Angela E Scheuerle

List of Publications by Year
in descending order

44 papers	1,133 citations	567281 15 h-index	414414 32 g-index
45 all docs	45 docs citations	45 times ranked	2275 citing authors

#	Article	IF	Citations
1	High Incidence of Noonan Syndrome Features Including Short Stature and Pulmonic Stenosis in Patients carrying NF1 Missense Mutations Affecting p.Arg1809: Genotypeâ€“Phenotype Correlation. Human Mutation, 2015, 36, 1052-1063.	2.5	143
2	Pregnancy Outcomes After Exposure to Certolizumab Pegol. Arthritis and Rheumatology, 2018, 70, 1399-1407.	5.6	129
3	De Novo Mutations in NALCN Cause a Syndrome Characterized by Congenital Contractures of the Limbs and Face, Hypotonia, and Developmental Delay. American Journal of Human Genetics, 2015, 96, 462-473.	6.2	124
4	Pregnancy outcomes in the omalizumab pregnancy registry and a disease-matched comparator cohort. Journal of Allergy and Clinical Immunology, 2020, 145, 528-536.e1.	2.9	91
5	Association Between Birth Defects and Cancer Risk Among Children and Adolescents in a Population-Based Assessment of 10 Million Live Births. JAMA Oncology, 2019, 5, 1150.	7.1	87
6	Populationâ€based birth defects data in the United States, 2008 to 2012: Presentation of stateâ€specific data and descriptive brief on variability of prevalence. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 972-993.	1.6	73
7	Birth defect classification by organ system: a novel approach to heighten teratogenic signalling in a pregnancy registry. Pharmacoepidemiology and Drug Safety, 2002, 11, 465-475.	1.9	57
8	Antiepileptic drug exposure and major congenital malformations: The role of pregnancy registries. Epilepsy and Behavior, 2007, 11, 277-282.	1.7	45
9	Mandibulofacial Dysostosis with Microcephaly: Mutation and Database Update. Human Mutation, 2016, 37, 148-154.	2.5	45
10	Expanding the clinical and molecular findings in RASA1 capillary malformation-arteriovenous malformation. European Journal of Human Genetics, 2018, 26, 1521-1536.	2.8	42
11	Expanding the Molecular and Clinical Phenotype of SSR4-CDG. Human Mutation, 2015, 36, 1048-1051.	2.5	22
12	Maternal occupational exposure to polycyclic aromatic hydrocarbons and craniosynostosis among offspring in the national birth defects prevention study. Birth Defects Research Part A: Clinical and Molecular Teratology, 2016, 106, 55-60.	1.6	22
13	Are birth defects among Hispanics related to maternal nativity or number of years lived in the United States?. Birth Defects Research Part A: Clinical and Molecular Teratology, 2009, 85, 755-763.	1.6	21
14	Clinical review procedures for the Antiretroviral Pregnancy Registry. Pharmacoepidemiology and Drug Safety, 2004, 13, 529-536.	1.9	17
15	Sociodemographic, health behavioral, and clinical risk factors for anotia/microtia in a population-based case-control study. International Journal of Pediatric Otorhinolaryngology, 2019, 122, 18-26.	1.0	17
16	Prune belly syndrome in surviving males can be caused by Hemizygous missense mutations in the X-linked Filamin A gene. BMC Medical Genetics, 2020, 21, 38.	2.1	16
17	Prenatal exposure to zidovudine and risk for ventricular septal defects and congenital heart defects: data from the Antiretroviral Pregnancy Registry. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2016, 197, 6-10.	1.1	15
18	Levetiracetam Pregnancy Registry: Final results and a review of the impact of registry methodology and definitions on the prevalence of major congenital malformations. Birth Defects Research, 2019, 111, 872-887.	1.5	14

