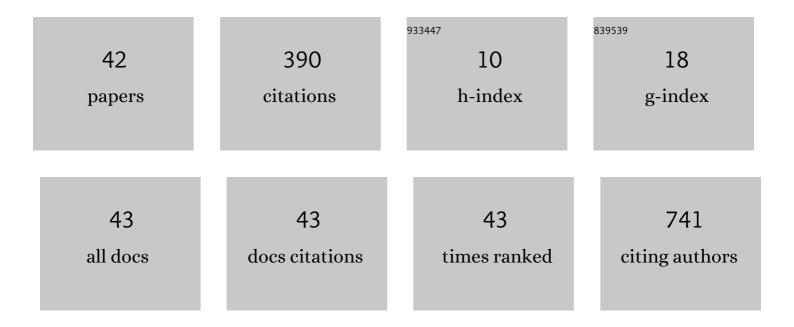
Fu-Sung Lo

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Neurological and endocrinological manifestations of 49 children with intracranial pure germinoma at initial diagnosis in Taiwan. Pediatrics and Neonatology, 2021, 62, 106-112.	0.9	11
2	Higher HbA1c may reduce axial length elongation in myopic children: a comparison cohort study. Acta Diabetologica, 2021, 58, 779-786.	2.5	4
3	Clinical and endocrinological manifestations of childhood-onset craniopharyngioma before surgical removal: A report from one medical center in Taiwan. Pediatrics and Neonatology, 2021, 62, 181-186.	0.9	4
4	Pathways of emotional autonomy, problemâ€solving ability, selfâ€efficacy, and selfâ€management on the glycemic control of adolescents with type 1 diabetes: A prospective study. Research in Nursing and Health, 2021, 44, 643-652.	1.6	4
5	Nasal Methicillin-Resistant Staphylococcus aureus Colonization in Patients with Type 1 Diabetes in Taiwan. Microorganisms, 2021, 9, 1296.	3.6	0
6	Data Homogeneity Effect in Deep Learning-Based Prediction of Type 1 Diabetic Retinopathy. Journal of Diabetes Research, 2021, 2021, 1-9.	2.3	2
7	Evaluation of Disease Complications Among Adults With Type 1 Diabetes and a Family History of Type 2 Diabetes in Taiwan. JAMA Network Open, 2021, 4, e2138775.	5.9	5
8	Association between cardiovascular anomalies and karyotypes in Turner syndrome patients in Taiwan: A local cohort study. Pediatrics and Neonatology, 2020, 61, 188-194.	0.9	6
9	Cardiac manifestations and gene mutations of patients with RASopathies in Taiwan. American Journal of Medical Genetics, Part A, 2020, 182, 357-364.	1.2	8
10	The Self-Management Experiences of Adolescents with Type 1 Diabetes: A Descriptive Phenomenology Study. International Journal of Environmental Research and Public Health, 2020, 17, 5132.	2.6	6
11	HLAâ€DQ genotype and biochemical characterization of antiâ€transglutaminase 2 antibodies in patients with type 1 diabetes mellitus in Taiwan. FASEB Journal, 2020, 34, 8459-8474.	0.5	4
12	Pathological and Incidental Findings in 403 Taiwanese Girls With Central Precocious Puberty at Initial Diagnosis. Frontiers in Endocrinology, 2020, 11, 256.	3.5	20
13	Roles of Emotional Autonomy, Problem-Solving Ability and Parent-Adolescent Relationships on Self-Management of Adolescents with Type 1 Diabetes in Taiwan. Journal of Pediatric Nursing, 2020, 55, e263-e269.	1.5	4
14	Emergence of a butterfly: the life experiences of type 1 diabetes Taiwanese patients during the 16–25 years old transition period. International Journal of Qualitative Studies on Health and Well-being, 2020, 15, 1748362.	1.6	5
15	MON-085 The Clinical and Endocrinologic Manifestations of Germinoma in Taiwanese Pediatric Population: One Medical Center Experience. Journal of the Endocrine Society, 2020, 4, .	0.2	0
16	MON-091 The Clinical and Endocrinologic Manifestations of Craniopharyngioma in Taiwanese Pediatric Population: One Medical Center Experience. Journal of the Endocrine Society, 2020, 4, .	0.2	0
17	SUN-058 Endocrine and Metabolic Complications in 16 Taiwanese Patients with Thalassaemia Major. Journal of the Endocrine Society, 2020, 4, .	0.2	0
18	Mutations in glucokinase and other genes detected in neonatal and type 1B diabetes patient using whole exome sequencing may lead to disease-causing changes in protein activity. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 428-433.	3.8	7

