# Ilse Wieland

## List of Publications by Citations



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|---|---|---|---|
| 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 | 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 |
| 13 | NFIB Haploinsufficiency Is Associated with Intellectual Disability and Macrocephaly. *American Journal of Human Genetics*, **2018**, 103, 752-768 | 11 | 19 |
| 12 | Surgery in Focal Congenital Hyperinsulinism (CHI) - The "Hyperinsulinism Germany International" Experience in 30 Children. *Pediatric Endocrinology Reviews*, **2016**, 14, 129-137 | 1.1 | 15 |
| 11 | Ectopic expression of DICE1 suppresses tumor cell growth. *Oncology Reports*, **2004**, 12, 207-11 | 3.5 | 12 |
| 10 | 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 |
| 9 | 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 |
| 8 | Mosaic Neurofibromatosis Type 1 With Multiple Cutaneous Diffuse and Plexiform Neurofibromas of the Lower Leg. *Anticancer Research*, **2020**, 40, 3423-3427 | 2.3 | 2 |
| 7 | Structural changes to primary visual cortex in the congenital absence of cone input in achromatopsia.. *NeuroImage: Clinical*, **2021**, 33, 102925 | 5.3 | 1 |
| 6 | 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 |
| 5 | 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 |
| 4 | 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 |
| 3 | Cutis marmorata telangiectatica congenita being caused by postzygotic GNA11 mutations.. *European Journal of Medical Genetics*, **2022**, 65, 104472 | 2.6 | 1 |
| 2 | 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 |
| 1 | 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 | |