Berenice Bilharinho Mendonca

List of Publications by Year
in descending order

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480 papers	19,444 citations	11608 70 h-index	20900 115 g-index
537 all docs	537 docs citations	537 times ranked	12450 citing authors

#	Article	IF	Citations
1	Primary Adrenal Insufficiency Due to Bilateral Adrenal Infarction in COVID-19. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e394-e400.	1.8	24
2	WT1 Pathogenic Variants are Associated with a Broad Spectrum of Differences in Sex Development Phenotypes and Heterogeneous Progression of Renal Disease. Sexual Development, 2022, 16, 46-54.	1.1	5
3	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
4	A Small Supernumerary Xp Marker Chromosome Including Genes <i>NR0B1</i> and <i>MAGEB</i> Causing Partial Gonadal Dysgenesis and Gonadoblastoma. Sexual Development, 2022, 16, 55-63.	1.1	1
5	Genetics of ovarian insufficiency and defects of folliculogenesis. Best Practice and Research in Clinical Endocrinology and Metabolism, 2022, 36, 101594.	2.2	36
6	Contribution of Clinical and Genetic Approaches for Diagnosing 209 Index Cases With 46,XY Differences of Sex Development. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1797-e1806.	1.8	11
7	Novel OTX2 loss of function variant associated with congenital hypopituitarism without eye abnormalities. Journal of Pediatric Endocrinology and Metabolism, 2022, .	0.4	0
8	Adult height of patients with SHOX haploinsufficiency with or without GH therapy: A Real-World Single-Center Study. Hormone Research in Paediatrics, 2022, , .	0.8	2
9	A truncating variant of RAD51B associated with primary ovarian insufficiency provides insights into its meiotic and somatic functions. Cell Death and Differentiation, 2022, 29, 2347-2361.	5.0	2
10	Mild androgen insensitivity syndrome: the current landscape. Endocrine Practice, 2022, , .	1.1	2
11	The Use of Genetics for Reaching a Diagnosis in XY DSD. Sexual Development, 2022, 16, 207-224.	1.1	5
12	High-throughput Sequencing to Identify Monogenic Etiologies in a Preselected Polycystic Ovary Syndrome Cohort. Journal of the Endocrine Society, 2022, 6, .	0.1	4
13	Genetics, clinical features and outcomes of non-syndromic pituitary gigantism: experience of a single center from Sao Paulo, Brazil. Pituitary, 2021, 24, 252-261.	1.6	4
14	Management of functioning pediatric adrenal tumors. Journal of Pediatric Surgery, 2021, 56, 768-771.	0.8	7
15	Real-World Estimates of Adrenal Insufficiencyâ€“Related Adverse Events in Children With Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e192-e203.	1.8	20
16	Insights from the genetic characterization of central precocious puberty associated with multiple anomalies. Human Reproduction, 2021, 36, 506-518.	0.4	16
17	Vasculometabolic effects in patients with congenital growth hormone deficiency with and without GH replacement therapy during adulthood. Pituitary, 2021, 24, 216-228.	1.6	6
18	Low Protein Expression of both ATRX and ZNRF3 as Novel Negative Prognostic Markers of Adult Adrenocortical Carcinoma. International Journal of Molecular Sciences, 2021, 22, 1238.	1.8	10

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19	SDHB large deletions are associated with absence of MIBG uptake in metastatic lesions of malignant paragangliomas. Endocrine, 2021, 72, 586-590.	1.1	4
20	Phosphodiesterase 2A and 3B variants are associated with primary aldosteronism. Endocrine-Related Cancer, 2021, 28, 1-13.	1.6	17
21	Performance of mutation pathogenicity prediction tools on missense variants associated with 46,XY differences of sex development. Clinics, 2021, 76, e2052.	0.6	10
22	Genetic and clinical aspects of paediatric pheochromocytomas and paragangliomas. Clinical Endocrinology, 2021, 95, 117-124.	1.2	10
