

Fernanda Borchers Coeli-Lacchini

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

170
citations

1478505

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1199594

12
g-index

19
all docs

19
docs citations

19
times ranked

277
citing authors

#	ARTICLE	IF	CITATIONS
1	How do sphingolipids play a role in epigenetic mechanisms and gene expression?. <i>Epigenomics</i> , 2022, 14, 219-222.	2.1	4
2	Vitamin D receptor hypermethylation as a biomarker for pediatric adrenocortical tumors. <i>European Journal of Endocrinology</i> , 2022, 186, 573-585.	3.7	4
3	Telomere length and Wnt/ β -catenin pathway in adamantinomatous craniopharyngiomas. <i>European Journal of Endocrinology</i> , 2022, 187, 219-230.	3.7	3
4	Tumor DNA Methylation Profiling Is a Robust and Independent Prognostic Marker for Pediatric Patients With Adrenocortical Tumors. <i>Journal of the Endocrine Society</i> , 2021, 5, A99-A100.	0.2	0
5	Effects of arginase genetic polymorphisms on nitric oxide formation in healthy pregnancy and in preeclampsia. <i>Nitric Oxide - Biology and Chemistry</i> , 2021, 109-110, 20-25.	2.7	6
6	Arginase II polymorphisms modify the hypotensive responses to propofol by affecting nitric oxide bioavailability. <i>European Journal of Clinical Pharmacology</i> , 2021, 77, 869-877.	1.9	6
7	Association of 11 β -hydroxysteroid dehydrogenase type1 (HSD11b1) gene polymorphisms with outcome of antidepressant therapy and suicide attempts. <i>Behavioural Brain Research</i> , 2020, 381, 112343.	2.2	5
8	Clinical, Molecular, Functional, and Structural Characterization of CYP17A1 Mutations in Brazilian Patients with 17-Hydroxylase Deficiency. <i>Hormone and Metabolic Research</i> , 2020, 52, 186-193.	1.5	5
9	Interaction Between NOS3 and HMOX1 on Antihypertensive Drug Responsiveness in Preeclampsia. <i>Revista Brasileira De Ginecologia E Obstetricia</i> , 2020, 42, 460-467.	0.8	5
10	Sphingolipids and Mitochondrial Dynamic. <i>Cells</i> , 2020, 9, 581.	4.1	46
11	USP8 Mutations and Cell Cycle Regulation in Corticotroph Adenomas. <i>Hormone and Metabolic Research</i> , 2020, 52, 117-123.	1.5	13
12	DDAH1 and DDAH2 polymorphisms associate with asymmetrical dimethylarginine plasma levels in erectile dysfunction patients but not in healthy controls. <i>Nitric Oxide - Biology and Chemistry</i> , 2019, 92, 11-17.	2.7	5
13	Circulating HO-1 levels are not associated with plasma sFLT-1 and GT _n <i>HMOX1</i> polymorphism in preeclampsia. <i>Hypertension in Pregnancy</i> , 2019, 38, 73-77.	1.1	3
14	Relationship between asymmetric dimethylarginine, nitrite and genetic polymorphisms: Impact on erectile dysfunction therapy. <i>Nitric Oxide - Biology and Chemistry</i> , 2017, 71, 44-51.	2.7	7
15	Clinical, hormonal, ovarian, and genetic aspects of 46,XX patients with congenital adrenal hyperplasia due to CYP17A1 defects. <i>Fertility and Sterility</i> , 2016, 105, 1612-1619.	1.0	27
16	Sonic Hedgehog mutations are not a common cause of congenital hypopituitarism in the absence of complex midline cerebral defects. <i>Clinical Endocrinology</i> , 2015, 82, 562-569.	2.4	12
17	Comments to a rational, non-radioactive strategy for the molecular diagnosis of congenital adrenal hyperplasia due to 21-hydroxylase deficiency. <i>Gene</i> , 2014, 534, 451-452.	2.2	0
18	A rational, non-radioactive strategy for the molecular diagnosis of congenital adrenal hyperplasia due to 21-hydroxylase deficiency. <i>Gene</i> , 2013, 526, 239-245.	2.2	15