Ming-Ching Shen

List of Publications by Year
in descending order

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35	802	840119 11	500791 28
papers	citations	h-index	g-index

35	35	35	769
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A distinct common p.Gln317* mutation among causative LMAN1 genetic mutations of combined factor V and factor VIII deficiency in five Taiwanese families. Haemophilia, 2022, 28, .	1.0	2
2	Deep vein thrombosis after major orthopedic surgery in Taiwan: A prospective cross-sectional study and literature review. Journal of the Formosan Medical Association, 2022, , .	0.8	4
3	Characterization of congenital factor XII deficiency in Taiwanese patients: identification of one novel and one common mutation. International Journal of Hematology, 2022, 116, 528-533.	0.7	1
4	Transfusionâ€transmitted infection and comorbidities in patients with severe haemophilia: A longitudinal birth cohort analysis. Haemophilia, 2021, 27, e458-e461.	1.0	0
5	Acquired FXIII inhibitor: Patient characteristics and treatment outcome, a case series in Taiwan. Journal of the Formosan Medical Association, 2021, 120, 411-414.	0.8	2
6	First reported case of congenital thrombotic thrombocytopenic purpura in Taiwan with novel mutation of ADAMTS13 gene. International Journal of Hematology, 2021, 113, 760-764.	0.7	0
7	Review article inferior vena cava thrombosis: a case series of patients observed in Taiwan and literature review. Thrombosis Journal, 2021, 19, 43.	0.9	8
8	Clinical features and genetic defect in six index patients with congenital fibrinogen disorders: Three novel mutations with one common mutation in Taiwan's population. Haemophilia, 2021, 27, 1022-1027.	1.0	1
9	A family with an MYH9-related disorder with different phenotypes masquerading as immune thrombocytopaenia: an underreported disorder in Taiwan. International Journal of Hematology, 2020, 112, 878-882.	0.7	2
10	Characterization of hereditary factor XI deficiency in Taiwanese patients: identification of three novel and two common mutations. International Journal of Hematology, 2020, 112, 169-175.	0.7	1
11	Segmental uniparental disomy as a rare cause of congenital severe factor XIII deficiency in a girl with only one heterozygous carrier parent. Pediatric Hematology and Oncology, 2018, 35, 442-446.	0.3	2
12	De novo mutation and somatic mosaicism of gene mutation in type 2A, 2B and 2M VWD. Thrombosis Journal, 2016, 14, 36.	0.9	6
13	A study of 65 patients with acquired hemophilia A in Taiwan. Journal of the Formosan Medical Association, 2015, 114, 321-327.	0.8	30
14	Functional characterization of a novel missense mutation, His147Arg, in A1 domain of FV protein causing type II deficiency. Thrombosis Research, 2014, 134, 153-159.	0.8	7
15	Evaluation of performance for automated differential leucocyte counting on Sysmex NE-8000 by NCCLS recommended protocol, H20-T. International Journal of Laboratory Hematology, 2008, 15, 287-299.	0.2	8
16	Pharmacokinetic Study of Recombinant Human Factor IX in Previously Treated Patients with Hemophilia B in Taiwan. Journal of the Formosan Medical Association, 2007, 106, 281-287.	0.8	7
17	A first Taiwanese Chinese family of type 2B von Willebrand disease with R1306W mutation. Thrombosis Research, 2003, 112, 291-295.	0.8	1
18	Methylation of the p15 INK4B gene in myelodysplastic syndrome: it can be detected early at diagnosis or during disease progression and is highly associated with leukaemic transformation. British Journal of Haematology, 2001, 112, 148-154.	1.2	140

#	Article	IF	Citations
19	Novel mutations in the Factor VII gene of Taiwanese Factor VII-deficient patients. British Journal of Haematology, 2001, 112, 566-571.	1.2	9
20	Epitope mapping of factor VIII inhibitor antibodies of Chinese origin. British Journal of Haematology, 2001, 113, 915-924.	1.2	10
21	Identification of a new CA dinucleotide repeat in the human factor VIII gene. British Journal of Haematology, 2000, 111, 1256-1259.	1.2	0
22	A longitudinal study of immunological status in Chinese Haemophiliacs: importance of the heat viral inactivation of factor concentrates. II. Improvements of CD4/CD8 ratio after treatments with heatâ€inactivated factor concentrates. Haemophilia, 1998, 4, 33-40.	1.0	95
23	Human immunodeficiency virus infection in Haemophiliacs in Taiwan: the importance of CD4 lymphocyte count in the progression to acquired immunodeficiency syndrome. Haemophilia, 1998, 4, 115-121.	1.0	1
24	Acute and chronic arsenic poisoning associated with treatment of acute promyelocytic leukaemia. British Journal of Haematology, 1998, 103, 1092-1095.	1.2	120
25	Clonal disease of natural killer large granular lymphocytes in Taiwan. British Journal of Haematology, 1998, 103, 1124-1128.	1.2	14
26	The Mutation at Position 20210 in the 3â€™-Untranslated Region of the Prothrombin Gene Is Extremely Rare in Taiwanese Chinese Patients with venous Thrombophilia. Thrombosis and Haemostasis, 1998, 80, 343-343.	1.8	34
27	HIGH PREVALENCE OF ANTITHROMBIN III, PROTEIN C AND PROTEIN S DEFICIENCY, BUT NO FACTOR V LEIDEN MUTATION IN VENOUS THROMBOPHILIC CHINESE PATIENTS IN TAIWAN. Thrombosis Research, 1997, 87, 377-385.	0.8	99
28	Genetic diagnosis of haemophilia A of Chinese origin. British Journal of Haematology, 1995, 91, 722-727.	1.2	11
29	Consistent presence of isochromosome 7q in hepatosplenic t Î³/Î´ lymphoma: A new cytogenetic-clinicopathologic entity. Genes Chromosomes and Cancer, 1995, 12, 161-164.	1.5	111
30	Analysis of Human Platelet Glycoprotein IIb-IIIa by Fluorescein Isothiocyanate-Conjugated Disintegrins with Flow Cytometry. Thrombosis and Haemostasis, 1994, 72, 919-925.	1.8	23
31	Genetic Basis and Carrier Detection of Hemophilia B of Chinese Origin. Thrombosis and Haemostasis, 1993, 69, 247-252.	1.8	15
32	Hodgkin's disease and nonâ€hodgkin's lymphoma containing reedâ€sternbergâ€like giant cells in Taiwan: A clinicopathologic analysis of 50 cases. Cancer, 1992, 69, 1254-1258.	2.0	6
33	PERIPHERAL TÎ³/Î´ LYMPHOMA PRESENTING WITH IDIOPATHIC THROMBOCYTOPENIC PURPURA-LIKE PICTURE. British Journal of Haematology, 1991, 78, 280-282.	1.2	17
34	Characterization of Genetic Defects of Hemophilia B of Chinese Origin. Thrombosis and Haemostasis, 1991, 66, 459-463.	1.8	10
35	A comparative study of carrier detection in haemophilia A by linear discriminant function. British Journal of Haematology, 1982, 52, 283-293.	1.2	5