Lavinia Schuler-Faccini

List of Publications by Year
in descending order

182 papers	5,864 citations	159585 30 h-index	91884 69 g-index
191 all docs	191 docs citations	191 times ranked	8606 citing authors

#	Article	IF	Citations
1	Recent dengue virus infection: epidemiological survey on risk factors associated with infection in a medium-sized city in Mato Grosso. Sao Paulo Medical Journal, 2022, 140, 33-41.	0.9	2
2	Neurodevelopment in Children Exposed to Zika in utero: Clinical and Molecular Aspects. Frontiers in Genetics, 2022, 13, 758715.	2.3	12
3	Disentangling Signatures of Selection Before and After European Colonization in Latin Americans. Molecular Biology and Evolution, 2022, 39, .	8.9	16
4	Microcephaly prevalence after the 2015 to 2016 Zika outbreak in TangarÃ¡ da Serra, Brazil: a population-based study. Reproductive and Developmental Medicine, 2022, 6, 98-103.	0.5	0
5	Prevalence of Congenital Anomaly and Its Relationship with Maternal Education and Age According to Local Development in the Extreme South of Brazil. International Journal of Environmental Research and Public Health, 2022, 19, 8079.	2.6	2
6	An invincible memory: what surname analysis tells us about history, health and population medical genetics in the Brazilian Northeast. Journal of Biosocial Science, 2021, 53, 183-198.	1.2	7
7	Genotypeâ€phenotype correlations on epidermolysis bullosa with congenital absence of skin: A comprehensive review. Clinical Genetics, 2021, 99, 29-41.	2.0	14
8	Zika virusâ€induced brain malformations in chicken embryos. Birth Defects Research, 2021, 113, 22-31.	1.5	9
9	Prevalence of congenital anomalies at birth among live births in the state of MaranhÃ£o from 2001 to 2016: temporal and spatial analysis. Revista Brasileira De Epidemiologia, 2021, 24, e210020.	0.8	7
10	Lista de anomalias congÃªnitas prioritÃ¡rias para vigilÃ¢ncia no Ã¢mbito do Sistema de InformaÃ§Ãµes sobre Nascidos Vivos do Brasil. Epidemiologia E Servicos De Saude: Revista Do Sistema Unico De Saude Do Brasil, 2021, 30, e2020835.	1.0	3
11	Aprosopia/holoprosencephaly in a stillborn puppy: when the face predicts the brain. International Journal of Veterinary Science and Medicine, 2021, 9, 7-10.	2.2	1
12	Mapeamento dinÃ¢mico da probabilidade de infestaÃ§Ã£o por vetores urbanos de arbovÃrus nos municÃpios do Rio Grande do Sul, 2016-2017. Epidemiologia E Servicos De Saude: Revista Do Sistema Unico De Saude Do Brasil, 2021, 30, e2020154.	1.0	2
13	A GWAS in Latin Americans identifies novel face shape loci, implicating VPS13B and a Denisovan introgressed region in facial variation. Science Advances, 2021, 7, .	10.3	32
14	Gene panel for the diagnosis of epidermolysis bullosa: proposal for a viable and efficient approach. Anais Brasileiros De Dermatologia, 2021, 96, 155-162.	1.1	6
15	Zika Brazilian Cohorts (ZBC) Consortium: Protocol for an Individual Participant Data Meta-Analysis of Congenital Zika Syndrome after Maternal Exposure during Pregnancy. Viruses, 2021, 13, 687.	3.3	9
16	Genetic Susceptibility to Drug Teratogenicity: A Systematic Literature Review. Frontiers in Genetics, 2021, 12, 645555.	2.3	11
17	Machine learning model on heart rate variability metrics identifies asymptomatic toddlers exposed to zika virus during pregnancy. Physiological Measurement, 2021, 42, 055008.	2.1	10
18	Molecular mechanisms of Zika virus teratogenesis from animal studies: a systematic review protocol. Systematic Reviews, 2021, 10, 160.	5.3	2