23	International practice of corticosteroid replacement therapy in congenital adrenal hyperplasia: data from the I-CAH registry. European Journal of Endocrinology, 2021, 184, 553-563.	1.9	21
24	The Cost-Effectiveness of Congenital Adrenal Hyperplasia Newborn Screening in Brazil: A Comparison Between Screened and Unscreened Cohorts. Frontiers in Pediatrics, 2021, 9, 659492.	0.9	4
25	Low Protein Expression of ATRX and ZNRF3 as a Novel Prognostic Marker of Adult Adrenocortical Carcinoma. Journal of the Endocrine Society, 2021, 5, A87-A88.	0.1	0
26	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
27	Allelic Variants in Established Hypopituitarism Genes Expand Our Knowledge of the Phenotypic Spectrum. Genes, 2021, 12, 1128.	1.0	0
28	The phenotypic spectrum associated with OTX2 mutations in humans. European Journal of Endocrinology, 2021, 185, 121-135.	1.9	15
29	Anthropometric, metabolic, and reproductive outcomes of patients with central precocious puberty treated with leuprorelin acetate 3-month depot (11.25Âmg). Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1371-1377.	0.4	2
30	Adrenal crisis and mortality rate in adrenal insufficiency and congenital adrenal hyperplasia. Archives of Endocrinology and Metabolism, 2021, 65, 488-494.	0.3	7
31	High-throughput splicing assays identify missense and silent splice-disruptive POU1F1 variants underlying pituitary hormone deficiency. American Journal of Human Genetics, 2021, 108, 1526-1539.	2.6	23
32	Genotypeâ€“Phenotype Correlations in Central Precocious Puberty Caused by MKRN3 Mutations. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1041-e1050.	1.8	31
33	Role of the Mevalonate Pathway in Adrenocortical Tumorigenesis. Hormone and Metabolic Research, 2021, 53, 124-131.	0.7	1
34	Plasma Renin Measurements are Unrelated to Mineralocorticoid Replacement Dose in Patients With Primary Adrenal Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 314-326.	1.8	30
35	Outcomes of Patients with Central Precocious Puberty Due to Loss-of-Function Mutations in the MKRN3 Gene after Treatment with Gonadotropin-Releasing Hormone Analog. Neuroendocrinology, 2020, 110, 705-713.	1.2	17
36	Clinical and Genetic Characterization of a Constitutional Delay of Growth and Puberty Cohort. Neuroendocrinology, 2020, 110, 959-966.	1.2	10

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37	SUN-061 Anthropometric and Reproductive Outcomes of Patients with Gonadotropin-Independent Precocious Puberty Due to McCune-Albright Syndrome After Treatment with Distinct Therapeutic Agents. Journal of the Endocrine Society, 2020, 4, .	0.1	0
38	Comprehensive Genetic Analysis of 128 Candidate Genes in a Cohort With Idiopathic, Severe, or Familial Osteoporosis. Journal of the Endocrine Society, 2020, 4, bvaa148.	0.1	11
39	A New Insight into the Surgical Treatment of Primary Macronodular Adrenal Hyperplasia. Journal of the Endocrine Society, 2020, 4, bvaa083.	0.1	14
40	A Bayesian Approach to Diagnose Growth Hormone Deficiency in Children: Insulin-Like Growth Factor Type 1 Is Valuable for Screening and IGF-Binding Protein Type 3 for Confirmation. Hormone Research in Paediatrics, 2020, 93, 197-205.	0.8	4
41	SAT-155 High Prevalence Alterations on DNA Mismatch Repair Genes Related to Lynch Syndrome in Pediatric Patients with Adrenocortical Tumor Carried of the Germline Mutation on TP53. Journal of the Endocrine Society, 2020, 4, .	0.1	0
42	Steroid Screening Tools Differentiating Nonclassical Congenital Adrenal Hyperplasia and Polycystic Ovary Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2895-e2902.	1.8	7
43	Long-term cardio-metabolic outcomes in patients with classical congenital adrenal hyperplasia: is the risk real?. Current Opinion in Endocrinology, Diabetes and Obesity, 2020, 27, 155-161.	1.2	18
