Joan K Morris

List of Publications by Year in descending order

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23879 8878 24,160 254 60 150 citations h-index g-index papers 263 263 263 30469 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Ten-Year Survival of Children With Congenital Anomalies: A European Cohort Study. Pediatrics, 2022, 149, .	1.0	18
2	COVIDâ€19 in pregnancyâ€"what study designs can we use to assess the risk of congenital anomalies in relation to COVIDâ€19 disease, treatment and vaccination?. Paediatric and Perinatal Epidemiology, 2022, 36, 493-507.	0.8	8
3	Survival of children with rare structural congenital anomalies: a multi-registry cohort study. Orphanet Journal of Rare Diseases, 2022, 17, 142.	1.2	8
4	Multi-marker risk-based screening for prostate cancer. Journal of Medical Screening, 2022, , 096914132210764.	1.1	0
5	Blood pressure meta-analysis highlights an implementation gap. Lancet, The, 2022, 399, 1379-1380.	6.3	1
6	Prescription of cardiovascular medication in children with congenital heart defects across six European Regions from 2000 to 2014: data from the EUROlinkCAT population-based cohort study. BMJ Open, 2022, 12, e057400.	0.8	2
7	Gastrostomy and congenital anomalies: a European population-based study. BMJ Paediatrics Open, 2022, 6, e001526.	0.6	1
8	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.	0.8	10
9	Size at birth, growth trajectory in early life, and cardiovascular and metabolic risks in early adulthood: EPICure study. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2021, 106, 149-155.	1.4	18
10	Maternal age in the epidemiology of common autosomal trisomies. Prenatal Diagnosis, 2021, 41, 573-583.	1.1	13
11	European trends in mortality in children with congenital anomalies: 2000–2015. Birth Defects Research, 2021, 113, 958-967.	0.8	15
12	Signal Detection in EUROmediCAT: Identification and Evaluation of Medication–Congenital Anomaly Associations and Use of VigiBase as a Complementary Source of Reference. Drug Safety, 2021, 44, 765-785.	1.4	11
13	Prevention of Neural Tube Defects in Europe: A Public Health Failure. Frontiers in Pediatrics, 2021, 9, 647038.	0.9	23
14	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.	0.8	31
15	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
16	Regional differences in short stature in England between 2006 and 2019: A cross-sectional analysis from the National Child Measurement Programme. PLoS Medicine, 2021, 18, e1003760.	3.9	8
17	Prevalence of microcephaly: the Latin American Network of Congenital Malformations 2010–2017. BMJ Paediatrics Open, 2021, 5, e001235.	0.6	2
18	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

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19	Bayesian hierarchical methods in the detection of potentially teratogenic firstâ€trimester medications. Pharmacoepidemiology and Drug Safety, 2020, 29, 337-346.	0.9	O
20	Two under-recognized limitations of number needed to treat. International Journal of Epidemiology, 2020, 49, 359-360.	0.9	3
21	Growth to early adulthood following extremely preterm birth: the EPICure study. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2020, 105, 496-503.	1.4	19
22	Urgent need for folic acid fortification of flour and grains: response to the 2019 UK Government's public consultation. Archives of Disease in Childhood, 2020, 105, 6-9.	1.0	8
23	A systematic review and meta-analyses of pregnancy and fetal outcomes in women with multiple sclerosis: a contribution from the IMI2 ConcePTION project. Journal of Neurology, 2020, 267, 2721-2731.	1.8	29
24	Use of infectious disease surveillance reports to monitor the Zika virus epidemic in Latin America and the Caribbean from 2015 to 2017: strengths and deficiencies. BMJ Open, 2020, 10, e042869.	0.8	9
25	Prevalence of Down's Syndrome in England, 1998–2013. International Journal of Population Data Science, 2020, 5, 1157.	0.1	8
