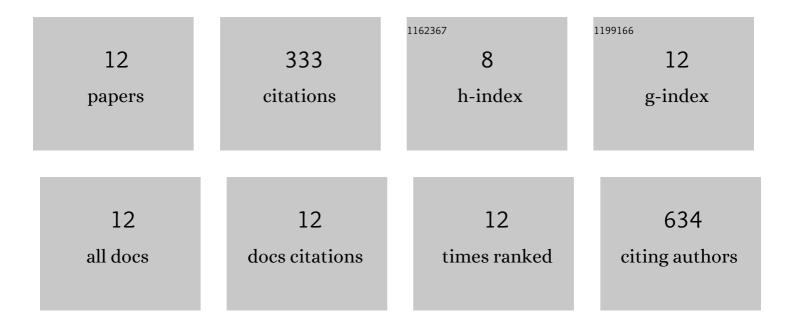
Yllka Kodra

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5700238/publications.pdf Version: 2024-02-01



YULKA KODDA

#	Article	IF	CITATIONS
1	Reflections on the Importance of Cost of Illness Analysis in Rare Diseases: A Proposal. International Journal of Environmental Research and Public Health, 2021, 18, 1101.	1.2	8
2	The Quality Evaluation of Rare Disease Registries—An Assessment of the Essential Features of a Disease Registry. International Journal of Environmental Research and Public Health, 2021, 18, 11968.	1.2	11
3	Shaping the Future of Rare Diseases after a Global Health Emergency: Organisational Points to Consider. International Journal of Environmental Research and Public Health, 2020, 17, 8694.	1.2	9
4	The EuRRECa Project as a Model for Data Access and Governance Policies for Rare Disease Registries That Collect Clinical Outcomes. International Journal of Environmental Research and Public Health, 2020, 17, 8743.	1.2	13
5	Social Economic Costs, Health-Related Quality of Life and Disability in Patients with Cri Du Chat Syndrome. International Journal of Environmental Research and Public Health, 2020, 17, 5951.	1.2	6
6	Rare Pathogenic Copy Number Variation in the 16p11.2 (BP4–BP5) Region Associated with Neurodevelopmental and Neuropsychiatric Disorders: A Review of the Literature. International Journal of Environmental Research and Public Health, 2020, 17, 9253.	1.2	1
7	Primary Sclerosing Cholangitis: Burden of Disease and Mortality Using Data from the National Rare Diseases Registry in Italy. International Journal of Environmental Research and Public Health, 2020, 17, 3095.	1.2	17
8	Recommendations for Improving the Quality of Rare Disease Registries. International Journal of Environmental Research and Public Health, 2018, 15, 1644.	1.2	116
9	Social/economic costs and health-related quality of life in patients with Duchenne muscular dystrophy in Europe. European Journal of Health Economics, 2016, 17, 19-29.	1.4	59
10	Social/economic costs and quality of life in patients with haemophilia in Europe. European Journal of Health Economics, 2016, 17, 53-65.	1.4	53
11	The social burden and quality of life of patients with haemophilia in Italy. Blood Transfusion, 2014, 12 Suppl 3, s567-75.	0.3	22
12	The Italian National Rare Diseases Registry. Blood Transfusion, 2014, 12 Suppl 3, s606-13.	0.3	18