Angelo Molinaro

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6481816/publications.pdf

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14	448	759233 12	1058476 14
papers	citations	h-index	g-index

14	14	14	919
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Chondroitin 6-sulphate is required for neuroplasticity and memory in ageing. Molecular Psychiatry, 2021, 26, 5658-5668.	7.9	36
2	Brain mitochondrial proteome alteration driven by creatine deficiency suggests novel therapeutic venues for creatine deficiency syndromes. Neuroscience, 2019, 409, 276-289.	2.3	8
3	A Nervous System-Specific Model of Creatine Transporter Deficiency Recapitulates the Cognitive Endophenotype of the Disease: a Longitudinal Study. Scientific Reports, 2019, 9, 62.	3.3	14
4	iPSC-derived neurons profiling reveals GABAergic circuit disruption and acetylated Î±-tubulin defect which improves after iHDAC6 treatment in Rett syndrome. Experimental Cell Research, 2018, 368, 225-235.	2.6	36
5	A Large Inversion Involving GNAS Exon A/B and All Exons Encoding GsÎ± Is Associated With Autosomal Dominant Pseudohypoparathyroidism Type Ib (PHP1B). Journal of Bone and Mineral Research, 2017, 32, 776-783.	2.8	22
6	Proteomic analysis of fine-needle aspiration in differential diagnosis of thyroid nodules. Translational Research, 2016, 176, 81-94.	5.0	25
7	A mouse model for creatine transporter deficiency reveals early onset cognitive impairment and neuropathology associated with brain aging. Human Molecular Genetics, 2016, 25, 4186-4200.	2.9	39
8	Analysis of Multiple Families With Single Individuals Affected by Pseudohypoparathyroidism Type Ib (PHP1B) Reveals Only One Novel Maternally Inherited GNAS Deletion. Journal of Bone and Mineral Research, 2016, 31, 796-805.	2.8	31
9	Similar frequency of paternal uniparental disomy involving chromosome 20q (patUPD20q) in Japanese and Caucasian patients affected by sporadic pseudohypoparathyroidism type Ib (sporPHP1B). Bone, 2015, 79, 15-20.	2.9	41
10	TSH Elevations as the First Laboratory Evidence for Pseudohypoparathyroidism Type Ib (PHP-Ib). Journal of Bone and Mineral Research, 2015, 30, 906-912.	2.8	28
11	BRAF mutation analysis in thyroid nodules with indeterminate cytology: our experience on surgical management of patients with thyroid nodules from an area of borderline iodine deficiency. Journal of Endocrinological Investigation, 2014, 37, 1009-1014.	3.3	21
12	Presence in the Pre-Surgical Fine-Needle Aspiration of Potential Thyroid Biomarkers Previously Identified in the Post-Surgical One. PLoS ONE, 2013, 8, e72911.	2.5	18
13	MicroRNA expression profile helps to distinguish benign nodules from papillary thyroid carcinomas starting from cells of fine-needle aspiration. European Journal of Endocrinology, 2012, 167, 393-400.	3.7	67
14	Identification and Functional Analysis of Novel Dual Oxidase 2 (DUOX2) Mutations in Children with Congenital or Subclinical Hypothyroidism. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E1335-E1339.	3.6	62