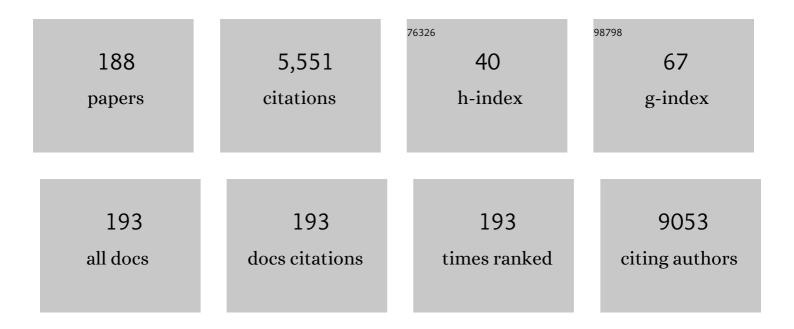
## **Gregory J Tsongalis**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Living the best of both worlds: A personal scientific journey. FASEB BioAdvances, 2022, 4, 95-101.	2.4	0
2	Plasmonic Nanoparticle Conjugation for Nucleic Acid Biosensing. Methods in Molecular Biology, 2022, 2393, 73-87.	0.9	0
3	DNA Genotyping As a Quality Assurance Measure in Surgical Pathology. Forensic Genomics, 2022, 2, 25-28.	0.5	0
4	Wastewater-Based SARS-CoV-2 Surveillance in Northern New England. Microbiology Spectrum, 2022, 10, e0220721.	3.0	8
5	Using Droplet Digital PCR to Detect Cyanobacteria in Human Lung Tissue. FASEB Journal, 2022, 36, .	0.5	0
6	Microsatellite Instability Testing for Lynch Syndrome in Colorectal Adenomas. FASEB Journal, 2022, 36,	0.5	0
7	Reducing dermal exposure to agrochemical carcinogens using a fluorescent dye-based intervention among subsistence farmers in rural Honduras. International Journal of Hygiene and Environmental Health, 2021, 234, 113734.	4.3	3
8	Evaluation of a Next-Generation Sequencing Metagenomics Assay to Detect and Quantify DNA Viruses in Plasma from Transplant Recipients. Journal of Molecular Diagnostics, 2021, 23, 719-731.	2.8	11
9	CRISPR-cas13 enzymology rapidly detects SARS-CoV-2 fragments in a clinical setting. Journal of Clinical Virology, 2021, 145, 105019.	3.1	18
10	Implementation of Reverse TranscriptaseÂPCR Testing for Severe Acute Respiratory Syndrome Coronavirus 2 under the US Food and Drug Administration Emergency Use Authorization. Clinical Chemistry, 2021, 67, 434-435.	3.2	9
11	Mixed Effects Machine Learning Models for Colon Cancer Metastasis Prediction using Spatially Localized Immuno-Oncology Markers. , 2021, , .		0
12	A case of molecularly confirmed <i>BAP1</i> inactivated melanocytic tumor with retention of immunohistochemical expression: A confounding factor. Journal of Cutaneous Pathology, 2020, 47, 485-489.	1.3	6
13	Maintaining Laboratory Services in a Rural Academic Medical Center During the Severe Acute Respiratory Syndrome Coronavirus 2 Pandemic. Advances in Molecular Pathology, 2020, 3, 5-11.	0.4	0
14	The 2020 Wild, Wild West of Diagnostics. Advances in Molecular Pathology, 2020, 3, 1-3.	0.4	0
15	Implementation of an Emergency Use Authorization Test During an Impending National Crisis. Journal of Molecular Diagnostics, 2020, 22, 844-846.	2.8	12
16	Molecular genetic profiling reveals novel association between FLT3 mutation and survival in glioma. Journal of Neuro-Oncology, 2020, 148, 473-480.	2.9	2
17	Comparison of Tissue Molecular Biomarker Testing Turnaround Times and Concordance Between Standard of Care and the Biocartis Idylla Platform in Patients With Colorectal Cancer. American Journal of Clinical Pathology, 2020, 154, 266-276.	0.7	10
18	HPV, vaccines, and cervical cancer in a low- and middle-income country. Current Problems in Cancer, 2020, 44, 100605.	2.0	6

