Anke Rissmann

List of Publications by Year
in descending order

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46	2,025	304743	254184
papers	citations	22 h-index	43 g-index

49	49	49	2611
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Twenty-year trends in the prevalence of Down syndrome and other trisomies in Europe: impact of maternal age and prenatal screening. European Journal of Human Genetics, 2013, 21, 27-33.	2.8	282
2	Long term trends in prevalence of neural tube defects in Europe: population based study. BMJ, The, 2015, 351, h5949.	6.0	180
3	Prenatal diagnosis and prevalence of critical congenital heart defects: an international retrospective cohort study. BMJ Open, 2019, 9, e028139.	1.9	126
4	Estimating Global Burden of Disease due to congenital anomaly: an analysis of European data. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2018, 103, F22-F28.	2.8	122
5	Prevalence of esophageal atresia among 18 international birth defects surveillance programs. Birth Defects Research Part A: Clinical and Molecular Teratology, 2012, 94, 893-899.	1.6	119
6	Paper 6: EUROCAT member registries: Organization and activities. Birth Defects Research Part A: Clinical and Molecular Teratology, 2011, 91, S51-S100.	1.6	107
7	Spectrum of congenital anomalies in pregnancies with pregestational diabetes. Birth Defects Research Part A: Clinical and Molecular Teratology, 2012, 94, 134-140.	1.6	97
8	Epidemiology of hypospadias in Europe: a registry-based study. World Journal of Urology, 2015, 33, 2159-2167.	2.2	88
9	Congenital anomalies associated with trisomy 18 or trisomy 13: A registryâ€based study in 16 european countries, 2000â€“2011. American Journal of Medical Genetics, Part A, 2015, 167, 3062-3069.	1.2	68
10	Selective serotonin reuptake inhibitor antidepressant use in first trimester pregnancy and risk of specific congenital anomalies: a European register-based study. European Journal of Epidemiology, 2015, 30, 1187-1198.	5.7	67
11	Trisomy 13 and 18â€”Prevalence and mortalityâ€”A multiâ€registry population based analysis. American Journal of Medical Genetics, Part A, 2019, 179, 2382-2392.	1.2	59
12	Prevalence of microcephaly in Europe: population based study. BMJ, The, 2016, 354, i4721.	6.0	57
13	Prevalence and mortality in children with congenital diaphragmatic hernia: a multicountry study. Annals of Epidemiology, 2021, 56, 61-69.e3.	1.9	52
14	Beta-Blocker Use in Pregnancy and Risk of Specific Congenital Anomalies: A European Case-Malformed Control Study. Drug Safety, 2018, 41, 415-427.	3.2	46
15	A sustainable solution for the activities of the European network for surveillance of congenital anomalies: EUROCAT as part of the EU Platform on Rare Diseases Registration. European Journal of Medical Genetics, 2018, 61, 513-517.	1.3	45
16	Epidemiology of achondroplasia: A populationâ€based study in Europe. American Journal of Medical Genetics, Part A, 2019, 179, 1791-1798.	1.2	33
17	Stillbirth and neonatal mortality in pregnancies complicated by major congenital anomalies: Findings from a large European cohort. Prenatal Diagnosis, 2017, 37, 1100-1111.	2.3	32
18	EUROlinkCAT protocol for a European population-based data linkage study investigating the survival, morbidity and education of children with congenital anomalies. BMJ Open, 2021, 11, e047859.	1.9	31

