S Faisal Ahmed

List of Publications by Year
in descending order

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154 papers	7,067 citations	87888 38 h-index	64796 79 g-index
154 all docs	154 docs citations	154 times ranked	5683 citing authors

#	Article	IF	Citations
1	I-DSD: The First 10 Years. Hormone Research in Paediatrics, 2023, 96, 238-246.	1.8	3
2	Congenital Adrenal Hyperplasiaâ€"Current Insights in Pathophysiology, Diagnostics, and Management. Endocrine Reviews, 2022, 43, 91-159.	20.1	182
3	A Nationwide Study of the Prevalence and Initial Management of Atypical Genitalia in the Newborn in Scotland. Sexual Development, 2022, 16, 11-18.	2.0	6
4	Elective hip arthroplasty rates and related complications in people with diabetes mellitus. HIP International, 2022, 32, 717-723.	1.7	3
5	Growth, puberty and testicular function in boys born small for gestational age with a nonspecific disorder of sex development. Clinical Endocrinology, 2022, 96, 165-174.	2.4	6
6	Update on the management of a newborn with a suspected difference of sex development. Archives of Disease in Childhood, 2022, 107, 866-871.	1.9	2
7	Congenital Micropenis: Etiology And Management. Journal of the Endocrine Society, 2022, 6, bvab172.	0.2	7
8	Treatment of congenital adrenal hyperplasia in children aged 0â€"3 years: a retrospective multicenter analysis of salt supplementation, glucocorticoid and mineralocorticoid medication, growth and blood pressure. European Journal of Endocrinology, 2022, 186, 587-596.	3.7	7
9	Vascular dysfunction and increased cardiovascular risk in hypospadias. European Heart Journal, 2022, 43, 1832-1845.	2.2	16
10	Treatment Adherence to Injectable Treatments in Pediatric Growth Hormone Deficiency Compared With Injectable Treatments in Other Chronic Pediatric Conditions: A Systematic Literature Review. Frontiers in Endocrinology, 2022, 13, 795224.	3.5	7
11	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
12	Prenatal dexamethasone treatment for classic 21-hydroxylase deficiency in Europe. European Journal of Endocrinology, 2022, 186, K17-K24.	3.7	7
13	Combined growth hormone and insulin-like growth factor-1 rescues growth retardation in glucocorticoid-treated mdxmice but does not prevent osteopenia. Journal of Endocrinology, 2022, 253, 63-74.	2.6	5
14	Differences or Disorders of sex development in Boys: impact on fertility. Annales D'Endocrinologie, 2022, , .	1.4	0
15	Delivery of multidisciplinary care in the field of differences and disorders of sex development (DSD). Expert Review of Endocrinology and Metabolism, 2022, 17, 225-234.	2.4	4
16	Management of Acute Adrenal Insufficiency-Related Adverse Events in Children with Congenital Adrenal Hyperplasia: Results of an International Survey of Specialist Centres. Hormone Research in Paediatrics, 2022, 95, 363-373.	1.8	2
17	The Use of Genetics for Reaching a Diagnosis in XY DSD. Sexual Development, 2022, 16, 207-224.	2.0	5
18	3T MRI-based age, sex and site-specific markers of musculoskeletal health in healthy children and young adults. Endocrine Connections, 2022, , .	1.9	1

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19	Analysis of therapy monitoring in the International Congenital Adrenal Hyperplasia Registry. Clinical Endocrinology, 2022, 97, 551-561.	2.4	4
20	A survey of the feasibility of developing osteoporosis clinical trials in Duchenne muscular dystrophy: Survey of the opinion of young people with Duchenne muscular dystrophy, families and clinicians. Clinical Trials, 2021, 18, 39-50.	1.6	0
21	Real-World Estimates of Adrenal Insufficiencyâ€“Related Adverse Events in Children With Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e192-e203.	3.6	20
