Maricilda Palandi de Mello

List of Publications by Year
in descending order

124 papers	1,818 citations	279487 23 h-index	414034 32 g-index
128 all docs	128 docs citations	128 times ranked	2127 citing authors

#	Article	IF	Citations
1	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
2	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
3	Bilateral Wilmsâ€™ tumor in a child with Denys-Drash syndrome: novel frameshift variant disrupts the WT1 nuclear location signaling region. Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 837-843.	0.4	3
4	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
5	APOL1 in an ethnically diverse pediatric population with nephrotic syndrome: implications in focal segmental glomerulosclerosis and other diagnoses. Pediatric Nephrology, 2021, 36, 2327-2336.	0.9	8
6	Promises and pitfalls of whole-exome sequencing exemplified by a nephrotic syndrome family. Molecular Genetics and Genomics, 2020, 295, 135-142.	1.0	2
7	Mutation update for theNR5A1gene involved in DSD and infertility. Human Mutation, 2020, 41, 58-68.	1.1	52
8	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
9	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
10	Androgens by immunoassay and mass spectrometry in children with 46,XY disorder of sex development. Endocrine Connections, 2020, 9, 1085-1094.	0.8	6
11	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
12	Novel non-classic CYP21A2 variants, including combined alleles, identified in patients with congenital adrenal hyperplasia. Clinical Biochemistry, 2019, 73, 50-56.	0.8	2
13	Clinical Findings and Follow-Up of 46,XY and 45,X/46,XY Testicular Dysgenesis. Sexual Development, 2019, 13, 171-177.	1.1	16
14	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
15	Functional characterization of fiveNR5A1gene mutations found in patients with 46,XY disorders of sex development. Human Mutation, 2018, 39, 114-123.	1.1	12
16	A Search for Disorders of Sex Development among Infertile Men. Sexual Development, 2018, 12, 275-280.	1.1	5
17	Association between serotonin 2C receptor gene (HTR2C) polymorphisms and psychopathological symptoms in children and adolescents. Brazilian Journal of Medical and Biological Research, 2018, 51, e7252.	0.7	4
18	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

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19	Dopamine D2 receptor gene polymorphisms and externalizing behaviors in children and adolescents. BMC Medical Genetics, 2018, 19, 65.	2.1	5
20	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
21	Discordant Genotypic Sex and Phenotype Variations in Two Spanish Siblings with 17Î±-Hydroxylase/17,20-Lyase Deficiency Carrying the Most Prevalent Mutated CYP17A1 Alleles of Brazilian Patients. Sexual Development, 2017, 11, 70-77.	1.1	8
22	<i>WT1 </i>Haploinsufficiency Supports Milder Renal Manifestation in Two Patients with Denys-Drash Syndrome. Sexual Development, 2017, 11, 34-39.	1.1	5
23	A Novel Homozygous Missense <i>FSHR</i> Variant Associated with Hypergonadotropic Hypogonadism in Two Siblings from a Brazilian Family. Sexual Development, 2017, 11, 137-142.	1.1	26
24	Development of CYP21A2 Genotyping Assay for the Diagnosis of Congenital Adrenal Hyperplasia. Molecular Diagnosis and Therapy, 2017, 21, 663-675.	1.6	15
25	Functional Impact of Novel Androgen Receptor Mutations on the Clinical Manifestation of Androgen Insensitivity Syndrome. Sexual Development, 2017, 11, 238-247.	1.1	9
26	A study of splicing mutations in disorders of sex development. Scientific Reports, 2017, 7, 16202.	1.6	5
27	Serum Concentration of Risperidone and Adverse Effects in Children and Adolescents. Journal of Child and Adolescent Psychopharmacology, 2017, 27, 211-212.	0.7	5
28	NPHS2Mutations: A Closer Look to Latin American Countries. BioMed Research International, 2017, 2017, 1-6.	0.9	2
29	Functional and Structural Consequences of NineCYP21A2Mutations Ranging from Very Mild to Severe Effects. International Journal of Endocrinology, 2016, 2016, 1-10.	0.6	8
30	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
31	Pharmacogenetics of Risperidone and Cardiovascular Risk in Children and Adolescents. International Journal of Endocrinology, 2016, 2016, 1-10.	0.6	18
32	NPHS1 gene mutations confirm congenital nephrotic syndrome in four Brazilian cases: A novel mutation is described. Nephrology, 2016, 21, 753-757.	0.7	10
33	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
34	<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
35	A Cytogenomic Approach in a Case of Syndromic XY Gonadal Dysgenesis. Sexual Development, 2016, 10, 23-27.	1.1	1
36	GAPO syndrome: a new syndromic cause of premature ovarian insufficiency. Climacteric, 2016, 19, 594-598.	1.1	13