#	Article	IF	CITATIONS
19	Feasibility of Brigade-Style, Multiphasic Cancer Screening in Rural Honduras. JCO Global Oncology, 2020, 6, 453-461.	1.8	3
20	Rapid EGFR mutation testing in lung cancer tissue samples using a fully automated system and single-use cartridge. Practical Laboratory Medicine, 2020, 20, e00156.	1.3	7
21	Molecular matching and treatment strategies for advanced stage lung cancer at Dartmouth-Hitchcock Medical Center: A three-year review of a Molecular Tumor Board. Practical Laboratory Medicine, 2020, 21, e00174.	1.3	1
22	Third-Generation Sequencing in the Clinical Laboratory: Exploring the Advantages and Challenges of Nanopore Sequencing. Journal of Clinical Microbiology, 2019, 58, .	3.9	146
23	BRAF V600E mutations are not an oncogenic driver of solitary xanthogranuloma and reticulohistiocytoma: Testing may be useful in screening for Erdheim-Chester disease. Experimental and Molecular Pathology, 2019, 111, 104320.	2.1	6
24	Design of peptide nucleic acid probes on plasmonic gold nanorods for detection of circulating tumor DNA point mutations. Biosensors and Bioelectronics, 2019, 130, 236-244.	10.1	56
25	Screening for Human Papillomavirus in a Low- and Middle-Income Country. Journal of Global Oncology, 2019, 5, JGO.18.00233.	0.5	6
26	Significant Expansion of Real-Time PCR Multiplexing with Traditional Chemistries using Amplitude Modulation. Scientific Reports, 2019, 9, 1053.	3.3	20
27	Ductal Carcinoma in Situ Biomarkers in a Precision Medicine Era. American Journal of Pathology, 2019, 189, 956-965.	3.8	15
28	Frequency of Somatic TP53 Mutations in Combination with Known Pathogenic Mutations in Colon Adenocarcinoma, Non–Small Cell Lung Carcinoma, and Gliomas as Identified by Next-Generation Sequencing. Neoplasia, 2018, 20, 256-262.	5.3	44
29	Rural distribution of human papilloma virus in low- and middle-income countries. Experimental and Molecular Pathology, 2018, 104, 146-150.	2.1	8
30	Implementation of Multicolor Melt Curve Analysis for High-Risk Human Papilloma Virus Detection in Low- and Middle-Income Countries: A Pilot Study for Expanded Cervical Cancer Screening in Honduras. Journal of Global Oncology, 2018, 4, 1-8.	0.5	6
31	Rapid Somatic Mutation Testing in Colorectal Cancer by Use of a Fully Automated System and Single-Use Cartridge: A Comparison with Next-Generation Sequencing. journal of applied laboratory medicine, The, 2018, 3, 178-184.	1.3	7
32	Maternally inherited 133kb deletion of 14q32 causing Kagami–Ogata syndrome. Journal of Human Genetics, 2018, 63, 1231-1239.	2.3	6
33	Pharmacogenetics of Opioid Use and Implications for Pain Management. journal of applied laboratory medicine, The, 2018, 2, 622-632.	1.3	9
34	Identifying aerosolized cyanobacteria in the human respiratory tract: A proposed mechanism for cyanotoxin-associated diseases. Science of the Total Environment, 2018, 645, 1003-1013.	8.0	44
35	Potential of STAT Somatic Mutation Testing at Resection. Clinical Chemistry, 2018, 64, 865-866.	3.2	4

