Aurore Carre

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5647831/publications.pdf

Version: 2024-02-01

28	1,063	394421 19	477307 29
papers	citations	h-index	g-index

30	30	30	1067
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetics of congenital hypothyroidism: Modern concepts. Pediatric Investigation, 2022, 6, 123-134.	1.4	8
2	New genetics in congenital hypothyroidism. Endocrine, 2021, 71, 696-705.	2.3	33
3	Ex vivo model for elucidating the functional and structural differentiation of the embryonic mouse thyroid. Molecular and Cellular Endocrinology, 2020, 510, 110834.	3.2	2
4	High Diagnostic Yield of Targeted Next-Generation Sequencing in a Cohort of Patients With Congenital Hypothyroidism Due to Dyshormonogenesis. Frontiers in Endocrinology, 2020, 11, 545339.	3.5	17
5	<scp>TUBB</scp> 1 mutations cause thyroid dysgenesis associated with abnormal platelet physiology. EMBO Molecular Medicine, 2018, 10, .	6.9	47
6	Thyroid Hypoplasia in Congenital Hypothyroidism Associated with Thyroid Peroxidase Mutations. Thyroid, 2018, 28, 941-944.	4.5	23
7	Mutations in BOREALIN cause thyroid dysgenesis. Human Molecular Genetics, 2017, 26, ddw419.	2.9	37
8	NADPH Oxidase NOX4 Is a Critical Mediator of BRAF^{V600E}-Induced Downregulation of the Sodium/Iodide Symporter in Papillary Thyroid Carcinomas. Antioxidants and Redox Signaling, 2017, 26, 864-877.	5.4	63
9	Update of Thyroid Developmental Genes. Endocrinology and Metabolism Clinics of North America, 2016, 45, 243-254.	3.2	29
10	When an Intramolecular Disulfide Bridge Governs the Interaction of DUOX2 with Its Partner DUOXA2. Antioxidants and Redox Signaling, 2015, 23, 724-733.	5.4	29
11	Molecular Insights into the Possible Role of Kir4.1 and Kir5.1 in Thyroid Hormone Biosynthesis. Hormone Research in Paediatrics, 2015, 83, 141-147.	1.8	5
12	Down Syndrome and Nonautoimmune Hypothyroidisms in Neonates and Infants. Hormone Research in Paediatrics, 2015, 83, 126-131.	1.8	23
13	Functional characterization of the novel sequence variant p.S304R in the hinge region of TSHR in a congenital hypothyroidism patients and analogy with other formerly known mutations of this gene portion. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 777-84.	0.9	4
14	DYRK1A BAC Transgenic Mouse: A New Model of Thyroid Dysgenesis in Down Syndrome. Endocrinology, 2015, 156, 1171-1180.	2.8	20
15	A NovelFOXE1Mutation (R73S) in Bamforthâ€“Lazarus Syndrome Causing Increased Thyroidal Gene Expression. Thyroid, 2014, 24, 649-654.	4.5	38
16	Further delineation of the phenotype of chromosome 14q13 deletions: (positional) involvement ofFOXG1appears the main determinant of phenotype severity, with no evidence for a holoprosencephaly locus. Journal of Medical Genetics, 2012, 49, 366-372.	3.2	24
17	Multiplex Ligation-Dependent Probe Amplification Improves the Detection Rate of <i>NKX2.1</i> Mutations in Patients Affected by Brain-Lung-Thyroid Syndrome. Hormone Research in Paediatrics, 2012, 77, 146-151.	1.8	20
18	Thyroid Function in Fetuses with Down Syndrome. Hormone Research in Paediatrics, 2012, 78, 88-93.	1.8	19

#	Article	IF	Citations
19	Hes1 Is Required for Appropriate Morphogenesis and Differentiation during Mouse Thyroid Gland Development. PLoS ONE, 2011, 6, e16752.	2.5	40
20	Pregnancy in women heterozygous for MCT8 mutations: risk of maternal hypothyroxinemia and fetal care. European Journal of Endocrinology, 2011, 164, 309-314.	3.7	19
21	NKX2-1mutations leading to surfactant protein promoter dysregulation cause interstitial lung disease in â€œBrain-Lung-Thyroid Syndromeâ€. Human Mutation, 2010, 31, E1146-E1162.	2.5	108
22	New Cases of Isolated Congenital Central Hypothyroidism Due to Homozygous Thyrotropin Beta Gene Mutations: A Pitfall to Neonatal Screening. Thyroid, 2010, 20, 639-645.	4.5	24
23	An Inactivating Mutation within the First Extracellular Loop of the Thyrotropin Receptor Impedes Normal Posttranslational Maturation of the Extracellular Domain. Endocrinology, 2009, 150, 1043-1050.	2.8	32
24	Five new TTF1/NKX2.1 mutations in brain-lung-thyroid syndrome: rescue by PAX8 synergism in one case. Human Molecular Genetics, 2009, 18, 2266-2276.	2.9	187
25	Sodium/Iodide Symporter (NIS) Gene Expression Is the Limiting Step for the Onset of Thyroid Function in the Human Fetus. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 70-76.	3.6	74
26	Polymorphic length of FOXE1 alanine stretch: evidence for genetic susceptibility to thyroid dysgenesis. Human Genetics, 2007, 122, 467-476.	3.8	61
27	Linkage and mutational analysis of familial thyroid dysgenesis demonstrate genetic heterogeneity implicating novel genes. European Journal of Human Genetics, 2005, 13, 232-239.	2.8	49
28	Molecular Mechanisms of Thyroid Dysgenesis. Hormone Research in Paediatrics, 2004, 62, 14-21.	1.8	27