

Jeremy Huw Jones

List of Publications by Year in descending order

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Version: 2024-02-01

13
papers

123
citations

1307594

7
h-index

1281871

11
g-index

13
all docs

13
docs citations

13
times ranked

199
citing authors

#	ARTICLE	IF	CITATIONS
1	Ectopic parathyroid hormone as a rare aetiology of hypercalcemia with rhabdomyosarcoma: a new treatment strategy with zoledronic acid and Denosumab. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2022, 35, 1107-1112.	0.9	1
2	Determination of thyroid volume in infants with suspected congenital hypothyroidism—the limitations of both subjective and objective evaluation. <i>BJR Open</i> , 2020, 2, 20200001.	0.6	1
3	Measured parental height in Turner syndrome—a valuable but underused diagnostic tool. <i>European Journal of Pediatrics</i> , 2018, 177, 171-179.	2.7	8
4	Neonatal Features of the Prader-Willi Syndrome; The Case for Making the Diagnosis During the First Week of Life. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2018, 10, 264-273.	0.9	16
5	Trends in Scottish newborn screening programme for congenital hypothyroidism 1980–2014: strategies for reducing age at notification after initial and repeat sampling. <i>Archives of Disease in Childhood</i> , 2017, 102, 936-941.	1.9	7
6	A further case of brain-lung-thyroid syndrome with deletion proximal to NKX2-1. <i>European Journal of Medical Genetics</i> , 2017, 60, 257-260.	1.3	13
7	How well does the capillary thyroid-stimulating hormone test for newborn thyroid screening predict the venous free thyroxine level?. <i>Archives of Disease in Childhood</i> , 2016, 101, 539-545.	1.9	11
8	Screening for Hypothyroidism in Down Syndrome Using the Capillary Thyroid Stimulating Hormone Method. <i>Journal of Pediatrics</i> , 2015, 166, 1013-1017.e2.	1.8	5
9	Diagnostic and Predictive Value of Ultrasound and Isotope Thyroid Scanning, Alone and in Combination, in Infants Referred with Thyroid-Stimulating Hormone Elevation on Newborn Screening. <i>Journal of Pediatrics</i> , 2014, 164, 846-854.	1.8	17
10	Novel heterozygous thyrotropin receptor mutation presenting with neonatal hyperthyrotropinaemia, mild thyroid hypoplasia and absent uptake on radioisotope scan. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2013, 26, 583-6.	0.9	5
11	Optimising outcome in congenital hypothyroidism; current opinions on best practice in initial assessment and subsequent management. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2012, 4, 13-22.	0.9	26
12	Visible thyroid ectopia. <i>Archives of Disease in Childhood: Fetal and Neonatal Edition</i> , 2012, 97, F482-F483.	2.8	1
13	Capillary TSH screening programme for Down's syndrome in Scotland, 1997-2009. <i>Archives of Disease in Childhood</i> , 2011, 96, 1113-1117.	1.9	12