Amali Mallawaarachchi

List of Publications by Year
in descending order

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9 papers	232 citations	1478505 6 h-index	1720034 7 g-index
9 all docs	9 docs citations	9 times ranked	515 citing authors

#	Article	IF	Citations
1	Massively parallel sequencing and targeted exomes in familial kidney disease can diagnose underlyingÂgenetic disorders. Kidney International, 2017, 92, 1493-1506.	5.2	74
2	Nemaline myopathy and distal arthrogryposis associated with an autosomal recessiveÂTNNT3Âsplice variant. Human Mutation, 2018, 39, 383-388.	2.5	48
3	Standardized practices for RNA diagnostics using clinically accessible specimens reclassifies 75% of putative splicing variants. Genetics in Medicine, 2022, 24, 130-145.	2.4	45
4	Attitudes and Practices of Australian Nephrologists Toward Implementation of Clinical Genomics. Kidney International Reports, 2021, 6, 272-283.	0.8	28
5	Renal genetics in Australia: Kidney medicine in the genomic age. Nephrology, 2019, 24, 279-286.	1.6	18
6	Australia and New Zealand renal gene panel testing in routine clinical practice of 542 families. Npj Genomic Medicine, 2021, 6, 20.	3.8	11
7	Comprehensive evaluation of a prospective Australian patient cohort with suspected genetic kidney disease undergoing clinical genomic testing: a study protocol. BMJ Open, 2019, 9, e029541.	1.9	6
8	The HIDDEN Protocol: An Australian Prospective Cohort Study to Determine the Utility of Whole Genome Sequencing in Kidney Failure of Unknown Aetiology. Frontiers in Medicine, 0, 9, .	2.6	2
9	Novel complement factor H gene mutation causing atypical haemolytic uraemic syndrome: early Eculizumab prevents acute dialysis. CKJ: Clinical Kidney Journal, 2017, 10, 263-265.	2.9	0