Emmanuel Spaggiari

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Pathological and sonographic review of early isolated severe lower urinary tract obstruction and implications for prenatal treatment. Ultrasound in Obstetrics and Gynecology, 2022, 59, 513-521.	1.7	3

2 Prenatal biochemical diagnosis of two forms of congenital diarrheal disorders (congenital chloride) Tj ETQq0 0 0 rgBT_3/Overlock 10 Tf 50

3	Contribution of threeâ€dimensional ultrasound and threeâ€dimensional helical computed tomography to prenatal diagnosis of Stickler syndrome. Ultrasound in Obstetrics and Gynecology, 2019, 54, 279-280.	1.7	1
4	Comparison of biochemical analysis of fetal serum and fetal urine in the prediction of postnatal renal outcome in lower urinary tract obstruction. Prenatal Diagnosis, 2018, 38, 555-560.	2.3	8
5	Preterm premature rupture of membranes is a collateral effect of improvement in perinatal outcomes following fetoscopic coagulation of chorionic vessels for twin–twin transfusion syndrome: a retrospective observational study of 1092 cases. BJOC: an International Journal of Obstetrics and Gynaecology. 2018. 125. 1154-1162.	2.3	40
6	Prenatal diagnosis of megacystis microcolon intestinal hypoperistalsis syndrome by biochemical analysis of fetal urine. Prenatal Diagnosis, 2018, 38, 585-590.	2.3	6
7	Pitfall in first-trimester diagnosis of chorionicity in twin pregnancy. Ultrasound in Obstetrics and Gynecology, 2017, 49, 277-278.	1.7	6
8	Sequential fetal serum \hat{l}^2 2-microglobulin to predict postnatal renal function in bilateral or low urinary tract obstruction. Ultrasound in Obstetrics and Gynecology, 2017, 49, 617-622.	1.7	22
9	Impact on spina bifida screening of shifting prenatal Down syndrome maternal serum screening from the second trimester to the first. Prenatal Diagnosis, 2017, 37, 673-679.	2.3	4
10	First fetal case of the 8q24.3 contiguous genes syndrome. American Journal of Medical Genetics, Part A, 2016, 170, 239-242.	1.2	17
11	Impact of Including or Removing Nuchal Translucency Measurement on the Detection and False-Positive Rates of First-Trimester Down Syndrome Screening. Fetal Diagnosis and Therapy, 2016, 40, 214-218.	1.4	6
12	Untreated maternal hyperthyroidism responsible of early neonatal demise. Journal of Obstetrics and Gynaecology, 2016, 36, 66-67.	0.9	0
13	Performance of prenatal diagnosis in esophageal atresia. Prenatal Diagnosis, 2015, 35, 888-893.	2.3	44
14	Prenatal prediction of pulmonary arterial hypertension in congenital diaphragmatic hernia. Ultrasound in Obstetrics and Gynecology, 2015, 45, 572-577.	1.7	29
15	Outcome and etiologies of fetal megacystis according to the gestational age at diagnosis. Prenatal Diagnosis, 2013, 33, 1162-1166.	2.3	37
16	Fetal obstructive uropathy complicated by urinary ascites: outcome and prognostic value of fetal serum β-2-microglobulin. Ultrasound in Obstetrics and Gynecology, 2013, 41, 185-189.	1.7	14
17	Mowat–Wilson syndrome in a fetus with antenatal diagnosis of short corpus callosum: Advocacy for standard autopsy. European Journal of Medical Genetics, 2013, 56, 297-300.	1.3	12
18	Management strategy in pregnancies with elevated second-trimester maternal serum alpha-fetoprotein based on a second assay. American Journal of Obstetrics and Gynecology, 2013, 208, 303.e1-303.e7.	1.3	15

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19	Ultrasound detection of eyelashes: a clue for prenatal diagnosis of Cornelia de Lange syndrome. Ultrasound in Obstetrics and Gynecology, 2013, 41, 341-342.	1.7	15
20	Outcome following prenatal diagnosis of severe bilateral renal hypoplasia. Prenatal Diagnosis, 2013, 33, 1167-1172.	2.3	14
21	Prognostic value of a hernia sac in congenital diaphragmatic hernia. Ultrasound in Obstetrics and Gynecology, 2013, 41, 286-290.	1.7	25
22	Prognosis and outcome of pregnancies exposed to renin–angiotensin system blockers. Prenatal Diagnosis, 2012, 32, 1071-1076.	2.3	29
23	Outcome in fetuses with isolated congenital diaphragmatic hernia with increased nuchal translucency thickness in first trimester. Prenatal Diagnosis, 2012, 32, 268-271.	2.3	10
24	Prenatal phenotype of congenital hyperparathyroidism. Prenatal Diagnosis, 2012, 32, 906-908.	2.3	2