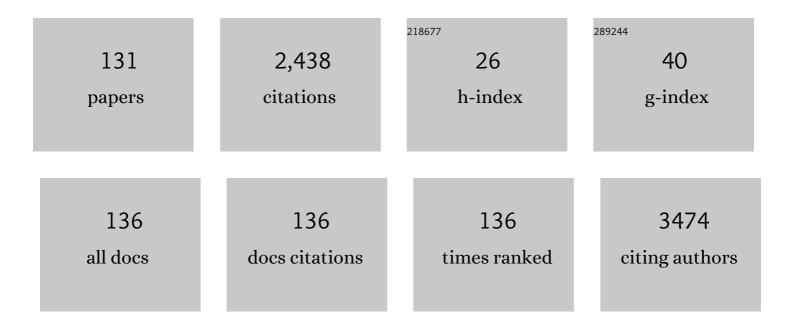
## Eny M Goloni-Bertollo

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Prenatal exposure to misoprostol and vascular disruption defects: A case-control study. American Journal of Medical Genetics Part A, 2000, 95, 302-306.	2.4	107
2	Head and neck cancer: causes, prevention and treatment. Brazilian Journal of Otorhinolaryngology, 2013, 79, 239-247.	1.0	105
3	Methylation as a biomarker for head and neck cancer. Oral Oncology, 2014, 50, 587-592.	1.5	89
4	VEGF gene alternative splicing: pro- and anti-angiogenic isoforms in cancer. Journal of Cancer Research and Clinical Oncology, 2012, 138, 363-370.	2.5	80
5	Unidentified bright objects on brain MRI in children as a diagnostic criterion for neurofibromatosis type 1. Pediatric Radiology, 2008, 38, 305-310.	2.0	70
6	Genetic polymorphisms involved in folate metabolism and elevated plasma concentrations of homocysteine: maternal risk factors for Down syndrome in Brazil. Genetics and Molecular Research, 2008, 7, 33-42.	0.2	63
7	Hepatocellular Carcinoma: a Comprehensive Review of Biomarkers, Clinical Aspects, and Therapy. Asian Pacific Journal of Cancer Prevention, 2017, 18, 863-872.	1.2	62
8	Meta-analysis and pooled analysis of GSTM1 and CYP1A1 polymorphisms and oral and pharyngeal cancers: a HuGE-GSEC review. Genetics in Medicine, 2008, 10, 369-384.	2.4	60
9	Influence of functional polymorphisms in TNF-1±, IL-8, and IL-10 cytokine genes on mRNA expression levels and risk of gastric cancer. Tumor Biology, 2015, 36, 9159-9170.	1.8	58
10	LHX6 is a sensitive methylation marker in head and neck carcinomas. Oncogene, 2006, 25, 5018-5026.	5.9	50
11	Systemic lupus erythematosus and microchimerism in autoimmunity. Transplantation Proceedings, 2002, 34, 2951-2952.	0.6	47
12	Effect of Whole Bone Marrow Cell Infusion in the Progression of Experimental Chronic Renal Failure. Transplantation Proceedings, 2008, 40, 853-855.	0.6	44
13	Validation of methylation markers for diagnosis of oral cavity cancer. European Journal of Cancer, 2015, 51, 632-641.	2.8	44
14	Vascular endothelial growth factor genetic variability and coronary artery disease in Brazilian population. Heart and Vessels, 2008, 23, 371-375.	1.2	40
15	Identification of dysregulated genes in lymphocytes from children with Down syndrome. Genome, 2008, 51, 19-29.	2.0	39
16	Maternal Risk for Down Syndrome Is Modulated by Genes Involved in Folate Metabolism. Disease Markers, 2012, 32, 73-81.	1.3	39
17	Sister chromatid exchanges and chromosome aberrations in lymphocytes of nurses handling antineoplastic drugs. International Journal of Cancer, 1992, 50, 341-344.	5.1	34
18	Prevalence of the GJB2 mutations and the del(GJB6-D13S1830) mutation in Brazilian patients with deafness. Hearing Research, 2004, 196, 87-93.	2.0	33

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19	Neurofibromatosis: chronological history and current issues. Anais Brasileiros De Dermatologia, 2013, 88, 329-343.	1.1	33
20	Genetic polymorphisms involved in folate metabolism and concentrations of methylmalonic acid and folate on plasma homocysteine and risk of coronary artery disease. Journal of Thrombosis and Thrombolysis, 2010, 29, 32-40.	2.1	32
21	Diffusion tensor MR imaging in neurofibromatosis type 1: expanding the knowledge of microstructural brain abnormalities. Pediatric Radiology, 2012, 42, 449-454.	2.0	32
22	Unidentified bright objects in neurofibromatosis type 1: Conventional MRI in the follow-up and correlation of microstructural lesions on diffusion tensor images. European Journal of Paediatric Neurology, 2012, 16, 42-47.	1.6	32
