

Oliver Blankenstein

List of Publications by Year in descending order

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Version: 2024-02-01

67
papers

2,414
citations

236912

25
h-index

206102

48
g-index

72
all docs

72
docs citations

72
times ranked

3107
citing authors

#	ARTICLE	IF	CITATIONS
1	Proopiomelanocortin Deficiency Treated with a Melanocortin-4 Receptor Agonist. <i>New England Journal of Medicine</i> , 2016, 375, 240-246.	27.0	358
2	MC4R agonism promotes durable weight loss in patients with leptin receptor deficiency. <i>Nature Medicine</i> , 2018, 24, 551-555.	30.7	219
3	Type 2 diabetes and impaired glucose tolerance in European children and adolescents with obesity – a problem that is no longer restricted to minority groups. <i>European Journal of Endocrinology</i> , 2004, 151, 199-206.	3.7	174
4	Somatostatin or octreotide as treatment options for chylothorax in young children: a systematic review. <i>Intensive Care Medicine</i> , 2006, 32, 650-657.	8.2	134
5	Somatic mutations and progressive monosomy modify SAMD9-related phenotypes in humans. <i>Journal of Clinical Investigation</i> , 2017, 127, 1700-1713.	8.2	129
6	Congenital hyperinsulinism. <i>Early Human Development</i> , 2010, 86, 287-294.	1.8	105
7	Evaluation of [¹⁸ F]Fluoro-DOPA Positron Emission Tomography-Computed Tomography for Surgery in Focal Congenital Hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008, 93, 869-875.	3.6	88
8	Hyperinsulinaemic hypoglycaemia: biochemical basis and the importance of maintaining normoglycaemia during management. <i>Archives of Disease in Childhood</i> , 2007, 92, 568-570.	1.9	75
9	Molecular characterization of the cfb gene encoding group B streptococcal CAMP-factor. <i>Medical Microbiology and Immunology</i> , 1994, 183, 239-256.	4.8	61
10	Treating patients not numbers: the benefit and burden of lowering TSH newborn screening cut-offs. <i>Archives of Disease in Childhood</i> , 2011, 96, 121-122.	1.9	57
11	Role of ⁶⁸ Ga somatostatin receptor PET/CT in the detection of endogenous hyperinsulinaemic focus: an explorative study. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2016, 43, 1593-1600.	6.4	48
12	The predictive value of preoperative fluorine-18-l-3,4-dihydroxyphenylalanine positron emission tomography-computed tomography scans in children with congenital hyperinsulinism of infancy. <i>Journal of Pediatric Surgery</i> , 2011, 46, 204-208.	1.6	47
13	Mean High-Dose l-Thyroxine Treatment Is Efficient and Safe to Achieve a Normal IQ in Young Adult Patients With Congenital Hypothyroidism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 1459-1469.	3.6	47
14	Absorption and tolerability of taste-masked hydrocortisone granules in neonates, infants and children under 6 years of age with adrenal insufficiency. <i>Clinical Endocrinology</i> , 2018, 88, 21-29.	2.4	46
15	Harmonization of growth hormone measurements with different immunoassays by data adjustment. <i>Clinical Chemistry and Laboratory Medicine</i> , 2011, 49, 1135-42.	2.3	40
16	Diabetes caused by insulin gene (INS) deletion: clinical characteristics of homozygous and heterozygous individuals. <i>European Journal of Endocrinology</i> , 2011, 165, 255-260.	3.7	38
17	Long-Term Lanreotide Treatment in Six Patients with Congenital Hyperinsulinism. <i>Hormone Research in Paediatrics</i> , 2012, 78, 106-112.	1.8	38
18	Comparison of response to 2-years' growth hormone treatment in children with isolated growth hormone deficiency, born small for gestational age, idiopathic short stature, or multiple pituitary hormone deficiency: combined results from two large observational studies. <i>International Journal of Pediatric Endocrinology (Springer)</i> , 2012, 2012, 22.	1.6	38