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19	Prevalence and antimicrobial resistance profile of Staphylococcus aureus in inherited epidermolysis bullosa: a crossâ€sectional multicenter study in Brazil. International Journal of Dermatology, 2021, 60, 1126-1130.	1.0	3
20	Zika Virus Infection Associated with Autism Spectrum Disorder: A Case Report. NeuroImmunoModulation, 2021, 28, 229-232.	1.8	8
21	Comparative Genomics Identifies Putative Interspecies Mechanisms Underlying Crbn-Sall4-Linked Thalidomide Embryopathy. Frontiers in Genetics, 2021, 12, 680217.	2.3	2
22	Functional Polymorphisms in the p53 Pathway Genes on the Genetic Susceptibility to Zika Virus Teratogenesis. Frontiers in Cellular and Infection Microbiology, 2021, 11, 641413.	3.9	1
23	Maternal outcomes and risk factors for COVID-19 severity among pregnant women. Scientific Reports, 2021, 11, 13898.	3.3	77
24	Prediction of eye, hair and skin colour in Latin Americans. Forensic Science International: Genetics, 2021, 53, 102517.	3.1	6
25	Possible Emergence of Zika Virus of African Lineage in Brazil and the Risk for New Outbreaks. Frontiers in Cellular and Infection Microbiology, 2021, 11, 680025.	3.9	4
26	Development of dentofacial characteristics related to Incontinentia Pigmenti syndrome: A repeated cross-sectional study. American Journal of Orthodontics and Dentofacial Orthopedics, 2021, 160, 66-76.	1.7	1
27	Evolutionary analysis of the anti-viral STAT2 gene of primates and rodents: Signature of different stages of an arms race. Infection, Genetics and Evolution, 2021, 95, 105030.	2.3	1
28	COVID-19 during pregnancy and adverse outcomes: Concerns and recommendations from The Brazilian Teratology Information Service. Genetics and Molecular Biology, 2021, 44, e20200224.	1.3	5
29	Knowledge and actions for the control of the vector Aedes aegypti in a municipality in the Legal Amazon. Revista Do Instituto De Medicina Tropical De Sao Paulo, 2021, 63, e64.	1.1	0
30	Evaluation of Polymorphisms in Toll-Like Receptor Genes as Biomarkers of the Response to Treatment of Erythema Nodosum Leprosum. Frontiers in Medicine, 2021, 8, 713143.	2.6	4
31	Genetic variants linked to folliculogenesis and successful pregnancy are not associated with twin births in a twinsâ€™ town. Journal of Maternal-Fetal and Neonatal Medicine, 2020, 33, 3431-3438.	1.5	1
32	HLA diversity in Brazil. Hla, 2020, 95, 3-14.	0.6	9
33	A large family with CYLD cutaneous syndrome: medical genetics at the community level. Journal of Community Genetics, 2020, 11, 279-284.	1.2	5
34	Neurodevelopment of Nonmicrocephalic Children, After 18 Months of Life, Exposed Prenatally to Zika Virus. Journal of Child Neurology, 2020, 35, 278-282.	1.4	22
35	Inherited epidermolysis bullosa: update on the clinical and genetic aspects. Anais Brasileiros De Dermatologia, 2020, 95, 551-569.	1.1	47
36	Epidermolysis bullosa with congenital absence of skin: Clinical and genetic characterization of a <scp>23â€case</scp> series. Clinical Genetics, 2020, 98, 99-101.	2.0	3

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37	CRL4-Cereblon complex in Thalidomide Embryopathy: a translational investigation. Scientific Reports, 2020, 10, 851.	3.3	8
38	Fetal Alcohol Spectrum Disorders: Health Needs Assessment in Brazil. Alcoholism: Clinical and Experimental Research, 2020, 44, 660-668.	2.4	2
39	Evolutionary analysis of the Musashi family: What can it tell us about Zika?. Infection, Genetics and Evolution, 2020, 84, 104364.	2.3	4
