Fernanda Borchers Coeli-Lacchini

List of Publications by Year
in descending order

18 papers	170 citations	1478505 6 h-index	1199594 12 g-index
19 all docs	19 docs citations	19 times ranked	277 citing authors

#	Article	IF	Citations
1	Sphingolipids and Mitochondrial Dynamic. Cells, 2020, 9, 581.	4.1	46
2	Clinical, hormonal, ovarian, and genetic aspects of 46,XX patients with congenital adrenal hyperplasia due to CYP17A1 defects. Fertility and Sterility, 2016, 105, 1612-1619.	1.0	27
3	A rational, non-radioactive strategy for the molecular diagnosis of congenital adrenal hyperplasia due to 21-hydroxylase deficiency. Gene, 2013, 526, 239-245.	2.2	15
4	USP8 Mutations and Cell Cycle Regulation in Corticotroph Adenomas. Hormone and Metabolic Research, 2020, 52, 117-123.	1.5	13
5	Sonic Hedgehog mutations are not a common cause of congenital hypopituitarism in the absence of complex midline cerebral defects. Clinical Endocrinology, 2015, 82, 562-569.	2.4	12
6	Relationship between asymmetric dimethylarginine, nitrite and genetic polymorphisms: Impact on erectile dysfunction therapy. Nitric Oxide - Biology and Chemistry, 2017, 71, 44-51.	2.7	7
7	Effects of arginase genetic polymorphisms on nitric oxide formation in healthy pregnancy and in preeclampsia. Nitric Oxide - Biology and Chemistry, 2021, 109-110, 20-25.	2.7	6
8	Arginase II polymorphisms modify the hypotensive responses to propofol by affecting nitric oxide bioavailability. European Journal of Clinical Pharmacology, 2021, 77, 869-877.	1.9	6
9	DDAH1 and DDAH2 polymorphisms associate with asymmetrical dimethylarginine plasma levels in erectile dysfunction patients but not in healthy controls. Nitric Oxide - Biology and Chemistry, 2019, 92, 11-17.	2.7	5
10	Association of 11Î²-hydroxysteroid dehydrogenase type1 (HSD11b1) gene polymorphisms with outcome of antidepressant therapy and suicide attempts. Behavioural Brain Research, 2020, 381, 112343.	2.2	5
11	Clinical, Molecular, Functional, and Structural Characterization of CYP17A1 Mutations in Brazilian Patients with 17-Hydroxylase Deficiency. Hormone and Metabolic Research, 2020, 52, 186-193.	1.5	5
12	Interaction Between NOS3 and HMOX1 on Antihypertensive Drug Responsiveness in Preeclampsia. Revista Brasileira De Ginecologia E Obstetricia, 2020, 42, 460-467.	0.8	5
13	How do sphingolipids play a role in epigenetic mechanisms and gene expression?. Epigenomics, 2022, 14, 219-222.	2.1	4
14	Vitamin D receptor hypermethylation as a biomarker for pediatric adrenocortical tumors. European Journal of Endocrinology, 2022, 186, 573-585.	3.7	4
15	Circulating HO-1 levels are not associated with plasma sFLT-1 and GT_n HMOX1 polymorphism in preeclampsia. Hypertension in Pregnancy, 2019, 38, 73-77.	1.1	3
16	Telomere length and Wnt/Î²-catenin pathway in adamantinomatous craniopharyngiomas. European Journal of Endocrinology, 2022, 187, 219-230.	3.7	3
17	Comments to â€œA rational, non-radioactive strategy for the molecular diagnosis of congenital adrenal hyperplasia due to 21-hydroxylase deficiencyâ€. Gene, 2014, 534, 451-452.	2.2	0
18	Tumor DNA Methylation Profiling Is a Robust and Independent Prognostic Marker for Pediatric Patients With Adrenocortical Tumors. Journal of the Endocrine Society, 2021, 5, A99-A100.	0.2	0