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19	Does the aromatic l-amino acid decarboxylase contribute to thyronamine biosynthesis?. <i>Molecular and Cellular Endocrinology</i> , 2012, 349, 195-201.	3.2	37
20	Quality of compounded hydrocortisone capsules used in the treatment of children. <i>European Journal of Endocrinology</i> , 2017, 177, 239-242.	3.7	37
21	Development and Testing in Healthy Adults of Oral Hydrocortisone Granules With Taste Masking for the Treatment of Neonates and Infants With Adrenal Insufficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 1681-1688.	3.6	35
22	Molecular Mechanisms of Neonatal Hyperinsulinism. <i>Hormone Research in Paediatrics</i> , 2006, 66, 289-296.	1.8	34
23	Familial Focal Congenital Hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011, 96, 24-28.	3.6	31
24	Two Novel GATA6 Mutations Cause Childhood-Onset Diabetes Mellitus, Pancreas Malformation and Congenital Heart Disease. <i>Hormone Research in Paediatrics</i> , 2013, 79, 250-256.	1.8	28
25	Longitudinal Imaging Reveals Pituitary Enlargement Preceding Hypoplasia in Two Brothers with Combined Pituitary Hormone Deficiency Attributable to PROP1 Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 4353-4357.	3.6	27
26	The Heterogeneity of Focal Forms of Congenital Hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012, 97, E94-E99.	3.6	26
27	Newborn screening by tandem mass spectrometry confirms the high prevalence of sickle cell disease among German newborns. <i>Annals of Hematology</i> , 2019, 98, 47-53.	1.8	23
28	Neonatal Screening for Congenital Metabolic and Endocrine Disorders. <i>Deutsches A&#x0308;rzteblatt International</i> , 2021, 118, 101-108.	0.9	23
29	A Prospective Study of Children Aged 0â€“8 Years with CAH and Adrenal Insufficiency Treated with Hydrocortisone Granules. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, e1433-e1440.	3.6	22
30	Incidence of sickle cell disease in an unselected cohort of neonates born in Berlin, Germany. <i>European Journal of Human Genetics</i> , 2014, 22, 1051-1053.	2.8	21
31	Is safety of childhood growth hormone therapy related to dose? Data from a large observational study. <i>European Journal of Endocrinology</i> , 2016, 174, 681-691.	3.7	21
32	International practice of corticosteroid replacement therapy in congenital adrenal hyperplasia: data from the I-CAH registry. <i>European Journal of Endocrinology</i> , 2021, 184, 553-563.	3.7	21
33	Functional Implications of LH/hCG Receptors in Pregnancy-Induced Cushing Syndrome. <i>Journal of the Endocrine Society</i> , 2017, 1, 57-71.	0.2	20
34	Real-World Estimates of Adrenal Insufficiencyâ€“Related Adverse Events in Children With Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, e192-e203.	3.6	20
35	EDTA in Dried Blood Spots Leads to False Results in Neonatal Endocrinologic Screening. <i>Clinical Chemistry</i> , 2008, 54, 602-605.	3.2	17
36	Positron emission tomography/computed tomography diagnostics by means of fluorine-18-L-dihydroxyphenylalanine in congenital hyperinsulinism. <i>Seminars in Pediatric Surgery</i> , 2011, 20, 23-27.	1.1	16

