

# Årika C Pavarino

## List of Publications by Year in descending order

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97  
papers

1,787  
citations

331670

21  
h-index

361022

35  
g-index

100  
all docs

100  
docs citations

100  
times ranked

2601  
citing authors

#	ARTICLE	IF	CITATIONS
1	Differential microRNA expression profile in blood of children with Down syndrome suggests a role in immunological dysfunction. <i>Human Cell</i> , 2022, 35, 639-648.	2.7	2
2	MiR-612, miR-637, and miR-874 can Regulate VEGFA Expression in Hepatocellular Carcinoma Cell Lines. <i>Genes</i> , 2022, 13, 282.	2.4	1
3	Regulation of VEGFA, KRAS, and NFE2L2 Oncogenes by MicroRNAs in Head and Neck Cancer. <i>International Journal of Molecular Sciences</i> , 2022, 23, 7483.	4.1	5
4	Alzheimer's Disease in the Down Syndrome: An Overview of Genetics and Molecular Aspects. <i>Neurology India</i> , 2021, 69, 32.	0.4	3
5	Polymorphisms in xenobiotic metabolism-related genes in patients with hepatocellular carcinoma: a case-control study. <i>Xenobiotica</i> , 2021, 51, 1-9.	1.1	5
6	Evaluation of molecular markers GSTM1 and GSTT1 and clinical factors in breast cancer: case-control study and literature review. <i>Xenobiotica</i> , 2021, 51, 1326-1334.	1.1	4
7	One-carbon metabolism and global DNA methylation in mothers of individuals with Down syndrome. <i>Human Cell</i> , 2021, 34, 1671-1681.	2.7	3
8	Association between folate metabolism polymorphisms and breast cancer: a case-control study. <i>Genetics and Molecular Biology</i> , 2021, 44, e20200485.	1.3	4
9	Psychosocial and Motor Characteristics of Patients With Hypermobility. <i>Frontiers in Psychiatry</i> , 2021, 12, 787822.	2.6	4
10	Role of Tropomyosin-related kinase B receptor and brain-derived neurotrophic factor in cancer. <i>Cytokine</i> , 2020, 136, 155270.	3.2	15
11	MicroRNAs as regulators of VEGFA and NFE2L2 in cancer. <i>Gene</i> , 2020, 759, 144994.	2.2	21
12	Differential expression of angiogenesis-related miRNAs and VEGFA in cirrhosis and hepatocellular carcinoma. <i>Archives of Medical Science</i> , 2020, 16, 1150-1157.	0.9	27
13	VEGFA and NFE2L2 Gene Expression and Regulation by MicroRNAs in Thyroid Papillary Cancer and Colloid Goiter. <i>Genes</i> , 2020, 11, 954.	2.4	18
14	Trends and predictions for survival and mortality in individuals with Down syndrome in Brazil: A 21-year analysis. <i>Journal of Intellectual Disability Research</i> , 2020, 64, 551-560.	2.0	5
15	Glutathione S-transferase Polymorphisms in Head and Neck Squamous Cell Carcinoma Treated with Chemotherapy and/or Radiotherapy. <i>Asian Pacific Journal of Cancer Prevention</i> , 2020, 21, 1637-1644.	1.2	5
16	Gene Polymorphisms Involved in Folate Metabolism and DNA Methylation with the Risk of Head and Neck Cancer. <i>Asian Pacific Journal of Cancer Prevention</i> , 2020, 21, 3751-3759.	1.2	8
17	Vitamin D3 increases the Caspase-3 p12, MTHFR, and P-glycoprotein reducing amyloid- $\beta$ 42 in the kidney of a mouse model for Down syndrome. <i>Life Sciences</i> , 2019, 231, 116537.	4.3	4
18	Polymorphisms in MTHFR, MTR, RFC1 and C $\beta$ YS genes involved in folate metabolism and thyroid cancer: a case-control study. <i>Archives of Medical Science</i> , 2019, 15, 522-530.	0.9	14

