Hepatocyte Nuclear Factor-4 Alfa Mutation Associated v
Hypoglycaemia and Atypical Renal Fanconi Syndrome:

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Citation Report

#	Article	IF	Citations
1	Fainting Fanconi syndrome clarified by proxy: a case report. BMC Nephrology, 2017, 18, 230.	0.8	12
2	Clinical heterogeneity of hyperinsulinism due to HNF1A and HNF4A mutations. Pediatric Diabetes, 2018, 19, 910-916.	1.2	23
3	Salt-Losing Tubulopathies in Children: Whatâ€™s New, Whatâ€™s Controversial?. Journal of the American Society of Nephrology: JASN, 2018, 29, 727-739.	3.0	57
4	HNF4A-related Fanconi syndrome in a Chinese patient: a case report and review of the literature. Journal of Medical Case Reports, 2018, 12, 203.	0.4	15
5	Challenging diagnosis of congenital hyperinsulinism in two infants of diabetic mothers with rare pathogenic KCNJ11 and HNF4A gene variants. International Journal of Pediatric Endocrinology (Springer), 2018, 2018, 5.	1.6	1
6	Congenital hyperinsulinism disorders: Genetic and clinical characteristics. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2019, 181, 682-692.	0.7	49
7	A Complicated Pregnancy in an Adult with HNF4A p.R63W-Associated Fanconi Syndrome. Case Reports in Medicine, 2019, 2019, 1-5.	0.3	4
8	Molecular Basis for Autosomal-Dominant Renal Fanconi Syndrome Caused by HNF4A. Cell Reports, 2019, 29, 4407-4421.e5.	2.9	31
10	Diazoxide-Responsive Forms of Congenital Hyperinsulinism. Contemporary Endocrinology, 2019, , 15-32.	0.3	1
11	Helping nephrologists find answers: hyperinsulinism and tubular dysfunction: Answers. Pediatric Nephrology, 2020, 35, 257-260.	0.9	0
12	Heterozygous recurrent <scp>HNF4A</scp> variant p.<scp>Arg85Trp</scp> causes Fanconi renotubular syndrome 4 with maturity onset diabetes of the young, an autosomal dominant phenocopy of Fanconi Bickel syndrome with colobomas. American Journal of Medical Genetics, Part A, 2021, 185, 566-570.	0.7	7
13	Novel Fanconi renotubular syndromes provide insights in proximal tubule pathophysiology. American Journal of Physiology - Renal Physiology, 2021, 320, F145-F160.	1.3	21
14	Hypoglycemia in the Newborn and Infant. , 2021, , 175-201.		2
15	Hyperinsulinaemic hypoglycaemia, renal Fanconi syndrome and liver disease due to a mutation in the HNF4A gene. Endocrinology, Diabetes and Metabolism Case Reports, 2017, 2017, .	0.2	13
16	Exceptional diazoxide sensitivity in hyperinsulinaemic hypoglycaemia due to a novel HNF4A mutation. Endocrinology, Diabetes and Metabolism Case Reports, 2019, 2019, .	0.2	3
17	Role of hepatocyte nuclear factor 4-alpha in gastrointestinal and liver diseases. World Journal of Gastroenterology, 2019, 25, 4074-4091.	1.4	55
18	HNF4A defines tissue-specific circadian rhythms by beaconing BMAL1::CLOCK chromatin binding and shaping theÂrhythmic chromatin landscape. Nature Communications, 2021, 12, 6350.	5.8	38
19	Transient congenital hyperinsulinism and hemolytic disease of a newborn despite rhesus D prophylaxis: a case report. Journal of Medical Case Reports, 2021, 15, 573.	0.4	0

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20	Hyperechoic Content of the Fetal Colon Is Not Always Cystinuriaâ€"Case Report. Frontiers in Pediatrics, 2021, 9, 822114.	0.9	0
21	Angiomotin mutation causes glomerulopathy and renal cysts by upregulating hepatocyte nuclear factor transcriptional activity. Clinical and Translational Medicine, 2022, 12, .	1.7	2
22	Hepatocyte nuclear factors play an important role in the pathogenesis of nephropathy. Clinical and Translational Discovery, 2022, 2, .	0.2	0
23	Renal Fanconi Syndromes and Other Proximal Tubular Disorders. , 2023, , 967-990.		0
24	Expanding the p.(Arg85Trp) Variant-Specific Phenotype of HNF4A: Features of Glycogen Storage Disease, Liver Cirrhosis, Impaired Mitochondrial Function, and Glomerular Changes. Molecular Syndromology, 2023, 14, 347-362.	0.3	1