44	High Prevalence of Alterations in DNA Mismatch Repair Genes of Lynch Syndrome in Pediatric Patients with Adrenocortical Tumors Carrying a Germline Mutation on TP53. Cancers, 2020, 12, 621.	1.7	4
45	XAF1 as a modifier of p53 function and cancer susceptibility. Science Advances, 2020, 6, eaba3231.	4.7	37
46	Allelic Variants of ARMC5 in Patients With Adrenal Incidentalomas and in Patients With Cushing's Syndrome Associated With Bilateral Adrenal Nodules. Frontiers in Endocrinology, 2020, 11, 36.	1.5	7
47	Adverse Outcomes and Economic Burden of Congenital Adrenal Hyperplasia Late Diagnosis in the Newborn Screening Absence. Journal of the Endocrine Society, 2020, 4, bvz013.	0.1	10
48	Sterol O-Acyl Transferase 1 as a Prognostic Marker of Adrenocortical Carcinoma. Cancers, 2020, 12, 247.	1.7	22
49	Genetics of Primary Ovarian Insufficiency in the Next-Generation Sequencing Era. Journal of the Endocrine Society, 2020, 4, bvz037.	0.1	45
50	Integrative and Analytical Review of the 5-Alpha-Reductase Type 2 Deficiency Worldwide . The Application of Clinical Genetics, 2020, Volume 13, 83-96.	1.4	28
51	SELAdb: A database of exonic variants in a Brazilian population referred to a quaternary medical center in SÃ£o Paulo. Clinics, 2020, 75, e1913.	0.6	15
52	Adrenal Insufficiency and Glucocorticoid Use During the COVID-19 Pandemic. Clinics, 2020, 75, e2022.	0.6	23
53	SUN-709 MiR-200c Expression Profiles in Plasma of 46,XY DSD Patients of Unknown Etiology. Journal of the Endocrine Society, 2020, 4, .	0.1	0
54	SUN-085 Clinical and Hormonal Features of 37 Families with Central Precocious Puberty Due to MKRN3 Loss-Of -Function Mutations. Journal of the Endocrine Society, 2020, 4, .	0.1	0

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55	OR15-04 Central Precocious Puberty without Central Nervous System Lesions: Is It Really Idiopathic?. Journal of the Endocrine Society, 2020, 4, .	0.1	0
56	SAT-560 Usefulness of Contralateral Suppression in Adrenal Venous Sampling to Define Lateralization in Primary Aldosteronism. Journal of the Endocrine Society, 2020, 4, .	0.1	0
57	SUN-178 Clinical and Anatomopathological Characteristics of Two Atypical Aldosterone-Producing Adenomas. Journal of the Endocrine Society, 2020, 4, .	0.1	0
58	Impact of schooling in the HIV/AIDS prevalence among Brazilian transgender women. Archives of Endocrinology and Metabolism, 2020, 64, 369-373.	0.3	3
59	SUN-049 Male Pattern Baldness and Waist-Hip Ratio as Markers of Arterial Stiffness in Transgender Men Undergoing Long-Term Testosterone Therapy. Journal of the Endocrine Society, 2020, 4, .	0.1	0
60	SUN-725 Clinical and Genetic Features of Families with Maternally Inherited Central Precocious Puberty. Journal of the Endocrine Society, 2020, 4, .	0.1	0
61	SUN-081 High Throughput Genetic Analysis Revealed Novel Genomic Loci and Candidate Genes Involved in Central Precocious Puberty Associated with Complex Phenotypes. Journal of the Endocrine Society, 2020, 4, .	0.1	0
62	Screening of targeted panel genes in Brazilian patients with primary ovarian insufficiency. PLoS ONE, 2020, 15, e0240795.	1.1	21
63	Anorexia as the first clinical manifestation of von Hippel-Lindau syndrome. Molecular and Clinical Oncology, 2020, 13, 65.	0.4	0
64	Anorexia as the first clinical manifestation of von Hippelâ€'Lindau syndrome. Molecular and Clinical Oncology, 2020, 13, 1-1.	0.4	0
65	Screening of targeted panel genes in Brazilian patients with primary ovarian insufficiency. , 2020, 15, e0240795.		0
66	Screening of targeted panel genes in Brazilian patients with primary ovarian insufficiency. , 2020, 15, e0240795.		0
67	Screening of targeted panel genes in Brazilian patients with primary ovarian insufficiency. , 2020, 15, e0240795.		0
68	Screening of targeted panel genes in Brazilian patients with primary ovarian insufficiency. , 2020, 15, e0240795.		0