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19	Molecular evaluation of glutathione S transferase family genes in patients with sporadic colorectal cancer. <i>World Journal of Gastroenterology</i> , 2018, 24, 4462-4471.	3.3	12
20	Differential Expression of Prostaglandin I2 Synthase Associated with Arachidonic Acid Pathway in the Oral Squamous Cell Carcinoma. <i>Journal of Oncology</i> , 2018, 2018, 1-13.	1.3	10
21	Interleukin 6 and 10 Serum Levels and Genetic Polymorphisms in Children with Down Syndrome. <i>Mediators of Inflammation</i> , 2018, 2018, 1-9.	3.0	5
22	Candidate Biomarkers for Oral Squamous Cell Carcinoma: Differential Expression of Oxidative Stress-Related Genes. <i>Asian Pacific Journal of Cancer Prevention</i> , 2018, 19, 1343-1349.	1.2	20
23	Relationship between CD44/CD133/CD117 cancer stem cells phenotype and Cetuximab and Paclitaxel treatment response in head and neck cancer cell lines. <i>American Journal of Cancer Research</i> , 2018, 8, 1633-1641.	1.4	10
24	Clinical, Epidemiological and Histopathological Aspects in Patients with Hepatocellular Carcinoma Undergoing Liver Transplantation. <i>Asian Pacific Journal of Cancer Prevention</i> , 2018, 19, 2795-2802.	1.2	5
25	Research Article Polymorphisms of interleukin 6 in Down syndrome individuals: a case-control study.. <i>Genetics and Molecular Research</i> , 2017, 16, .	0.2	1
26	Hepatocellular Carcinoma: a Comprehensive Review of Biomarkers, Clinical Aspects, and Therapy. <i>Asian Pacific Journal of Cancer Prevention</i> , 2017, 18, 863-872.	1.2	62
27	Differential Expression of Inflammation-Related Genes in Children with Down Syndrome. <i>Mediators of Inflammation</i> , 2016, 2016, 1-8.	3.0	12
28	Role of MTHFR C677T and MTR A2756G polymorphisms in thyroid and breast cancer development. <i>Genetics and Molecular Research</i> , 2016, 15, .	0.2	17
29	Trisomy 21 Alters DNA Methylation in Parent-of-Origin-Dependent and -Independent Manners. <i>PLoS ONE</i> , 2016, 11, e0154108.	2.5	52
30	Variables associated to fetal microchimerism in systemic lupus erythematosus patients. <i>Clinical Rheumatology</i> , 2016, 35, 107-111.	2.2	9
31	<i>CYP1A1</i> , <i>CYP2E1</i> and <i>EPHX1</i> polymorphisms in sporadic colorectal neoplasms. <i>World Journal of Gastroenterology</i> , 2016, 22, 9974.	3.3	16
32	Polymorphisms of folate metabolism genes in patients with cirrhosis and hepatocellular carcinoma. <i>World Journal of Hepatology</i> , 2016, 8, 1234.	2.0	18
33	A case-control study of <i>CYP2E1</i> (PstI) and <i>CYP1A1</i> (MspI) polymorphisms in colorectal cancer. <i>Genetics and Molecular Research</i> , 2015, 14, 17856-17863.	0.2	4
34	Neurofibromatosis: part 2 – clinical management. <i>Arquivos De Neuro-Psiquiatria</i> , 2015, 73, 531-543.	0.8	10
35	Is Magnetic Resonance Spectroscopy Capable of Detecting Metabolic Abnormalities in Neurofibromatosis Type 1 That Are Not Revealed in Brain Parenchyma of Normal Appearance?. <i>Pediatric Neurology</i> , 2015, 52, 314-319.	2.1	8
36	Influence of functional polymorphisms in TNF- $\alpha$ , IL-8, and IL-10 cytokine genes on mRNA expression levels and risk of gastric cancer. <i>Tumor Biology</i> , 2015, 36, 9159-9170.	1.8	58

