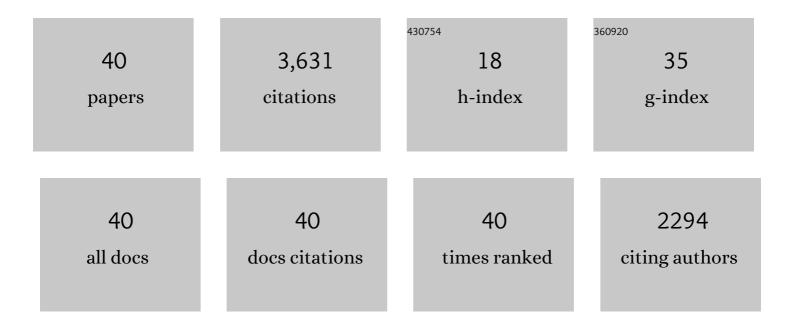
Nancy F Olivieri

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	Survival and complications in patients with haemoglobin E thalassaemia in Sri Lanka: a prospective, longitudinal cohort study. The Lancet Global Health, 2022, 10, e134-e141.	2.9	6
2	Consequences to patients, clinicians, and manufacturers when very serious adverse drug reactions are identified (1997–2019): A qualitative analysis from the Southern Network on Adverse Reactions (SONAR). EClinicalMedicine, 2021, 31, 100693.	3.2	3
3	Oxidative status in the β-thalassemia syndromes in Sri Lanka; a cross-sectional survey. Free Radical Biology and Medicine, 2021, 166, 337-347.	1.3	6
4	A "One-Stop―Screening Protocol for Haemoglobinopathy Traits and Iron Deficiency in Sri Lanka. Frontiers in Molecular Biosciences, 2019, 6, 66.	1.6	3
5	Single-center retrospective study of the effectiveness and toxicity of the oral iron chelating drugs deferiprone and deferasirox. PLoS ONE, 2019, 14, e0211942.	1.1	29
6	Headache: an important symptom possibly linked to white matter lesions in thalassaemia. British Journal of Haematology, 2019, 185, 541-548.	1.2	7
7	Iron status and anaemia in Sri Lankan secondary school children: A cross-sectional survey. PLoS ONE, 2017, 12, e0188110.	1.1	15
8	Improving Laboratory and Clinical Hematology Services in Resource Limited Settings. Hematology/Oncology Clinics of North America, 2016, 30, 497-512.	0.9	4
9	Dysregulated arginine metabolism and cardiopulmonary dysfunction in patients with thalassaemia. British Journal of Haematology, 2015, 169, 887-898.	1.2	22
10	Treatment of heart failure in adults with thalassemia major: response in patients randomised to deferoxamine with or without deferiprone. Journal of Cardiovascular Magnetic Resonance, 2013, 15, 38.	1.6	47
11	Chelation use and iron burden in North American and British thalassemia patients: a report from the Thalassemia Longitudinal Cohort. Blood, 2012, 119, 2746-2753.	0.6	78
12	Treatment strategies for hemoglobin E beta-thalassemia. Blood Reviews, 2012, 26, S28-S30.	2.8	15
13	Hb E/beta-thalassaemia: a common & clinically diverse disorder. Indian Journal of Medical Research, 2011, 134, 522-31.	0.4	37
14	Emerging insights in the management of hemoglobin E beta thalassemia. Annals of the New York Academy of Sciences, 2010, 1202, 155-157.	1.8	12
15	HbE/β-Thalassemia: Basis of Marked Clinical Diversity. Hematology/Oncology Clinics of North America, 2010, 24, 1055-1070.	0.9	34
16	Phenomenon of Pain In Thalassemia: A Prospective Analysis by the Thalassemia Clinical Research Network (TCRN). Blood, 2010, 116, 256-256.	0.6	0
17	Rates of Non-Transfusional Iron Accumulation (NTIA) In Hemoglobin E Thalassemia. Blood, 2010, 116, 5147-5147.	0.6	1
18	Pulmonary Hypertension in Thalassemia Assessed by Echocardiography: A Report From Baseline Data of the Thalassemia Clinical Research Network Longitudinal Cohort Study Blood, 2009, 114, 2016-2016.	0.6	3

