Eny M Goloni-Bertollo

List of Publications by Year
in descending order

131	2,438	218677	289244
papers	citations	26 h-index	40 g-index

136	136	136	3474
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	MiR-612, miR-637, and miR-874 can Regulate VEGFA Expression in Hepatocellular Carcinoma Cell Lines. Genes, 2022, 13, 282.	2.4	1
3	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
4	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
5	Alzheimer's Disease in the Down Syndrome: An Overview of Genetics and Molecular Aspects. Neurology India, 2021, 69, 32.	0.4	3
6	Polymorphisms in xenobiotic metabolism-related genes in patients with hepatocellular carcinoma: a caseâ€“control study. Xenobiotica, 2021, 51, 1-9.	1.1	5
7	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
8	One-carbon metabolism and global DNA methylation in mothers of individuals with Down syndrome. Human Cell, 2021, 34, 1671-1681.	2.7	3
9	Anti-EGFR treatment effects on laryngeal cancer stem cells. American Journal of Translational Research (discontinued), 2021, 13, 143-155.	0.0	3
10	Association between folate metabolism polymorphisms and breast cancer: a case-control study. Genetics and Molecular Biology, 2021, 44, e20200485.	1.3	4
11	Role of Tropomyosin-related kinase B receptor and brain-derived neurotrophic factor in cancer. Cytokine, 2020, 136, 155270.	3.2	15
12	MicroRNAs as regulators of VEGFA and NFE2L2 in cancer. Gene, 2020, 759, 144994.	2.2	21
13	Differential expression of angiogenesis-related miRNAs and VEGFA in cirrhosis and hepatocellular carcinoma. Archives of Medical Science, 2020, 16, 1150-1157.	0.9	27
14	VEGFA and NFE2L2 Gene Expression and Regulation by MicroRNAs in Thyroid Papillary Cancer and Colloid Goiter. Genes, 2020, 11, 954.	2.4	18
15	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
16	Skin wound healing triggers epigenetic modifications of histone H4. Journal of Translational Medicine, 2020, 18, 138.	4.4	13
17	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
18	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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19	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
20	Hypoxic niches are endowed with a protumorigenic mechanism that supersedes the protective function of PTEN. FASEB Journal, 2019, 33, 13435-13449.	0.5	17
21	Vitamin D3 increases the Caspase-3 p12, MTHFR, and P-glycoprotein reducing amyloid-Î²42 in the kidney of a mouse model for Down syndrome. Life Sciences, 2019, 231, 116537.	4.3	4
22	Atorvastatin increases oxidative stress and inhibits cell migration of oral squamous cell carcinoma in vitro. Oral Oncology, 2019, 90, 109-114.	1.5	6
23	Polymorphisms in MTHFR, MTR, RFC1 and CÃŸS genes involved in folate metabolism and thyroid cancer: a case-control study. Archives of Medical Science, 2019, 15, 522-530.	0.9	14
24	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
25	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
26	Interleukin 6 and 10 Serum Levels and Genetic Polymorphisms in Children with Down Syndrome. Mediators of Inflammation, 2018, 2018, 1-9.	3.0	5
27	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
28	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
29	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
30	Research Article Polymorphisms of interleukin 6 in Down syndrome individuals: a case-control study.. Genetics and Molecular Research, 2017, 16, .	0.2	1
31	Hepatocellular Carcinoma: a Comprehensive Review of Biomarkers, Clinical Aspects, and Therapy. Asian Pacific Journal of Cancer Prevention, 2017, 18, 863-872.	1.2	62
32	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
33	Differential Expression of Inflammation-Related Genes in Children with Down Syndrome. Mediators of Inflammation, 2016, 2016, 1-8.	3.0	12
34	Role of MTHFR C677T and MTR A2756G polymorphisms in thyroid and breast cancer development. Genetics and Molecular Research, 2016, 15, .	0.2	17
35	Variables associated to fetal microchimerism in systemic lupus erythematosus patients. Clinical Rheumatology, 2016, 35, 107-111.	2.2	9
36	CYP1A1,CYP2E1andEPHX1polymorphisms in sporadic colorectal neoplasms. World Journal of Gastroenterology, 2016, 22, 9974.	3.3	16