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37	<i>TLR2</i> and <i>TLR4</i> polymorphisms influence mRNA and protein expression in colorectal cancer. <i>World Journal of Gastroenterology</i> , 2015, 21, 7730.	3.3	31
38	Neurofibromatoses: part 1 ? diagnosis and differential diagnosis. <i>Arquivos De Neuro-Psiquiatria</i> , 2014, 72, 241-250.	0.8	27
39	Genetic Polymorphisms Involved in Folate Metabolism and Maternal Risk for Down Syndrome: A Meta-Analysis. <i>Disease Markers</i> , 2014, 2014, 1-12.	1.3	18
40	DNMT3B C46359T and SHMT1 C1420T polymorphisms in the folate pathway in carcinogenesis of head and neck. <i>Molecular Biology Reports</i> , 2014, 41, 581-589.	2.3	17
41	Meta-analysis of Methylenetetrahydrofolate reductase maternal gene in Down syndrome: increased susceptibility in women carriers of the MTHFR 677T allele. <i>Molecular Biology Reports</i> , 2014, 41, 5491-5504.	2.3	13
42	Gene expression profile of 5-fluorouracil metabolic enzymes in laryngeal cancer cell line: Predictive parameters for response to 5-fluorouracil-based chemotherapy. <i>Biomedicine and Pharmacotherapy</i> , 2014, 68, 515-519.	5.6	5
43	Altered Expression of Immune-Related Genes in Children with Down Syndrome. <i>PLoS ONE</i> , 2014, 9, e107218.	2.5	23
44	Alterations in the expression pattern of MTHFR, DHFR, TYMS, and SLC19A1 genes after treatment of laryngeal cancer cells with high and low doses of methotrexate. <i>Tumor Biology</i> , 2013, 34, 3765-3771.	1.8	15
45	Association between GSTP1, GSTM1 and GSTT1 polymorphisms involved in xenobiotic metabolism and head and neck cancer development. <i>Molecular Biology Reports</i> , 2013, 40, 4181-4188.	2.3	10
46	<i>DHFR</i> 19-bp Deletion and <i>SHMT</i> C1420T Polymorphisms and Metabolite Concentrations of the Folate Pathway in Individuals with Down Syndrome. <i>Genetic Testing and Molecular Biomarkers</i> , 2013, 17, 274-277.	0.7	7
47	No evidence for association of the CD40, CD40L and BLYS polymorphisms, B-cell co-stimulatory molecules, with Brazilian endemic <i>Plasmodium vivax</i> malaria. <i>Transactions of the Royal Society of Tropical Medicine and Hygiene</i> , 2013, 107, 377-383.	1.8	9
48	Head and neck cancer: causes, prevention and treatment. <i>Brazilian Journal of Otorhinolaryngology</i> , 2013, 79, 239-247.	1.0	105
49	<i>BHMT</i> G742A and <i>MTHFD1</i> G1958A Polymorphisms and Down Syndrome Risk in the Brazilian Population. <i>Genetic Testing and Molecular Biomarkers</i> , 2012, 16, 628-631.	0.7	14
50	Head and neck cancer: genetic polymorphisms and folate metabolism. <i>Brazilian Journal of Otorhinolaryngology</i> , 2012, 78, 132-139.	1.0	14
51	Association between 11 genetic polymorphisms in folate-metabolising genes and head and neck cancer risk. <i>European Journal of Cancer</i> , 2012, 48, 1525-1531.	2.8	27
52	Genetic polymorphisms modulate the folate metabolism of Brazilian individuals with Down syndrome. <i>Molecular Biology Reports</i> , 2012, 39, 9277-9284.	2.3	12
53	Maternal Risk for Down Syndrome Is Modulated by Genes Involved in Folate Metabolism. <i>Disease Markers</i> , 2012, 32, 73-81.	1.3	39
54	Clinical and epidemiological characteristics of patients in the head and neck surgery department of a university hospital. <i>Sao Paulo Medical Journal</i> , 2012, 130, 307-313.	0.9	22