40	Novel <i>AHDC1</i> Gene Mutation in a Brazilian Individual: Implications of Xia-Gibbs Syndrome. Molecular Syndromology, 2020, 11, 24-29.	0.8	12
41	Cancer-related worry and risk perception in Brazilian individuals seeking genetic counseling for hereditary breast cancer. Genetics and Molecular Biology, 2020, 43, e20190097.	1.3	6
42	Site Occupancy by Aedes aegypti in a Subtropical City is Most Sensitive to Control during Autumn and Winter Months. American Journal of Tropical Medicine and Hygiene, 2020, 103, 445-454.	1.4	3
43	Motor development in non-microcephalic infants born to mothers with Zika Virus infection during pregnancy. Fisioterapia E Pesquisa, 2020, 27, 174-179.	0.1	1
44	Anomalias congÃªnitas na perspectiva da vigilÃ¢ncia em saÃºde: compilaÃ§Ã£o de uma lista com base na CID-10. Epidemiologia E Servicos De Saude: Revista Do Sistema Unico De Saude Do Brasil, 2020, 29, e2020164.	1.0	1
45	Clusters of genetic diseases in Brazil. Journal of Community Genetics, 2019, 10, 121-128.	1.2	17
46	Prevalence and causes of congenital microcephaly in the absence of a Zika virus outbreak in southern Brazil. Jornal De Pediatria, 2019, 95, 600-606.	2.0	12
47	The role of ESCO2, SALL4 and TBX5 genes in the susceptibility to thalidomide teratogenesis. Scientific Reports, 2019, 9, 11413.	3.3	11
48	NR3C1,ABCB1,TNFandCYP2C19polymorphisms association with the response to the treatment of erythema nodosum leprosum. Pharmacogenomics, 2019, 20, 503-516.	1.3	1
49	Assembling systems biology, embryo development and teratogenesis: What do we know so far and where to go next?. Reproductive Toxicology, 2019, 88, 67-75.	2.9	7
50	Rare Diseases in Uruguay: Focus on Infants with Abnormal Newborn Screening. Journal of Inborn Errors of Metabolism and Screening, 2019, 7, .	0.3	3
51	Whole-exome sequencing in familial keratoconus: the challenges of a genetically complex disorder. Arquivos Brasileiros De Oftalmologia, 2019, 82, 453-459.	0.5	7
52	Prevalence and causes of congenital microcephaly in the absence of a Zika virus outbreak in southern Brazil. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2019, 95, 600-606.	0.2	0
53	A GWAS in Latin Americans highlights the convergent evolution of lighter skin pigmentation in Eurasia. Nature Communications, 2019, 10, 358.	12.8	130
54	Analysis of a Protein Network Related to Copy Number Variations in Autism Spectrum Disorder. Journal of Molecular Neuroscience, 2019, 69, 140-149.	2.3	7

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55	Information and Diagnosis Networks â€“ tools to improve diagnosis and treatment for patients with rare genetic diseases. Genetics and Molecular Biology, 2019, 42, 155-164.	1.3	9
56	An overview of the genetic basis of epidermolysis bullosa in Brazil: discovery of novel and recurrent diseaseâ€causing variants. Clinical Genetics, 2019, 96, 189-198.	2.0	22
57	Population medical genetics: translating science to the community. Genetics and Molecular Biology, 2019, 42, 312-320.	1.3	8
58	Genetic analysis of patients with fructose-1,6-bisphosphatase deficiency. Gene, 2019, 699, 102-109.	2.2	9
59	From abortion-inducing medications to Zika Virus Syndrome: 27 years experience of the First Teratogen Information Service in Latin America. Genetics and Molecular Biology, 2019, 42, 297-304.	1.3	1
60	The role of FAS, FAS-L, BAX, and BCL-2 gene polymorphisms in determining susceptibility to unexplained recurrent pregnancy loss. Journal of Assisted Reproduction and Genetics, 2019, 36, 995-1002.	2.5	9
