Anwar Baban

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6906809/publications.pdf

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82	1,967	279798	289244
papers	citations	23 h-index	40 g-index

85	85	85	3457
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The role of the electrocardiographic phenotype in risk stratification for sudden cardiac death in childhood hypertrophic cardiomyopathy. European Journal of Preventive Cardiology, 2022, 29, 645-653.	1.8	20
2	Genetics in Congenital Heart Diseases. Heart Failure Clinics, 2022, 18, 139-153.	2.1	8
3	ICD Outcome in Pediatric Cardiomyopathies. Journal of Cardiovascular Development and Disease, 2022, 9, 33.	1.6	3
4	Cardiomyopathies in Children and Systemic Disorders When Is It Useful to Look beyond the Heart?. Journal of Cardiovascular Development and Disease, 2022, 9, 47.	1.6	5
5	Arrhythmogenic cardiomyopathy in children according to â€œPadua criteriaâ€: Single pediatric center experience. International Journal of Cardiology, 2022, 350, 83-89.	1.7	12
6	Syndromic and Non-Syndromic Patients with Repaired Tetralogy of Fallot: Does It Affect the Long-Term Outcome?. Journal of Clinical Medicine, 2022, 11, 850.	2.4	3
7	8p23.1 deletion : Look out for left ventricular hypertrabeculation and not only congenital heart diseases . Single â€•center experience and literature revision. American Journal of Medical Genetics, Part A, 2022, 188, 883-895.	1.2	4
8	Congenital heart defects in molecularly confirmed <scp>KBG</scp> syndrome patients. American Journal of Medical Genetics, Part A, 2022, 188, 1149-1159.	1.2	5
9	Relationship Between Maximal Left Ventricular Wall Thickness and Sudden Cardiac Death in Childhood Onset Hypertrophic Cardiomyopathy. Circulation: Arrhythmia and Electrophysiology, 2022, 15, CIRCEP121010075.	4.8	8
10	Clinical Features and Natural History of Preadolescent Nonsyndromic HypertrophicÂCardiomyopathy. Journal of the American College of Cardiology, 2022, 79, 1986-1997.	2.8	20
11	Discovering a new part of the phenotypic spectrum of Coffin-Siris syndrome in a fetal cohort. Genetics in Medicine, 2022, 24, 1753-1760.	2.4	6
12	Persistent myocardial atrophy despite LV reverse remodeling in Duchenne cardiomyopathy treated by LVAD. Pediatric Transplantation, 2021, 25, e13890.	1.0	0
13	Deciphering Genetic Variants of Warfarin Metabolism in Children With Ventricular Assist Devices. Pediatric Cardiology, 2021, 42, 1082-1087.	1.3	2
14	Troponin T Mutation as a Cause of Left Ventricular Systolic Dysfunction in a Young Patient with Previous Surgical Correction of Aortic Coarctation. Biomolecules, 2021, 11, 696.	4.0	4
15	1â€…The role of the electrocardiographic phenotype in risk stratification for sudden cardiac death in childhood hypertrophic cardiomyopathy. , 2021, , .		2
16	Cardiac Defects and Genetic Syndromes: Old Uncertainties and New Insights. Genes, 2021, 12, 1047.	2.4	12
17	Clinical presentation and longâ€term outcomes of infantile hypertrophic cardiomyopathy: a European multicentre study. ESC Heart Failure, 2021, 8, 5057-5067.	3.1	22
18	Medical conditions of children and young people with Down syndrome. Journal of Intellectual Disability Research, 2021, 65, 199-209.	2.0	11

