

Laura C G De Graaff

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

287
citations

1039406

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996533

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docs citations

28
times ranked

266
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	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
3	Genetic screening of a Dutch population with isolated GH deficiency (IGHD). <i>Clinical Endocrinology</i> , 2009, 70, 742-750.	1.2	24
4	A value-based healthcare approach: Health-related quality of life and psychosocial functioning in women with Turner syndrome. <i>Clinical Endocrinology</i> , 2020, 92, 434-442.	1.2	20
5	Three years of growth hormone treatment in young adults with Prader-Willi syndrome: sustained positive effects on body composition. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 163.	1.2	17
6	Body proportions before and during growth hormone therapy in children with chronic renal failure. <i>Pediatric Nephrology</i> , 2003, 18, 679-684.	0.9	16
7	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
8	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
9	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
10	Three years of growth hormone treatment in young adults with Prader-Willi Syndrome previously treated with growth hormone in childhood: Effects on glucose homeostasis and metabolic syndrome. <i>Clinical Endocrinology</i> , 2020, 93, 439-448.	1.2	12
11	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
12	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
13	Genetic Polymorphisms in the Locus Control Region and Promoter of <i>GH1</i> Are Related to Serum IGF-I Levels and Height in Patients with Isolated Growth Hormone Deficiency and Healthy Controls. <i>Hormone Research in Paediatrics</i> , 2010, 73, 25-34.	0.8	8
14	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
15	Unique near-complete deletion of <i>GLI2</i> in a patient with combined pituitary hormone deficiency and post-axial polydactyly. <i>Growth Hormone and IGF Research</i> , 2020, 50, 35-41.	0.5	7
16	What Every Internist-Endocrinologist Should Know about Rare Genetic Syndromes in Order to Prevent Needless Diagnostics, Missed Diagnoses and Medical Complications: Five Years of <i>Internal Medicine for Rare Genetic Syndromes</i> [™] . <i>Journal of Clinical Medicine</i> , 2021, 10, 5457.	1.0	7
17	Transition readiness among adolescents with rare endocrine conditions. <i>Endocrine Connections</i> , 2021, 10, 432-446.	0.8	5
18	Bone mineral density during 3 years of growth hormone in previously GH-treated young adults with PWS. <i>European Journal of Endocrinology</i> , 2021, 184, 773-782.	1.9	4

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19	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
20	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
21	Hyperprolactinemia in Adults with Prader-Willi Syndrome. <i>Journal of Clinical Medicine</i> , 2021, 10, 3613.	1.0	4
22	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
23	Genetic analysis of IRF6, a gene involved in craniofacial midline formation, in relation to pituitary and facial morphology of patients with idiopathic growth hormone deficiency. <i>Pituitary</i> , 2017, 20, 499-508.	1.6	3
24	Case Report: A Detailed Phenotypic Description of Patients and Relatives with Combined Central Hypothyroidism and Growth Hormone Deficiency Carrying IGSF1 Mutations. <i>Genes</i> , 2022, 13, 623.	1.0	3
25	Congenital hypopituitarism in two brothers with a duplication of the acro gigantism gene™ GPR101: clinical findings and review of the literature. <i>Pituitary</i> , 2021, 24, 229-241.	1.6	2
26	Family Matters: Trauma and Quality of Life in Family Members of Individuals With Prader-Willi Syndrome. <i>Frontiers in Psychiatry</i> , 0, 13, .	1.3	2
27	Thyroid Function in Adults With Prader-Willi Syndrome. <i>Journal of the Endocrine Society</i> , 2021, 5, A853-A853.	0.1	0
28	What Every Internist Should Know About Rare Genetic Syndromes in Order to Prevent Needless Diagnostics, Missed Diagnoses and Medical Complications: Five-Year Experience of Internal Medicine for Complex Rare Genetic Syndromes. <i>Journal of the Endocrine Society</i> , 2021, 5, A513-A514.	0.1	0