61	Microcephaly infant mortality in Brazil before zika outbreak.. Revista De La Facultad De Ciencias Medicas De Cordoba, 2019, 76, 217-221.	0.3	1
62	ADGRL3 rs6551665 as a Common Vulnerability Factor Underlying Attention-Deficit/Hyperactivity Disorder and Autism Spectrum Disorder. NeuroMolecular Medicine, 2019, 21, 60-67.	3.4	19
63	Erythema Nodosum Leprosum: Update and challenges on the treatment of a neglected condition. Acta Tropica, 2018, 183, 134-141.	2.0	44
64	ZIKA Virus and Neuroscience: the Need for a Translational Collaboration. Molecular Neurobiology, 2018, 55, 1551-1555.	4.0	7
65	Genetic susceptibility to thalidomide embryopathy in humans: Study of candidate development genes. Birth Defects Research, 2018, 110, 456-461.	1.5	4
66	Lack of association between genetic polymorphisms in IGF1 and IGFBP3 with twin births in a Brazilian population (CÃ¢ndido GodÃ³i, Rio Grande do Sul). Genetics and Molecular Biology, 2018, 41, 775-780.	1.3	2
67	Perfil das anomalias congÃªnitas em nascidos vivos de TangarÃ¡ da Serra, Mato Grosso, 2006-2016*. Epidemiologia E Servicos De Saude: Revista Do Sistema Unico De Saude Do Brasil, 2018, 27, e2018008.	1.0	7
68	Zika Virus as a Possible Risk Factor for Autism Spectrum Disorder: Neuroimmunological Aspects. NeuroImmunoModulation, 2018, 25, 320-327.	1.8	33
69	Collagen I Defect Corneal Profiles in Osteogenesis Imperfecta. Cornea, 2018, 37, 1561-1565.	1.7	12
70	Is intrauterine exposure to acetaminophen associated with emotional and hyperactivity problems during childhood? Findings from the 2004 Pelotas birth cohort. BMC Psychiatry, 2018, 18, 368.	2.6	31
71	Latin Americans show wide-spread Converso ancestry and imprint of local Native ancestry on physical appearance. Nature Communications, 2018, 9, 5388.	12.8	123
72	Genome-wide association studies and CRISPR/Cas9-mediated gene editing identify regulatory variants influencing eyebrow thickness in humans. PLoS Genetics, 2018, 14, e1007640.	3.5	20

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73	Spatial analyzes of HLA data in Rio Grande do Sul, south Brazil: genetic structure and possible correlation with autoimmune diseases. International Journal of Health Geographics, 2018, 17, 34.	2.5	7
74	Angiogenesis and oxidative stress-related gene variants in recurrent pregnancy loss. Reproduction, Fertility and Development, 2018, 30, 498.	0.4	11
75	Evidence for Association Between OXTR Gene and ASD Clinical Phenotypes. Journal of Molecular Neuroscience, 2018, 65, 213-221.	2.3	19
76	Genetic evaluation in TP53 and MDM2 as modifier genes for congenital aniridia. Gene Reports, 2018, 13, 235-236.	0.8	0
77	Why is congenital Zika syndrome asymmetrically distributed among human populations?. PLoS Biology, 2018, 16, e2006592.	5.6	32
78	Intrafamilial clinical variability in four families with incontinentia pigmenti. American Journal of Medical Genetics, Part A, 2018, 176, 2318-2324.	1.2	5
79	Twin Peaks: A spatial and temporal study of twinning rates in Brazil. PLoS ONE, 2018, 13, e0200885.	2.5	6
80	Zika rash and increased risk of congenital brain abnormalities â€“ Authors' reply. Lancet, The, 2017, 389, 152.	13.7	1
81	Angiogenesis-related genes and thalidomide teratogenesis in humans: an approach on genetic variation and review of past in vitro studies. Reproductive Toxicology, 2017, 70, 133-140.	2.9	5
82	Analysis of Polymorphism rs1042522 in TP53 Gene in the Mothers of Twins and of Singletons: A Population-Based Study in Rio Grande do Sul, Brazil. Twin Research and Human Genetics, 2017, 20, 132-136.	0.6	4
