

# Cinthya Zepeda-Mendoza

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2602448/publications.pdf>

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#	ARTICLE	IF	CITATIONS
1	The Iceberg under Water: Unexplored Complexity of Chromoanagenesis in Congenital Disorders. American Journal of Human Genetics, 2019, 104, 565-577.	6.2	46
2	Variable expressivity of syndromic BMP4-related eye, brain, and digital anomalies: A review of the literature and description of three new cases. European Journal of Human Genetics, 2019, 27, 1379-1388.	2.8	8
3	Concomitant 1p/19q co-deletion and IDH1/2, ATRX, and TP53 mutations within a single clone of Æœdual-genotypeÆœ-IDH-mutant infiltrating gliomas. Acta Neuropathologica, 2020, 139, 1105-1107.	7.7	8
4	Myeloid malignancies with 5q and 7q deletions are associated with extreme genomic complexity, biallelic TP53 variants, and very poor prognosis. Blood Cancer Journal, 2021, 11, 18.	6.2	8
5	An intragenic duplication of TRPS1 leading to abnormal transcripts and causing trichorhinophalangeal syndrome type I. Journal of Physical Education and Sports Management, 2019, 5, a004655.	1.2	5
6	Familial segregation of a 5q15Æœq21.2 deletion associated with facial dysmorphism and speech delay. Clinical Case Reports (discontinued), 2019, 7, 1154-1160.	0.5	4
7	Computational Prediction of Position Effects of Human Chromosome Rearrangements. Current Protocols in Human Genetics, 2018, 97, e57.	3.5	2
8	Prenatal characterization of a novel inverted <i>SMAD2</i> duplication by mate pair sequencing in a fetus with dextrocardia. Clinical Case Reports (discontinued), 2021, 9, 769-774.	0.5	0