22	Assessing the health-related management of people with differences of sex development. Endocrine, 2021, 71, 675-680.	2.3	9
23	Supporting international networks through platforms for standardised data collectionâ€”the European Registries for Rare Endocrine Conditions (EuRRECa) model. Endocrine, 2021, 71, 555-560.	2.3	9
24	Surgical Practice in Girls with Congenital Adrenal Hyperplasia: An International Registry Study. Sexual Development, 2021, 15, 229-235.	2.0	4
25	Testosterone Therapy and Its Monitoring in Adolescent Boys with Hypogonadism: Results of an International Survey from the I-DSD Registry. Sexual Development, 2021, 15, 236-243.	2.0	4
26	Experience of health care at a reference centre as reported by patients and parents of children with rare conditions. Orphanet Journal of Rare Diseases, 2021, 16, 65.	2.7	3
27	CPMSâ€“improving patient care in Europe via virtual case discussions. Endocrine, 2021, 71, 549-554.	2.3	13
28	Outcome Squares Integrating Efficacy and Safety, as Applied to Functioning Pituitary Adenoma Surgery. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3300-e3311.	3.6	7
29	Therapy options for adrenal insufficiency and recommendations for the management of adrenal crisis. Endocrine, 2021, 71, 586-594.	2.3	31
30	International practice of corticosteroid replacement therapy in congenital adrenal hyperplasia: data from the I-CAH registry. European Journal of Endocrinology, 2021, 184, 553-563.	3.7	21
31	Gonadectomy in conditions affecting sex development: a registry-based cohort study. European Journal of Endocrinology, 2021, 184, 791-801.	3.7	9
32	Society for Endocrinology UK Guidance on the initial evaluation of a suspected difference or disorder of sex development (Revised 2021). Clinical Endocrinology, 2021, 95, 818-840.	2.4	29
33	Observer Agreement of Vertebral Fracture Grading Using Dual Energy Absorptiometry Vertebral Fracture Assessment in Duchenne Muscular Dystrophy. Journal of Clinical Densitometry, 2021, 24, 622-629.	1.2	0
34	The Quality Evaluation of Rare Disease Registriesâ€”An Assessment of the Essential Features of a Disease Registry. International Journal of Environmental Research and Public Health, 2021, 18, 11968.	2.6	11
35	A comparison of the bone and growth phenotype of mdx, mdx:cmahâ^'/â^' and mdx:utrn+/â^' murine models with the C57BL10 wildtype mouse. DMM Disease Models and Mechanisms, 2020, 13, .	2.4	7
36	Plasma Renin Measurements are Unrelated to Mineralocorticoid Replacement Dose in Patients With Primary Adrenal Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 314-326.	3.6	30

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37	The External Genitalia Score (EGS): A European Multicenter Validation Study. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e222-e230.	3.6	51
38	The EuRRECa Project as a Model for Data Access and Governance Policies for Rare Disease Registries That Collect Clinical Outcomes. International Journal of Environmental Research and Public Health, 2020, 17, 8743.	2.6	13
39	Recommendations for Diagnosis and Treatment of Pseudohypoparathyroidism and Related Disorders: An Updated Practical Tool for Physicians and Patients. Hormone Research in Paediatrics, 2020, 93, 182-196.	1.8	42
40	Muscle deficits with normal bone microarchitecture and geometry in young adults with well-controlled childhood-onset Crohnâ€™s disease. European Journal of Gastroenterology and Hepatology, 2020, 32, 1497-1506.	1.6	7
41	Serum Anti-MÃ¼llerian Hormone in the Prediction of Response to hCG Stimulation in Children With DSD. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1608-1616.	3.6	14
42	Parent-reported outcomes in young children with disorders/differences of sex development. International Journal of Pediatric Endocrinology (Springer), 2020, 2020, 3.	1.6	2
