Citation Report
List of articles citing

Prenatal detection of trisomy 9 mosaicism

DOI: 10.1002/pd.1970090803
Prenatal Diagnosis, 1989, 9, 549-54.

#	Paper	IF	Citations
27	Chromosomal Mosaicism in Chorionic Villus Sampling. *Clinics in Perinatology*, **1990**, 17, 867-888	2.8	26
26	An infant with trisomy 9 mosaicism presenting as a complete trisomy 9 by amniocentesis. *Prenatal Diagnosis*, **1992**, 12, 31-7	3.2	23
25	Prenatal sonography in trisomy 9. *Prenatal Diagnosis*, **1992**, 12, 175-81	3.2	29
24	Proposed guidelines for diagnosis of chromosome mosaicism in amniocytes based on data derived from chromosome mosaicism and pseudomosaicism studies. *Prenatal Diagnosis*, **1992**, 12, 555-73	3.2	119
23	Chromosomal abnormalities associated with a single umbilical artery. *Prenatal Diagnosis*, **1992**, 12, 965-8	3.2	32
22	Prenatal diagnosis of trisomy 9 mosaic presenting as a case of Dandy-Walker malformation. *Prenatal Diagnosis*, **1993**, 13, 79-85	3.2	17
21	Prenatal diagnosis of trisomy 9 mosaicism: two new cases. *Prenatal Diagnosis*, **1993**, 13, 1001-7	3.2	17
20	Characteristics of structural heart defects in trisomy 9 and their relationship to those in trisomy 13, 18, and 21. *American Heart Journal*, **1993**, 125, 1681-90	4.9	6
19	Trisomy 9: review and report of two new cases. *American Journal of Medical Genetics Part A*, **1995**, 56, 252-7		56
18	Trisomy 9 syndrome: report of a case with Crohn disease and review of the literature. *American Journal of Medical Genetics Part A*, **1995**, 56, 258-64		35
17	Prenatal diagnosis of trisomy 9. Six cases and a review of the literature. *Prenatal Diagnosis*, **1995**, 15, 609-14	3.2	29
16	Rare trisomy mosaicism diagnosed in amniocytes, involving an autosome other than chromosomes 13, 18, 20, and 21: karyotype/phenotype correlations. *Prenatal Diagnosis*, **1997**, 17, 201-42	3.2	153
15	Prenatal diagnosis of trisomy 9: cytogenetic, FISH, and DNA studies. *Prenatal Diagnosis*, **1997**, 17, 933-940	3.2	22
14	Prenatal confirmation of true fetal trisomy 22 mosaicism by fetal skin biopsy following normal fetal blood sampling. **1998**, 18, 384-389		25
13	Antenatal detection of mosaic trisomy 9 by ultrasound: a case report and literature review. *Journal of Maternal-Fetal and Neonatal Medicine*, **2003**, 14, 65-9	2	8
12	Trisomy 9 screened positive for trisomy 18 by maternal serum screening. *Prenatal Diagnosis*, **2004**, 24, 836-8	3.2	5
11	Discrepancies in cytogenetic results between amniocytes and postnatally obtained blood: trisomy 9 mosaicism. *Congenital Anomalies (discontinued)*, **2006**, 46, 115-7	1.1	8

10	Diagnóstico prenatal de trisomía 9 en mosaico en el segundo trimestre de la gestación. *Progresos En Obstetricia Y Ginecologia*, **2008**, 51, 635-640	0	
9	Mosaic trisomy 9 at amniocentesis: prenatal diagnosis and molecular genetic analyses. *Taiwanese Journal of Obstetrics and Gynecology*, **2010**, 49, 341-50	1.6	18
8	New report of two patients with mosaic trisomy 9 presenting unusual features and longer survival. *Sao Paulo Medical Journal*, **2011**, 129, 428-32	1.6	10
7	Presenting physical characteristics, medical conditions, and developmental status of long-term survivors with trisomy 9 mosaicism. *American Journal of Medical Genetics, Part A*, **2011**, 155A, 1033-9	2.5	17
6	Current knowledge of prenatal diagnosis of mosaic autosomal trisomy in amniocytes: karyotype/phenotype correlations. *Prenatal Diagnosis*, **2015**, 35, 841-7	3.2	10
5	Birth of a child with trisomy 9 mosaicism syndrome associated with paternal isodisomy 9: case of a positive noninvasive prenatal test result unconfirmed by invasive prenatal diagnosis. *Molecular Cytogenetics*, **2015**, 8, 44	2	7
4	Twenty-five additional cases of trisomy 9 mosaic: Birth information, medical conditions, and developmental status. *American Journal of Medical Genetics, Part A*, **2015**, 167A, 997-1007	2.5	10
3	Prenatal Medicine. **2016**,		
2	Partial trisomy 9 (9pter->9q22.1) and partial monosomy 14 (14pter- >14q11.2) due to paternal translocation t(9;14)(q22.1;q11.2) in a case of Dysmorphic features. *Intractable and Rare Diseases Research*, **2019**, 8, 72-77	1.4	1
1	Fetal Karyotyping Using Chorionic Villus Samples. **1991**, 153-167		1