36 Integrative Systems Biology. , 2018, , 205-215.

#	Article	IF	CITATIONS
37	Molecular Assessment of Human Diseases in the Clinical Laboratory. , 2018, , 709-730.		2
38	HS3ST1 genotype regulates antithrombin's inflammomodulatory tone and associates with atherosclerosis. Matrix Biology, 2017, 63, 69-90.	3.6	19
39	Variant call concordance between two laboratory-developed, solid tumor targeted genomic profiling assays using distinct workflows and sequencing instruments. Experimental and Molecular Pathology, 2017, 102, 215-218.	2.1	4
40	miRNA analysis in pancreatic cancer: the Dartmouth experience. Clinical Chemistry and Laboratory Medicine, 2017, 55, 755-762.	2.3	55
41	Not all good things come in big packages. Clinical Chemistry and Laboratory Medicine, 2017, 55, 605-607.	2.3	2
42	A Phase II Trial of Dovitinib in BCG-Unresponsive Urothelial Carcinoma with <i>FGFR3</i> Mutations or Overexpression: Hoosier Cancer Research Network Trial HCRN 12-157. Clinical Cancer Research, 2017, 23, 3003-3011.	7.0	59
43	Immune modulation associated with vascular endothelial growth factor (VEGF) blockade in patients with glioblastoma. Cancer Immunology, Immunotherapy, 2017, 66, 379-389.	4.2	20
44	Somatic mutation analysis in melanoma using targeted next generation sequencing. Experimental and Molecular Pathology, 2017, 103, 172-177.	2.1	19
45	The Case for Laboratory Developed Procedures. Academic Pathology, 2017, 4, 2374289517708309.	1.1	24
46	Somatic Mutation Analysis of Human Cancers: Challenges in Clinical Practice. Journal of Clinical Pharmacology, 2017, 57, S60-S66.	2.0	4
47	Development of HLA-B*57:01 Genotyping Real-Time PCR with Optimized Hydrolysis Probe Design. Journal of Molecular Diagnostics, 2017, 19, 742-754.	2.8	7
48	Triple-Negative Breast Cancer. American Journal of Pathology, 2017, 187, 2133-2138.	3.8	60
49	Improving Adequacy of Small Biopsy and Fine-Needle Aspiration Specimens for Molecular Testing by Next-Generation Sequencing in Patients With Lung Cancer: A Quality Improvement Study at Dartmouth-Hitchcock Medical Center. Archives of Pathology and Laboratory Medicine, 2017, 141, 402-409.	2.5	33
50	Hyper-Methylated Loci Persisting from Sessile Serrated Polyps to Serrated Cancers. International Journal of Molecular Sciences, 2017, 18, 535.	4.1	33
51	Use of Biosynthetic Controls as Performance Standards for Next-Generation Sequencing Assays of Somatic Tumors: A Multilaboratory Study. journal of applied laboratory medicine, The, 2017, 2, 138-149.	1.3	Ο
52	Personalized Medicine for the Treatment of Human Cancer. , 2017, , 843-855.		0
53	The Use of Targeted Therapies for Precision Medicine in Oncology. Clinical Chemistry, 2016, 62, 1556-1564.	3.2	10
54	Somatic gene mutation analysis of triple negative breast cancers. Breast, 2016, 29, 202-207.	2.2	23

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55	An analysis of human papillomavirus testing and endocervical component on pap tests: A pilot study using the Roche Cobas <sup>®</sup> assay. Diagnostic Cytopathology, 2016, 44, 280-282.	1.0	0
56	Targeted next-generation sequencing detects a high frequency of potentially actionable mutations in metastatic breast cancers. Experimental and Molecular Pathology, 2016, 100, 421-425.	2.1	28
57	Detection ofCALRMutation in Clonal and Nonclonal Hematologic Diseases Using Fragment Analysis and Next-Generation Sequencing. American Journal of Clinical Pathology, 2016, 146, 448-455.	0.7	12
58	Clinical Genotyping of Non–Small Cell Lung Cancers Using Targeted Next-Generation Sequencing: Utility of Identifying Rare and Co-mutations in Oncogenic Driver Genes. Neoplasia, 2016, 18, 577-583.	5.3	19
59	Role of microRNAs in regulation of the TNF/TNFR gene superfamily in chronic lymphocytic leukemia. Clinical Biochemistry, 2016, 49, 1307-1310.	1.9	24
