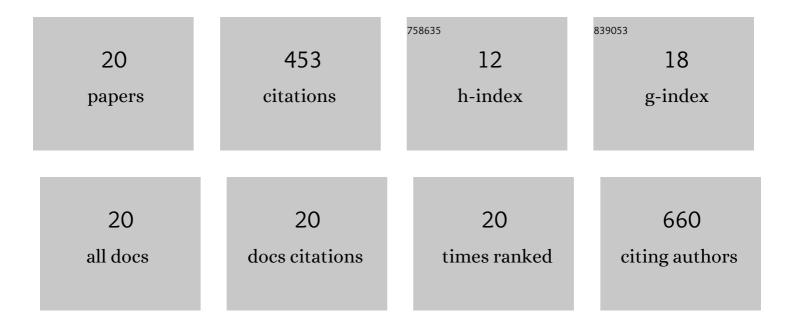
Erika Maka

List of Publications by Year in descending order

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Ερικλ Μλκλ

#	Article	IF	CITATIONS
1	TUBB4B gene mutation in Leber phenotype of congenital amaurosis syndrome associated with early-onset deafness. European Journal of Medical Genetics, 2022, 65, 104471.	0.7	3
2	The Global Retinoblastoma Outcome Study: a prospective, cluster-based analysis of 4064 patients from 149 countries. The Lancet Global Health, 2022, 10, e1128-e1140.	2.9	24
3	Travel burden and clinical presentation of retinoblastoma: analysis of 1024 patients from 43 African countries and 518 patients from 40 European countries. British Journal of Ophthalmology, 2021, 105, 1435-1443.	2.1	19
4	Adjuvant therapy of histopathological risk factors of retinoblastoma in Europe: A survey by the European Retinoblastoma Group (EURbG). Pediatric Blood and Cancer, 2021, 68, e28963.	0.8	9
5	Incidence of Retinoblastoma Has Increased: Results from 40 European Countries. Ophthalmology, 2021, 128, 1369-1371.	2.5	26
6	Pseudouridylation defect due to <i>DKC1</i> and <i>NOP10</i> mutations causes nephrotic syndrome with cataracts, hearing impairment, and enterocolitis. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 15137-15147.	3.3	32
7	Examination of corneal deposits in nephropathic cystinosis using in vivo confocal microscopy and anterior segment optical coherence tomography: an age-dependent cross sectional study. BMC Ophthalmology, 2020, 20, 73.	0.6	4
8	Global Retinoblastoma Presentation and Analysis by National Income Level. JAMA Oncology, 2020, 6, 685.	3.4	192
9	The validity of telemedicine-based screening for retinopathy of prematurity in the Premature Eye Rescue Program in Hungary. Journal of Telemedicine and Telecare, 2019, 27, 1357633X1988011.	1.4	4
10	Clinicopathological Review of 547 Bulbar Enucleations in Hungary (2006–2017). Journal of Ophthalmology, 2019, 2019, 1-7.	0.6	3
11	Preoperative biometry data of eyes with unilateral congenital cataract. Journal of Cataract and Refractive Surgery, 2018, 44, 1198-1202.	0.7	12
12	Bedside ROP screening and telemedicine interpretation integrated to a neonatal transport system: Economic aspects and return on investment analysis. Early Human Development, 2017, 106-107, 1-5.	0.8	29
13	Involving the parents of preterm babies. Community Eye Health Journal, 2017, 30, 60-61.	0.4	0
14	Mitochondrial Ca2+ uptake correlates with the severity of the symptoms in autosomal dominant optic atrophy. Cell Calcium, 2015, 57, 49-55.	1.1	16
15	Measurements of Orbital Volume using Cone-Beam Computed Tomography in Eye Movement Abnormalities. European Journal of Ophthalmology, 2014, 24, 101-106.	0.7	0
16	NPHS2 homozygous p.R229Q variant: potential modifier instead of causal effect in focal segmental glomerulosclerosis. Pediatric Nephrology, 2013, 28, 2061-2064.	0.9	21
17	Measurement of Orbital Volume after Enucleation and Orbital Implantation. PLoS ONE, 2012, 7, e50333.	1.1	17
18	Duodenal Ulceration in a Patient With Celiac Disease and Plasminogen I Deficiency: Coincidence or Cofactors?. Pediatrics, 2011, 128, e1302-e1306.	1.0	4

#	Article	IF	CITATIONS
19	Recently Recognized Cases of Ophthalmofilariasis in Hungary. European Journal of Ophthalmology, 2009, 19, 675-678.	0.7	19
20	Orbital Tumour as Initial Manifestation of Acute Myeloid Leukemia: Granulocytic Sarcoma: Case Report. Pathology and Oncology Research, 2008, 14, 209-211.	0.9	19