23	Iron deficiency anemia in children: a challenge for public health and for society. Sao Paulo Medical Journal, 2005, 123, 88-92.	0.9	30
24	Epidemiologic evaluation of head and neck patients in a university hospital of Northwestern São Paulo State. Brazilian Journal of Otorhinolaryngology, 2008, 74, 68-73.	1.0	30
25	Association between 11 genetic polymorphisms in folate-metabolising genes and head and neck cancer risk. European Journal of Cancer, 2012, 48, 1525-1531.	2.8	27
26	Neurofibromatoses: part 1 ? diagnosis and differential diagnosis. Arquivos De Neuro-Psiquiatria, 2014, 72, 241-250.	0.8	27
27	Differential expression of angiogenesis-related miRNAs and VECFA in cirrhosis and hepatocellular carcinoma. Archives of Medical Science, 2020, 16, 1150-1157.	0.9	27
28	Maternal risk for Down syndrome is modulated by genes involved in folate metabolism. Disease Markers, 2012, 32, 73-81.	1.3	27
29	Altered Expression of Immune-Related Genes in Children with Down Syndrome. PLoS ONE, 2014, 9, e107218.	2.5	23
30	Clinical and epidemiological characteristics of patients in the head and neck surgery department of a university hospital. Sao Paulo Medical Journal, 2012, 130, 307-313.	0.9	22
31	A80G polymorphism of reduced folate carrier 1 (RFC1) and C776G polymorphism of transcobalamin 2 (TC2) genes in Down's syndrome etiology. Sao Paulo Medical Journal, 2008, 126, 329-332.	0.9	22
32	The MTR A2756G polymorphism is associated with an increase of plasma homocysteine concentration in Brazilian individuals with Down syndrome. Brazilian Journal of Medical and Biological Research, 2008, 41, 34-40.	1.5	21
33	MicroRNAs as regulators of VEGFA and NFE2L2 in cancer. Gene, 2020, 759, 144994.	2.2	21
34	Clinical profile of children with down syndrome treated in a genetics outpatient service in the southeast of Brazil. Revista Da Associação Médica Brasileira, 2009, 55, 547-552.	0.7	20
35	Análise dos genes GSTM1 e GSTT1 em pacientes com câncer de cabeça e pescoço. Revista Da Associação Médica Brasileira, 2010, 56, 299-303.	0.7	20
36	Polymorphisms and haplotypes in methylenetetrahydrofolate reductase gene and head and neck squamous cell carcinoma risk. Molecular Biology Reports, 2012, 39, 635-643.	2.3	20

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37	Candidate Biomarkers for Oral Squamous Cell Carcinoma: Differential Expression of Oxidative Stress-Related Genes. Asian Pacific Journal of Cancer Prevention, 2018, 19, 1343-1349.	1.2	20
38	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
39	GAPO syndrome: Three new Brazilian cases, additional osseous manifestations, and review of the literature. American Journal of Medical Genetics, Part A, 2008, 146A, 1523-1529.	1.2	19
40	5-Methyltetrahydrofolate-homocysteine methyltransferase gene polymorphism (MTR) and risk of head and neck cancer. Brazilian Journal of Medical and Biological Research, 2010, 43, 445-450.	1.5	19
41	Polymorphisms of the CYP1A1 and CYP2E1 genes in head and neck squamous cell carcinoma risk. Molecular Biology Reports, 2012, 39, 1055-1063.	2.3	19
42	Genetic Polymorphisms Involved in Folate Metabolism and Maternal Risk for Down Syndrome: A Meta-Analysis. Disease Markers, 2014, 2014, 1-12.	1.3	18
43	VEGFA and NFE2L2 Gene Expression and Regulation by MicroRNAs in Thyroid Papillary Cancer and Colloid Goiter. Genes, 2020, 11, 954.	2.4	18
44	Tetrasomy 15q11-q13 identified by fluorescence in situ hybridization in a patient with autistic disorder. Arquivos De Neuro-Psiquiatria, 2002, 60, 290-294.	0.8	18
45	Polymorphisms of folate metabolism genes in patients with cirrhosis and hepatocellular carcinoma. World Journal of Hepatology, 2016, 8, 1234.	2.0	18
