Andrew A Dwyer

List of Publications by Year
in descending order

130	8,206	66234	48187
papers	citations	42 h-index	88 g-index

133	133	133	5968
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Coping response and family communication of cancer risk in men harboring aBRCAmutation: A mixed methods study. Psycho-Oncology, 2022, 31, 486-495.	1.0	5
2	What's missing in sex chromosome aneuploidies? Representation and inclusion. Journal of Pediatric Nursing, 2022, 62, 202-204.	0.7	1
3	A Comparison of the Blood Glucose, Growth Hormone, and Cortisol Responses to Two Doses of Insulin (0.15â€‰U/kg vs. 0.10â€‰U/kg) in the Insulin Tolerance Test: A Single-Centre Audit of 174 Cases. International Journal of Endocrinology, 2022, 2022, 1-8.	0.6	1
4	Impact of BRCA Status on Reproductive Decision-Making and Self-Concept: A Mixed-Methods Study Informing the Development of Tailored Interventions. Cancers, 2022, 14, 1494.	1.7	2
5	Exploring Rare Disease Patient Attitudes and Beliefs regarding Genetic Testing: Implications for Person-Centered Care. Journal of Personalized Medicine, 2022, 12, 477.	1.1	3
6	Correspondence on â€œEnsuring best practice in genomics education and evaluation: Reporting item standards for education and its evaluation in genomics (RISE2 Genomics)â€•by Nisselle etÂal. Genetics in Medicine, 2022, 24, 962-963.	1.1	2
7	A Developmental Perspective Sheds Light on Reproductive Differences Between Congenital and Acquired Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e3955-e3956.	1.8	0
8	Congenital Hypogonadotropic Hypogonadism with Anosmia and Gorlin Features Caused by a PTCH1 Mutation Reveals a New Candidate Gene for Kallmann Syndrome. Neuroendocrinology, 2021, 111, 99-114.	1.2	20
9	Evaluating coâ€created patientâ€facing materials to increase understanding of genetic test results. Journal of Genetic Counseling, 2021, 30, 598-605.	0.9	19
10	Symptom perception in heart failure â€“ Interventions and outcomes: A scoping review. International Journal of Nursing Studies, 2021, 116, 103524.	2.5	17
11	Congenital hyperinsulinism: 2 case reports with different rare variants in ABCC8. Annals of Pediatric Endocrinology and Metabolism, 2021, 26, 60-65.	0.8	2
12	Validating online approaches for rare disease research using latent class mixture modeling. Orphanet Journal of Rare Diseases, 2021, 16, 209.	1.2	4
13	Insulin-like Growth Factor 1, but Not Insulin-Like Growth Factor-Binding Protein 3, Predicts Central Precocious Puberty in Girls 6â€“8 Years Old: A Retrospective Study. Hormone Research in Paediatrics, 2021, 94, 1-8.	0.8	5
14	Framing Effects on Decision-Making for Diagnostic Genetic Testing: Results from a Randomized Trial. Genes, 2021, 12, 941.	1.0	3
15	Endocrine Nurses Society Position Statement on Transgender and Gender Diverse Care. Journal of the Endocrine Society, 2021, 5, bvab105.	0.1	3
16	A rare disease patient-reported outcome measure: revision and validation of the German version of the Systemic Sclerosis Quality of Life Questionnaire (SScQoL) using the Rasch model. Orphanet Journal of Rare Diseases, 2021, 16, 356.	1.2	2
17	Patient and healthcare professional eHealth literacy and needs for systemic sclerosis support: a mixed methods study. RMD Open, 2021, 7, e001783.	1.8	9
18	Parent of origin differences in psychosocial burden and approach to BRCA risk management. Breast Journal, 2020, 26, 734-738.	0.4	7

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19	Minipuberty: A Primer for Pediatric Nurses. Journal of Pediatric Nursing, 2020, 50, 138-139.	0.7	1
20	Neuron-Derived Neurotrophic Factor Is Mutated in Congenital Hypogonadotropic Hypogonadism. American Journal of Human Genetics, 2020, 106, 58-70.	2.6	39
21	Symptom perception in heart failure: a scoping review on definition, factors and instruments. European Journal of Cardiovascular Nursing, 2020, 19, 100-117.	0.4	25
