

Daniel T Swarr

List of Publications by Year in descending order

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Version: 2024-02-01

30
papers

1,009
citations

567281

15
h-index

580821

25
g-index

34
all docs

34
docs citations

34
times ranked

1776
citing authors

#	ARTICLE	IF	CITATIONS
1	Detection and impact of genetic disease in a level IV neonatal intensive care unit. <i>Journal of Perinatology</i> , 2022, 42, 580-588.	2.0	6
2	Inflammatory blockade prevents injury to the developing pulmonary gas exchange surface in preterm primates. <i>Science Translational Medicine</i> , 2022, 14, eabl8574.	12.4	10
3	Perinatal Outcomes of Fetuses and Infants Diagnosed with Trisomy 13 or Trisomy 18. <i>Journal of Pediatrics</i> , 2022, 247, 116-123.e5.	1.8	14
4	Beyond diagnostic yield: prenatal exome sequencing results in maternal, neonatal, and familial clinical management changes. <i>Genetics in Medicine</i> , 2021, 23, 909-917.	2.4	21
5	In utero and post-natal development of the human lung and its defence mechanisms. , 2021, , 1-20.		2
6	Identifying Out New Epigenetic Markers in Lung Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 1029-1030.	5.6	0
7	Cell- and tissue-based therapies for lung disease. , 2020, , 1253-1272.		0
8	Molecular Determinants of Lung Morphogenesis. , 2019, , 26-39.e4.		4
9	Making a Genetic Diagnosis in a Level IV Neonatal Intensive Care Unit Population: Who, When, How, and at What Cost?. <i>Journal of Pediatrics</i> , 2019, 213, 211-217.e4.	1.8	26
10	The long noncoding RNA Falcor regulates Foxa2 expression to maintain lung epithelial homeostasis and promote regeneration. <i>Genes and Development</i> , 2019, 33, 656-668.	5.9	30
11	Congenital Cystic Lung Lesions. <i>American Journal of Surgical Pathology</i> , 2019, 43, 47-55.	3.7	43
12	Novel Molecular and Phenotypic Insights into Congenital Lung Malformations. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, 1328-1339.	5.6	42
13	Systematic Review and Meta-analysis: Gene Association Studies in Neonatal Sepsis. <i>American Journal of Perinatology</i> , 2017, 34, 684-692.	1.4	16
14	Accelerating Scientific Advancement for Pediatric Rare Lung Disease Research. Report from a National Institutes of Healthâ€NHLBI Workshop, September 3 and 4, 2015. <i>Annals of the American Thoracic Society</i> , 2016, 13, 385-393.	3.2	9
15	Detection of mutually exclusive mosaicism in a girl with genotypeâ€phenotype discrepancies. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 3091-3095.	1.2	4
16	Mutations in SPECC1L, encoding sperm antigen with calponin homology and coiled-coil domains 1-like, are found in some cases of autosomal dominant Opitz G/BBB syndrome. <i>Journal of Medical Genetics</i> , 2015, 52, 104-110.	3.2	40
17	Lung Endoderm Morphogenesis: Gasping for Form and Function. <i>Annual Review of Cell and Developmental Biology</i> , 2015, 31, 553-573.	9.4	80
18	Long noncoding RNAs are spatially correlated with transcription factors and regulate lung development. <i>Genes and Development</i> , 2014, 28, 1363-1379.	5.9	148

#	ARTICLE	IF	CITATIONS
19	Novel <i>FREM1</i> mutations expand the phenotypic spectrum associated with manitoba-ocular-trichoanal (MOTA) syndrome and bifid nose renal agenesis anorectal malformations (BNAR) syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 473-478.	1.2	23
20	Expanding the differential diagnosis of fetal hydrops: an unusual prenatal presentation of megalencephaly-capillary malformation syndrome. Prenatal Diagnosis, 2013, 33, 1010-1012.	2.3	9
21	Clinical features of three girls with mosaic genome-wide paternal uniparental isodisomy. American Journal of Medical Genetics, Part A, 2013, 161, 1929-1939.	1.2	63
22	Novel <i>FOXF1</i> Mutations in Sporadic and Familial Cases of Alveolar Capillary Dysplasia with Misaligned Pulmonary Veins Imply a Role for its DNA Binding Domain. Human Mutation, 2013, 34, 801-811.	2.5	97
23	Unusual Cardiac Masses in a Newborn with Infantile Pompe Disease. JIMD Reports, 2011, 5, 17-20.	1.5	4
24	Unraveling the complex genetic underpinnings of asthma and allergic disorders. Current Opinion in Allergy and Clinical Immunology, 2010, 10, 434-442.	2.3	17
25	Potocki-Shaffer syndrome: Comprehensive clinical assessment, review of the literature, and proposals for medical management. American Journal of Medical Genetics, Part A, 2010, 152A, 565-572.	1.2	31
26	Fate of the Residual Distal and Proximal Aorta After Acute Type A Dissection Repair Using a Contemporary Surgical Reconstruction Algorithm. Annals of Thoracic Surgery, 2007, 84, 1955-1964.	1.3	231
27	Lewis Acid-Base, Molecular Modeling, and Isotopic Labeling in a Sophomore Inorganic Chemistry Laboratory. Journal of Chemical Education, 2004, 81, 722.	2.3	7
28	Comparison of Alternative Diagnostic Approaches for Managing Appendicitis in Children: The Effect of Disease Prevalence and Spectrum. Pediatrics, 2004, 114, 513-514.	2.1	19
29	The Oxa-Pictet-Spengler Reaction of 1-(3-Furyl)alkan-2-ols. Synthesis, 2002, 2002, 1541-1545.	2.3	13
30	VALIDATION OF THE CRITICAL CARE FAMILY SATISFACTION SURVEY (CCFSS) IN 7 ADULT AND 2 PEDIATRIC ICU ENVIRONMENTS. Critical Care Medicine, 2002, 30, A139.	0.9	0