Phenotype and Molecular Characterizations of 30 Child Mutations

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Citation Report

#	Article	IF	Citations
1	NR5A1 Gene Variants: Variable Phenotypes, New Variants, Different Outcomes. Sexual Development, 2019, 13, 258-263.	1.1	8
2	Mutation update for theNR5A1gene involved in DSD and infertility. Human Mutation, 2020, 41, 58-68.	1.1	52
3	Disorders of sex development. , 2020, , 107-119.		0
4	A non-surgical approach to 46,XY differences in sex development through hormonal suppression at puberty: a single-center case series study. Endocrine, 2020, 70, 170-177.	1.1	4
5	Variants of STAR, AMH and ZFPM2/FOG2 May Contribute towards the Broad Phenotype Observed in 46,XY DSD Patients with Heterozygous Variants of NR5A1. International Journal of Molecular Sciences, 2020, 21, 8554.	1.8	9
6	Identification and functional analysis of fourteen NR5A1 variants in patients with the 46 XY disorders of sex development. Gene, 2020, 760, 145004.	1.0	7
7	NR5A1 c.991â€1GÂ>ÂC spliceâ€site variant causes familial 46,XY partial gonadal dysgenesis with incomplete penetrance. Clinical Endocrinology, 2021, 94, 656-666.	1.2	9
8	Approach to the Newborn with Disorders of Sex Development. , 0, , .		1
9	Epididymis cell atlas in a patient with a sex development disorder and a novel NR5A1 gene mutation. Asian Journal of Andrology, 2022, .	0.8	0
10	Case Report: Severe Gonadal Dysgenesis Causing 46,XY Disorder of Sex Development Due to a Novel NR5A1 Variant. Frontiers in Genetics, 0, 13, .	1.1	2
11	Next-generation sequencing of 500 POI patients identified novel responsible monogenic and oligogenic variants. Journal of Ovarian Research, 2023, 16, .	1.3	4
12	A Novel NR5A1 Mutation in a Thai Boy with 46, XY DSD. Journal of Pediatric Genetics, 0, , .	0.3	0
13	Nuclear Receptor Gene Variants Underlying Disorders/Differences of Sex Development through Abnormal Testicular Development. Biomolecules, 2023, 13, 691.	1.8	2