26	Epidemiology of congenital cerebral anomalies in Europe: a multicentre, population-based EUROCAT study. Archives of Disease in Childhood, 2019, 104, 1181-1187.	1.0	24
27	ZikaPLAN: addressing the knowledge gaps and working towards a research preparedness network in the Americas. Global Health Action, 2019, 12, 1666566.	0.7	13
28	Did advice on the prescription of sodium valproate reduce prescriptions to women? An observational study in three European countries between 2007 and 2016. Pharmacoepidemiology and Drug Safety, 2019, 28, 1519-1528.	0.9	15
29	Prescription of antiepileptic medicines including valproate in pregnant women: A study in three European countries. Pharmacoepidemiology and Drug Safety, 2019, 28, 1510-1518.	0.9	18
30	Trisomy 13 and 18—Prevalence and mortality—A multiâ€registry population based analysis. American Journal of Medical Genetics, Part A, 2019, 179, 2382-2392.	0.7	59
31	Haematological management of major bleeding associated with direct oral anticoagulants – UK experience. British Journal of Haematology, 2019, 185, 514-522.	1.2	15
32	Identifying signals of potentially harmful medications in pregnancy: use of the double false discovery rate method to adjust for multiple testing. British Journal of Clinical Pharmacology, 2019, 85, 356-365.	1.1	3
33	Trajectories of behavior, attention, social and emotional problems from childhood to early adulthood following extremely preterm birth: a prospective cohort study. European Child and Adolescent Psychiatry, 2019, 28, 531-542.	2.8	79
34	Insulin analogues use in pregnancy among women with pregestational diabetes mellitus and risk of congenital anomaly: a retrospective population-based cohort study. BMJ Open, 2018, 8, e014972.	0.8	19
35	Early echocardiography does not predict subsequent treatment of symptomatic patent ductus arteriosus in extremely preterm infants. Acta Paediatrica, International Journal of Paediatrics, 2018, 107, 1909-1916.	0.7	5
36	A three-year prospective study of the presentation and clinical outcomes of major bleeding episodes associated with oral anticoagulant use in the UK (ORANGE study). Haematologica, 2018, 103, 738-745.	1.7	30

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37	Low cigarette consumption and risk of coronary heart disease and stroke: meta-analysis of 141 cohort studies in 55 study reports. BMJ: British Medical Journal, 2018, 360, j5855.	2.4	393
38	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.	0.7	45
39	The impact of providing blood to the scene of an accident on transfusion laboratory practice. Transfusion Medicine, 2018, 28, 56-59.	0.5	1
40	An overview of concepts and approaches used in estimating the burden of congenital disorders globally. Journal of Community Genetics, 2018, 9, 347-362.	0.5	19
41	Chromosomal disorders: estimating baseline birth prevalence and pregnancy outcomes worldwide. Journal of Community Genetics, 2018, 9, 377-386.	0.5	33
42	Prenatal reflex DNA screening for trisomies 21, 18, and 13. Genetics in Medicine, 2018, 20, 825-830.	1.1	24
43	Cognitive trajectories from infancy to early adulthood following birth before 26 weeks of gestation: a prospective, population-based cohort study. Archives of Disease in Childhood, 2018, 103, 363-370.	1.0	140
44	Evaluation of stability of directly standardized rates for sparse data using simulation methods. Population Health Metrics, 2018, 16, 19.	1.3	10
45	Estimating the birth prevalence and pregnancy outcomes of congenital malformations worldwide. Journal of Community Genetics, 2018, 9, 387-396.	0.5	48
46	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.	0.7	3
47	Trends in congenital anomalies in Europe from 1980 to 2012. PLoS ONE, 2018, 13, e0194986.	1.1	106
48	Investigation of EEG Activity Compared with Mean Arterial Blood Pressure in Extremely Preterm Infants. Frontiers in Neurology, 2018, 9, 87.	1.1	1
49	Projected number of children with isolated spina bifida or down syndrome in England and Wales by 2020. European Journal of Medical Genetics, 2018, 61, 539-545.	0.7	5