69	Two rare loss-of-function variants in the STAG3 gene leading to primary ovarian insufficiency. European Journal of Medical Genetics, 2019, 62, 186-189.	0.7	30
70	Impact of Long-Term Dexamethasone Therapy on the Metabolic Profile of Patients With 21-Hydroxylase Deficiency. Journal of the Endocrine Society, 2019, 3, 1574-1582.	0.1	7
71	Combined pituitary hormone deficiency caused by PROP1 mutations: update 20 years post-discovery. Archives of Endocrinology and Metabolism, 2019, 63, 167-174.	0.3	23
72	Genetic Evidence of the Association of DEAH-Box Helicase 37 Defects With 46,XY Gonadal Dysgenesis Spectrum. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5923-5934.	1.8	26

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73	Management of 46,XY Differences/Disorders of Sex Development (DSD) Throughout Life. Endocrine Reviews, 2019, 40, 1547-1572.	8.9	68
74	KCNJ5 Somatic Mutation Is a Predictor of Hypertension Remission After Adrenalectomy for Unilateral Primary Aldosteronism. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4695-4702.	1.8	42
75	New Insights Into Pheochromocytoma Surveillance of Young Patients With VHL Missense Mutations. Journal of the Endocrine Society, 2019, 3, 1682-1692.	0.1	15
76	IGF-1 assessed by pubertal status has the best positive predictive power for GH deficiency diagnosis in peripubertal children. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 173-179.	0.4	24
77	Premature Pubarche due to Exogenous Testosterone Gel or Intense Diaper Rash Prevention Cream Use: A Case Series. Hormone Research in Paediatrics, 2019, 91, 411-415.	0.8	10
78	Persistent Poor Metabolic Profile in Postmenopausal Women With Ovarian Hyperandrogenism After Testosterone Level Normalization. Journal of the Endocrine Society, 2019, 3, 1087-1096.	0.1	9
79	Evaluation of SHOX defects in the era of nextâ€generation sequencing. Clinical Genetics, 2019, 96, 261-265.	1.0	9
80	Exome Sequencing Reveals the POLR3H Gene as a Novel Cause of Primary Ovarian Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2827-2841.	1.8	28
81	Androgen Biosynthetic Defects: 17Î²-Hydroxysteroid Dehydrogenadse Type 3 and 5Î±-Reductase Type 2 Deficiencies. , 2019, , 486-491.		0
82	DLK1 Is a Novel Link Between Reproduction and Metabolism. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2112-2120.	1.8	75
83	Psychosexual Aspects, Effects of Prenatal Androgen Exposure, and Gender Change in 46,XY Disorders of Sex Development. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 1160-1170.	1.8	22
84	Clinical spectrum of Li-Fraumeni syndrome/Li-Fraumeni-like syndrome in Brazilian individuals with the TP53 p.R337H mutation. Journal of Steroid Biochemistry and Molecular Biology, 2019, 190, 250-255.	1.2	23
85	Targeted Assessment of G0S2 Methylation Identifies a Rapidly Recurrent, Routinely Fatal Molecular Subtype of Adrenocortical Carcinoma. Clinical Cancer Research, 2019, 25, 3276-3288.	3.2	51
86	Mobile DNA in Endocrinology: LINE-1 Retrotransposon Causing Partial Androgen Insensitivity Syndrome. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 6385-6390.	1.8	10
87	Classic congenital adrenal hyperplasia and its impact on reproduction. Fertility and Sterility, 2019, 111, 7-12.	0.5	39
88	Mutations in MAP3K1 that cause 46,XY disorders of sex development disrupt distinct structural domains in the protein. Human Molecular Genetics, 2019, 28, 1620-1628.	1.4	21
89	A 46,XX testicular disorder of sex development caused by a Wilms' tumour Factorâ€1 (WT1) pathogenic variant. Clinical Genetics, 2019, 95, 172-176.	1.0	24
90	Genetic diagnosis of congenital hypopituitarism by a target gene panel: novel pathogenic variants in GLI2, OTX2 and GHRHR. Endocrine Connections, 2019, 8, 590-595.	0.8	10

