

# Tommaso F Nicoletti

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6862222/publications.pdf>

Version: 2024-02-01

28  
papers

305  
citations

1163117

8  
h-index

888059

17  
g-index

28  
all docs

28  
docs citations

28  
times ranked

624  
citing authors

#	ARTICLE	IF	CITATIONS
1	Muscle magnetic resonance imaging in myotonic dystrophy type 1 (DM1): Refining muscle involvement and implications for clinical trials. <i>European Journal of Neurology</i> , 2022, 29, 843-854.	3.3	10
2	Elevated serum Neurofilament Light chain (NfL) as a potential biomarker of neurological involvement in Myotonic Dystrophy type 1 (DM1). <i>Journal of Neurology</i> , 2022, .	3.6	3
3	Clinical characteristics of metabolic associated fatty liver disease (MAFLD) in subjects with myotonic dystrophy type 1 (DM1). <i>Digestive and Liver Disease</i> , 2021, 53, 1451-1457.	0.9	6
4	Application of a Clinical Workflow May Lead to Increased Diagnostic Precision in Hereditary Spastic Paraplegias and Cerebellar Ataxias: A Single Center Experience. <i>Brain Sciences</i> , 2021, 11, 246.	2.3	10
5	Eyelid closing and opening disorders in patients with unilateral brain lesions: A case report with video neuroimage and a systematic review of the literature. <i>Journal of Clinical Neuroscience</i> , 2021, 87, 69-73.	1.5	3
6	Sex differences in Lemierre syndrome: Individual patient-level analysis. <i>Thrombosis Research</i> , 2021, 202, 36-39.	1.7	3
7	Patients with Lemierre syndrome have a high risk of new thromboembolic complications, clinical sequelae and death: an analysis of 712 cases. <i>Journal of Internal Medicine</i> , 2021, 289, 325-339.	6.0	61
8	Comment on: Clinico-radiologic features and therapeutic strategies in tumefactive demyelination: a retrospective analysis of 50 consecutive cases. <i>Therapeutic Advances in Neurological Disorders</i> , 2021, 14, 175628642110500.	3.5	2
9	The role of the neurologist in the diagnostic route of HSP and cerebellar ataxias in the next generation sequencing era: A single center experience. <i>Journal of the Neurological Sciences</i> , 2021, 429, 118282.	0.6	0
10	A man with sarcoidosis and slurred speech. <i>European Journal of Neurology</i> , 2020, 27, e7-e8.	3.3	1
11	Letter: prevalence and patterns of gastrointestinal symptoms in a large Western cohort of patients with COVID-19. <i>Alimentary Pharmacology and Therapeutics</i> , 2020, 52, 902-903.	3.7	9
12	Response to "Autosomal recessive axonal neuropathy caused by HINT1 mutation: New association of a psychiatric disorder to the neurological phenotype". <i>Neuromuscular Disorders</i> , 2020, 30, 265-266.	0.6	4
13	Assessment of neurological manifestations in hospitalized patients with COVID-19. <i>European Journal of Neurology</i> , 2020, 27, 2322-2328.	3.3	36
14	NGS-based detection of a novel mutation in PRKCG (SCA14) in sporadic adult-onset ataxia plus dystonic tremor. <i>Neurological Sciences</i> , 2020, 41, 2989-2991.	1.9	3
15	High Prevalence and Gender-Related Differences of Gastrointestinal Manifestations in a Cohort of DM1 Patients: A Perspective, Cross-Sectional Study. <i>Frontiers in Neurology</i> , 2020, 11, 394.	2.4	12
16	Dominus effect: challenging complications of alemtuzumab-related thyroid autoimmunity. <i>Journal of Endocrinological Investigation</i> , 2020, 43, 1159-1161.	3.3	2
17	Myotonic dystrophy type 1 cosegregating with autosomal dominant polycystic kidney disease type 2. <i>Neurological Sciences</i> , 2020, 41, 3761-3763.	1.9	0
18	Compound heterozygosity for an expanded (GAA) and a (GAAGGA) repeat at FXN locus: from a diagnostic pitfall to potential clues to the pathogenesis of Friedreich ataxia. <i>Neurogenetics</i> , 2020, 21, 279-287.	1.4	2

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19	Liver involvement is not associated with mortality: results from a large cohort of SARS-CoV-2 positive patients. <i>Alimentary Pharmacology and Therapeutics</i> , 2020, 52, 1060-1068.	3.7	76
20	Gastrointestinal symptoms and digestive comorbidities in an Italian cohort of patients with COVID-19. <i>European Review for Medical and Pharmacological Sciences</i> , 2020, 24, 7506-7511.	0.7	19
21	Cerebral Fat Embolism: A Rare Cause of Juvenile Stroke. <i>Neurology India</i> , 2020, 68, 1263.	0.4	0
22	Reply to the letter entitled "Predictors of respiratory impairment in patients with myotonic dystrophy type 1". <i>Journal of the Neurological Sciences</i> , 2019, 403, 166-167.	0.6	0
23	Prevalence and predictor factors of respiratory impairment in a large cohort of patients with Myotonic Dystrophy type 1 (DM1): A retrospective, cross sectional study. <i>Journal of the Neurological Sciences</i> , 2019, 399, 118-124.	0.6	31
24	Muscle hypertrophy in amyloid myopathy. <i>Neuromuscular Disorders</i> , 2019, 29, 150-151.	0.6	3
25	Teaching NeuroImages. <i>Neurology</i> , 2019, 92, e1000-e1001.	1.1	4
26	Ventral Longitudinal Intraspinous Fluid Collection Presenting as Upper Limb Amyotrophy. <i>European Neurology</i> , 2018, 80, 126-127.	1.4	3
27	Secondary hypokalemic periodic paralysis as a rare clinical presentation of Conn syndrome. <i>Clinical Neurophysiology</i> , 2018, 129, 2505-2506.	1.5	2
28	Imaging Features of Varicella Zoster Virus Cranial Multiple Mononeuropathies. <i>European Neurology</i> , 2018, 79, 315-316.	1.4	0