

Zhenping Chen

List of Publications by Year in descending order

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papers

460
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840119

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#	ARTICLE	IF	CITATIONS
1	Single Nucleotide Polymorphisms of the HIF1A Gene are Associated With Sensitivity of Glucocorticoid Treatment in Pediatric ITP Patients. <i>Journal of Pediatric Hematology/Oncology</i> , 2023, 45, 195-199.	0.3	2
2	Enhanced pharmacokinetics and reduced bleeds in boys with hemophilia A after switching to Kovaltry from other standard half-life factor VIII concentrates. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12686.	1.0	4
3	Inter-individual variability in pharmacokinetics and clinical features in pediatric patients with severe hemophilia A. <i>Thrombosis Research</i> , 2022, 213, 71-77.	0.8	5
4	Nephrotic syndrome in two haemophilia B children with inhibitor under low-dose immune tolerance induction combined with rituximab-based immunosuppressant protocol. <i>Haemophilia</i> , 2022, 28, .	1.0	1
5	Eradication of FIX inhibitor in haemophilia B children using low-dose immune tolerance induction with rituximab-based immunosuppressive agent(s) in China. <i>Haemophilia</i> , 2022, , .	1.0	3
6	F8 gene mutation spectrum in severe hemophilia A with inhibitors: A large cohort data analysis from a single center in China. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12723.	1.0	3
7	Individualised prophylaxis based on personalised target trough FVIII level optimised clinical outcomes in paediatric patients with severe haemophilia A. <i>Haemophilia</i> , 2022, 28, .	1.0	7
8	A low-dose immune tolerance induction (ITI) protocol incorporating immunosuppressive agents in haemophilia A children with high-titre factor VIII inhibitor and poor ITI prognostic risk. <i>Haemophilia</i> , 2021, 27, e469-e472.	1.0	2
9	Comparative pharmacokinetics of Kogenate FS and Kovaltry in 14 Chinese paediatric patients with haemophilia A: A single-centre study. <i>Haemophilia</i> , 2021, 27, e287-e290.	1.0	2
10	Sirolimus is effective in autoimmune lymphoproliferative syndrome-type III: A pedigree case report with homozygous variation PRKCD. <i>International Journal of Immunopathology and Pharmacology</i> , 2021, 35, 205873842110259.	1.0	5
11	A novel mutation in GP1BA gene in a family with autosomal dominant Bernard Soulier syndrome variant: A case report. <i>Experimental and Therapeutic Medicine</i> , 2021, 21, 360.	0.8	2
12	Pharmacokinetic variability of factor VIII concentrates in Chinese pediatric patients with moderate or severe hemophilia A. <i>Pediatric Investigation</i> , 2021, 5, 38-45.	0.6	7
13	Pharmacokinetic study of Kovaltry in thirty-five pediatric patients aged <12 years with severe hemophilia A. <i>Haemophilia</i> , 2021, 27, e340-e346.	1.0	0
14	Pharmacokinetic-guided prophylaxis improved clinical outcomes in paediatric patients with severe haemophilia A. <i>Haemophilia</i> , 2021, 27, e450-e457.	1.0	10
15	Pharmacokinetics and complementary evaluation system-based guidance on prophylaxis of paediatric patients with haemophilia A in China with Kovaltry: protocol of the LEAP study. <i>BMJ Open</i> , 2021, 11, e048432.	0.8	1
16	Low-dose immune tolerance induction alone or with immunosuppressants according to prognostic risk factors in Chinese children with hemophilia A inhibitors. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12562.	1.0	6
17	Cost-effectiveness Analysis of Prophylaxis Versus On-demand Treatment for Children With Hemophilia B Without Inhibitors in China. <i>Clinical Therapeutics</i> , 2021, 43, 1536-1546.	1.1	1
18	Significant reduction in hemarthrosis in boys with severe hemophilia A: The China hemophilia individualized low-dose secondary prophylaxis study. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12552.	1.0	5