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91	ESR1 polymorphism (rs2234693) influences femoral bone mass in patients with Turner syndrome. Endocrine Connections, 2019, 8, 1513-1519.	0.8	12
92	New genetic findings in a large cohort of congenital hypogonadotropic hypogonadism. European Journal of Endocrinology, 2019, 181, 103-119.	1.9	70
93	OR06-6 Whole-Exome Sequencing of Patients with Pituitary Stalk Interruption Syndrome (PSIS) Reveals Probably Pathogenic Variants in Novel Candidate Genes.. Journal of the Endocrine Society, 2019, 3, .	0.1	0
94	OR04-6 Predictors of Clinical Outcome after Adrenalectomy for Unilateral Primary Aldosteronism. Journal of the Endocrine Society, 2019, 3, .	0.1	0
95	SUN-359 Preserved Bone Mineral Density In Adults With Classical Forms Of Congenital Adrenal Hyperplasia Submitted To Long-term Low Glucocorticoid Doses. Journal of the Endocrine Society, 2019, 3, .	0.1	0
96	SAT-064 Validation of Furosemide Upright Test in Primary Aldosteronism Diagnosis Using Direct Renin Assay. Journal of the Endocrine Society, 2019, 3, .	0.1	0
97	MON-251 Clinical Features of a Large Cohort of Patients with Familial Central Precocious Puberty Caused by Loss-of-Function Mutations in MKRN3. Journal of the Endocrine Society, 2019, 3, .	0.1	0
98	Longâ€term outcomes and molecular analysis of a large cohort of patients with 46,<scp>XY</scp> disorder of sex development due to partial gonadal dysgenesis. Clinical Endocrinology, 2018, 89, 164-177.	1.2	13
99	Partial androgen insensitivity syndrome due to somatic mosaicism of the androgen receptor. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 223-228.	0.4	9
100	Pathogenic copy number variants in patients with congenital hypopituitarism associated with complex phenotypes. Clinical Endocrinology, 2018, 88, 425-431.	1.2	11
101	Spontaneous fertility in a male patient with testotoxicosis despite suppression of FSH levels. Human Reproduction, 2018, 33, 914-918.	0.4	3
102	Long-Term Outcomes of Patients with Central Precocious Puberty due to Hypothalamic Hamartoma after GnRHa Treatment: Anthropometric, Metabolic, and Reproductive Aspects. Neuroendocrinology, 2018, 106, 203-210.	1.2	21
103	The role of ARMC5 in human cell cultures from nodules of primary macronodular adrenocortical hyperplasia (PMAH). Molecular and Cellular Endocrinology, 2018, 460, 36-46.	1.6	38
104	Biallelic and monoallelic ESR2 variants associated with 46,XY disorders of sex development. Genetics in Medicine, 2018, 20, 717-727.	1.1	28
105	Identification of the first homozygous 1â€bp deletion in GDF9 gene leading to primary ovarian insufficiency by using targeted massively parallel sequencing. Clinical Genetics, 2018, 93, 408-411.	1.0	29
106	A severe phenotype of Kennedy disease associated with a very large CAG repeat expansion. Muscle and Nerve, 2018, 57, E95-E97.	1.0	11
107	Methylome profiling of healthy and central precocious puberty girls. Clinical Epigenetics, 2018, 10, 146.	1.8	34
108	Primary malignant tumors of the adrenal glands. Clinics, 2018, 73, e756s.	0.6	27

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109	Androgen receptor mRNA analysis from whole blood: a lowâ€cost strategy for detection of androgen receptor gene splicing defects. Clinical Genetics, 2018, 94, 489-490.	1.0	2
110	Central Precocious Puberty Caused by a Heterozygous Deletion in the MKRN3 Promoter Region. Neuroendocrinology, 2018, 107, 127-132.	1.2	23
111	Androgen insensitivity syndrome: a review. Archives of Endocrinology and Metabolism, 2018, 62, 227-235.	0.3	100
112	Applicability of a novel mathematical model for the prediction of adult height and age at menarche in girls with idiopathic central precocious puberty. Clinics, 2018, 73, e480.	0.6	4
113	Low estrogen doses normalize testosterone and estradiol levels to the female range in transgender women. Clinics, 2018, 73, e86.	0.6	15