22	GnRH stimulation testing and serum inhibin B in males: insufficient specificity for discriminating between congenital hypogonadotropic hypogonadism from constitutional delay of growth and puberty. Human Reproduction, 2020, 35, 2312-2322.	0.4	13
23	Parent of Origin Effects on Family Communication of Risk in BRCA+ Women: A Qualitative Investigation of Human Factors in Cascade Screening. Cancers, 2020, 12, 2316.	1.7	11
24	Editorial: New Aspects in Hypogonadism. Frontiers in Endocrinology, 2020, 11, 426.	1.5	0
25	A sexual health course for advanced practice registered nurses: Effect on preparedness, comfort, and confidence in delivering comprehensive care. Nurse Education Today, 2020, 92, 104506.	1.4	10
26	Non-invasive assessment of coronary endothelial function in children and adolescents with type 1 diabetes mellitus using isometric handgrip exerciseâ€"MRI: A feasibility study. PLoS ONE, 2020, 15, e0228569.	1.1	5
27	Insight Into the Ontogeny of GnRH Neurons From Patients Born Without a Nose. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1538-1551.	1.8	7
28	Psychosexual effects resulting from delayed, incomplete, or absent puberty. Current Opinion in Endocrine and Metabolic Research, 2020, 14, 15-21.	0.6	14
29	Congenital hypogonadotropic hypogonadism/Kallmann syndrome is associated with statural gain in both men and women: a monocentric study. European Journal of Endocrinology, 2020, 182, 185.	1.9	21
30	Transitional Care in Endocrinology. , 2020, , 281-317.		0
31	Fertility and the Hypogonadal Male. , 2019, , 94-105.		0
32	Psychological Aspects of Congenital Hypogonadotropic Hypogonadism. Frontiers in Endocrinology, 2019, 10, 353.	1.5	26
33	Genetic Competencies for Effective Pediatric Endocrine Nursing Practice. Journal of Pediatric Nursing, 2019, 48, 127-128.	0.7	2
34	Developing a rare disease chronic care model: Management of systemic sclerosis (MANOSS) study protocol. Journal of Advanced Nursing, 2019, 75, 3774-3791.	1.5	4
35	Functional Hypogonadotropic Hypogonadism in Men: Underlying Neuroendocrine Mechanisms and Natural History. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3403-3414.	1.8	28
36	Patient perceptions of peripheral artery disease: A cross-sectional study of hospitalized adults. Journal of Vascular Nursing, 2019, 37, 188-193.	0.2	3

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37	Classification of Hypothalamic-Pituitary-Gonadal (HPG) Axis Endocrine Disorders. , 2019, , 853-870.		2
38	Spermatogenesis and Assisted Fertility Treatment. , 2019, , 903-923.		0
39	Anatomy and Physiology of theÂHypothalamic-Pituitary-Gonadal (HPG) Axis. , 2019, , 839-852.		20
40	Evaluation of Endocrine Disorders of the Hypothalamic-Pituitary-Gonadal (HPG) Axis. , 2019, , 871-883.		0
41	Thigh and abdominal adipose tissue depot associations with testosterone levels in postmenopausal females. Clinical Endocrinology, 2019, 90, 433-439.	1.2	12
42	A novel CHD7 mutation in an adolescent presenting with growth and pubertal delay. Annals of Pediatric Endocrinology and Metabolism, 2019, 24, 49-54.	0.8	5
43	The Metabolic Syndrome in Central Hypogonadotrophic Hypogonadism. Frontiers of Hormone Research, 2018, 49, 156-169.	1.0	19
44	PENS Position Statement on Bullying Prevention. Journal of Pediatric Nursing, 2018, 39, 91-93.	0.7	1
45	Congenital hypogonadotropic hypogonadism and constitutional delay of growth and puberty have distinct genetic architectures. European Journal of Endocrinology, 2018, 178, 377-388.	1.9	95
46	GENETICS IN ENDOCRINOLOGY: Genetic counseling for congenital hypogonadotropic hypogonadism and Kallmann syndrome: new challenges in the era of oligogenism and next-generation sequencing. European Journal of Endocrinology, 2018, 178, R55-R80.	1.9	128