60	Clinical Trials in Precision Oncology. Clinical Chemistry, 2016, 62, 442-448.	3.2	8
61	The potential utility of re-mining results of somatic mutation testing: KRAS status in lung adenocarcinoma. Cancer Genetics, 2016, 209, 195-198.	0.4	77
62	Effective quality management practices in routine clinical next-generation sequencing. Clinical Chemistry and Laboratory Medicine, 2016, 54, 761-71.	2.3	22
63	Molecular Genetic Analysis of Ovarian Brenner Tumors and Associated Mucinous Epithelial Neoplasms. American Journal of Pathology, 2016, 186, 671-677.	3.8	40
64	The Pitfalls of Companion Diagnostics. Journal of Molecular Diagnostics, 2016, 18, 331-335.	2.8	1
65	Genomic characterization of patient-derived xenograft models established from fine needle aspirate biopsies of a primary pancreatic ductal adenocarcinoma and from patient-matched metastatic sites. Oncotarget, 2016, 7, 17087-17102.	1.8	40
66	Simultaneous extraction of DNA and total RNA from varying specimen types to enhance tissue utilization for molecular analysis Journal of Clinical Oncology, 2016, 34, e23209-e23209.	1.6	0
67	Rat-bite fever: An uncommon cause of fever and rash in a 9-year-old patient. JAAD Case Reports, 2015, 1, 371-374.	0.8	8
68	Rapid fluorescence in situ hybridisation (FISH) for <i>HER2 (ERBB2)</i> assessment in breast and gastro-oesophageal cancer. Journal of Clinical Pathology, 2015, 68, 306-308.	2.0	6
69	Variation in pre-PCR processing of FFPE samples leads to discrepancies in <i>BRAF</i> and <i>EGFR</i> mutation detection: a diagnostic RING trial. Journal of Clinical Pathology, 2015, 68, 111-118.	2.0	34
70	A multiplex PCR assay for the simultaneous detection of Chlamydia trachomatis, Neisseria gonorrhoeae, and Trichomonas vaginalis. Experimental and Molecular Pathology, 2015, 98, 214-218.	2.1	22
71	Regulatory T cells are not a strong predictor of survival for patients with glioblastoma. Neuro-Oncology, 2015, 17, 801-809.	1.2	43
72	Molecular profiling of intrahepatic and extrahepatic cholangiocarcinoma using next generation sequencing. Experimental and Molecular Pathology, 2015, 99, 240-244.	2.1	39

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73	A 78-Year-Old Woman with Brain Metastases. Clinical Chemistry, 2015, 61, 584-586.	3.2	1
74	Implementation of a Molecular Tumor Board: The Impact on Treatment Decisions for 35 Patients Evaluated at Dartmouth-Hitchcock Medical Center. Oncologist, 2015, 20, 1011-1018.	3.7	81
75	Potential driver mutations in ovarian Brenner tumors Journal of Clinical Oncology, 2015, 33, e22087-e22087.	1.6	0
76	Democratizing Molecular Diagnostics for the Developing World. American Journal of Clinical Pathology, 2014, 141, 17-24.	0.7	42
77	Lynch Syndrome Presenting as Endometrial Cancer. Clinical Chemistry, 2014, 60, 111-121.	3.2	32
78	Distinct patterns of DNA methylation in conventional adenomas involving the right and left colon. Modern Pathology, 2014, 27, 145-155.	5.5	40
79	Validation of interleukin 28B genotyping assay for clinical use. Clinical Biochemistry, 2014, 47, 478-480.	1.9	3
80	A MicroRNA-Based Test Improves Endoscopic Ultrasound–Guided Cytologic Diagnosis of Pancreatic Cancer. Clinical Gastroenterology and Hepatology, 2014, 12, 1717-1723.	4.4	34
81	Development of a rapid clinical TPMT genotyping assay. Clinical Biochemistry, 2014, 47, 126-129.	1.9	15
82	Framework for development of physician competencies in genomic medicine: report of the Competencies Working Group of the Inter-Society Coordinating Committee for Physician Education in Genomics. Genetics in Medicine, 2014, 16, 804-809.	2.4	123
83	Automated processing of fluorescence in-situ hybridization slides for HER2 testing in breast and gastro-esophageal carcinomas. Experimental and Molecular Pathology, 2014, 97, 116-119.	2.1	7
