

Caroline D Robson

List of Publications by Year in descending order

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Version: 2024-02-01

89
papers

3,379
citations

159525

30
h-index

155592

55
g-index

91
all docs

91
docs citations

91
times ranked

3961
citing authors

#	ARTICLE	IF	CITATIONS
1	High dose dexmedetomidine as the sole sedative for pediatric MRI. <i>Paediatric Anaesthesia</i> , 2008, 18, 403-411.	0.6	296
2	Mutations in LRP2, which encodes the multiligand receptor megalin, cause Donnai-Barrow and facio-oculo-acoustico-renal syndromes. <i>Nature Genetics</i> , 2007, 39, 957-959.	9.4	284
3	The spectrum of vascular anomalies in patients with PTEN mutations: implications for diagnosis and management. <i>Journal of Medical Genetics</i> , 2007, 44, 594-602.	1.5	233
4	The Presentation and Management of Nasal Dermoid. <i>JAMA Otolaryngology</i> , 2003, 129, 464.	1.5	163
5	Volumetric MRI Study of Brain in Children With Intrauterine Exposure to Cocaine, Alcohol, Tobacco, and Marijuana. <i>Pediatrics</i> , 2008, 121, 741-750.	1.0	140
6	Nasal Glioma and Encephalocele: Diagnosis and Management. <i>Laryngoscope</i> , 2003, 113, 2069-2077.	1.1	117
7	Transnasal Endoscopic Excision of Midline Nasal Dermoid from the Anterior Cranial Base. <i>Plastic and Reconstructive Surgery</i> , 1998, 102, 2119-2123.	0.7	89
8	How Accurately Does Current Fetal Imaging Identify Posterior Fossa Anomalies?. <i>American Journal of Roentgenology</i> , 2008, 190, 1637-1643.	1.0	83
9	Level of In Utero Cocaine Exposure and Neonatal Ultrasound Findings. <i>Pediatrics</i> , 1999, 104, 1101-1105.	1.0	75
10	HOXB1 Founder Mutation in Humans Recapitulates the Phenotype of Hoxb1 Mice. <i>American Journal of Human Genetics</i> , 2012, 91, 171-179.	2.6	72
11	A defect in myoblast fusion underlies Carey-Fineman-Ziter syndrome. <i>Nature Communications</i> , 2017, 8, 16077.	5.8	72
12	NEUROGENIC TUMORS OF THE NECK. <i>Radiologic Clinics of North America</i> , 2000, 38, 1077-1090.	0.9	67
13	Loss of MAFB Function in Humans and Mice Causes Duane Syndrome, Aberrant Extraocular Muscle Innervation, and Inner-Ear Defects. <i>American Journal of Human Genetics</i> , 2016, 98, 1220-1227.	2.6	66
14	Mucoepidermoid Carcinoma of the Parotid Gland in Children. <i>JAMA Otolaryngology</i> , 2006, 132, 375.	1.5	64
15	Lingual Thyroid in Children: A Rare Clinical Entity. <i>Laryngoscope</i> , 2008, 118, 1174-1179.	1.1	64
16	Biallelic mutations in human DCC cause developmental split-brain syndrome. <i>Nature Genetics</i> , 2017, 49, 606-612.	9.4	62
17	Foregut Duplication Cysts in the Head and Neck. <i>JAMA Otolaryngology</i> , 2010, 136, 778.	1.5	55
18	Imaging of head and neck neoplasms in children. <i>Pediatric Radiology</i> , 2010, 40, 499-509.	1.1	51

