

# Tiffany Busa

## List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , <b>2017</b> , 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. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 1308-1318	8.1	22
8	Prenatal findings in children with early postnatal diagnosis of CHARGE syndrome. <i>Prenatal Diagnosis</i> , <b>2016</b> , 36, 561-7	3.2	18
7	Prenatal diagnosis of micrognathia in 41 fetuses: Retrospective analysis of outcome and genetic etiologies. <i>American Journal of Medical Genetics, Part A</i> , <b>2019</b> , 179, 2365-2373	2.5	13
6	Novel PTEN germline mutation in a family with mild phenotype: difficulties in genetic counseling. <i>Gene</i> , <b>2013</b> , 512, 194-7	3.8	8
5	FOXC1 haploinsufficiency due to 6p25 deletion in a patient with rapidly progressing aortic valve disease. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 2489-2493	2.5	5
4	Confirmation that RIPK4 mutations cause not only Bartsocas-Papas syndrome but also CHAND syndrome. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 3114-3117	2.5	4
3	Growth charts in Kabuki syndrome 1. <i>American Journal of Medical Genetics, Part A</i> , <b>2020</b> , 182, 446-453	2.5	4
2	Esophageal atresia with tracheoesophageal fistula in a patient with 7q35-36.3 deletion including SHH gene. <i>European Journal of Medical Genetics</i> , <b>2016</b> , 59, 546-8	2.6	3
1	WNT10B variants in split hand/foot malformation: Report of three novel families and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , <b>2019</b> , 179, 1351-1356	2.5	2