84	A rapid RT-PCR assay for the detection of HIV-1 in human plasma specimens. Experimental and Molecular Pathology, 2014, 97, 111-115.	2.1	5
85	Determining methylation status of methylguanine DNA methyl transferase (MGMT) from formalin-fixed, paraffin embedded tumor tissue. MethodsX, 2014, 1, 42-48.	1.6	2
86	Biomarker Testing for Breast, Lung, and Gastroesophageal Cancers at NCI Designated Cancer Centers. Journal of the National Cancer Institute, 2014, 106, .	6.3	18
87	Molecular Profiling of Appendiceal Epithelial Tumors Using Massively Parallel Sequencing to Identify Somatic Mutations. Clinical Chemistry, 2014, 60, 1004-1011.	3.2	80
88	Routine use of the Ion Torrent AmpliSeqâ,,¢ Cancer Hotspot Panel for identification of clinically actionable somatic mutations. Clinical Chemistry and Laboratory Medicine, 2014, 52, 707-14.	2.3	123
89	Personalized therapy for breast cancer. Clinical Genetics, 2014, 86, 62-67.	2.0	52
90	Effects of dose reduction on gemcitabine-based neoadjuvant chemoradiotherapy for localized pancreatic cancer Journal of Clinical Oncology, 2014, 32, e15262-e15262.	1.6	0

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91	Prognostic value of serum carbohydrate 19-9 in patients receiving gemcitabine-based neoadjuvant therapy for pancreatic cancer Journal of Clinical Oncology, 2014, 32, e15189-e15189.	1.6	1
92	The Emerging Role of the Molecular Diagnostics Laboratory in Breast Cancer Personalized Medicine. American Journal of Pathology, 2013, 183, 1075-1083.	3.8	37
93	MicroRNAs as diagnostic markers for pancreatic ductal adenocarcinoma and its precursor, pancreatic intraepithelial neoplasm. Cancer Genetics, 2013, 206, 217-221.	0.4	69
94	Epidermal growth factor receptor tyrosine kinase inhibitors as initial therapy for non-small cell lung cancer: Focus on epidermal growth factor receptor mutation testing and mutation-positive patients. Cancer Treatment Reviews, 2013, 39, 839-850.	7.7	100
95	Clinical utility of miRNAs in diagnosis and prognosis. Clinical Biochemistry, 2013, 46, 839.	1.9	1
96	Pancreatic Cyst Prevalence and the Risk of Mucin-Producing Adenocarcinoma in US Adults. American Journal of Gastroenterology, 2013, 108, 1546-1550.	0.4	115
97	A clinical PCR fragment analysis assay for TA repeat sizing in the UGT1A1 promoter region. Clinica Chimica Acta, 2013, 422, 1-4.	1.1	7
98	Validation of a solid-phase electrochemical array for genotyping hepatitis C virus. Experimental and Molecular Pathology, 2013, 95, 18-22.	2.1	19
99	Bioinformatics: What the Clinical Laboratorian Needs to Know and Prepare For. Clinical Chemistry, 2013, 59, 1301-1305.	3.2	4
100	A Comprehensive Assay for CFTR Mutational Analysis Using Next-Generation Sequencing. Clinical Chemistry, 2013, 59, 1481-1488.	3.2	44
101	A multiplex real-time polymerase chain reaction assay with two internal controls for the detection of <i>Brucella</i> species in tissues, blood, and feces from marine mammals. Journal of Veterinary Diagnostic Investigation, 2013, 25, 72-81.	1.1	19
102	MicroRNA Analysis: Is It Ready for Prime Time?. Clinical Chemistry, 2013, 59, 343-347.	3.2	10
103	Correlation Between Red Blood Cell Survival and Cytochrome P450 1A2 Enzyme Activity. Blood, 2013, 122, 3658-3658.	1.4	4
104	Are oncologists involved in cancer biomarker decisions at their institutions?. Journal of Clinical Oncology, 2013, 31, 6617-6617.	1.6	1
105	What are NCI-designated cancer centers using for gastric and esophageal cancer HER2 testing?. Journal of Clinical Oncology, 2013, 31, e15010-e15010.	1.6	3
106	Biomarker testing methods in breast, gastric, and lung cancers: A benchmarking survey of NCI cancer centers Journal of Clinical Oncology, 2013, 31, e22093-e22093.	1.6	2