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37	A de novo mutation in CYP21A2 gene in a case of in vitro fertilization. Molecular Genetics and Metabolism Reports, 2015, 5, 98-102.	0.4	2
38	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
39	NPHS2 mutations account for only 15Â% of nephrotic syndrome cases. BMC Medical Genetics, 2015, 16, 88.	2.1	18
40	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
41	In vitro functional studies of rare <scp>CYP</scp>21A2 mutations and establishment of an activity gradient for nonclassic mutations improve phenotype predictions in congenital adrenal hyperplasia. Clinical Endocrinology, 2015, 82, 37-44.	1.2	22
42	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
43	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
44	Homozygous Inactivating Mutation inNANOS3in Two Sisters with Primary Ovarian Insufficiency. BioMed Research International, 2014, 2014, 1-8.	0.9	36
45	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
46	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
47	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
48	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
49	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
50	New mutation in the myocilin gene segregates with juvenile-onset open-angle glaucoma in a Brazilian family. Gene, 2013, 523, 50-57.	1.0	20
51	The effect of fetal androgen metabolismâ€related gene variants on external genitalia virilization in congenital adrenal hyperplasia. Clinical Genetics, 2013, 84, 482-488.	1.0	16
52	Leydig Cell Tumour in a 46,XX Child with Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency. Hormone Research in Paediatrics, 2013, 79, 179-184.	0.8	4
53	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
54	Accuracy of anthropometric measurements in estimating fat mass in individuals with 21-hydroxylase deficiency. Nutrition, 2012, 28, 984-990.	1.1	7

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55	Clinical and molecular spectrum of patients with 17Î²-hydroxysteroid dehydrogenase type 3 (17-Î²-HSD3) deficiency. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 533-539.	1.3	29
56	Frasier syndrome: four new cases with unusual presentations. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 525-532.	1.3	14
57	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
58	Functional studies of CYP21A2 mutants complement structural and clinical predictions of disease severity in CAH. Clinical Endocrinology, 2012, 76, 766-768.	1.2	4
59	Prevalence of genital abnormalities in neonates. Jornal De Pediatria, 2012, 88, 489-95.	0.9	10
60	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
61	Multifunctional role of steroidogenic factor 1 and disorders of sex development. Arquivos Brasileiros De Endocrinologia E Metabologia, 2011, 55, 607-612.	1.3	14
62	Metabolic evaluation of young women with congenital adrenal hyperplasia. Arquivos Brasileiros De Endocrinologia E Metabologia, 2011, 55, 646-652.	1.3	5
63	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
64	A de novo cryptic 5p deletion and 9p duplication detected by subtelomeric MLPA in a boy with cri du chat syndrome. American Journal of Medical Genetics, Part A, 2011, 155, 450-454.	0.7	2
65	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
66	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
67	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
68	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
69	The Novel WT1 Gene Mutation p.H377N Associated to Denys-Drash Syndrome. Journal of Pediatric Hematology/Oncology, 2010, 32, 486-488.	0.3	2
70	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
71	Novel deletion alleles carrying CYP21A1P/A2chimeric genes in Brazilian patients with 21-hydroxylase deficiency. BMC Medical Genetics, 2010, 11, 104.	2.1	14
72	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

