Laura Lane

List of Publications by Year in descending order

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1478505 1372567 16 194 6 10 citations h-index g-index papers 16 16 16 238 citing authors all docs docs citations times ranked

#	Article	IF	CITATIONS
1	Graves' disease: moving forwards. Archives of Disease in Childhood, 2023, 108, 276-281.	1.9	6
2	A survey of the young person's experience of Graves' disease and its management. Clinical Endocrinology, 2021, 94, 330-340.	2.4	8
3	New Therapeutic Horizons for Graves' Hyperthyroidism. Endocrine Reviews, 2020, 41, 873-884.	20.1	56
4	An Intronic HCP5 Variant Is Associated With Age of Onset and Susceptibility to Graves Disease in UK and Polish Cohorts. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3277-e3284.	3.6	12
5	Graves' disease: developments in first-line antithyroid drugs in the young. Expert Review of Endocrinology and Metabolism, 2020, 15, 59-69.	2.4	3
6	Puberty: Normal physiology (brief overview). Best Practice and Research in Clinical Endocrinology and Metabolism, 2019, 33, 101265.	4.7	74
7	Analysis of BAFF gene polymorphisms in UK Graves' disease patients. Clinical Endocrinology, 2019, 90, 170-174.	2.4	19
8	Adult height in patients with familial male-limited precocious puberty and the role of an aromatase inhibitor in patient management. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 551-560.	0.9	12
9	Graves' disease. Time to move on. Archives of Disease in Childhood, 2018, 103, archdischild-2017-314486.	1.9	3
10	Who needs thyroid function testing at birth?. Archives of Disease in Childhood, 2017, 102, 212-215.	1.9	1
11	Novel Immunomodulatory Treatment Modalities. , 2017, , 177-192.		0
12	Congenital hypothyroidism – what's new?. Paediatrics and Child Health (United Kingdom), 2015, 25, 302-307.	0.4	0
13	An audit of paediatric obesity in secondary care. Endocrine Abstracts, 0, , .	0.0	0
14	Association of a promoter BAFF polymorphism in Graves' disease. Endocrine Abstracts, 0, , .	0.0	0
15	Adult height in patients with testotoxicosis. Endocrine Abstracts, 0, , .	0.0	0
16	The SH2B3 tryptophan 262 variant is associated with Graves' disease and Addison's disease. Endocrine Abstracts, 0, , .	0.0	0