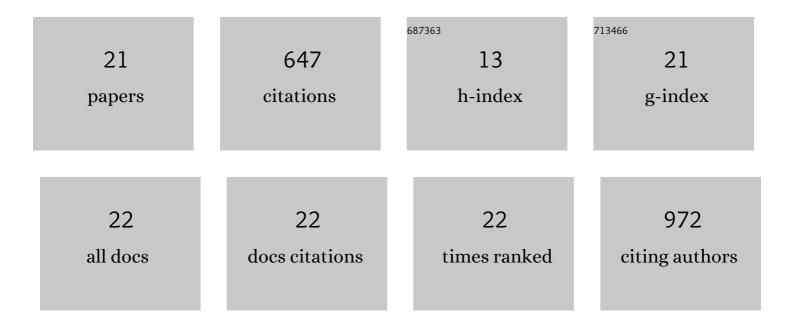
Claudia Stancanelli

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5889436/publications.pdf

Version: 2024-02-01



| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, . | 6.0 | 118 |
| 2 | Monitoring effectiveness and safety of Tafamidis in transthyretin amyloidosis in Italy: a longitudinal multicenter study in a non-endemic area. Journal of Neurology, 2016, 263, 916-924. | 3.6 | 76 |
| 3 | Transthyretin-Related Familial Amyloid Polyneuropathy (TTR-FAP): A Single-Center Experience in Sicily, an Italian Endemic Area. Journal of Neuromuscular Diseases, 2015, 2, S39-S48. | 2.6 | 67 |
| 4 | Endocardial and Epicardial Deformations in Cardiac Amyloidosis and Hypertrophic Cardiomyopathy. Circulation Journal, 2011, 75, 1200-1208. | 1.6 | 54 |
| 5 | A study of the neuropathy associated with transthyretin amyloidosis (ATTR) in the UK. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 620-627. | 1.9 | 52 |
| 6 | ATTRv amyloidosis Italian Registry: clinical and epidemiological data. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 259-265. | 3.0 | 51 |
| 7 | Transthyretinâ€related familial amyloidotic polyneuropathy: description of a cohort of patients with Leu64 mutation and late onset. Journal of the Peripheral Nervous System, 2012, 17, 385-390. | 3.1 | 41 |
| 8 | Genetic neuromuscular disorders: living the era of a therapeutic revolution. Part 1: peripheral neuropathies. Neurological Sciences, 2019, 40, 661-669. | 1.9 | 32 |
| 9 | Novel outcome measures for Charcotâ~'Marieâ^'Tooth disease: validation and reliability of the 6â€min walk test and StepWatch ^{â,,¢} Activity Monitor and identification of the walking features related to higher quality of life. European Journal of Neurology, 2016, 23, 1343-1350. | 3.3 | 26 |
| 10 | Non-coding variants contribute to the clinical heterogeneity of TTR amyloidosis. European Journal of Human Genetics, 2017, 25, 1055-1060. | 2.8 | 23 |
| 11 | Charcot–Marie–Tooth 2F: phenotypic presentation of the Arg136Leu HSP27 mutation in a multigenerational family. Neurological Sciences, 2015, 36, 1003-1006. | 1.9 | 18 |
| 12 | Phenotypic variability of TTR Val122Ile mutation: a Caucasian patient with axonal neuropathy and normal heart. Neurological Sciences, 2017, 38, 525-526. | 1.9 | 15 |
| 13 | 6MWT performance correlates with peripheral neuropathy but not with cardiac involvement in patients with hereditary transthyretin amyloidosis (hATTR). Neuromuscular Disorders, 2019, 29, 213-220. | 0.6 | 14 |
| 14 | Unusual features of central nervous system involvement in <scp>CMTX</scp> associated with a novel mutation of <scp><i>GJB1</i></scp> gene. Journal of the Peripheral Nervous System, 2012, 17, 407-411. | 3.1 | 13 |
| 15 | Parenteral nutrition improves nutritional status, autonomic symptoms and quality of life in transthyretin amyloid polyneuropathy. Neuromuscular Disorders, 2016, 26, 374-377. | 0.6 | 13 |
| 16 | Subacute inflammatory demyelinating polyneuropathy disclosed by massive nerve root enhancement in CMT1A. Muscle and Nerve, 2012, 45, 451-452. | 2.2 | 10 |
| 17 | Psychosocial impact of sport activity in neuromuscular disorders. Neurological Sciences, 2020, 41, 2561-2567. | 1.9 | 8 |
| 18 | Autonomic Involvement in Subacute and Chronic Immune-Mediated Neuropathies. Autoimmune Diseases, 2013, 2013, 1-7. | 0.6 | 6 |

| # | Article | IF | CITATIONS |
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| 19 | Are novel outcome measures for Charcot–Marie–Tooth disease sensitive to change? The 6-minute walk test and StepWatchâ,¢ Activity Monitor in a 12-month longitudinal study. Neuromuscular Disorders, 2019, 29, 310-316. | 0.6 | 6 |
| 20 | Unilateral hyperhidrosis as persistently isolated feature of syringomyelia and Arnold Chiari type 1. Neurological Sciences, 2018, 39, 1607-1608. | 1.9 | 3 |
| 21 | Considerable post-partum worsening in a patient with CMT2E. Neurological Sciences, 2013, 34, 1813-1814. | 1.9 | 1 |