50	EUROCAT: an update on its functions and activities. Journal of Community Genetics, 2018, 9, 407-410.	0.5	28
51	Public health failure in the prevention of neural tube defects: time to abandon the tolerable upper intake level of folate. Public Health Reviews, 2018, 39, 2.	1.3	68
52	Epidemiology of septo-optic dysplasia with focus on prevalence and maternal age – A EUROCAT study. European Journal of Medical Genetics, 2018, 61, 483-488.	0.7	26
53	Beckwith Wiedemann syndrome: A population-based study on prevalence, prenatal diagnosis, associated anomalies and survival in Europe. European Journal of Medical Genetics, 2018, 61, 499-507.	0.7	28
54	Response to Walker. Genetics in Medicine, 2018, 20, 1295-1295.	1.1	4

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55	Starting the polypill: the use of a single age cut-off in males and females. Journal of Medical Screening, 2017, 24, 50-53.	1.1	5
56	Mortality from aortic stenosis: prospective study of serum calcium and phosphate. Journal of Internal Medicine, 2017, 281, 407-411.	2.7	8
57	Risk Factor Models for Neurodevelopmental Outcomes in Children Born Very Preterm or With Very Low Birth Weight: A Systematic Review of Methodology and Reporting. American Journal of Epidemiology, 2017, 185, 601-612.	1.6	55
58	Revised estimates of the risk of fetal loss following a prenatal diagnosis of trisomy 13 or trisomy 18. American Journal of Medical Genetics, Part A, 2017, 173, 953-958.	0.7	18
59	OC-59â€Born at the threshold of viability: the impact on cognitive development into adulthood. , 2017, , .		0
60	The association between Ehlersâ€Danlos syndromeâ€"hypermobility type and gastrointestinal symptoms in university students: a crossâ€sectional study. Neurogastroenterology and Motility, 2017, 29, e12942.	1.6	21
61	The Need to Interpret the Results of a Clinical Trial in the Context of Other Evidence. American Journal of Medicine, 2017, 130, 251-252.	0.6	0
62	Prognostic Factors for Behavioral Problems and Psychiatric Disorders in Children Born Very Preterm or Very Low Birth Weight. Journal of Developmental and Behavioral Pediatrics, 2016, 37, 88-102.	0.6	53
63	Prevention of Late Onset Sepsis and Central-line Associated Blood Stream Infection in Preterm Infants. Pediatric Infectious Disease Journal, 2016, 35, 401-406.	1.1	17
64	Management and outcomes of neonates with down syndrome admitted to neonatal units. Birth Defects Research Part A: Clinical and Molecular Teratology, 2016, 106, 468-474.	1.6	10
65	Prognostic factors for cerebral palsy and motor impairment in children born very preterm or very low birthweight: a systematic review. Developmental Medicine and Child Neurology, 2016, 58, 554-569.	1.1	99
66	Use of hierarchical models to analyze European trends in congenital anomaly prevalence. Birth Defects Research Part A: Clinical and Molecular Teratology, 2016, 106, 480-488.	1.6	3
67	The Reply. American Journal of Medicine, 2016, 129, e303.	0.6	0
68	Cost–benefit analysis of the polypill in the primary prevention of myocardial infarction and stroke. European Journal of Epidemiology, 2016, 31, 415-426.	2.5	22
69	Reply. Journal of Allergy and Clinical Immunology, 2016, 137, 1624-1625.	1.5	0
70	Lamotrigine use in pregnancy and risk of orofacial cleft and other congenital anomalies. Neurology, 2016, 86, 1716-1725.	1.5	59
71	EUROmediCAT signal detection: an evaluation of selected congenital anomalyâ€medication associations. British Journal of Clinical Pharmacology, 2016, 82, 1094-1109.	1.1	17
72	Is a prenatal diagnosis detrimental to the survival of a fetus with trisomy 18?. American Journal of Medical Genetics, Part A, 2016, 170, 850-851.	0.7	0

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73	EUROmediCAT signal detection: a systematic method for identifying potential teratogenic medication. British Journal of Clinical Pharmacology, 2016, 82, 1110-1122.	1.1	10
74	Child–Parent Familial Hypercholesterolemia Screening in Primary Care. New England Journal of Medicine, 2016, 375, 1628-1637.	13.9	250