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19	Craniofacial, Temporal Bone, and Audiologic Abnormalities in the Spectrum of Hemifacial Microsomia. <i>JAMA Otolaryngology</i> , 2001, 127, 265.	1.5	50
20	Primary Thyroid Teratomas in Children. <i>American Journal of Surgical Pathology</i> , 2005, 29, 700-706.	2.1	48
21	Pleomorphic adenoma of the parotid gland in children. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2007, 71, 1717-1723.	0.4	44
22	MR imaging of fetal head and neck anomalies. <i>Neuroimaging Clinics of North America</i> , 2004, 14, 273-291.	0.5	43
23	Congenital hearing impairment. <i>Pediatric Radiology</i> , 2006, 36, 309-324.	1.1	43
24	Pediatric Chronic Nonbacterial Osteomyelitis of the Jaw: Clinical, Radiographic, and Histopathologic Features. <i>Journal of Oral and Maxillofacial Surgery</i> , 2016, 74, 2393-2402.	0.5	41
25	The Biology and Management of Laryngeal Neurofibroma. <i>JAMA Otolaryngology</i> , 2004, 130, 1400.	1.5	40
26	Frequency and Cause of Disagreements in Diagnoses for Fetuses Referred for Ventriculomegaly. <i>Radiology</i> , 2008, 247, 516-527.	3.6	40
27	Encephalocraniocutaneous lipomatosis accompanied by the formation of bone cysts: Harboring clues to pathogenesis?. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 2973-2980.	0.7	38
28	Update on current evaluation and management of pediatric nasal dermoid. <i>Laryngoscope</i> , 2016, 126, 2151-2160.	1.1	37
29	Cerebral Vein Malformations Result from Loss of Twist1 Expression and BMP Signaling from Skull Progenitor Cells and Dura. <i>Developmental Cell</i> , 2017, 42, 445-461.e5.	3.1	37
30	Intracranial anomalies detected by imaging studies in 30 patients with Apert syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 1337-1338.	0.7	32
31	CT versus MR in neonatal brain imaging at term. <i>Pediatric Radiology</i> , 2003, 33, 442-449.	1.1	31
32	A prospective study of 113 deep neck infections managed using a clinical practice guideline. <i>Laryngoscope</i> , 2013, 123, 3211-3218.	1.1	31
33	Congenital nasal neuroglial heterotopia and encephaloceles: An update on current evaluation and management. <i>Laryngoscope</i> , 2016, 126, 2161-2167.	1.1	30
34	Success of Nonsedated Neuroradiologic MRI in Children 1â€“7 Years Old. <i>American Journal of Roentgenology</i> , 2021, 216, 1370-1377.	1.0	30
35	The presentation and management of mandibular tumors in the pediatric population. <i>Laryngoscope</i> , 2013, 123, 2035-2042.	1.1	29
36	Neurologic complications of cerebral angiography in childhood moyamoya syndrome. <i>Pediatric Radiology</i> , 1998, 28, 824-829.	1.1	29

