Gil Guerra-Junior

List of Publications by Year
in descending order

203	4,171	270111 25	182931 54
papers	citations	h-index	g-index

238	238	238	4963
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Variants in 46,XY DSD-Related Genes in Syndromic and Non-Syndromic Small for Gestational Age Children with Hypospadias. Sexual Development, 2022, 16, 27-33.	1.1	3
2	Growth, puberty and testicular function in boys born small for gestational age with a nonspecific disorder of sex development. Clinical Endocrinology, 2022, 96, 165-174.	1.2	6
3	Effect of Physical Training on Body Composition in Brazilian Military. International Journal of Environmental Research and Public Health, 2022, 19, 1732.	1.2	5
4	Leydig and Sertoli cell function in individuals with genital ambiguity, 46,XY karyotype, palpable gonads and normal testosterone secretion: a case-control study. Sao Paulo Medical Journal, 2022, , .	0.4	1
5	Estimation of body fat in children with intellectual disability: development and cross-validation of a simple anthropometric method. Jornal De Pediatria, 2022, , .	0.9	2
6	Trends in Time Regarding Sex Assignment of Patients with Disorders of Sex Development: Experience of an Interdisciplinary Service. Sexual Development, 2022, 16, 236-241.	1.1	0
7	Rationale and design of the Brazilian diabetes study: a prospective cohort of type 2 diabetes. Current Medical Research and Opinion, 2022, 38, 523-529.	0.9	3
8	Suggested Cutoff Point for Testosterone by Liquid Chromatography with Tandem Mass Spectrometry (LC-MS/MS) after Stimulation with Recombinant Human Chorionic Gonadotropin. Sexual Development, 2022, 16, 266-269.	1.1	2
9	Can the sedentary behavior of basketball and volleyball players impact bone mass and bone geometry?. International Journal of Sports Science and Coaching, 2022, 17, 1428-1434.	0.7	0
10	Can Non-Coding NR5A1 Gene Variants Explain Phenotypes of Disorders of Sex Development?. Sexual Development, 2022, 16, 252-260.	1.1	4
11	Neck circumference and excess weight: proposal of cutoff points for Brazilian adolescents. Jornal De Pediatria, 2021, 97, 191-196.	0.9	4
12	Fat Distribution and Lipid Profile of Young Adults with Congenital Adrenal Hyperplasia Due to 21â€Hydroxylase Enzyme Deficiency. Lipids, 2021, 56, 101-110.	0.7	5
13	Insulin Resistance in Congenital Adrenal Hyperplasia is Compensated for by Reduced Insulin Clearance. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1574-e1585.	1.8	7
14	Bioelectrical Impedance Phase Angle and Its Determinants in Patients with Classic Congenital Adrenal Hyperplasia. Journal of the American College of Nutrition, 2021, , 1-8.	1.1	1
15	Cardiovascular dysfunction risk in young adults with congenital adrenal hyperplasia caused by 21â€hydroxylase enzyme deficiency. International Journal of Clinical Practice, 2021, 75, e14233.	0.8	4
16	Dapagliflozin increases the lean-to total mass ratio in type 2 diabetes mellitus. Nutrition and Diabetes, 2021, 11, 17.	1.5	8
17	Adult Height in 299 Patients with Turner Syndrome with or without Growth Hormone Therapy: Results and Literature Review. Hormone Research in Paediatrics, 2021, 94, 63-70.	0.8	6
18	Normal bone health in young adults with 21-hydroxylase enzyme deficiency undergoing glucocorticoid replacement therapy. Osteoporosis International, 2021, , 1.	1.3	1

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19	The bone densitometry is normal in Turner syndrome prepubertal patients after height age correction. Revista Brasileira De Saude Materno Infantil, 2021, 21, 1129-1134.	0.2	0
20	Promises and pitfalls of whole-exome sequencing exemplified by a nephrotic syndrome family. Molecular Genetics and Genomics, 2020, 295, 135-142.	1.0	2
21	Mutation update for theNR5A1gene involved in DSD and infertility. Human Mutation, 2020, 41, 58-68.	1.1	52
22	Why pediatricians need to know the disorders of sex development: experience of 709 cases in a specialized service. Jornal De Pediatria, 2020, 96, 607-613.	0.9	3
