

Sara Cabet

List of Publications by Year in descending order

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9
papers

114
citations

1684188
5
h-index

1474206
9
g-index

9
all docs

9
docs citations

9
times ranked

396
citing authors

#	ARTICLE	IF	CITATIONS
1	Prenatal diagnosis of Aicardi syndrome based on a suggestive imaging pattern: A multicenter case series. <i>Prenatal Diagnosis</i> , 2022, 42, 484-494.	2.3	8
2	Prenatal imaging features related to <i>RAC3</i> pathogenic variant and differential diagnoses. <i>Prenatal Diagnosis</i> , 2022, 42, 478-481.	2.3	2
3	Prenatal diagnosis of congenital perineal lipoma: tip of urorectal septum malformation sequence?. <i>Ultrasound in Obstetrics and Gynecology</i> , 2022, 60, 139-141.	1.7	1
4	Two different prenatal imaging cerebral patterns of tubulinopathy. <i>Ultrasound in Obstetrics and Gynecology</i> , 2021, 57, 493-497.	1.7	14
5	SPEN haploinsufficiency causes a neurodevelopmental disorder overlapping proximal 1p36 deletion syndrome with an epismature of X chromosomes in females. <i>American Journal of Human Genetics</i> , 2021, 108, 502-516.	6.2	48
6	Prenatal cerebral imaging features of a new syndromic entity related to KIAA1109 pathogenic variants mimicking tubulinopathy. <i>Prenatal Diagnosis</i> , 2020, 40, 276-281.	2.3	4
7	Novel truncating and missense variants extending the spectrum of EMC1-related phenotypes, causing autism spectrum disorder, severe global development delay and visual impairment. <i>European Journal of Medical Genetics</i> , 2020, 63, 103897.	1.3	11
8	A novel truncating variant p.(Arg297*) in the GRM1 gene causing autosomal-recessive cerebellar ataxia with juvenile-onset. <i>European Journal of Medical Genetics</i> , 2019, 62, 103726.	1.3	7
9	A novel lethal recognizable polymicrogyric syndrome caused by ATP1A2 homozygous truncating variants. <i>Brain</i> , 2019, 142, 3367-3374.	7.6	19