Bruno Ferraz-de-Souza

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9509479/publications.pdf

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39	1,430	430442	433756
papers	citations	18 h-index	31 g-index

39	39	39	1896
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetic Analysis of Pediatric Primary Adrenal Insufficiency of Unknown Etiology: 25 Yearsâ€™ Experience in the UK. Journal of the Endocrine Society, 2021, 5, bvab086.	0.1	34
2	Real-world impact of glucocorticoid replacement therapy on bone mineral density: retrospective experience of a large single-center CAH cohort spanning 24Âyears. Osteoporosis International, 2020, 31, 905-912.	1.3	7
3	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
4	The molecular landscape of osteogenesis imperfecta in a Brazilian tertiary service cohort. Osteoporosis International, 2020, 31, 1341-1352.	1.3	5
5	Prevention and treatment of oral adverse effects of antiresorptive medications for osteoporosis â€“ A position paper of the Brazilian Society of Endocrinology and Metabolism (SBEM), Brazilian Society of Stomatology and Oral Pathology (Sobep), and Brazilian Association for Bone Evaluation and Osteometabolism (Abrasso). Archives of Endocrinology and Metabolism, 2020, . .	0.3	2
6	Acute and long-term kidney function after parathyroidectomy for primary hyperparathyroidism. PLoS ONE, 2020, 15, e0244162.	1.1	8
7	Acute and long-term kidney function after parathyroidectomy for primary hyperparathyroidism. , 2020, 15, e0244162.		0
8	Acute and long-term kidney function after parathyroidectomy for primary hyperparathyroidism. , 2020, 15, e0244162.		0
9	Acute and long-term kidney function after parathyroidectomy for primary hyperparathyroidism. , 2020, 15, e0244162.		0
10	Acute and long-term kidney function after parathyroidectomy for primary hyperparathyroidism. , 2020, 15, e0244162.		0
11	Transcriptomic Response to 1,25-Dihydroxyvitamin D in Human Fibroblasts with or without a Functional Vitamin D Receptor (VDR): Novel Target Genes and Insights into VDR Basal Transcriptional Activity. Cells, 2019, 8, 318.	1.8	9
12	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
13	Diagnosis and treatment of hypoparathyroidism: a position statement from the Brazilian Society of Endocrinology and Metabolism. Archives of Endocrinology and Metabolism, 2018, 62, 106-124.	0.3	18
14	The elusive clinical significance of osteocalcin actions in energy metabolism in humans. Archives of Endocrinology and Metabolism, 2018, 62, 271-272.	0.3	0
15	Genetics of osteoporosis: searching for candidate genes for bone fragility. Archives of Endocrinology and Metabolism, 2016, 60, 391-401.	0.3	37
16	25-Hydroxyvitamin D and TSH as Risk Factors or Prognostic Markers in Thyroid Carcinoma. PLoS ONE, 2016, 11, e0164550.	1.1	26
17	Nonspecific binding of a frequently used vitamin D receptor (VDR) antibody: important implications for vitamin D research in human health. Endocrine, 2016, 54, 556-559.	1.1	3
18	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

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19	Wide spectrum of NR5A1â€related phenotypes in 46,XY and 46,XX individuals. Birth Defects Research Part C: Embryo Today Reviews, 2016, 108, 309-320.	3.6	76
20	POD-1/TCF21Reduces SHP Expression, AffectingLRH-1Regulation and Cell Cycle Balance in Adrenocortical and Hepatocarcinoma Tumor Cells. BioMed Research International, 2015, 2015, 1-9.	0.9	10
21	Normal bone mass and normocalcemia in adulthood despite homozygous vitamin D receptor mutations. Osteoporosis International, 2015, 26, 1819-1823.	1.3	12
22	The role of enteric hormone GLP-2 in the response of bone markers to a mixed meal in postmenopausal women with type 2 diabetes mellitus. Diabetology and Metabolic Syndrome, 2015, 7, 13.	1.2	20
23	The evolution of primary hyperparathyroidism. Archives of Endocrinology and Metabolism, 2015, 59, 381-382.	0.3	5
24	Homozygous Inactivating Mutation inNANOS3in Two Sisters with Primary Ovarian Insufficiency. BioMed Research International, 2014, 2014, 1-8.	0.9	36
25	Symptomatic intracranial hypertension and prolonged hypocalcemia following treatment of Pagetâ€™s disease of the skull with zoledronic acid. Journal of Bone and Mineral Metabolism, 2013, 31, 360-365.	1.3	9
26	POD-1 binding to the E-box sequence inhibits SF-1 and StAR expression in human adrenocortical tumor cells. Molecular and Cellular Endocrinology, 2013, 371, 140-147.	1.6	28
27	Diagnosis and treatment of Paget's disease of bone: a mini-review. Arquivos Brasileiros De Endocrinologia E Metabologia, 2013, 57, 577-582.	1.3	7
28	Potential Effects of Alendronate on Fibroblast Growth Factor 23 Levels and Effective Control of Hypercalciuria in an Adult with Jansen's Metaphyseal Chondrodysplasia. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 1098-1103.	1.8	19
29	Mutations in the PCNA-binding domain of CDKN1C cause IMAGe syndrome. Nature Genetics, 2012, 44, 788-792.	9.4	169
30	Role of DAX-1 <i>(NR0B1)</i> and Steroidogenic Factor-1 <i>(NR5A1)</i> in Human Adrenal Function. Endocrine Development, 2011, 20, 38-46.	1.3	40
31	Steroidogenic factor-1 (SF-1, NR5A1) and human disease. Molecular and Cellular Endocrinology, 2011, 336, 198-205.	1.6	143
32	ChIPâ€onâ€chip analysis reveals angiopoietin 2 (Ang2, ANGPT2) as a novel target of steroidogenic factorâ€1 (SFâ€1, NR5A1) in the human adrenal gland. FASEB Journal, 2011, 25, 1166-1175.	0.2	27
33	Sterol O-Acyltransferase 1 (SOAT1, ACAT) Is a Novel Target of Steroidogenic Factor-1 (SF-1, NR5A1,) Tj ETQq1 1 0.784314 rgBT /Over	1.8	45
34	Human Male Infertility Associated with Mutations in NR5A1 Encoding Steroidogenic Factor 1. American Journal of Human Genetics, 2010, 87, 505-512.	2.6	210
35	CBP/p300-Interacting Transactivator, with Glu/Asp-Rich C-Terminal Domain, 2, and Pre-B-Cell Leukemia Transcription Factor 1 in Human Adrenal Development and Disease. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 678-683.	1.8	35
36	Five novel mutations in steroidogenic factor 1 (SF1,NR5A1) in 46,XY patients with severe underandrogenization but without adrenal insufficiency. Human Mutation, 2008, 29, 59-64.	1.1	141

#	Article	IF	Citations
37	Disorders of Adrenal Development. , 2008, 13, 19-32.		41
38	Heterozygous Missense Mutations in Steroidogenic Factor 1 (SF1/Ad4BP, NR5A1) Are Associated with 46,XY Disorders of Sex Development with Normal Adrenal Function. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 991-999.	1.8	189
39	Genetic Disorders Involving Adrenal Development. , 2007, 11, 36-46.		5