

# Michele Santoro

## List of Publications by Year in descending order

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Version: 2024-02-01

38  
papers

1,019  
citations

430754

18  
h-index

454834

30  
g-index

47  
all docs

47  
docs citations

47  
times ranked

1699  
citing authors

#	ARTICLE	IF	CITATIONS
1	Recommendations for Improving the Quality of Rare Disease Registries. International Journal of Environmental Research and Public Health, 2018, 15, 1644.	1.2	116
2	Environmental and individual exposure and the risk of congenital anomalies: a review of recent epidemiological evidence. Epidemiologia E Prevenzione, 2018, 42, 1-34.	1.1	93
3	Hazardous waste and health impact: a systematic review of the scientific literature. Environmental Health, 2017, 16, 107.	1.7	90
4	Long-term survival of children born with congenital anomalies: A systematic review and meta-analysis of population-based studies. PLoS Medicine, 2020, 17, e1003356.	3.9	63
5	Cancer Mortality in an Area of Campania (Italy) Characterized by Multiple Toxic Dumping Sites. Annals of the New York Academy of Sciences, 2006, 1076, 449-461.	1.8	58
6	Data Quality in Rare Diseases Registries. Advances in Experimental Medicine and Biology, 2017, 1031, 149-164.	0.8	56
7	Malignant mesothelioma due to non-occupational asbestos exposure from the Italian national surveillance system (ReNaM): epidemiology and public health issues. Occupational and Environmental Medicine, 2015, 72, 648-655.	1.3	52
8	Epidemiological patterns of asbestos exposure and spatial clusters of incident cases of malignant mesothelioma from the Italian national registry. BMC Cancer, 2015, 15, 286.	1.1	45
9	Epidemiology of achondroplasia: A population-based study in Europe. American Journal of Medical Genetics, Part A, 2019, 179, 1791-1798.	0.7	33
10	Risk perception and access to environmental information in four areas in Italy affected by natural or anthropogenic pollution. Environment International, 2016, 95, 8-15.	4.8	32
11	Cluster analysis of mortality and malformations in the Provinces of Naples and Caserta (Campania) Tj ETQq1 1 0.784314 rgBT /Overlock 0.2 26	0.7	26
12	Lifestyle and sociodemographic risk factors for gastroschisis: a systematic review and meta-analysis. Archives of Disease in Childhood, 2020, 105, 756-764.	1.0	25
13	Sex differences for major congenital heart defects in Down Syndrome: A population based study. European Journal of Medical Genetics, 2018, 61, 546-550.	0.7	23
14	Epidemiology of Dandy-Walker Malformation in Europe: A EUROCAT Population-Based Registry Study. Neuroepidemiology, 2019, 53, 169-179.	1.1	23
15	Congenital heart disease in live-born children: incidence, distribution, and yearly changes in the Campania Region. Journal of Cardiovascular Medicine, 2008, 9, 368-374.	0.6	22
16	Rare Disease Registries Classification and Characterization: A Data Mining Approach. Public Health Genomics, 2015, 18, 113-122.	0.6	21
17	Linking a European cohort of children born with congenital anomalies to vital statistics and mortality records: A EUROLINKCAT study. PLoS ONE, 2021, 16, e0256535.	1.1	21
18	Participatory health impact assessment used to support decision-making in waste management planning: A replicable experience from Italy. Waste Management, 2017, 59, 557-566.	3.7	20

#	ARTICLE	IF	CITATIONS
19	Ten-Year Survival of Children With Congenital Anomalies: A European Cohort Study. <i>Pediatrics</i> , 2022, 149, .	1.0	18
20	The Quality of Rare Disease Registries: Evaluation and Characterization. <i>Public Health Genomics</i> , 2016, 19, 108-115.	0.6	16
21	Epidemiology of Pierreâ€™Robin sequence in Europe: A populationâ€™based EUROCAT study. <i>Paediatric and Perinatal Epidemiology</i> , 2021, 35, 530-539.	0.8	13
22	Adverse reproductive outcomes associated with exposure to a municipal solid waste incinerator. <i>Annali Dell'Istituto Superiore Di Sanita</i> , 2016, 52, 576-581.	0.2	12
23	Survival of patients with rare diseases: a population-based study in Tuscany (Italy). <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 275.	1.2	11
24	Mesothelioma incidence in the neighbourhood of an asbestos-cement plant located in a national priority contaminated site. <i>Annali Dell'Istituto Superiore Di Sanita</i> , 2014, 50, 322-7.	0.2	11
25	Temporal and geographical variations in survival of children born with congenital anomalies in Europe: A multiâ€™registry cohort study. <i>Paediatric and Perinatal Epidemiology</i> , 2022, 36, 792-803.	0.8	10
26	Epidemiology of systemic sclerosis: a multi-database population-based study in Tuscany (Italy). <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 90.	1.2	9
27	Prevalence Estimates of Rare Congenital Anomalies by Integrating Two Population-Based Registries in Tuscany, Italy. <i>Public Health Genomics</i> , 2017, 20, 229-234.	0.6	8
28	Congenital Anomalies in Contaminated Sites: A Multisite Study in Italy. <i>International Journal of Environmental Research and Public Health</i> , 2017, 14, 292.	1.2	8
29	Survival of children with rare structural congenital anomalies: a multi-registry cohort study. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, 142.	1.2	8
30	Respiratory Symptoms in Relation to Living near a Crude Oil First Treatment Plant in Italy: A Cross-Sectional Study. <i>International Journal of Environmental Research and Public Health</i> , 2018, 15, 2636.	1.2	4
31	Orphan Drug Use in Patients With Rare Diseases: A Population-Based Cohort Study. <i>Frontiers in Pharmacology</i> , 2022, 13, .	1.6	4
32	Methods and data needs to assess health impacts of chemicals in industrial contaminated sites. <i>Epidemiologia E Prevenzione</i> , 2019, 43, 223-237.	1.1	3
33	Environmental and health data needed to develop national surveillance systems in industrially contaminated sites. <i>Epidemiologia E Prevenzione</i> , 2018, 42, 11-20.	1.1	3
34	Association between maternal body mass index and congenital anomalies: A caseâ€™control study in Tuscany (Italy). <i>Birth Defects Research</i> , 2022, 114, 116-123.	0.8	2
35	Healthcare Burden of Rare Diseases: A Population-Based Study in Tuscany (Italy). <i>International Journal of Environmental Research and Public Health</i> , 2022, 19, 7553.	1.2	2
36	Characterization and classification of Rare Disease Registries by using exploratory data analyses. <i>Orphanet Journal of Rare Diseases</i> , 2014, 9, P4.	1.2	1

#	ARTICLE	IF	CITATIONS
37	Sociodemographic Differences in Prenatal Diagnosis of Chromosomal Anomalies: A Population-Based Study. <i>Frontiers in Pediatrics</i> , 2021, 9, 630363.	0.9	1
38	Cluster Analysis of Mortality in an Area of Campania Region (Italy), with Intense Environmental Pressure due to Waste. <i>Epidemiology</i> , 2009, 20, S85.	1.2	0