9	8
137	Taphonomy of the fossil hominid bones from the Acheulean site of Castel di Guido near Rome, Italy. Journal of Human Evolution, 2001, 41, 211-225.	1.3	15
138	APC gene mutations and colorectal adenomatosis in familial adenomatous polyposis. British Journal of Cancer, 2000, 82, 348-353.	2.9	101
139	Mutational analysis of OB gene in obese and type 2 diabetes affected subjects International Journal of Molecular Medicine, 2000, 6, 97-9.	1.8	3
140	Interaction between the G1057D variant of IRS-2 and overweight in the pathogenesis of type 2 diabetes. Human Molecular Genetics, 2000, 9, 2517-2521.	1.4	88
141	Microsatellite instability in thyroid tumours and tumour-like lesions. British Journal of Cancer, 1999, 79, 340-345.	2.9	35
142	Transcript dosage effect in familial adenomatous polyposis: Model offered by two kindreds with exon 9APC gene mutations. , 1998, 11, 197-201.		22
143	Transcripts with splicings of exons 15 and 16 of the hMLH1 gene in normal lymphocytes: implications in RNA-based mutation screening of hereditary non-polyposis colorectal cancer. European Journal of Cancer, 1998, 34, 927-930.	1.3	12
144	Optimized PCR labeling in mutational and microsatellite analysis. Clinical Chemistry, 1998, 44, 1381-1387.	1.5	7

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145	Origin and Gender Determination of Dried Blood on a Statue of the Virgin Mary. Journal of Forensic Sciences, 1998, 43, 431-434.	0.9	7
146	The 72-kDa and the 92-kDa gelatinases, but not their inhibitors TIMP-1 and TIMP-2, are expressed in early psoriatic lesions. Experimental Dermatology, 1997, 6, 321-327.	1.4	15
147	Use of a multiplex polymerase chain reaction assay in the sex typing of DNA extracted from archaeological bone. International Journal of Osteoarchaeology, 1997, 7, 605-609.	0.6	15
148	Microsatellite instability in sporadic mucinous colorectal carcinomas: relationship to clinico-pathological variables. , 1997, 182, 380-384.		72
149	Microsatellite instability in sporadic mucinous colorectal carcinomas: relationship to clinico-pathological variables. , 1997, 182, 380.		7
150	Genetic instability in early gastric cancer. Cancer Genetics and Cytogenetics, 1996, 91, 119.	1.0	0
151	Microsatellite alterations pathological aspects of breast cancer. Cancer Genetics and Cytogenetics, 1996, 91, 131.	1.0	0
152	Congenital hypertrophy of the retinal pigment epithelium in familial adenomatous polyposis: Novel criteria of assessment and correlations with constitutional adenomatous polyposis coli gene mutations. , 1996, 78, 2400-2410.		31
153	Multiplex PCR analysis and genotype-phenotype correlations of frequentAPC mutations. Human Mutation, 1995, 5, 144-152.	1.1	24
154	Microsatellite instability and pathological aspects of breast cancer. International Journal of Cancer, 1995, 64, 264-268.	2.3	50
155	Novel mutations and inactivation of both alleles of the APC gene in desmoid tumors. Human Molecular Genetics, 1995, 4, 1979-1981.	1.4	54
156	Alterations of DNA-repeats in human pancreatic cancer. Gastroenterology, 1995, 108, A461.	0.6	0
157	A novel deletion in exon 15 of the adenomatous polyposis coli gene in an Italian kindred. Human Mutation, 1994, 3, 301-304.	1.1	15
158	A novel mutation at the splice junction of exon 9 of the APC gene in familial adenomatous polyposis. Human Mutation, 1994, 3, 305-308.	1.1	11
159	Analysis of adenomatous polyposis coli gene in thyroid tumours. British Journal of Cancer, 1994, 70, 1085-1088.	2.9	47