#	Article	IF	Citations
19	A physician survey regarding diagnostic variability among birth defects. American Journal of Medical Genetics, Part A, 2010, 152A, 1594-1598.	1.2	12
20	An additional case of Hennekam lymphangiectasiaâ€“lymphedema syndrome caused by lossâ€ofâ€function mutation in ADAMTS3. American Journal of Medical Genetics, Part A, 2018, 176, 2858-2861.	1.2	12
21	Coâ€occurring defect analysis: A platform for analyzing birth defect coâ€occurrence in registries. Birth Defects Research, 2019, 111, 1356-1364.	1.5	12
22	Clinical Exome Studies Have Inconsistent Coverage. Clinical Chemistry, 2020, 66, 199-206.	3.2	12
23	Preâ€Zika descriptive epidemiology of microcephaly in Texas, 2008â€“2012. Birth Defects Research, 2018, 110, 395-405.	1.5	11
24	Overcoming presynaptic effects of VAMP2 mutations with 4â€aminopyridine treatment. Human Mutation, 2020, 41, 1999-2011.	2.5	11
25	Birth defects that coâ€occur with nonâ€syndromic gastroschisis and omphalocele. American Journal of Medical Genetics, Part A, 2020, 182, 2581-2593.	1.2	9
26	Birth defects and neonatal morbidity caused by teratogen exposure after the embryonic period. Birth Defects Research Part A: Clinical and Molecular Teratology, 2016, 106, 935-939.	1.6	8
27	Likelihood of meeting defined VATER/VACTERL phenotype in infants with esophageal atresia with or without tracheoesophageal fistula. American Journal of Medical Genetics, Part A, 2019, 179, 2202-2206.	1.2	8
28	Selected acculturation factors and birth defects in the National Birth Defects Prevention Study, 1997â€“2011. Birth Defects Research, 2019, 111, 598-612.	1.5	8
29	An additional case of NÃ©storâ€Guillermo progeria syndrome diagnosed in early childhood. American Journal of Medical Genetics, Part A, 2020, 182, 2399-2402.	1.2	8
30	Birth defect co-occurrence patterns in the Texas Birth Defects Registry. Pediatric Research, 2022, 91, 1278-1285.	2.3	8
31	Risk factors and time trends for isolated craniosynostosis. Birth Defects Research, 2021, 113, 43-54.	1.5	7
32	Incontinentia pigmenti in adults. American Journal of Medical Genetics, Part A, 2019, 179, 1415-1419.	1.2	4
33	Patterns of co-occurring birth defects among infants with hypospadias. Journal of Pediatric Urology, 2021, 17, 64.e1-64.e8.	1.1	4
34	A Comprehensive Assessment of Co-occurring Birth Defects among Infants with Non-Syndromic Anophthalmia or Microphthalmia. Ophthalmic Epidemiology, 2021, 28, 428-435.	1.7	4
35	Asymmetric faces: Symbolic, spiritual, and representative. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2021, 187, 278-282.	1.6	3
36	Pharmacovigilance pregnancy data in a large population of patients with chronic inflammatory disease exposed to certolizumab pegol. Therapeutic Advances in Musculoskeletal Disease, 2022, 14, 1759720X2210876.	2.7	3

#	Article	IF	Citations
37	Descriptive epidemiology of birth defects thought to arise by new mutation. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 913-927.	1.6	2
38	Acculturation and selected birth defects among nonâ€Hispanic Blacks in a populationâ€based caseâ€“control study. Birth Defects Research, 2020, 112, 535-554.	1.5	2
39	Patterns of congenital anomalies among individuals with trisomy 13 in Texas. American Journal of Medical Genetics, Part A, 2021, 185, 1787-1793.	1.2	2
40	Temporal trends in diagnoses of congenital microcephaly, Texas Hospital Discharge Diagnoses, 2000â€“2015. Birth Defects Research, 2019, 111, 584-590.	1.5	1
41	74. Maternal Dolutegravir (DTG) Use During Pregnancy and Birth Outcomes: The Antiretroviral Pregnancy Registry (APR). Open Forum Infectious Diseases, 2021, 8, S48-S49.	0.9	1
42	Some Intensification and Refining. Journal of Craniofacial Surgery, 2017, 28, 308.	0.7	0
43	Defect evaluation by infant photographs in a multicenter pharmaceutical clinical trial. Birth Defects Research, 2020, 112, 118-121.	1.5	0
44	Enamel Pit Defects and Taurodontism in a Patient with Ring Chromosome 14 and 47,XXX. Journal of Dentistry for Children, 2017, 84, 39-43.	0.2	0