114	Assembling the jigsaw puzzle: <scp>CBX</scp>2 isoform 2 and its targets in disorders/differences of sex development. Molecular Genetics & Genomic Medicine, 2018, 6, 785-795.	0.6	16
115	An update of genetic basis of PCOS pathogenesis. Archives of Endocrinology and Metabolism, 2018, 62, 352-361.	0.3	88
116	Testosterone replacement in androgen insensitivity: is there an advantage?. Annals of Translational Medicine, 2018, 6, S85-S85.	0.7	1
117	High Frequency of <i>MKRN3</i> Mutations in Male Central Precocious Puberty Previously Classified as Idiopathic. Neuroendocrinology, 2017, 105, 17-25.	1.2	65
118	46,XY disorder of sex development (DSD) due to 17Î²-hydroxysteroid dehydrogenase type 3 deficiency. Journal of Steroid Biochemistry and Molecular Biology, 2017, 165, 79-85.	1.2	66
119	Persistent MÃ¼llerian duct syndrome due to a mutation in the anti-MÃ¼llerian hormone receptor gene (AMHR2). Anales De PediatrÃa (English Edition), 2017, 86, 94-95.	0.1	0
120	Clinical, genetic, and structural basis of congenital adrenal hyperplasia due to 11Î²-hydroxylase deficiency. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E1933-E1940.	3.3	106
121	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
122	Triple A Syndrome: Preliminary Response to the Antioxidant N-Acetylcysteine Treatment in a Child. Hormone Research in Paediatrics, 2017, 88, 167-171.	0.8	4
123	Discriminating between virilizing ovary tumors and ovary hyperthecosis in postmenopausal women: clinical data, hormonal profiles and image studies. European Journal of Endocrinology, 2017, 177, 93-102.	1.9	37
124	Paternally Inherited DLK1 Deletion Associated With Familial Central Precocious Puberty. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1557-1567.	1.8	145
125	Heterozygous Nonsense Mutation in the Androgen Receptor Gene Associated with Partial Androgen Insensitivity Syndrome in an Individual with 47,XXY Karyotype. Sexual Development, 2017, 11, 78-81.	1.1	8
126	Assessment of stress levels in girls with central precocious puberty before and during long-acting gonadotropin-releasing hormone agonist treatment: a pilot study. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 657-662.	0.4	14

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127	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
128	Physiology of Male Gonadotropic Axis and Disorders of Sex Development. , 2017, , 75-96.		0
129	Neonatal 17â€hydroxyprogesterone levels adjusted according to age at sample collection and birthweight improve the efficacy of congenital adrenal hyperplasia newborn screening. Clinical Endocrinology, 2017, 86, 480-487.	1.2	34
130	A novel homozygous 1-bp deletion in the NOBOX gene in two Brazilian sisters with primary ovarian failure. Endocrine, 2017, 58, 442-447.	1.1	17
131	Global Application of the Assessment of Communication Skills of Paediatric Endocrinology Fellows in the Management of Differences in Sex Development Using the ESPE E-Learning.Org Portal. Hormone Research in Paediatrics, 2017, 88, 127-139.	0.8	13
132	MP37-08 A NEW INSIGHT FOR THE TREATMENT OF PRIMARY MACRONODULAR ADRENAL HYPERPLASIA: ADRENAL SPARING SURGERY EARLY OUTCOMES. Journal of Urology, 2017, 197, .	0.2	0
133	A recurrent synonymous mutation in the human androgen receptor gene causing complete androgen insensitivity syndrome. Journal of Steroid Biochemistry and Molecular Biology, 2017, 174, 14-16.	1.2	16
134	Molecular analysis of brazilian patients with combined pituitary hormone deficiency and orthotopic posterior pituitary lobe reveals eight different <scp>PROP</scp>1 alterations with three novel mutations. Clinical Endocrinology, 2017, 87, 725-732.	1.2	13
135	Nonâ€coding variation in disorders of sex development. Clinical Genetics, 2017, 91, 163-172.	1.0	39
