

# Renchi Yang

## List of Publications by Year in descending order

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95  
papers

1,715  
citations

331670

21  
h-index

361022

35  
g-index

96  
all docs

96  
docs citations

96  
times ranked

1599  
citing authors

#	ARTICLE	IF	CITATIONS
1	Efficacy and safety of eltrombopag in Chinese patients with chronic immune thrombocytopenia: stage 2 results from a multicenter phase III study. <i>Platelets</i> , 2022, 33, 82-88.	2.3	17
2	Dose tapering to withdrawal stage and long-term efficacy and safety of hetrombopag for the treatment of immune thrombocytopenia: Results from an open-label extension study. <i>Journal of Thrombosis and Haemostasis</i> , 2022, 20, 716-728.	3.8	6
3	Mapping the Haem-A-QoL to the EQ-5D-5L in patients with hemophilia. <i>Quality of Life Research</i> , 2022, 31, 1533-1544.	3.1	7
4	Demographics, clinical profile and treatment landscape of patients with haemophilia B in China. <i>Haemophilia</i> , 2022, 28, .	2.1	3
5	Prophylactic emicizumab for hemophilia A in the Asia-Pacific region: A randomized study (HAVEN 5). <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12670.	2.3	24
6	Pharmacokinetics, Safety, and Pharmacodynamics of Romiplostim in Chinese Subjects With Immune Thrombocytopenia: A Phase I/II Trial. <i>Clinical Pharmacology in Drug Development</i> , 2022, 11, 379-387.	1.6	2
7	Inhibitors: Diagnostic challenges, unknowns of inhibitor development, treatment of bleeding and surgery, and insights into diagnosis and treatment in China. <i>Haemophilia</i> , 2022, 28, 111-118.	2.1	0
8	Health-related quality of life among adults with haemophilia in China: A comparison with age-matched general population. <i>Haemophilia</i> , 2022, 28, 776-783.	2.1	2
9	Efficacy, safety and pharmacokinetics of recombinant human coagulation factor VIII (omifloctocog) Tj ETQq1 1 0.784314 rgBJT /Overlo	2.1	2
10	An overview of patients with haemophilia A in China: Epidemiology, disease severity and treatment strategies. <i>Haemophilia</i> , 2021, 27, e51-e59.	2.1	15
11	A multicenter, randomized phase III trial of hetrombopag: a novel thrombopoietin receptor agonist for the treatment of immune thrombocytopenia. <i>Journal of Hematology and Oncology</i> , 2021, 14, 37.	17.0	33
12	Patients with haemophilia A with inhibitors in China: a national real-world analysis and follow-up. <i>British Journal of Haematology</i> , 2021, 192, 900-908.	2.5	7
13	Association of <i>FOXP3</i> gene polymorphisms with chronic immune thrombocytopenia in a Chinese Han population. <i>International Journal of Laboratory Hematology</i> , 2021, 43, 1104-1109.	1.3	6
14	Evaluating the psychometric properties of the EQ-5D-5L and SF-6D among patients with haemophilia. <i>European Journal of Health Economics</i> , 2021, 22, 547-557.	2.8	18
15	First open-label, single-arm, prospective study of real-world use of FIX replacement therapy in a predominantly pediatric hemophilia B population in China. <i>Medicine (United States)</i> , 2021, 100, e26077.	1.0	0
16	Pharmacokinetic, efficacy and safety evaluation of B-domain-deleted recombinant FVIII (SCT800) for prophylactic treatment in adolescent and adult patients with severe haemophilia A. <i>Haemophilia</i> , 2021, 27, 814-822.	2.1	6
17	Current status of haemophilia inhibitor management in mainland China: a haemophilia treatment centres survey on treatment preferences and real-world clinical practices. <i>British Journal of Haematology</i> , 2021, 194, 750-758.	2.5	2
18	Investigating the Added Value of the EQ-5D-5L With Two Bolt-On Items in Patients With Hemophilia. <i>Frontiers in Medicine</i> , 2021, 8, 707998.	2.6	3