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19	Clinical Manifestations of 22q11.2 Deletion Syndrome. Heart Failure Clinics, 2021, 18, 155-164.	2.1	15
20	Myocardial and Arrhythmic Spectrum of Neuromuscular Disorders in Children. Biomolecules, 2021, 11, 1578.	4.0	5
21	Delayed appearance of 3â€methylglutaconic aciduria in neonates with early onset metabolic cardiomyopathies: A potential pitfall for the diagnosis. American Journal of Medical Genetics, Part A, 2020, 182, 64-70.	1.2	4
22	Cardiovascular Involvement in Pediatric Laminopathies. Report of Six Patients and Literature Revision. Frontiers in Pediatrics, 2020, 8, 374.	1.9	9
23	Clinical Profile of Cardiac Involvement in Danon Disease. Circulation Genomic and Precision Medicine, 2020, 13, e003117.	3.6	29
24	Atypical cardiac defects in patients with RASopathies: Updated data on CARNET study. Birth Defects Research, 2020, 112, 725-731.	1.5	17
25	CantÃº syndrome versus Zimmermann-Laband syndrome: Report of nine individuals with ABCC9 variants. European Journal of Medical Genetics, 2020, 63, 103996.	1.3	7
26	KBG syndrome: Common and uncommon clinical features based on 31 new patients. American Journal of Medical Genetics, Part A, 2020, 182, 1073-1083.	1.2	27
27	An International Multicenter Evaluation of Type 5 Long QT Syndrome. Circulation, 2020, 141, 429-439.	1.6	39
28	Differences in morbidity and mortality in Down syndrome are related to the type of congenital heart defect. American Journal of Medical Genetics, Part A, 2020, 182, 1342-1350.	1.2	19
29	Development of a Novel Risk Prediction Model for Sudden Cardiac Death in Childhood Hypertrophic Cardiomyopathy (HCM Risk-Kids). JAMA Cardiology, 2019, 4, 918.	6.1	147
30	SOS1 mutations in Noonan syndrome: Cardiomyopathies and not only congenital heart defects! Report of six patients including two novel variants and literature review. American Journal of Medical Genetics, Part A, 2019, 179, 2083-2090.	1.2	10
31	Palmoplantar Keratoderma and Woolly Hair Revealing Asymptomatic Arrhythmogenic Cardiomyopathy. Acta Dermato-Venereologica, 2019, 99, 831-832.	1.3	4
32	Hidden Complexity in Routine Adult and Pediatric Arrhythmias Interpretation. Cardiac Electrophysiology Clinics, 2019, 11, 391-404.	1.7	0
33	Targeted panel sequencing in pediatric primary cardiomyopathy supports a critical role of TNNI3. Clinical Genetics, 2019, 96, 549-559.	2.0	28
34	Heart rate reduction strategy using ivabradine in end-stage Duchenne cardiomyopathy. International Journal of Cardiology, 2019, 280, 99-103.	1.7	17
35	A heterozygous, intragenic deletion of CNOT2 recapitulates the phenotype of 12q15 deletion syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 1615-1621.	1.2	10
36	Left pulmonary artery in 22q11.2 deletion syndrome. Echocardiographic evaluation in patients without cardiac defects and role of Tbx1 in mice. PLoS ONE, 2019, 14, e0211170.	2.5	13

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37	LTBP2â€related â€œMarfanâ€likeâ€phenotype in two Roma/Gypsy subjects with the LTBP2 homozygous p.R299X variant. American Journal of Medical Genetics, Part A, 2019, 179, 104-112.	1.2	10
38	The ARID1B spectrum in 143 patients: from nonsyndromic intellectual disability to Coffinâ€“Siris syndrome. Genetics in Medicine, 2019, 21, 1295-1307.	2.4	80
39	Atrioventricular canal defect and genetic syndromes: The unifying role of sonic hedgehog. Clinical Genetics, 2019, 95, 268-276.	2.0	20
40	Data on cardiac defects, morbidity and mortality in patients affected by RASopathies. CARNET study results. Data in Brief, 2018, 16, 649-654.	1.0	6
41	Clinical Presentation and Natural History of Hypertrophic Cardiomyopathy in RASopathies. Heart Failure Clinics, 2018, 14, 225-235.	2.1	44
42	First evidence of maternally inherited mosaicism in TGFBR1 and subtle primary myocardial changes in Loeys-Dietz syndrome: a case report. BMC Medical Genetics, 2018, 19, 170.	2.1	4
43	Long-term survival and phenotypic spectrum in heterotaxy syndrome: A 25-year follow-up experience. International Journal of Cardiology, 2018, 268, 100-105.	1.7	24
44	Pharmacological resources, diagnostic approach and coordination of care in joint hypermobility-related disorders. Expert Review of Clinical Pharmacology, 2018, 11, 689-703.	3.1	6
45	Some Isolated Cardiac Malformations Can Be Related to Laterality Defects. Journal of Cardiovascular Development and Disease, 2018, 5, 24.	1.6	15
46	Bradyarrhythmias in Repaired Atrioventricular Septal Defects: Single-Center Experience Based on 34ÂYears of Follow-Up of 522 Patients. Pediatric Cardiology, 2018, 39, 1590-1597.	1.3	12
47	Heterotaxy syndrome with and without spleen: Different infection risk and management. Journal of Allergy and Clinical Immunology, 2017, 139, 1981-1984.e1.	2.9	14
48	Congenital heart defects in molecularly proven Kabuki syndrome patients. American Journal of Medical Genetics, Part A, 2017, 173, 2912-2922.	1.2	60
49	Congenital heart disease and genetic syndromes: new insights into molecular mechanisms. Expert Review of Molecular Diagnostics, 2017, 17, 861-870.	3.1	39
50	Cardiac defects, morbidity and mortality in patients affected by RASopathies. CARNET study results. International Journal of Cardiology, 2017, 245, 92-98.	1.7	75
51	Human Bâ€cell memory is shaped by ageâ€and tissueâ€specific Tâ€independent and GCâ€dependent events. European Journal of Immunology, 2017, 47, 327-344.	2.9	62
52	Assessment of copy number variations in 120 patients with Poland syndrome. BMC Medical Genetics, 2016, 17, 89.	2.1	20
53	Congenital heart defects in Noonan syndrome and RIT1 mutation. Genetics in Medicine, 2016, 18, 1320.	2.4	16
54	Gershoniâ€Baruch syndrome: First report of a surviving child. American Journal of Medical Genetics, Part A, 2016, 170, 707-711.	1.2	1