75	Prevalence of microcephaly in Europe: population based study. BMJ, The, 2016, 354, i4721.	3.0	57
76	Risk of congenital anomalies after exposure to asthma medication in the first trimester of pregnancy $\hat{a} \in \text{``a cohort linkage study. BJOG: an International Journal of Obstetrics and Gynaecology, 2016, 123, 1609-1618.}$	1.1	32
77	Prevention of neural tube defects in the UK: a missed opportunity. Archives of Disease in Childhood, 2016, 101, 604-607.	1.0	46
78	A Meta-analysis of Individual Participant Data Reveals an Association between Circulating Levels of IGF-I and Prostate Cancer Risk. Cancer Research, 2016, 76, 2288-2300.	0.4	117
79	Accuracy of reporting abortions with Down syndrome in England and Wales: a data linkage study. Journal of Public Health, 2016, 38, 170-174.	1.0	4
80	Grapefruit Juice and Statins. American Journal of Medicine, 2016, 129, 26-29.	0.6	57
81	Selective Serotonin Reuptake Inhibitor (SSRI) Antidepressants in Pregnancy and Congenital Anomalies: Analysis of Linked Databases in Wales, Norway and Funen, Denmark. PLoS ONE, 2016, 11, e0165122.	1.1	42
82	Prevalence and severity of dental fluorosis in four English cities. Community Dental Health, 2016, 33, 292-296.	0.2	8
83	Down syndrome birth weight in England and Wales: Implications for clinical practice. American Journal of Medical Genetics, Part A, 2015, 167, 3070-3075.	0.7	19
84	The Association of H1N1 Pandemic Influenza with Congenital Anomaly Prevalence in Europe. Epidemiology, 2015, 26, 853-861.	1.2	5
85	Increasing body weight. International Journal of Clinical Practice, 2015, 69, 148-149.	0.8	0
86	Estimation of sodium excretion should be made as simple as possible, but not simpler. Journal of Hypertension, 2015, 33, 884-886.	0.3	14
87	Congenital anomalies associated with trisomy 18 or trisomy 13: A registryâ€based study in 16 european countries, 2000–2011. American Journal of Medical Genetics, Part A, 2015, 167, 3062-3069.	0.7	68
88	Community water fluoridation and health outcomes in England: a crossâ€sectional study. Community Dentistry and Oral Epidemiology, 2015, 43, 550-559.	0.9	16
89	Occupational Exposure to Hydrazine and Subsequent Risk of Lung Cancer: 50-Year Follow-Up. PLoS ONE, 2015, 10, e0138884.	1.1	13
90	Use of asthma medication during pregnancy and risk of specific congenital anomalies: AÂEuropean case-malformed control study. Journal of Allergy and Clinical Immunology, 2015, 136, 1496-1502.e7.	1.5	67

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91	Detection and investigation of temporal clusters of congenital anomaly in Europe: seven years of experience of the EUROCAT surveillance system. European Journal of Epidemiology, 2015, 30, 1153-1164.	2.5	29
92	Haemoglobin level at birth is associated with short term outcomes and mortality in preterm infants. BMC Medicine, 2015, 13, 16.	2.3	52
93	Prognostic Factors for Poor Cognitive Development in Children Born Very Preterm or With Very Low Birth Weight. JAMA Pediatrics, 2015, 169, 1162.	3.3	271
94	Functional gastrointestinal disorders are associated with the joint hypermobility syndrome in secondary care: a case–control study. Neurogastroenterology and Motility, 2015, 27, 569-579.	1.6	79
95	Prevention of Neural Tube Defects: A Cross-Sectional Study of the Uptake of Folic Acid Supplementation in Nearly Half a Million Women. PLoS ONE, 2014, 9, e89354.	1.1	82
96	Antenatal detection of Edwards (Trisomy 18) and Patau (Trisomy 13) syndrome: England and Wales 2005-2012. Journal of Medical Screening, 2014, 21, 113-119.	1.1	12
97	Quantifying the health benefits of chronic disease prevention: a fresh approach using cardiovascular disease as an example. European Journal of Epidemiology, 2014, 29, 605-612.	2.5	19
98	An Emmonsia Species Causing Disseminated Infection in South Africa. New England Journal of Medicine, 2014, 370, 283-284.	13.9	29