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19	Spectrum of congenital anomalies among VACTERL cases: a EUROCAT population-based study. Pediatric Research, 2020, 87, 541-549.	2.3	30
20	From Inception to ConcePTION: Genesis of a Network to Support Better Monitoring and Communication of Medication Safety During Pregnancy and Breastfeeding. Clinical Pharmacology and Therapeutics, 2022, 111, 321-331.	4.7	30
21	Epidemiology of septo-optic dysplasia with focus on prevalence and maternal age â€“ A EUROCAT study. European Journal of Medical Genetics, 2018, 61, 483-488.	1.3	26
22	Assisted reproductive techniques and the risk of anorectal malformations: a German case-control study. Orphanet Journal of Rare Diseases, 2012, 7, 65.	2.7	24
23	Epidemiology of congenital cerebral anomalies in Europe: a multicentre, population-based EUROCAT study. Archives of Disease in Childhood, 2019, 104, 1181-1187.	1.9	24
24	Congenital clubfoot in Europe: A populationâ€based study. American Journal of Medical Genetics, Part A, 2019, 179, 595-601.	1.2	24
25	Epidemiology of Dandy-Walker Malformation in Europe: A EUROCAT Population-Based Registry Study. Neuroepidemiology, 2019, 53, 169-179.	2.3	23
26	Analysis of Mortality among Neonates and Children with Spina Bifida: An International Registryâ€Based Study, 2001â€2012. Paediatric and Perinatal Epidemiology, 2019, 33, 436-448.	1.7	23
27	Prevention of Neural Tube Defects in Europe: A Public Health Failure. Frontiers in Pediatrics, 2021, 9, 647038.	1.9	23
28	Linking a European cohort of children born with congenital anomalies to vital statistics and mortality records: A EUROlinkCAT study. PLoS ONE, 2021, 16, e0256535.	2.5	21
29	Multicentre approach to epidemiological aspects of craniosynostosis in Germany. British Journal of Oral and Maxillofacial Surgery, 2018, 56, 881-886.	0.8	16
30	HSPA6: A new autosomal recessive candidate gene for the VATER/VACTERL malformation spectrum. Birth Defects Research, 2019, 111, 591-597.	1.5	15
31	A multiâ€country study of prevalence and early childhood mortality among children with omphalocele. Birth Defects Research, 2020, 112, 1787-1801.	1.5	14
32	Maternal risk factors for the <scp>VACTERL</scp> association: A <scp>EUROCAT</scp> caseâ€“control study. Birth Defects Research, 2020, 112, 688-698.	1.5	14
33	Epidemiology of Pierreâ€Robin sequence in Europe: A populationâ€based EUROCAT study. Paediatric and Perinatal Epidemiology, 2021, 35, 530-539.	1.7	13
34	Signal Detection in EUROmediCAT: Identification and Evaluation of Medicationâ€“Congenital Anomaly Associations and Use of VigiBase as a Complementary Source of Reference. Drug Safety, 2021, 44, 765-785.	3.2	11
35	Temporal and geographical variations in survival of children born with congenital anomalies in Europe: A multiâ€registry cohort study. Paediatric and Perinatal Epidemiology, 2022, 36, 792-803.	1.7	10
36	Population-based cross-sectional study to assess newborn hearing screening program in central Germany. International Journal of Pediatric Otorhinolaryngology, 2018, 107, 110-120.	1.0	9

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37	Congenital cytomegalovirus infection in Central Germany: an underestimated risk. Archives of Gynecology and Obstetrics, 2017, 296, 231-240.	1.7	8
38	Survival of infants born with esophageal atresia among 24 international birth defects surveillance programs. Birth Defects Research, 2021, 113, 945-957.	1.5	8
39	Periconceptional folic acid supplement use among women of reproductive age and its determinants in central rural Germany: Results from a cross sectional study. Birth Defects Research, 2020, 112, 1057-1066.	1.5	8
40	Methadone, Pierre Robin sequence and other congenital anomalies: caseâ€“control study. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2020, 105, 151-157.	2.8	7
41	Prevalence of valproate syndrome in Europe from 2005 to 2014: A registry based multi-centre study. European Journal of Medical Genetics, 2018, 61, 479-482.	1.3	3
42	Estimated Prevalence of Harmful Alcohol Consumption in Pregnant and Nonpregnant Women in Saxonyâ€Anhalt (NorthEast Germany) Using Biomarkers. Alcoholism: Clinical and Experimental Research, 2021, 45, 819-827.	2.4	3
43	The impact of neuropsychiatric disease on fetal growth: a caseâ€“control study. Archives of Gynecology and Obstetrics, 2019, 300, 1591-1600.	1.7	1
44	Analysis of early neonatal case fatality rate among newborns with congenital hydrocephalus, a 2000â€“2014 <scp>multiâ€country registryâ€based</scp> study. Birth Defects Research, 2022, 114, 631-644.	1.5	1
45	Gastrostomy and congenital anomalies: a European population-based study. BMJ Paediatrics Open, 2022, 6, e001526.	1.4	1
46	A Multicountry Analysis of Prevalence and Mortality among Neonates and Children with Bladder Exstrophy. American Journal of Perinatology, 0, , .	1.4	0