

Phenotypic profiling of parents with cryptic nonclassic findings in 145 unrelated families

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society Clinical Practice Guideline. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 4133-4160.	3.6	1,117
2	Relationship of CYP21A2 genotype and serum 17-hydroxyprogesterone and cortisol levels in a large cohort of Italian children with premature pubarche. <i>European Journal of Endocrinology</i> , 2011, 165, 307-314.	3.7	39
3	Management of CAH during pregnancy. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2012, 19, 489-496.	2.3	32
5	Absence of Clinically Relevant Growth Acceleration in Untreated Children with Non-Classical Congenital Adrenal Hyperplasia. <i>Hormone Research in Paediatrics</i> , 2012, 77, 164-169.	1.8	13
6	Nonclassic congenital adrenal hyperplasia. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2012, 19, 151-158.	2.3	31
8	Prenatal Treatment of Congenital Adrenal Hyperplasiaâ€”Not Standard of Care. <i>Journal of Genetic Counseling</i> , 2012, 21, 615-624.	1.6	29
9	Non-classic congenital adrenal hyperplasia. <i>Steroids</i> , 2013, 78, 747-750.	1.8	45
10	Approach to the Patient: The Adult With Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2013, 98, 2645-2655.	3.6	107
11	Management of adolescents with congenital adrenal hyperplasia. <i>Lancet Diabetes and Endocrinology</i> , 2013, 1, 341-352.	11.4	90
12	Excited state dynamics of DNA bases. <i>International Reviews in Physical Chemistry</i> , 2013, 32, 308-342.	2.3	185
13	Endocrine Disturbances Affecting Reproduction. , 2014, , 551-564.e4.		3
16	Fetal endocrine therapy for congenital adrenal hyperplasia should not be done. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2015, 29, 469-483.	4.7	32
17	Inadequate Cortisol Response to the Tetracosactide (SynacthenÂ®) Test in Non-Classic Congenital Adrenal Hyperplasia: An Exception to the Rule?. <i>Hormone Research in Paediatrics</i> , 2015, 83, 262-267.	1.8	25
18	The Classic and Nonclassic Congenital Adrenal Hyperplasias. <i>Endocrine Practice</i> , 2015, 21, 383-389.	2.1	29
20	Adrenal Steroid Metabolites Accumulating in Congenital Adrenal Hyperplasia Lead to Transactivation of the Glucocorticoid Receptor. <i>Endocrinology</i> , 2015, 156, 3504-3510.	2.8	32
21	Substitution therapy in adult patients with congenital adrenal hyperplasia. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2015, 29, 33-45.	4.7	20
22	The spectrum of clinical, hormonal and molecular findings in 280 individuals with nonclassical congenital adrenal hyperplasia caused by mutations of the <i>CYP21A2</i> gene. <i>Clinical Endocrinology</i> , 2015, 82, 543-549.	2.4	68
23	Cortisol response to adrenocorticotropin testing in non-classical congenital adrenal hyperplasia (NCCAH). <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2016, 29, 1365-1371.	0.9	9

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24	Intrafamilial Phenotypic Variability and Consequences of Non-Compliance with Treatment in Congenital Adrenal Hyperplasia and Congenital Hypothyroidism within a Single Family. <i>Hormone Research in Paediatrics</i> , 2017, 88, 172-178.	1.8	2
25	Congenital Adrenal Hyperplasia. <i>Journal of Pediatric and Adolescent Gynecology</i> , 2017, 30, 520-534.	0.7	117
26	The spectrum of CYP21A2 mutations in Congenital Adrenal Hyperplasia in an Indian cohort. <i>Clinica Chimica Acta</i> , 2017, 464, 189-194.	1.1	21
27	Revisiting the prevalence of nonclassic congenital adrenal hyperplasia in US Ashkenazi Jews and Caucasians. <i>Genetics in Medicine</i> , 2017, 19, 1276-1279.	2.4	90
28	Congenital adrenal hyperplasia. <i>Lancet, The</i> , 2017, 390, 2194-2210.	13.7	534
29	Non-classic congenital adrenal hyperplasia due to 21-hydroxylase deficiency revisited: an update with a special focus on adolescent and adult women. <i>Human Reproduction Update</i> , 2017, 23, 580-599.	10.8	136
30	Clinical, biochemical and genetic features with nonclassical 21-hydroxylase deficiency and final height. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2017, 30, 759-766.	0.9	16
31	Gender Reversal And Bilateral Giant Adrenal Myelolipomas In A 46,XX Patient With 21-Hydroxylase Deficiency: Case Report And Review Of The Literature. <i>AACE Clinical Case Reports</i> , 2017, 3, e217-e224.	1.1	4
33	Genetics and Pathophysiology of Congenital Adrenal Hyperplasia. <i>Contemporary Endocrinology</i> , 2018, , 109-127.	0.1	0
34	Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society* Clinical Practice Guideline. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 4043-4088.	3.6	667
35	An overview of inborn errors of metabolism manifesting with primary adrenal insufficiency. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2018, 19, 53-67.	5.7	13
36	Longitudinal Assessment of Illnesses, Stress Dosing, and Illness Sequelae in Patients With Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 2336-2345.	3.6	51
37	Endocrine Disturbances Affecting Reproduction. , 2019, , 594-608.e5.		1
38	Polycystic Ovary Syndrome and NC-CAH: Distinct Characteristics and Common Findings. A Systematic Review. <i>Frontiers in Endocrinology</i> , 2019, 10, 388.	3.5	36
40	Management of the Female With Non-classical Congenital Adrenal Hyperplasia (NCCAH): A Patient-Oriented Approach. <i>Frontiers in Endocrinology</i> , 2019, 10, 366.	3.5	36
41	Cortisol and Aldosterone Responses to Hypoglycemia and Na Depletion in Women With Non-Classic 21-Hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 55-64.	3.6	7
42	Clinical and hormonal characteristics in heterozygote carriers of congenital adrenal hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2020, 198, 105554.	2.5	15
43	Molecular diagnosis of patients with congenital adrenal hyperplasia due to 21-hydroxylase deficiency. <i>BMC Endocrine Disorders</i> , 2020, 20, 165.	2.2	6