47	Patient Perspectives on Nurse-led Consultations Within a Pilot Structured Transition Program for Young Adults Moving From an Academic Tertiary Setting to Community-based Type 1 Diabetes Care. Journal of Pediatric Nursing, 2018, 38, 99-105.	0.7	17
48	DCC/NTN1 complex mutations in patients with congenital hypogonadotropic hypogonadism impair GnRH neuron development. Human Molecular Genetics, 2018, 27, 359-372.	1.4	42
49	International practice settings, interventions and outcomes of nurse practitioners in geriatric care: A scoping review. International Journal of Nursing Studies, 2018, 78, 61-75.	2.5	54
50	Metaâ€analysis of the effectiveness of nursing discharge planning interventions for older inpatients discharged home. Journal of Advanced Nursing, 2018, 74, 788-799.	1.5	37
51	Evaluating CHARGE syndrome in congenital hypogonadotropic hypogonadism patients harboring CHD7 variants. Genetics in Medicine, 2018, 20, 872-881.	1.1	38
52	Fertility induction in hypogonadotropic hypogonadal men. Clinical Endocrinology, 2018, 89, 712-718.	1.2	32
53	Acceptance of the Advanced Practice Nurse in Lung Cancer Role by Healthcare Professionals and Patients: A Qualitative Exploration. Journal of Nursing Scholarship, 2018, 50, 540-548.	1.1	18
54	Accuracy, satisfaction and usability of a flash glucose monitoring system among children and adolescents with type 1 diabetes attending a summer camp. Pediatric Diabetes, 2018, 19, 1276-1284.	1.2	20

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55	Transition of Care from Childhood to Adulthood: Congenital Hypogonadotropic Hypogonadism. Endocrine Development, 2018, 33, 82-98.	1.3	6
56	Genetic testing facilitates prepubertal diagnosis of congenital hypogonadotropic hypogonadism. Clinical Genetics, 2017, 92, 213-216.	1.0	14
57	Feasibility of advanced practice nursing in lung cancer consultations during early treatment: A phase II study. European Journal of Oncology Nursing, 2017, 29, 106-114.	0.9	14
58	<scp>KLB</scp> , encoding Î²â€Klotho, is mutated in patients with congenital hypogonadotropic hypogonadism. EMBO Molecular Medicine, 2017, 9, 1379-1397.	3.3	77
59	Beyond hormone replacement: quality of life in women with congenital hypogonadotropic hypogonadism. Endocrine Connections, 2017, 6, 404-412.	0.8	31
60	Developing and evaluating rare disease educational materials co-created by expert clinicians and patients: the paradigm of congenital hypogonadotropic hypogonadism. Orphanet Journal of Rare Diseases, 2017, 12, 57.	1.2	26
61	Society for Endocrinology <scp>UK</scp> guidance on the evaluation of suspected disorders of sexual development: emphasizing the opportunity to predict adolescent pubertal failure through a neonatal diagnosis of absent minipuberty. Clinical Endocrinology, 2017, 86, 305-306.	1.2	21
62	Adherence to treatment in men with hypogonadotrophic hypogonadism. Clinical Endocrinology, 2017, 86, 377-383.	1.2	32
63	004-Caring Matters: A Cross-Sectional Study of Patient Perspectives on Nurse-led Consultations as Part of Structured Transitional Care for Type 1 Diabetes. Journal of Pediatric Nursing, 2017, 37, 144.	0.7	0
64	Î²-Klotho deficiency protects against obesity through a crosstalk between liver, microbiota, and brown adipose tissue. JCI Insight, 2017, 2, .	2.3	41
65	Multidisciplinary management of diabetic kidney disease. JBI Database of Systematic Reviews and Implementation Reports, 2016, 14, 169-207.	1.7	26
66	Effectiveness of nursing discharge planning interventions on health-related outcomes in discharged elderly inpatients. JBI Database of Systematic Reviews and Implementation Reports, 2016, 14, 217-260.	1.7	21
67	MANAGEMENT OF ENDOCRINE DISEASE: Reversible hypogonadotropic hypogonadism. European Journal of Endocrinology, 2016, 174, R267-R274.	1.9	66
