Gene mutations and clinical phenotypes in Chinese chil

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Citation Report

#	Article	IF	Citations
1	eRAM: encyclopedia of rare disease annotations for precision medicine. Nucleic Acids Research, 2018, 46, D937-D943.	6.5	56
2	A Case of Sporadic Blau Syndrome with an Uncommon Clinical Course. Case Reports in Rheumatology, 2018, 2018, 1-5.	0.2	10
3	PedAM: a database for Pediatric Disease Annotation and Medicine. Nucleic Acids Research, 2018, 46, D977-D983.	6.5	27
4	Application of genome analysis strategies in the clinical testing for pediatric diseases. Pediatric Investigation, 2018, 2, 72-81.	0.6	29
5	Genotypeâ€“Phenotype Association Analysis Reveals New Pathogenic Factors for Osteogenesis Imperfecta Disease. Frontiers in Pharmacology, 2019, 10, 1200.	1.6	11
6	Systematically Analyzing the Pathogenic Variations for Acute Intermittent Porphyria. Frontiers in Pharmacology, 2019, 10, 1018.	1.6	14
7	Hereditary systemic autoinflammatory diseases and Schnitzlerâ€™s syndrome. Rheumatology, 2019, 58, vi31-vi43.	0.9	21
8	Whole-exome sequencing in three children with sporadic Blau syndrome, one of them co-presenting with recurrent polyserositis. Autoimmunity, 2020, 53, 344-352.	1.2	2
9	A young female with early onset arthritis, uveitis, hepatic, and renal granulomas: a clinical tryst with Blau syndrome over 20Âyears and case-based review. Rheumatology International, 2021, 41, 173-181.	1.5	22
10	Genetic Basis of Myocarditis: Myth or Reality?. , 2020, , 45-89.		15
11	Recent advances in the epidemiology and genetics of acute intermittent porphyria. Intractable and Rare Diseases Research, 2020, 9, 196-204.	0.3	13
12	Blau Syndrome Complicated by Atypical Type IIa Takayasu Arteritis. Journal of Child Science, 2021, 11, e313-e316.	0.1	1
13	Palmitoylation restricts SQSTM1/p62-mediated autophagic degradation of NOD2 to modulate inflammation. Cell Death and Differentiation, 2022, 29, 1541-1551.	5.0	20
14	Genetic and Clinical Features of Blau Syndrome among Chinese Patients with Uveitis. Ophthalmology, 2022, 129, 821-828.	2.5	5
15	Progress in the genetics of uveitis. Genes and Immunity, 2022, 23, 57-65.	2.2	17
25	Kidney Involvement in Autoinflammatory Diseases. Kidney Diseases (Basel, Switzerland), 2023, 9, 157-172.	1.2	1
26	A Chinese girl of Blau syndrome with renal arteritis and a literature review. Pediatric Rheumatology, 2023, 21, .	0.9	1