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19	Chelation Choices and Iron Burden Among Patients with Thalassemia in the 21st Century: a Report From the Thalassemia Clinical Research Network (TCRN) Longitudinal Cohort Blood, 2009, 114, 4056-4056.	0.6	5
20	Renal Dysfunction in Thalassemia Blood, 2009, 114, 2008-2008.	0.6	1
21	Studies in haemoglobin E betaâ€ŧhalassaemia. British Journal of Haematology, 2008, 141, 388-397.	1.2	103
22	Hemoglobin H-Constant Spring in North America: A Common Alpha Thalassemia with Frequent Complications Blood, 2008, 112, 1880-1880.	0.6	0
23	Body Composition and Its Relationship to Growth and Bone Mass in Patients with Thalassemia. Blood, 2008, 112, 3890-3890.	0.6	0
24	Increased Nucleosomal DNA Fragmentation in Leukocytes of Thalassemia Patients Blood, 2008, 112, 1868-1868.	0.6	0
25	Progression of Avascular Necrosis of the Hip in Sickle Cell Disease: 2 Year Follow-Up of Randomized Trial of Aggressive Physical Therapy and Hip Coring Decompression Blood, 2004, 104, 1685-1685.	0.6	1
26	A Phase II Study with ICL670 (Exjade®), a Once-Daily Oral Iron Chelator, in Patients with Various Transfusion-Dependent Anemias and Iron Overload Blood, 2004, 104, 3193-3193.	0.6	17
27	Low Bone Mass in Thalassemia: The Thalassemia Clinical Research Network (TCRN) Experience Blood, 2004, 104, 3613-3613.	0.6	0
28	Patients' health or company profits? The commercialisation of academic research. Science and Engineering Ethics, 2003, 9, 29-41.	1.7	60
29	Deferiprone and hepatic fibrosis. Blood, 2003, 101, 5089-5091.	0.6	18
30	Long-Term Trials of Deferiprone in Cooley's Anemiaa. Annals of the New York Academy of Sciences, 1998, 850, 217-222.	1.8	13
31	Long-Term Safety and Effectiveness of Iron-Chelation Therapy with Deferiprone for Thalassemia Major. New England Journal of Medicine, 1998, 339, 417-423.	13.9	389
32	Iron-Chelating Therapy and the Treatment of Thalassemia. Blood, 1997, 89, 739-761.	0.6	971
33	Iron-Chelation Therapy with Oral Deferiprone in Patients with Thalassemia Major. New England Journal of Medicine, 1995, 332, 918-922.	13.9	306
34	Critical Comparison of Novel and Existing Methods of Compliance Assessment During a Clinical Trial of an Oral Iron Chelator. Journal of Clinical Pharmacology, 1994, 34, 944-949.	1.0	79
35	Iron overload cardiomyopathies: New insights into an old disease. Cardiovascular Drugs and Therapy, 1994, 8, 101-110.	1.3	128
36	Survival in Medically Treated Patients with Homozygous β-Thalassemia. New England Journal of Medicine, 1994, 331, 574-578.	13.9	829

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#	Article	IF	CITATIONS
37	Pain Perception and Effectiveness of the Eutectic Mixture of Local Anesthetics in Children Undergoing Venipuncture. Pediatric Research, 1992, 32, 520-523.	1.1	35
38	Reduction in tissue iron stores with a new regimen of continuous ambulatory intravenous deferoxamine. American Journal of Hematology, 1992, 41, 61-63.	2.0	34
39	Relationship between the pharmacokinetics and iron excretion pharmacodynamics of the new oral iron chelator 1,2-dimethyl-3-hydroxypyrid-4-one in patients with thalassemia. Clinical Pharmacology and Therapeutics, 1991, 50, 294-298.	2.3	51
40	Prevention of Cardiac Disease by Subcutaneous Deferoxamine in Patients with Thalassemia Major. New England Journal of Medicine, 1985, 312, 1600-1603.	13.9	269