68	Congenital hypogonadotropic hypogonadism: implications of absent mini-puberty. Minerva Endocrinologica, 2016, 41, 188-95.	1.7	12
69	Natural History of Growth Hormone Deficiency in a Pediatric Cohort. Hormone Research in Paediatrics, 2015, 83, 252-261.	0.8	12
70	European-South Africa collaboration on the genetic basis of gonadotropin-releasing hormone deficiency in failure to progress through puberty and infertility. Journal of Endocrinology Metabolism and Diabetes of South Africa, 2015, 20, 67-68.	0.4	4
71	Psychosexual Development in Men with Congenital Hypogonadotropic Hypogonadism on Long-Term Treatment: A Mixed Methods Study. Sexual Medicine, 2015, 3, 32-41.	0.9	34
72	TRANSITION IN ENDOCRINOLOGY: Hypogonadism in adolescence. European Journal of Endocrinology, 2015, 173, R15-R24.	1.9	59

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73	European Consensus Statement on congenital hypogonadotropic hypogonadismâ€"pathogenesis, diagnosis and treatment. Nature Reviews Endocrinology, 2015, 11, 547-564.	4.3	664
74	Gonadotrophin replacement for induction of fertility in hypogonadal men. Best Practice and Research in Clinical Endocrinology and Metabolism, 2015, 29, 91-103.	2.2	96
75	Congenital hypogonadotropic hypogonadism with split hand/foot malformation: a clinical entity with a high frequency of FGFR1 mutations. Genetics in Medicine, 2015, 17, 651-659.	1.1	55
76	Comment on reversal of hypogonadotropic hypogonadism in a Chinese cohort. Asian Journal of Andrology, 2015, 17, 508.	0.8	3
77	Reversal and Relapse of Hypogonadotropic Hypogonadism: Resilience and Fragility of the Reproductive Neuroendocrine System. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 861-870.	1.8	144
78	Absence of Central Circadian Pacemaker Abnormalities in Humans With Loss of Function Mutation in Prokineticin 2. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E561-E566.	1.8	8
79	Hormonal control of spermatogenesis in men: Therapeutic aspects in hypogonadotropic hypogonadism. Annales D'Endocrinologie, 2014, 75, 98-100.	0.6	21
80	Skeletal Muscle Mitochondria in the Elderly: Effects of Physical Fitness and Exercise Training. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1852-1861.	1.8	114
81	Identifying the unmet health needs of patients with congenital hypogonadotropic hypogonadism using a web-based needs assessment: implications for online interventions and peer-to-peer support. Orphanet Journal of Rare Diseases, 2014, 9, 83.	1.2	63
82	Comparative functional analysis of two fibroblast growth factor receptor 1 (FGFR1) mutations affecting the same residue (R254W and R254Q) in isolated hypogonadotropic hypogonadism (IHH). Gene, 2013, 516, 146-151.	1.0	19
83	Abrupt decrease in serum testosterone levels after an oral glucose load in men: implications for screening for hypogonadism. Clinical Endocrinology, 2013, 78, 291-296.	1.2	91
84	Mutations in FGF17, IL17RD, DUSP6, SPRY4, and FLRT3 Are Identified in Individuals with Congenital Hypogonadotropic Hypogonadism. American Journal of Human Genetics, 2013, 92, 725-743.	2.6	227
85	Ataxia, Dementia, and Hypogonadotropism Caused by Disordered Ubiquitination. New England Journal of Medicine, 2013, 368, 1992-2003.	13.9	208
86	Trial of Recombinant Follicle-Stimulating Hormone Pretreatment for GnRH-Induced Fertility in Patients with Congenital Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E1790-E1795.	1.8	124
87	Prioritizing Genetic Testing in Patients With Kallmann Syndrome Using Clinical Phenotypes. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E943-E953.	1.8	157
88	Testosterone restoration using enclomiphene citrate in men with secondary hypogonadism: a pharmacodynamic and pharmacokinetic study. BJU International, 2013, 112, 1188-1200.	1.3	35
