Guy Van Vliet

List of Publications by Year
in descending order

28 papers	1,432 citations	566801 15 h-index	580395 25 g-index
30 all docs	30 docs citations	30 times ranked	1444 citing authors

#	Article	IF	Citations
1	Is the Incidence of Congenital Hypothyroidism Really Increasing? A 20-Year Retrospective Population-Based Study in QuÃ©bec. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 2422-2429.	1.8	196
2	European Society for Paediatric Endocrinology Consensus Guidelines on Screening, Diagnosis, and Management of Congenital Hypothyroidism. Hormone Research in Paediatrics, 2014, 81, 80-103.	0.8	193
3	A Search for the Possible Molecular Mechanisms of Thyroid Dysgenesis: Sex Ratios and Associated Malformations. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 2502-2506.	1.8	172
4	Prevention of intellectual disability through screening for congenital hypothyroidism: how much and at what level?. Archives of Disease in Childhood, 2011, 96, 374-379.	1.0	169
5	Primary Adrenal Insufficiency in Children: Twenty Years Experience at the Sainte-Justine Hospital, Montreal. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 3243-3250.	1.8	152
6	Sudden death in growth hormoneâ€“treated children with Prader-Willi syndrome. Journal of Pediatrics, 2004, 144, 129-131.	0.9	102
7	Mutations in NFKB2and potential genetic heterogeneity in patients with DAVID syndrome, having variable endocrine and immune deficiencies. BMC Medical Genetics, 2014, 15, 139.	2.1	84
8	Are lower TSH cutoffs in neonatal screening for congenital hypothyroidism warranted?. European Journal of Endocrinology, 2017, 177, D1-D12.	1.9	81
9	How Many Deaths Can Be Prevented by Newborn Screening for Congenital Adrenal Hyperplasia?. Hormone Research in Paediatrics, 2007, 67, 284-291.	0.8	63
10	Screening for neonatal endocrinopathies: rationale, methods and results. Seminars in Fetal and Neonatal Medicine, 2004, 9, 75-85.	2.8	48
11	A Search for the Possible Molecular Mechanisms of Thyroid Dysgenesis: Sex Ratios and Associated Malformations. , 0, .		31
12	Treating fetal thyroid and adrenal disorders through the mother. Nature Clinical Practice Endocrinology and Metabolism, 2008, 4, 675-682.	2.9	24
13	Non-Immune Goiter and Hypothyroidism in a 19-Week Fetus: A Plea for Conservative Treatment. Journal of Pediatrics, 2010, 156, 1026-1029.	0.9	20
14	Nine novel mutations inNR0B1 (DAX1) causing adrenal hypoplasia congenita. Human Mutation, 2001, 18, 547-547.	1.1	16
15	Wide Spectrum of DUOX2 Deficiency: From Life-Threatening Compressive Goiter in Infancy to Lifelong Euthyroidism. Thyroid, 2019, 29, 1018-1022.	2.4	16
16	Demonstration of Autosomal Monoallelic Expression in Thyroid Tissue Assessed by Whole-Exome and Bulk RNA Sequencing. Thyroid, 2016, 26, 852-859.	2.4	15
17	Challenges in Assessing the Cost-Effectiveness of Newborn Screening: The Example of Congenital Adrenal Hyperplasia. International Journal of Neonatal Screening, 2020, 6, 82.	1.2	11
18	Avoiding the Overdiagnosis of Congenital Hypothyroidism in Premature Newborns. Pediatrics, 2019, 144, e20191706.	1.0	8

#	Article	IF	Citations
19	Interpreting Minor Variations in Thyroid Function or Echostructure. Pediatric Clinics of North America, 2015, 62, 929-942.	0.9	6
20	Redefining Congenital Hypothyroidism?. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1463-e1465.	1.8	5
21	Similar age-dependent levothyroxine requirements of schoolchildren with congenital or acquired hypothyroidism. European Journal of Pediatrics, 2016, 175, 869-872.	1.3	4
22	Quality of referral of short children to the paediatric endocrinologist and impact of a fax communication system. Paediatrics and Child Health, 2013, 18, 533-537.	0.3	3
23	Whole-exome sequencing: opportunities in pediatric endocrinology. Personalized Medicine, 2014, 11, 63-78.	0.8	3
24	Severe Congenital Hypothyroidism Due to a Novel Deep Intronic Mutation in the TSH Receptor Gene Causing Intron Retention. Journal of the Endocrine Society, 2021, 5, bvaa183.	0.1	3
25	Growth Hormone Supplementation and Psychosocial Functioning to Adult Height in Turner Syndrome: A Questionnaire Study of Participants in the Canadian Randomized Trial. Frontiers in Endocrinology, 2019, 10, 125.	1.5	1
26	Population-based TSH Screening of Newborns for Hyperthyroidism: It May Be Feasible, but Is It Justified?. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2630-e2631.	1.8	1
27	Longitudinal study of thyroid function in Down's syndrome in the first two decades. Journal of Pediatrics, 2005, 147, 707-708.	0.9	0
28	Conserved Telomere Length in Human Ectopic Thyroids: An Argument Against Premature Differentiation Causing Arrested Migration. Thyroid, 2015, 25, 1050-1054.	2.4	0