

# Noreen Karim

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5124049/publications.pdf>

Version: 2024-02-01

16  
papers

157  
citations

1163117

8  
h-index

1199594

12  
g-index

16  
all docs

16  
docs citations

16  
times ranked

205  
citing authors

#	ARTICLE	IF	CITATIONS
1	Human stratum corneum proteomics reveals cross-linking of a broad spectrum of proteins in cornified envelopes. <i>Experimental Dermatology</i> , 2019, 28, 618-622.	2.9	27
2	A Novel Missense Mutation in the CLPP Gene Causing Perrault Syndrome Type 3 in a Turkish Family. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2016, 8, 472-477.	0.9	19
3	Proteomic genotyping of fingermark donors with genetically variant peptides. <i>Forensic Science International: Genetics</i> , 2019, 42, 21-30.	3.1	18
4	Relationship between the molecular composition, visible light absorption, and health-related properties of smoldering woodsmoke aerosols. <i>Atmospheric Chemistry and Physics</i> , 2020, 20, 539-559.	4.9	18
5	Optimal processing for proteomic genotyping of single human hairs. <i>Forensic Science International: Genetics</i> , 2020, 47, 102314.	3.1	17
6	Age-Related Changes in Hair Shaft Protein Profiling and Genetically Variant Peptides. <i>Forensic Science International: Genetics</i> , 2020, 47, 102309.	3.1	13
7	Proteomic genotyping: Using mass spectrometry to infer SNP genotypes in pigmented and non-pigmented hair. <i>Forensic Science International</i> , 2020, 310, 110200.	2.2	13
8	Proteomic manifestations of genetic defects in autosomal recessive congenital ichthyosis. <i>Journal of Proteomics</i> , 2019, 201, 104-109.	2.4	10
9	Novel TGM1 mutation in a Pakistani family affected with severe lamellar ichthyosis. <i>Pediatrics and Neonatology</i> , 2018, 59, 628-629.	0.9	7
10	Elucidation of familial relationships using hair shaft proteomics. <i>Forensic Science International: Genetics</i> , 2021, 54, 102564.	3.1	6
11	Proteomic genotyping: Using mass spectrometry to infer SNP genotypes in a forensic context. <i>Forensic Science International: Genetics Supplement Series</i> , 2019, 7, 664-666.	0.3	5
12	Molecular Genetic Study of a Large Inbred Pakistani Family Affected with Autosomal Recessive Congenital Ichthyosis Through Whole Exome Sequencing. <i>Genetic Testing and Molecular Biomarkers</i> , 2019, 23, 428-432.	0.7	4
13	Co-Occurrence of Autosomal Recessive Lamellar Ichthyosis and X-Linked Recessive Ichthyosis in a Consanguineous Pakistani Family. <i>Annals of Dermatology</i> , 2019, 31, 581.	0.9	0
14	Identification of two novel variants in GNPTAB underlying mucopolidosis II in a Pakistani family. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2020, 33, 647-651.	0.9	0
15	Epidermal cell cultures from white and green sturgeon ( <i>Acipenser transmontanus</i> and <i>medirostris</i> ): Expression of TGM1-like transglutaminases and CYP4501A. <i>PLoS ONE</i> , 2022, 17, e0265218.	2.5	0
16	Identification of Frameshift Variants in POLH Gene Causing Xeroderma Pigmentosum in Two Consanguineous Pakistani Families. <i>Genes</i> , 2022, 13, 543.	2.4	0