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19	Phase 4 Safety and Efficacy Study of Antihemophilic Factor (Recombinant) in Previously Treated Chinese Patients With Severe/Moderately Severe Hemophilia A. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2021, 27, 107602962198981.	1.7	2
20	A novel SERPINC1 frameshift mutation in two antithrombin deficiency families. <i>International Journal of Laboratory Hematology</i> , 2020, 42, e48-e51.	1.3	2
21	Role of bone marrow-derived mesenchymal stem cell defects in CD8 <sup>+</sup> CD28 <sup>+</sup> suppressor T lymphocyte induction in patients with immune thrombocytopenia and associated mechanisms. <i>British Journal of Haematology</i> , 2020, 191, 852-862.	2.5	14
22	Safety and Efficacy of Moroctocog Alfa (AF-CC) in Chinese Patients with Hemophilia A: Results of Two Open-Label Studies. <i>Journal of Blood Medicine</i> , 2020, Volume 11, 439-448.	1.7	1
23	Haemophilia care in China: Achievements in the past decade. <i>Haemophilia</i> , 2020, 26, 759-767.	2.1	20
24	Management of haemophilia patients in the COVID-19 pandemic: Experience in Wuhan and Tianjin, two differently affected cities in China. <i>Haemophilia</i> , 2020, 26, 1031-1037.	2.1	4
25	CD70-silenced dendritic cells induce immune tolerance in immune thrombocytopenia patients. <i>British Journal of Haematology</i> , 2020, 191, 466-475.	2.5	7
26	Real-world analysis of haemophilia patients in China: A single centre's experience. <i>Haemophilia</i> , 2020, 26, 584-590.	2.1	10
27	Molecular analysis in 12 factor XI deficiency patients from China: Identification of three novel splicing mutations. <i>Thrombosis Research</i> , 2020, 192, 100-102.	1.7	2
28	Integrin $\alpha$ 23 Deficiency Results in Hypertriglyceridemia via Disrupting LPL (Lipoprotein Lipase) Secretion. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2020, 40, 1296-1310.	2.4	4
29	Levels of Soluble CD30 and CD26 and Their Clinical Significance in Patients with Primary Immune Thrombocytopenia. <i>BioMed Research International</i> , 2020, 2020, 1-8.	1.9	0
30	Outcome of CARE: a 6-year national registry of acquired haemophilia A in China. <i>British Journal of Haematology</i> , 2019, 187, 653-665.	2.5	28
31	Carbapenem-resistant Enterobacteriaceae in hematological patients: Outcome of patients with Carbapenem-resistant Enterobacteriaceae infection and risk factors for progression to infection after rectal colonization. <i>International Journal of Antimicrobial Agents</i> , 2019, 54, 527-529.	2.5	24
32	Current and emerging treatments for immune thrombocytopenia. <i>Expert Review of Hematology</i> , 2019, 12, 723-732.	2.2	13
33	IP-10 and MCP-1 gene polymorphisms in Chinese patients with chronic immune thrombocytopenia. <i>Autoimmunity</i> , 2019, 52, 235-241.	2.6	0
34	Health-related quality of life and health status in persons with haemophilia A with inhibitors: A prospective, multicentre, non-interventional study (NIS). <i>Haemophilia</i> , 2019, 25, 382-391.	2.1	28
35	Efficacy and safety of prophylaxis with BAY 81-8973 in Chinese patients with severe haemophilia A enrolled in the LEOPOLD II trial. <i>Haemophilia</i> , 2019, 25, e153-e158.	2.1	7
36	Building a network for hemophilia care in China: 15 years of achievement for the Hemophilia Treatment Center Collaborative Network of China. <i>Blood Advances</i> , 2019, 3, 34-37.	5.2	16

