Ilse Wieland

List of Publications by Year in Descending Order

14 papers	89 citations	5 h-index	9 g-index
14 ext. papers	115 ext. citations	3.1 avg, IF	1.8 L-index

#	Paper	IF	Citations
14	Correlation of PET-MRI, Pathology, LOH, and Surgical Success in a Case of CHI With Atypical Large Pancreatic Focus.. *Journal of the Endocrine Society*, **2022**, 6, bvac056	0.4	0
13	Cutis marmorata telangiectatica congenita being caused by postzygotic GNA11 mutations.. *European Journal of Medical Genetics*, **2022**, 65, 104472	2.6	1
12	Oral HRAS Mutation in Orofacial Nevus Sebaceous Syndrome (Schimmelpenning-Feuerstein-Mims-Syndrome): A Case Report With a Literature Survey.. *In Vivo*, **2022**, 36, 274-293	2.3	
11	Structural changes to primary visual cortex in the congenital absence of cone input in achromatopsia.. *NeuroImage: Clinical*, **2021**, 33, 102925	5.3	1
10	Structural Differences Across Multiple Visual Cortical Regions in the Absence of Cone Function in Congenital Achromatopsia. *Frontiers in Neuroscience*, **2021**, 15, 718958	5.1	1
9	Mutation in an Implant-associated Peripheral Giant Cell Granuloma of the Jaw: Implications of Genetic Analysis of the Lesion for Treatment Concept and Surveillance. *In Vivo*, **2021**, 35, 947-953	2.3	1
8	Neurofibromatosis Type 1 With Cherubism-like Phenotype, Multiple Osteolytic Bone Lesions of Lower Extremities, and Alagille-syndrome: Case Report With Literature Survey. *In Vivo*, **2021**, 35, 1711-1736	2.3	1
7	Mosaic Neurofibromatosis Type 1 With Multiple Cutaneous Diffuse and Plexiform Neurofibromas of the Lower Leg. *Anticancer Research*, **2020**, 40, 3423-3427	2.3	2
6	Expansion of the phenotypic spectrum and description of molecular findings in a cohort of patients with oculocutaneous mosaic RASopathies. *Molecular Genetics & Genomic Medicine*, **2019**, 7, e625	2.3	25
5	NFIB Haploinsufficiency Is Associated with Intellectual Disability and Macrocephaly. *American Journal of Human Genetics*, **2018**, 103, 752-768	11	19
4	Surgery in Focal Congenital Hyperinsulinism (CHI) - The "Hyperinsulinism Germany International" Experience in 30 Children. *Pediatric Endocrinology Reviews*, **2016**, 14, 129-137	1.1	15
3	A cryptic unbalanced translocation der(4)t(4;17)(p16.1;q25.3) identifies Wittwer syndrome as a variant of Wolf-Hirschhorn syndrome. *American Journal of Medical Genetics, Part A*, **2014**, 164A, 3213-4	2.5	3
2	High-level somatic mosaicism of AKT1 c.49G>A mutation in skin scrapings from epidermal nevi enables non-invasive molecular diagnosis in patients with Proteus syndrome. *American Journal of Medical Genetics, Part A*, **2013**, 161A, 889-91	2.5	8
1	Ectopic expression of DICE1 suppresses tumor cell growth. *Oncology Reports*, **2004**, 12, 207-11	3.5	12