# Karen Nuytemans

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8459590/publications.pdf

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| 23 | 511 | 933447 | 888059 |
|---|---|---|---|
| papers | citations | 10 h-index | 17 g-index |

| 23 | 23 | 23 | 1109 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Identifying differential regulatory control of APOE É›4 on African versus European haplotypes as potential therapeutic targets. Alzheimer's and Dementia, 2022, 18, 1930-1942. | 0.8 | 12 |
| 2 | Genetic architecture of RNA editing regulation in Alzheimerâ€™s disease across diverse ancestral populations. Human Molecular Genetics, 2022, 31, 2876-2886. | 2.9 | 2 |
| 3 | Increased APOE Îµ4 expression is associated with the difference in Alzheimer's disease risk from diverse ancestral backgrounds. Alzheimer's and Dementia, 2021, 17, 1179-1188. | 0.8 | 33 |
| 4 | Successful Management of Catastrophic Thrombotic Storm in a Young Boy. Journal of Pediatric Hematology/Oncology, 2021, Publish Ahead of Print, e1132-e1135. | 0.6 | 1 |
| 5 | Derivation of stem cell line UMi028-A-2 containing a CRISPR/Cas9 induced Alzheimerâ€™s disease risk variant p.S1038C in the TTC3 gene. Stem Cell Research, 2021, 52, 102258. | 0.7 | 7 |
| 6 | Genomewide Association Studies of <scp>LRRK2</scp> Modifiers of Parkinson's Disease. Annals of Neurology, 2021, 90, 76-88. | 5.3 | 30 |
| 7 | A novel duplication involving in a Turkish family supports its role in North Carolina macular dystrophy (NCMD/MCDR1). Molecular Vision, 2021, 27, 518-527. | 1.1 | 2 |
| 8 | Ancestryâ€specific intronic variants on the APOE É›4 haplotype influence enhancer activity and interaction with APOE promoter. Alzheimer's and Dementia, 2021, 17, e055266. | 0.8 | 0 |
| 9 | Derivation of a CRISPR genome edited stem cell line containing a risk variant in TTC3.. Alzheimer's and Dementia, 2021, 17 Suppl 3, e056331. | 0.8 | 0 |
| 10 | Novel Variants in LRRK2 and GBA Identified in Latino Parkinson Disease Cohort Enriched for Caribbean Origin. Frontiers in Neurology, 2020, 11, 573733. | 2.4 | 6 |
| 11 | Increased APOEâ€e4 expression is associated with reactive A1 astrocytes and may confer the difference in Alzheimer disease risk from different ancestral backgrounds. Alzheimer's and Dementia, 2020, 16, e045415. | 0.8 | 0 |
| 12 | Development of a massively parallel reporter assay to identify functional regulatory variants in the PICALM Alzheimer disease associated locus. Alzheimer's and Dementia, 2020, 16, e045908. | 0.8 | 0 |
| 13 | Identification of differential regulation of European versus African local ancestry haplotypes surrounding ApoEÎµ4. Alzheimer's and Dementia, 2020, 16, e046016. | 0.8 | 0 |
| 14 | Motivations for Participation in Parkinson Disease Genetic Research Among Hispanics versus Non-Hispanics. Frontiers in Genetics, 2019, 10, 658. | 2.3 | 10 |
| 15 | Variants in chondroitin sulfate metabolism genes in thrombotic storm. Thrombosis Research, 2018, 161, 43-51. | 1.7 | 5 |
| 16 | Transcriptomic analysis of synovial extracellular RNA following knee trauma: A pilot study. Journal of Orthopaedic Research, 2018, 36, 1659-1665. | 2.3 | 11 |
| 17 | Generation of disease-specific autopsy-confirmed iPSCs lines from postmortem isolated Peripheral Blood Mononuclear Cells. Neuroscience Letters, 2017, 637, 201-206. | 2.1 | 6 |
| 18 | Identification of TMEM230 mutations in familial Parkinson's disease. Nature Genetics, 2016, 48, 733-739. | 21.4 | 146 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Overlap between Parkinson disease and Alzheimer disease in ABCA7 functional variants. Neurology: Genetics, 2016, 2, e44. | 1.9 | 31 |
| 20 | PARK10 is a major locus for sporadic neuropathologically confirmed Parkinson disease. Neurology, 2015, 84, 972-980. | 1.1 | 48 |
| 21 | Absence of C9ORF72 expanded or intermediate repeats in autopsyâ€confirmed Parkinson's disease. Movement Disorders, 2014, 29, 827-830. | 3.9 | 24 |
| 22 | C9ORF72 Intermediate Repeat Copies Are a Significant Risk Factor for Parkinson Disease. Annals of Human Genetics, 2013, 77, 351-363. | 0.8 | 69 |
| 23 | Whole exome sequencing of rare variants in EIF4G1 and VPS35 in Parkinson disease. Neurology, 2013, 80, 982-989. | 1.1 | 68 |