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37	Mutation profiling by targeted sequencing of "triple-negative" essential thrombocythaemia patients. <i>British Journal of Haematology</i> , 2018, 181, 857-860.	2.5	8
38	Reduced PTEN involved in primary immune thrombocytopenia via contributing to B cell hyper-responsiveness. <i>Molecular Immunology</i> , 2018, 93, 144-151.	2.2	8
39	Bleeding events and safety outcomes in persons with haemophilia A with inhibitors: A prospective, multi-centre, non-interventional study. <i>Haemophilia</i> , 2018, 24, 921-929.	2.1	20
40	Abnormal Distribution and Function of Monocyte Subsets in Patients With Primary Immune Thrombocytopenia. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 786-792.	1.7	11
41	Upregulation of CD72 expression on CD19 <sup>+</sup> CD27 <sup>+</sup> memory B cells by CD40L in primary immune thrombocytopenia. <i>British Journal of Haematology</i> , 2017, 178, 308-318.	2.5	13
42	The State of Skewed X Chromosome Inactivation is Retained in the Induced Pluripotent Stem Cells from a Female with Hemophilia B. <i>Stem Cells and Development</i> , 2017, 26, 1003-1011.	2.1	3
43	Decreased TLR4 expression on monocytes may cause regulatory T cells abnormality in patients with primary immune thrombocytopenia. <i>Autoimmunity</i> , 2017, 50, 283-292.	2.6	9
44	Numerical and functional defects in CD8 <sup>+</sup> CD28 <sup>~</sup> T suppressor lymphocytes from patients with primary immune thrombocytopenia. <i>British Journal of Haematology</i> , 2017, 178, 292-301.	2.5	9
45	Status and trend analysis of prophylactic usage of recombinant factor VIII in Chinese pediatric patients with hemophilia A: ReCare " a retrospective, phase IV, non-interventional study. <i>Current Medical Research and Opinion</i> , 2017, 33, 1571-1578.	1.9	1
46	Long-term efficacy and safety of prophylaxis with recombinant factor VIII in Chinese pediatric patients with hemophilia A: a multi-center, retrospective, non-interventional, phase IV (ReCARE) study. <i>Current Medical Research and Opinion</i> , 2017, 33, 1223-1230.	1.9	9
47	The Efficacy of Recombinant FVIII Low-Dose Prophylaxis in Chinese Pediatric Patients With Severe Hemophilia A: A Retrospective Analysis From the ReCARE Study. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 851-858.	1.7	14
48	Increased plasma sCXCL16 levels may have a relationship with Th1/Th2 imbalance in primary immune thrombocytopenia. <i>Cytokine</i> , 2017, 99, 124-131.	3.2	5
49	Efficacy of standard prophylaxis versus on-demand treatment with bayer's sucrose-formulated recombinant FVIII (rFVIII-FS) in Chinese children with severe hemophilia A. <i>Pediatric Hematology and Oncology</i> , 2017, 34, 146-156.	0.8	12
50	Multicentre, randomised phase III study of the efficacy and safety of eltrombopag in Chinese patients with chronic immune thrombocytopenia. <i>British Journal of Haematology</i> , 2017, 176, 101-110.	2.5	55
51	Long-term results of splenectomy in adult chronic immune thrombocytopenia. <i>European Journal of Haematology</i> , 2017, 98, 235-241.	2.2	36
52	Aberrant expression of long noncoding RNA TMEVPG1 in patients with primary immune thrombocytopenia. <i>Autoimmunity</i> , 2016, 49, 496-502.	2.6	20
53	Effects of the multidrug resistance-1 gene on drug resistance in primary immune thrombocytopenia. <i>Autoimmunity</i> , 2016, 49, 486-495.	2.6	5
54	Different dosages of intravenous immunoglobulin (IVIg) in treating immune thrombocytopenia with long-term follow-up of three years: Results of a prospective study including 167 cases. <i>Autoimmunity</i> , 2016, 49, 50-57.	2.6	13

