## Adrian K Charles

List of Publications by Year in descending order

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99 papers 3,268 citations

126708 33 h-index 53 g-index

102 all docs

102 docs citations

102 times ranked

5061 citing authors

#	Article	IF	CITATIONS
1	Congenital iRHOM2 deficiency causes ADAM17 dysfunction and environmentally directed immunodysregulatory disease. Nature Immunology, 2022, 23, 75-85.	7.0	3
2	Human AGR2 Deficiency Causes Mucus Barrier Dysfunction and Infantile Inflammatory Bowel Disease. Cellular and Molecular Gastroenterology and Hepatology, 2021, 12, 1809-1830.	2.3	26
3	Death Certification., 2021,, 9-17.		O
4	BCG vaccination–induced emergency granulopoiesis provides rapid protection from neonatal sepsis. Science Translational Medicine, 2020, 12, .	5.8	76
5	Long-term fate of the incised urethral plate in Snodgrass procedure; A real concern does exist. Urology Case Reports, 2020, 32, 101216.	0.1	9
6	Congenital Mid Ureteric Valve Stenosis Revisited: Case Report and Review of the Literature. Frontiers in Pediatrics, 2019, 7, 108.	0.9	2
7	Histological chorioamnionitis and developmental outcomes in very preterm infants. Journal of Perinatology, 2019, 39, 321-330.	0.9	19
8	Differentiation of Islet Progenitors Regulated by Nicotinamide into Transcriptome-Verified $\hat{l}^2$ Cells That Ameliorate Diabetes. Stem Cells, 2017, 35, 1341-1354.	1.4	15
9	Maternal Chorioamnionitis and Postneonatal Respiratory Tract Infection in Ex-Preterm Infants. Journal of Pediatrics, 2017, 184, 62-67.e2.	0.9	11
10	Methods to decrease variability in histological scoring in placentas from a cohort of preterm infants. BMJ Open, 2017, 7, e013877.	0.8	4
11	Preclinical evaluation of drugs to block inflammation-driven preterm birth. Innate Immunity, 2017, 23, 20-33.	1.1	14
12	Phylogeny of Wilms tumor?. EBioMedicine, 2016, 9, 11-12.	2.7	2
13	Evolution of Renal Cysts to Anaplastic Sarcoma of Kidney in a Child With DICER1 Syndrome. Pediatric Blood and Cancer, 2016, 63, 1272-1275.	0.8	32
14	Gynecologic Manifestations of the DICER1 Syndrome. Surgical Pathology Clinics, 2016, 9, 227-241.	0.7	43
15	<i>Haemophilus influenzae</i> : a potent perinatal pathogen disproportionately isolated from Indigenous women and their neonates. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2016, 56, 75-81.	0.4	9
16	Surgically managed perinatal testicular torsion: a single centre experience. Journal of Maternal-Fetal and Neonatal Medicine, 2016, 29, 1265-1268.	0.7	12
17	Risk of cancer among children with birth defects: A novel approach. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 284-291.	1.6	16
18	Ovarian Sex Cord-Stromal Tumors in Patients With Probable or Confirmed Germline DICER1 Mutations. International Journal of Gynecological Pathology, 2015, 34, 266-274.	0.9	39