43	Peptide hormone analysis in diagnosis and treatment of Differences of Sex Development: joint position paper of EU COST Action â€˜DSDnetâ€™ and European Reference Network on Rare Endocrine Conditions. European Journal of Endocrinology, 2020, 182, P1-P15.	3.7	20
44	An overview of clinical activities in Endo-ERN: the need for alignment of future network criteria. European Journal of Endocrinology, 2020, 183, 141-148.	3.7	9
45	MON-170 Real World Estimates of Adrenal Insufficiency Related Adverse Events in Children with Congenital Adrenal Hyperplasia: On Behalf of the I-CAH Consortium. Journal of the Endocrine Society, 2020, 4, .	0.2	1
46	Peer Review of Paediatric Endocrine Services in the UK: A Template for Quality and Service Improvement. Hormone Research in Paediatrics, 2020, 93, 616-621.	1.8	0
47	Addressing gaps in care of people with conditions affecting sex development and maturation. Nature Reviews Endocrinology, 2019, 15, 615-622.	9.6	30
48	Measurement of Salivary Adrenal-Specific Androgens as Biomarkers of Therapy Control in 21-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 6417-6429.	3.6	31
49	Skeletal Fragility and Its Clinical Determinants in Children With Type 1 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3585-3594.	3.6	29
50	The Role of International Databases in Understanding the Aetiology and Consequences of Differences/Disorders of Sex Development. International Journal of Molecular Sciences, 2019, 20, 4405.	4.1	23
51	Clinical but Not Histological Outcomes in Males With 45,X/46,XY Mosaicism Vary Depending on Reason for Diagnosis. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4366-4381.	3.6	27
52	Association Between Extra-Genital Congenital Anomalies and Hypospadias Outcome. Sexual Development, 2019, 13, 67-73.	2.0	10
53	Response to Letter to the Editor: â€œClinical but Not Histological Outcomes in Males With 45,X/46,XY Mosaicism Vary Depending on Reason for Diagnosisâ€• Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5812-5813.	3.6	0
54	Cardiac disorders and structural brain abnormalities are commonly associated with hypospadias in children with neurodevelopmental disorders. Clinical Dysmorphology, 2019, 28, 112-117.	0.3	5

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55	Testosterone Therapy in Adolescent Boys: The Need for a Structured Approach. Hormone Research in Paediatrics, 2019, 92, 215-228.	1.8	34
56	Proteomic Evidence of Biological Aging in a Child with a Compound Heterozygous ZMPSTE24 Mutation. Proteomics - Clinical Applications, 2019, 13, 1800135.	1.6	8
57	The current landscape of European registries for rare endocrine conditions. European Journal of Endocrinology, 2019, 180, 89-98.	3.7	25
58	Standardised data collection for clinical follow-up and assessment of outcomes in differences of sex development (DSD): recommendations from the COST action DSDnet. European Journal of Endocrinology, 2019, 181, 545-564.	3.7	21
59	Suppressor of cytokine signaling 2 (SOCS2) deletion protects bone health of mice with DSS induced inflammatory bowel disease. DMM Disease Models and Mechanisms, 2018, 11, .	2.4	15
60	Long-Term Skeletal Disproportion in Childhood-Onset Crohnâ€™s Disease. Hormone Research in Paediatrics, 2018, 89, 132-135.	1.8	5
61	Longitudinal changes in bone parameters in young girls with anorexia nervosa. Bone, 2018, 116, 22-27.	2.9	10
62	Bone protective agents in children. Archives of Disease in Childhood, 2018, 103, 503-508.	1.9	8
63	Animal models to explore the effects of glucocorticoids on skeletal growth and structure. Journal of Endocrinology, 2018, 236, R69-R91.	2.6	38
64	Childhood-onset growth hormone deficiency and the transition to adulthood: current perspective. Therapeutics and Clinical Risk Management, 2018, Volume 14, 2283-2291.	2.0	14