136	Differential Expression of Stem Cell Markers in Human Adamantinomatous Craniopharyngioma and Pituitary Adenoma. Neuroendocrinology, 2017, 104, 183-193.	1.2	19
137	Reprint of â€œSteroid 5Î±-reductase 2 deficiencyâ€. Journal of Steroid Biochemistry and Molecular Biology, 2017, 165, 95-100.	1.2	9
138	Malignant testicular germ cell tumors in postpubertal individuals with androgen insensitivity: prevalence, pathology and relevance of single nucleotide polymorphism-based susceptibility profiling. Human Reproduction, 2017, 32, 2561-2573.	0.4	50
139	Successful Pregnancies After Adequate Hormonal Replacement in Patients With Combined Pituitary Hormone Deficiencies. Journal of the Endocrine Society, 2017, 1, 1322-1330.	0.1	14
140	An activating mutation in the CRHR1 gene is rarely associated with pituitary-dependent hyperadrenocorticism in poodles. Clinics, 2017, 72, 575-581.	0.6	1
141	Growth hormone deficiency with advanced bone age: phenotypic interaction between GHRH receptor and CYP21A2 mutations diagnosed by sanger and whole exome sequencing. Archives of Endocrinology and Metabolism, 2017, 61, 633-636.	0.3	4
142	Adjuvant radiotherapy for the primary treatment of adrenocortical carcinoma: are we offering the best?. International Braz J Urol: Official Journal of the Brazilian Society of Urology, 2017, 43, 841-848.	0.7	17
143	Long-term follow-up of a female with congenital adrenal hyperplasia due to P450-oxidoreductase deficiency. Archives of Endocrinology and Metabolism, 2016, 60, 500-504.	0.3	15
144	Long-term Results after CT-Guided Percutaneous Ethanol Ablation for the Treatment of Hyperfunctioning Adrenal Disorders. Clinics, 2016, 71, 600-605.	0.6	6

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145	Surgical Treatment after Failed Primary Correction of Urogenital Sinus in Female Patients with Virilizing Congenital Adrenal Hyperplasia: Are Good Results Possible?. Frontiers in Pediatrics, 2016, 4, 118.	0.9	10
146	Steroid 5Î±-reductase 2 deficiency. Journal of Steroid Biochemistry and Molecular Biology, 2016, 163, 206-211.	1.2	123
147	<scp>HESX</scp>1 mutations in patients with congenital hypopituitarism: variable phenotypes with the same genotype. Clinical Endocrinology, 2016, 85, 408-414.	1.2	24
148	Negative correlation between tumour size and cortisol/ACTH ratios in patients with Cushingâ€™s disease harbouring microadenomas or macroadenomas. Journal of Endocrinological Investigation, 2016, 39, 1401-1409.	1.8	16
149	MP12-14 A META-ANALYSIS OF THE ROLE OF ADJUVANT RADIOTHERAPY AFTER SURGERY FOR ADRENOCORTICAL CARCINOMA. Journal of Urology, 2016, 195, .	0.2	0
150	Molecular CYP21A2 diagnosis in 480 Brazilian patients with congenital adrenal hyperplasia before newborn screening introduction. European Journal of Endocrinology, 2016, 175, 107-116.	1.9	60
151	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	7.7	482
152	A homozygous point mutation in the GH1 promoter (c.-223C>T) leads to reduced GH1 expression in siblings with isolated GH deficiency (IGHD). European Journal of Endocrinology, 2016, 175, K7-K15.	1.9	5
153	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
154	Effects of Type 1 Insulin-Like Growth Factor Receptor Silencing in a Human Adrenocortical Cell Line. Hormone and Metabolic Research, 2016, 48, 484-488.	0.7	3
155	The Use of Three-dimensional Printers for Partial Adrenalectomy: Estimating the Resection Limits. Urology, 2016, 90, 217-221.	0.5	31
156	Successful Live Birth in a Woman With 17Î±-Hydroxylase Deficiency Through IVF Frozen-Thawed Embryo Transfer. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 345-348.	1.8	45
157	High 18F-FDG uptake in PMAH correlated with normal expression of Glut1, HK1, HK2, and HK3. Acta Radiologica, 2016, 57, 370-377.	0.5	9
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