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37	Polymorphisms of folate metabolism genes in patients with cirrhosis and hepatocellular carcinoma. World Journal of Hepatology, 2016, 8, 1234.	2.0	18
38	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
39	Neurofibromatosis: part 2 â€“ clinical management. Arquivos De Neuro-Psiquiatria, 2015, 73, 531-543.	0.8	10
40	Effect ofHelicobacter pyloriEradication on TLR2 and TLR4 Expression in Patients with Gastric Lesions. Mediators of Inflammation, 2015, 2015, 1-9.	3.0	16
41	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
42	Validation of methylation markers for diagnosis of oral cavity cancer. European Journal of Cancer, 2015, 51, 632-641.	2.8	44
43	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
44	Influence of functional polymorphisms in TNF-Î±, 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
45	Neurofibromatoses: part 1 ? diagnosis and differential diagnosis. Arquivos De Neuro-Psiquiatria, 2014, 72, 241-250.	0.8	27
46	Genetic Polymorphisms Involved in Folate Metabolism and Maternal Risk for Down Syndrome: A Meta-Analysis. Disease Markers, 2014, 2014, 1-12.	1.3	18
47	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
48	Methylation as a biomarker for head and neck cancer. Oral Oncology, 2014, 50, 587-592.	1.5	89
49	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
50	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
51	Altered Expression of Immune-Related Genes in Children with Down Syndrome. PLoS ONE, 2014, 9, e107218.	2.5	23
52	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
53	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
54	PP082. Oral Oncology, 2013, 49, S122.	1.5	0

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55	DHFR 19-bp Deletion and SHMT 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
56	Head and neck cancer: causes, prevention and treatment. Brazilian Journal of Otorhinolaryngology, 2013, 79, 239-247.	1.0	105
57	Neurofibromatosis: chronological history and current issues. Anais Brasileiros De Dermatologia, 2013, 88, 329-343.	1.1	33
58	BHMTG742A andMTHFD1G1958A Polymorphisms and Down Syndrome Risk in the Brazilian Population. Genetic Testing and Molecular Biomarkers, 2012, 16, 628-631.	0.7	14
59	Head and neck cancer: genetic polymorphisms and folate metabolism. Brazilian Journal of Otorhinolaryngology, 2012, 78, 132-139.	1.0	14
60	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
61	Genetic polymorphisms modulate the folate metabolism of Brazilian individuals with Down syndrome. Molecular Biology Reports, 2012, 39, 9277-9284.	2.3	12
62	Maternal Risk for Down Syndrome Is Modulated by Genes Involved in Folate Metabolism. Disease Markers, 2012, 32, 73-81.	1.3	39
63	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
64	Q36R polymorphism of KiSS-1 gene in Brazilian head and neck cancer patients. Molecular Biology Reports, 2012, 39, 6029-6034.	2.3	4
65	Diffusion tensor MR imaging in neurofibromatosis type 1: expanding the knowledge of microstructural brain abnormalities. Pediatric Radiology, 2012, 42, 449-454.	2.0	32
66	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
67	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
68	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
69	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
70	Polymorphism C1420T of Serine hydroxymethyltransferase gene on maternal risk for Down syndrome. Molecular Biology Reports, 2012, 39, 2561-2566.	2.3	16
71	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
72	Maternal risk for Down syndrome is modulated by genes involved in folate metabolism. Disease Markers, 2012, 32, 73-81.	1.3	27

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73	Head and neck carcinogenesis: impact of MTHFD1 G1958A polymorphism. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira (English Edition), 2011, 57, 188-193.	0.1	3
74	CarcinogÃªnese de cabeÃ§a e pescoÃ§o: impacto do polimorfismo MTHFD1 G1958A. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2011, 57, 194-199.	0.7	10
75	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
76	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
77	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
78	Head and neck carcinogenesis: impact of MTHFD1 G1958A polymorphism. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2011, 57, 188-193.	0.7	0
79	AnÃ¡lise do gene TAX1BP1 em pacientes com cÃ¢ncer de cabeÃ§a e pescoÃ§o. Brazilian Journal of Otorhinolaryngology, 2010, 76, 193-198.	1.0	3
80	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
81	Polimorfismo do gene metilenotetra-hidrofolato redutase (MTHFR) e o risco de carcinoma espinocelular de cabeÃ§a e pescoÃ§o. Brazilian Journal of Otorhinolaryngology, 2010, 76, 776-782.	1.0	10
82	The association between CBS 844ins68 polymorphism and head and neck squamous cell carcinoma risk â€“ a case-control analysis. Archives of Medical Science, 2010, 5, 772-779.	0.9	12
83	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
84	AnÃ¡lise dos genes GSTM1 e GSTT1 em pacientes com cÃ¢ncer de cabeÃ§a e pescoÃ§o. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2010, 56, 299-303.	0.7	20
85	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
86	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
87	Analysis of the TAX1BP1 gene in head and neck cancer patients. Brazilian Journal of Otorhinolaryngology, 2010, 76, 193-8.	1.0	4
88	[GSTM1 and GSTT1 genes analysis in head and neck cancer patients]. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2010, 56, 299-303.	0.7	8
89	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
90	Clinical profile of children with down syndrome treated in a genetics outpatient service in the southeast of Brazil. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2009, 55, 547-552.	0.7	20