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19	Regulator of G protein signaling 5 is a determinant of gestational hypertension and preeclampsia. Science Translational Medicine, 2015, 7, 290ra88.	5.8	39
20	NUT protein immunoreactivity in ovarian germ cell tumours. Pathology, 2015, 47, 118-122.	0.3	8
21	Inflammatory myofibroblastic tumour of the bladder in children: A review. Journal of Pediatric Urology, 2015, 11, 239-245.	0.6	20
22	Infection with Toxin A-Negative, Toxin B-Negative, Binary Toxin-Positive Clostridium difficile in a Young Patient with Ulcerative Colitis. Journal of Clinical Microbiology, 2015, 53, 3702-3704.	1.8	36
23	Congenital Tumors. , 2015, , 449-479.		0
24	Comparative drug screening in NUT midline carcinoma. British Journal of Cancer, 2014, 110, 1189-1198.	2.9	44
25	Connective tissue growth factor is expressed in bone marrow stromal cells and promotes interleukin-7-dependent B lymphopoiesis. Haematologica, 2014, 99, 1149-1156.	1.7	18
26	Multinodular Goiter in Children: An Important Pointer to a Germline DICER1 Mutation. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1947-1948.	1.8	30
27	Expression Profile of Wnt/ $\hat{l}^2$ -Catenin Signalling Molecules and the Wnt Antagonist Secreted Frizzled-Related Protein 4 in Apoptosis in Breast Cancer Tissue Micro-Arrays. Journal of Analytical Oncology, 2014, 3, 205-212.	0.1	1
28	Mutations in KLHL40 Are a Frequent Cause of Severe Autosomal-Recessive Nemaline Myopathy. American Journal of Human Genetics, 2013, 93, 6-18.	2.6	186
29	Overexpression of Aromatase Associated With Loss of Heterozygosity of the <i>STK11 &lt; /i&gt;Gene Accounts for Prepubertal Gynecomastia in Boys with Peutz-Jeghers Syndrome. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E1979-E1987.</i>	1.8	29
30	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.	1.1	97
31	Novel BRD4–NUT fusion isoforms increase the pathogenic complexity in NUT midline carcinoma. Oncogene, 2013, 32, 4664-4674.	2.6	41
32	Collaborating with consumers: the key to achieving statutory notification for birth defects and cerebral palsy in Western Australia. Journal of Registry Management, 2013, 40, 9-13.	0.1	1
33	Histologic Chorioamnionitis Is Associated With Reduced Risk of Late-Onset Sepsis in Preterm Infants. Pediatrics, 2012, 129, e134-e141.	1.0	115
34	Responsiveness of human monocytes to the commensal bacterium Staphylococcus epidermidis develops late in gestation. Pediatric Research, 2012, 72, 10-18.	1.1	53
35	Placental measurements associated with intelligence quotient at age 7 years. Journal of Developmental Origins of Health and Disease, 2012, 3, 190-197.	0.7	20
36	SNP-based arrays complement classic cytogenetics in the detection of chromosomal aberrations in Wilms' tumor. Cancer Genetics, 2012, 205, 80-93.	0.2	10

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37	Characterization of 17.94, a novel anaplastic Wilms' tumor cell line. Cancer Genetics, 2012, 205, 319-326.	0.2	16
38	Changes in thymic regulatory T-cell maturation from birth to puberty: Differences in atopic children. Journal of Allergy and Clinical Immunology, 2012, 129, 199-206.e4.	1.5	56
39	Inflammatory and Haematological Markers in the Maternal, Umbilical Cord and Infant Circulation in Histological Chorioamnionitis. PLoS ONE, 2012, 7, e51836.	1.1	48
40	Solitary, multifocal and generalized myofibromas: clinicopathological and immunohistochemical features of 114 cases. Histopathology, 2012, 60, E1-11.	1.6	59
41	Fetal akinesia: review of the genetics of the neuromuscular causes. Journal of Medical Genetics, 2011, 48, 793-801.	1.5	65
42	Extending the phenotypes associated with <i>DICER1 </i> hi>mutations. Human Mutation, 2011, 32, 1381-1384.	1.1	173
43	Fetal oromandibular limb hypogenesis syndrome following uterine curettage in early pregnancy. Birth Defects Research Part A: Clinical and Molecular Teratology, 2011, 91, 226-229.	1.6	9
44	Pathology, genetics and cytogenetics of Wilms' tumour. Pathology, 2011, 43, 302-312.	0.3	29
45	Perilobar Nephrogenic Rests and Chromosome 22. Pediatric and Developmental Pathology, 2011, 14, 485-492.	0.5	5
46	Infection and Fetal Loss in the Mid-Second Trimester of Pregnancy. Obstetrical and Gynecological Survey, 2010, 65, 613-614.	0.2	0
47	Cytogenetic findings in Wilms' tumour: a single institute study. Pathology, 2010, 42, 643-649.	0.3	6
48	Caudal dysgenesis and sirenomeliaâ€single centre experience suggests common pathogenic basis. American Journal of Medical Genetics, Part A, 2010, 152A, 2578-2587.	0.7	46
49	Familial pleuropulmonary blastoma in Australia. Pediatric Blood and Cancer, 2010, 55, 1417-1419.	0.8	15
50	Uptake of the Perinatal Society of Australia and New Zealand perinatal mortality audit guideline. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2010, 50, 138-143.	0.4	20
51	Infection and fetal loss in the midâ€second trimester of pregnancy. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2010, 50, 221-225.	0.4	31
52	Celiac Disease and Eosinophilic Esophagitis: A True Association. Journal of Pediatric Gastroenterology and Nutrition, 2010, 50, 397-399.	0.9	67
53	Gross Placental Structure in a Low-Risk Population of Singleton, Term, First-Born Infants. Pediatric and Developmental Pathology, 2009, 12, 200-210.	0.5	18
54	The genotype of the NK cell receptor, KIR2DL4, influences INFÂ secretion by decidual natural killer cells. Molecular Human Reproduction, 2009, 15, 489-497.	1.3	31

