Elisabetta Bolognesi

List of Publications by Year
in descending order

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30 papers	718 citations	430874 18 h-index	552781 26 g-index
30 all docs	30 docs citations	30 times ranked	1111 citing authors

#	Article	IF	Citations
1	HLA Allele Frequencies and Association with Severity of COVID-19 Infection in Northern Italian Patients. Cells, 2022, 11, 1792.	4.1	5
2	Oligomeric Î±-Syn and SNARE complex proteins in peripheral extracellular vesicles of neural origin are biomarkers for Parkinson's disease. Neurobiology of Disease, 2021, 148, 105185.	4.4	62
3	NK Cell Subpopulations and Receptor Expression in Recovering SARS-CoV-2 Infection. Molecular Neurobiology, 2021, 58, 6111-6120.	4.0	10
4	SNAP-25 Single Nucleotide Polymorphisms, Brain Morphology and Intelligence in Children With Borderline Intellectual Functioning: A Mediation Analysis. Frontiers in Neuroscience, 2021, 15, 715048.	2.8	1
5	The VDR FokI (rs2228570) polymorphism is involved in Parkinson's disease. Journal of the Neurological Sciences, 2021, 428, 117606.	0.6	7
6	Intervening on the Developmental Course of Children With Borderline Intellectual Functioning With a Multimodal Intervention: Results From a Randomized Controlled Trial. Frontiers in Psychology, 2020, 11, 679.	2.1	2
7	Immune regulation of neurodevelopment at the motherâ€“foetus interface: the case of autism. Clinical and Translational Immunology, 2020, 9, e1211.	3.8	24
8	Vitamin D Receptor Polymorphisms Associated with Autism Spectrum Disorder. Autism Research, 2020, 13, 680-690.	3.8	22
9	Serum miRNAs Expression and SNAP-25 Genotype in Alzheimerâ€™s Disease. Frontiers in Aging Neuroscience, 2019, 11, 52.	3.4	19
10	HLA-G allelic distribution in Sardinian children with Autism spectrum disorders: A replication study. Brain, Behavior, and Immunity, 2019, 79, 314-318.	4.1	9
11	The Syntaxin-1A gene single nucleotide polymorphism rs4717806 associates with the risk of ischemic heart disease. Medicine (United States), 2019, 98, e15846.	1.0	2
12	HLA-Gâ^—14bp Insertion and the KIR2DS1-HLAC2 Complex Impact on Behavioral Impairment in Children with Autism Spectrum Disorders. Neuroscience, 2018, 370, 163-169.	2.3	13
13	HLA-G coding region polymorphism is skewed in autistic spectrum disorders. Brain, Behavior, and Immunity, 2018, 67, 308-313.	4.1	21
14	Vitamin D-binding protein gene polymorphisms are not associated with MS risk in an Italian cohort. Journal of Neuroimmunology, 2017, 305, 92-95.	2.3	15
15	ApoE and SNAP-25 Polymorphisms Predict the Outcome of Multidimensional Stimulation Therapy Rehabilitation in Alzheimerâ€™s Disease. Neurorehabilitation and Neural Repair, 2016, 30, 883-893.	2.9	19
16	Association between SNAP-25 gene polymorphisms and cognition in autism: functional consequences and potential therapeutic strategies. Translational Psychiatry, 2015, 5, e500-e500.	4.8	76
17	An HLA-Gâ^—14bp insertion/deletion polymorphism associates with the development of autistic spectrum disorders. Brain, Behavior, and Immunity, 2015, 44, 207-212.	4.1	32
18	Genetic adaptation of the human circadian clock to day-length latitudinal variations and relevance for affective disorders. Genome Biology, 2014, 15, 499.	8.8	28

#	Article	IF	Citations
19	Activating KIR molecules and their cognate ligands prevail in children with a diagnosis of ASD and in their mothers. Brain, Behavior, and Immunity, 2014, 36, 54-60.	4.1	28
20	Possible Association between SNAP-25 Single Nucleotide Polymorphisms and Alterations of Categorical Fluency and Functional MRI Parameters in Alzheimer's Disease. Journal of Alzheimer's Disease, 2014, 42, 1015-1028.	2.6	31
21	SNAP-25 single nucleotide polymorphisms are associated with hyperactivity in autism spectrum disorders. Pharmacological Research, 2011, 64, 283-288.	7.1	54
22	Vitamin D receptor (VDR) gene SNPs influence VDR expression and modulate protection from multiple sclerosis in HLA-DRB1*15-positive individuals. Brain, Behavior, and Immunity, 2011, 25, 1460-1467.	4.1	73
23	KIR-HLA Genotypes in HIV-Infected Patients Lacking Immunological Recovery despite Effective Antiretroviral Therapy. PLoS ONE, 2011, 6, e27349.	2.5	22
24	HLA polymorphisms in Italian children with autism spectrum disorders: Results of a family based linkage study. Journal of Neuroimmunology, 2011, 230, 135-142.	2.3	25
25	An Evolutionary Analysis of RAC2 Identifies Haplotypes Associated with Human Autoimmune Diseases. Molecular Biology and Evolution, 2011, 28, 3319-3329.	8.9	19
26	Family-based transmission analysis of HLA genetic markers in Sardinian children with autistic spectrum disorders. Human Immunology, 2009, 70, 184-190.	2.4	27
27	Neuropsycological gender differences in healthy individuals and in pediatric neurodevelopmental disorders. A role for SNAP-25. Medical Hypotheses, 2009, 73, 978-980.	1.5	16
28	A collaborative European search for non-DQA1*05-DQB1*02 celiac disease loci on HLA-DR3 haplotypes: analysis of transmission from homozygous parents. Human Immunology, 2003, 64, 350-358.	2.4	27
29	Association study of a new polymorphism in the PECAM-1 gene in multiple sclerosis. Journal of Neuroimmunology, 2000, 104, 174-178.	2.3	19
30	The natural history of an HLA haplotype and its recombinants. Immunogenetics, 1998, 48, 8-15.	2.4	10