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73	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
74	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
75	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
76	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
77	46,XX Male - Testicular Disorder of Sexual Differentiation (DSD): hormonal, molecular and cytogenetic studies. Arquivos Brasileiros De Endocrinologia E Metabologia, 2010, 54, 685-689.	1.3	16
78	Impairment in Anthropometric Parameters and Body Composition in Females with Classical 21-Hydroxylase Deficiency. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 519-29.	0.4	7
79	Novel Mutations inCYP11B1Gene Leading to 11Î²-Hydroxylase Deficiency in Brazilian Patients. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 3481-3485.	1.8	28
80	T allele of â^'344C/T polymorphism in aldosterone synthase gene is not associated with resistant hypertension. Hypertension Research, 2009, 32, 159-162.	1.5	13
81	PIP4KIIA and Î²â€globin: transcripts differentially expressed in reticulocytes and associated with high levels of Hb H in two siblings with Hb H disease. European Journal of Haematology, 2009, 83, 490-493.	1.1	9
82	Complement 4 phenotypes and genotypes in Brazilian patients with classical 21-hydroxylase deficiency. Clinical and Experimental Immunology, 2009, 155, 182-188.	1.1	6
83	Inhibition of CYP21A2 Enzyme Activity Caused by Novel Missense Mutations Identified in Brazilian and Scandinavian Patients. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 2416-2420.	1.8	59
84	21-Hydroxylase Deficiency Transiently Mimicking Combined 21- and 11Î²-Hydroxylase Deficiency. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 487-94.	0.4	3
85	XX Maleness and XX True Hermaphroditism inSRY-Negative Monozygotic Twins: Additional Evidence for a Common Origin. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 339-343.	1.8	44
86	Apparent mineralocorticoid excess syndrome in a Brazilian boy caused by the homozygous missense mutation p.R186C in the HSD11B2 gene. Arquivos Brasileiros De Endocrinologia E Metabologia, 2008, 52, 1277-1281.	1.3	10
87	Absence of mutations in Pax6 gene in three cases of Morning Glory syndrome associated with isolated growth hormone deficiency. Arquivos Brasileiros De Endocrinologia E Metabologia, 2008, 52, 1221-1227.	1.3	12
88	Clinical and genetic findings of five patients with WT1-related disorders. Arquivos Brasileiros De Endocrinologia E Metabologia, 2008, 52, 1236-1243.	1.3	16
89	Mutations in the vitamin D receptor gene in four patients with hereditary 1,25-dihydroxyvitamin D-resistant rickets. Arquivos Brasileiros De Endocrinologia E Metabologia, 2008, 52, 1244-1251.	1.3	24
90	Heterozygosis for CYP21A2 mutation considered as 21-hydroxylase deficiency in neonatal screening. Arquivos Brasileiros De Endocrinologia E Metabologia, 2008, 52, 1388-1392.	1.3	5

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91	An illustrative case of LÃ©ri-Weill dyschondrosteosis. Genetics and Molecular Biology, 2008, 31, 839-842.	0.6	1
92	Pipkiia and Beta-Globin: Transcripts Differentially Expressed in Reticulocytes and Associated with High Levels of Hb H in Two Siblings with Hb H Disease.. Blood, 2007, 110, 3821-3821.	0.6	0
93	Association between theMTHFRA1298C polymorphism and increased risk of acute myeloid leukemia in Brazilian children. Leukemia and Lymphoma, 2006, 47, 2070-2075.	0.6	20
94	Q289P Mutation in FGFR2 Gene Causes Saethre-Chotzen Syndrome: Some Considerations About Familial Heterogeneity. Cleft Palate-Craniofacial Journal, 2006, 43, 142-147.	0.5	15
95	Mutations in SRY and WT1 genes required for gonadal development are not responsible for XY partial gonadal dysgenesis. Brazilian Journal of Medical and Biological Research, 2005, 38, 17-25.	0.7	17
96	A naturally occurring deletion in the SRY promoter region affecting the Sp1 binding site is associated with sex reversal. Journal of Endocrinological Investigation, 2005, 28, 651-656.	1.8	19
97	Genetic variation and relationships at six VNTR loci in two distinct sample populations in Brazil. Annals of Human Biology, 2004, 31, 660-668.	0.4	3
98	Clinical Findings in Four Brazilian Families Affected by Saethre-Chotzen Syndrome without TWIST Mutations. Cleft Palate-Craniofacial Journal, 2004, 41, 250-255.	0.5	7
99	Classical congenital adrenal hyperplasia due to 21-hydroxylase deficiency: a cross-sectional study of factors involved in bone mineral density. Journal of Bone and Mineral Metabolism, 2003, 21, 396-401.	1.3	46
100	Iron-responsive genes of Phanerochaete chrysosporium isolated by differential display reverse transcription polymerase chain reaction. Environmental Microbiology, 2003, 5, 777-786.	1.8	11
101	Role of Metals in Wood Biodegradation. ACS Symposium Series, 2003, , 154-174.	0.5	14
102	Phenotype and genotype correlation of the microconversion from the CYP21A1P to the CYP21A2 gene in congenital adrenal hyperplasia. Brazilian Journal of Medical and Biological Research, 2003, 36, 1311-1318.	0.7	36
103	Molecular and Biochemical Screening for the Diagnosis and Management of Medullary Thyroid Carcinoma in Multiple Endocrine Neoplasia Type 2A. Hormone and Metabolic Research, 2002, 34, 202-206.	0.7	14
104	Differential gene expression in response to copper inAcidithiobacillus ferrooxidans analyzed by RNA arbitrarily primed polymerase chain reaction. Electrophoresis, 2002, 23, 520-527.	1.3	35
105	Iron-regulated proteins inPhanerochaete chrysosporium andLentinula edodes: Differential analysis by sodium dodecyl sulfate polyacrylamide gel electrophoresis and two-dimensional polyacrylamide gel electrophoresis profiles. Electrophoresis, 2002, 23, 655-661.	1.3	31
106	Identification of a neocentromere in a rearranged Y chromosome with no detectable DYZ3 centromeric sequence. American Journal of Medical Genetics Part A, 2002, 113, 263-267.	2.4	8
107	Novel mutations affecting SRY DNA-binding activity: the HMG box N65H associated with 46,XY pure gonadal dysgenesis and the familial non-HMG box R30I associated with variable phenotypes. Journal of Molecular Medicine, 2002, 80, 782-790.	1.7	68
108	Molecular characterization of Acidithiobacillus ferrooxidans and A. thiooxidans strains isolated from mine wastes in Brazil. Antonie Van Leeuwenhoek, 2001, 80, 65-75.	0.7	20

