Tiffany Busa

List of Publications by Year
in descending order

10 papers	202 citations	1477746 6 h-index	1281420 11 g-index
12 all docs	12 docs citations	12 times ranked	389 citing authors

#	Article	IF	Citations
1	Phenotype and genotype analysis of a French cohort of 119 patients with CHARGE syndrome. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2017, 175, 417-430.	0.7	65
2	The phenotypic spectrum of WWOX-related disorders: 20 additional cases of WOREE syndrome and review of the literature. Genetics in Medicine, 2019, 21, 1308-1318.	1.1	48
3	Prenatal findings in children with early postnatal diagnosis of CHARGE syndrome. Prenatal Diagnosis, 2016, 36, 561-567.	1.1	28
4	Prenatal diagnosis of micrognathia in 41 fetuses: Retrospective analysis of outcome and genetic etiologies. American Journal of Medical Genetics, Part A, 2019, 179, 2365-2373.	0.7	21
5	Novel PTEN germline mutation in a family with mild phenotype: Difficulties in genetic counseling. Gene, 2013, 512, 194-197.	1.0	9
6	FOXC1 haploinsufficiency due to 6p25 deletion in a patient with rapidly progressing aortic valve disease. American Journal of Medical Genetics, Part A, 2017, 173, 2489-2493.	0.7	7
7	Growth charts in Kabuki syndrome 1. American Journal of Medical Genetics, Part A, 2020, 182, 446-453.	0.7	7
8	Confirmation that RIPK4 mutations cause not only Bartsocasâ€Papas syndrome but also CHAND syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 3114-3117.	0.7	6
9	Esophageal atresia with tracheoesophageal fistula in a patient with 7q35â€“36.3 deletion including SHH gene. European Journal of Medical Genetics, 2016, 59, 546-548.	0.7	4
10	WNT10B variants in split hand/foot malformation: Report of three novel families and review of the literature. American Journal of Medical Genetics, Part A, 2019, 179, 1351-1356.	0.7	4