Rafael Loch Batista

List of Publications by Year
in descending order

42 papers	593 citations	933447 10 h-index	642732 23 g-index
43 all docs	43 docs citations	43 times ranked	768 citing authors

#	Article	IF	Citations
1	Steroid 5Î±-reductase 2 deficiency. Journal of Steroid Biochemistry and Molecular Biology, 2016, 163, 206-211.	2.5	123
2	Androgen insensitivity syndrome: a review. Archives of Endocrinology and Metabolism, 2018, 62, 227-235.	0.6	100
3	Management of 46,XY Differences/Disorders of Sex Development (DSD) Throughout Life. Endocrine Reviews, 2019, 40, 1547-1572.	20.1	68
4	Cabergoline in the Management of Residual Nonfunctioning Pituitary Adenoma. American Journal of Clinical Oncology: Cancer Clinical Trials, 2019, 42, 221-227.	1.3	41
5	Integrative and Analytical Review of the 5-Alpha-Reductase Type 2 Deficiency Worldwide . The Application of Clinical Genetics, 2020, Volume 13, 83-96.	3.0	28
6	A 46,XX testicular disorder of sex development caused by a Wilms' tumour Factorâ€1 (WT1) pathogenic variant. Clinical Genetics, 2019, 95, 172-176.	2.0	24
7	Nonfunctioning Pituitary Adenoma Recurrence and Its Relationship with Sex, Size, and Hormonal Immunohistochemical Profile. World Neurosurgery, 2018, 120, e241-e246.	1.3	22
8	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.	3.6	22
9	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.	2.5	16
10	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.	2.4	13
11	A severe phenotype of Kennedy disease associated with a very large CAG repeat expansion. Muscle and Nerve, 2018, 57, E95-E97.	2.2	11
12	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.	3.6	11
13	Two cases of Kallmann syndrome associated with empty sella. Pituitary, 2008, 11, 109-112.	2.9	10
14	Severe Psychotic Disorder as the Main Manifestation of Adrenal Insufficiency. Case Reports in Psychiatry, 2015, 2015, 1-4.	0.5	10
15	Mobile DNA in Endocrinology: LINE-1 Retrotransposon Causing Partial Androgen Insensitivity Syndrome. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 6385-6390.	3.6	10
16	Reprint of â€œSteroid 5Î±-reductase 2 deficiencyâ€. Journal of Steroid Biochemistry and Molecular Biology, 2017, 165, 95-100.	2.5	9
17	Partial androgen insensitivity syndrome due to somatic mosaicism of the androgen receptor. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 223-228.	0.9	9
18	Hyperprolactinemia and immunohistochemical expression of intracellular prolactin and prolactin receptor in primary central nervous system tumors and their relationship with cellular replication. Brain Tumor Pathology, 2007, 24, 41-46.	1.7	8

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19	Thickened Pituitary Stalk Associated with a Mass in the Sphenoidal Sinus: An Alarm to Suspect Hypophysitis by Immunoglobulin G4?. International Archives of Otorhinolaryngology, 2015, 19, 273-276.	0.8	8
20	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.	2.0	8
21	Association between KISS1 rs5780218 promoter polymorphism and onset of growth hormone secreting pituitary adenoma. Annales D'Endocrinologie, 2019, 80, 96-100.	1.4	6
22	Vasculometabolic effects in patients with congenital growth hormone deficiency with and without GH replacement therapy during adulthood. Pituitary, 2021, 24, 216-228.	2.9	6
23	False positive results using calcitonin as a screening method for medullary thyroid carcinoma. Indian Journal of Endocrinology and Metabolism, 2013, 17, 524.	0.4	5
24	Progression of an Invasive ACTH Pituitary Macroadenoma with Cushingâ€™s Disease to Pituitary Carcinoma. Case Reports in Oncological Medicine, 2015, 2015, 1-4.	0.3	5
25	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.	2.0	5
26	The Use of Genetics for Reaching a Diagnosis in XY DSD. Sexual Development, 2022, 16, 207-224.	2.0	5
27	Impact of schooling in the HIV/AIDS prevalence among Brazilian transgender women. Archives of Endocrinology and Metabolism, 2020, 64, 369-373.	0.6	3
28	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.	2.0	2
29	Mild androgen insensitivity syndrome: the current landscape. Endocrine Practice, 2022, , .	2.1	2
30	Central adrenal insufficiency: who, when, and how? From the evidence to the controversies â€“ an exploratory review. Archives of Endocrinology and Metabolism, 2022, , .	0.6	2
31	Testosterone replacement in androgen insensitivity: is there an advantage?. Annals of Translational Medicine, 2018, 6, S85-S85.	1.7	1
32	AlteraÃ§Ãµes neuroendÃ³crinas em pacientes com traumatismo cranioencefÃ¡lico. Brazilian Neurosurgery, 2013, 32, 74-79.	0.1	0
33	Androgen Biosynthetic Defects: 17Î²-Hydroxysteroid Dehydrogenadse Type 3 and 5Î±-Reductase Type 2 Deficiencies. , 2019, , 486-491.		0
34	Complete Androgen Insensitivity in Girls with Inguinal Hernias: A Serendipity Opportunity for Early Diagnosis. Journal of Investigative Surgery, 2021, 34, 234-235.	1.3	0
35	Possibilidade de associaÃ§Ã£o de melanoma e acromegalia. Anais Brasileiros De Dermatologia, 2008, 83, 369-371.	1.1	0
36	Clinically nonfunctioning pituitary adenoma growth after radiosurgery. Arquivos De Neuro-Psiquiatria, 2012, 70, 643-644.	0.8	0

#	Article	IF	Citations
37	SUN-709 MiR-200c Expression Profiles in Plasma of 46,XY DSD Patients of Unknown Etiology. Journal of the Endocrine Society, 2020, 4, .	0.2	0
38	SUN-095 Understanding and Communication Around DSD According to the Mothers and Patientsâ€™ Perspectives. Journal of the Endocrine Society, 2020, 4, .	0.2	0
39	OR15-06 Integrative and Analytical Review of the 5 Alpha Reductase Type 2 Deficiency Worldwide. Journal of the Endocrine Society, 2020, 4, .	0.2	0
40	SUN-071 Prenatal and Post-Natal Influence of Androgens in the Psychosexual Development in Individuals with 21-hydroxylase Congenital Adrenal Hyperplasia. Journal of the Endocrine Society, 2020, 4, .	0.2	0
41	SUN-078 Clinical, Hormonal, Psychosexual Aspects, Gonadal Tumors and Genetic Background of an Androgen Insensitivity Syndrome Cohort. Journal of the Endocrine Society, 2020, 4, .	0.2	0
42	Anorexia as the first clinical manifestation of von Hippel-Lindau syndrome. Molecular and Clinical Oncology, 2020, 13, 65.	1.0	0