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55	Q36R polymorphism of KISS-1 gene in Brazilian head and neck cancer patients. <i>Molecular Biology Reports</i> , 2012, 39, 6029-6034.	2.3	4
56	Diffusion tensor MR imaging in neurofibromatosis type 1: expanding the knowledge of microstructural brain abnormalities. <i>Pediatric Radiology</i> , 2012, 42, 449-454.	2.0	32
57	Unidentified bright objects in neurofibromatosis type 1: Conventional MRI in the follow-up and correlation of microstructural lesions on diffusion tensor images. <i>European Journal of Paediatric Neurology</i> , 2012, 16, 42-47.	1.6	32
58	Polymorphisms and haplotypes in methylenetetrahydrofolate reductase gene and head and neck squamous cell carcinoma risk. <i>Molecular Biology Reports</i> , 2012, 39, 635-643.	2.3	20
59	MTHFD1 G1958A, BHMT G742A, TC2 C776G and TC2 A67G polymorphisms and head and neck squamous cell carcinoma risk. <i>Molecular Biology Reports</i> , 2012, 39, 887-893.	2.3	16
60	Polymorphisms of the CYP1A1 and CYP2E1 genes in head and neck squamous cell carcinoma risk. <i>Molecular Biology Reports</i> , 2012, 39, 1055-1063.	2.3	19
61	Polymorphism C1420T of Serine hydroxymethyltransferase gene on maternal risk for Down syndrome. <i>Molecular Biology Reports</i> , 2012, 39, 2561-2566.	2.3	16
62	VEGF gene alternative splicing: pro- and anti-angiogenic isoforms in cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2012, 138, 363-370.	2.5	80
63	Maternal risk for Down syndrome is modulated by genes involved in folate metabolism. <i>Disease Markers</i> , 2012, 32, 73-81.	1.3	27
64	Carcinogênese de cabeça e pescoço: impacto do polimorfismo MTHFD1 G1958A. <i>Revista Da Associação Médica Brasileira</i> , 2011, 57, 194-199.	0.7	10
65	A80G polymorphism of reduced folate carrier 1 (RFC1) gene and head and neck squamous cell carcinoma etiology in Brazilian population. <i>Molecular Biology Reports</i> , 2011, 38, 1071-1078.	2.3	15
66	Análise do gene TAX1BP1 em pacientes com câncer de cabeça e pescoço. <i>Brazilian Journal of Otorhinolaryngology</i> , 2010, 76, 193-198.	1.0	3
67	Genetic polymorphisms involved in folate metabolism and concentrations of methylmalonic acid and folate on plasma homocysteine and risk of coronary artery disease. <i>Journal of Thrombosis and Thrombolysis</i> , 2010, 29, 32-40.	2.1	32
68	Polimorfismo do gene metilenotetra-hidrofolato redutase (MTHFR) e o risco de carcinoma espinocelular de cabeça e pescoço. <i>Brazilian Journal of Otorhinolaryngology</i> , 2010, 76, 776-782.	1.0	10
69	The association between CBS 844ins68 polymorphism and head and neck squamous cell carcinoma risk â€” a case-control analysis. <i>Archives of Medical Science</i> , 2010, 5, 772-779.	0.9	12
70	19-base pair deletion polymorphism of the dihydrofolate reductase (DHFR) gene: maternal risk of Down syndrome and folate metabolism. <i>Sao Paulo Medical Journal</i> , 2010, 128, 215-218.	0.9	7
71	Análise dos genes GSTM1 e GSTT1 em pacientes com câncer de cabeça e pescoço. <i>Revista Da Associação Médica Brasileira</i> , 2010, 56, 299-303.	0.7	20
72	5-Methyltetrahydrofolate-homocysteine methyltransferase gene polymorphism (MTR) and risk of head and neck cancer. <i>Brazilian Journal of Medical and Biological Research</i> , 2010, 43, 445-450.	1.5	19