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37	Outcome of Fetuses With Cerebral Ventriculomegaly and Septum Pellucidum Leaflet Abnormalities. American Journal of Roentgenology, 2011, 196, W83-W92.	1.0	28
38	Zygomatocotemporal Synostosis: A Rare Cause of Progressive Facial Asymmetry. Cleft Palate-Craniofacial Journal, 2007, 44, 106-111.	0.5	27
39	Evaluation of Real-Time Single-Shot Fast Spin-Echo MRI for Visualization of the Fetal Midline Corpus Callosum and Secondary Palate. American Journal of Roentgenology, 2006, 187, 1505-1511.	1.0	26
40	Apert syndrome: what prenatal radiographic findings should prompt its consideration?. Prenatal Diagnosis, 2006, 26, 966-972.	1.1	25
41	Imaging Features of Juvenile Xanthogranuloma of the Pediatric Head and Neck. American Journal of Neuroradiology, 2016, 37, 910-916.	1.2	24
42	Using spectral-domain optical coherence tomography to detect optic neuropathy in patients with craniosynostosis. Journal of AAPOS, 2014, 18, 543-549.	0.2	23
43	Optic Atrophy and a Leigh-Like Syndrome Due to Mutations in the C12orf65 Gene. Journal of Neuro-Ophthalmology, 2014, 34, 39-43.	0.4	22
44	Marked Regression of Metastatic Pilocytic Astrocytoma During Treatment With Imatinib Mesylate (STI-571, Gleevec): A Case Report and Laboratory Investigation. Journal of Pediatric Hematology/Oncology, 2003, 25, 644-648.	0.3	21
45	Additional clinical manifestations in children with sensorineural hearing loss and biallelic GJB2 mutations: Who should be offered GJB2 testing?. American Journal of Medical Genetics, Part A, 2007, 143A, 1560-1566.	0.7	21
46	Ultrasound and MRI of Fetuses With Ventriculomegaly: Can Cortical Development Be Used to Predict Postnatal Outcome?. American Journal of Roentgenology, 2011, 196, 1457-1467.	1.0	21
47	Temporal bone abnormalities in children with GJB2 mutations. Laryngoscope, 2011, 121, 630-635.	1.1	18
48	Imaging of Pediatric Orbital Diseases. Neuroimaging Clinics of North America, 2015, 25, 477-501.	0.5	18
49	Pleomorphic adenoma of the head and neck in children: presentation and management. Laryngoscope, 2019, 129, 2603-2609.	1.1	18
50	Prenatal Imaging of Craniosynostosis Syndromes. Seminars in Ultrasound, CT and MRI, 2015, 36, 453-464.	0.7	17
51	Cockayne syndrome: The developing phenotype. American Journal of Medical Genetics, Part A, 2005, 135A, 214-216.	0.7	16
52	Mucoepidermoid carcinoma of the head and neck in children. International Journal of Pediatric Otorhinolaryngology, 2019, 120, 93-99.	0.4	16
53	Vallecular cyst in the pediatric population: Evaluation and management. International Journal of Pediatric Otorhinolaryngology, 2018, 113, 198-203.	0.4	15
54	Safety and efficacy of gamma-secretase inhibitor nirogacestat (PF03084014) in desmoid tumor: Report of four pediatric/young adult cases. Pediatric Blood and Cancer, 2020, 67, e28636.	0.8	15

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55	Acute hydrocephalus secondary to obstruction of the foramen of Monro and cerebral aqueduct caused by a choroid plexus cyst in the lateral ventricle. <i>Journal of Neurosurgery: Pediatrics</i> , 2007, 107, 236-239.	0.8	14
56	Cysts and tumors of the oral cavity, oropharynx, and nasopharynx in children. <i>Neuroimaging Clinics of North America</i> , 2003, 13, 427-442.	0.5	13
57	Imaging Optimization in Children. <i>Journal of the American College of Radiology</i> , 2018, 15, 440-443.	0.9	13
58	Altered White Matter Organization in the TUBB3 E410K Syndrome. <i>Cerebral Cortex</i> , 2019, 29, 3561-3576.	1.6	13
59	Novel variants in TUBA1A cause congenital fibrosis of the extraocular muscles with or without malformations of cortical brain development. <i>European Journal of Human Genetics</i> , 2021, 29, 816-826.	1.4	13
60	TUBB3 Arg262His causes a recognizable syndrome including CFEOM3, facial palsy, joint contractures, and early-onset peripheral neuropathy. <i>Human Genetics</i> , 2021, 140, 1709-1731.	1.8	13
61	The role of chromosomal translocation (15;19) in the carcinoma of the upper aerodigestive tract in children. <i>Otolaryngology - Head and Neck Surgery</i> , 2003, 129, 698-704.	1.1	12
62	Dentigerous cyst associated with a displaced tooth in the maxillary sinus: an unusual cause of recurrent sinusitis in an adolescent. <i>Pediatric Radiology</i> , 2009, 39, 1102-1104.	1.1	12
63	Meningoencephalocele and Other Dural Disruptions. <i>Journal of Craniofacial Surgery</i> , 2011, 22, 182-186.	0.3	12
64	Enlargement of the Internal Auditory Canal and Associated Posterior Fossa Anomalies in PHACES Association. <i>American Journal of Neuroradiology</i> , 2015, 36, 2159-2162.	1.2	12
65	Sinus Computed Tomography Imaging in Pediatric Cystic Fibrosis. <i>Otolaryngology - Head and Neck Surgery</i> , 2016, 155, 160-165.	1.1	12
66	Guidelines for magnetic resonance imaging in pediatric head and neck pathologies: a multicentre international consensus paper. <i>Neuroradiology</i> , 2022, 64, 1081-1100.	1.1	12
67	Recessive MYF5 Mutations Cause External Ophthalmoplegia, Rib, and Vertebral Anomalies. <i>American Journal of Human Genetics</i> , 2018, 103, 115-124.	2.6	11
68	Role of Surgery in Rhabdomyosarcoma of the Head and Neck in Children. <i>Laryngoscope</i> , 2021, 131, E984-E992.	1.1	11
69	Neuroimaging of Children With Surgically Treated Hydrocephalus: A Practical Approach. <i>American Journal of Roentgenology</i> , 2017, 208, 413-419.	1.0	10
70	Diagnostic equivalency of fast T2 and FLAIR sequences for pediatric brain MRI: a pilot study. <i>Pediatric Radiology</i> , 2020, 50, 550-559.	1.1	10
71	Ultrasonographic detection of intracranial melanocytosis in an infant. <i>Pediatric Radiology</i> , 2010, 40, 210-214.	1.1	9
72	Anomalous superior oblique muscles and tendons in congenital fibrosis of the extraocular muscles. <i>Journal of AAPOS</i> , 2019, 23, 325.e1-325.e6.	0.2	9

