

# Armaghan Alam

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2121595/publications.pdf>

Version: 2024-02-01

9  
papers

245  
citations

1478280

6  
h-index

1474057

9  
g-index

12  
all docs

12  
docs citations

12  
times ranked

510  
citing authors

| # | ARTICLE  | IF  | CITATIONS |
|---|--|-----|-----------|
| 1 | The impact of brain invasion criteria on the incidence and distribution of WHO grade 1, 2, and 3 meningiomas. <i>Neuro-Oncology</i> , 2022, 24, 1524-1532.   | 0.6 | 9         |
| 2 | Survival and Recurrence Outcomes Following Adjuvant Radiotherapy for Grade 2 Intracranial Meningiomas: 13-Year Experience in a Tertiary-Care Center. <i>World Neurosurgery</i> , 2022, , .                           | 0.7 | 3         |
| 3 | Choice and Trade-offs: Parent Decision Making for Neurotechnologies for Pediatric Drug-Resistant Epilepsy. <i>Journal of Child Neurology</i> , 2021, 36, 943-949.  | 0.7 | 11        |
| 4 | Using Peer Support to Strengthen Mental Health During the COVID-19 Pandemic: A Review. <i>Frontiers in Psychiatry</i> , 2021, 12, 714181.  | 1.3 | 62        |
| 5 | Fineâ€Mapping of <i>SNCA</i> in Rapid Eye Movement Sleep Behavior Disorder and Overt Synucleinopathies. <i>Annals of Neurology</i> , 2020, 87, 584-598.  | 2.8 | 39        |
| 6 | SMPD1 variants do not have a major role in rapid eye movement sleep behavior disorder. <i>Neurobiology of Aging</i> , 2020, 93, 142.e5-142.e7.   | 1.5 | 4         |
| 7 | Increased yield of full GBA sequencing in Ashkenazi Jews with Parkinson's disease. <i>European Journal of Medical Genetics</i> , 2019, 62, 65-69.  | 0.7 | 49        |
| 8 | Carriers of both GBA and LRRK2 mutations, compared to carriers of either, in Parkinson's disease: Risk estimates and genotype-phenotype correlations. <i>Parkinsonism and Related Disorders</i> , 2019, 62, 179-184. | 1.1 | 58        |
| 9 | The <i>GBA</i> p.Trp378Gly mutation is a probable Frenchâ€Canadian founder mutation causing Gaucher disease and synucleinopathies. <i>Clinical Genetics</i> , 2018, 94, 339-345.                                     | 1.0 | 9         |