99	Oral anticoagulant agentâ€associated bleeding events reporting system (<scp>ORANGE</scp>) study. British Journal of Haematology, 2014, 167, 274-276.	1.2	1
100	Major congenital anomalies in babies born with Down syndrome: A EUROCAT populationâ€based registry study. American Journal of Medical Genetics, Part A, 2014, 164, 2979-2986.	0.7	57
101	A Prospective Evaluation of Undiagnosed Joint Hypermobility Syndrome in Patients With Gastrointestinal Symptoms. Clinical Gastroenterology and Hepatology, 2014, 12, 1680-1687.e2.	2.4	85
102	Analysis of the Born in Bradford birth cohort. Lancet, The, 2014, 383, 122-123.	6.3	0
103	Birth prevalence and survival of exomphalos in england and wales: 2005 to 2011. Birth Defects Research Part A: Clinical and Molecular Teratology, 2014, 100, 721-725.	1.6	33
104	Preventive Angioplasty in Myocardial Infarction. New England Journal of Medicine, 2014, 370, 280-283.	13.9	13
105	Prevalence and risk of <scp>D</scp> own syndrome in monozygotic and dizygotic multiple pregnancies in <scp>E</scp> urope: implications for prenatal screening. BJOG: an International Journal of Obstetrics and Gynaecology, 2014, 121, 809-820.	1.1	60
106	Epidemiology of multiple congenital anomalies in Europe: A EUROCAT populationâ€based registry study. Birth Defects Research Part A: Clinical and Molecular Teratology, 2014, 100, 270-276.	1.6	64
107	Refining the American guidelines for prevention of cardiovascular disease. Lancet, The, 2014, 383, 598.	6.3	2
108	Randomized Trial of Preventive Angioplasty in Myocardial Infarction. New England Journal of Medicine, 2013, 369, 1115-1123.	13.9	871

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109	Survival of trisomy 18 (Edwards syndrome) and trisomy 13 (Patau Syndrome) in England and Wales: 2004–2011. American Journal of Medical Genetics, Part A, 2013, 161, 2512-2518.	0.7	51
110	Patient safety and estimation of renal function in patients prescribed new oral anticoagulants for stroke prevention in atrial fibrillation: a cross-sectional study. BMJ Open, 2013, 3, e003343.	0.8	67
111	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.	1.4	282
112	Trends in maternal age distribution and the live birth prevalence of Down's syndrome in England and Wales: 1938–2010. European Journal of Human Genetics, 2013, 21, 943-947.	1.4	50
113	The population prevalence of Down's syndrome in England and Wales in 2011. European Journal of Human Genetics, 2013, 21, 1016-1019.	1.4	88
114	Antenatal screening for Down syndrome: A quantitative demonstration of the improvements over the past 20 years. Journal of Health Services Research and Policy, 2013, 18, 195-201.	0.8	4
115	PTU-124â€The Association of the Joint Hypermobility Syndrome with Functional Gastrointestinal Disorders – an Interesting New finding that may Explain Aetiology. Gut, 2013, 62, A97.1-A97.	6.1	О
116	Letters to the editor. Journal of the Royal College of Physicians of Edinburgh, The, 2013, 43, 94-95.	0.2	1
117	Letters to the editor. Journal of the Royal College of Physicians of Edinburgh, The, 2013, 43, 283-285.	0.2	О
118	De novo deletions and duplications detected by array CGH: a study of parental origin in relation to mechanisms of formation and size of imbalance. European Journal of Human Genetics, 2012, 20, 155-160.	1.4	23
119	The value of early second trimester PAPP-A and ADAM12 in screening for pre-eclampsia. Journal of Medical Screening, 2012, 19, 51-54.	1.1	13
120	Cost-effectiveness of diagnosis of high blood pressure in primary care. Lancet, The, 2012, 379, 709-710.	6.3	3
121	Newer anticonvulsants: Lamotrigine. Birth Defects Research Part A: Clinical and Molecular Teratology, 2012, 94, 959-959.	1.6	1
122	Screening for pre-eclampsia using serum placental growth factor and endoglin with Down's syndrome Quadruple test markers. Journal of Medical Screening, 2012, 19, 60-67.	1.1	6
123	Personalized medicine: hope or hype. European Heart Journal, 2012, 33, 1553-1554.	1.0	3
124	Trisomy 21 mosaicism and maternal age. American Journal of Medical Genetics, Part A, 2012, 158A, 2482-2484.	0.7	8