65	Management of Gonads in Adults with Androgen Insensitivity: An International Survey. Hormone Research in Paediatrics, 2018, 90, 236-246.	1.8	34
66	Genetic testing of XY newborns with a suspected disorder of sex development. Current Opinion in Pediatrics, 2018, 30, 548-557.	2.0	9
67	Involving Individuals with Disorders of Sex Development and Their Parents in Exploring New Models of Shared Learning: Proceedings from a DSDnet COST Action Workshop. Sexual Development, 2018, 12, 225-231.	2.0	13
68	Diagnosis and management of pseudohypoparathyroidism and related disorders: first international Consensus Statement. Nature Reviews Endocrinology, 2018, 14, 476-500.	9.6	224
69	GENETICS IN ENDOCRINOLOGY: Approaches to molecular genetic diagnosis in the management of differences/disorders of sex development (DSD): position paper of EU COST Action BM 1303 â€˜DSDnetâ€™. European Journal of Endocrinology, 2018, 179, R197-R206.	3.7	105
70	Effects of Recombinant Human Growth Hormone in Children with Crohnâ€™s Disease on the Muscle-Bone Unit: A Preliminary Study. Hormone Research in Paediatrics, 2018, 90, 128-131.	1.8	2
71	Recommendations for Improving the Quality of Rare Disease Registries. International Journal of Environmental Research and Public Health, 2018, 15, 1644.	2.6	116
72	The evaluation and management of the boy with DSD. Best Practice and Research in Clinical Endocrinology and Metabolism, 2018, 32, 445-453.	4.7	4

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73	Integrating clinical and genetic approaches in the diagnosis of 46,XY disorders of sex development. Endocrine Connections, 2018, 7, 1480-1490.	1.9	18
74	De novo mutations in SMCHD1 cause Bosma arhinia microphthalmia syndrome and abrogate nasal development. Nature Genetics, 2017, 49, 249-255.	21.4	88
75	Use of Cortisol and Adrenal Weight at Pediatric Postmortem. Fetal and Pediatric Pathology, 2017, 36, 246-255.	0.7	0
76	A retrospective analysis of longitudinal changes in bone mineral content in cystic fibrosis. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 807-814.	0.9	9
77	Prevalence of endocrine and genetic abnormalities in boys evaluated systematically for a disorder of sex development. Human Reproduction, 2017, 32, 2130-2137.	0.9	30
78	Androgen-responsive non-coding small RNAs extend the potential of HCG stimulation to act as a bioassay of androgen sufficiency. European Journal of Endocrinology, 2017, 177, 339-346.	3.7	6
79	Data Quality in Rare Diseases Registries. Advances in Experimental Medicine and Biology, 2017, 1031, 149-164.	1.6	56
80	Genomic and non-genomic effects of androgens in the cardiovascular system: clinical implications. Clinical Science, 2017, 131, 1405-1418.	4.3	91
81	The relationship between adiposity, bone density and microarchitecture is maintained in young women irrespective of diabetes status. Clinical Endocrinology, 2017, 87, 327-335.	2.4	15
82	An assessment of the quality of the I-DSD and the I-CAH registries - international registries for rare conditions affecting sex development. Orphanet Journal of Rare Diseases, 2017, 12, 56.	2.7	31
83	Metformin suppresses adipogenesis through both AMP-activated protein kinase (AMPK)-dependent and AMPK-independent mechanisms. Molecular and Cellular Endocrinology, 2017, 440, 57-68.	3.2	105
84	The current state of diagnostic genetics for conditions affecting sex development. Clinical Genetics, 2017, 91, 157-162.	2.0	36
85	Amalgamated Reference Data for Size-Adjusted Bone Densitometry Measurements in 3598 Children and Young Adultsâ€"the ALPHABET Study. Journal of Bone and Mineral Research, 2017, 32, 172-180.	2.8	98
86	Shorter anogenital and anoscrotal distances correlate with the severity of hypospadias: A prospective study. Journal of Pediatric Urology, 2017, 13, 57.e1-57.e5.	1.1	33