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37	Newborn Screening for Sickle Cell Disease: Technical and Legal Aspects of a German Pilot Study with 38,220 Participants. <i>BioMed Research International</i> , 2014, 2014, 1-10.	1.9	16
38	Gender Influences Short-Term Growth Hormone Treatment Response in Children. <i>Hormone Research in Paediatrics</i> , 2012, 77, 188-194.	1.8	15
39	Occurrence of giant focal forms of congenital hyperinsulinism with incorrect visualization by ¹⁸ F-DOPA- ¹⁸ F-PET/CT scanning. <i>Clinical Endocrinology</i> , 2014, 81, 847-854.	2.4	14
40	Comparative meta-analysis of Kabuki syndrome with and without hyperinsulinaemic hypoglycaemia. <i>Clinical Endocrinology</i> , 2020, 93, 346-354.	2.4	14
41	Congenital disorder of glycosylation type 1a in a macrosomic 16-month-old boy with an atypical phenotype and homozygosity of the N216I mutation. <i>European Journal of Pediatrics</i> , 2003, 162, 710-713.	2.7	13
42	Visualization of the focus in congenital hyperinsulinism by intraoperative sonography. <i>Seminars in Pediatric Surgery</i> , 2011, 20, 28-31.	1.1	13
43	Real-life GH dosing patterns in children with GHD, TS or born SGA: a report from the NordiNet [®] International Outcome Study. <i>European Journal of Endocrinology</i> , 2017, 177, 145-155.	3.7	13
44	Positive correlation of thyroid hormones and serum copper in children with congenital hypothyroidism. <i>Journal of Trace Elements in Medicine and Biology</i> , 2016, 37, 90-95.	3.0	11
45	Primary sulphonylurea therapy in a newborn with transient neonatal diabetes attributable to a paternal uniparental disomy 6q24 (⁶UPD). <i>Diabetes, Obesity and Metabolism</i> , 2018, 20, 474-475.	4.4	11
46	Management and interpretation of heterogeneous observational data: Using insulin-like growth factor-I data from the NordiNet [®] International Outcome Study. <i>Growth Hormone and IGF Research</i> , 2015, 25, 41-46.	1.1	9
47	Characterization of diabetes following pancreatic surgery in patients with congenital hyperinsulinism. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 230.	2.7	9
48	Open-Source Technology for Real-Time Continuous Glucose Monitoring in the Neonatal Intensive Care Unit: Case Study in a Neonate With Transient Congenital Hyperinsulinism. <i>Journal of Medical Internet Research</i> , 2020, 22, e21770.	4.3	9
49	How to improve information extraction from German medical records. <i>IT - Information Technology</i> , 2017, 59, 171-179.	0.9	8
50	Sleep Timing in Patients with Precocious and Delayed Pubertal Development. <i>Clocks & Sleep</i> , 2019, 1, 140-150.	2.0	8
51	Rationale of a lower dexamethasone dose in prenatal congenital adrenal hyperplasia therapy based on pharmacokinetic modelling. <i>European Journal of Endocrinology</i> , 2021, 185, 365-374.	3.7	8
52	Treatment of congenital adrenal hyperplasia in children aged 0-3 years: a retrospective multicenter analysis of salt supplementation, glucocorticoid and mineralocorticoid medication, growth and blood pressure. <i>European Journal of Endocrinology</i> , 2022, 186, 587-596.	3.7	7
53	Idiopathic Growth Hormone Deficiency: A Vanishing Diagnosis?. <i>Hormone Research in Paediatrics</i> , 2000, 53, 1-8.	1.8	6
54	Height Gain with Combined Growth Hormone and Gonadotropin-Releasing Hormone Analog Therapy in Two Pubertal Siblings with a Growth Hormone-Releasing Hormone Receptor Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008, 93, 204-207.	3.6	6

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55	Normative Thyroid-Stimulating Hormone Values for Healthy Nigerian Newborns. <i>Hormone Research in Paediatrics</i> , 2016, 85, 22-28.	1.8	6
56	Severe Acquired Primary Hypothyroidism in Children and its Influence on Growth: A Retrospective Analysis of 43 Cases. <i>Experimental and Clinical Endocrinology and Diabetes</i> , 2022, 130, 217-222.	1.2	3
57	Towards the Development of Smart and Reliable Health Assistance Networks Exemplified by an Apnea Detection System. , 2014, , .		2
58	Hydrocortisone Replacement in Disorders of Sex Development. <i>Endocrine Development</i> , 2014, 27, 160-171.	1.3	2
59	Is safety of childhood growth hormone therapy related to dose? Data from a large observational study. <i>European Journal of Endocrinology</i> , 2017, 176, X1.	3.7	2
60	Incidence of Daytime Sleepiness and Associated Factors in Two First Nations Communities in Saskatchewan, Canada. <i>Clocks & Sleep</i> , 2018, 1, 13-25.	2.0	2
61	Exploring Dried Blood Spot Cortisol Concentrations as an Alternative for Monitoring Pediatric Adrenal Insufficiency Patients: A Model-Based Analysis. <i>Frontiers in Pharmacology</i> , 2022, 13, 819590.	3.5	2
62	Congenital Adrenal Hyperplasia with Non-functional Mutations in Both Alleles in a Clinically Unaffected Infant. <i>Journal of Tropical Pediatrics</i> , 2016, 62, 158-160.	1.5	0
63	Hyperinsulinismus. <i>Springer Reference Medizin</i> , 2018, , 1-10.	0.0	0
64	Neuroendokrine Tumoren. , 2011, , 893-944.		0
65	Neuroendokrine Tumoren. , 2016, , 465-505.		0
66	Hyperinsulinismus. <i>Springer Reference Medizin</i> , 2020, , 149-158.	0.0	0
67	Simultaneous newborn screening for sickle cell disease, biotinidase deficiency, and hereditary tyrosinemia type 1 with an optimized tandem mass spectrometry protocol. <i>Annals of Hematology</i> , 2022, , .	1.8	0