

# Guy Van Vliet

## List of Publications by Citations

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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	A search for the possible molecular mechanisms of thyroid dysgenesis: sex ratios and associated malformations. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>1999</b> , 84, 2502-6	5.6	145
26	Is the incidence of congenital hypothyroidism really increasing? A 20-year retrospective population-based study in QuÉbec. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2011</b> , 96, 2422-9	5.6	143
25	European Society for Paediatric Endocrinology consensus guidelines on screening, diagnosis, and management of congenital hypothyroidism. <i>Hormone Research in Paediatrics</i> , <b>2014</b> , 81, 80-103	3.3	136
24	Prevention of intellectual disability through screening for congenital hypothyroidism: how much and at what level?. <i>Archives of Disease in Childhood</i> , <b>2011</b> , 96, 374-9	2.2	134
23	Primary adrenal insufficiency in children: twenty years experience at the Sainte-Justine Hospital, Montreal. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2005</b> , 90, 3243-50	5.6	125
22	Sudden death in growth hormone-treated children with Prader-Willi syndrome. <i>Journal of Pediatrics</i> , <b>2004</b> , 144, 129-31	3.6	86
21	Mutations in NFKB2 and potential genetic heterogeneity in patients with DAVID syndrome, having variable endocrine and immune deficiencies. <i>BMC Medical Genetics</i> , <b>2014</b> , 15, 139	2.1	56
20	Are lower TSH cutoffs in neonatal screening for congenital hypothyroidism warranted?. <i>European Journal of Endocrinology</i> , <b>2017</b> , 177, D1-D12	6.5	51
19	How many deaths can be prevented by newborn screening for congenital adrenal hyperplasia?. <i>Hormone Research in Paediatrics</i> , <b>2007</b> , 67, 284-91	3.3	45
18	Screening for neonatal endocrinopathies: rationale, methods and results. <i>Seminars in Fetal and Neonatal Medicine</i> , <b>2004</b> , 9, 75-85		36
17	A Search for the Possible Molecular Mechanisms of Thyroid Dysgenesis: Sex Ratios and Associated Malformations		17
16	Treating fetal thyroid and adrenal disorders through the mother. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , <b>2008</b> , 4, 675-82		18
15	Non-immune goiter and hypothyroidism in a 19-week fetus: a plea for conservative treatment. <i>Journal of Pediatrics</i> , <b>2010</b> , 156, 1026-1029	3.6	17
14	Nine novel mutations in NR0B1 (DAX1) causing adrenal hypoplasia congenita. <i>Human Mutation</i> , <b>2001</b> , 18, 547	4.7	11
13	Wide Spectrum of Deficiency: From Life-Threatening Compressive Goiter in Infancy to Lifelong Euthyroidism. <i>Thyroid</i> , <b>2019</b> , 29, 1018-1022	6.2	10
12	Demonstration of Autosomal Monoallelic Expression in Thyroid Tissue Assessed by Whole-Exome and Bulk RNA Sequencing. <i>Thyroid</i> , <b>2016</b> , 26, 852-9	6.2	10
11	Interpreting Minor Variations in Thyroid Function or Echostructure: Treating Patients, Not Numbers or Images. <i>Pediatric Clinics of North America</i> , <b>2015</b> , 62, 929-42	3.6	5

10	Avoiding the Overdiagnosis of Congenital Hypothyroidism in Premature Newborns. <i>Pediatrics</i> , <b>2019</b> , 144,	7.4	4
9	Similar age-dependent levothyroxine requirements of schoolchildren with congenital or acquired hypothyroidism. <i>European Journal of Pediatrics</i> , <b>2016</b> , 175, 869-72	4.1	3
8	Whole-exome sequencing: opportunities in pediatric endocrinology. <i>Personalized Medicine</i> , <b>2014</b> , 11, 63-78	2.2	2
7	Quality of referral of short children to the paediatric endocrinologist and impact of a fax communication system. <i>Paediatrics and Child Health</i> , <b>2013</b> , 18, 533-7	0.7	2
6	Challenges in Assessing the Cost-Effectiveness of Newborn Screening: The Example of Congenital Adrenal Hyperplasia. <i>International Journal of Neonatal Screening</i> , <b>2020</b> , 6,	2.6	2
5	Growth Hormone Supplementation and Psychosocial Functioning to Adult Height in Turner Syndrome: A Questionnaire Study of Participants in the Canadian Randomized Trial. <i>Frontiers in Endocrinology</i> , <b>2019</b> , 10, 125	5.7	1
4	Severe Congenital Hypothyroidism Due to a Novel Deep Intronic Mutation in the TSH Receptor Gene Causing Intron Retention. <i>Journal of the Endocrine Society</i> , <b>2021</b> , 5, bvaa183	0.4	1
3	Redefining Congenital Hypothyroidism?. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2021</b> , 106, e1463-e1465	5.6	0
2	Conserved Telomere Length in Human Ectopic Thyroids: An Argument Against Premature Differentiation Causing Arrested Migration. <i>Thyroid</i> , <b>2015</b> , 25, 1050-4	6.2	
1	Longitudinal study of thyroid function in Down's syndrome in the first two decades. <i>Journal of Pediatrics</i> , <b>2005</b> , 147, 707-8	3.6	