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109	H28+C Insertion in the CYP21 Gene: A Novel Frameshift Mutation in a Brazilian Patient with the Classical Form of 21-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5877-5880.	1.8	26
110	Morphometry and Histology of Gonads From 13 Children With Dysgenetic Male Pseudohermaphroditism. Archives of Pathology and Laboratory Medicine, 2001, 125, 652-656.	1.2	25
111	Molecular mapping of an idic(Yp) chromosome in an Ullrich-Turner patient. , 2000, 91, 95-98.		8
112	Recurrence of a Nonsense Mutation in the Conserved Domain of SRY in a Brazilian Patient with 46,XY Gonadal Dysgenesis. Journal of Pediatric Endocrinology and Metabolism, 1999, 12, 455-7.	0.4	2
113	Mutation distribution and CYP21/C4 locus variability in Brazilian families with the classical form of the 21-hydroxylase deficiency. Acta Paediatrica, International Journal of Paediatrics, 1999, 88, 275-83.	0.7	24
114	Molecular Variability of Cyp21/C4 Gene Organization and Cyp21B Mutations in Disease-Causing Alleles of Brazilian Families with 21-Hydroxylase Deficiency. Pediatric Research, 1999, 45, 443-443.	1.1	0
115	Two cases of Y; autosome translocations: A 45, X male and a clinically trisomy 18 patient. American Journal of Medical Genetics Part A, 1994, 49, 388-392.	2.4	13
116	2 CLINICAL AND GENETICAL CHARACTERISTICS IN CLASSIC CONGENITAL ADRENAL HYPERPLASIA DUE TO. 21-HYDROXYLASE DEFICIENCY. PRELIMINARY DATA IN 12 BRAZILIAN FAMILIES. Pediatric Research, 1994, 36, 678-678.	1.1	0
117	Repetitive sequences in the ribosomal intergenic spacer of Trypanosoma cruzi. Molecular and Biochemical Parasitology, 1993, 60, 273-280.	0.5	28
118	Identification of new world Leishmania using ribosomal gene spacer probes. Molecular and Biochemical Parasitology, 1992, 56, 15-26.	0.5	46
119	True agonadism: Report of a case analyzed with Y-specific DNA probes. American Journal of Medical Genetics Part A, 1991, 41, 444-445.	2.4	23
120	EXCITATION OF CHLOROPLASTS IN Euglena gracilis IN THE ABSENCE OF LIGHT. Photochemistry and Photobiology, 1988, 47, 457-461.	1.3	18
121	Enzymatically generated electronically excited molecules induce transformation of 4-thiouridine to uridine. Biochemical and Biophysical Research Communications, 1983, 117, 923-929.	1.0	4
122	PEROXIDASE-GENERATED TRIPLET INDOLE-3-ALDEHYDE ADDS TO URIDINE BASES AND EXCITES THE 4-THIOURIDINE GROUP IN t-RNAPhe. Photochemistry and Photobiology, 1982, 36, 21-24.	1.3	28
123	Excited indole-3-aldehyde from the peroxidase-catalyzed aerobic oxidation of indole-3-acetic acid. Reaction with and energy transfer to transfer ribonucleic acid. Biochemistry, 1980, 19, 5270-5275.	1.2	55
124	Busca de novas variantes no gene HSD17B3 em pacientes com distÃºrbios da diferenciaÃ§Ã£o do sexo. , 0, , .		0