23	Why pediatricians need to know the disorders of sex development: experience of 709 cases in a specialized service. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2020, 96, 607-613.	0.2	0
24	Geometric indices of femur bone strength in female handball players. Women and Health, 2020, 60, 1118-1128.	0.4	2
25	Phase Angle is Determined by Body Composition and Cardiorespiratory Fitness in Adolescents. International Journal of Sports Medicine, 2020, 41, 610-615.	0.8	15
26	Predicting the rate of oxygen uptake from step counts using ActiGraph waistâ€worn accelerometers in adults with Down syndrome. Journal of Intellectual Disability Research, 2020, 64, 602-611.	1.2	3
27	Prevalence of overweight in adolescents from a Southern Brazilian city according to different anthropometric indexes. Revista Paulista De Pediatria, 2020, 39, e2019277.	0.4	2
28	Androgens by immunoassay and mass spectrometry in children with 46,XY disorder of sex development. Endocrine Connections, 2020, 9, 1085-1094.	0.8	6
29	Association between Down Syndrome and Disorders of Sex Development: Report of Three Cases and Review of 188 Cases in the Literature. Sexual Development, 2020, 14, 3-11.	1.1	0
30	Adipokines in young survivors of childhood acute lymphocytic leukemia revisited: beyond fat mass. Annals of Pediatric Endocrinology and Metabolism, 2020, 25, 174-181.	0.8	2
31	Comparing Motor Competence of Sex- and Age-Matched Youth With Intellectual Disability From Brazil and the United States. Adapted Physical Activity Quarterly, 2020, 37, 423-440.	0.6	3
32	Methods for data analysis of resting energy expenditure measured using indirect calorimetry. Nutrition, 2019, 59, 44-49.	1.1	12
33	Assessment of dapagliflozin effect on diabetic endothelial dysfunction of brachial artery (ADDENDA-BHS2 trial): rationale, design, and baseline characteristics of a randomized controlled trial. Diabetology and Metabolic Syndrome, 2019, 11, 62.	1.2	9
34	Novel non-classic CYP21A2 variants, including combined alleles, identified in patients with congenital adrenal hyperplasia. Clinical Biochemistry, 2019, 73, 50-56.	0.8	2
35	A group of Brazilian Turner syndrome patients: Better quality of life than the control group. American Journal of Medical Genetics, Part A, 2019, 179, 2196-2201.	0.7	3
36	EFFECTS OF COMBAT SPORTS ON BONE MASS: SYSTEMATIC REVIEW. Revista Brasileira De Medicina Do Esporte, 2019, 25, 240-244.	0.1	3

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37	Physical training over 6 months is associated with improved changes in phase angle, body composition, and blood glucose in healthy young males. American Journal of Human Biology, 2019, 31, e23275.	0.8	14
38	Effect of 7 Months of Physical Training and Military Routine on the Bone Mass of Young Adults. Military Medicine, 2019, 184, e353-e359.	0.4	3
39	Clinical Findings and Follow-Up of 46,XY and 45,X/46,XY Testicular Dysgenesis. Sexual Development, 2019, 13, 171-177.	1.1	16
40	Initial evidence for hypothalamic gliosis in children with obesity by quantitative T2 MRI and implications for blood oxygenâ€level dependent response to glucose ingestion. Pediatric Obesity, 2019, 14, e12486.	1.4	30
41	Influence of physical training on bone mineral density in healthy young adults: a systematic review. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2019, 65, 1102-1106.	0.3	2
42	Imaging Techniques in the Diagnostic Journey of Disorders of Sex Development. Sexual Development, 2018, 12, 95-99.	1.1	18
43	Accuracy of Bioelectrical Impedance Analysis in Estimated Longitudinal Fat-Free Mass Changes in Male Army Cadets. Military Medicine, 2018, 183, e324-e331.	0.4	9
44	Systematic review of quality of life in Turner syndrome. Quality of Life Research, 2018, 27, 1985-2006.	1.5	16
45	Clinical and cytogenetic features of 516 patients with suspected Turner syndrome â€“ a single-center experience. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 167-173.	0.4	3
