Noreen Karim

List of Publications by Year
in descending order

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

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		1163117	1199594
16	157	8	12
papers	citations	h-index	g-index

16	16	16	205
all docs	docs citations	times ranked	citing authors

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