Ettore Piro

List of Publications by Year
in descending order

52 papers	660 citations	516215 16 h-index	676716 22 g-index
53 all docs	53 docs citations	53 times ranked	678 citing authors

#	Article	IF	Citations
1	The world of twins: an update. Journal of Maternal-Fetal and Neonatal Medicine, 2010, 23, 59-62.	0.7	46
2	Kabuki make-up (Niikawa-Kuroki) syndrome: Clinical and radiological observations in two sicilian children. Pediatric Radiology, 1991, 21, 428-431.	1.1	33
3	Clinical and molecular characterization of 112 single-center patients with Neurofibromatosis type 1. Italian Journal of Pediatrics, 2018, 44, 45.	1.0	32
4	Prematurity and twinning. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 6-10.	0.7	31
5	Interstitial deletion of chromosome 2p15-16.1: Report of two patients and critical review of current genotypeâ€“phenotype correlation. European Journal of Medical Genetics, 2012, 55, 238-244.	0.7	30
6	Congenital hepatic mesenchymal hamartoma associated with mesenchymal stem villous hyperplasia of the placenta: case report. Journal of Pediatric Surgery, 2005, 40, e37-e39.	0.8	28
7	Intrauterine growth pattern and birthweight discordance in twin pregnancies: a retrospective study. Italian Journal of Pediatrics, 2014, 40, 43.	1.0	28
8	NF1 microdeletion syndrome: case report of two new patients. Italian Journal of Pediatrics, 2019, 45, 138.	1.0	27
9	Autosomal recessive polycystic kidney disease: case report of a newborn with rare PKHD1 mutation, rapid renal enlargement and early fatal outcome. Italian Journal of Pediatrics, 2020, 46, 154.	1.0	20
10	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
11	Oxidative Stress in Preterm Infants: Overview of Current Evidence and Future Prospects. Pharmaceuticals, 2020, 13, 145.	1.7	20
12	Dyke-Davidoff-Masson syndrome: case report of fetal unilateral ventriculomegaly and hypoplastic left middle cerebral artery. Italian Journal of Pediatrics, 2013, 39, 32.	1.0	19
13	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
14	Esophageal atresia and Beckwithâ€“Wiedemann syndrome in one of the naturally conceived discordant newborn twins: first report. Clinical Case Reports (discontinued), 2018, 6, 399-401.	0.2	18
15	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
16	Perlman syndrome: Clinical report and nine-year follow-up. American Journal of Medical Genetics, Part A, 2005, 139A, 131-135.	0.7	17
17	Intrauterine growth restriction and congenital malformations: a retrospective epidemiological study. Italian Journal of Pediatrics, 2013, 39, 23.	1.0	16
18	Predictive Factors of Abdominal Compartment Syndrome in Neonatal Age. American Journal of Perinatology, 2014, 31, 049-054.	0.6	16