46	Comparison between two inhibin B ELISA assays in 46,XY testicular disorders of sex development (DSD) with normal testosterone secretion. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 191-194.	0.4	0
47	Functional characterization of fiveNR5A1gene mutations found in patients with 46,XY disorders of sex development. Human Mutation, 2018, 39, 114-123.	1.1	12
48	Can anthropometric, body composition, and bone variables be considered risk factors for musculoskeletal injuries in Brazilian military students?. BMC Musculoskeletal Disorders, 2018, 19, 377.	0.8	5
49	The heterogeneity of autoimmune polyendocrine syndrome type 1: Clinical features, new mutations and cytokine autoantibodies in a Brazilian cohort from tertiary care centers. Clinical Immunology, 2018, 197, 231-238.	1.4	9
50	A Search for Disorders of Sex Development among Infertile Men. Sexual Development, 2018, 12, 275-280.	1.1	5
51	Adaptive thermogenesis and changes in body composition and physical fitness in army cadets. Journal of Sports Medicine and Physical Fitness, 2018, 59, 94-101.	0.4	5
52	Three new Brazilian cases of 17Î±-hydroxylase deficiency: clinical, molecular, hormonal, and treatment features. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 937-942.	0.4	9
53	Dopamine D2 receptor gene polymorphisms and externalizing behaviors in children and adolescents. BMC Medical Genetics, 2018, 19, 65.	2.1	5
54	FSH may be a useful tool to allow early diagnosis of Turner syndrome. BMC Endocrine Disorders, 2018, 18, 8.	0.9	11

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55	Prevalence of Testicular Adrenal Rest Tumor and Factors Associated with Its Development in Congenital Adrenal Hyperplasia. Hormone Research in Paediatrics, 2018, 90, 161-168.	0.8	17
56	Visfatin is a positive predictor of bone mineral density in young survivors of acute lymphocytic leukemia. Journal of Bone and Mineral Metabolism, 2017, 35, 73-82.	1.3	9
57	Body mass index reference charts for the individuals with Down syndrome aged 2â€18 years. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2017, 93, 94-99.	0.2	0
58	Dual-energy x-ray absorptiometry metabolic maps to resting energy expenditure estimation, and body size dependence in Brazilian young men. Nutrition, 2017, 39-40, 43-49.	1.1	1
59	Growth charts for Brazilian children with Down syndrome: Birth to 20 years of age. Journal of Epidemiology, 2017, 27, 265-273.	1.1	12
60	Quantitative Ultrasonography Measurements of the Phalanges in Adolescents: A Mixed Longitudinal Study. Ultrasound in Medicine and Biology, 2017, 43, 2934-2938.	0.7	2
61	Functional Impact of Novel Androgen Receptor Mutations on the Clinical Manifestation of Androgen Insensitivity Syndrome. Sexual Development, 2017, 11, 238-247.	1.1	9
62	A study of splicing mutations in disorders of sex development. Scientific Reports, 2017, 7, 16202.	1.6	5
63	Serum Concentration of Risperidone and Adverse Effects in Children and Adolescents. Journal of Child and Adolescent Psychopharmacology, 2017, 27, 211-212.	0.7	5
64	Body mass index reference charts for individuals with Down syndrome aged 2â€“18 years. Jornal De Pediatria, 2017, 93, 94-99.	0.9	12
65	NPHS2Mutations: A Closer Look to Latin American Countries. BioMed Research International, 2017, 2017, 1-6.	0.9	2
66	The effects of an after-school intervention program on physical activity level, sedentary time, and cardiovascular risk factors in adolescents. Motriz Revista De Educacao Fisica, 2017, 23, .	0.3	4
67	Functional and Structural Consequences of NineCYP21A2Mutations Ranging from Very Mild to Severe Effects. International Journal of Endocrinology, 2016, 2016, 1-10.	0.6	8
68	408 Cases of Genital Ambiguity Followed by Single Multidisciplinary Team during 23 Years: Etiologic Diagnosis and Sex of Rearing. International Journal of Endocrinology, 2016, 2016, 1-9.	0.6	23
69	Pharmacogenetics of Risperidone and Cardiovascular Risk in Children and Adolescents. International Journal of Endocrinology, 2016, 2016, 1-10.	0.6	18