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44	Growth Trajectory and Adult Height in Children with Nonclassical Congenital Adrenal Hyperplasia. <i>Hormone Research in Paediatrics</i> , 2020, 93, 173-181.	1.8	12
45	Screening for Nonclassic Congenital Adrenal Hyperplasia in the Era of Liquid Chromatography-Tandem Mass Spectrometry. <i>Journal of the Endocrine Society</i> , 2020, 4, bvz030.	0.2	6
46	EMQN best practice guidelines for molecular genetic testing and reporting of 21-hydroxylase deficiency. <i>European Journal of Human Genetics</i> , 2020, 28, 1341-1367.	2.8	50
47	Genetic characterization of a large cohort of Argentine 21-hydroxylase Deficiency. <i>Clinical Endocrinology</i> , 2020, 93, 19-27.	2.4	3
48	Genotype-Phenotype Correlation in Patients with Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency in Cuba. <i>International Journal of Endocrinology</i> , 2021, 2021, 1-6.	1.5	3
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51	Targeted gene panel sequencing for molecular diagnosis of congenital adrenal hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2021, 211, 105899.	2.5	12
52	Diagnostic Challenges in Nonclassical Congenital Adrenal Hyperplasia. , 2021, , 53-61.		0
53	Russian clinical practice guidelines «congenital adrenal hyperplasia». <i>Obesity and Metabolism</i> , 2021, 18, 345-382.	1.2	5
54	Adrenal Gland Diseases. , 2017, , 195-214.		0
56	17-Hydroxyprogesterone Response to Standard Dose Synacthene Stimulation Test in CYP21A2 Heterozygous Carriers and Non-Carriers in Symptomatic and Asymptomatic Groups: Meta-Analyses. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021, .	0.9	0
57	Nebennierentumoren mit Androgen-Ä–strogen-Äœberproduktion. <i>Springer Reference Medizin</i> , 2021, , 1-7.	0.0	0
58	Transition in Endocrinology. , 2022, , 1227-1246.		0
59	Genetics of Congenital Adrenal Hyperplasia. , 2022, , 932-941.		0
60	17-Hydroxyprogesterone Response to Standard Dose Synacthen Stimulation Test in CYP21A2 Heterozygous Carriers and Non-carriers in Symptomatic and Asymptomatic Groups: Meta-analyses. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2022, 14, 56-68.	0.9	0
61	Genetic analysis and novel variation identification in Chinese patients with congenital adrenal hyperplasia due to 21-hydroxylase deficiency. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2022, 222, 106156.	2.5	2
62	Disorders of the adrenal cortex: Genetic and molecular aspects. <i>Frontiers in Endocrinology</i> , 0, 13, .	3.5	7

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63	Hypogonadismus, InfertilitÄt und sexuelle Dysfunktion bei systemischen Erkrankungen. Springer Reference Medizin, 2021, , 1-43.	0.0	0
64	Nebennierentumoren mit Androgen-/Ä–strogen-Äœberproduktion. Springer Reference Medizin, 2023, , 385-391.	0.0	0
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68	Hyperandrogenism and Its Possible Effects on Endometrial Receptivity: A Review. International Journal of Molecular Sciences, 2023, 24, 12026.	4.1	3
69	Testicular Dysfunction in Systemic Diseases. , 2023, , 503-542.		1