## Hind Alsharhan

## List of Publications by Year in descending order

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2258059 2053705 6 24 3 5 citations h-index g-index papers 6 6 6 37 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	<i>ALG13</i> Xâ€linked intellectual disability: New variants, glycosylation analysis, and expanded phenotypes. Journal of Inherited Metabolic Disease, 2021, 44, 1001-1012.	3.6	9
2	Disorders of phenylalanine and tyrosine metabolism. Translational Science of Rare Diseases, 2020, 5, 3-58.	1.5	4
3	Expanding the phenotype, genotype and biochemical knowledge of <scp>ALG3â€CDG</scp> . Journal of Inherited Metabolic Disease, 2021, 44, 987-1000.	3.6	4
4	Early Diagnosis of Classic Homocystinuria in Kuwait through Newborn Screening: A 6-Year Experience. International Journal of Neonatal Screening, 2021, 7, 56.	<b>3.</b> 2	4
5	Urinary Uracil: A Useful Marker for Ornithine Transcarbamylase Deficiency in Affected Males. Clinical Chemistry, 2020, 66, 988-989.	3.2	3
6	COXPD9 in an individual from Puerto Rico and literature review. American Journal of Medical Genetics, Part A, 2021, 185, 2519-2525.	1.2	0