70	Accuracy of Body Mass Index Cutoffs for Classifying Obesity in Chilean Children and Adolescents. International Journal of Environmental Research and Public Health, 2016, 13, 472.	1.2	2
71	Validity of Bioelectrical Impedance Analysis to Estimation Fat-Free Mass in the Army Cadets. Nutrients, 2016, 8, 121.	1.7	27
72	Central precocious puberty: revisiting the diagnosis and therapeutic management. Archives of Endocrinology and Metabolism, 2016, 60, 163-172.	0.3	76

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73	NPHS1 gene mutations confirm congenital nephrotic syndrome in four Brazilian cases: A novel mutation is described. Nephrology, 2016, 21, 753-757.	0.7	10
74	A Single Nucleotide Variant in the Promoter Region of 17Î²-HSD Type 5 Gene Influences External Genitalia Virilization in Females with 21-Hydroxylase Deficiency. Hormone Research in Paediatrics, 2016, 85, 333-338.	0.8	5
75	Competitive Swimming and Handball Participation Have a Positive Influence on Bone Parameters as Assessed by Phalangeal Quantitative Ultrasound in Female Adolescents. Pediatric Exercise Science, 2016, 28, 423-430.	0.5	3
76	<i>NR5A1</i> Loss-of-Function Mutations Lead to 46,XY Partial Gonadal Dysgenesis Phenotype: Report of Three Novel Mutations. Sexual Development, 2016, 10, 191-199.	1.1	18
77	The Long-Term Outcome of Boys With Partial Androgen Insensitivity Syndrome and a Mutation in the Androgen Receptor Gene. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 3959-3967.	1.8	81
78	Overweight and obesity in children and adolescents with Down syndromeâ€"prevalence, determinants, consequences, and interventions: A literature review. Research in Developmental Disabilities, 2016, 57, 181-192.	1.2	196
79	Global Disorders of Sex Development Update since 2006: Perceptions, Approach and Care. Hormone Research in Paediatrics, 2016, 85, 158-180.	0.8	852
80	Lipomatous nevus and urethral caruncle mistaken for ambiguous genitalia in a female infant. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 233-6.	0.4	1
81	New approach to phenotypic variability and karyotype-phenotype correlation in Turner syndrome. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 475-9.	0.4	23
82	Minimum Time to Achieve the Steady State and Optimum Abbreviated Period to Estimate the Resting Energy Expenditure by Indirect Calorimetry in Healthy Young Adults. Nutrition in Clinical Practice, 2016, 31, 349-354.	1.1	32
83	A new compound heterozygosis for inactivating mutations in the glucokinase gene as cause of permanent neonatal diabetes mellitus (PNDM) in double-first cousins. Diabetology and Metabolic Syndrome, 2015, 7, 101.	1.2	12
84	NPHS2 mutations account for only 15Â% of nephrotic syndrome cases. BMC Medical Genetics, 2015, 16, 88.	2.1	18
85	Subcutaneous adipose tissue plays a beneficial effect on subclinical atherosclerosis in young survivors of acute lymphocytic leukemia. Vascular Health and Risk Management, 2015, 11, 479.	1.0	10
86	Brazilian Pediatric Reference Data for Quantitative Ultrasound of Phalanges According to Gender, Age, Height and Weight. PLoS ONE, 2015, 10, e0127294.	1.1	8
87	Hyperprolactinemia in Children and Adolescents with Use of Risperidone: Clinical and Molecular Genetics Aspects. Journal of Child and Adolescent Psychopharmacology, 2015, 25, 738-748.	0.7	30
88	Effects of programmed physical activity on body composition in post-pubertal schoolchildren. Jornal De Pediatria, 2015, 91, 122-129.	0.9	24
89	Ovotesticular disorder of sex development with unusual karyotype: patient report. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 677-80.	0.4	7
90	Two Novel Mutations in the Thyroid Hormone Receptor Î² in Patients with Resistance to Thyroid Hormone (RTH Î²): Clinical, Biochemical, and Molecular Data. Hormone and Metabolic Research, 2015, 47, 889-894.	0.7	5

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91	Parentsâ€™ experiences of having a baby with ambiguous genitalia. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 833-8.	0.4	13
