Ekram M Fateen

List of Publications by Year
in descending order

14	144	1307366 7	1199470 12
papers	citations	h-index	g-index

14	14	14	268
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical, Biochemical, and Molecular Characterization of Metachromatic Leukodystrophy Among Egyptian Pediatric Patients: Expansion of the ARSA Mutational Spectrum. Journal of Molecular Neuroscience, 2021, 71, 1112-1130.	1.1	1
2	Mucopolysaccharidoses diagnosis in the era of enzyme replacement therapy in Egypt. Heliyon, 2021, 7, e07830.	1.4	3
3	Twenty- five years of biochemical diagnosis of Gaucher disease: the Egyptian experience. Heliyon, 2019, 5, e02574.	1.4	10
4	Four novel mutations in the N-acetylgalactosamine-6-sulfate sulfatase gene among Egyptian patients with Morquio A disease. Gene, 2017, 600, 48-54.	1.0	3
5	PEX6 is Expressed in Photoreceptor Cilia and Mutated in Deafblindness with Enamel Dysplasia and Microcephaly. Human Mutation, 2016, 37, 170-174.	1.1	36
6	Prenatal genetic testing, counseling and follow-up of 33 Egyptian pregnant females with history of mucopolysaccharidoses. Egyptian Journal of Medical Human Genetics, 2015, 16, 159-163.	0.5	5
7	Fifteen years experience: Egyptian metabolic lab. Egyptian Journal of Medical Human Genetics, 2014, 15, 379-385.	0.5	7
8	Glycogen storage disease type III in Egyptian children: A single centre clinico-laboratory study. Arab Journal of Gastroenterology, 2014, 15, 63-67.	0.4	9
9	Outcome of enzyme replacement therapy in children with Gaucher disease: The Egyptian experience. Egyptian Journal of Medical Human Genetics, 2011, 12, 9-14.	0.5	19
10	Phenotypical variability in glycogen storage disease type III with a recurrent AGL mutation c.750â€“753delAGAC. Muscle and Nerve, 2011, 43, 451-451.	1.0	3
11	Egyptian glycogen storage disease type III â€“ identification of six novel AGL mutations, including a large 1.5Âkb deletion and a missense mutation p.L620P with subtype IIId. Clinical Chemistry and Laboratory Medicine, 2009, 47, 1233-8.	1.4	10
12	Dyggveâ€“Melchiorâ€“Clausen syndrome: clinical, genetic, and radiological study of 15 Egyptian patients from nine unrelated families. Journal of Children's Orthopaedics, 2009, 3, 451-458.	0.4	19
13	The Effect of Diet on Antioxidant Status in Patients with Galactosemia. Journal of Medical Sciences (Faisalabad, Pakistan), 2006, 6, 452-457.	0.0	2
14	Molecular characterization of Egyptian patients with glycogen storage disease type IIIa. Journal of Human Genetics, 2005, 50, 538-542.	1.1	17