107	Clinical and Genetic Determinants of Blood Pressure Under Ambulatory Conditions on Days With and Without Acute Exercise. FASEB Journal, 2013, 27, 910.15.	0.5	0
108	Microrna Expression In Patients With Myelodysplastic Syndromes Treated With Demethylating Agents. Blood, 2013, 122, 3758-3758.	1.4	0

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109	Polymorphisms in the Brain-Derived Neurotrophic Factor Gene Influence Memory and Processing Speed One Month after Brain Injury. Journal of Neurotrauma, 2012, 29, 1111-1118.	3.4	72
110	The human epidermal growth factor receptor 2 (HER2). Clinical Chemistry and Laboratory Medicine, 2012, 50, 23-30.	2.3	21
111	Evaluating the thermostability of commercial fast real-time PCR master mixes. Experimental and Molecular Pathology, 2012, 93, 261-263.	2.1	4
112	Evaluation of the Cobas 4800 HPV Test for Detecting High-Risk Human Papilloma-Virus in Cervical Cytology Specimens. Pathogens, 2012, 1, 30-36.	2.8	6
113	KRAS Detection in Colonic Tumors by DNA Extraction From FTA Paper. Diagnostic Molecular Pathology, 2011, 20, 189-193.	2.1	10
114	Cancer and Leukemia Group B Pathology Committee Guidelines for Tissue Microarray Construction Representing Multicenter Prospective Clinical Trial Tissues. Journal of Clinical Oncology, 2011, 29, 2282-2290.	1.6	25
115	Bexarotene Plus Erlotinib Suppress Lung Carcinogenesis Independent of <i>KRAS</i> Mutations in Two Clinical Trials and Transgenic Models. Cancer Prevention Research, 2011, 4, 818-828.	1.5	50
116	COMT Val158Met Genotype and Individual Differences in Executive Function in Healthy Adults. Journal of the International Neuropsychological Society, 2011, 17, 174-180.	1.8	70
117	Multilaboratory Evaluation of Real-Time PCR Tests for Hepatitis B Virus DNA Quantification. Journal of Clinical Microbiology, 2011, 49, 2854-2858.	3.9	51
118	Primary Melanoma of the Spinal Cord: A Case Report, Molecular Footprint, and Review of the Literature. Journal of Clinical Oncology, 2011, 29, e499-e502.	1.6	22
119	A validation study of a new molecular diagnostic assay: The Dartmouth-Hitchcock Medical Center experience with the GeneSearchâ,,¢ BLN assay in breast sentinel lymph nodes. Experimental and Molecular Pathology, 2010, 88, 1-6.	2.1	13
120	Endothelial nitric oxide gene polymorphisms, nitric oxide production and coronary artery disease risk in a South Indian population. Experimental and Molecular Pathology, 2010, 89, 205-208.	2.1	48
121	Molecular diagnostics: parallels between infectious disease and emerging oncology testing. Expert Opinion on Medical Diagnostics, 2010, 4, 185-188.	1.6	1
122	MicroRNA-21 Is Induced Early in Pancreatic Ductal Adenocarcinoma Precursor Lesions. Clinical Chemistry, 2010, 56, 603-612.	3.2	197
123	The Silencing of MicroRNA 148a Production by DNA Hypermethylation Is an Early Event in Pancreatic Carcinogenesis. Clinical Chemistry, 2010, 56, 1107-1118.	3.2	139
124	The additive blood pressure lowering effects of exercise intensity on post-exercise hypotension. American Heart Journal, 2010, 160, 513-520.	2.7	120
125	Molecular Assessment of Human Disease in the Clinical Laboratory. , 2010, , 413-420.		0
126	Molecular Determination of Primary Versus Metastatic Squamous Cell Carcinoma (SCC) of the Lung in the Context of SCC of the Head and Neck (H/N). FASEB Journal, 2010, 24, lb443.	0.5	0

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127	A Standardized BCR-ABL Monitoring Test: Assessment of Potential Adoption Impacts In Healthcare In the United States. Blood, 2010, 116, 4754-4754.	1.4	21
128	Use of a Linear Array for the Detection of Human Papillomavirus Genotypes in Head and Neck Cancer. Archives of Pathology and Laboratory Medicine, 2010, 134, 1813-1817.	2.5	17
