Guy Van Vliet

List of Publications by Year in Descending Order

27 papers	1,071 citations	13 h-index	30 g-index
30 ext. papers	1,283 ext. citations	4.1 avg, IF	4.13 L-index

#	Paper	IF	Citations
27	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	0.4	1
26	Redefining Congenital Hypothyroidism?. *Journal of Clinical Endocrinology and Metabolism*, **2021**, 106, e1463-e1465	5.6	0
25	Challenges in Assessing the Cost-Effectiveness of Newborn Screening: The Example of Congenital Adrenal Hyperplasia. *International Journal of Neonatal Screening*, **2020**, 6,	2.6	2
24	Wide Spectrum of Deficiency: From Life-Threatening Compressive Goiter in Infancy to Lifelong Euthyroidism. *Thyroid*, **2019**, 29, 1018-1022	6.2	10
23	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	5.7	1
22	Avoiding the Overdiagnosis of Congenital Hypothyroidism in Premature Newborns. *Pediatrics*, **2019** , 144,	7.4	4
21	Are lower TSH cutoffs in neonatal screening for congenital hypothyroidism warranted?. *European Journal of Endocrinology*, **2017**, 177, D1-D12	6.5	51
20	Similar age-dependent levothyroxine requirements of schoolchildren with congenital or acquired hypothyroidism. *European Journal of Pediatrics*, **2016**, 175, 869-72	4.1	3
19	Demonstration of Autosomal Monoallelic Expression in Thyroid Tissue Assessed by Whole-Exome and Bulk RNA Sequencing. *Thyroid*, **2016**, 26, 852-9	6.2	10
18	Interpreting Minor Variations in Thyroid Function or Echostructure: Treating Patients, Not Numbers or Images. *Pediatric Clinics of North America*, **2015**, 62, 929-42	3.6	5
17	Conserved Telomere Length in Human Ectopic Thyroids: An Argument Against Premature Differentiation Causing Arrested Migration. *Thyroid*, **2015**, 25, 1050-4	6.2	
16	Whole-exome sequencing: opportunities in pediatric endocrinology. *Personalized Medicine*, **2014**, 11, 63-78	2.2	2
15	European Society for Paediatric Endocrinology consensus guidelines on screening, diagnosis, and management of congenital hypothyroidism. *Hormone Research in Paediatrics*, **2014**, 81, 80-103	3.3	136
14	Mutations in NFKB2 and potential genetic heterogeneity in patients with DAVID syndrome, having variable endocrine and immune deficiencies. *BMC Medical Genetics*, **2014**, 15, 139	2.1	56
13	Quality of referral of short children to the paediatric endocrinologist and impact of a fax communication system. *Paediatrics and Child Health*, **2013**, 18, 533-7	0.7	2
12	Is the incidence of congenital hypothyroidism really increasing? A 20-year retrospective population-based study in QuÞec. *Journal of Clinical Endocrinology and Metabolism*, **2011**, 96, 2422-9	5.6	143
11	Prevention of intellectual disability through screening for congenital hypothyroidism: how much and at what level?. *Archives of Disease in Childhood*, **2011**, 96, 374-9	2.2	134

10	Non-immune goiter and hypothyroidism in a 19-week fetus: a plea for conservative treatment. *Journal of Pediatrics*, **2010**, 156, 1026-1029	3.6	17
9	Treating fetal thyroid and adrenal disorders through the mother. *Nature Clinical Practice Endocrinology and Metabolism*, **2008**, 4, 675-82		18
8	How many deaths can be prevented by newborn screening for congenital adrenal hyperplasia?. *Hormone Research in Paediatrics*, **2007**, 67, 284-91	3.3	45
7	Longitudinal study of thyroid function in Down's syndrome in the first two decades. *Journal of Pediatrics*, **2005**, 147, 707-8	3.6	
6	Primary adrenal insufficiency in children: twenty years experience at the Sainte-Justine Hospital, Montreal. *Journal of Clinical Endocrinology and Metabolism*, **2005**, 90, 3243-50	5.6	125
5	Sudden death in growth hormone-treated children with Prader-Willi syndrome. *Journal of Pediatrics* , **2004**, 144, 129-31	3.6	86
4	Screening for neonatal endocrinopathies: rationale, methods and results. *Seminars in Fetal and Neonatal Medicine*, **2004**, 9, 75-85		36
3	Nine novel mutations in NR0B1 (DAX1) causing adrenal hypoplasia congenita. *Human Mutation*, **2001**, 18, 547	4.7	11
2	A search for the possible molecular mechanisms of thyroid dysgenesis: sex ratios and associated malformations. *Journal of Clinical Endocrinology and Metabolism*, **1999**, 84, 2502-6	5.6	145
1	A Search for the Possible Molecular Mechanisms of Thyroid Dysgenesis: Sex Ratios and Associated Malformations		47