# Yongli Guo

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2879955/publications.pdf

Version: 2024-02-01

| 59 | 1,207 | 489802 | 488211 |
|---|---|---|---|
| papers | citations | 18 h-index | 31 g-index |

| 62 | 62 | 62 | 1733 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Predictive Significance of Enhanced Level of Angiogenesis and Tissue Neutrophils for Antrochoanal Polyps Recurrence in Children. Ear, Nose and Throat Journal, 2022, 101, NP284-NP290. | 0.4 | 4 |
| 2 | A group of sclerosing epithelioid fibrosarcomas with low-level amplified EWSR1-CREB3L1 fusion gene in children. Pathology Research and Practice, 2022, 230, 153754. | 1.0 | 5 |
| 3 | Epidemiology of extrapulmonary tuberculosis among pediatric inpatients in mainland China: a descriptive, multicenter study. Emerging Microbes and Infections, 2022, 11, 1090-1102. | 3.0 | 12 |
| 4 | Clinical Features and Survival of Chinese Children With Trilateral Retinoblastoma During 2006-2019: A Retrospective Multicenter Study. American Journal of Ophthalmology, 2021, 223, 184-192. | 1.7 | 4 |
| 5 | The efficacy and safety of montelukast in children with obstructive sleep apnea: a systematic review and meta-analysis. Sleep Medicine, 2021, 78, 193-201. | 0.8 | 18 |
| 6 | Effects of obstructive sleep apnoea severity on neurocognitive and brain white matter alterations in children according to sex: aÂtract-based spatial statistics study. Sleep Medicine, 2021, 82, 134-143. | 0.8 | 11 |
| 7 | MYC-associated protein X binding with the variant rs72780850 in RNA helicase DEAD box 1 for susceptibility to neuroblastoma. Science China Life Sciences, 2021, 64, 991-999. | 2.3 | 3 |
| 8 | Efficacy of Initial Sirolimus Therapy for 27 Patients with Intractable Lymphatic Malformations. Laryngoscope, 2021, 131, 1902-1908. | 1.1 | 8 |
| 9 | Clinical implications of TPO and AOX1 in pediatric papillary thyroid carcinoma. Translational Pediatrics, 2021, 10, 723-732. | 0.5 | 4 |
| 10 | A somatic mutation in PIK3CD unravels a novel candidate gene for lymphatic malformation. Orphanet Journal of Rare Diseases, 2021, 16, 208. | 1.2 | 8 |
| 11 | Primary cardiac CICâ€rearranged undifferentiated sarcoma in an infant. Pediatric Investigation, 2021, 5, 313-317. | 0.6 | 3 |
| 12 | DCX and CRABP2 are candidate genes for differential diagnosis between preâ€chemotherapy embryonic and alveolar rhabdomyosarcoma in pediatric patients. Pediatric Investigation, 2021, 5, 106-111. | 0.6 | 1 |
| 13 | Clinical Heterogeneity of Differentiated Thyroid Cancer between Children Less than 10 Years of Age and Those Older than 10 Years: A Retrospective Study of 70 Cases. European Thyroid Journal, 2021, 10, 364-371. | 1.2 | 6 |
| 14 | Characterization of the Blood and Cerebrospinal Fluid Microbiome in Children with Bacterial Meningitis and Its Potential Correlation with Inflammation. MSystems, 2021, 6, e0004921. | 1.7 | 10 |
| 15 | A Novel Germline Compound Heterozygous Mutation of BRCA2 Gene Associated With Familial Peripheral Neuroblastic Tumors in Two Siblings. Frontiers in Genetics, 2021, 12, 652718. | 1.1 | 0 |
| 16 | The Impacts of Obstructive Sleep Apnea Severity on Brain White Matter Integrity and Cognitive Functions in Children: A Diffusion Tensor Imaging Study. Nature and Science of Sleep, 2021, Volume 13, 2125-2135. | 1.4 | 3 |
| 17 | The optimal surgical approach for papillary thyroid carcinoma with pathological n1 metastases: An analysis from the SEER database. Laryngoscope, 2020, 130, 269-273. | 1.1 | 1 |
| 18 | DNMIVD: DNA methylation interactive visualization database. Nucleic Acids Research, 2020, 48, D856-D862. | 6.5 | 86 |