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55	The Effect of Danazol in Primary Immune Thrombocytopenia. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2016, 22, 727-733.	1.7	31
56	Stromal cell-derived factor-1 rs2297630 polymorphism associated with platelet production and treatment response in Chinese patients with chronic immune thrombocytopenia. <i>Platelets</i> , 2016, 27, 338-343.	2.3	9
57	The polymorphisms of tumor necrosis factor-induced protein 3 gene may contribute to the susceptibility of chronic primary immune thrombocytopenia in Chinese population. <i>Platelets</i> , 2016, 27, 26-31.	2.3	11
58	Abnormal lipid rafts related ganglioside expression and signaling in T lymphocytes in immune thrombocytopenia patients. <i>Autoimmunity</i> , 2016, 49, 58-68.	2.6	0
59	Acquired Hemophilia A. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2015, 21, 35-40.	1.7	19
60	Interleukin 35 may contribute to the loss of immunological self-tolerance in patients with primary immune thrombocytopenia. <i>British Journal of Haematology</i> , 2015, 169, 278-285.	2.5	19
61	Prophylaxis vs. on-demand treatment with BAY 81-8973, a full-length plasma protein-free recombinant factor VIII product: results from a randomized trial (LEOPOLD II). <i>Journal of Thrombosis and Haemostasis</i> , 2015, 13, 360-369.	3.8	74
62	Elevated Semaphorin 5A correlated with Th1 polarization in patients with chronic immune thrombocytopenia. <i>Thrombosis Research</i> , 2015, 136, 859-864.	1.7	19
63	Interleukin-7 is decreased and maybe plays a pro-inflammatory function in primary immune thrombocytopenia. <i>Platelets</i> , 2015, 26, 243-249.	2.3	7
64	Reduced <i>MIR130A</i> is involved in primary immune thrombocytopenia via targeting <i>TGFB1</i> and <i>IL18</i> . <i>British Journal of Haematology</i> , 2014, 166, 767-773.	2.5	19
65	Absence of Association of <i>Interleukin-18</i> Gene Polymorphisms with Primary Immune Thrombocytopenia in a Chinese Han Population. <i>DNA and Cell Biology</i> , 2014, 33, 537-542.	1.9	4
66	Retrospective analysis of 1,226 Chinese patients with haemophilia in a single medical centre. <i>Journal of Thrombosis and Thrombolysis</i> , 2014, 38, 92-97.	2.1	8
67	The defective bone marrow-derived mesenchymal stem cells in patients with chronic immune thrombocytopenia. <i>Autoimmunity</i> , 2014, 47, 519-529.	2.6	28
68	External validation and clinical evaluation of the International Prognostic Score of Thrombosis for Essential Thrombocythemia (IPSET-thrombosis) in a large cohort of Chinese patients. <i>European Journal of Haematology</i> , 2014, 92, 502-509.	2.2	24
69	Decreased IL-35 levels in patients with immune thrombocytopenia. <i>Human Immunology</i> , 2014, 75, 909-913.	2.4	21
70	The polymorphisms of T cell-specific TBX21 gene may contribute to the susceptibility of chronic immune thrombocytopenia in Chinese population. <i>Human Immunology</i> , 2014, 75, 129-133.	2.4	11
71	Association of ABCB1 gene polymorphisms and haplotypes with therapeutic efficacy of glucocorticoids in Chinese patients with immune thrombocytopenia. <i>Human Immunology</i> , 2014, 75, 317-321.	2.4	25
72	Effect of Eltrombopag on Platelet Response and Safety Results in Chinese Adults with Chronic ITP-Primary Result of a Phase III Study. <i>Blood</i> , 2014, 124, 1464-1464.	1.4	6