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55	Allometric Metabolic Scaling and Fetal and Placental Weight. Placenta, 2009, 30, 355-360.	0.7	57
56	Non-Linear and Gender-Specific Relationships Among Placental Growth Measures and The Fetoplacental Weight Ratioa~†. Placenta, 2009, 30, 1052-1057.	0.7	80
57	Causes of death and associated conditions (Codac) – a utilitarian approach to the classification of perinatal deaths. BMC Pregnancy and Childbirth, 2009, 9, 22.	0.9	100
58	Unexplained fetal death: Are women with a history of fetal loss at higher risk?. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2009, 49, 151-157.	0.4	9
59	An evaluation of classification systems for stillbirth. BMC Pregnancy and Childbirth, 2009, 9, 24.	0.9	131
60	Cells of Epithelial Lineage Are Present in Blood, Engraft the Bronchial Epithelium, and Are Increased in Human Lung Transplantation. Journal of Heart and Lung Transplantation, 2009, 28, 550-557.	0.3	7
61	Function of caspase-14 in trophoblast differentiation. Reproductive Biology and Endocrinology, 2009, 7, 98.	1.4	8
62	Thymic Indoleamine 2,3-Dioxygenase-Positive Eosinophils in Young Children. American Journal of Pathology, 2009, 175, 2043-2052.	1.9	35
63	RCPA QAP paediatric/perinatal session. Pathology, 2009, 41, 8.	0.3	1
64	Fetal growth and the risk of childhood CNS tumors and lymphomas in Western Australia. International Journal of Cancer, 2008, 123, 436-443.	2.3	24
65	Fetal growth and the risk of childhood non-CNS solid tumours in Western Australia. British Journal of Cancer, 2008, 99, 179-181.	2.9	20
66	Chronic diarrhoea: a presentation of immature neuroblastoma. ANZ Journal of Surgery, 2008, 78, 218-219.	0.3	2
67	Placental characteristics and birthweight. Paediatric and Perinatal Epidemiology, 2008, 22, 229-239.	0.8	111
68	Frequency and Timing of Loss of Imprinting at 11p13 and 11p15 in Wilms' Tumor Development. Molecular Cancer Research, 2008, 6, 1114-1123.	1.5	20
69	Renal tumours of childhood: an update. Pathology, 2008, 40, 217-227.	0.3	12
70	Placental growth patterns affect birth weight for given placental weight. Birth Defects Research Part A: Clinical and Molecular Teratology, 2007, 79, 281-288.	1.6	76
71	Aetiology of stillbirth: unexplored is not unexplained. Australian and New Zealand Journal of Public Health, 2007, 31, 444-449.	0.8	31
72	The role of autopsy following pregnancy termination for fetal abnormality. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2007, 47, 445-449.	0.4	36