83	The phenotypic spectrum of congenital Zika syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 841-857.	1.2	167
84	Multiethnic GWAS Reveals Polygenic Architecture of Earlobe Attachment. American Journal of Human Genetics, 2017, 101, 913-924.	6.2	29
85	The impact of thalidomide use in birth defects in Brazil. European Journal of Medical Genetics, 2017, 60, 12-15.	1.3	16
86	Teratogens: a public health issue â€“ a Brazilian overview. Genetics and Molecular Biology, 2017, 40, 387-397.	1.3	26
87	Search for DQ2.5 and DQ8 alleles using a lower cost technique in patients with type 1 diabetes and celiac disease in a population of southern Brazil. Archives of Endocrinology and Metabolism, 2017, 61, 550-555.	0.6	1
88	Socioeconomic Status Is Not Related with Facial Fluctuating Asymmetry: Evidence from Latin-American Populations. PLoS ONE, 2017, 12, e0169287.	2.5	17
89	Leprosy in Southern Brazil: a twenty-year epidemiological profile. Revista Da Sociedade Brasileira De Medicina Tropical, 2017, 50, 251-255.	0.9	11
90	Music genetics research: Association with musicality of a polymorphism in the AVPR1A gene. Genetics and Molecular Biology, 2017, 40, 421-429.	1.3	12

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91	Zika virus infection and congenital anomalies in the Americas: opportunities for regional action. Revista Panamericana De Salud Publica/Pan American Journal of Public Health, 2017, 41, 1-8.	1.1	5
92	RelaciÃ³n entre tratamiento hormonal, cirugÃa-ortodoncia maxilofacial, traumatismos y malformaciones craneofaciales y la asimetrÃa fluctuante. Revista Argentina De Antropologia Biologica, 2017, 20, 6.	0.4	0
93	Screening for germline BRCA1, BRCA2, TP53 and CHEK2 mutations in families at-risk for hereditary breast cancer identified in a population-based study from Southern Brazil. Genetics and Molecular Biology, 2016, 39, 210-222.	1.3	21
94	Congenital Zika virus syndrome in Brazil: a case series of the first 1501 livebirths with complete investigation. Lancet, The, 2016, 388, 891-897.	13.7	515
95	New Findings in eNOS gene and Thalidomide Embryopathy Suggest pre-transcriptional effect variants as susceptibility factors. Scientific Reports, 2016, 6, 23404.	3.3	12
96	A genome-wide association scan in admixed Latin Americans identifies loci influencing facial and scalp hair features. Nature Communications, 2016, 7, 10815.	12.8	159
97	Psychomotor agitation and mood instability in patients with autism spectrum disorders: A possible effect of SLC6A4 gene?. Research in Autism Spectrum Disorders, 2016, 26, 48-56.	1.5	1
98	Zika virus: A new human teratogen? Implications for women of reproductive age. Clinical Pharmacology and Therapeutics, 2016, 100, 28-30.	4.7	29
99	A genome-wide association scan implicates DCHS2, RUNX2, GLI3, PAX1 and EDAR in human facial variation. Nature Communications, 2016, 7, 11616.	12.8	171
100	Genomic and in silico analyses of CRBN gene and thalidomide embryopathy in humans. Reproductive Toxicology, 2016, 66, 99-106.	2.9	8
101	Retinoblastoma in a pediatric oncology reference center in Southern Brazil. BMC Pediatrics, 2016, 16, 48.	1.7	16
102	Spinocerebellar ataxia type 3/Machadoâ€“Joseph disease: segregation patterns and factors influencing instability of expanded <scp>CAG</scp> transmissions. Clinical Genetics, 2016, 90, 134-140.	2.0	36
103	Microcephaly in Brazil: how to interpret reported numbers?. Lancet, The, 2016, 387, 621-624.	13.7	193
