

Sara Fitzgerald-Butt

List of Publications by Year in descending order

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

Version: 2024-02-01

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	The Responsibility to Recontact Research Participants after Reinterpretation of Genetic and Genomic Research Results. <i>American Journal of Human Genetics</i> , 2019, 104, 578-595.	6.2	91
2	Rare GATA5 sequence variants identified in individuals with bicuspid aortic valve. <i>Pediatric Research</i> , 2014, 76, 211-216.	2.3	74
3	Utilization of Whole Exome Sequencing to Identify Causative Mutations in Familial Congenital Heart Disease. <i>Circulation: Cardiovascular Genetics</i> , 2016, 9, 320-329.	5.1	71
4	Understanding of informed consent by parents of children enrolled in a genetic biobank. <i>Genetics in Medicine</i> , 2014, 16, 141-148.	2.4	43
5	Novel familial dilated cardiomyopathy mutation in <i>MYL2</i> affects the structure and function of myosin regulatory light chain. <i>FEBS Journal</i> , 2015, 282, 2379-2393.	4.7	42
6	Measuring genetic knowledge: a brief survey instrument for adolescents and adults. <i>Clinical Genetics</i> , 2016, 89, 235-243.	2.0	33
7	Reclassification of Variants of Uncertain Significance in Children with Inherited Arrhythmia Syndromes is Predicted by Clinical Factors. <i>Pediatric Cardiology</i> , 2019, 40, 1679-1687.	1.3	24
8	Lifetime Prevalence of Sexual Intercourse and Contraception Use at Last Sex Among Adolescents and Young Adults With Congenital Heart Disease. <i>Journal of Adolescent Health</i> , 2015, 56, 396-401.	2.5	22
9	Genetic knowledge and attitudes of parents of children with congenital heart defects. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 3069-3075.	1.2	18
10	Assessment of large copy number variants in patients with apparently isolated congenital left-sided cardiac lesions reveals clinically relevant genomic events. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 2176-2188.	1.2	17
11	Parental Knowledge and Attitudes Toward Hypertrophic Cardiomyopathy Genetic Testing. <i>Pediatric Cardiology</i> , 2010, 31, 195-202.	1.3	16
12	Novel frameshift variant in MYL2 reveals molecular differences between dominant and recessive forms of hypertrophic cardiomyopathy. <i>PLoS Genetics</i> , 2020, 16, e1008639.	3.5	16
13	Clinical exome sequencing reports: current informatics practice and future opportunities. <i>Journal of the American Medical Informatics Association: JAMIA</i> , 2017, 24, 1184-1191.	4.4	12
14	Genetic counseling for congenital heart disease – Practice resource of the National Society of Genetic Counselors. <i>Journal of Genetic Counseling</i> , 2022, 31, 9-33.	1.6	9
15	Exome sequencing in multiplex families with left-sided cardiac defects has high yield for disease gene discovery. <i>PLoS Genetics</i> , 2022, 18, e1010236.	3.5	8
16	At the Heart of the Pregnancy: What Prenatal and Cardiovascular Genetic Counselors Need to Know about Maternal Heart Disease. <i>Journal of Genetic Counseling</i> , 2017, 26, 669-688.	1.6	5
17	A cohort study of multiple families with <i>FBN1</i> p.R650C variant, ectopia lentis, and low but not absent risk for aortopathy. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 2995-3002.	1.2	4
18	Abnormal Longitudinal Growth of the Aorta in Children with Familial Bicuspid Aortic Valve. <i>Pediatric Cardiology</i> , 2017, 38, 1709-1715.	1.3	4

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19	Management of amended variant classification laboratory reports by genetic counselors in the United States and Canada: An exploratory study. <i>Journal of Genetic Counseling</i> , 2021, , .	1.6	3
20	Assessing genetic counselors'™ graduate school education and training in congenital heart defects. <i>Journal of Genetic Counseling</i> , 2022, 31, 735-745.	1.6	2
21	Comparison of willingness and preference for genetic counseling via telemedicine: before vs. during the COVID-19 pandemic. <i>Journal of Community Genetics</i> , 0, , .	1.2	2