Tiffany Busa

List of Publications by Citations

10 papers	121 citations	5 h-index	11 g-index
12 ext. papers	165 ext. citations	3.2 avg, IF	1.65 L-index

#	Paper	IF	Citations
10	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	3.1	41
9	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	8.1	22
8	Prenatal findings in children with early postnatal diagnosis of CHARGE syndrome. *Prenatal Diagnosis*, **2016**, 36, 561-7	3.2	18
7	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	2.5	13
6	Novel PTEN germline mutation in a family with mild phenotype: difficulties in genetic counseling. *Gene*, **2013**, 512, 194-7	3.8	8
5	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	2.5	5
4	Confirmation that RIPK4 mutations cause not only Bartsocas-Papas syndrome but also CHAND syndrome. *American Journal of Medical Genetics, Part A*, **2017**, 173, 3114-3117	2.5	4
3	Growth charts in Kabuki syndrome 1. *American Journal of Medical Genetics, Part A*, **2020**, 182, 446-453	2.5	4
2	Esophageal atresia with tracheoesophageal fistula in a patient with 7q35-36.3 deletion including SHH gene. *European Journal of Medical Genetics*, **2016**, 59, 546-8	2.6	3
1	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	2.5	2