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73	Squamous cell carcinoma of the head and neck in children. International Journal of Pediatric Otorhinolaryngology, 2019, 117, 131-137.	0.4	9
74	Characteristic Cochlear Hypoplasia in Patients with Walker-Warburg Syndrome: A Radiologic Study of the Inner Ear in Î±-Dystroglycanâ€“Related Muscular Disorders. American Journal of Neuroradiology, 2021, 42, 167-172.	1.2	9
75	Diffusion imaging in neonates. Neuroimaging Clinics of North America, 2002, 12, 55-70.	0.5	8
76	Neuroimaging in Kabuki syndrome and another <scp><i>KMT2D</i></scp>â€“related disorder. American Journal of Medical Genetics, Part A, 2021, 185, 3770-3783.	0.7	7
77	Magnetic resonance features and cranial nerve involvement in pediatric head and neck rhabdomyosarcomas. Neuroradiology, 2021, 63, 1925-1934.	1.1	6
78	Developmental conjunctival cyst of the eyelid in a child. Journal of AAPOS, 2012, 16, 196-198.	0.2	5
79	Magnetic Resonance Imaging Features and Clinical Findings in Pediatric Idiopathic Intracranial Hypertension: A Caseâ€“Control Study. Life, 2021, 11, 487.	1.1	5
80	Re-Examining the Cochlea in Branchio-Oto-Renal Syndrome: Genotype-Phenotype Correlation. American Journal of Neuroradiology, 2022, 43, 309-314.	1.2	5
81	An unresponsive infant in the emergency room. Seminars in Pediatric Neurology, 1999, 6, 225-228.	1.0	4
82	Desmoid tumors of the head and neck in the pediatric population: Has anything changed?. International Journal of Pediatric Otorhinolaryngology, 2021, 140, 110511.	0.4	4
83	Facial Paralysis From Post-Transplant Lymphoproliferative Disorder. Otolaryngology and Neurotology, 2021, Publish Ahead of Print, e605-e608.	0.7	3
84	Basilar Meningitis Associated With Ethmoid and Sphenoid Cephaloceles. Pediatric Neurology, 2005, 33, 57-60.	1.0	2
85	Denervation-related fatty muscle infiltration. Pediatric Radiology, 2009, 39, 1379-1379.	1.1	1
86	Association of the Long QT Syndrome With Goiter and Deafness. American Journal of Cardiology, 2010, 105, 681-686.	0.7	1
87	Radiology Quiz Case 3. JAMA Otolaryngology, 2012, 138, 775.	1.5	1
88	Recommendations for Neuroradiology Training during Radiology Residency by the American Society of Neuroradiology Section Chiefs Leadership Group. American Journal of Neuroradiology, 2021, 42, E7-E9.	1.2	1
89	Radiology Quiz Case 1. JAMA Otolaryngology - Head and Neck Surgery, 2013, 139, 643.	1.2	0