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19	Risk Factors Associated With the Development of Nephropathy 10 Years After Diagnosis in Taiwanese Children With Juvenile-Onset Type 1 Diabetes—A Cohort Study From the CGJDES. Frontiers in Endocrinology, 2018, 9, 429.	3.5	7
20	A 26-week, randomized trial of insulin detemir versus NPH insulin in children and adolescents with type 2 diabetes (iDEAt2). European Journal of Pediatrics, 2018, 177, 1497-1503.	2.7	4
21	Nomogram for prediction of non-proliferative diabetic retinopathy in juvenile-onset type 1 diabetes: a cohort study in an Asian population. Scientific Reports, 2018, 8, 12164.	3.3	15
22	Mutation screening of INS and KCNJ11 genes in Taiwanese children with type 1B diabetic onset before the age of 5 years. Journal of the Formosan Medical Association, 2018, 117, 734-737.	1.7	3
23	Autoantibodies against islet cell antigens in children with type 1 diabetes mellitus. Oncotarget, 2018, 9, 16275-16283.	1.8	13
24	Factors associated with diabetic nephropathy in children, adolescents, and adults with type 1 diabetes. Journal of the Formosan Medical Association, 2017, 116, 924-932.	1.7	11
25	A comprehensive analysis of the association of common variants of ABCG2 with gout. Scientific Reports, 2017, 7, 9988.	3.3	19
26	Pubertal Timing, Smoking, and Sexual Activity. Pediatrics and Neonatology, 2017, 58, 293-294.	0.9	0
27	Functional Analysis of VDR Gene Mutation R343H in A Child with Vitamin D-Resistant Rickets with Alopecia. Scientific Reports, 2017, 7, 15337.	3.3	6
28	Pathways of emotional autonomy, self-care behaviors, and depressive symptoms on health adaptation in adolescents with type 1 diabetes. Nursing Outlook, 2017, 65, 68-76.	2.6	17
29	Detection of Rare Somatic GNAS Mutation in McCune-Albright Syndrome Using a Novel Peptide Nucleic Acid Probe in a Single Tube. Molecules, 2017, 22, 1874.	3.8	4
30	Risk factors associated with the development of retinopathy 10 yr after the diagnosis of juvenile-onset type 1 diabetes in Taiwan: a cohort study from the CCJDES. Pediatric Diabetes, 2016, 17, 407-416.	2.9	17
31	A neonate with recurrent tetany: questions. Pediatric Nephrology, 2016, 31, 753-753.	1.7	4
32	A neonate with recurrent tetany: Answers. Pediatric Nephrology, 2016, 31, 755-757.	1.7	1
33	Association of Cytotoxic T-Lymphocyte-Associated Protein 4 (CTLA4) Gene Polymorphisms with Autoimmune Thyroid Disease in Children and Adults: Case-Control Study. PLoS ONE, 2016, 11, e0154394.	2.5	59
34	Mutations in Pseudohypoparathyroidism 1a and Pseudopseudohypoparathyroidism in Ethnic Chinese. PLoS ONE, 2014, 9, e90640.	2.5	8
35	Noonan syndrome caused by germline KRAS mutation in Taiwan: report of two patients and a review of the literature. European Journal of Pediatrics, 2009, 168, 919-923.	2.7	29
36	High resolution melting analysis for mutation detection for PTPN11 gene: Applications of this method for diagnosis of Noonan syndrome. Clinica Chimica Acta, 2009, 409, 75-77.	1.1	3

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37	Hodgkin's lymphoma in a patient with Noonan syndrome with germ-line PTPN11 mutations. International Journal of Hematology, 2008, 88, 287-290.	1.6	8
38	MUTATIONAL ANALYSIS OF <i>PTPN11</i> AND <i>KRAS</i> GENES IN TAIWANESE CHILDREN WITH NOONAN SYNDROME. Pediatrics, 2008, 121, S116-S117.	2.1	0
39	Two Novel PHEX Mutations in Taiwanese Patients with X-Linked Hypophosphatemic Rickets. Nephron Physiology, 2006, 103, p157-p163.	1.2	10
40	Clinical features of type 1 diabetic children at initial diagnosis. Acta Paediatrica Taiwanica = Taiwan Er Ke Yi Xue Hui Za Zhi, 2004, 45, 218-23.	0.1	3
41	Polymorphism in the Transmembrane Region of the Major Histocompatibility Complex Class I Chain-Related Gene A: Association of Five GCT Repetitions with Graves' Disease in Children. Thyroid, 2003, 13, 839-842.	4.5	12
42	Association of CTLA4 gene A-G polymorphism with type 1 diabetes in Chinese children. Clinical Endocrinology, 2000, 52, 153-157.	2.4	47