46	DNMT3B C46359T and SHMT1 C1420T polymorphisms in the folate pathway in carcinogenesis of head and neck. Molecular Biology Reports, 2014, 41, 581-589.	2.3	17
47	Role of MTHFR C677T and MTR A2756C polymorphisms in thyroid and breast cancer development. Genetics and Molecular Research, 2016, 15, .	0.2	17
48	Hypoxic niches are endowed with a protumorigenic mechanism that supersedes the protective function of PTEN. FASEB Journal, 2019, 33, 13435-13449.	0.5	17
49	HomocisteÃna e polimorfismos dos genes MTHFR e VEGF: impacto na doença arterial coronariana. Arquivos Brasileiros De Cardiologia, 2009, 92, 263-268.	0.8	16
50	MTHFD1 G1958A, BHMT G742A, TC2 C776G and TC2 A67G polymorphisms and head and neck squamous cell carcinoma risk. Molecular Biology Reports, 2012, 39, 887-893.	2.3	16
51	Polymorphism C1420T of Serine hydroxymethyltransferase gene on maternal risk for Down syndrome. Molecular Biology Reports, 2012, 39, 2561-2566.	2.3	16
52	Effect of <i>Helicobacter pylori</i> Eradication on TLR2 and TLR4 Expression in Patients with Gastric Lesions. Mediators of Inflammation, 2015, 2015, 1-9.	3.0	16
53	<i>CYP1A1</i> , <i>CYP2E1</i> and <i>EPHX1</i> polymorphisms in sporadic colorectal neoplasms. World Journal of Gastroenterology, 2016, 22, 9974.	3.3	16
54	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

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55	CSTT1 and GSTM1 polymorphism in cigarette smokers with head and neck squamous cell carcinoma. Brazilian Journal of Otorhinolaryngology, 2006, 72, 654-658.	1.0	15
56	Influence of UDP-Glucuronosyltransferase Polymorphisms on Mycophenolate Mofetil-Induced Side Effects in Kidney Transplant Patients. Transplantation Proceedings, 2008, 40, 708-710.	0.6	15
57	A80G polymorphism of reduced folate carrier 1 (RFC1) gene and head and neck squamous cell carcinoma etiology in Brazilian population. Molecular Biology Reports, 2011, 38, 1071-1078.	2.3	15
58	Effect of stem cells seeded onto biomaterial on the progression of experimental chronic kidney disease. Experimental Biology and Medicine, 2011, 236, 746-754.	2.4	15
59	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. Tumor Biology, 2013, 34, 3765-3771.	1.8	15
60	Comparative effects of mesenchymal stem cell therapy in distinct stages of chronic renal failure. Clinical and Experimental Nephrology, 2015, 19, 783-789.	1.6	15
61	Role of Tropomyosin-related kinase B receptor and brain-derived neurotrophic factor in cancer. Cytokine, 2020, 136, 155270.	3.2	15
62	<i>BHMT</i> G742A and <i>MTHFD1</i> G1958A Polymorphisms and Down Syndrome Risk in the Brazilian Population. Genetic Testing and Molecular Biomarkers, 2012, 16, 628-631.	0.7	14
63	Head and neck cancer: genetic polymorphisms and folate metabolism. Brazilian Journal of Otorhinolaryngology, 2012, 78, 132-139.	1.0	14
64	Polymorphisms in MTHFR, MTR, RFC1 and CßS genes involved in folate metabolism and thyroid cancer: a case-control study. Archives of Medical Science, 2019, 15, 522-530.	0.9	14
65	Microscopical evaluation of extracellular matrix and its relation to the palatopharyngeal muscle in obstructive sleep apnea. Microscopy Research and Technique, 2011, 74, 430-439.	2.2	13
66	Meta-analysis of Methylenetetrahydrofolate reductase maternal gene in Down syndrome: increased susceptibility in women carriers of the MTHFR 677T allele. Molecular Biology Reports, 2014, 41, 5491-5504.	2.3	13
67	Skin wound healing triggers epigenetic modifications of histone H4. Journal of Translational Medicine, 2020, 18, 138.	4.4	13