104	Primary prevention of neural tube defects in Brazil: insights into anencephaly. Journal of Community Genetics, 2016, 7, 97-105.	1.2	2
105	KAMUTHE video microanalysis system for use in Brazil: translation, cross-cultural adaptation and evidence of validity and reliability. Health Psychology Report, 2016, 5, 125-137.	0.9	3
106	Possible Association Between Zika Virus Infection and Microcephaly â€" Brazil, 2015. Morbidity and Mortality Weekly Report, 2016, 65, 59-62.	15.1	859
107	Thalidomide embryopathy: Followâ€up of cases born between 1959 and 2010. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 794-803.	1.6	26
108	Clinical and molecular characterization of a Brazilian cohort of campomelic dysplasia patients, and identification of seven new SOX9 mutations. Genetics and Molecular Biology, 2015, 38, 14-20.	1.3	8

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109	Heart and blood medications. , 2015, , 193-223.		0
110	Self-Assessment of Color Categories and Its Relationship with HLA Profiling in Brazilian Bone Marrow Donors. Biology of Blood and Marrow Transplantation, 2015, 21, 1140-1144.	2.0	13
111	A genome-wide association study identifies multiple loci for variation in human ear morphology. Nature Communications, 2015, 6, 7500.	12.8	80
112	Maple syrup urine disease in Brazil: a panorama of the last two decades. Jornal De Pediatria, 2015, 91, 292-298.	2.0	18
113	Project REENCONTRO: ethical aspects of genetic identification in families separated by the compulsory isolation of leprosy patients in Brazil. Journal of Community Genetics, 2015, 6, 215-222.	1.2	2
114	Pharmacoepidemiology and thalidomide embryopathy surveillance in Brazil. Reproductive Toxicology, 2015, 53, 63-67.	2.9	17
115	Ethics, genetics and public policies in Uruguay: newborn and infant screening as a paradigm. Journal of Community Genetics, 2015, 6, 241-249.	1.2	6
116	Ocular and craniofacial phenotypes in a large Brazilian family with congenital aniridia. Clinical Genetics, 2015, 87, 68-73.	2.0	2
117	The Genetic Basis of Autism Spectrum Disorder. , 2015, , 39-63.		1
118	Impact on Pregnancies in South Brazil from the Influenza A (H1N1) Pandemic: Cohort Study. PLoS ONE, 2014, 9, e88624.	2.5	18
119	Implications of the Admixture Process in Skin Color Molecular Assessment. PLoS ONE, 2014, 9, e96886.	2.5	22
120	[NO TITLE AVAILABLE]. Genetics and Molecular Biology, 2014, 37, 263-270.	1.3	16
121	[NO TITLE AVAILABLE]. Genetics and Molecular Biology, 2014, 37, 186-193.	1.3	8
122	Genetics and human rights: Two histories: restoring genetic identity after forced disappearance and identity suppression in Argentina and after compulsory isolation for leprosy in Brazil. Genetics and Molecular Biology, 2014, 37, 299-304.	1.3	8
123	Glycogen storage disease type I: clinical and laboratory profile. Jornal De Pediatria, 2014, 90, 572-579.	2.0	22
124	A DNA repair variant in POLQ (c.-1060A > G) is associated to hereditary breast cancer patients: a caseâ€“control study. BMC Cancer, 2014, 14, 850.	2.6	12
125	Admixture in Latin America: Geographic Structure, Phenotypic Diversity and Self-Perception of Ancestry Based on 7,342 Individuals. PLoS Genetics, 2014, 10, e1004572.	3.5	350
126	p53 signaling pathway polymorphisms associated to recurrent pregnancy loss. Molecular Biology Reports, 2014, 41, 1871-1877.	2.3	33

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127	Interaction between TP63 and MDM2 genes and the risk of recurrent pregnancy loss. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2014, 182, 7-10.	1.1	11