125	Randomized Polypill Crossover Trial in People Aged 50 and Over. PLoS ONE, 2012, 7, e41297.	1.1	128
126	Blood pressures in subjects for life assurance medical examination and the effect of ten minutes recumbent rest. Journal of the Royal College of Physicians of Edinburgh, The, 2012, 42, 205-210.	0.2	0

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127	Body weight reduction to avoid the excess risk of type 2 diabetes. British Journal of General Practice, 2012, 62, e411-e414.	0.7	10
128	The population impact of screening for Down syndrome: audit of 19 326 invasive diagnostic tests in England and Wales in 2008. Prenatal Diagnosis, 2012, 32, 596-601.	1.1	15
129	Cytogenetic and epidemiological findings in Down syndrome: England and Wales 1989–2009. American Journal of Medical Genetics, Part A, 2012, 158A, 1151-1157.	0.7	33
130	Cytological and epidemiological findings in trisomies 13, 18, and 21: England and Wales 2004–2009. American Journal of Medical Genetics, Part A, 2012, 158A, 1145-1150.	0.7	19
131	The evaluation of cascade testing for familial hypercholesterolemia. American Journal of Medical Genetics, Part A, 2012, 158A, 78-84.	0.7	63
132	Aspirin in the prevention of cancer. Lancet, The, 2011, 377, 1649.	6.3	9
133	Assessing Risk Factors as Potential Screening Tests. Archives of Internal Medicine, 2011, 171, 286.	4.3	40
134	Reconciling the Evidence on Serum Homocysteine and Ischaemic Heart Disease: A Meta-Analysis. PLoS ONE, 2011, 6, e16473.	1.1	67
135	Screening for Future Cardiovascular Disease Using Age Alone Compared with Multiple Risk Factors and Age. PLoS ONE, 2011, 6, e18742.	1.1	95
136	Valproic Acid Monotherapy in Pregnancy and Major Congenital Malformations. Obstetrical and Gynecological Survey, 2010, 65, 619-620.	0.2	3
137	The maternal ageâ€specific live birth prevalence of trisomies 13 and 18 compared to trisomy 21 (Down) Tj ETQq1	1.9.78431 1.9.78431	.4 rgBT /Ov 109
138	Increasing body weight: 1989–2008. International Journal of Clinical Practice, 2010, 64, 408-409.	0.8	1
139	Down's syndrome: screening and antenatal diagnosis regionally in England and Wales 1989–2008. Journal of Medical Screening, 2010, 17, 170-175.	1.1	9
140	Case-control analysis of paternal age and trisomic anomalies. Archives of Disease in Childhood, 2010, 95, 893-897.	1.0	27
141	De novo apparently balanced translocations in man are predominantly paternal in origin and associated with a significant increase in paternal age. Journal of Medical Genetics, 2010, 47, 112-115.	1.5	38
142	Intrauterine exposure to carbamazepine and specific congenital malformations: systematic review and case-control study. BMJ: British Medical Journal, 2010, 341, c6581-c6581.	2.4	179
143	Valproic Acid Monotherapy in Pregnancy and Major Congenital Malformations. New England Journal of Medicine, 2010, 362, 2185-2193.	13.9	473
144	Trends in Down's syndrome live births and antenatal diagnoses in England and Wales from 1989 to 2008: analysis of data from the National Down Syndrome Cytogenetic Register. BMJ: British Medical Journal, 2009, 339, b3794-b3794.	2.4	168

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145	Down syndrome and paternal age, a new analysis of case–control data collected in the 1960s. American Journal of Medical Genetics, Part A, 2009, 149A, 1205-1208.	0.7	17
146	Recurrence risks for trisomies 13, 18, and 21. American Journal of Medical Genetics, Part A, 2009, 149A, 2716-2722.	0.7	49
147	Maternal ageâ€specific risk of nonâ€chromosomal anomalies. BJOG: an International Journal of Obstetrics and Gynaecology, 2009, 116, 1111-1119.	1.1	74
148	Combination Therapy Versus Monotherapy in Reducing Blood Pressure: Meta-analysis on 11,000 Participants from 42 Trials. American Journal of Medicine, 2009, 122, 290-300.	0.6	747
149	Use of blood pressure lowering drugs in the prevention of cardiovascular disease: meta-analysis of 147 randomised trials in the context of expectations from prospective epidemiological studies. BMJ: British Medical Journal, 2009, 338, b1665-b1665.	2.4	2,170