87	Understanding the needs of professionals who provide psychosocial care for children and adults with disorders of sex development. BMJ Paediatrics Open, 2017, 1, e000132.	1.4	19
88	The measurement of urinary gonadotropins for assessment and management of pubertal disorder. Hormones, 2016, 15, 377-384.	1.9	12
89	Frequency and aetiology of hypercalcaemia. Archives of Disease in Childhood, 2016, 101, 344-347.	1.9	19
90	Society for Endocrinology <scp>UK</scp> guidance on the initial evaluation of an infant or an adolescent with a suspected disorder of sex development (Revised 2015). Clinical Endocrinology, 2016, 84, 771-788.	2.4	196

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91	Growth hormone deficiency during young adulthood and the benefits of growth hormone replacement. Endocrine Connections, 2016, 5, R1-R11.	1.9	33
92	Identification of an AR Mutation-Negative Class of Androgen Insensitivity by Determining Endogenous AR Activity. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 4468-4477.	3.6	64
93	Response to the Council of Europe Human Rights Commissioner's Issue Paper on Human Rights and Intersex People. European Urology, 2016, 70, 407-409.	1.9	35
94	Prevalence of Vertebral Fractures in Children with Suspected Osteoporosis. Journal of Pediatrics, 2016, 179, 219-225.	1.8	14
95	An Unbalanced Rearrangement of Chromosomes 4:20 is Associated with Childhood Osteoporosis and Reduced Caspase-3 Levels. Journal of Pediatric Genetics, 2016, 05, 167-173.	0.7	2
96	The Long-Term Outcome of Boys With Partial Androgen Insensitivity Syndrome and a Mutation in the Androgen Receptor Gene. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 3959-3967.	3.6	81
97	Assessing the feasibility of injectable growth-promoting therapy in Crohnâ€™s disease. Pilot and Feasibility Studies, 2016, 2, 71.	1.2	2
98	Current models of care for disorders of sex development â€“ results from an International survey of specialist centres. Orphanet Journal of Rare Diseases, 2016, 11, 155.	2.7	63
99	An audit of the management of childhood-onset growth hormone deficiency during young adulthood in Scotland. International Journal of Pediatric Endocrinology (Springer), 2016, 2016, 6.	1.6	12
100	Global Disorders of Sex Development Update since 2006: Perceptions, Approach and Care. Hormone Research in Paediatrics, 2016, 85, 158-180.	1.8	852
101	The outcome of prenatal identification of sex chromosome abnormalities. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2016, 101, F423-F427.	2.8	8
102	Growth and the Growth Hormone-Insulin Like Growth Factor 1 Axis in Children With Chronic Inflammation: Current Evidence, Gaps in Knowledge, and Future Directions. Endocrine Reviews, 2016, 37, 62-110.	20.1	104
103	Deficits in Trabecular Bone Microarchitecture in Young Women With Type 1 Diabetes Mellitus. Journal of Bone and Mineral Research, 2015, 30, 1386-1393.	2.8	82
104	Assessing Osteoporosis in the Young Adult. European Endocrinology, 2015, 11, 43.	1.5	0
105	Turner syndrome-issues to consider for transition to adulthood. British Medical Bulletin, 2015, 113, 45-58.	6.9	26
106	A critical appraisal of vertebral fracture assessment in paediatrics. Bone, 2015, 81, 255-259.	2.9	31
107	The pitfalls associated with urinary steroid metabolite ratios in children undergoing investigations for suspected disorders of steroid synthesis. International Journal of Pediatric Endocrinology (Springer), 2015, 2015, 10.	1.6	8
108	DNA copy number variations are important in the complex genetic architecture of mÃ¼llerian disorders. Fertility and Sterility, 2015, 103, 1021-1030.e1.	1.0	21