128	The role of Î²3 integrin gene variants in Autism Spectrum Disorders â€" Diagnosis and symptomatology. Gene, 2014, 553, 24-30.	2.2	26
129	Lack of association between thrombophilic gene variants and recurrent pregnancy loss. Human Fertility, 2014, 17, 99-105.	1.7	17
130	Health needs assessment for congenital anomalies in middle-income countries: Examining the case for neural tube defects in Brazil. Journal of Community Genetics, 2014, 5, 147-155.	1.2	9
131	Spatial and temporal analysis of infant mortality from congenital malformations in Brazil (1996â€"2010). Journal of Community Genetics, 2014, 5, 269-282.	1.2	6
132	Polymorphisms in the endothelial nitric oxide synthase gene in thalidomide embryopathy. Nitric Oxide - Biology and Chemistry, 2013, 35, 89-92.	2.7	13
133	So Close, So Far Away: Analysis of Surnames in a Town of Twins (CÃ¢ndido GodÃ³i, Brazil). Annals of Human Genetics, 2013, 77, 125-136.	0.8	7
134	Recognition of the phenotype of thalidomide embryopathy in countries endemic for leprosy. Clinical Dysmorphology, 2013, 22, 59-63.	0.3	31
135	MSX1 and PAX9 Investigation in Monozygotic Twins With Variable Expression of Tooth Agenesis. Twin Research and Human Genetics, 2013, 16, 1112-1116.	0.6	3
136	Maternal SNPs in the p53 Pathway: Risk Factors for Trisomy 21?. Disease Markers, 2013, 34, 41-49.	1.3	7
137	High twinning rate in Candido Godoi: a new role for p53 in human fertility. Human Reproduction, 2012, 27, 2866-2871.	0.9	19
138	Prevalence of ERÎ±-397 PvuII C/T, ERÎ±-351 XbaI A/G and PGR PROGINS polymorphisms in Brazilian breast cancer-unaffected women. Brazilian Journal of Medical and Biological Research, 2012, 45, 891-897.	1.5	5
139	Maternal drinking behavior and Fetal Alcohol Spectrum Disorders in adolescents with criminal behavior in southern Brazil. Genetics and Molecular Biology, 2012, 35, 960-965.	1.3	21
140	GSTM1, GSTT1, and GSTP1 polymorphisms, breast cancer risk factors and mammographic density in women submitted to breast cancer screening. Revista Brasileira De Epidemiologia, 2012, 15, 246-255.	0.8	20
141	Presymptomatic Testing for Neurogenetic Diseases in Brazil: Assessing Who Seeks and Who Follows through with Testing. Journal of Genetic Counseling, 2012, 21, 101-112.	1.6	25
142	Prevalence of the STK15 F31I polymorphism and its relationship with mammographic density. Brazilian Journal of Medical and Biological Research, 2011, 44, 291-296.	1.5	6
143	Epidemiological Surveillance of Birth Defects Compatible with Thalidomide Embryopathy in Brazil. PLoS ONE, 2011, 6, e21735.	2.5	30
144	Twin Town in South Brazil: A Nazi's Experiment or a Genetic Founder Effect?. PLoS ONE, 2011, 6, e20328.	2.5	15

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145	MTHFR C677T is not a risk factor for autism spectrum disorders in South Brazil. Psychiatric Genetics, 2010, 20, 187-189.	1.1	31
146	Maternal Gene Polymorphisms Involved in Folate Metabolism as Risk Factors for Down Syndrome Offspring in Southern Brazil. Disease Markers, 2010, 29, 95-101.	1.3	25
147	Maternal gene polymorphisms involved in folate metabolism as risk factors for Down syndrome offspring in Southern Brazil. Disease Markers, 2010, 29, 95-101.	1.3	18
148	Population prevalence of hereditary breast cancer phenotypes and implementation of a genetic cancer risk assessment program in southern Brazil. Genetics and Molecular Biology, 2009, 32, 447-455.	1.3	17