68	The association between CBS 844ins68 polymorphism and head and neck squamous cell carcinoma risk – a case-control analysis. Archives of Medical Science, 2010, 5, 772-779.	0.9	12
69	Genetic polymorphisms modulate the folate metabolism of Brazilian individuals with Down syndrome. Molecular Biology Reports, 2012, 39, 9277-9284.	2.3	12
70	Differential Expression of Inflammation-Related Genes in Children with Down Syndrome. Mediators of Inflammation, 2016, 2016, 1-8.	3.0	12
71	Molecular evaluation of glutathione S transferase family genes in patients with sporadic colorectal cancer. World Journal of Gastroenterology, 2018, 24, 4462-4471.	3.3	12
72	Characterization and strong risk association of TLR2 del -196 to -174 polymorphism and Helicobacter pylori and their influence on mRNA expression in gastric cancer. World Journal of Gastrointestinal Oncology, 2020, 12, 535-548.	2.0	12

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73	Angiotensin-Converting Enzyme Gene Polymorphism in Chronic Allograft Nephropathy. Transplantation Proceedings, 2006, 38, 1327-1328.	0.6	11
74	Combination of Angiotensin-Converting Enzyme and Methylenetetrahydrofolate Reductase Gene Polymorphisms as Determinant Risk Factors for Chronic Allograft Dysfunction. Transplantation Proceedings, 2007, 39, 78-80.	0.6	11
75	Methylenetetrahydrofolate reductase gene polymorphism and its association with coronary artery disease. Sao Paulo Medical Journal, 2007, 125, 4-8.	0.9	10
76	Role of Glutathione S-Transferase Polymorphisms and Chronic Allograft Dysfunction. Transplantation Proceedings, 2008, 40, 743-745.	0.6	10
77	Effectiveness of two programs of intermittent ferrous supplementation for treating iron-deficiency anemia in infants: randomized clinical trial. Sao Paulo Medical Journal, 2008, 126, 314-318.	0.9	10
78	Polimorfismo do gene metilenotetra-hidrofolato redutase (MTHFR) e o risco de carcinoma espinocelular de cabeça e pescoço. Brazilian Journal of Otorhinolaryngology, 2010, 76, 776-782.	1.0	10
79	Carcinogênese de cabeça e pescoço: impacto do polimorfismo MTHFD1 G1958A. Revista Da Associação Médica Brasileira, 2011, 57, 194-199.	0.7	10
80	Association between GSTP1, GSTM1 and GSTT1 polymorphisms involved in xenobiotic metabolism and head and neck cancer development. Molecular Biology Reports, 2013, 40, 4181-4188.	2.3	10
81	Neurofibromatosis: part 2 – clinical management. Arquivos De Neuro-Psiquiatria, 2015, 73, 531-543.	0.8	10
82	Differential Expression of Prostaglandin I2 Synthase Associated with Arachidonic Acid Pathway in the Oral Squamous Cell Carcinoma. Journal of Oncology, 2018, 2018, 1-13.	1.3	10
83	Genetic variability of vascular endothelial growth factor and prognosis of head and neck cancer in a Brazilian population. Brazilian Journal of Medical and Biological Research, 2010, 43, 127-133.	1.5	10
84	Relationship between CD44/CD133/CD117 cancer stem cells phenotype and Cetuximab and Paclitaxel treatment response in head and neck cancer cell lines. American Journal of Cancer Research, 2018, 8, 1633-1641.	1.4	10
85	Presence of the R1748X Mutation in the <i>NF1</i> Gene in a Brazilian Patient with Ectropion uveae. Ophthalmic Research, 2004, 36, 349-352.	1.9	9
86	Variables associated to fetal microchimerism in systemic lupus erythematosus patients. Clinical Rheumatology, 2016, 35, 107-111.	2.2	9
87	Is Magnetic Resonance Spectroscopy Capable of Detecting Metabolic Abnormalities in Neurofibromatosis Type 1 That Are Not Revealed in Brain Parenchyma of Normal Appearance?. Pediatric Neurology, 2015, 52, 314-319.	2.1	8