92	ComparaÃ§Ã£o do estado nutricional de meninas de acordo com diferentes referÃªncias para Ãndice de massa corporal. Revista Brasileira De Saude Materno Infantil, 2015, 15, 121-129.	0.2	2
93	Efetividade de programas de intervenÃ§Ã£o escolar para reduzir fatores de risco Ã saÃºde em adolescentes: uma revisÃ£o sistemÃ¡tica. Revista Brasileira De Cineantropometria E Desempenho Humano, 2015, 17, 485.	0.5	2
94	Growth Curves for Girls with Turner Syndrome. BioMed Research International, 2014, 2014, 1-8.	0.9	16
95	Prader-Willi syndrome: a case report with atypical developmental features. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 983-8.	0.4	1
96	Long-Term Follow-Up of Patients with 46,XY Partial Gonadal Dysgenesis Reared as Males. International Journal of Endocrinology, 2014, 2014, 1-8.	0.6	21
97	Homozygous Inactivating Mutation inNANOS3in Two Sisters with Primary Ovarian Insufficiency. BioMed Research International, 2014, 2014, 1-8.	0.9	36
98	Genetic Predictors of Long-Term Response to Growth Hormone (GH) Therapy in Children With GH Deficiency and Turner Syndrome: The Influence of a SOCS2 Polymorphism. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E1808-E1813.	1.8	21
99	Preserved Fertility in a Patient with Gynecomastia Associated with the p.Pro695Ser Mutation in the Androgen Receptor. Sexual Development, 2014, 8, 350-355.	1.1	8
100	The novel p.Cys65Tyr mutation in NR5A1gene in three 46,XY siblings with normal testosterone levels and their mother with primary ovarian insufficiency. BMC Medical Genetics, 2014, 15, 7.	2.1	25
101	Growth curves in Down syndrome: Implications for clinical practice. American Journal of Medical Genetics, Part A, 2014, 164, 844-847.	0.7	27
102	Growth and body composition in Brazilian female rhythmic gymnastics athletes. Journal of Sports Sciences, 2014, 32, 1790-1796.	1.0	14
103	Central Precocious Puberty That Appears to Be Sporadic Caused by Paternally Inherited Mutations in the Imprinted Gene Makorin Ring Finger 3. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E1097-E1103.	1.8	126
104	Performance of Phalangeal Quantitative Ultrasound Parameters in the Evaluation of Reduced Bone Mineral Density Assessed By DX in Patients with 21 Hydroxylase Deficiency. Ultrasound in Medicine and Biology, 2014, 40, 1414-1419.	0.7	3
105	Cranial radiotherapy predisposes to abdominal adiposity in survivors of childhood acute lymphocytic leukemia. Radiation Oncology, 2013, 8, 39.	1.2	15
106	Cross-sectional study of the association of body composition and physical fitness with bone status in children and adolescents from 11 to 16Âyears old. BMC Pediatrics, 2013, 13, 117.	0.7	14
107	Screening of Y chromosome microdeletions in 46,XY partial gonadal dysgenesis and in patients with a 45,X/46,XY karyotype or its variants. BMC Medical Genetics, 2013, 14, 115.	2.1	17
108	Secular trends of height, weight and BMI in young adult Brazilian military students in the 20th century. Annals of Human Biology, 2013, 40, 554-556.	0.4	5

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109	Two distinct WT1 mutations identified in patients and relatives with isolated nephrotic proteinuria. Biochemical and Biophysical Research Communications, 2013, 441, 371-376.	1.0	12
110	Estimation of percent body fat based on anthropometric measurements in children and adolescents with congenital adrenal hyperplasia due to 21-hydroxylase deficiency. Clinical Nutrition, 2013, 32, 45-50.	2.3	0
111	The Sitting Height/Height Ratio for Age in Healthy and Short Individuals and Its Potential Role in Selecting Short Children for <i>SHOX</i> Analysis. Hormone Research in Paediatrics, 2013, 80, 449-456.	0.8	45
112	Clinical and Laboratorial Features That May Differentiate 46,XY DSD due to Partial Androgen Insensitivity and 5Î±-Reductase Type 2 Deficiency. International Journal of Endocrinology, 2012, 2012, 1-7.	0.6	25
113	Disorders of Sex Development and Hypogonadism: Genetics, Mechanism, and Therapies. International Journal of Endocrinology, 2012, 2012, 1-2.	0.6	2