149	Development and validation of a simple questionnaire for the identification of hereditary breast cancer in primary care. BMC Cancer, 2009, 9, 283.	2.6	61
150	Influence of the 5-HTTLPR polymorphism and environmental risk factors in a Brazilian sample of patients with autism spectrum disorders. Brain Research, 2009, 1267, 9-17.	2.2	27
151	Evaluation of C677T and A1298C polymorphisms of the <i>MTHFR</i> gene as maternal risk factors for Down syndrome and congenital heart defects. American Journal of Medical Genetics, Part A, 2009, 149A, 2080-2087.	1.2	59
152	Dipyrone use during pregnancy and adverse perinatal events. Archives of Gynecology and Obstetrics, 2009, 279, 293-297.	1.7	26
153	Consistency of self-reported first-degree family history of cancer in a population-based study. Familial Cancer, 2009, 8, 195-202.	1.9	19
154	Association of the HLAâ€G 14â€bp insertion/deletion polymorphism with juvenile idiopathic arthritis and rheumatoid arthritis. Tissue Antigens, 2008, 71, 440-446.	1.0	64
155	Prospective evaluation of pregnant women vaccinated against rubella in southern Brazil. Reproductive Toxicology, 2008, 25, 120-123.	2.9	32
156	Detection of R337H, a germline TP53 mutation predisposing to multiple cancers, in asymptomatic women participating in a breast cancer screening program in Southern Brazil. Cancer Letters, 2008, 261, 21-25.	7.2	94
157	A exposiÃ§Ã£o prÃ©-natal ao Ã¡lcool como fator de risco para comportamentos disfuncionais: o papel do pediatra. Jornal De Pediatria, 2008, 84, S76-S79.	2.0	6
158	Prenatal alcohol exposure as a risk factor for dysfunctional behaviors: the role of the pediatrician. Jornal De Pediatria, 2008, 84, S76-9.	2.0	4
159	Cancer Genetic Counseling in Public Health Care Hospitals: The Experience of Three Brazilian Services. Public Health Genomics, 2007, 10, 110-119.	1.0	13
160	New cases of thalidomide embryopathy in Brazil. Birth Defects Research Part A: Clinical and Molecular Teratology, 2007, 79, 671-672.	1.6	61
161	Clinical Characterization and Risk Profile of Individuals Seeking Genetic Counseling for Hereditary Breast Cancer in Brazil. Journal of Genetic Counseling, 2007, 16, 363-371.	1.6	22
162	Increased nuchal translucency in arthrogryposis, renal dysfunction and cholestasis (ARC) syndrome and discovery of a Portuguese specific mutation in theVPS33B gene. Ultrasound in Obstetrics and Gynecology, 2006, 28, 233-234.	1.7	10

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163	Reproductive results associated with misoprostol and other substances utilized for interruption of pregnancy. European Journal of Clinical Pharmacology, 2005, 61, 71-72.	1.9	8
164	The beliefs of mothers in southern Brazil regarding risk-factors associated with congenital abnormalities. Genetics and Molecular Biology, 2004, 27, 147-153.	1.3	9
165	Fetal safety of loratadine use in the first trimester of pregnancy: A multicenter study. Journal of Allergy and Clinical Immunology, 2003, 111, 479-483.	2.9	71
166	Community Diagnosis of Maternal Exposure to Risk Factors for Congenital Defects. Public Health Genomics, 2003, 6, 96-103.	1.0	0
167	Reproductive Risk Factors Related to Socioeconomic Status in Pregnant Women in Southern Brazil. Public Health Genomics, 2003, 6, 77-83.	1.0	2
168	Metoclopramide for Nausea and Vomiting of Pregnancy: A Prospective Multicenter International Study. American Journal of Perinatology, 2002, 19, 311-316.	1.4	65
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