88	Gene Polymorphisms Involved in Folate Metabolism and DNA Methylation with the Risk of Head and Neck Cancer. Asian Pacific Journal of Cancer Prevention, 2020, 21, 3751-3759.	1.2	8
89	[CSTM1 and GSTT1 genes analysis in head and neck cancer patients]. Revista Da Associação Médica Brasileira, 2010, 56, 299-303.	0.7	8
90	19-base pair deletion polymorphism of the dihydrofolate reductase (DHFR) gene: maternal risk of Down syndrome and folate metabolism. Sao Paulo Medical Journal, 2010, 128, 215-218.	0.9	7

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91	<i>DHFR</i> 19-bp Deletion and <i>SHMT</i> C1420T Polymorphisms and Metabolite Concentrations of the Folate Pathway in Individuals with Down Syndrome. Genetic Testing and Molecular Biomarkers, 2013, 17, 274-277.	0.7	7
92	Biomarcadores de suscetibilidade à endometriose. Revista Brasileira De Ginecologia E Obstetricia, 2004, 26, 299-304.	0.8	7
93	Effect of Folate, Vitamin B6, and Vitamin B12 Intake and MTHFR C677T Polymorphism on Homocysteine Concentrations of Renal Transplant Recipients. Transplantation Proceedings, 2007, 39, 3163-3165.	0.6	6
94	Atorvastatin increases oxidative stress and inhibits cell migration of oral squamous cell carcinoma in vitro. Oral Oncology, 2019, 90, 109-114.	1.5	6
95	CFTR Molecular Analysis Reveals Infrequent Allele Frequencies in Nine Cystic Fibrosis Patients from Sao Paulo State, Brazil. Human Biology, 2003, 75, 393-398.	0.2	5
96	Gene expression profile of 5-fluorouracil metabolic enzymes in laryngeal cancer cell line: Predictive parameters for response to 5-fluorouracil-based chemotherapy. Biomedicine and Pharmacotherapy, 2014, 68, 515-519.	5.6	5
97	Interleukin 6 and 10 Serum Levels and Genetic Polymorphisms in Children with Down Syndrome. Mediators of Inflammation, 2018, 2018, 1-9.	3.0	5
98	Trends and predictions for survival and mortality in individuals with Down syndrome in Brazil: A 21â€year analysis. Journal of Intellectual Disability Research, 2020, 64, 551-560.	2.0	5
99	Polymorphisms in xenobiotic metabolism-related genes in patients with hepatocellular carcinoma: a case–control study. Xenobiotica, 2021, 51, 1-9.	1.1	5
100	Glutathione S-transferase Polymorphisms in Head and Neck Squamous Cell Carcinoma Treated with Chemotherapy and/or Radiotherapy. Asian Pacific Journal of Cancer Prevention, 2020, 21, 1637-1644.	1.2	5
101	Clinical, Epidemiological and Histopathological Aspects in Patients with Hepatocellular Carcinoma Undergoing Liver Transplantation. Asian Pacific Journal of Cancer Prevention, 2018, 19, 2795-2802.	1.2	5
102	Polymorphism of methylenetetrahydrofolate reductase (MTHFR) gene and risk of head and neck squamous cell carcinoma. Brazilian Journal of Otorhinolaryngology, 2010, 76, 776-82.	1.0	5
103	Regulation of VEGFA, KRAS, and NFE2L2 Oncogenes by MicroRNAs in Head and Neck Cancer. International Journal of Molecular Sciences, 2022, 23, 7483.	4.1	5
104	Polimorfismos GSTT1 e GSTM1 em indivÃduos tabagistas com carcinoma espinocelular de cabeça e pescoço. Revista Brasileira De Otorrinolaringologia, 2006, 72, 654-658.	0.2	4
105	Q36R polymorphism of KiSS-1 gene in Brazilian head and neck cancer patients. Molecular Biology Reports, 2012, 39, 6029-6034.	2.3	4
106	A case-control study of CYP2E1 (PstI) and CYP1A1 (MspI) polymorphisms in colorectal cancer. Genetics and Molecular Research, 2015, 14, 17856-17863.	0.2	4
107	Vitamin D3 increases the Caspase-3 p12, MTHFR, and P-glycoprotein reducing amyloid-Î <sup>2</sup> 42 in the kidney of a mouse model for Down syndrome. Life Sciences, 2019, 231, 116537.	4.3	4
108	Evaluation of molecular markers GSTM1 and GSTT1 and clinical factors in breast cancer: case-control study and literature review. Xenobiotica, 2021, 51, 1326-1334.	1.1	4