114	Accuracy of anthropometric measurements in estimating fat mass in individuals with 21-hydroxylase deficiency. Nutrition, 2012, 28, 984-990.	1.1	7
115	Analysis of anti-MÃ¼llerian hormone (AMH) and its receptor (AMHR2) genes in patients with persistent MÃ¼llerian duct syndrome. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 473-478.	1.3	21
116	Frasier syndrome: four new cases with unusual presentations. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 525-532.	1.3	14
117	Klinefelter syndrome: an unusual diagnosis in pediatric patients. Jornal De Pediatria, 2012, 88, 323-7.	0.9	9
118	46,XX DSD and Antley-Bixler syndrome due to novel mutations in the cytochrome P450 oxidoreductase gene. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 578-585.	1.3	13
119	Sexual difference, identification and object choice in individuals with sex differentiation disorders. Revista Latinoamericana De Psicopatologia Fundamental, 2012, 15, 464-481.	0.0	1
120	The use of fluorescence in situ hybridization in the diagnosis of hidden mosaicism: apropos of three cases of sex chromosome anomalies. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 545-551.	1.3	11
121	Excesso de peso e fatores associados em adolescentes. Revista De Nutricao, 2012, 25, 229-236.	0.4	8
122	Growth hormone pharmacogenetics: the interactive effect of a microsatellite in the IGF1 promoter region with the GHR-exon 3 and â^'202 A/C IGFBP3 variants on treatment outcomes of children with severe GH deficiency. Pharmacogenomics Journal, 2012, 12, 439-445.	0.9	22
123	Turner syndrome and metabolic derangements: Another example of fetal programming. Early Human Development, 2012, 88, 99-102.	0.8	13
124	Spontaneous puberty in girls with early diagnosis of Turner syndrome. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 653-657.	1.3	17
125	Prevalence of genital abnormalities in neonates. Jornal De Pediatria, 2012, 88, 489-95.	0.9	10
126	Influence of body composition on bone mass in children and adolescents. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira (English Edition), 2011, 57, 648-653.	0.1	0

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127	InfluÃªncia da composiÃ§Ã£o corporal sobre a massa Ã³ssea em crianÃ§as e adolescentes. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2011, 57, 662-667.	0.3	14
128	Complete gonadal dysgenesis in clinical practice: the 46,XY karyotype accounts for more than one third of cases. Fertility and Sterility, 2011, 96, 1431-1434.	0.5	42
129	Multifunctional role of steroidogenic factor 1 and disorders of sex development. Arquivos Brasileiros De Endocrinologia E Metabologia, 2011, 55, 607-612.	1.3	14
130	Metabolic evaluation of young women with congenital adrenal hyperplasia. Arquivos Brasileiros De Endocrinologia E Metabologia, 2011, 55, 646-652.	1.3	5
131	Efeito da atividade fÃsica programada sobre a aptidÃ£o fÃsica em escolares adolescentes DOI:10.5007/1980-0037.2010v12n2p98. Revista Brasileira De Cineantropometria E Desempenho Humano, 2011, 12, .	0.5	4
132	Social skills in women with Turner Syndrome. Scandinavian Journal of Psychology, 2011, 52, 440-447.	0.8	13
133	Severe forms of partial androgen insensitivity syndrome due to p.L830F novel mutation in androgen receptor gene in a Brazilian family. BMC Research Notes, 2011, 4, 173.	0.6	11
134	Growth hormone effect on body composition in Turner syndrome. Endocrine, 2011, 40, 486-491.	1.1	10
135	OCT4 immunohistochemistry may be necessary to identify the real risk of gonadal tumors in patients with Turner syndrome and Y chromosome sequences. Human Reproduction, 2011, 26, 3450-3455.	0.4	25
136	Molecular Diagnosis of 5Î±-Reductase Type II Deficiency in Brazilian Siblings with 46,XY Disorder of Sex Development. International Journal of Molecular Sciences, 2011, 12, 9471-9480.	1.8	3
137	Abnormalities in body composition and nutritional status in HIV-infected children and adolescents on antiretroviral therapy. International Journal of STD and AIDS, 2011, 22, 453-456.	0.5	16