150	Body-mass index and cause-specific mortality in 900â€^000 adults: collaborative analyses of 57 prospective studies. Lancet, The, 2009, 373, 1083-1096.	6.3	3,779
151	Maternal Age-Specific Risk of Nonchromosomal Anomalies. Obstetrical and Gynecological Survey, 2009, 64, 650-651.	0.2	0
152	Prenatal screening for serious congenital heart defects using nuchal translucency: a metaâ€analysis. Prenatal Diagnosis, 2008, 28, 1094-1104.	1.1	29
153	The risk of fetal loss following a prenatal diagnosis of trisomy 13 or trisomy 18. American Journal of Medical Genetics, Part A, 2008, 146A, 827-832.	0.7	103
154	Reply to Herlihy and Halliday. European Journal of Human Genetics, 2008, 16, 1174-1174.	1.4	1
155	Is the prevalence of Klinefelter syndrome increasing?. European Journal of Human Genetics, 2008, 16, 163-170.	1.4	189
156	A 16-week, randomized, double-blind, placebo-controlled, crossover trial to quantify the combined effect of an angiotensin-converting enzyme inhibitor and a \hat{l}^2 -blocker on blood pressure reduction. Clinical Therapeutics, 2008, 30, 2030-2039.	1.1	4
157	Meta-analysis audit trail. Lancet, The, 2008, 371, 558.	6.3	0
158	Does lamotrigine use in pregnancy increase orofacial cleft risk relative to other malformations?. Neurology, 2008, 71, 714-722.	1.5	151
159	Commentary: Clustering in Down syndrome. International Journal of Epidemiology, 2008, 37, 1179-1180.	0.9	4
160	Multiple-marker screening for Down's syndrome: a method of assessing the statistical robustness of proposed tests. Journal of Medical Screening, 2008, 15, 55-61.	1,1	4
161	Ascertainment and accuracy of Down syndrome cases reported in congenital anomaly registers in England and Wales. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2008, 94, F23-F27.	1.4	23
162	Paternal age and birth defects: how strong is the association. Human Reproduction, 2007, 22, 2349-2350.	0.4	84

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163	Estimating the risk of Down's syndrome in antenatal screening and the gestation at which this risk applies. Journal of Medical Screening, 2007, 14, 5-7.	1.1	6
164	Prevalence of neural tube defect pregnancies in England and Wales from 1964 to 2004. Journal of Medical Screening, 2007, 14, 55-59.	1.1	33
165	The effect of correlations between screening markers on screening performance. Journal of Medical Screening, 2007, 14, 151-157.	1.1	4
166	Blood cholesterol and vascular mortality by age, sex, and blood pressure: a meta-analysis of individual data from 61 prospective studies with 55â€^000 vascular deaths. Lancet, The, 2007, 370, 1829-1839.	6.3	1,907
167	Truncation limits for CT marker ratios in prenatal screening for Down syndrome. Prenatal Diagnosis, 2007, 27, 187-188.	1.1	1
168	Calcium channel blockers and headache. British Journal of Clinical Pharmacology, 2007, 63, 157-158.	1.1	8
169	The proportions of Down's syndrome pregnancies detected prenatally in England and Wales from 1989 to 2004. Journal of Medical Screening, 2006, 13, 163-165.	1.1	3
170	Validation plots in antenatal screening for Down's syndrome. Journal of Medical Screening, 2006, 13, 166-171.	1.1	7
171	Maternal age-specific fetal loss rates in Down syndrome pregnancies. Prenatal Diagnosis, 2006, 26, 499-504.	1.1	79
172	Cross-trimester marker ratios in prenatal screening for Down syndrome. Prenatal Diagnosis, 2006, 26, 514-523.	1.1	20
173	Screening in early pregnancy for pre-eclampsia using down syndrome quadruple test markers. Prenatal Diagnosis, 2006, 26, 559-564.	1.1	38
174	Cross trimester marker ratios: parameter estimates valid with no inconsistency. Prenatal Diagnosis, 2006, 26, 994-994.	1.1	1
175	Rates of Down syndrome at the upper extreme of maternal age: considerations and recommendations in analysis. Prenatal Diagnosis, 2006, 26, 1091-1091.	1.1	2
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