# Angela E Scheuerle

List of Publications by Year
in descending order



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|---|---|---|---|
| 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 | Birth defect co-occurrence patterns in the Texas Birth Defects Registry. Pediatric Research, 2022, 91, 1278-1285. | 2.3 | 8 |
| 2 | 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 |
| 3 | Patterns of co-occurring birth defects among infants with hypospadias. Journal of Pediatric Urology, 2021, 17, 64.e1-64.e8. | 1.1 | 4 |
| 4 | Risk factors and time trends for isolated craniosynostosis. Birth Defects Research, 2021, 113, 43-54. | 1.5 | 7 |
| 5 | 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 |
| 6 | Asymmetric faces: Symbolic, spiritual, and representative. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2021, 187, 278-282. | 1.6 | 3 |
| 7 | 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 |
| 8 | 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 |
| 9 | 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 |
| 10 | Clinical Exome Studies Have Inconsistent Coverage. Clinical Chemistry, 2020, 66, 199-206. | 3.2 | 12 |
| 11 | Defect evaluation by infant photographs in a multicenter pharmaceutical clinical trial. Birth Defects Research, 2020, 112, 118-121. | 1.5 | 0 |
| 12 | Overcoming presynaptic effects of VAMP2 mutations with 4â€aminopyridine treatment. Human Mutation, 2020, 41, 1999-2011. | 2.5 | 11 |
| 13 | 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 |
| 14 | 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 |
| 15 | 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 |
| 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 | Coâ€occurring defect analysis: A platform for analyzing birth defect coâ€occurrence in registries. Birth Defects Research, 2019, 111, 1356-1364. | 1.5 | 12 |
| 18 | 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 |

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|---|---|---|---|
| 19 | Incontinentia pigmenti in adults. American Journal of Medical Genetics, Part A, 2019, 179, 1415-1419. | 1.2 | 4 |
| 20 | 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 |
| 21 | 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 |
| 22 | 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 |
| 23 | Temporal trends in diagnoses of congenital microcephaly, Texas Hospital Discharge Diagnoses, 2000â€“2015. Birth Defects Research, 2019, 111, 584-590. | 1.5 | 1 |
| 24 | 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 |
| 25 | Pregnancy Outcomes After Exposure to Certolizumab Pegol. Arthritis and Rheumatology, 2018, 70, 1399-1407. | 5.6 | 129 |
| 26 | Preâ€Zika descriptive epidemiology of microcephaly in Texas, 2008â€“2012. Birth Defects Research, 2018, 110, 395-405. | 1.5 | 11 |
| 27 | 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 |
| 28 | Expanding the clinical and molecular findings in RASA1 capillary malformation-arteriovenous malformation. European Journal of Human Genetics, 2018, 26, 1521-1536. | 2.8 | 42 |
| 29 | Some Intensification and Refining. Journal of Craniofacial Surgery, 2017, 28, 308. | 0.7 | 0 |
| 30 | 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 |
| 31 | Mandibulofacial Dysostosis with Microcephaly: Mutation and Database Update. Human Mutation, 2016, 37, 148-154. | 2.5 | 45 |
| 32 | 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 |
| 33 | 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 |
| 34 | 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 |
| 35 | 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 |
| 36 | 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 |

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|---|---|---|---|
| 37 | 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 |
| 38 | Expanding the Molecular and Clinical Phenotype of SSR4-CDG. Human Mutation, 2015, 36, 1048-1051. | 2.5 | 22 |
| 39 | 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 |
| 40 | A physician survey regarding diagnostic variability among birth defects. American Journal of Medical Genetics, Part A, 2010, 152A, 1594-1598. | 1.2 | 12 |
| 41 | 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 |
| 42 | Antiepileptic drug exposure and major congenital malformations: The role of pregnancy registries. Epilepsy and Behavior, 2007, 11, 277-282. | 1.7 | 45 |
| 43 | Clinical review procedures for the Antiretroviral Pregnancy Registry. Pharmacoepidemiology and Drug Safety, 2004, 13, 529-536. | 1.9 | 17 |
| 44 | 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 |