

# Tiffany Busa

## List of Publications by Year in Descending Order

**Source:** <https://exaly.com/author-pdf/9314125/tiffany-busa-publications-by-year.pdf>

**Version:** 2024-04-26

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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	Growth charts in Kabuki syndrome 1. <i>American Journal of Medical Genetics, Part A</i> , <b>2020</b> , 182, 446-453	2.5	4
9	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
8	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
7	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
6	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
5	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
4	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
3	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
2	Prenatal findings in children with early postnatal diagnosis of CHARGE syndrome. <i>Prenatal Diagnosis</i> , <b>2016</b> , 36, 561-7	3.2	18
1	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