129	Warfarin genotyping using three different platforms. American Journal of Translational Research (discontinued), 2010, 2, 441-6.	0.0	10
130	Evaluation of the Nanosphere Verigene® System and the Verigene® F5/F2/MTHFR Nucleic Acid Tests. Experimental and Molecular Pathology, 2009, 87, 105-108.	2.1	29
131	MicroRNAs: Novel Biomarkers for Human Cancer. Clinical Chemistry, 2009, 55, 623-631.	3.2	485
132	The endothelial nitric oxide synthase â~'786 T>C polymorphism and the exercise-induced blood pressure and nitric oxide responses among men with elevated blood pressure. Atherosclerosis, 2009, 204, e28-e34.	0.8	38
133	Molecular Assessment of Human Disease in the Clinical Laboratory. , 2009, , 605-612.		0
134	Pharmacogenomics and Personalized Medicine in the Treatment of Human Diseases. , 2009, , 613-622.		0
135	+ACA BRCA1 promoter polymorphism genotypic frequency among unaffected individuals and breast disease patients. FASEB Journal, 2009, 23, 925.9.	0.5	0
136	Peripheral Blood Chimerism Can Replace Marrow Chimerism Analyses Following Adult Allogeneic Stem Cell Transplant Blood, 2009, 114, 4316-4316.	1.4	0
137	Analysis of MicroRNAs in Pancreatic Fine-Needle Aspirates Can Classify Benign and Malignant Tissues. Clinical Chemistry, 2008, 54, 1716-1724.	3.2	194
138	Single nucleotide polymorphisms in ANKK1 and the dopamine D2 receptor gene affect cognitive outcome shortly after traumatic brain injury: A replication and extension study. Brain Injury, 2008, 22, 705-714.	1.2	75
139	Expression of BRCA1 and FAC1 in primary sporadic breast cancers FASEB Journal, 2008, 22, 898.26.	0.5	0
140	Global histone H3 lysine 4 methylation patterns in human breast cancer. FASEB Journal, 2008, 22, 470.1.	0.5	0
141	Transgenic cyclin E triggers dysplasia and multiple pulmonary adenocarcinomas. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 4089-4094.	7.1	73
142	Pharmacogenetics: where are we with respect to personalized medicine?. Expert Opinion on Medical Diagnostics, 2007, 1, 117-128.	1.6	0
143	Development of an Integrated Assay for Detection of BCR-ABL RNA. Clinical Chemistry, 2007, 53, 1593-1600.	3.2	47
144	Validation of a CYP2D6 Genotyping Panel on the NanoChip Molecular Biology Workstation. Clinical Chemistry, 2007, 53, 823-828.	3.2	11

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145	Dietary calcium intake and renin angiotensin system polymorphisms alter the blood pressure response to aerobic exercise: a randomized control design. Nutrition and Metabolism, 2007, 4, 1.	3.0	68
146	Search for evidence of recurring or persistent viruses in Crohn's disease. Apmis, 2007, 115, 962-968.	2.0	22
147	MTHFR Gene polymorphisms, B-vitamins and hyperhomocystinemia in young and middle-aged acute myocardial infarction patients. Experimental and Molecular Pathology, 2007, 82, 227-233.	2.1	22
148	Quantitative Assessment of the BCR-ABL Transcript Using the Cepheid Xpert BCR-ABL Monitor Assay. Archives of Pathology and Laboratory Medicine, 2007, 131, 947-950.	2.5	17
149	Apolipoprotein A1 genotype affects the change in high density lipoprotein cholesterol subfractions with exercise training. Atherosclerosis, 2006, 185, 65-69.	0.8	42
150	The effect of apolipoprotein E genotype on serum lipoprotein particle response to exercise. Atherosclerosis, 2006, 188, 126-133.	0.8	25
151	Negative urine opioid screening caused by rifampin-mediated induction of oxycodone hepatic metabolism. Clinica Chimica Acta, 2006, 367, 196-200.	1.1	36
152	Molecular diagnostics: A historical perspective. Clinica Chimica Acta, 2006, 369, 188-192.	1.1	17
153	Clinical genotyping: The need for interrogation of single nucleotide polymorphisms and mutations in the clinical laboratory. Clinica Chimica Acta, 2006, 363, 127-137.	1.1	20