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73	Distinct Molecular Abnormalities Underlie Unique Clinical Features of Essential Thrombocythemia in Children. <i>Blood</i> , 2014, 124, 4579-4579.	1.4	0
74	Integrin $\alpha$ IIb $\beta$ 3-Mediated Outside-in Signaling: Brake or Amplifier of Platelet Activation?. <i>Blood</i> , 2014, 124, 4161-4161.	1.4	0
75	Reduced expression of <i>MIR409</i> in primary immune thrombocytopenia. <i>British Journal of Haematology</i> , 2013, 161, 128-135.	2.5	23
76	DNA Methylation and Primary Immune Thrombocytopenia. <i>Seminars in Hematology</i> , 2013, 50, S116-S126.	3.4	11
77	Paediatric essential thrombocythaemia: clinical and molecular features, diagnosis and treatment. <i>British Journal of Haematology</i> , 2013, 163, 295-302.	2.5	25
78	A multicenter randomized controlled trial of recombinant human thrombopoietin treatment in patients with primary immune thrombocytopenia. <i>International Journal of Hematology</i> , 2012, 96, 222-228.	1.6	77
79	Immunosuppressive function of mesenchymal stem cells from human umbilical cord matrix in immune thrombocytopenia patients. <i>Thrombosis and Haemostasis</i> , 2012, 107, 937-950.	3.4	37
80	Meeting the challenges of haemophilia care and patient support in China and Brazil. <i>Haemophilia</i> , 2012, 18, 33-38.	2.1	23
81	Hemophilic Pseudotumor in Chinese Patients: A Retrospective Single-Centered Analysis of 14 Cases. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2011, 17, 279-282.	1.7	28
82	Thrombopoietic growth factors in the treatment of immune thrombocytopenic purpura. <i>Critical Reviews in Oncology/Hematology</i> , 2011, 77, 172-183.	4.4	15
83	Factor VIII gene mutations profile in 148 Chinese hemophilia A subjects. <i>European Journal of Haematology</i> , 2010, 85, 264-272.	2.2	20
84	An Fc $\gamma$ RIIb transmembrane polymorphism in Chinese ITP patients. <i>Platelets</i> , 2010, 21, 479-485.	2.3	8
85	The expression of IFN- $\gamma$ , IL-4, Foxp3 and perforin genes are not correlated with DNA methylation status in patients with immune thrombocytopenic purpura. <i>Platelets</i> , 2010, 21, 137-143.	2.3	18
86	Th1 (CXCL10) and Th2 (CCL2) chemokine expression in patients with immune thrombocytopenia. <i>Human Immunology</i> , 2010, 71, 586-591.	2.4	30
87	BAFF and BAFF-R of peripheral blood and spleen mononuclear cells in idiopathic thrombocytopenic purpura. <i>Autoimmunity</i> , 2009, 42, 112-119.	2.6	41
88	Rituximab treatment for chronic refractory idiopathic thrombocytopenic purpura. <i>Critical Reviews in Oncology/Hematology</i> , 2008, 65, 21-31.	4.4	28
89	CD72 Polymorphism Associated with Child-Onset of Idiopathic Thrombocytopenic Purpura in Chinese Patients. <i>Journal of Clinical Immunology</i> , 2008, 28, 214-219.	3.8	14
90	Antithrombin, Protein C, Protein S and Activated Protein C Resistance in the General Healthy Chinese Population: Normal Plasmatic Ranges and Genetic Defects.. <i>Blood</i> , 2007, 110, 3982-3982.	1.4	1

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91	Splenectomy for Chronic Idiopathic Thrombocytopenic Purpura in Children: A Single Center Study in China. <i>Acta Haematologica</i> , 2006, 115, 39-45.	1.4	28
92	Abnormality of CD4+CD25+regulatory T cells in idiopathic thrombocytopenic purpura. <i>European Journal of Haematology</i> , 2006, 78, 061213212227003-???	2.2	199
93	Splenectomy for adult chronic idiopathic thrombocytopenic purpura: experience from a single center in China. <i>European Journal of Haematology</i> , 2005, 75, 424-429.	2.2	18
94	Type 1 and type 2 T-cell profiles in idiopathic thrombocytopenic purpura. <i>Haematologica</i> , 2005, 90, 914-23.	3.5	129
95	Angiogenesis in Hematologic Malignancies and Its Clinical Implications. <i>International Journal of Hematology</i> , 2002, 75, 246-256.	1.6	19