Julia Bijok

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1601539/publications.pdf

Version: 2024-02-01

		1163117	1125743
27	199	8	13
papers	citations	h-index	g-index
32	32	32	262
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	First trimester pregnancy loss: Clinical implications of genetic testing. Journal of Obstetrics and Gynaecology Research, 2017, 43, 23-29.	1.3	23
2	Prenatal diagnosis of congenital myopathies and muscular dystrophies. Clinical Genetics, 2016, 90, 199-210.	2.0	20
3	Triploidy - variability of sonographic phenotypes. Prenatal Diagnosis, 2017, 37, 774-780.	2.3	18
4	Clinical significance of the prenatal double bubble sign - single institution experience. Prenatal Diagnosis, 2015, 35, 1093-1096.	2.3	14
5	Targeted prenatal diagnosis of Pallister-Killian syndrome. Prenatal Diagnosis, 2017, 37, 446-452.	2.3	13
6	Firstâ€ŧrimester spontaneous pregnancy loss – molecular analysis using multiplex ligationâ€dependent probe amplification. Clinical Genetics, 2016, 89, 620-624.	2.0	11
7	Multiplex Ligation-dependent Probe Amplification (MLPA) – new possibilities of prenatal diagnosis. Ginekologia Polska, 2013, 84, 461-4.	0.7	11
8	Complex malformations involving the fetal body wall $\hat{a} \in \text{``definition and classification issues. Prenatal Diagnosis, 2017, 37, 1033-1039.}$	2.3	10
9	Extended genetic testing in fetuses with sonographic skeletal system abnormalities. Ultrasound in Obstetrics and Gynecology, 2022, 59, 660-667.	1.7	9
10	Maternal complications in molecularly confirmed diandric and digynic triploid pregnancies: single institution experience and literature review. Archives of Gynecology and Obstetrics, 2020, 301, 1139-1145.	1.7	8
11	Triploid pregnancy–Clinical implications. Clinical Genetics, 2021, 100, 368-375.	2.0	8
12	Prenatal diagnosis of craniosynostosis (compound Saethre-Chotzen syndrome phenotype) caused by a de novo complex chromosomal rearrangement (1; 4; 7) with a microdeletion of 7p21.3–7p15.3, including TWIST1 gene – a case report. Ginekologia Polska, 2014, 85, 541-4.	0.7	8
13	Distribution of diandric and digynic triploidy depending on gestational age. Journal of Assisted Reproduction and Genetics, 2021, 38, 2391-2395.	2.5	6
14	Usefulness of methylation-specific multiplex ligation-dependent probe amplification for identification of parental origin of triploidy. Journal of Human Genetics, 2020, 65, 889-894.	2.3	4
15	Implementation of Exome Sequencing in Prenatal Diagnosis and Impact on Genetic Counseling: The Polish Experience. Genes, 2022, 13, 724.	2.4	4
16	Prenatal diagnosis and clinical significance of cephalocele—A single institution experience and literature review. Prenatal Diagnosis, 2020, 40, 612-617.	2.3	3
17	Prenatal diagnosis of acrania/exencephaly/anencephaly sequence (AEAS): additional structural and genetic anomalies. Archives of Gynecology and Obstetrics, 2023, 307, 293-299.	1.7	3
18	Skinâ€Covered Bladder Exstrophy Diagnosed Antenatally. Journal of Ultrasound in Medicine, 2013, 32, 2043-2045.	1.7	2

#	Article	IF	CITATIONS
19	The location of the fetal ears: A hint for prenatal diagnosis of agnathiaâ€otocephaly complex. Journal of Clinical Ultrasound, 2019, 47, 369-371.	0.8	2
20	In-house genetic counseling increases the detection of abnormal karyotypes—a 26-year experience in prenatal diagnosis in a single tertiary referral hospital in Poland. Journal of Assisted Reproduction and Genetics, 2020, 37, 1999-2006.	2.5	2
21	Prenatal diagnosis of Duchenne and Becker muscular dystrophies: Underestimated problem of the secondary prevention of monogenetic disorders. Journal of Obstetrics and Gynaecology Research, 2017, 43, 1111-1121.	1.3	1
22	Twin pregnancies discordant for digynic triploidy – A case series. Taiwanese Journal of Obstetrics and Gynecology, 2021, 60, 139-141.	1.3	1
23	Prenatal diagnosis of glutaric acidemia type 2 with the use of exome sequencing — an up-to-date review and new case report. Ginekologia Polska, 2021, 92, 51-56.	0.7	1
24	Maternal blood intrauterine transfusions in the therapy of red-cell alloimmunization performed in three difficult cases. Ginekologia Polska, 2014, 85, .	0.7	1
25	Ultrasound evaluation of a bilobed placenta with †battledore cord insertion' — a report of an unusual case. Ginekologia Polska, 2020, 91, 100-100.	0.7	1
26	Foeto–maternal haemorrhage: An unexpected challenge. Journal of Obstetrics and Gynaecology, 2017, 37, 818-820.	0.9	0
27	Reply. Ultrasound in Obstetrics and Gynecology, 2022, 59, 698-699.	1.7	0