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19	Spotlight on eltrombopag in pediatric ITP in China: a long-term observational study in real-world practice. <i>Blood Advances</i> , 2021, 5, 3799-3806.	2.5	12
20	Detection of Drug-Induced Thrombocytopenia Signals in Children Using Routine Electronic Medical Records. <i>Frontiers in Pharmacology</i> , 2021, 12, 756207.	1.6	4
21	The Association of Circulating T Follicular Helper Cells and Regulatory Cells with Acute Myeloid Leukemia Patients. <i>Acta Haematologica</i> , 2020, 143, 19-25.	0.7	3
22	Screening for Genetic Mutations for the Early Diagnosis of Common Variable Immunodeficiency in Children With Refractory Immune Thrombocytopenia: A Retrospective Data Analysis From a Tertiary Children's Center. <i>Frontiers in Pediatrics</i> , 2020, 8, 595135.	0.9	7
23	Maternal microchimerism protects hemophilia A patients from inhibitor development. <i>Blood Advances</i> , 2020, 4, 1867-1869.	2.5	3
24	Bleeds and imaging scoring scales in relation to pharmacokinetics of coagulation factor VIII in Chinese pediatric patients with severe hemophilia A. <i>Thrombosis Research</i> , 2020, 193, 83-85.	0.8	1
25	A previously treated severe haemophilia A patient developed high-titre inhibitor after vaccinations. <i>International Journal of Immunopathology and Pharmacology</i> , 2020, 34, 205873842093461.	1.0	3
26	Case report of a novel MPIG6B gene mutation in a Chinese boy with pancytopenia and splenomegaly. <i>Gene</i> , 2019, 715, 143957.	1.0	9
27	Low-dose immune tolerance induction for children with hemophilia A with poor-risk high-titer inhibitors: A pilot study in China. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2019, 3, 741-748.	1.0	11
28	Efficacy and safety of eltrombopag in the treatment of severe chronic immune thrombocytopenia in children of China: A single-center observational study. <i>International Journal of Immunopathology and Pharmacology</i> , 2019, 33, 205873841987212.	1.0	7
29	Low-Dose Immune Tolerance Induction for Hemophilia a Children with Poor-Risk High-Titer Inhibitors. <i>Blood</i> , 2019, 134, 1122-1122.	0.6	0
30	Application of High-Throughput Sequencing in the Diagnosis of Inherited Immune-Thrombocytopenia from Children Chronic/Refractory ITP. <i>Blood</i> , 2019, 134, 86-86.	0.6	0
31	Break-through bleeding in relation to pharmacokinetics of Factor VIII in paediatric patients with severe haemophilia A. <i>Haemophilia</i> , 2018, 24, 120-125.	1.0	3
32	Synergistic defects of novo FAS and homozygous UNC13D leading to autoimmune lymphoproliferative syndrome-like disease: A 10-year-old Chinese boy case report. <i>Gene</i> , 2018, 672, 45-49.	1.0	3
33	STAT1 single nucleotide polymorphisms and susceptibility to immune thrombocytopenia. <i>Autoimmunity</i> , 2015, 48, 305-312.	1.2	9
34	Foxp3 methylation status in children with primary immune thrombocytopenia. <i>Human Immunology</i> , 2014, 75, 1115-1119.	1.2	17
35	An Fc γ RIIb transmembrane polymorphism in Chinese ITP patients. <i>Platelets</i> , 2010, 21, 479-485.	1.1	8
36	Th1 (CXCL10) and Th2 (CCL2) chemokine expression in patients with immune thrombocytopenia. <i>Human Immunology</i> , 2010, 71, 586-591.	1.2	30

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37	BAFF and BAFF-R of peripheral blood and spleen mononuclear cells in idiopathic thrombocytopenic purpura. <i>Autoimmunity</i> , 2009, 42, 112-119.	1.2	41
38	Fetal BM-derived mesenchymal stem cells promote the expansion of human Th17 cells, but inhibit the production of Th1 cells. <i>European Journal of Immunology</i> , 2009, 39, 2840-2849.	1.6	63
39	Raised expression of APRIL in Chinese patients with immune thrombocytopenia and its clinical implications. <i>Autoimmunity</i> , 2009, 42, 692-698.	1.2	17
40	CD72 Polymorphism Associated with Child-Onset of Idiopathic Thrombocytopenic Purpura in Chinese Patients. <i>Journal of Clinical Immunology</i> , 2008, 28, 214-219.	2.0	14
41	Single Nucleotide Polymorphism in DNMT3B Promoter and the Risk for Idiopathic Thrombocytopenic Purpura in Chinese Population. <i>Journal of Clinical Immunology</i> , 2008, 28, 399-404.	2.0	18
42	Decreased DNA Methyltransferase 3A and 3B mRNA Expression in Peripheral Blood Mononuclear Cells and Increased Plasma SAH Concentration in Adult Patients with Idiopathic Thrombocytopenic Purpura. <i>Journal of Clinical Immunology</i> , 2008, 28, 432-439.	2.0	33
43	Health-related quality of life measured by the Short Form 36 in immune thrombocytopenic purpura: a cross-sectional survey in China. <i>European Journal of Haematology</i> , 2007, 78, 518-523.	1.1	53
44	Interferon- γ +874A/T and interleukin-4 intron3 VNTR gene polymorphisms in Chinese patients with idiopathic thrombocytopenic purpura. <i>European Journal of Haematology</i> , 2007, 79, 191-197.	1.1	25