89	Effectiveness of nursing discharge planning interventions on health-related outcomes in elderly inpatients discharged home: a systematic review protocol. JBI Database of Systematic Reviews and Implementation Reports, 2013, 11, 1-12.	1.7	4
90	Genetic Overlap in Kallmann Syndrome, Combined Pituitary Hormone Deficiency, and Septo-Optic Dysplasia. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E694-E699.	1.8	136

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91	The lack of effect of insulin on luteinizing hormone pulsatility in healthy male volunteers provides evidence of a sexual dimorphism in the metabolic regulation of reproductive hormones. American Journal of Clinical Nutrition, 2012, 96, 283-288.	2.2	6
92	An ancient founder mutation in PROKR2 impairs human reproduction. Human Molecular Genetics, 2012, 21, 4314-4324.	1.4	31
93	GnRH-Deficient Phenotypes in Humans and Mice With Heterozygous Variants in KISS1/Kiss1. Obstetrical and Gynecological Survey, 2012, 67, 546-547.	0.2	0
94	Role of fibroblast growth factor (FGF) signaling in the neuroendocrine control of human reproduction. Molecular and Cellular Endocrinology, 2011, 346, 37-43.	1.6	24
95	Genetic basis and variable phenotypic expression of Kallmann syndrome: towards a unifying theory. Trends in Endocrinology and Metabolism, 2011, 22, 249-58.	3.1	127
96	The Role of the Prokineticin 2 Pathway in Human Reproduction: Evidence from the Study of Human and Murine Gene Mutations. Endocrine Reviews, 2011, 32, 225-246.	8.9	95
97	GnRH-Deficient Phenotypes in Humans and Mice with Heterozygous Variants inKISS1/Kiss1. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E1771-E1781.	1.8	59
98	Expanding the Phenotype and Genotype of Female GnRH Deficiency. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E566-E576.	1.8	97
99	TAC3/TACR3 Mutations Reveal Preferential Activation of GnRH Release by Neurokinin B in Neonatal Life Followed by Reversal in Adulthood. Endocrinology, 2010, 151, 1970-1971.	1.4	0
100	Congenital Idiopathic Hypogonadotropic Hypogonadism: Evidence of Defects in the Hypothalamus, Pituitary, and Testes. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3019-3027.	1.8	115
101	TAC3/TACR3 Mutations Reveal Preferential Activation of Gonadotropin-Releasing Hormone Release by Neurokinin B in Neonatal Life Followed by Reversal in Adulthood. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 2857-2867.	1.8	250
102	Impact of Acute Biochemical Castration on Insulin Sensitivity in Healthy Adult Men. Endocrine Research, 2010, 35, 71-84.	0.6	15
103	Oligogenic basis of isolated gonadotropin-releasing hormone deficiency. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 15140-15144.	3.3	313
104	Human GnRH Deficiency: A Unique Disease Model to Unravel the Ontogeny of GnRH Neurons. Neuroendocrinology, 2010, 92, 81-99.	1.2	87
105	The Long-Term Clinical Follow-Up and Natural History of Men with Adult-Onset Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 4235-4243.	1.8	45
106	Role of Seminiferous Tubular Development in Determining the FSH versus LH Responsiveness to GnRH in Early Sexual Maturation. Neuroendocrinology, 2009, 90, 260-268.	1.2	9
107	Impaired Fibroblast Growth Factor Receptor 1 Signaling as a Cause of Normosmic Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 4380-4390.	1.8	82
108	Mutations inProkineticin 2andProkineticin receptor 2genes in Human Gonadotrophin-Releasing Hormone Deficiency: Molecular Genetics and Clinical Spectrum. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 3551-3559.	1.8	190

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109	Relative Roles of Inhibin B and Sex Steroids in the Negative Feedback Regulation of Follicle-Stimulating Hormone in Men across the Full Spectrum of Seminiferous Epithelium Function. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 1809-1814.	1.8	40
