Reuven Sharony

List of Publications by Year
in descending order

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40	553	687220 13	713332 21
papers	citations	h-index	g-index

42	42	42	903
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Spectrum of PEX1 and PEX6 variants in Heimler syndrome. European Journal of Human Genetics, 2016, 24, 1565-1571.	1.4	49
2	When genotype is not predictive of phenotype: implications for genetic counseling based on 21,594 chromosomal microarray analysis examinations. Genetics in Medicine, 2018, 20, 128-131.	1.1	47
3	Cutâ€œoff value of nuchal translucency as indication for chromosomal microarray analysis. Ultrasound in Obstetrics and Gynecology, 2017, 50, 332-335.	0.9	44
4	Oocyte activation by calcium ionophore and congenital birth defects: a retrospective cohort study. Fertility and Sterility, 2016, 106, 590-596.e2.	0.5	40
5	Prenatal diagnosis of dacryocystocele: a possible marker for syndromes. Ultrasound in Obstetrics and Gynecology, 1999, 14, 71-73.	0.9	34
6	Chromosomal microarray analysis in fetuses with aberrant right subclavian artery. Ultrasound in Obstetrics and Gynecology, 2017, 49, 337-341.	0.9	34
7	Prenatal diagnosis of fetal cerebellar lesions: a case report and review of the literature. Prenatal Diagnosis, 1999, 19, 1077-1080.	1.1	28
8	Four USH2A Founder Mutations Underlie the Majority of Usher Syndrome Type 2 Cases among Non-Ashkenazi Jews. Genetic Testing and Molecular Biomarkers, 2008, 12, 289-294.	1.7	28
9	Granulosa cell tumors of the ovary: Do they have any unique ultrasonographic and color Doppler flow features?. International Journal of Gynecological Cancer, 2001, 11, 229-233.	1.2	20
10	Automated image analysis of placental villi and syncytial knots in histological sections. Placenta, 2017, 53, 113-118.	0.7	18
11	Fetal lateral neck cysts: the significance of associated findings. Prenatal Diagnosis, 2005, 25, 507-510.	1.1	17
12	Spectrum of genes for inherited hearing loss in the Israeli Jewish population, including the novel human deafness gene <scp>ATOH1</scp>. Clinical Genetics, 2020, 98, 353-364.	1.0	15
13	Spinocerebellar ataxia type 3 in Israel: phenotype and genotype of a Jew Yemenite subpopulation. Journal of Neurology, 2016, 263, 2207-2214.	1.8	13
14	Agenesis of the corpus callosum. An autopsy study in fetuses. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2016, 468, 219-230.	1.4	13
15	Ophthalmic manifestations of Heimler syndrome due to PEX6 mutations. Ophthalmic Genetics, 2018, 39, 384-390.	0.5	13
16	Familial lethal skeletal dysplasia with cloverleaf skull and multiple anomalies of brain, eye, face and heart: a new autosomal recessive multiple congenital anomalies syndrome. Clinical Genetics, 2002, 61, 369-374.	1.0	12
17	Telomere homeostasis in placentas from pregnancies with uncontrolled diabetes. Placenta, 2016, 44, 13-18.	0.7	10
18	Is the ratio of maternal serum to amniotic fluid AFP superior to serum levels as a predictor of pregnancy complications?. Archives of Gynecology and Obstetrics, 2016, 293, 767-770.	0.8	10

#	Article	IF	Citations
19	Morphometric characteristics of the umbilical cord and vessels in fetal growth restriction and pre-eclampsia. Early Human Development, 2016, 92, 57-62.	0.8	9
20	Correlation between prenatal and postnatal penile and clitoral measurements. Journal of Clinical Ultrasound, 2012, 40, 394-398.	0.4	8
21	Predictors of twin pregnancy after ovarian stimulation and intrauterine insemination in women with unexplained infertility. Human Fertility, 2017, 20, 200-203.	0.7	7
22	Prenatal course of metaphyseal anadysplasia associated with homozygous mutation in <scp>MMP9</scp> identified by exome sequencing. Clinical Genetics, 2017, 92, 645-648.	1.0	7
23	The association between maternal serum first trimester free Î²hCG, second trimester intact hCG levels and foetal growth restriction and preeclampsia. Journal of Obstetrics and Gynaecology, 2018, 38, 363-366.	0.4	7
24	Yemenite-Jewish families with Machadoâ€“Joseph disease (MJD/SCA3) share a recent common ancestor. European Journal of Human Genetics, 2019, 27, 1731-1737.	1.4	7
25	Chromosomal microarray findings in pregnancies with an isolated pelvic kidney. Journal of Perinatal Medicine, 2018, 47, 30-34.	0.6	6
26	Non-visualization of fetal gallbladder in microarray era â€“ a retrospective cohort study and review of the literature. Journal of Maternal-Fetal and Neonatal Medicine, 2019, 32, 2643-2648.	0.7	6
27	Low maternal serum concentrations of human chorionic gonadotropin as part of the triple test screening: a follow-up study. Journal of Maternal-Fetal and Neonatal Medicine, 2003, 13, 300-304.	0.7	5
28	Prenatal Diagnosis of Pericentric Inversion in Homologues of Chromosome 9: A Decision Dilemma. American Journal of Perinatology, 2007, 24, 137-140.	0.6	5
29	Genomic Alterations Are Enhanced in Placentas from Pregnancies with Fetal Growth Restriction and Preeclampsia: Preliminary Results. Molecular Syndromology, 2015, 6, 276-280.	0.3	5
30	Prenatal diagnosis of fetal cerebellar lesions: a case report and review of the literature. Prenatal Diagnosis, 1999, 19, 1077-80.	1.1	5
31	Prenatal sonographic detection of a lipomeningocele as a sacral lesion. , 2000, 28, 150-152.		4
32	Congenital deficiency of alpha-fetoprotein and associated chromosomal abnormality in the placenta. American Journal of Medical Genetics Part A, 2003, 121A, 113-117.	2.4	4
33	Adverse Outcome of Pregnancies with Extremely High Levels of Maternal Serum Human Chorionic Gonadotropin. Fetal Diagnosis and Therapy, 2008, 23, 233-236.	0.6	4
34	Three peaks in the polymerase chain reaction fragile X analysis. Journal of Medical Screening, 2012, 19, 112-115.	1.1	4
35	The magnitude of elevated maternal serum human chorionic gonadotropin and pregnancy complications. Journal of Obstetrics and Gynaecology, 2017, 37, 576-579.	0.4	4
36	The yield of the prenatal work-up in intrauterine growth restriction and the spectrum of fetal abnormalities detected postnatally. Journal of Maternal-Fetal and Neonatal Medicine, 2019, 32, 753-759.	0.7	3

#	Article	IF	Citations
37	Based on a cohort of 52,879 microarrays, recurrent intragenic FBN2 deletion encompassing exons 1â€“8 does not cause Beals syndrome. European Journal of Medical Genetics, 2020, 63, 104008.	0.7	3
38	Down Syndrome Screening: Evidence that Test Results Differ According to Phenotype. Journal of Fetal Medicine, 2016, 03, 137-141.	0.1	2
39	The impact of third-trimester genetic counseling. Archives of Gynecology and Obstetrics, 2018, 297, 659-665.	0.8	1
40	The mid-gestation triple test profile among women diagnosed with vasa previa. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 1402-1406.	0.7	1