

Telma Francisco

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6335118/publications.pdf>

Version: 2024-02-01

12
papers

110
citations

1937685

4
h-index

1281871

11
g-index

12
all docs

12
docs citations

12
times ranked

150
citing authors

#	ARTICLE	IF	CITATIONS
1	Treatment and long-term outcome in primary distal renal tubular acidosis. <i>Nephrology Dialysis Transplantation</i> , 2019, 34, 981-991.	0.7	75
2	Aetiology, course and treatment of acute tubulointerstitial nephritis in paediatric patients: a cross-sectional web-based survey. <i>BMJ Open</i> , 2021, 11, e047059.	1.9	11
3	Multiple haemangiomas, diaphragmatic eventration and Beckwith-Wiedemann syndrome: an unusual association. <i>BMJ Case Reports</i> , 2013, 2013, bcr2013010077-bcr2013010077.	0.5	6
4	Parathyroid hormone and phosphate homeostasis in patients with Bartter and Gitelman syndrome: an international cross-sectional study. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 2474-2486.	0.7	5
5	Bilateral facial palsy: a form of neuroborreliosis presentation in paediatric age. <i>BMJ Case Reports</i> , 2013, 2013, bcr2012008060-bcr2012008060.	0.5	3
6	Beyond polycystic kidney disease. <i>BMJ Case Reports</i> , 2017, 2017, bcr-2017-220766.	0.5	2
7	Value-based decision-making for orphan drugs with multiple criteria decision analysis: burosumab for the treatment of X-linked hypophosphatemia. <i>Current Medical Research and Opinion</i> , 2021, 37, 1021-1030.	1.9	2
8	Challenges in orthopaedic management of Parkes-Weber syndrome. <i>BMJ Case Reports</i> , 2013, 2013, bcr2013008800-bcr2013008800.	0.5	2
9	X-linked hypophosphatemic rickets: a new mutation. <i>Jornal Brasileiro De Nefrologia: Orgao Oficial De Sociedades Brasileira E Latino-Americana De Nefrologia</i> , 2021, 43, 279-282.	0.9	2
10	Immunological Reconstitution Inflammatory Syndrome and Thrombotic Microangiopathy: Severe Complications in a Child With Acquired Immunodeficiency Syndrome. <i>Clinical Pediatrics</i> , 2019, 58, 1022-1026.	0.8	1
11	The urine as a diagnostic key for a homozygous EGFR mutation. <i>Portuguese Journal of Nephrology & Hypertension</i> , 2022, 36, 15-19.	0.1	1
12	The Kidney Genetics Clinic: delivering precision medicine for kidney patients. <i>Portuguese Journal of Nephrology & Hypertension</i> , 2021, 35, .	0.1	0