Laura C G De Graaff

List of Publications by Year
in descending order

28	287	1040056 9	996975 15
papers	citations	h-index	g-index

28	28	28	266
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Missed Diagnoses and Health Problems in Adults With Prader-Willi Syndrome: Recommendations for Screening and Treatment. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e4671-e4687.	3.6	40
2	Central Adrenal Insufficiency Is Rare in Adults With Praderâ€“Willi Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2563-e2571.	3.6	27
3	Genetic screening of a Dutch population with isolated GH deficiency (IGHD). Clinical Endocrinology, 2009, 70, 742-750.	2.4	24
4	A valueâ€based healthcare approach: Healthâ€related quality of life and psychosocial functioning in women with Turner syndrome. Clinical Endocrinology, 2020, 92, 434-442.	2.4	20
5	Three years of growth hormone treatment in young adults with Prader-Willi syndrome: sustained positive effects on body composition. Orphanet Journal of Rare Diseases, 2020, 15, 163.	2.7	17
6	Body proportions before and during growth hormone therapy in children with chronic renal failure. Pediatric Nephrology, 2003, 18, 679-684.	1.7	16
7	Hypogonadism in Adult Males with Prader-Willi Syndromeâ€”Clinical Recommendations Based on a Dutch Cohort Study, Review of the Literature and an International Expert Panel Discussion. Journal of Clinical Medicine, 2021, 10, 4361.	2.4	16
8	Growth Hormone Treatment for Adults With Prader-Willi Syndrome: A Meta-Analysis. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 3068-3091.	3.6	15
9	Thyroid Function in Adults with Praderâ€“Willi Syndrome; a Cohort Study and Literature Review. Journal of Clinical Medicine, 2021, 10, 3804.	2.4	13
10	Three years of growth hormone treatment in young adults with Praderâ€Willi Syndrome previously treated with growth hormone in childhood: Effects on glucose homeostasis and metabolic syndrome. Clinical Endocrinology, 2020, 93, 439-448.	2.4	12
11	Hypogonadism in Women with Prader-Willi Syndromeâ€”Clinical Recommendations Based on a Dutch Cohort Study, Review of the Literature and an International Expert Panel Discussion. Journal of Clinical Medicine, 2021, 10, 5781.	2.4	12
12	Effects of Childhood Multidisciplinary Care and Growth Hormone Treatment on Health Problems in Adults with Prader-Willi Syndrome. Journal of Clinical Medicine, 2021, 10, 3250.	2.4	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. Hormone Research in Paediatrics, 2010, 73, 25-34.	1.8	8
14	Health Problems in Adults with Praderâ€“Willi Syndrome of Different Genetic Subtypes: Cohort Study, Meta-Analysis and Review of the Literature. Journal of Clinical Medicine, 2022, 11, 4033.	2.4	8
15	Unique near-complete deletion of GLI2 in a patient with combined pituitary hormone deficiency and post-axial polydactyly. Growth Hormone and IGF Research, 2020, 50, 35-41.	1.1	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 â€˜Internal Medicine for Rare Genetic Syndromesâ€™. Journal of Clinical Medicine, 2021, 10, 5457.	2.4	7
17	Transition readiness among adolescents with rare endocrine conditions. Endocrine Connections, 2021, 10, 432-446.	1.9	5
18	Bone mineral density during 3 years of growth hormone in previously GH-treated young adults with PWS. European Journal of Endocrinology, 2021, 184, 773-782.	3.7	4

#	Article	IF	Citations
19	The Diagnostic Journey of a Patient with Praderâ€“Willi-Like Syndrome and a Unique Homozygous SNURF-SNRPN Variant; Bio-Molecular Analysis and Review of the Literature. Genes, 2021, 12, 875.	2.4	4
20	Hyponatremia in Children and Adults with Praderâ€“Willi Syndrome: A Survey Involving Seven Countries. Journal of Clinical Medicine, 2021, 10, 3555.	2.4	4
21	Hyperprolactinemia in Adults with Prader-Willi Syndrome. Journal of Clinical Medicine, 2021, 10, 3613.	2.4	4
22	We mind your step: understanding and preventing drop-out in the transfer from paediatric to adult tertiary endocrine healthcare. Endocrine Connections, 2022, 11, .	1.9	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. Pituitary, 2017, 20, 499-508.	2.9	3
24	Case Report: A Detailed Phenotypic Description of Patients and Relatives with Combined Central Hypothyroidism and Growth Hormone Deficiency Carrying IGSF1 Mutations. Genes, 2022, 13, 623.	2.4	3
25	Congenital hypopituitarism in two brothers with a duplication of the â€˜acrogigantism geneâ€™ GPR101: clinical findings and review of the literature. Pituitary, 2021, 24, 229-241.	2.9	2
26	Family Matters: Trauma and Quality of Life in Family Members of Individuals With Prader-Willi Syndrome. Frontiers in Psychiatry, 0, 13, .	2.6	2
27	Thyroid Function in Adults With Prader-Willi Syndrome. Journal of the Endocrine Society, 2021, 5, A853-A853.	0.2	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. Journal of the Endocrine Society, 2021, 5, A513-A514.	0.2	0