

Karlijn Pellikaan

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

232
citations

1162367

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h-index

1058022

14
g-index

21
all docs

21
docs citations

21
times ranked

255
citing authors

#	ARTICLE	IF	CITATIONS
1	We mind your step: understanding and preventing drop-out in the transfer from paediatric to adult tertiary endocrine healthcare. <i>Endocrine Connections</i> , 2022, 11, .	0.8	4
2	Health Problems in Adults with Prader-Willi Syndrome of Different Genetic Subtypes: Cohort Study, Meta-Analysis and Review of the Literature. <i>Journal of Clinical Medicine</i> , 2022, 11, 4033.	1.0	8
3	Transition readiness among adolescents with rare endocrine conditions. <i>Endocrine Connections</i> , 2021, 10, 432-446.	0.8	5
4	Thyroid Function in Adults With Prader-Willi Syndrome. <i>Journal of the Endocrine Society</i> , 2021, 5, A853-A853.	0.1	0
5	Growth Hormone Treatment for Adults With Prader-Willi Syndrome: A Meta-Analysis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, 3068-3091.	1.8	15
6	The Diagnostic Journey of a Patient with Prader-Willi-Like Syndrome and a Unique Homozygous SNURF-SNRPN Variant; Bio-Molecular Analysis and Review of the Literature. <i>Genes</i> , 2021, 12, 875.	1.0	4
7	Effects of Childhood Multidisciplinary Care and Growth Hormone Treatment on Health Problems in Adults with Prader-Willi Syndrome. <i>Journal of Clinical Medicine</i> , 2021, 10, 3250.	1.0	10
8	Thyroid Function in Adults with Prader-Willi Syndrome; a Cohort Study and Literature Review. <i>Journal of Clinical Medicine</i> , 2021, 10, 3804.	1.0	13
9	Hyponatremia in Children and Adults with Prader-Willi Syndrome: A Survey Involving Seven Countries. <i>Journal of Clinical Medicine</i> , 2021, 10, 3555.	1.0	4
10	Hyperprolactinemia in Adults with Prader-Willi Syndrome. <i>Journal of Clinical Medicine</i> , 2021, 10, 3613.	1.0	4
11	Hypogonadism in Adult Males with Prader-Willi Syndrome—Clinical Recommendations Based on a Dutch Cohort Study, Review of the Literature and an International Expert Panel Discussion. <i>Journal of Clinical Medicine</i> , 2021, 10, 4361.	1.0	16
12	What Every Internist-Endocrinologist Should Know about Rare Genetic Syndromes in Order to Prevent Needless Diagnostics, Missed Diagnoses and Medical Complications: Five Years of Internal Medicine for Rare Genetic Syndromes™. <i>Journal of Clinical Medicine</i> , 2021, 10, 5457.	1.0	7
13	Hypogonadism in Women with Prader-Willi Syndrome—Clinical Recommendations Based on a Dutch Cohort Study, Review of the Literature and an International Expert Panel Discussion. <i>Journal of Clinical Medicine</i> , 2021, 10, 5781.	1.0	12
14	Missed Diagnoses and Health Problems in Adults With Prader-Willi Syndrome: Recommendations for Screening and Treatment. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e4671-e4687.	1.8	40
15	Central Adrenal Insufficiency Is Rare in Adults With Prader-Willi Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e2563-e2571.	1.8	27
16	MON-284 Systematic Screening Reveals Large Number of Undiagnosed and Untreated Cardiovascular Risk Factors in Adults with Prader-Willi Syndrome. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.1	0
17	SUN-080 We Mind Your Step: Understanding and Preventing Drop-Out in the Transition from Paediatric to Adult Tertiary Endocrine Healthcare. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.1	0
18	SUN-308 Central Adrenal Insufficiency Is Rare in Adults with Prader-Willi Syndrome. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.1	0

#	ARTICLE	IF	CITATIONS
19	Intracranial Carotid Artery Calcification From Infancy to Old Age. <i>Journal of the American College of Cardiology</i> , 2018, 72, 582-584.	1.2	17
20	Automatic segmentation and quantification of the cardiac structures from non-contrast-enhanced cardiac CT scans. <i>Physics in Medicine and Biology</i> , 2017, 62, 3798-3813.	1.6	26
21	Intracranial Carotid Artery Calcification Relates to Recanalization and Clinical Outcome After Mechanical Thrombectomy. <i>Stroke</i> , 2017, 48, 342-347.	1.0	20