BariÅŽilmaz

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5526406/publications.pdf

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#	Article	IF	CITATIONS
1	Homozygous c.130–131 ins A (pW44X) mutation in the HAX1 gene as the most common cause of congenital neutropenia in Turkey: Report from the Turkish Severe Congenital Neutropenia Registry. Pediatric Blood and Cancer, 2019, 66, e27923.	1.5	16
2	Outcomes of Osteosarcoma in Children Without High-Dose Methotrexate: Could It Be Less Toxic Without Effecting Survival Rates?. Journal of Adolescent and Young Adult Oncology, 2022, 11, 252-258.	1.3	2
3	The Relationship of Blood Prolactin, Cortisol and Thyroid Hormones in Prematural Newborns with Respiratory Distress Syndrome. Research in Pediatrics & Neonatology, 2019, 3, .	0.2	2
4	Childhood Myelodysplastic Syndrome Progressing to Pre-B Acute Lymphoblastic Leukemia in Two Children with Trisomy 5 and Trisomy 8. Turkiye Klinikleri Pediatri, 2021, 30, 156-159.	0.0	0
5	Severe Necrotizing Perianal Cellulitis in a Child with Acute Lymphoblastic Leukemia that Developed After Rectal Enema Use. Turkish Journal of Pediatric Disease, 0, , .	0.0	O
6	Successful Treatment of Two Children's with Refractory Immune Thrombocytopenic Purpura with Eltrombopag. Academic Journal of Pediatrics & Neonatology, 2018, 7, .	0.1	0
7	A Rare Cause of Cytopenias in Childhood: Paroxysmal Nocturnal Hemoglobinuria. Research in Pediatrics & Neonatology, 2018, 3, .	0.2	0