154	Angiotensin-Converting Enzyme Genotype and Adherence to Aerobic Exercise Training. Preventive Cardiology, 2006, 9, 21-24.	1.1	18
155	RAAS polymorphisms alter the acute blood pressure response to aerobic exercise among men with hypertension. European Journal of Applied Physiology, 2006, 97, 26-33.	2.5	54
156	Branched DNA Technology in Molecular Diagnostics. American Journal of Clinical Pathology, 2006, 126, 448-453.	0.7	78
157	<i>BRCA1</i> and <i>BRCA2</i> Mutation Screening Using SmartCycler II High-Resolution Melt Curve Analysis. Archives of Pathology and Laboratory Medicine, 2006, 130, 185-187.	2.5	31
158	Genetic testing: current and future trends. Medical Laboratory Observer, 2006, 38, 42, 44.	0.1	0
159	Parameters involved in the conversion of real-time PCR assays from the ABI prism 7700 to the Cepheid SmartCycler® II. Clinical Biochemistry, 2005, 38, 183-186.	1.9	8
160	Prevalence of the Factor V G1691A and the Factor II/prothrombin G20210A gene polymorphisms among Tamilians. Experimental and Molecular Pathology, 2005, 79, 9-13.	2.1	24
161	DNA STAT: the ability to perform clinical analysis in real time. Expert Review of Molecular Diagnostics, 2005, 5, 627-628.	3.1	2
162	Characterization of " Candidatus Piscichlamydia salmonis―(Order Chlamydiales ), a Chlamydia-Like Bacterium Associated With Epitheliocystis in Farmed Atlantic Salmon ( Salmo salar ). Journal of Clinical Microbiology, 2004, 42, 5286-5297.	3.9	126

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163	Prevalence of MTHFR gene polymorphisms (C677T and A1298C) among Tamilians. Experimental and Molecular Pathology, 2004, 77, 85-88.	2.1	53
164	Apolipoprotein e genotype and changes in serum lipids and maximal oxygen uptake with exercise training. Metabolism: Clinical and Experimental, 2004, 53, 193-202.	3.4	70
165	Breast cancer as a model of realistic challenges in pharmacogenomics. Clinical Biochemistry, 2003, 36, 89-94.	1.9	6
166	A reality check for molecular diagnostics in clinical practice. Pharmacogenomics, 2003, 4, 667-668.	1.3	3
167	A novel mutation in exon 5 of the ALAS2 gene results in X-linked sideroblastic anemia. Clinica Chimica Acta, 2002, 321, 49-53.	1.1	7
168	HER2: The Neu Prognostic Marker for Breast Cancer. Critical Reviews in Clinical Laboratory Sciences, 2001, 38, 167-182.	6.1	11
169	Correlation of polymorphisms to coagulation and biochemical risk factors for cardiovascular diseases. American Journal of Cardiology, 2001, 87, 1361-1366.	1.6	116
170	Single nucleotide polymorphisms in the methylenetetrahydrofolate reductase gene are common in US Caucasian and Hispanic American populations. International Journal of Molecular Medicine, 2001, 8, 509-11.	4.0	28
171	<i>Mycobacterium Marinum</i> Dermatitis and Panniculitis with Chronic Pleuritis in a Captive White Whale ( <i>Delphinapterus Leucas</i> ) with Aortic Rupture. Journal of Veterinary Diagnostic Investigation, 2001, 13, 524-530.	1.1	18
172	Changes in Academic Productivity: Implications for Clinical Laboratory Research and Development. Clinical Chemistry, 2000, 46, 303-305.	3.2	0
173	p185HER2Overexpression in Human Breast Cancer Using Molecular and Immunohistochemical Methods. Cancer Investigation, 2000, 18, 336-342.	1.3	18
174	Rapid Detection of HSV from Cytologic Specimens Collected into ThinPrep Fixative. Acta Cytologica, 1999, 43, 1034-1038.	1.3	17
175	Polymorphisms in the Genes for Coagulation Factors II, V, and VII in Patients With Ischemic Heart Disease. Archives of Pathology and Laboratory Medicine, 1999, 123, 1230-1235.	2.5	19
176	Molecular Oncology: Diagnostic and Prognostic Assessment of Human Cancers in the Clinical Laboratory. Cancer Investigation, 1998, 16, 485-502.	1.3	5
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