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Fetal holoprosencephaly: associated malformations and chromosomal defects

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#	Paper	IF	Citations
51	Ultrasonographically detectable markers of fetal chromosomal abnormalities. <i>Lancet, The</i> , <b>1992</b> , 340, 704-7	40	314
50	HIV infection at outcome of pregnancy in the Paris area, France. Lancet, The, 1992, 340, 707-9	40	22
49	Descriptive epidemiology of holoprosencephaly and arhinencephaly in metropolitan Atlanta, 1968-1992. <i>American Journal of Medical Genetics Part A</i> , <b>1996</b> , 66, 320-33		49
48	Molecular cytogenetic analysis of patients with holoprosencephaly and structural rearrangements of 7q. <i>American Journal of Medical Genetics Part A</i> , <b>1998</b> , 76, 51-57		14
47	First-trimester ultrasound diagnosis of holoprosencephaly: three case reports. <i>Ultrasound in Obstetrics and Gynecology</i> , <b>1999</b> , 13, 356-9	5.8	25
46	Holoprosencephaly in a Klinefelter fetus. American Journal of Medical Genetics Part A, <b>1999</b> , 85, 511-2		6
45	Investigation of the epidemiology and prenatal diagnosis of holoprosencephaly in the North of England. <i>American Journal of Obstetrics and Gynecology</i> , <b>2001</b> , 184, 1256-62	6.4	71
44	Prenatal diagnosis of 13q-syndrome in a fetus with holoprosencephaly and thumb agenesis. <i>Ultrasound in Obstetrics and Gynecology</i> , <b>2001</b> , 17, 166-8	5.8	20
43	Holoprosencephaly. Fetal and Maternal Medicine Review, <b>2001</b> , 12, 1-21		2
42	Brains and faces in holoprosencephaly: pre- and postnatal description of 30 cases. <i>Ultrasound in Obstetrics and Gynecology</i> , <b>2002</b> , 19, 24-38	5.8	87
41	Prenatal diagnosis of trisomy 18p and distal 21q22.3 deletion. <i>Prenatal Diagnosis</i> , <b>2003</b> , 23, 758-61	3.2	9
40	First-trimester sonographic diagnosis of holoprosencephaly: value of the "butterfly" sign. <i>Journal of Ultrasound in Medicine</i> , <b>2004</b> , 23, 761-5; quiz 766-7	2.9	61
39	Sonographic diagnosis of fetal adrenal hyperplasia: utility for prenatal corticotherapy. <i>Prenatal Diagnosis</i> , <b>2005</b> , 25, 1060-1	3.2	6
38	Prenatal diagnosis of cyclopia and proboscis in a fetus with normal chromosome at 13 weeks of gestation by three-dimensional transabdominal sonography. <i>Prenatal Diagnosis</i> , <b>2005</b> , 25, 1059-60	3.2	9
37	Prenatal Ultrasound: Brain. <b>2005</b> , 1157-1218		
36	Magnetic resonance evaluation of fetal ventriculomegaly-associated congenital malformations and lesions. <i>Seminars in Fetal and Neonatal Medicine</i> , <b>2005</b> , 10, 429-43	3.7	26
35	Neuroimaging evaluation of cerebral palsy. <i>Clinics in Perinatology</i> , <b>2006</b> , 33, 517-44	2.8	18

34	Prenatal diagnosis of trisomy 13: analysis of 28 cases. <i>Journal of Ultrasound in Medicine</i> , <b>2006</b> , 25, 429-	352.9	28
33	Prenatal findings and molecular cytogenetic analyses of partial trisomy 12q (12q24.32>qter) and partial monosomy 21q (21q22.2>qter). <i>Prenatal Diagnosis</i> , <b>2006</b> , 26, 313-20	3.2	8
32	Holoprosencephaly. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2008, 87, 13-	<b>37</b> 3	1
31	Holoprosencephaly. Orphanet Journal of Rare Diseases, 2007, 2, 8	4.2	247
30	An epidemiological study of holoprosencephaly from a regional congenital anomaly register: 1995-2004. <i>Prenatal Diagnosis</i> , <b>2007</b> , 27, 340-7	3.2	30
29	First trimester three-dimensional transvaginal imaging of alobar holoprosencephaly associated with proboscis and hypotelorism (ethmocephaly) in a 46,XX fetus. <i>Congenital Anomalies</i> (discontinued), 2008, 48, 51-5	1.1	13
28	Monosomy 18p presenting with holoprosencephaly and increased nuchal translucency in the first trimester: report of 2 cases. <i>Journal of Ultrasound in Medicine</i> , <b>2009</b> , 28, 1077-80	2.9	12
27	Prenatal sonographic features of fetuses in trisomy 13 pregnancies (I). <i>Taiwanese Journal of Obstetrics and Gynecology</i> , <b>2009</b> , 48, 210-7	1.6	15
26	Prenatal sonographic features of fetuses in trisomy 13 pregnancies (II). <i>Taiwanese Journal of Obstetrics and Gynecology</i> , <b>2009</b> , 48, 218-24	1.6	19
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24	Prenatal ultrasound diagnosis in 51 cases of holoprosencephaly: craniofacial anatomy, associated malformations, and genetics. <i>Cleft Palate-Craniofacial Journal</i> , <b>2010</b> , 47, 15-21	1.9	10
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22	Holoprosencephaly. <b>2012</b> , 1045-1063		
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17	[Frequency of holoprosencephaly in Chile]. <i>Revista Medica De Chile</i> , <b>2015</b> , 143, 874-9	0.5	1

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6	The Neural Crest and Craniofacial Malformations. <b>2014</b> , 219-269		
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1	Molecular cytogenetic analysis of patients with holoprosencephaly and structural rearrangements of 7q. <i>American Journal of Medical Genetics Part A</i> , <b>1998</b> , 76, 51-7		1