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109	An update on diabetes related skeletal fragility. Expert Review of Endocrinology and Metabolism, 2015, 10, 193-210.	2.4	3
110	Serum <scp>YKL</scp>â€40 in psoriasis with and without arthritis; correlation with disease activity and highâ€resolution power Doppler ultrasonographic joint findings. Journal of the European Academy of Dermatology and Venereology, 2015, 29, 682-688.	2.4	21
111	International Networks for Supporting Research and Clinical Care in the Field of Disorders of Sex Development. Endocrine Development, 2014, 27, 284-292.	1.3	26
112	Direct stimulation of bone mass by increased GH signalling in the osteoblasts of Socs2â^'/â^' mice. Journal of Endocrinology, 2014, 223, 93-106.	2.6	14
113	The Prevalence and Determinants of Short Stature in HIV-Infected Children. Journal of the International Association of Providers of AIDS Care, 2014, 13, 529-533.	1.5	7
114	<scp>MRI</scp>â€based abnormalities in young adults at risk of adverse bone health due to childhoodâ€onset metabolic & endocrine conditions. Clinical Endocrinology, 2014, 80, 811-817.	2.4	10
115	Management of children with disorders of sex development: new care standards explained. Psychology and Sexuality, 2014, 5, 5-14.	1.9	5
116	Endocrine Therapy for Growth Retardation in Paediatric Inflammatory Bowel Disease. Paediatric Drugs, 2014, 16, 29-42.	3.1	7
117	Novel Associations in Disorders of Sex Development: Findings From the I-DSD Registry. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E348-E355.	3.6	85
118	Contemporary Risk of Hip Fracture in Type 1 and Type 2 Diabetes: A National Registry Study From Scotland. Journal of Bone and Mineral Research, 2014, 29, 1054-1060.	2.8	111
119	Changes Over Time in Sex Assignment for Disorders of Sex Development. Pediatrics, 2014, 134, e710-e715.	2.1	98
120	PWE-072â€...The Effects Of Anti-tnf Therapy On Growth In Ibd In Scottish Children. Gut, 2014, 63, A155.1-A155.	12.1	0
121	Deficiency of the bone mineralization inhibitor NPP1 protects against obesity and diabetes. DMM Disease Models and Mechanisms, 2014, 7, 1341-50.	2.4	21
122	Understanding the genetic aetiology in patients with XY DSD. British Medical Bulletin, 2013, 106, 67-89.	6.9	79
123	An electronic surveillance system for monitoring the hospital presentation of nutritional vitamin D deficiency in children in Scotland. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 1053-8.	0.9	2
124	Pathophysiology and Management of Abnormal Growth in Children with Chronic Inflammatory Bowel Disease. World Review of Nutrition and Dietetics, 2013, 106, 142-148.	0.3	9
125	Management of boys and men with disorders of sex development. Current Opinion in Endocrinology, Diabetes and Obesity, 2012, 19, 190-196.	2.3	3
126	Urinary gonadotrophins: a useful non-invasive marker of activation of the hypothalamic pituitary-gonadal axis. International Journal of Pediatric Endocrinology (Springer), 2012, 2012, 10.	1.6	15

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127	A multidisciplinary approach to understanding skeletal dysplasias. Expert Review of Endocrinology and Metabolism, 2011, 6, 731-743.	2.4	2
128	UK guidance on the initial evaluation of an infant or an adolescent with a suspected disorder of sex development. Clinical Endocrinology, 2011, 75, 12-26.	2.4	124
129	Factors That Influence the Decision to Perform a Karyotype in Suspected Disorders of Sex Development: Lessons from the Scottish Genital Anomaly Network Register. Sexual Development, 2011, 5, 103-108.	2.0	24
130	The concordance between serum antiâ€Mullerian hormone and testosterone concentrations depends on duration of hCG stimulation in boys undergoing investigation of gonadal function. Clinical Endocrinology, 2010, 72, 814-819.	2.4	42
