

# Nuha Alrayes

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3539584/publications.pdf>

Version: 2024-02-01

10  
papers

154  
citations

1478505

6  
h-index

1372567

10  
g-index

10  
all docs

10  
docs citations

10  
times ranked

387  
citing authors

| #  | ARTICLE  | IF  | CITATIONS |
|----|--|-----|-----------|
| 1  | Exome analysis identified a novel missense mutation in the CLPP gene in a consanguineous Saudi family expanding the clinical spectrum of Perrault Syndrome type-3. <i>Journal of the Neurological Sciences</i> , 2015, 353, 149-154.     | 0.6 | 37        |
| 2  | A missense mutation in TRAPPC6A leads to build-up of the protein, in patients with a neurodevelopmental syndrome and dysmorphic features. <i>Scientific Reports</i> , 2018, 8, 2053.   | 3.3 | 30        |
| 3  | The alkylglycerol monooxygenase (AGMO) gene previously involved in autism also causes a novel syndromic form of primary microcephaly in a consanguineous Saudi family. <i>Journal of the Neurological Sciences</i> , 2016, 363, 240-244. | 0.6 | 18        |
| 4  | Truncating mutation in intracellular phospholipase A1 gene (DDHD2) in hereditary spastic paraplegia with intellectual disability (SPG54). <i>BMC Research Notes</i> , 2015, 8, 271.  | 1.4 | 17        |
| 5  | Myocardial infarction biomarker discovery with integrated gene expression, pathways and biological networks analysis. <i>Genomics</i> , 2020, 112, 5072-5085.  | 2.9 | 17        |
| 6  | Molecular insights into the coding region mutations of low-density lipoprotein receptor adaptor protein 1 (LDLRAP1) linked to familial hypercholesterolemia. <i>Journal of Gene Medicine</i> , 2020, 22, e3176.                          | 2.8 | 12        |
| 7  | Whole exome sequencing of a Saudi family and systems biology analysis identifies CPED1 as a putative causative gene to Celiac Disease. <i>Saudi Journal of Biological Sciences</i> , 2020, 27, 1494-1502.                                | 3.8 | 8         |
| 8  | Exome Analysis Identifies a Novel Compound Heterozygous Alteration in TGM1 Gene Leading to Lamellar Ichthyosis in a Child From Saudi Arabia: Case Presentation. <i>Frontiers in Pediatrics</i> , 2019, 7, 44.                            | 1.9 | 7         |
| 9  | Novel missense alteration in <i>LRP4</i> gene underlies Cenani-Lenz syndactyly syndrome in a consanguineous family. <i>Journal of Gene Medicine</i> , 2020, 22, e3143.   | 2.8 | 5         |
| 10 | Exome Sequencing Identifies the Extremely Rare ITGAV and FN1 Variants in Early Onset Inflammatory Bowel Disease Patients. <i>Frontiers in Pediatrics</i> , 2022, 10, .   | 1.9 | 3         |