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73	Timing of morphologic and apoptotic changes in the sheep fetal kidney in response to bladder outflow obstruction. Journal of Pediatric Urology, 2006, 2, 216-224.	0.6	5
74	Placenta and Fetal Growth Restriction. Clinical Obstetrics and Gynecology, 2006, 49, 236-256.	0.6	188
75	Discordant Fetal Infection for Parvovirus B19 in a Dichorionic Twin Pregnancy. Twin Research and Human Genetics, 2006, 9, 456-459.	0.3	18
76	Pentoxifylline Reduces the Incidence and Severity of Necrotizing Enterocolitis in a Neonatal Rat Model. Pediatric Research, 2006, 60, 185-189.	1.1	57
77	Discordant fetal infection for parvovirus B19 in a dichorionic twin pregnancy. Twin Research and Human Genetics, 2006, 9, 456-9.	0.3	9
78	Advanced abdominal pregnancy: still an occurrence in modern medicine. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2005, 45, 518-521.	0.4	34
79	Interpretation of recent sudden infant death syndrome rates in Western Australia Journal of Paediatrics and Child Health, 2005, 41, 669-670.	0.4	3
80	The effect of terminations of pregnancy for fetal abnormalities on trends in mortality to one year of age in Western Australia. Paediatric and Perinatal Epidemiology, 2005, 19, 284-293.	0.8	25
81	The expression of apoptosis related genes in the first trimester human placenta using a short term in vitro model. Apoptosis: an International Journal on Programmed Cell Death, 2005, 10, 135-140.	2.2	7
82	Diamniotic Conjoined Fetuses in a Triplet Pregnancy: An Insight into Embryonic Topology. Pediatric and Developmental Pathology, 2005, 8, 666-672.	0.5	15
83	Morphometry of the Basal Plate Superficial Uteroplacental Vasculature in Normal Midtrimester and at Term. Pediatric and Developmental Pathology, 2005, 8, 639-646.	0.5	6
84	Caspase-14 expression in the human placenta. Reproductive BioMedicine Online, 2005, 11, 236-243.	1.1	12
85	Functional Deletion of the Calcium-Sensing Receptor in a Case of Neonatal Severe Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 3721-3730.	1.8	41
86	Neonatal congenital microvillus atrophy. Postgraduate Medical Journal, 2004, 80, 80-83.	0.9	11
87	Screening for defective DNA mismatch repair in stage II and III colorectal cancer patients. Clinical Gastroenterology and Hepatology, 2004, 2, 1017-1025.	2.4	41
88	Wilms tumor in a pediatric renal transplant recipient with unexpected Denys–Drash syndrome. Transplantation Proceedings, 2002, 34, 3203-3204.	0.3	2
89	The role of the pathologist in the management of neuroblastoma. Pediatric Surgery International, 2002, 18, 295-298.	0.6	0
90	The role of comparative genomic hybridisation in prenatal diagnosis. British Journal of Obstetrics and Gynaecology, 2001, 108, 642-648.	0.9	3

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91	Central nervous system metastasis in Wilms' tumor. Cancer, 1998, 83, 2023-2029.	2.0	37
92	Antisense WT1 transcription parallels sense mRNA and protein expression in fetal kidney and can elevate protein levelsin vitro., 1998, 185, 352-359.		45
93	Renal tumours of childhood. Histopathology, 1998, 32, 293-309.	1.6	64
94	Microdissecting the Genetic Events in Nephrogenic Rests and Wilms' Tumor Development. American Journal of Pathology, 1998, 153, 991-1000.	1.9	78
95	Focal dermal hypoplasia (Goltz syndrome) presenting as a severe fetal malformation syndrome. Clinical Dysmorphology, 1997, 6, 267-272.	0.1	21
96	Letter to the Editor: Ossifying Renal Tumor of Infancy. Pediatric Pathology & Laboratory Medicine: Journal of the Society for Pediatric Pathology, Affiliated With the International Paediatric Pathology Association, 1997, 17, 332-334.	0.3	5
97	Nephrogenic Rest and Mesoblastic Nephroma. Pediatric Pathology & Laboratory Medicine: Journal of the Society for Pediatric Pathology, Affiliated With the International Paediatric Pathology Association, 1996, 16, 695-696.	0.3	1
98	Early morphological evidence of autosomal recessive polycystic kidney disease. Lancet, The, 1995, 345, 987.	6.3	3
99	Stillbirth and intrauterine growth restriction. , 0, , 62-82.		3