Rivka Sukenik-Halevy

List of Publications by Year
in descending order

39	545	687363 13	713466 21
papers	citations	h-index	g-index

41	41	41	942
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The yield of chromosomal microarray analysis among pregnancies terminated due to fetal malformations. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 336-340.	1.5	4
2	Challenges in variant interpretation in prenatal exome sequencing. European Journal of Medical Genetics, 2022, 65, 104410.	1.3	3
3	Prenatal and postnatal chromosomal microarray analysis in 885 cases of various congenital heart defects. Archives of Gynecology and Obstetrics, 2022, 306, 1007-1013.	1.7	4
4	The prevalence of prenatal sonographic findings in postnatal diagnostic exome sequencing performed for neurocognitive phenotypes: A cohort study. Prenatal Diagnosis, 2022, , .	2.3	4
5	The Yield of Chromosomal Microarray in Pregnancies Complicated with Fetal Growth Restriction Can Be Predicted According to Clinical Parameters. Fetal Diagnosis and Therapy, 2021, 48, 140-148.	1.4	4
6	Chromosomal microarray should be performed for cases of fetal short long bones detected prenatally. Archives of Gynecology and Obstetrics, 2021, 303, 85-92.	1.7	5
7	The diagnostic efficacy of exome data analysis using fixed neurodevelopmental gene lists: Implications for prenatal setting. Prenatal Diagnosis, 2021, 41, 701-707.	2.3	3
8	The diagnostic potential of targeted imaging of the fetal pancreas. Prenatal Diagnosis, 2021, 41, 828-834.	2.3	5
9	Chromosomal Microarray Analysis in Pregnancies With Corpus Callosum or Posterior Fossa Anomalies. Neurology: Genetics, 2021, 7, e585.	1.9	4
10	Does parity affect pregnancy outcomes in the elderly gravida?. Archives of Gynecology and Obstetrics, 2020, 301, 85-91.	1.7	12
11	Ten points to consider when providing genetic counseling for variants of incomplete penetrance and variable expressivity detected in a prenatal setting. Acta Obstetricia Et Gynecologica Scandinavica, 2020, 99, 1427-1429.	2.8	3
12	Should We Report 15q11.2 BP1-BP2 Deletions and Duplications in the Prenatal Setting?. Journal of Clinical Medicine, 2020, 9, 2602.	2.4	8
13	Smithâ€“Lemliâ€“Opitz syndrome: what is the actual risk for couples carriers of the DHCR7:c.964-1G>C variant?. European Journal of Human Genetics, 2020, 28, 938-942.	2.8	8
14	Bi-allelic ADARB1 Variants Associated with Microcephaly, Intellectual Disability, and Seizures. American Journal of Human Genetics, 2020, 106, 467-483.	6.2	31
15	The yield of chromosomal microarray testing for cases of abnormal fetal head circumference. Journal of Perinatal Medicine, 2020, 48, 553-558.	1.4	7
16	The rare 13q33â€“q34 microdeletions: eight new patients and review of the literature. Human Genetics, 2019, 138, 1145-1153.	3.8	11
17	A Rare Case of 7 Simultaneous Arterial Dissections and Review of The Literature. Vascular and Endovascular Surgery, 2019, 53, 617-622.	0.7	4
18	A new method for evaluating short fetal corpus callosum. Prenatal Diagnosis, 2019, 39, 1283-1290.	2.3	7

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19	Microarray findings in pregnancies with oligohydramnios â€“ a retrospective cohort study and literature review. Journal of Perinatal Medicine, 2019, 48, 53-58.	1.4	6
20	Telomere Homeostasis and Senescence Markers Are Differently Expressed in Placentas From Pregnancies With Early- Versus Late-Onset Preeclampsia. Reproductive Sciences, 2019, 26, 1203-1209.	2.5	20
21	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.	1.5	3
22	What are the prevalence, characteristics and significance of fetal lateral neck cysts detected in an early anatomical scan?. Archives of Gynecology and Obstetrics, 2018, 298, 51-58.	1.7	2
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.	0.9	7
24	The impact of third-trimester genetic counseling. Archives of Gynecology and Obstetrics, 2018, 297, 659-665.	1.7	1
25	Prediction of the efficacy of dinoprostone slow release vaginal insert (Propess) for cervical ripening: A prospective cohort study. Journal of Obstetrics and Gynaecology Research, 2018, 44, 1739-1746.	1.3	23
26	Skin closure at cesarean delivery, glue vs subcuticular sutures: aÂrandomized controlled trial. American Journal of Obstetrics and Gynecology, 2017, 216, 406.e1-406.e5.	1.3	24
27	Deficiency of the sphingosine-1-phosphate lyase SGPL1 is associated with congenital nephrotic syndrome and congenital adrenal calcifications. Human Mutation, 2017, 38, 365-372.	2.5	71
28	Diagnostic accuracy, work-up, and outcomes of pregnancies with clubfoot detected by prenatal sonography. Prenatal Diagnosis, 2017, 37, 754-763.	2.3	28
29	The yield and complications of amniocentesis performed after 24 weeks of gestation. Archives of Gynecology and Obstetrics, 2017, 296, 69-75.	1.7	8
30	Oocyte activation by calcium ionophore and congenital birth defects: a retrospective cohort study. Fertility and Sterility, 2016, 106, 590-596.e2.	1.0	40
31	Clinical aspects of prenatally detected congenital heart malformations and the yield of chromosomal microarray analysis. Prenatal Diagnosis, 2016, 36, 1185-1191.	2.3	20
32	Telomere homeostasis in trophoblasts and in cord blood cells from pregnancies complicated with preeclampsia. American Journal of Obstetrics and Gynecology, 2016, 214, 283.e1-283.e7.	1.3	40
33	The time-consuming demands of the practice of medical genetics in the era of advanced genomic testing. Genetics in Medicine, 2016, 18, 372-377.	2.4	35
34	Amniocytes from aneuploidy embryos have enhanced random aneuploidy and signs of senescence â€" Can these findings be related to medical problems?. Gene, 2015, 562, 232-235.	2.2	16
35	Microscopic chromosome Xp distal deletions â€“ a challenging issue in prenatal genetic counseling. Prenatal Diagnosis, 2014, 34, 592-597.	2.3	2
36	Telomere shortening in intra uterine growth restriction placentas. Early Human Development, 2014, 90, 465-469.	1.8	28

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37	Compliance for genetic screening in the Arab population in Israel. Israel Medical Association Journal, 2012, 14, 538-42.	0.1	9
38	Large-Scale Population Carrier Screening for Spinal Muscular Atrophy in Israelâ€"Effect of Ethnicity on the False-Negative Rate. Genetic Testing and Molecular Biomarkers, 2010, 14, 319-324.	0.7	7
39	Telomere aggregate formation in placenta specimens of pregnancies complicated with pre-eclampsia. Cancer Genetics and Cytogenetics, 2009, 195, 27-30.	1.0	28