

# Rafael Loch Batista

## List of Publications by Year in descending order

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Version: 2024-02-01

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	WT1 Pathogenic Variants are Associated with a Broad Spectrum of Differences in Sex Development Phenotypes and Heterogeneous Progression of Renal Disease. <i>Sexual Development</i> , 2022, 16, 46-54.	2.0	5
2	Contribution of Clinical and Genetic Approaches for Diagnosing 209 Index Cases With 46,XY Differences of Sex Development. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e1797-e1806.	3.6	11
3	Mild androgen insensitivity syndrome: the current landscape. <i>Endocrine Practice</i> , 2022, , .	2.1	2
4	The Use of Genetics for Reaching a Diagnosis in XY DSD. <i>Sexual Development</i> , 2022, 16, 207-224.	2.0	5
5	Central adrenal insufficiency: who, when, and how? From the evidence to the controversies – an exploratory review. <i>Archives of Endocrinology and Metabolism</i> , 2022, , .	0.6	2
6	Vasculometabolic effects in patients with congenital growth hormone deficiency with and without GH replacement therapy during adulthood. <i>Pituitary</i> , 2021, 24, 216-228.	2.9	6
7	Complete Androgen Insensitivity in Girls with Inguinal Hernias: A Serendipity Opportunity for Early Diagnosis. <i>Journal of Investigative Surgery</i> , 2021, 34, 234-235.	1.3	0
8	&lt;p&gt;Integrative and Analytical Review of the 5-Alpha-Reductase Type 2 Deficiency Worldwide&lt;/p&gt;. <i>The Application of Clinical Genetics</i> , 2020, Volume 13, 83-96.	3.0	28
9	SUN-709 MiR-200c Expression Profiles in Plasma of 46,XY DSD Patients of Unknown Etiology. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
10	SUN-095 Understanding and Communication Around DSD According to the Mothers and Patients’s™ Perspectives. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
11	Impact of schooling in the HIV/AIDS prevalence among Brazilian transgender women. <i>Archives of Endocrinology and Metabolism</i> , 2020, 64, 369-373.	0.6	3
12	OR15-06 Integrative and Analytical Review of the 5 Alpha Reductase Type 2 Deficiency Worldwide. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
13	SUN-071 Prenatal and Post-Natal Influence of Androgens in the Psychosexual Development in Individuals with 21-hydroxylase Congenital Adrenal Hyperplasia. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
14	SUN-078 Clinical, Hormonal, Psychosexual Aspects, Gonadal Tumors and Genetic Background of an Androgen Insensitivity Syndrome Cohort. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
15	Anorexia as the first clinical manifestation of von Hippel-Lindau syndrome. <i>Molecular and Clinical Oncology</i> , 2020, 13, 65.	1.0	0
16	Management of 46,XY Differences/Disorders of Sex Development (DSD) Throughout Life. <i>Endocrine Reviews</i> , 2019, 40, 1547-1572.	20.1	68
17	Cabergoline in the Management of Residual Nonfunctioning Pituitary Adenoma. <i>American Journal of Clinical Oncology: Cancer Clinical Trials</i> , 2019, 42, 221-227.	1.3	41
18	Androgen Biosynthetic Defects: 17 <sup>β</sup> -Hydroxysteroid Dehydrogenase Type 3 and 5 $\alpha$ -Reductase Type 2 Deficiencies. , 2019, , 486-491.		0

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19	Psychosexual Aspects, Effects of Prenatal Androgen Exposure, and Gender Change in 46,XY Disorders of Sex Development. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 1160-1170.	3.6	22
20	Mobile DNA in Endocrinology: LINE-1 Retrotransposon Causing Partial Androgen Insensitivity Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 6385-6390.	3.6	10
21	Association between KISS1 rs5780218 promoter polymorphism and onset of growth hormone secreting pituitary adenoma. <i>Annales D'Endocrinologie</i> , 2019, 80, 96-100.	1.4	6
22	A 46,XX testicular disorder of sex development caused by a Wilms' tumour Factor (WT1) pathogenic variant. <i>Clinical Genetics</i> , 2019, 95, 172-176.	2.0	24
23	Long-term outcomes and molecular analysis of a large cohort of patients with 46,XY disorder of sex development due to partial gonadal dysgenesis. <i>Clinical Endocrinology</i> , 2018, 89, 164-177.	2.4	13
24	Partial androgen insensitivity syndrome due to somatic mosaicism of the androgen receptor. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2018, 31, 223-228.	0.9	9
25	A severe phenotype of Kennedy disease associated with a very large CAG repeat expansion. <i>Muscle and Nerve</i> , 2018, 57, E95-E97.	2.2	11
26	Nonfunctioning Pituitary Adenoma Recurrence and Its Relationship with Sex, Size, and Hormonal Immunohistochemical Profile. <i>World Neurosurgery</i> , 2018, 120, e241-e246.	1.3	22
27	Androgen receptor mRNA analysis from whole blood: a low-cost strategy for detection of androgen receptor gene splicing defects. <i>Clinical Genetics</i> , 2018, 94, 489-490.	2.0	2
28	Androgen insensitivity syndrome: a review. <i>Archives of Endocrinology and Metabolism</i> , 2018, 62, 227-235.	0.6	100
29	Testosterone replacement in androgen insensitivity: is there an advantage?. <i>Annals of Translational Medicine</i> , 2018, 6, S85-S85.	1.7	1
30	Heterozygous Nonsense Mutation in the Androgen Receptor Gene Associated with Partial Androgen Insensitivity Syndrome in an Individual with 47,XXY Karyotype. <i>Sexual Development</i> , 2017, 11, 78-81.	2.0	8
31	A recurrent synonymous mutation in the human androgen receptor gene causing complete androgen insensitivity syndrome. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 174, 14-16.	2.5	16
32	Reprint of "Steroid 5 $\alpha$ -reductase 2 deficiency". <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 165, 95-100.	2.5	9
33	Steroid 5 $\alpha$ -reductase 2 deficiency. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2016, 163, 206-211.	2.5	123
34	Progression of an Invasive ACTH Pituitary Macroadenoma with Cushing's Disease to Pituitary Carcinoma. <i>Case Reports in Oncological Medicine</i> , 2015, 2015, 1-4.	0.3	5
35	Severe Psychotic Disorder as the Main Manifestation of Adrenal Insufficiency. <i>Case Reports in Psychiatry</i> , 2015, 2015, 1-4.	0.5	10
36	Thickened Pituitary Stalk Associated with a Mass in the Sphenoidal Sinus: An Alarm to Suspect Hypophysitis by Immunoglobulin G4?. <i>International Archives of Otorhinolaryngology</i> , 2015, 19, 273-276.	0.8	8

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37	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
38	Alterações neuroendócrinas em pacientes com traumatismo cranioencefálico. Brazilian Neurosurgery, 2013, 32, 74-79.	0.1	0
39	Clinically nonfunctioning pituitary adenoma growth after radiosurgery. Arquivos De Neuro-Psiquiatria, 2012, 70, 643-644.	0.8	0
40	Two cases of Kallmann syndrome associated with empty sella. Pituitary, 2008, 11, 109-112.	2.9	10
41	Possibilidade de associação de melanoma e acromegalia. Anais Brasileiros De Dermatologia, 2008, 83, 369-371.	1.1	0
42	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