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55	Coronary artery ectasia in Noonan syndrome: Report of an individual with SOS1 mutation and literature review. American Journal of Medical Genetics, Part A, 2016, 170, 665-669.	1.2	13
56	Cardiovascular malformations in Adamsâ€“Oliver syndrome. American Journal of Medical Genetics, Part A, 2015, 167, 1175-1177.	1.2	11
57	Hypoplastic left heart syndrome and 21q22.3 deletion. American Journal of Medical Genetics, Part A, 2015, 167, 579-586.	1.2	8
58	Diagnosis of Noonan syndrome and related disorders using target next generation sequencing. BMC Medical Genetics, 2014, 15, 14.	2.1	59
59	Holtâ€“Oram syndrome with intermediate atrioventricular canal defect, and aortic coarctation: Functional characterization of a de novo TBX5 mutation. American Journal of Medical Genetics, Part A, 2014, 164, 1419-1424.	1.2	21
60	Identification of TBX5 mutations in a series of 94 patients with Tetralogy of Fallot. American Journal of Medical Genetics, Part A, 2014, 164, 3100-3107.	1.2	47
61	Exome sequencing identifies a novel mutation in PIK3R1 as the cause of SHORT syndrome. BMC Medical Genetics, 2014, 15, 51.	2.1	34
62	Coffin-Siris Syndrome and the BAF Complex: Genotype-Phenotype Study in 63 Patients. Human Mutation, 2013, 34, 1519-1528.	2.5	178
63	Congenital heart defects in recurrent reciprocal 1q21.1 deletion and duplication syndromes: Rare association with pulmonary valve stenosis. European Journal of Medical Genetics, 2013, 56, 144-149.	1.3	33
64	Atrioventricular canal defect in patients with RASopathies. European Journal of Human Genetics, 2013, 21, 200-204.	2.8	22
65	Syndromic nonâ€compaction of the left ventricle: associated chromosomal anomalies. Clinical Genetics, 2013, 84, 362-367.	2.0	30
66	RDDR: a dysmorphology diagnostic network for newborns in central Italy. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 113-115.	1.5	1
67	Hand and Upper Limb Anomalies in Poland Syndrome. Journal of Pediatric Orthopaedics, 2012, 32, 722-726.	1.2	29
68	Familial Poland anomaly revisited. American Journal of Medical Genetics, Part A, 2012, 158A, 140-149.	1.2	29
69	Atrioventricular Canal Defect and Associated Genetic Disorders: New Insights into Polydactyly Syndromes. Neurology International, 2011, 1, e7.	0.5	6
70	Ebstein anomaly: Genetic heterogeneity and association with microdeletions 1p36 and 8p23.1. American Journal of Medical Genetics, Part A, 2011, 155, 2196-2202.	1.2	38
71	Hypoplastic Left Heart Syndrome in Patients With Kabuki Syndrome. Pediatric Cardiology, 2010, 31, 1111-1113.	1.3	11
72	Dextrocardia in patients with Poland syndrome: Phenotypic characterization provides insight into the pathogenesis. Journal of Thoracic and Cardiovascular Surgery, 2010, 139, 1177-1182.	0.8	43

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73	A t(7;12) balanced translocation with breakpoints overlapping those of the Williamsâ€“Beuren and 12q14 microdeletion syndromes. American Journal of Medical Genetics, Part A, 2010, 152A, 1285-1294.	1.2	1
74	RASopathies: Clinical Diagnosis in the First Year of Life. Molecular Syndromology, 2010, 1, 282-289.	0.8	73
75	A spectrum of LMX1B mutations in Nail-Patella syndrome: New point mutations, deletion, and evidence of mosaicism in unaffected parents. Genetics in Medicine, 2010, 12, 431-439.	2.4	26
76	Poland syndrome with bilateral features: Case description with review of the literature. American Journal of Medical Genetics, Part A, 2009, 149A, 1597-1602.	1.2	62
77	Response to Klinger and Merlob re: Case description with review of the literature. Am J Med Genet Part A 149A:1597â€“1602, 2009. American Journal of Medical Genetics, Part A, 2009, 149A, 2899-2899.	1.2	2
78	Recurrent microdeletion at 17q12 as a cause of Mayer-Rokitansky-Kuster-Hauser (MRKH) syndrome: two case reports. Orphanet Journal of Rare Diseases, 2009, 4, 25.	2.7	105
79	Pituitary hypoplasia and growth hormone deficiency in Coffinâ€Siris syndrome. American Journal of Medical Genetics, Part A, 2008, 146A, 384-388.	1.2	15
80	Further case of metaphyseal acroscyphodysplasia with cone-shaped epiphyses (Bellini disease or) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 4	0.3	1
81	Unilateral radio-ulnar synostosis associated with hypotonia, developmental delay, and facial dysmorphism. American Journal of Medical Genetics, Part A, 2005, 137A, 106-108.	1.2	1
82	How wide is the ocular spectrum of Delleman syndrome?. Clinical Dysmorphology, 2004, 13, 33-34.	0.3	8