

Rabia Habib

List of Publications by Year in descending order

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15
papers

206
citations

1040056

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1058476

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times ranked

328
citing authors

#	ARTICLE	IF	CITATIONS
1	Chronic Exposure to Organophosphates Pesticides and Risk of Metabolic Disorder in Cohort from Pakistan and Cameroon. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 2310.	2.6	23
2	Analysis of PON1 gene polymorphisms (rs662 and rs854560) and inflammatory markers in organophosphate pesticides exposed cohorts from two distinct populations.. <i>Environmental Research</i> , 2020, 191, 110210.	7.5	11
3	Oxidative Stress and Analysis of Selected SNPs of ACHE (rs 2571598), BCHE (rs 3495), CAT (rs 7943316), SIRT1 (rs 10823108), GSTP1 (rs 1695), and Gene GSTM1, GSTT1 in Chronic Organophosphates Exposed Groups from Cameroon and Pakistan. <i>International Journal of Molecular Sciences</i> , 2020, 21, 6432.	4.1	7
4	Mixture of Organophosphates Chronic Exposure and Pancreatic Dysregulations in Two Different Population Samples. <i>Frontiers in Public Health</i> , 2020, 8, 534902.	2.7	10
5	Cannabis Constituents and Acetylcholinesterase Interaction: Molecular Docking, In Vitro Studies and Association with CNR1 rs806368 and ACHE rs17228602. <i>Biomolecules</i> , 2020, 10, 758.	4.0	18
6	Biochemical Analysis and Association of Butyrylcholinesterase SNPs rs3495 and rs1803274 with Substance Abuse Disorder. <i>Journal of Molecular Neuroscience</i> , 2019, 67, 445-455.	2.3	3
7	Association of status of acetylcholinesterase and ACHE gene 3â€™ UTR variants (rs17228602, rs17228616) with drug addiction vulnerability in pakistani population. <i>Chemico-Biological Interactions</i> , 2019, 308, 130-136.	4.0	6
8	Decoding Common Features of Neurodegenerative Disorders: From Differentially Expressed Genes to Pathways. <i>Current Genomics</i> , 2018, 19, 300-312.	1.6	21
9	Protein Quantity and Quality of Safflower Seed Improved by NP Fertilizer and Rhizobacteria (<i>Azospirillum</i> and <i>Azotobacter</i> spp.). <i>Frontiers in Plant Science</i> , 2016, 7, 104.	3.6	38
10	Expansion of the spectrum of ITGB6-related disorders to adolescent alopecia, dentogingival abnormalities and intellectual disability. <i>European Journal of Human Genetics</i> , 2016, 24, 1223-1227.	2.8	20
11	A Novel Locus for Ectodermal Dysplasia of Hair, Nail and Skin Pigmentation Anomalies Maps to Chromosome 18p11.32-p11.31. <i>PLoS ONE</i> , 2015, 10, e0129811.	2.5	2
12	A nonsense mutation in the gene ROR2 underlying autosomal dominant brachydactyly type B. <i>Clinical Dysmorphology</i> , 2013, 22, 47-50.	0.3	5
13	A novel splice site mutation in gene C2orf37 underlying Woodhouseâ€™Sakati syndrome (WSS) in a consanguineous family of Pakistani origin. <i>Gene</i> , 2011, 490, 26-31.	2.2	22
14	DFNB89, a novel autosomal recessive nonsyndromic hearing impairment locus on chromosome 16q21-q23.2. <i>Human Genetics</i> , 2011, 129, 379-385.	3.8	11
15	Mutations in Lipase H Gene Underlie Autosomal Recessive Hypotrichosis in Five Pakistani Families. <i>Acta Dermato-Venereologica</i> , 2010, 90, 93-94.	1.3	9