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|---|---|---|---|
| 19 | Clinical Features of Children with Retinoblastoma and Neuroblastoma. Journal of Ophthalmology, 2020, 2020, 1-8. | 0.6 | 7 |
| 20 | Two novel mutations of PAX3 and SOX10 were characterized as genetic causes of Waardenburg Syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1217. | 0.6 | 13 |
| 21 | Genotype-phenotype correlations of Berardinelli-Seip congenital lipodystrophy and novel candidate genes prediction. Orphanet Journal of Rare Diseases, 2020, 15, 108. | 1.2 | 4 |
| 22 | Application of Gastroscopy in the Diagnosis of Congenital Pyriform Sinus Fistula in Children. Frontiers in Pediatrics, 2020, 8, 541249. | 0.9 | 1 |
| 23 | Bioinformatics analysis to screen key genes in papillary thyroid carcinoma. Oncology Letters, 2020, 19, 195-204. | 0.8 | 10 |
| 24 | Downregulated NORAD in neuroblastoma promotes cell proliferation via chromosomal instability and predicts poor prognosis. Acta Biochimica Polonica, 2020, 67, 595-603. | 0.3 | 5 |
| 25 | Correlation between TERT C228T and clinic-pathological features in pediatric papillary thyroid carcinoma. Science China Life Sciences, 2019, 62, 1563-1571. | 2.3 | 16 |
| 26 | Functional Polymorphisms in BARD1 Association with Neuroblastoma in a regional Han Chinese Population. Journal of Cancer, 2019, 10, 2153-2160. | 1.2 | 10 |
| 27 | Genotypeâ€“Phenotype Association Analysis Reveals New Pathogenic Factors for Osteogenesis Imperfecta Disease. Frontiers in Pharmacology, 2019, 10, 1200. | 1.6 | 11 |
| 28 | Systematically Analyzing the Pathogenic Variations for Acute Intermittent Porphyria. Frontiers in Pharmacology, 2019, 10, 1018. | 1.6 | 14 |
| 29 | Clinical analysis of surgical treatment for head and neck lymphatic malformations in children: a series of 128 cases. Acta Oto-Laryngologica, 2019, 139, 713-719. | 0.3 | 11 |
| 30 | lncRNA SNHG16 is associated with proliferation and poor prognosis of pediatric neuroblastoma. International Journal of Oncology, 2019, 55, 93-102. | 1.4 | 22 |
| 31 | Sequencing XMET genes to promote genotype-guided risk assessment and precision medicine. Science China Life Sciences, 2019, 62, 895-904. | 2.3 | 5 |
| 32 | eRAM: encyclopedia of rare disease annotations for precision medicine. Nucleic Acids Research, 2018, 46, D937-D943. | 6.5 | 56 |
| 33 | RRS1 gene expression involved in the progression of papillary thyroid carcinoma. Cancer Cell International, 2018, 18, 20. | 1.8 | 18 |
| 34 | CO 2 laser cauterization approach to congenital pyriform sinus fistula. Journal of Pediatric Surgery, 2018, 53, 1313-1317. | 0.8 | 21 |
| 35 | Multiple microRNAs function as self-protective modules in acetaminophen-induced hepatotoxicity in humans. Archives of Toxicology, 2018, 92, 845-858. | 1.9 | 42 |
| 36 | Whole-Genome Sequencing Identifies a Novel Variation of WAS Gene Coordinating With Heterozygous Germline Mutation of APC to Enhance Hepatoblastoma Oncogenesis. Frontiers in Genetics, 2018, 9, 668. | 1.1 | 11 |

