Sara Fitzgerald-Butt

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8289897/publications.pdf

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21 papers	516 citations	759233 12 h-index	752698 20 g-index
22 all docs	22 docs citations	22 times ranked	1078 citing authors

#	Article	IF	Citations
1	Genetic counseling for congenital heart disease â€“ Practice resource of the National Society of Genetic Counselors. Journal of Genetic Counseling, 2022, 31, 9-33.	1.6	9
2	Assessing genetic counselorsâ€™ graduate school education and training in congenital heart defects. Journal of Genetic Counseling, 2022, 31, 735-745.	1.6	2
3	Exome sequencing in multiplex families with left-sided cardiac defects has high yield for disease gene discovery. PLoS Genetics, 2022, 18, e1010236.	3.5	8
4	Management of amended variant classification laboratory reports by genetic counselors in the United States and Canada: An exploratory study. Journal of Genetic Counseling, 2021, , .	1.6	3
5	Novel frameshift variant in MYL2 reveals molecular differences between dominant and recessive forms of hypertrophic cardiomyopathy. PLoS Genetics, 2020, 16, e1008639.	3.5	16
6	Reclassification of Variants of Uncertain Significance in Children with Inherited Arrhythmia Syndromes is Predicted by Clinical Factors. Pediatric Cardiology, 2019, 40, 1679-1687.	1.3	24
7	The Responsibility to Recontact Research Participants after Reinterpretation of Genetic and Genomic Research Results. American Journal of Human Genetics, 2019, 104, 578-595.	6.2	91
8	At the Heart of the Pregnancy: What Prenatal and Cardiovascular Genetic Counselors Need to Know about Maternal Heart Disease. Journal of Genetic Counseling, 2017, 26, 669-688.	1.6	5
9	Clinical exome sequencing reports: current informatics practice and future opportunities. Journal of the American Medical Informatics Association: JAMIA, 2017, 24, 1184-1191.	4.4	12
10	A cohort study of multiple families with FBN1 p.R650C variant, ectopia lentis, and low but not absent risk for aortopathy. American Journal of Medical Genetics, Part A, 2017, 173, 2995-3002.	1.2	4
11	Abnormal Longitudinal Growth of the Aorta in Children with Familial Bicuspid Aortic Valve. Pediatric Cardiology, 2017, 38, 1709-1715.	1.3	4
12	Assessment of large copy number variants in patients with apparently isolated congenital leftâ€sided cardiac lesions reveals clinically relevant genomic events. American Journal of Medical Genetics, Part A, 2017, 173, 2176-2188.	1.2	17
13	Measuring genetic knowledge: a brief survey instrument for adolescents and adults. Clinical Genetics, 2016, 89, 235-243.	2.0	33
14	Utilization of Whole Exome Sequencing to Identify Causative Mutations in Familial Congenital Heart Disease. Circulation: Cardiovascular Genetics, 2016, 9, 320-329.	5.1	71
15	Lifetime Prevalence of Sexual Intercourse and Contraception Use at Last Sex Among Adolescents and Young Adults With Congenital Heart Disease. Journal of Adolescent Health, 2015, 56, 396-401.	2.5	22
16	Novel familial dilated cardiomyopathy mutation in <scp>MYL</scp>2 affects the structure and function of myosin regulatory light chain. FEBS Journal, 2015, 282, 2379-2393.	4.7	42
17	Rare GATA5 sequence variants identified in individuals with bicuspid aortic valve. Pediatric Research, 2014, 76, 211-216.	2.3	74
18	Understanding of informed consent by parents of children enrolled in a genetic biobank. Genetics in Medicine, 2014, 16, 141-148.	2.4	43

#	Article	IF	Citations
19	Genetic knowledge and attitudes of parents of children with congenital heart defects. American Journal of Medical Genetics, Part A, 2014, 164, 3069-3075.	1.2	18
20	Parental Knowledge and Attitudes Toward Hypertrophic Cardiomyopathy Genetic Testing. Pediatric Cardiology, 2010, 31, 195-202.	1.3	16
21	Comparison of willingness and preference for genetic counseling via telemedicine: before vs. during the COVID-19 pandemic. Journal of Community Genetics, 0, , .	1.2	2