Mona Hafez

List of Publications by Year in descending order

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1307594 1281871 12 316 7 11 citations g-index h-index papers 12 12 12 397 citing authors all docs docs citations times ranked

#	Article	IF	CITATIONS
1	Liraglutide in Children and Adolescents with Type 2 Diabetes. New England Journal of Medicine, 2019, 381, 637-646.	27.0	209
2	Abnormal Glucose Tolerance in \hat{I}^2 -Thalassemia: Assessment of Risk Factors. Hemoglobin, 2009, 33, 101-108.	0.8	30
3	Monogenic leptin deficiency in early childhood obesity. Pediatric Obesity, 2020, 15, e12574.	2.8	18
4	Clinical Phenotypes and Immunological Characteristics of 18 Egyptian LRBA Deficiency Patients. Journal of Clinical Immunology, 2020, 40, 820-832.	3.8	17
5	Changes of cardiac functions after hemodialysis session in pediatric patients with end-stage renal disease: conventional echocardiography and two-dimensional speckle tracking study. Pediatric Nephrology, 2020, 35, 861-870.	1.7	13
6	Prediction of Residual Valvular Lesions in Rheumatic Heart Disease: Role of Adhesion Molecules. Pediatric Cardiology, 2013, 34, 583-590.	1.3	11
7	Evaluation of carotid artery access in comparison with femoral artery access in neonatal percutaneous stenting of ductus arteriosus. Cardiology in the Young, 2021, 31, 1465-1471.	0.8	7
8	Cardiac implications of multisystem inflammatory syndrome associated with COVID-19 in children under the age of 5 years. Cardiology in the Young, 2022, 32, 800-805.	0.8	5
9	Urinary C-peptide and urinary C-peptide creatinine ratio as markers for insulin resistance in obese children and adolescents. Pediatric Research, 2022, 92, 805-809.	2.3	3
10	A novel fibrillin-1 mutation in an egyptian marfan family: A proband showing nephrotic syndrome due to focal segmental glomerulosclerosis. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2017, 28, 141.	0.3	2
11	Multi-Modal Imaging of a Large-sized Right Atrial Appendage Aneurysm in Infancy. Journal of Cardiovascular Imaging, 2022, 30, 92.	0.7	1
12	A novel <scp><i>POU1F1</i></scp> pathogenic variant: Two familial case reports with phenotype expansion. Clinical Genetics, 2021, 100, 641-642.	2.0	0