110	Inhibition of Luteinizing Hormone Secretion by Testosterone in Men Requires Aromatization for Its Pituitary But Not Its Hypothalamic Effects: Evidence from the Tandem Study of Normal and Gonadotropin-Releasing Hormone-Deficient Men. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 784-791.	1.8	119
111	The Relative Role of Gonadal Sex Steroids and Gonadotropin-Releasing Hormone Pulse Frequency in the Regulation of Follicle-Stimulating Hormone Secretion in Men. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 2686-2692.	1.8	55
112	Decreased FGF8 signaling causes deficiency of gonadotropin-releasing hormone in humans and mice. Journal of Clinical Investigation, 2008, 118, 2822-2831.	3.9	348
113	Reversal of Idiopathic Hypogonadotropic Hypogonadism. New England Journal of Medicine, 2007, 357, 863-873.	13.9	362
114	Acute Sex Steroid Withdrawal Reduces Insulin Sensitivity in Healthy Men with Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 4254-4259.	1.8	208
115	Digenic mutations account for variable phenotypes in idiopathic hypogonadotropic hypogonadism. Journal of Clinical Investigation, 2007, 117, 457-463.	3.9	338
116	Mutations in fibroblast growth factor receptor 1 cause Kallmann syndrome with a wide spectrum of reproductive phenotypes. Molecular and Cellular Endocrinology, 2006, 254-255, 60-69.	1.6	176
117	Testis morphology in patients with idiopathic hypogonadotropic hypogonadism. Human Reproduction, 2006, 21, 1033-1040.	0.4	19
118	Coding sequence analysis of GNRHR and GPR54 in patients with congenital and adult-onset forms of hypogonadotropic hypogonadism. European Journal of Endocrinology, 2006, 155, S3-S10.	1.9	72
119	Mutations in fibroblast growth factor receptor 1 cause both Kallmann syndrome and normosmic idiopathic hypogonadotropic hypogonadism. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 6281-6286.	3.3	225
120	Reversible Kallmann Syndrome, Delayed Puberty, and Isolated Anosmia Occurring in a Single Family with a Mutation in the Fibroblast Growth Factor Receptor 1 Gene. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 1317-1322.	1.8	144
121	Relationship Between Testosterone Levels, Insulin Sensitivity, and Mitochondrial Function in Men. Diabetes Care, 2005, 28, 1636-1642.	4.3	392
122	Increasing Insulin Resistance Is Associated with a Decrease in Leydig Cell Testosterone Secretion in Men. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 2636-2641.	1.8	424
123	Acute Stress Masking the Biochemical Phenotype of Partial Androgen Insensitivity Syndrome in a Patient with a Novel Mutation in the Androgen Receptor. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 1053-1058.	1.8	4
124	Predictors of Outcome of Long-Term GnRH Therapy in Men with Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2002, 87, 4128-4136.	1.8	210
125	Adherence to treatment for chronic hypogonadism: the role of illness perceptions and depressive symptoms. Endocrine Abstracts, 0, , .	0.0	0
126	Research topic: identifying the needs of patients with Congenital Hypogonadotrophic Hypogonadism, implications for nursing practice. Endocrine Abstracts, 0, , .	0.0	0

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127	Unmet health and information needs of women with hypogonadotropic hypogonadism. Endocrine Abstracts, 0, , .	0.0	0
128	Secondary Infertility. Endocrine Abstracts, 0, , .	0.0	0
129	Clinical practice overlap and seamless care - links between hypogonadism, the metabolic syndrome and type 2 diabetes. Endocrine Abstracts, 0, , .	0.0	0
130	Navigating Disrupted Puberty: Development and Evaluation of a Mobile-Health Transition Passport for Klinefelter Syndrome. Frontiers in Endocrinology, 0, 13, .	1.5	4