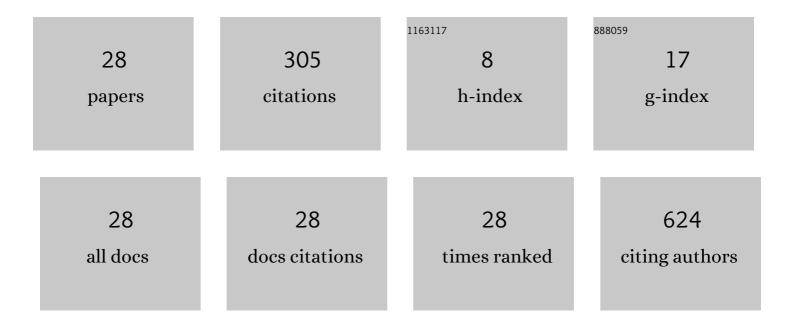
Tommaso F Nicoletti

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6862222/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Liver involvement is not associated with mortality: results from a large cohort of SARSâ€CoVâ€2â€positive patients. Alimentary Pharmacology and Therapeutics, 2020, 52, 1060-1068.	3.7	76
2	Patients with Lemierre syndrome have a high risk of new thromboembolic complications, clinical sequelae and death: an analysis of 712 cases. Journal of Internal Medicine, 2021, 289, 325-339.	6.0	61
3	Assessment of neurological manifestations in hospitalized patients with COVIDâ€19. European Journal of Neurology, 2020, 27, 2322-2328.	3.3	36
4	Prevalence and predictor factors of respiratory impairment in a large cohort of patients with Myotonic Dystrophy type 1 (DM1): A retrospective, cross sectional study. Journal of the Neurological Sciences, 2019, 399, 118-124.	0.6	31
5	Gastrointestinal symptoms and digestive comorbidities in an Italian cohort of patients with COVID-19. European Review for Medical and Pharmacological Sciences, 2020, 24, 7506-7511.	0.7	19
6	High Prevalence and Gender-Related Differences of Gastrointestinal Manifestations in a Cohort of DM1 Patients: A Perspective, Cross-Sectional Study. Frontiers in Neurology, 2020, 11, 394.	2.4	12
7	Application of a Clinical Workflow May Lead to Increased Diagnostic Precision in Hereditary Spastic Paraplegias and Cerebellar Ataxias: A Single Center Experience. Brain Sciences, 2021, 11, 246.	2.3	10
8	Muscle magnetic resonance imaging in myotonic dystrophy type 1 (DM1): Refining muscle involvement and implications for clinical trials. European Journal of Neurology, 2022, 29, 843-854.	3.3	10
9	Letter: prevalence and patterns of gastrointestinal symptoms in a large Western cohort of patients with COVID-19. Alimentary Pharmacology and Therapeutics, 2020, 52, 902-903.	3.7	9
10	Clinical characteristics of metabolic associated fatty liver disease (MAFLD) in subjects with myotonic dystrophy type 1 (DM1). Digestive and Liver Disease, 2021, 53, 1451-1457.	0.9	6
11	Response to "Autosomal recessive axonal neuropathy caused by HINT1 mutation: New association of a psychiatric disorder to the neurological phenotype― Neuromuscular Disorders, 2020, 30, 265-266.	0.6	4
12	Teaching NeuroImages. Neurology, 2019, 92, e1000-e1001.	1.1	4
13	Ventral Longitudinal Intraspinal Fluid Collection Presenting as Upper Limb Amyotrophy. European Neurology, 2018, 80, 126-127.	1.4	3
14	Muscle hypertrophy in amyloid myopathy. Neuromuscular Disorders, 2019, 29, 150-151.	0.6	3
15	NGS-based detection of a novel mutation in PRKCG (SCA14) in sporadic adult-onset ataxia plus dystonic tremor. Neurological Sciences, 2020, 41, 2989-2991.	1.9	3
16	Eyelid closing and opening disorders in patients with unilateral brain lesions: A case report with video neuroimage and a systematic review of the literature. Journal of Clinical Neuroscience, 2021, 87, 69-73.	1.5	3
17	Sex differences in Lemierre syndrome: Individual patient-level analysis. Thrombosis Research, 2021, 202, 36-39.	1.7	3
18	Elevated serum Neurofilament Light chain (NfL) as a potential biomarker of neurological involvement in Myotonic Dystrophy type 1 (DM1). Journal of Neurology, 2022, , .	3.6	3

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#	Article	IF	CITATIONS
19	Secondary hypokalemic periodic paralysis as a rare clinical presentation of Conn syndrome. Clinical Neurophysiology, 2018, 129, 2505-2506.	1.5	2
20	Dominus effect: challenging complications of alemtuzumab-related thyroid autoimmunity. Journal of Endocrinological Investigation, 2020, 43, 1159-1161.	3.3	2
21	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. Neurogenetics, 2020, 21, 279-287.	1.4	2
22	Comment on: Clinico-radiologic features and therapeutic strategies in tumefactive demyelination: a retrospective analysis of 50 consecutive cases. Therapeutic Advances in Neurological Disorders, 2021, 14, 175628642110500.	3.5	2
23	A man with sarcoidosis and slurred speech. European Journal of Neurology, 2020, 27, e7-e8.	3.3	1
24	Imaging Features of Varicella Zoster Virus Cranial Multiple Mononeuropathies. European Neurology, 2018, 79, 315-316.	1.4	0
25	Reply to the letter entitled "Predictors of respiratory impairment in patients with myotonic dystrophy type 1― Journal of the Neurological Sciences, 2019, 403, 166-167.	0.6	0
26	Myotonic dystrophy type 1 cosegregating with autosomal dominant polycystic kidney disease type 2. Neurological Sciences, 2020, 41, 3761-3763.	1.9	0
27	The role of the neurologist in the diagnostic route of HSP and cerebellar ataxias in the next generation sequencing era: A single center experience. Journal of the Neurological Sciences, 2021, 429, 118282.	0.6	0
28	Cerebral Fat Embolism: A Rare Cause of Juvenile Stroke. Neurology India, 2020, 68, 1263.	0.4	0