Phyllis W Speiser

List of Publications by Year in Descending Order

45 papers	5,977 citations	24 h-index	48 g-index
48 ext. papers	6,780 ext. citations	8.2 avg, IF	5.36 L-index

#	Paper	IF	Citations
45	Congenital adrenal hyperplasia - current insights in pathophysiology, diagnostics and management. *Endocrine Reviews*, **2021**,	27.2	28
44	Newborn Screening Protocols and Positive Predictive Value for Congenital Adrenal Hyperplasia Vary across the United States. *International Journal of Neonatal Screening*, **2020**, 6,	2.6	11
43	Invited Commentary: A Phase 2, Multicenter Study of Nevanimibe for the Treatment of Congenital Adrenal Hyperplasia. *Journal of Clinical Endocrinology and Metabolism*, **2020**, 105,	5.6	3
42	Response to Letter to the Editor: "Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society Clinical Practice Guideline". *Journal of Clinical Endocrinology and Metabolism*, **2019**, 104, 1928	5.6	1
41	Demographics and anthropometrics impact benefits of health intervention: data from the Reduce Obesity and Diabetes Project. *Obesity Science and Practice*, **2019**, 5, 46-58	2.6	
40	Emerging medical therapies for congenital adrenal hyperplasia. *F1000Research*, **2019**, 8, 363	3.6	4
39	Iatrogenic Cushing Syndrome in a Child With Congenital Adrenal Hyperplasia: Erroneous Compounding of Hydrocortisone. *Journal of Clinical Endocrinology and Metabolism*, **2018**, 103, 7-11	5.6	15
38	Psychosocial Screening in Disorders/Differences of Sex Development: Psychometric Evaluation of the Psychosocial Assessment Tool. *Hormone Research in Paediatrics*, **2018**, 90, 368-380	3.3	10
37	Cardiovascular and Metabolic Outcomes in Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. *Journal of Clinical Endocrinology and Metabolism*, **2018**, 103, 4097-4103	5.6	38
36	Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society Clinical Practice Guideline. *Journal of Clinical Endocrinology and Metabolism*, **2018**, 103, 4043-4088	5.6	371
35	Genital Reconstructive Surgery in Females With Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. *Journal of Clinical Endocrinology and Metabolism*, **2018**, 103, 4089-4096	5.6	33
34	Plasma advanced glycation end products (AGEs), receptors for AGEs and their correlation with inflammatory markers in middle school-age children. *Hormone Research in Paediatrics*, **2013**, 80, 318-27	3.3	19
33	The Reduce Obesity and Diabetes (ROAD) Project: Design and Methodological Considerations. *Childhood Obesity*, **2011**, 7, 223-234	2.5	7
32	Medical treatment of classic and nonclassic congenital adrenal hyperplasia. *Advances in Experimental Medicine and Biology*, **2011**, 707, 41-5	3.6	6
31	Guidelines for the Development of Comprehensive Care Centers for Congenital Adrenal Hyperplasia: Guidance from the CARES Foundation Initiative. *International Journal of Pediatric Endocrinology (Springer)*, **2010**, 2010, 275213	1.5	20
30	A Summary of the Endocrine Society Clinical Practice Guidelines on Congenital Adrenal Hyperplasia due to Steroid 21-Hydroxylase Deficiency. *International Journal of Pediatric Endocrinology (Springer)* , **2010**, 2010, 494173	1.5	20
29	Congenital adrenal hyperplasia due to steroid 21-hydroxylase deficiency: an Endocrine Society clinical practice guideline. *Journal of Clinical Endocrinology and Metabolism*, **2010**, 95, 4133-60	5.6	783