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73	Genetic variability of vascular endothelial growth factor and prognosis of head and neck cancer in a Brazilian population. <i>Brazilian Journal of Medical and Biological Research</i> , 2010, 43, 127-133.	1.5	10
74	Homocisteína e polimorfismos dos genes MTHFR e VEGF: impacto na doença arterial coronariana. <i>Arquivos Brasileiros De Cardiologia</i> , 2009, 92, 263-268.	0.8	16
75	The maspin expression in canine mammary tumors: an immunohistochemical and molecular study. <i>Pesquisa Veterinaria Brasileira</i> , 2009, 29, 167-173.	0.5	1
76	Unidentified bright objects on brain MRI in children as a diagnostic criterion for neurofibromatosis type 1. <i>Pediatric Radiology</i> , 2008, 38, 305-310.	2.0	70
77	Vascular endothelial growth factor genetic variability and coronary artery disease in Brazilian population. <i>Heart and Vessels</i> , 2008, 23, 371-375.	1.2	40
78	GAP0 syndrome: Three new Brazilian cases, additional osseous manifestations, and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 1523-1529.	1.2	19
79	Influence of UDP-Glucuronosyltransferase Polymorphisms on Mycophenolate Mofetil-Induced Side Effects in Kidney Transplant Patients. <i>Transplantation Proceedings</i> , 2008, 40, 708-710.	0.6	15
80	Role of Glutathione S-Transferase Polymorphisms and Chronic Allograft Dysfunction. <i>Transplantation Proceedings</i> , 2008, 40, 743-745.	0.6	10
81	Effect of Whole Bone Marrow Cell Infusion in the Progression of Experimental Chronic Renal Failure. <i>Transplantation Proceedings</i> , 2008, 40, 853-855.	0.6	44
82	Identification of dysregulated genes in lymphocytes from children with Down syndrome. <i>Genome</i> , 2008, 51, 19-29.	2.0	39
83	The MTR A2756G polymorphism is associated with an increase of plasma homocysteine concentration in Brazilian individuals with Down syndrome. <i>Brazilian Journal of Medical and Biological Research</i> , 2008, 41, 34-40.	1.5	21
84	Effectiveness of two programs of intermittent ferrous supplementation for treating iron-deficiency anemia in infants: randomized clinical trial. <i>Sao Paulo Medical Journal</i> , 2008, 126, 314-318.	0.9	10
85	A80G polymorphism of reduced folate carrier 1 (RFC1) and C776G polymorphism of transcobalamin 2 (TC2) genes in Down's syndrome etiology. <i>Sao Paulo Medical Journal</i> , 2008, 126, 329-332.	0.9	22
86	Genetic polymorphisms involved in folate metabolism and elevated plasma concentrations of homocysteine: maternal risk factors for Down syndrome in Brazil. <i>Genetics and Molecular Research</i> , 2008, 7, 33-42.	0.2	63
87	Combination of Angiotensin-Converting Enzyme and Methylenetetrahydrofolate Reductase Gene Polymorphisms as Determinant Risk Factors for Chronic Allograft Dysfunction. <i>Transplantation Proceedings</i> , 2007, 39, 78-80.	0.6	11
88	Effect of Folate, Vitamin B6, and Vitamin B12 Intake and MTHFR C677T Polymorphism on Homocysteine Concentrations of Renal Transplant Recipients. <i>Transplantation Proceedings</i> , 2007, 39, 3163-3165.	0.6	6
89	Methylenetetrahydrofolate reductase gene polymorphism and its association with coronary artery disease. <i>Sao Paulo Medical Journal</i> , 2007, 125, 4-8.	0.9	10
90	Genetic relatedness among clinical strains of <i>Stenotrophomonas maltophilia</i> in tertiary care hospital settings in São Paulo State, Brazil. <i>Brazilian Journal of Microbiology</i> , 2007, 38, 278-284.	2.0	2

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91	Angiotensin-Converting Enzyme Gene Polymorphism in Chronic Allograft Nephropathy. Transplantation Proceedings, 2006, 38, 1327-1328.	0.6	11
92	High frequencies of plexiform neurofibromas, mental retardation, learning difficulties, and scoliosis in Brazilian patients with neurofibromatosis type 1. Brazilian Journal of Medical and Biological Research, 2005, 38, 1441-1447.	1.5	19
93	Hyperhomocysteinemia and MTHFR C677T and A1298C polymorphisms are associated with chronic allograft nephropathy in renal transplant recipients. Transplantation Proceedings, 2004, 36, 2979-2981.	0.6	15
94	Systemic lupus erythematosus and microchimerism in autoimmunity. Transplantation Proceedings, 2002, 34, 2951-2952.	0.6	47
95	Analysis of the TP53 Gene in Normal Skin and Hair Follicle Samples From Sun-Exposed and Non-Sun-Exposed Sites on Normal and Albino Individuals Living in Southeast Brazil. Archives of Dermatology, 1999, 135, 1559-1560.	1.4	0
96	Chromosome breakpoint distribution in nonmelanoma skin cancers. Cancer Genetics and Cytogenetics, 1997, 99, 81-84.	1.0	1
97	Cytogenetic study of neoplastic and nonneoplastic cells of the skin. Cancer Genetics and Cytogenetics, 1995, 85, 16-19.	1.0	13