Gregorio Serra

List of Publications by Year
in descending order

21 papers	400 citations	840119 11 h-index	794141 19 g-index
28 all docs	28 docs citations	28 times ranked	165 citing authors

#	Article	IF	Citations
1	Smartphone use and addiction during the coronavirus disease 2019 (COVID-19) pandemic: cohort study on 184 Italian children and adolescents. Italian Journal of Pediatrics, 2021, 47, 150.	1.0	99
2	Clinical and molecular characterization of 112 single-center patients with Neurofibromatosis type 1. Italian Journal of Pediatrics, 2018, 44, 45.	1.0	32
3	NF1 microdeletion syndrome: case report of two new patients. Italian Journal of Pediatrics, 2019, 45, 138.	1.0	27
4	Recommendations for neonatologists and pediatricians working in first level birthing centers on the first communication of genetic disease and malformation syndrome diagnosis: consensus issued by 6 Italian scientific societies and 4 parentsâ€™ associations. Italian Journal of Pediatrics, 2021, 47, 94.	1.0	25
5	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
6	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
7	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
8	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
9	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
10	Jacobsen syndrome and neonatal bleeding: report on two unrelated patients. Italian Journal of Pediatrics, 2021, 47, 147.	1.0	18
11	Novel SCNN1A gene splicing-site mutation causing autosomal recessive pseudohypoaldosteronism type 1 (PHA1) in two Italian patients belonging to the same small town. Italian Journal of Pediatrics, 2021, 47, 138.	1.0	14
12	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
13	The social role of pediatrics in the past and present times. Italian Journal of Pediatrics, 2021, 47, 239.	1.0	13
14	2q13 microdeletion syndrome: Report on a newborn with additional features expanding the phenotype. Clinical Case Reports (discontinued), 2021, 9, e04289.	0.2	12
15	Infant developmental profile of Crisponi syndrome due to compound heterozygosity for CRLF1 deletion. Clinical Dysmorphology, 2020, 29, 141-143.	0.1	11
16	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
17	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
18	Perinatal and newborn care in a two years retrospective study in a first level peripheral hospital in Sicily (Italy). Italian Journal of Pediatrics, 2019, 45, 152.	1.0	7

#	Article	IF	Citations
19	Growth patterns and associated risk factors of congenital malformations in twins. Italian Journal of Pediatrics, 2020, 46, 73.	1.0	5
20	Quando lâ€™amnios si rompe troppo presto eâ€¦ da solo. Medico E Bambino, 2022, 41, 387-389.	0.1	5
21	Il neonato che â€œsa di saleâ€. Medico E Bambino, 2021, 40, 119-122.	0.1	0