28	Prevention and treatment of pediatric obesity: an endocrine society clinical practice guideline based on expert opinion. *Journal of Clinical Endocrinology and Metabolism*, **2008**, 93, 4576-99	5.6	343
27	Reproductive outcome of women with 21-hydroxylase-deficient nonclassic adrenal hyperplasia. *Journal of Clinical Endocrinology and Metabolism*, **2006**, 91, 3451-6	5.6	113
26	Childhood obesity. *Journal of Clinical Endocrinology and Metabolism*, **2005**, 90, 1871-87	5.6	404
25	Congenital adrenal hyperplasia. *New England Journal of Medicine*, **2003**, 349, 776-88	59.2	669
24	Consensus statement on 21-hydroxylase deficiency from the European Society for Paediatric Endocrinology and the Lawson Wilkins Pediatric Endocrine Society. *Hormone Research in Paediatrics* , **2002**, 58, 188-95	3.3	110
23	Molecular diagnosis of CYP21 mutations in congenital adrenal hyperplasia: implications for genetic counseling. *Molecular Diagnosis and Therapy*, **2001**, 1, 101-10		22
22	Congenital adrenal hyperplasia: transition from chil dhood to adulthood. *Journal of Endocrinological Investigation*, **2001**, 24, 681-91	5.2	18
21	Profiles of obese children presenting for metabolic evaluation. *Journal of Pediatric Endocrinology and Metabolism*, **2001**, 14, 1145-50	1.6	12
20	21-Hydroxylase-deficient nonclassic adrenal hyperplasia is a progressive disorder: a multicenter study. *American Journal of Obstetrics and Gynecology*, **2000**, 183, 1468-74	6.4	128
19	Distinct missense mutations of the FGFR3 lys650 codon modulate receptor kinase activation and the severity of the skeletal dysplasia phenotype. *American Journal of Human Genetics*, **2000**, 67, 1411-21	11	131
18	Congenital adrenal hyperplasia due to 21-hydroxylase deficiency. *Endocrine Reviews*, **2000**, 21, 245-91	27.2	834
17	Genotyping of CYP21, linked chromosome 6p markers, and a sex-specific gene in neonatal screening for congenital adrenal hyperplasia. *Journal of Clinical Endocrinology and Metabolism*, **1999** , 84, 960-6	5.6	81
16	Prenatal treatment of congenital adrenal hyperplasia. *Journal of Urology*, **1999**, 162, 534-6	2.5	3
15	Congenital adrenal hyperplasia due to steroid 21-hydroxylase deficiency. *Clinical Endocrinology*, **1998**, 49, 411-7	3.4	20
14	Magnetic resonance imaging in the congenital adrenal hyperplasia population: increased frequency of white-matter abnormalities and temporal lobe atrophy. *Journal of Child Neurology*, **1997**, 12, 181-6	2.5	48
13	Steroid 21-hydroxylase expression and activity in human lymphocytes. *Molecular and Cellular Endocrinology*, **1997**, 127, 11-8	4.4	25
12	Prenatal diagnosis and treatment of congenital adrenal hyperplasia. *Journal of Pediatric Endocrinology and Metabolism*, **1994**, 7, 183-91	1.6	14
11	Mutations in steroid 21-hydroxylase (CYP21). *Human Mutation*, **1994**, 3, 373-8	4.7	95

10	Prenatal diagnosis and management of congenital adrenal hyperplasia. *Clinics in Perinatology*, **1994**, 21, 631-45	2.8	2
9	Mutations in the CYP11B1 gene causing congenital adrenal hyperplasia and hypertension cluster in exons 6, 7, and 8. *Proceedings of the National Academy of Sciences of the United States of America*, **1993**, 90, 4552-6	11.5	171
8	Glucocorticoid-suppressible hyperaldosteronism results from hybrid genes created by unequal crossovers between CYP11B1 and CYP11B2. *Proceedings of the National Academy of Sciences of the United States of America*, **1992**, 89, 8327-31	11.5	213
7	Disease expression and molecular genotype in congenital adrenal hyperplasia due to 21-hydroxylase deficiency. *Journal of Clinical Investigation*, **1992**, 90, 584-95	15.9	375
6	Insulin insensitivity in adrenal hyperplasia due to nonclassical steroid 21-hydroxylase deficiency. *Journal of Clinical Endocrinology and Metabolism*, **1992**, 75, 1421-1424	5.6	86
5	A mutation (Pro-30 to Leu) in CYP21 represents a potential nonclassic steroid 21-hydroxylase deficiency allele. *Molecular Endocrinology*, **1991**, 5, 685-92		162
4	Growth and final height in classical and nonclassical 21-hydroxylase deficiency. *Journal of Endocrinological Investigation*, **1989**, 12, 91-5	5.2	10
3	Molecular genetic analysis of nonclassic steroid 21-hydroxylase deficiency associated with HLA-B14,DR1. *New England Journal of Medicine*, **1988**, 319, 19-23	59.2	170
2	High Frequency of Nonclassical Steroid 21-Hydroxylase Deficiency. *Obstetrical and Gynecological Survey*, **1986**, 41, 244-245	2.4	9
1	High frequency of nonclassical steroid 21-hydroxylase deficiency. *American Journal of Human Genetics*, **1985**, 37, 650-67	11	339