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19	Paternal uniparental disomy chromosome 14â€like syndrome due a maternal de novo 160â€‰kb deletion at the 14q32.2 region not encompassing the IGâ€•and the MEG3â€DMRs: Patient report and genotypeâ€“phenotype correlation. American Journal of Medical Genetics, Part A, 2015, 167, 3130-3138.	0.7	15
20	A premature infant with Costello syndrome due to a rare G13C HRAS mutation. American Journal of Medical Genetics, Part A, 2009, 149A, 487-489.	0.7	14
21	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
22	Cardio-facio-cutaneous syndrome and gastrointestinal defects: report on a newborn with 19p13.3 deletion including the MAPÂ2â€‰K2 gene. Italian Journal of Pediatrics, 2022, 48, 65.	1.0	14
23	The social role of pediatrics in the past and present times. Italian Journal of Pediatrics, 2021, 47, 239.	1.0	13
24	Recognizable neonatal clinical features of aplasia cutis congenita. Italian Journal of Pediatrics, 2020, 46, 25.	1.0	12
25	2q13 microdeletion syndrome: Report on a newborn with additional features expanding the phenotype. Clinical Case Reports (discontinued), 2021, 9, e04289.	0.2	12
26	Infant developmental profile of Crisponi syndrome due to compound heterozygosity for CRLF1 deletion. Clinical Dysmorphology, 2020, 29, 141-143.	0.1	11
27	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
28	A novel mutation in KCNQ3 â€related benign familial neonatal epilepsy: electroclinical features and neurodevelopmental outcome. Epileptic Disorders, 2019, 21, 87-91.	0.7	10
29	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
30	Congenital heart defects in newborns with apparently isolated single gastrointestinal malformation: A retrospective study. Early Human Development, 2016, 103, 43-47.	0.8	8
31	A Case of Cardiomyopathy Due to Premature Ductus Arteriosus Closure: The Flip Side of Paracetamol. Pediatrics, 2018, 141, .	1.0	8
32	Etiological heterogeneity and clinical variability in newborns with esophageal atresia. Italian Journal of Pediatrics, 2018, 44, 19.	1.0	8
33	10qter deletion: A new case. American Journal of Medical Genetics, Part A, 2008, 146A, 2435-2438.	0.7	6
34	Characterization of a complex rearrangement involving chromosomes 1, 4 and 8 by fish and array-CGH. Journal of Applied Genetics, 2012, 53, 285-288.	1.0	6
35	Benign familial infantile epilepsy associated with KCNQ3 mutation: a rare occurrence or an underestimated event?. Epileptic Disorders, 2020, 22, 807-810.	0.7	6
36	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

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37	Methemoglobinemia Associated with Late-Onset Neonatal Sepsis: A Single-Center Experience. American Journal of Perinatology, 2019, 36, 1510-1513.	0.6	5
38	Growth patterns and associated risk factors of congenital malformations in twins. Italian Journal of Pediatrics, 2020, 46, 73.	1.0	5
39	Case Report: Unusual Clinical Presentation of a Rare Cardiac Inflammatory Myofibroblastic Tumor in Children: The Differential Diagnosis With Pediatric Emergencies. Frontiers in Pediatrics, 2021, 9, 718157.	0.9	5
40	Migraine in Children Under 7ÂYears of Age: a Review. Current Pain and Headache Reports, 2020, 24, 79.	1.3	4
41	Dilated azygos arch mimicking an aortic arch anomaly during thoracic surgery. Early Human Development, 2017, 111, 20-22.	0.8	3
42	Congenital pelvic skeletal anomalies: Clinical and radiographic evaluation of newborns with gastrointestinal malformation. Early Human Development, 2020, 141, 104945.	0.8	3
43	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
44	Delayed neonatal visual evoked potentials are associated to asymmetric growth pattern in twins. Clinical Neurophysiology, 2020, 131, 744-749.	0.7	3
45	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
46	Screening of Endothelial Expression Libraries for the Identification of Novel Autoantigens Involved in Distinct Autoimmune Diseases Characterized by Endothelial Dysfunction. Annals of the New York Academy of Sciences, 2007, 1109, 178-184.	1.8	2
47	From Neonatal Intensive Care to Neurocritical Care: Is It Still a Mirage? The Sicilian Multicenter Project. Critical Care Research and Practice, 2021, 2021, 1-9.	0.4	2
48	Spontaneous Resorption of an Occipital Meningocele: Computed Tomography and Magnetic Resonance Imaging Evaluation. Journal of Pediatric Neurology, 2016, 14, 043-046.	0.0	1
49	Clinical cardiac assessment in newborns with prenatally diagnosed intrathoracic masses. Italian Journal of Pediatrics, 2018, 44, 98.	1.0	1
50	Management of multiple pregnancy with an affected twin. Italian Journal of Pediatrics, 2015, 41, .	1.0	0
51	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
52	Developments in pediatrics in 2020: choices in allergy, autoinflammatory disorders, critical care, endocrinology, genetics, infectious diseases, microbiota, neonatology, neurology, nutrition, ortopedics, respiratory tract illnesses and rheumatology. Italian Journal of Pediatrics, 2021, 47, 232.	1.0	0