

# Osama K. Zaki

## List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

17  
papers

219  
citations

8  
h-index

14  
g-index

19  
ext. papers

271  
ext. citations

2.1  
avg, IF

2.64  
L-index

#	Paper	IF	Citations
17	Rapid progression and mortality of lysosomal acid lipase deficiency presenting in infants. <i>Genetics in Medicine</i> , <b>2016</b> , 18, 452-8	8.1	42
16	l-Carnitine supplementation improves the behavioral symptoms in autistic children. <i>Research in Autism Spectrum Disorders</i> , <b>2013</b> , 7, 159-166	3	36
15	Genotype-phenotype correlation in 18 Egyptian patients with glutaric acidemia type I. <i>Metabolic Brain Disease</i> , <b>2017</b> , 32, 1417-1426	3.9	35
14	Glutaric aciduria type 1: neuroimaging features with clinical correlation. <i>Pediatric Radiology</i> , <b>2015</b> , 45, 1696-705	2.8	32
13	Cytogenetic study in cases with recurrent abortion in Saudi Arabia. <i>Annals of Saudi Medicine</i> , <b>2000</b> , 20, 233-6	1.6	20
12	A survey of 1,000 cases referred for cytogenetic study to King Khalid University Hospital, Saudi Arabia. <i>Human Heredity</i> , <b>1999</b> , 49, 208-14	1.1	12
11	Structural chromosomal abnormalities in couples with recurrent abortion in Egypt. <i>Turkish Journal of Medical Sciences</i> , <b>2015</b> , 45, 208-13	2.7	11
10	Novel mutation in an Egyptian patient with infantile Canavan disease. <i>Metabolic Brain Disease</i> , <b>2016</b> , 31, 573-7	3.9	8
9	Screening of diseases associated with abnormal metabolites for evaluation of HPLC in organic aciduria profiling. <i>Egyptian Journal of Medical Human Genetics</i> , <b>2014</b> , 15, 69-78	2	5
8	Mutation analysis of methylmalonyl CoA mutase gene exon 2 in Egyptian families: Identification of 25 novel allelic variants. <i>Meta Gene</i> , <b>2015</b> , 3, 71-88	0.7	4
7	Effect of L-carnitine on behavioral disorder in autistic children. <i>Value in Health</i> , <b>2013</b> , 16, A15	3.3	4
6	Demographic and clinical features of glutaric acidemia type 1; a high frequency among isolates in Upper Egypt. <i>Egyptian Journal of Medical Human Genetics</i> , <b>2014</b> , 15, 187-192	2	2
5	A clinical study of mentally retarded children with fragile X syndrome in Saudi Arabia. <i>Annals of Saudi Medicine</i> , <b>2000</b> , 20, 16-9	1.6	1
4	Challenges in diagnosis and counseling of a family with two recessive neurometabolic disorders. <i>Egyptian Journal of Medical Human Genetics</i> , <b>2016</b> , 17, 247-250	2	
3	Environmental risk factors in isolated limb reduction defects. <i>Middle East Journal of Medical Genetics</i> , <b>2014</b> , 3, 53-64		
2	Genotoxicity of cadmium in rat lung cells assessed by an alkaline comet assay and the possible protective role of selenium. <i>Egyptian Journal of Histology</i> , <b>2012</b> , 35, 853-861	0.8	
1	Deletion of short arm of chromosome 3 in a Saudi girl. <i>Annals of Saudi Medicine</i> , <b>1996</b> , 16, 455-7	1.6	

