

PÃ©ter Gergics

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4062213/publications.pdf>

Version: 2024-02-01

10
papers

251
citations

1307594

7
h-index

1372567

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g-index

12
all docs

12
docs citations

12
times ranked

473
citing authors

#	ARTICLE	IF	CITATIONS
1	The phenotypic spectrum associated with OTX2 mutations in humans. <i>European Journal of Endocrinology</i> , 2021, 185, 121-135.	3.7	15
2	High-throughput splicing assays identify missense and silent splice-disruptive POU1F1 variants underlying pituitary hormone deficiency. <i>American Journal of Human Genetics</i> , 2021, 108, 1526-1539.	6.2	23
3	Murine SEC24D can substitute functionally for SEC24C during embryonic development. <i>Scientific Reports</i> , 2021, 11, 21100.	3.3	3
4	Otx2b mutant zebrafish have pituitary, eye and mandible defects that model mammalian disease. <i>Human Molecular Genetics</i> , 2020, 29, 1648-1657.	2.9	6
5	Pituitary Transcription Factor Mutations Leading to Hypopituitarism. <i>Experientia Supplementum</i> (2012), 2019, 111, 263-298.	0.9	4
6	Gene Expression in Mouse Thyrotrope Adenoma: Transcription Elongation Factor Stimulates Proliferation. <i>Endocrinology</i> , 2016, 157, 3631-3646.	2.8	13
7	ISL1-based LIM complexes control Slit2 transcription in developing cranial motor neurons. <i>Scientific Reports</i> , 2016, 6, 36491.	3.3	16
8	Genetics of Combined Pituitary Hormone Deficiency: Roadmap into the Genome Era. <i>Endocrine Reviews</i> , 2016, 37, 636-675.	20.1	147
9	Lhx4 Deficiency: Increased Cyclin-Dependent Kinase Inhibitor Expression and Pituitary Hypoplasia. <i>Molecular Endocrinology</i> , 2015, 29, 597-612.	3.7	11
10	The pattern of congenital heart defects arising from reduced Tbx5 expression is altered in a Down syndrome mouse model. <i>BMC Developmental Biology</i> , 2015, 15, 30.	2.1	12