138	Influence of body composition on bone mass in children and adolescents. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2011, 57, 648-653.	0.3	1
139	The novel p.E89K mutation in the SRY gene inhibits DNA binding and causes the 46,XY disorder of sex development. Brazilian Journal of Medical and Biological Research, 2011, 44, 361-365.	0.7	8
140	Fatores associados a atraso no diagnÃ³stico da sÃndrome de Turner. Revista Paulista De Pediatria, 2011, 29, 67-72.	0.4	5
141	Normalization of height and excess body fat in children with salt-wasting 21-hydroxylase deficiency. Jornal De Pediatria, 2011, 87, 263-268.	0.9	7
142	Puberdade precoce central como Ãºnica manifestaÃ§Ã£o de cisto aracnoide supraselar. Revista Paulista De Pediatria, 2011, 29, 126-129.	0.4	0
143	The novel p.E89K mutation in the SRY gene inhibits DNA binding and causes the 46,XY disorder of sex development. Brazilian Journal of Medical and Biological Research, 2011, 44, 361-365.	0.7	3
144	Complete XY gonadal dysgenesis due to p.D293N homozygous mutation in theNR5A1 gene: a case study. Journal of Applied Genetics, 2010, 51, 223-224.	1.0	21

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145	Novel deletion alleles carrying CYP21A1P/A2chimeric genes in Brazilian patients with 21-hydroxylase deficiency. BMC Medical Genetics, 2010, 11, 104.	2.1	14
146	Screening of autosomal gene deletions in patients with hypogonadotropic hypogonadism using multiplex ligationâ€dependent probe amplification: detection of a hemizygosis for the fibroblast growth factor receptor 1. Clinical Endocrinology, 2010, 72, 371-376.	1.2	30
147	46,XY and 45,X/46,XY testicular dysgenesis: similar gonadal and genital phenotype, different prognosis. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 331-334.	1.3	20
148	Structural aspects of the p.P222Q homozygous mutation of HSD3B2 gene in a patient with congenital adrenal hyperplasia. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 768-774.	1.3	10
149	Long-term follow-up of an 8-year-old boy with insulinoma as the first manifestation of a familial form of multiple endocrine neoplasia type 1. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 754-760.	1.3	18
150	Phenotypic variability in a family with x-linked adrenoleukodystrophy caused by the p.Trp132Ter mutation. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 738-743.	1.3	8
151	Novel DMRT1 3'UTR+11insT mutation associated to XY partial gonadal dysgenesis. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 749-753.	1.3	14
152	Cardiovascular and renal anomalies in Turner syndrome. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2010, 56, 655-659.	0.3	35
153	Six new cases confirm the clinical molecular profile of complete combined 17Î±-hydroxylase/ 17,20-lyase deficiency in Brazil. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 711-716.	1.3	12
154	Comparison of bone quantity by ultrasound measurements of phalanges between white and black children living in ParanÃ¡, Brazil, with Europeans. Brazilian Journal of Medical and Biological Research, 2010, 43, 976-981.	0.7	4
155	SÃndrome de Turner: um diagnÃ³stico pediÃ¡trico frequentemente realizado por nÃ£o pediatras. Jornal De Pediatria, 2010, 86, 121-125.	0.9	1
156	A novel homozygous splice acceptor site mutation of KISS1R in two siblings with normosmic isolated hypogonadotropic hypogonadism. European Journal of Endocrinology, 2010, 163, 29-34.	1.9	50
157	Effects of growth hormone on body proportions in Turner syndrome compared with non-treated patients and normal women. Journal of Endocrinological Investigation, 2010, 33, 691-695.	1.8	6
158	Turner syndrome: a pediatric diagnosis frequently made by non-pediatricians. Jornal De Pediatria, 2010, 86, 121-125.	0.9	11
159	Adiposity in childhood cancer survivors: insights into obesity physiopathology. Arquivos Brasileiros De Endocrinologia E Metabologia, 2009, 53, 190-200.	1.3	38
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