131	The effect of GH and IGF1 on linear growth and skeletal development and their modulation by SOCS proteins. Journal of Endocrinology, 2010, 206, 249-259.	2.6	114
132	Biologic therapy and its effect on skeletal development in children with chronic inflammation. Expert Review of Endocrinology and Metabolism, 2010, 5, 733-740.	2.4	0
133	Investigation and initial management of ambiguous genitalia. Best Practice and Research in Clinical Endocrinology and Metabolism, 2010, 24, 197-218.	4.7	88
134	The European Disorder of Sex Development Registry: A Virtual Research Environment. Sexual Development, 2010, 4, 192-198.	2.0	43
135	Congenital adrenal hyperplasia in a Nigerian child with a novel compound heterozygote mutation in CYP11B1. Clinical Endocrinology, 2007, 66, 070208104737006-???.	2.4	7
136	Prolonged human chorionic gonadotrophin stimulation as a tool for investigating and managing undescended testes. Clinical Endocrinology, 2007, 67, 816-821.	2.4	21
137	The psychological impact of genital anomalies on the parents of affected children. Acta Paediatrica, International Journal of Paediatrics, 2007, 96, 348-352.	1.5	73
138	Consensus statement on management of intersex disorders. Archives of Disease in Childhood, 2005, 91, 554-563.	1.9	900
139	Prevalence of hypospadias and other genital anomalies among singleton births, 1988-1997, in Scotland. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2004, 89, 149F-151.	2.8	100
140	Bone Mineral Assessment by Dual Energy X-ray Absorptiometry in Children With Inflammatory Bowel Disease: Evaluation by Age or Bone Area. Journal of Pediatric Gastroenterology and Nutrition, 2004, 38, 276-280.	1.8	65
141	Testosterone measurements in early infancy. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2004, 89, F558-F559.	2.8	48
142	Erroneous testosterone assay causing diagnostic confusion in a newborn infant with intersex anomalies. Acta Paediatrica, International Journal of Paediatrics, 2004, 93, 1004-1005.	1.5	7
143	Shortâ€term growth and bone turnover in children undergoing occlusive steroid (â€˜Wetâ€Wrapâ€™) dressings for treatment of atopic eczema. Journal of Dermatological Treatment, 2003, 14, 149-152.	2.2	21
144	The genetics of male undermasculinization. Clinical Endocrinology, 2002, 56, 1-18.	2.4	59

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145	The role of a clinical score in the assessment of ambiguous genitalia. BJU International, 2000, 85, 120-124.	2.5	218
146	The testosterone:androstenedione ratio in male undermasculinization. Clinical Endocrinology, 2000, 53, 697-702.	2.4	65
147	Pituitary-gonadal axis in male undermasculinisation. Archives of Disease in Childhood, 2000, 82, 54-58.	1.9	28
148	Phenotypic Features, Androgen Receptor Binding, and Mutational Analysis in 278 Clinical Cases Reported as Androgen Insensitivity Syndrome1. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 658-665.	3.6	269
149	Phenotypic Features, Androgen Receptor Binding, and Mutational Analysis in 278 Clinical Cases Reported as Androgen Insensitivity Syndrome. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 658-665.	3.6	244
150	Clinical Experience during the Paediatric Undergraduate Course. Journal of the Royal Society of Medicine, 1999, 92, 293-298.	2.0	6
151	Assessment of the gonadotrophin-gonadal axis in androgen insensitivity syndrome. Archives of Disease in Childhood, 1999, 80, 324-329.	1.9	89
152	GNAS1 mutational analysis in pseudohypoparathyroidism. Clinical Endocrinology, 1998, 49, 525-531.	2.4	68
153	GNAS1 mutational analysis in pseudohypoparathyroidism. Clinical Endocrinology, 1998, 49, 525-31.	2.4	22
154	Initial evaluation of congenital hypothyroidism: a survey of general paediatricians in East Anglia. Archives of Disease in Childhood, 1997, 77, 339-341.	1.9	6