Ingrid Anne Mandy Schierz

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1170180/publications.pdf

Version: 2024-02-01

23	191	1039406 9	1125271 13
papers	citations	h-index	g-index

23	23	23	131
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical and genetic approach in the characterization of newborns with anorectal malformation. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 4513-4520.	0.7	3
2	Hypertrophic pyloric stenosis masked by kidney failure in a male infant with a contiguous gene deletion syndrome at Xp22.31 involving the steroid sulfatase gene: case report. Italian Journal of Pediatrics, 2022, 48, 19.	1.0	6
3	Interstitial deletions of chromosome 1p: novel 1p31.3p22.2 microdeletion in a newborn with craniosynostosis, coloboma and cleft palate, and review of the genomic and phenotypic profiles. Italian Journal of Pediatrics, 2022, 48, 38.	1.0	9
4	2q13 microdeletion syndrome: Report on a newborn with additional features expanding the phenotype. Clinical Case Reports (discontinued), 2021, 9, e04289.	0.2	12
5	Novel missense mutation of the TP63 gene in a newborn with Hay-Wells/Ankyloblepharon-Ectodermal defects-Cleft lip/palate (AEC) syndrome: clinical report and follow-up. Italian Journal of Pediatrics, 2021, 47, 196.	1.0	19
6	Congenital pelvic skeletal anomalies: Clinical and radiographic evaluation of newborns with gastrointestinal malformation. Early Human Development, 2020, 141, 104945.	0.8	3
7	Novel LRPPRC compound heterozygous mutation in a child with early-onset Leigh syndrome French-Canadian type: case report of an Italian patient. Italian Journal of Pediatrics, 2020, 46, 140.	1.0	18
8	Infant developmental profile of Crisponi syndrome due to compound heterozygosity for CRLF1 deletion. Clinical Dysmorphology, 2020, 29, 141-143.	0.1	11
9	A refugee newborn with heart failure and initial hydrops: Diagnostic clues of spectral Doppler examinations. Journal of Paediatrics and Child Health, 2020, 56, 1315-1315.	0.4	0
10	Total colonic aganglionosis and cleft palate in a newborn with Janus-cysteine 618 mutation of RET proto-oncogene: a case report. Italian Journal of Pediatrics, 2020, 46, 135.	1.0	3
11	Neonatal hyperinsulinemic hypoglycemia: case report of kabuki syndrome due to a novel KMT2D splicing-site mutation. Italian Journal of Pediatrics, 2020, 46, 136.	1.0	20
12	Growth patterns and associated risk factors of congenital malformations in twins. Italian Journal of Pediatrics, 2020, 46, 73.	1.0	5
13	Recognizable neonatal clinical features of aplasia cutis congenita. Italian Journal of Pediatrics, 2020, 46, 25.	1.0	12
14	Neonatal ten-year retrospective study on neural tube defects in a second level University Hospital. Italian Journal of Pediatrics, 2020, 46, 72.	1.0	10
15	A Case of Cardiomyopathy Due to Premature Ductus Arteriosus Closure: The Flip Side of Paracetamol. Pediatrics, 2018, 141, .	1.0	8
16	Transitional hemodynamics in infants of diabetic mothers by targeted neonatal echocardiography, electrocardiography and peripheral flow study. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 1578-1585.	0.7	14
17	An unusual association of leftâ€sided gastroschisis and persistent right umbilical vein. Clinical Case Reports (discontinued), 2018, 6, 2511-2512.	0.2	1
18	Clinical cardiac assessment in newborns with prenatally diagnosed intrathoracic masses. Italian Journal of Pediatrics, 2018, 44, 98.	1.0	1

#	Article	IF	Citations
19	Etiological heterogeneity and clinical variability in newborns with esophageal atresia. Italian Journal of Pediatrics, 2018, 44, 19.	1.0	8
20	Dilated azygos arch mimicking an aortic arch anomaly during thoracic surgery. Early Human Development, 2017, 111, 20-22.	0.8	3
21	Early intestinal perforation secondary to congenital mesenteric defects. Journal of Pediatric Surgery Case Reports, 2016, 8, 10-12.	0.1	1
22	Congenital heart defects in newborns with apparently isolated single gastrointestinal malformation: A retrospective study. Early Human Development, 2016, 103, 43-47.	0.8	8
23	Predictive Factors of Abdominal Compartment Syndrome in Neonatal Age. American Journal of Perinatology, 2014, 31, 049-054.	0.6	16