## Frances Bu'lock

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11089152/publications.pdf

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#	Article	IF	CITATIONS
1	Contribution of Global Rare Copy-Number Variants to the Risk of Sporadic Congenital Heart Disease. American Journal of Human Genetics, 2012, 91, 489-501.	6.2	272
2	Development Of Coâ€Ordination Of Sucking, Swallowing And Breathing: Ultrasound Study Of Term And Preterm Infants. Developmental Medicine and Child Neurology, 1990, 32, 669-678.	2.1	267
3	α-Cardiac myosin heavy chain (MYH6) mutations affecting myofibril formation are associated with congenital heart defects. Human Molecular Genetics, 2010, 19, 4007-4016.	2.9	131
4	Alpha-cardiac actin mutations produce atrial septal defects. Human Molecular Genetics, 2008, 17, 256-265.	2.9	128
5	Phenotype-specific effect of chromosome 1q21.1 rearrangements and GJA5 duplications in 2436 congenital heart disease patients and 6760 controls. Human Molecular Genetics, 2012, 21, 1513-1520.	2.9	101
6	Tropomyosin 1: Multiple roles in the developing heart and in the formation of congenital heart defects. Journal of Molecular and Cellular Cardiology, 2017, 106, 1-13.	1.9	40