

Xiaomei Lu

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/217146/publications.pdf>

Version: 2024-02-01

10
papers

44
citations

1937685

4
h-index

1872680

6
g-index

10
all docs

10
docs citations

10
times ranked

69
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular epidemiological and hematological profile of thalassemia in the Dongguan Region of Guangdong Province, Southern China. <i>Journal of Clinical Laboratory Analysis</i> , 2021, 35, e23596.	2.1	16
2	Prevalence of S gene mutations within the major hydrophilic region of hepatitis B virus in patients in Dongguan, southern China. <i>Archives of Virology</i> , 2017, 162, 2949-2957.	2.1	5
3	Mutations within the major hydrophilic region (MHR) of Hepatitis B virus from individuals with simultaneous HBsAg and anti-HBs in Guangzhou, Southern China. <i>Journal of Medical Virology</i> , 2018, 90, 1337-1342.	5.0	5
4	A novel variant in the CDH23 gene is associated with non-syndromic hearing loss in a Chinese family. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2018, 104, 108-112.	1.0	5
5	A novel missense mutation in the SLC26A4 gene causes nonsyndromic hearing loss and enlarged vestibular aqueduct. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2017, 95, 104-108.	1.0	3
6	A novel nonsense mutation of <i>ZEB2</i> gene in a Chinese patient with Mowat-Wilson syndrome. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23413.	2.1	3
7	A Novel β -Thalassemia Nonsense Mutation on the β -Globin Gene: <i>HBA2</i> : c.184A>T. <i>Hemoglobin</i> , 2017, 41, 306-307.	0.8	2
8	A novel mutation in the SLC26A4 gene in a Chinese family with non-syndromic hearing loss and enlarged vestibular aqueduct. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2018, 107, 97-100.	1.0	2
9	A molecular beacon-based asymmetric PCR assay for detecting polymorphisms related to folate metabolism. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23337.	2.1	2
10	Coinheritance of β - and β -Thalassemia with a Novel Mutation (<i>HBB</i> : Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 382 Td (c.268_281de	0.8	1