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|---|---|---|---|
| 37 | RDAD: A Machine Learning System to Support Phenotype-Based Rare Disease Diagnosis. Frontiers in Genetics, 2018, 9, 587. | 1.1 | 33 |
| 38 | Deep Learning-Based Multi-Omics Data Integration Reveals Two Prognostic Subtypes in High-Risk Neuroblastoma. Frontiers in Genetics, 2018, 9, 477. | 1.1 | 146 |
| 39 | Association between mild or moderate obstructive sleep apnea-hypopnea syndrome and cognitive dysfunction in children. Sleep Medicine, 2018, 50, 132-136. | 0.8 | 22 |
| 40 | MiR-20a-5p suppresses tumor proliferation by targeting autophagy-related gene 7 in neuroblastoma. Cancer Cell International, 2018, 18, 5. | 1.8 | 41 |
| 41 | PedAM: a database for Pediatric Disease Annotation and Medicine. Nucleic Acids Research, 2018, 46, D977-D983. | 6.5 | 27 |
| 42 | Application of genome analysis strategies in the clinical testing for pediatric diseases. Pediatric Investigation, 2018, 2, 72-81. | 0.6 | 29 |
| 43 | Efficacy and safety of oral sildenafil in treatment of pediatric head and neck lymphatic malformations. Acta Oto-Laryngologica, 2017, 137, 674-678. | 0.3 | 20 |
| 44 | Investigation of IGF2, IGFBP2 and p63 proteins in rhabdomyosarcoma tumors. Growth Hormone and IGF Research, 2017, 33, 17-22. | 0.5 | 5 |
| 45 | Whole Genome Sequencing Identifies Novel Compound Heterozygous Lysosomal Trafficking Regulator Gene Mutations Associated with Autosomal Recessive Chediak-Higashi Syndrome. Scientific Reports, 2017, 7, 41308. | 1.6 | 9 |
| 46 | Correlation between BRAF V600E mutation and clinicopathological features in pediatric papillary thyroid carcinoma. Science China Life Sciences, 2017, 60, 729-738. | 2.3 | 28 |
| 47 | Detection of FOXO1 break-apart status by fluorescence in situ hybridization in atypical alveolar rhabdomyosarcoma. Science China Life Sciences, 2017, 60, 721-728. | 2.3 | 6 |
| 48 | The Feasibility of Xpert MTB/RIF Testing to Detect Rifampicin Resistance among Childhood Tuberculosis for Prevalence Surveys in Northern China. BioMed Research International, 2017, 2017, 1-10. | 0.9 | 9 |
| 49 | MicroRNA-365a-3p promotes tumor growth and metastasis in laryngeal squamous cell carcinoma. Oncology Reports, 2016, 35, 2017-2026. | 1.2 | 36 |
| 50 | MicroRNA hsa-miR-25-3p suppresses the expression and drug induction of CYP2B6 in human hepatocytes. Biochemical Pharmacology, 2016, 113, 88-96. | 2.0 | 45 |
| 51 | Deafness gene mutations in newborns in Beijing. Acta Oto-Laryngologica, 2016, 136, 475-479. | 0.3 | 10 |
| 52 | Single nucleotide polymorphism rs11669203 in TGFBR3L is associated with the risk of neuroblastoma in a Chinese population. Tumor Biology, 2016, 37, 3739-3747. | 0.8 | 11 |
| 53 | Maternal smoking during pregnancy and risk of childhood neuroblastoma: Systematic review and meta-analysis. Journal of Cancer Research and Therapeutics, 2016, 12, 999. | 0.3 | 18 |
| 54 | Modulation of ALDH5A1 and SLC22A7 by microRNA hsa-miR-29a-3p in human liver cells. Biochemical Pharmacology, 2015, 98, 671-680. | 2.0 | 21 |

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|---|---|---|---|
| 55 | Suppression of CYP2C9 by MicroRNA hsa-miR-128-3p in Human Liver Cells and Association with Hepatocellular Carcinoma. Scientific Reports, 2015, 5, 8534. | 1.6 | 92 |
| 56 | MicroRNA hsa-miR-29a-3p modulates CYP2C19 in human liver cells. Biochemical Pharmacology, 2015, 98, 215-223. | 2.0 | 51 |
| 57 | Candidate Gene Association Analysis of Neuroblastoma in Chinese Children Strengthens the Role of LMO1. PLoS ONE, 2015, 10, e0127856. | 1.1 | 23 |
| 58 | Functional Genetic Variants of TNFSF15 and Their Association with Gastric Adenocarcinoma: A Case-Control Study. PLoS ONE, 2014, 9, e108321. | 1.1 | 14 |
| 59 | Gene Expression Variability in Human Hepatic Drug Metabolizing Enzymes and Transporters. PLoS ONE, 2013, 8, e60368. | 1.1 | 43 |