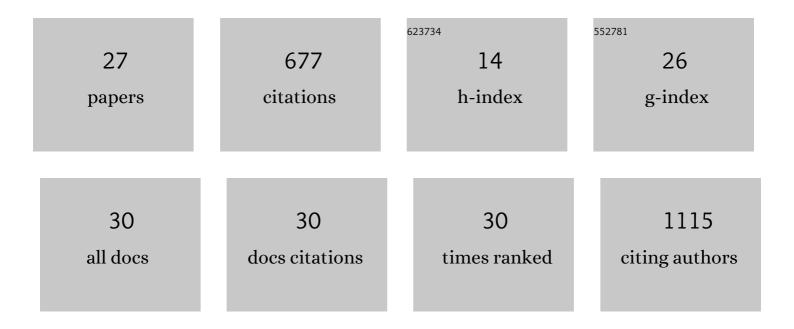
Giuseppe Cardillo

List of Publications by Year in descending order

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CHISEDDE CARDILLO

#	Article	IF	CITATIONS
1	Antithrombotic and Anti-Inflammatory Effects of Fondaparinux and Enoxaparin in Hospitalized COVID-19 Patients: The FONDENOXAVID Study. Journal of Blood Medicine, 2021, Volume 12, 69-75.	1.7	16
2	COVID-19 After Lung Resection in Northern Italy. Seminars in Thoracic and Cardiovascular Surgery, 2021, , .	0.6	5
3	Clinical Differences between COVID-19 and a COVID-Like Syndrome. Journal of Clinical Medicine, 2021, 10, 2519.	2.4	6
4	The Impact of Risk-Adjusted Heparin Regimens on the Outcome of Patients with COVID-19 Infection. A Prospective Cohort Study. Viruses, 2021, 13, 1720.	3.3	9
5	Pathophysiology of Vaccine-Induced Prothrombotic Immune Thrombocytopenia (VIPIT) and Vaccine-Induced Thrombocytopenic Thrombosis (VITT) and Their Diagnostic Approach in Emergency. Medicina (Lithuania), 2021, 57, 997.	2.0	9
6	Possible Adrenal Involvement in Long COVID Syndrome. Medicina (Lithuania), 2021, 57, 1087.	2.0	22
7	Fondaparinux Use in Patients With COVID-19: A Preliminary Multicenter Real-World Experience. Journal of Cardiovascular Pharmacology, 2020, 76, 369-371.	1.9	24
8	Thromboprofilaxys With Fondaparinux vs. Enoxaparin in Hospitalized COVID-19 Patients: A Multicenter Italian Observational Study. Frontiers in Medicine, 2020, 7, 569567.	2.6	21
9	Clotting Factors in COVID-19: Epidemiological Association and Prognostic Values in Different Clinical Presentations in an Italian Cohort. Journal of Clinical Medicine, 2020, 9, 1371.	2.4	63
10	Prognostic Value of Fibrinogen among COVID-19 Patients Admitted to an Emergency Department: An Italian Cohort Study. Journal of Clinical Medicine, 2020, 9, 4134.	2.4	28
11	Twelve Novel Mutations in the <i>SLC26A3</i> Gene in 17 Sporadic Cases of Congenital Chloride Diarrhea. Journal of Pediatric Gastroenterology and Nutrition, 2017, 65, 26-30.	1.8	9
12	Anthropometric characteristics of young Italian tennis players. Journal of Human Sport and Exercise, 2017, 12, .	0.4	4
13	Intracranial pressure in unresponsive chronic migraine. Journal of Neurology, 2014, 261, 1365-1373.	3.6	44
14	Erythrocytosis after liver transplantation: The experience of a university hospital. Liver Transplantation, 2013, 19, 420-424.	2.4	1
15	Extensive Molecular Analysis of Patients Bearing CFTR-Related Disorders. Journal of Molecular Diagnostics, 2012, 14, 81-89.	2.8	52
16	High prevalence of bilateral transverse sinus stenosis-associated IIHWOP in unresponsive chronic headache sufferers: Pathogenetic implications in primary headache progression. Cephalalgia, 2011, 31, 763-765.	3.9	22
17	Congenital Diarrheal Disorders: Improved Understanding of Gene Defects Is Leading to Advances in Intestinal Physiology and Clinical Management. Journal of Pediatric Gastroenterology and Nutrition, 2010, 50, 360-366.	1.8	73
18	PA55 GENOTYPE PREDICTS THE RESPONSE TO THERAPY IN CHILDREN WITH CONGENITAL CHLORIDE DIARRHEA. Digestive and Liver Disease, 2010, 42, S363-S364.	0.9	0

GIUSEPPE CARDILLO

#	Article	IF	CITATIONS
19	Three Novel CFTR Polymorphic Repeats Improve Segregation Analysis for Cystic Fibrosis. Clinical Chemistry, 2009, 55, 1372-1379.	3.2	36
20	Functional and structural characterization of novel mutations and genotype–phenotype correlation in 51 phenylalanine hydroxylase deficient families from Southern Italy. FEBS Journal, 2009, 276, 2048-2059.	4.7	29
21	Buerger`s Disease and Hyperhomocysteinemia: Is there a Relationship?. The Open Atherosclerosis & Thrombosis Journal, 2009, 1, 6-8.	0.4	2
22	Different outcome of six homozygotes for prothrombin A20210A gene variant. Journal of Translational Medicine, 2008, 6, 36.	4.4	4
23	Epidemiology and a novel procedure for large scale analysis of CFTR rearrangements in classic and atypical CF patients: A multicentric Italian study. Journal of Cystic Fibrosis, 2008, 7, 347-351.	0.7	47
24	Five human phenylalanine hydroxylase proteins identified in mild hyperphenylalaninemia patients are disease-causing variants. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2008, 1782, 378-384.	3.8	11
25	MOLECULAR ANALYSIS OF GENES ENCODING CFTR INTERACTORS OF SLC26 FAMILY IN CF PATIENTS: PRELIMINARY RESULTS. Journal of Cystic Fibrosis, 2008, 7, S12-S13.	0.7	0
26	Molecular Epidemiology of Phenylalanine Hydroxylase Deficiency in Southern Italy: a 96% Detection Rate with Ten Novel Mutations. Annals of Human Genetics, 2007, 71, 185-193.	0.8	35
27	Butyrate as an effective treatment of congenital chloride diarrhea. Gastroenterology, 2004, 127, 630-634.	1.3	102