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109	Mutational analysis of the GAP-related domain of the neurofibromatosis type 1 gene in Brazilian NF1 patients. Genetics and Molecular Biology, 2004, 27, 326-330.	1.3	4
110	Association between folate metabolism polymorphisms and breast cancer: a case-control study. Genetics and Molecular Biology, 2021, 44, e20200485.	1.3	4
111	Analysis of the TAX1BP1 gene in head and neck cancer patients. Brazilian Journal of Otorhinolaryngology, 2010, 76, 193-8.	1.0	4
112	From Tissue Physoxia to Cancer Hypoxia, Cost-Effective Methods to Study Tissue-Specific O2 Levels in Cellular Biology. International Journal of Molecular Sciences, 2022, 23, 5633.	4.1	4
113	Prevalência de achados radiográficos da neurofibromatose tipo 1: estudo de 82 casos. Radiologia Brasileira, 2002, 35, 65-70.	0.7	3
114	Apoptosis in tongue squamous cell carcinoma and its correlation with clinically occult cervical metastasis. Micron, 2008, 39, 910-914.	2.2	3
115	Análise do gene TAX1BP1 em pacientes com câncer de cabeça e pescoço. Brazilian Journal of Otorhinolaryngology, 2010, 76, 193-198.	1.0	3
116	Head and neck carcinogenesis: impact of MTHFD1 G1958A polymorphism. Revista Da Associação Médica Brasileira (English Edition), 2011, 57, 188-193.	0.1	3
117	Alzheimer's Disease in the Down Syndrome: An Overview of Genetics and Molecular Aspects. Neurology India, 2021, 69, 32.	0.4	3
118	One-carbon metabolism and global DNA methylation in mothers of individuals with Down syndrome. Human Cell, 2021, 34, 1671-1681.	2.7	3
119	Overexpression of Antiangiogenic Vascular Endothelial Growth Factor Isoform and Splicing Regulatory Factors in Oral, Laryngeal and Pharyngeal Squamous Cell Carcinomas. Asian Pacific Journal of Cancer Prevention, 2017, 18, 2171-2177.	1.2	3
120	Anti-EGFR treatment effects on laryngeal cancer stem cells. American Journal of Translational Research (discontinued), 2021, 13, 143-155.	0.0	3
121	Double aneuploidy (48,XXY,+21) of maternal origin in a child born to a 13-year-old mother: evaluation of the maternal folate metabolism. Genetic Counseling, 2009, 20, 225-34.	0.1	3
122	The chromosome 5q21 band minisatellite and head and neck cancer. Cancer Genetics and Cytogenetics, 2003, 147, 87-88.	1.0	2
123	Werner's syndrome and restrictive cardiomyopathy. International Journal of Cardiology, 2006, 108, 284-285.	1.7	2
124	Genetic relatedness among clinical strains of Stenotrophomonas maltophilia in tertiary care hospital settings in São Paulo State, Brazil. Brazilian Journal of Microbiology, 2007, 38, 278-284.	2.0	2
125	Comparing techniques for the identification of the MTHFR A1298C polymorphism. Journal of Biomolecular Techniques, 2008, 19, 103-5.	1.5	2
126	Differential microRNA expression profile in blood of children with Down syndrome suggests a role in immunological dysfunction. Human Cell, 2022, 35, 639-648.	2.7	2

#	Article	IF	CITATIONS
127	Research Article Polymorphisms of interleukin 6 in Down syndrome individuals: a case-control study Genetics and Molecular Research, 2017, 16, .	0.2	1
128	The maspin expression in canine mammary tumors: an immunohistochemical and molecular study. Pesquisa Veterinaria Brasileira, 2009, 29, 167-173.	0.5	1
129	MiR-612, miR-637, and miR-874 can Regulate VEGFA Expression in Hepatocellular Carcinoma Cell Lines. Genes, 2022, 13, 282.	2.4	1
130	PP082. Oral Oncology, 2013, 49, S122.	1.5	0
131	Head and neck carcinogenesis: impact of MTHFD1 G1958A polymorphism. Revista Da Associação Médica Brasileira, 2011, 57, 188-193.	0.7	0