CÃ©cile Brachet

List of Publications by Year
in descending order

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25	821	687335	580810
papers	citations	13 h-index	25 g-index

27	27	27	1265
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Pubertal induction and transition to adult sex hormone replacement in patients with congenital pituitary or gonadal reproductive hormone deficiency: an Endo-ERN clinical practice guideline. European Journal of Endocrinology, 2022, 186, G9-G49.	3.7	25
2	SRYâ€negative 46,XX testicular/ovotesticular DSD: Longâ€term outcomes and early blockade of gonadotropic axis. Clinical Endocrinology, 2021, 94, 667-676.	2.4	10
3	The Retina in Patients With Triple A Syndrome: A Window Into Neurodegeneration?. Frontiers in Endocrinology, 2021, 12, 729056.	3.5	0
4	Increasing knowledge in IGF1R defects: lessons from 35 new patients. Journal of Medical Genetics, 2020, 57, 160-168.	3.2	20
5	Delayed diagnosis of congenital hypopituitarism associated with low socio-economic status and/or migration. European Journal of Pediatrics, 2020, 179, 151-155.	2.7	2
6	A novel approach in the intraoperative management of ovotesticular DSD. Journal of Pediatric Urology, 2020, 16, 768-770.	1.1	4
7	Homozygous p.R31H GNRH1 mutation and normosmic congenital hypogonadotropic hypogonadism in a patient and self-limited delayed puberty in his relatives. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1237-1240.	0.9	5
8	Contribution of functionally assessed GHRHR mutations to idiopathic isolated growth hormone deficiency in patients without GH1 mutations. Human Mutation, 2019, 40, 2033-2043.	2.5	9
9	Truncating RAX Mutations: Anophthalmia, Hypopituitarism, Diabetes Insipidus, and Cleft Palate in Mice and Men. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2925-2930.	3.6	10
10	Adult Height after Growth Hormone Treatment at Pubertal Onset in Short Adolescents Born Small for Gestational Age: Results from a Belgian Registry-Based Study. International Journal of Endocrinology, 2018, 2018, 1-8.	1.5	12
11	Congenital Hypothyroidism: Long-Term Experience with Early and High Levothyroxine Dosage. Hormone Research in Paediatrics, 2016, 85, 188-197.	1.8	11
12	Isolated and Syndromic Retinal Dystrophy Caused by Biallelic Mutations in RCBTB1 , a Gene Implicated in Ubiquitination. American Journal of Human Genetics, 2016, 99, 470-480.	6.2	39
13	Live birth after autograft of ovarian tissue cryopreserved during childhood: Figure 1. Human Reproduction, 2015, 30, 2107-2109.	0.9	310
14	Nephrogenic Syndrome of Inappropriate Antidiuresis in a Female Neonate: Review of the Clinical Presentation in Females. Hormone Research in Paediatrics, 2015, 84, 65-67.	1.8	8
15	Complex Tissue-Specific Epigenotypes in Russell-Silver Syndrome Associated with 11p15 ICR1 Hypomethylation. Human Mutation, 2014, 35, 1211-1220.	2.5	34
16	Hearing Loss Is Part of the Clinical Picture of ENPP1 Loss of Function Mutation. Hormone Research in Paediatrics, 2014, 81, 63-66.	1.8	22
17	Haematopoietic stem cell transplantation for severe sickle cell disease in childhood: a single centre experience of 50 patients. British Journal of Haematology, 2014, 165, 402-408.	2.5	63
18	Symptomatic Heterozygotes and Prenatal Diagnoses in a Nonconsanguineous Family with Syndromic Combined Pituitary Hormone Deficiency Resulting from Two NovelLHX3Mutations. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E503-E509.	3.6	25

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19	Long-Term Treatment of Hyponatremic Patients with Nephrogenic Syndrome of Inappropriate Antidiuresis: Personal Experience and Review of Published Case Reports. Nephron Clinical Practice, 2012, 120, c168-c172.	2.3	35
20	Central precocious puberty after interpersonal transfer of testosterone gel: just a coincidence?. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 757-60.	0.9	13
21	Growth retardation in untreated autosomal dominant familial neurohypophyseal diabetes insipidus caused by one recurring and two novel mutations in the vasopressin-neurophysin II gene. European Journal of Endocrinology, 2011, 164, 179-187.	3.7	25
22	Hematopoietic Stem Cell Transplantation for Sickle Cell Disease in Childhood: A Single Center Experience with 45 Patients. Blood, 2011, 118, 3103-3103.	1.4	2
23	Association of parathyroid adenoma and familial hypocalciuric hypercalcaemia in a teenager. European Journal of Endocrinology, 2009, 161, 207-210.	3.7	42
24	Children With Sickle Cell Disease: Growth and Gonadal Function After Hematopoietic Stem Cell Transplantation. Journal of Pediatric Hematology/Oncology, 2007, 29, 445-450.	0.6	49
25	Childrenâ€™s virilization and the use of a testosterone gel by their fathers. European Journal of Pediatrics, 2005, 164, 646-647.	2.7	43