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91	HomocisteÃna e polimorfismos dos genes MTHFR e VEGF: impacto na doenÃ§a arterial coronariana. Arquivos Brasileiros De Cardiologia, 2009, 92, 263-268.	0.8	16
92	The maspin expression in canine mammary tumors: an immunohistochemical and molecular study. Pesquisa Veterinaria Brasileira, 2009, 29, 167-173.	0.5	1
93	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
94	Apoptosis in tongue squamous cell carcinoma and its correlation with clinically occult cervical metastasis. Micron, 2008, 39, 910-914.	2.2	3
95	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
96	Vascular endothelial growth factor genetic variability and coronary artery disease in Brazilian population. Heart and Vessels, 2008, 23, 371-375.	1.2	40
97	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
98	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
99	Influence of UDP-Glucuronosyltransferase Polymorphisms on Mycophenolate Mofetil-Induced Side Effects in Kidney Transplant Patients. Transplantation Proceedings, 2008, 40, 708-710.	0.6	15
100	Role of Glutathione S-Transferase Polymorphisms and Chronic Allograft Dysfunction. Transplantation Proceedings, 2008, 40, 743-745.	0.6	10
101	Effect of Whole Bone Marrow Cell Infusion in the Progression of Experimental Chronic Renal Failure. Transplantation Proceedings, 2008, 40, 853-855.	0.6	44
102	Identification of dysregulated genes in lymphocytes from children with Down syndrome. Genome, 2008, 51, 19-29.	2.0	39
103	Epidemiologic evaluation of head and neck patients in a university hospital of Northwestern SÃ£o Paulo State. Brazilian Journal of Otorhinolaryngology, 2008, 74, 68-73.	1.0	30
104	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
105	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
106	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
107	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
108	Comparing techniques for the identification of the MTHFR A1298C polymorphism. Journal of Biomolecular Techniques, 2008, 19, 103-5.	1.5	2

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109	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
110	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
111	Methylenetetrahydrofolate reductase gene polymorphism and its association with coronary artery disease. Sao Paulo Medical Journal, 2007, 125, 4-8.	0.9	10
112	Genetic relatedness among clinical strains of Stenotrophomonas maltophilia in tertiary care hospital settings in SÃ£o Paulo State, Brazil. Brazilian Journal of Microbiology, 2007, 38, 278-284.	2.0	2
113	Werner's syndrome and restrictive cardiomyopathy. International Journal of Cardiology, 2006, 108, 284-285.	1.7	2
114	Angiotensin-Converting Enzyme Gene Polymorphism in Chronic Allograft Nephropathy. Transplantation Proceedings, 2006, 38, 1327-1328.	0.6	11
115	Polimorfismos GSTT1 e GSTM1 em indivÃduos tabagistas com carcinoma espinocelular de cabeÃ§a e pescoÃ§o. Revista Brasileira De Otorrinolaringologia, 2006, 72, 654-658.	0.2	4
116	GSTT1 and GSTM1 polymorphism in cigarette smokers with head and neck squamous cell carcinoma. Brazilian Journal of Otorhinolaryngology, 2006, 72, 654-658.	1.0	15
117	LHX6 is a sensitive methylation marker in head and neck carcinomas. Oncogene, 2006, 25, 5018-5026.	5.9	50
118	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
119	Iron deficiency anemia in children: a challenge for public health and for society. Sao Paulo Medical Journal, 2005, 123, 88-92.	0.9	30
120	Presence of the R1748X Mutation in the NF1 Gene in a Brazilian Patient with Ectropion uveae. Ophthalmic Research, 2004, 36, 349-352.	1.9	9
121	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
122	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
123	Biomarcadores de suscetibilidade Ã endometriose. Revista Brasileira De Ginecologia E Obstetricia, 2004, 26, 299-304.	0.8	7
124	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
125	The chromosome 5q21 band minisatellite and head and neck cancer. Cancer Genetics and Cytogenetics, 2003, 147, 87-88.	1.0	2
126	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

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127	Systemic lupus erythematosus and microchimerism in autoimmunity. Transplantation Proceedings, 2002, 34, 2951-2952.	0.6	47
128	PrevalÃªncia de achados radiogrÃ¡ficos da neurofibromatose tipo 1: estudo de 82 casos. Radiologia Brasileira, 2002, 35, 65-70.	0.7	3
129	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
130	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
131	Sister chromatid exchanges and chromosome aberrations in lymphocytes of nurses handling antineoplastic drugs. International Journal of Cancer, 1992, 50, 341-344.	5.1	34