## Nikolaos Settas

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1343199/publications.pdf

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13	220	1307594 <b>7</b>	1125743
papers	citations	h-index	g-index
14	14	14	387
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Somatic USP8 Gene Mutations Are a Common Cause of Pediatric Cushing Disease. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 2836-2843.	3.6	81
2	Succinate dehydrogenase (SDH) deficiency, Carney triad and the epigenome. Molecular and Cellular Endocrinology, 2018, 469, 107-111.	3.2	45
3	SGPL1 Deficiency: A Rare Cause of Primary Adrenal Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 1484-1490.	3.6	27
4	Carney Triad, Carney-Stratakis Syndrome, 3PAS and Other Tumors Due to SDH Deficiency. Frontiers in Endocrinology, 2021, 12, 680609.	3.5	11
5	Variants in PRKAR1B cause a neurodevelopmental disorder with autism spectrum disorder, apraxia, and insensitivity to pain. Genetics in Medicine, 2021, 23, 1465-1473.	2.4	10
6	The E3 ubiquitin ligase Siah1 regulates adrenal gland organization and aldosterone secretion. JCI Insight, 2017, 2, .	5.0	9
7	Genomic and sequence variants of protein kinase A regulatory subunit type $1\hat{l}^2$ (PRKAR1B) in patients with adrenocortical disease and Cushing syndrome. Genetics in Medicine, 2021, 23, 174-182.	2.4	8
8	ARMC5 variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. Endocrine-Related Cancer, 2020, 27, 509-517.	3.1	7
9	PRKACB variants in skeletal disease or adrenocortical hyperplasia: effects on protein kinase A. Endocrine-Related Cancer, 2020, 27, 647-656.	3.1	7
10	USP13 genetics and expression in a family with thyroid cancer. Endocrine, 2022, 77, 281-290.	2.3	5
11	Medullary thyroid cancer, leukemia, mesothelioma and meningioma associated with germline APC and RASAL1 variants: a new syndrome?. Hormones, 2018, 16, 423-428.	1.9	3
12	Kisspeptin deficiency leads to abnormal adrenal glands and excess steroid hormone secretion. Human Molecular Genetics, 2020, 29, 3443-3450.	2.9	3
13	The PRKAR1B p.R115K Variant is Associated with Lipoprotein Profile in African American Youth with Metabolic Challenges. Journal of the Endocrine Society, 2021, 5, bvab071.	0.2	3