Laura C G De Graaff

List of Publications by Year
in descending order

28	287	1039406 9	996533 15
papers	citations	h-index	g-index

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

#	Article	IF	Citations
1	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.	1.0	3
2	We mind your step: understanding and preventing drop-out in the transfer from paediatric to adult tertiary endocrine healthcare. Endocrine Connections, 2022, 11, .	0.8	4
3	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.	1.0	8
4	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.	1.6	2
5	Transition readiness among adolescents with rare endocrine conditions. Endocrine Connections, 2021, 10, 432-446.	0.8	5
6	Thyroid Function in Adults With Prader-Willi Syndrome. Journal of the Endocrine Society, 2021, 5, A853-A853.	0.1	0
7	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.	0.1	0
8	Growth Hormone Treatment for Adults With Prader-Willi Syndrome: A Meta-Analysis. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 3068-3091.	1.8	15
9	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.	1.9	4
10	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.	1.0	4
11	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.	1.0	10
12	Thyroid Function in Adults with Praderâ€“Willi Syndrome; a Cohort Study and Literature Review. Journal of Clinical Medicine, 2021, 10, 3804.	1.0	13
13	Hyponatremia in Children and Adults with Praderâ€“Willi Syndrome: A Survey Involving Seven Countries. Journal of Clinical Medicine, 2021, 10, 3555.	1.0	4
14	Hyperprolactinemia in Adults with Prader-Willi Syndrome. Journal of Clinical Medicine, 2021, 10, 3613.	1.0	4
15	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.	1.0	16
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.	1.0	7
17	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.	1.0	12
18	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.	0.5	7

#	Article	IF	Citations
19	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.	1.8	40
20	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.	1.2	12
21	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.	1.2	17
22	A valueâ€based healthcare approach: Healthâ€related quality of life and psychosocial functioning in women with Turner syndrome. Clinical Endocrinology, 2020, 92, 434-442.	1.2	20
23	Central Adrenal Insufficiency Is Rare in Adults With Praderâ€“Willi Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2563-e2571.	1.8	27
24	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.	1.6	3
25	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.	0.8	8
26	Genetic screening of a Dutch population with isolated GH deficiency (IGHD). Clinical Endocrinology, 2009, 70, 742-750.	1.2	24
27	Body proportions before and during growth hormone therapy in children with chronic renal failure. Pediatric Nephrology, 2003, 18, 679-684.	0.9	16
28	Family Matters: Trauma and Quality of Life in Family Members of Individuals With Prader-Willi Syndrome. Frontiers in Psychiatry, 0, 13, .	1.3	2