Cristina Cheroni

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2793270/publications.pdf

Version: 2024-02-01

26 papers 1,684 citations

394421 19 h-index 25 g-index

28 all docs 28 docs citations

28 times ranked

2907 citing authors

#	Article	IF	CITATIONS
1	From cohorts to molecules: Adverse impacts of endocrine disrupting mixtures. Science, 2022, 375, eabe8244.	12.6	129
2	CHD8 haploinsufficiency links autism to transient alterations in excitatory and inhibitory trajectories. Cell Reports, 2022, 39, 110615.	6.4	40
3	ADAM10 hyperactivation acts on piccolo to deplete synaptic vesicle stores in Huntington's disease. Human Molecular Genetics, 2021, 30, 1175-1187.	2.9	11
4	A09â€ADAM10 activity at the huntington's disease presynapse. , 2021, , .		1
5	Autism spectrum disorder at the crossroad between genes and environment: contributions, convergences, and interactions in ASD developmental pathophysiology. Molecular Autism, 2020, 11, 69.	4.9	125
6	Modulating eIF6 levels unveils the role of translation in ecdysone biosynthesis during Drosophila development. Developmental Biology, 2019, 455, 100-111.	2.0	6
7	Human Cortical Organoids Expose a Differential Function of GSK3 on Cortical Neurogenesis. Stem Cell Reports, 2019, 13, 847-861.	4.8	48
8	SBDS-Deficient Cells Have an Altered Homeostatic Equilibrium due to Translational Inefficiency Which Explains their Reduced Fitness and Provides a Logical Framework for Intervention. PLoS Genetics, 2017, 13, e1006552.	3.5	31
9	DEPDC5 variants increase fibrosis progression in Europeans with chronic hepatitis C virus infection. Hepatology, 2016, 63, 418-427.	7.3	31
10	Differences in protein quality control correlate with phenotype variability in 2 mouse models of familial amyotrophic lateral sclerosis. Neurobiology of Aging, 2015, 36, 492-504.	3.1	63
11	Hepatitis C Virus Deletion Mutants Are Found in Individuals Chronically Infected with Genotype 1 Hepatitis C Virus in Association with Age, High Viral Load and Liver Inflammatory Activity. PLoS ONE, 2015, 10, e0138546.	2.5	14
12	The Association of Il28b Genotype with the Histological Features of Chronic Hepatitis C Is HCV Genotype Dependent. International Journal of Molecular Sciences, 2014, 15, 7213-7224.	4.1	19
13	Genome-Wide Analysis of DNA Methylation, Copy Number Variation, and Gene Expression in Monozygotic Twins Discordant for Primary Biliary Cirrhosis. Frontiers in Immunology, 2014, 5, 128.	4.8	57
14	Interleukin 28B Genotype and Insulin Resistance in Chronic Hepatitis C Patients. Antiviral Therapy, 2014, 19, 747-753.	1.0	6
15	Interaction between PNPLA3 I148M Variant and Age at Infection in Determining Fibrosis Progression in Chronic Hepatitis C. PLoS ONE, 2014, 9, e106022.	2.5	9
16	IL28B polymorphisms predict interferon-related hepatitis B surface antigen seroclearance in genotype D hepatitis B e antigen-negative patients with chronic hepatitis B. Hepatology, 2013, 57, 890-896.	7.3	153
17	Cirrhosis and Rapid Virological Response to Peginterferon Plus Ribavirin Determine Treatment Outcome in HCV-1 IL28B rs12979860 CC Patients. BioMed Research International, 2013, 2013, 1-6.	1.9	3
18	Intracellular Modulation, Extracellular Disposal and Serum Increase of MiR-150 Mark Lymphocyte Activation. PLoS ONE, 2013, 8, e75348.	2.5	66

#	Article	IF	CITATION
19	Identification of New Autoantigens by Protein Array Indicates a Role for IL4 Neutralization in Autoimmune Hepatitis. Molecular and Cellular Proteomics, 2012, 11, 1885-1897.	3.8	38
20	Dysfunction of constitutive and inducible ubiquitin-proteasome system in amyotrophic lateral sclerosis: Implication for protein aggregation and immune response. Progress in Neurobiology, 2012, 97, 101-126.	5.7	129
21	Interleukin 28B polymorphism predicts pegylated interferon plus ribavirin treatment outcome in chronic hepatitis C genotype 4. Hepatology, 2012, 55, 336-342.	7.3	81
22	Genetic variation in the <i>interleukin</i> - <i>28B</i> gene is not associated with fibrosis progression in patients with chronic hepatitis C and known date of infection. Hepatology, 2011, 54, 1127-1134.	7. 3	115
23	Characterization of Detergent-Insoluble Proteins in ALS Indicates a Causal Link between Nitrative Stress and Aggregation in Pathogenesis. PLoS ONE, 2009, 4, e8130.	2.5	101
24	Functional alterations of the ubiquitin-proteasome system in motor neurons of a mouse model of familial amyotrophic lateral sclerosisâ€. Human Molecular Genetics, 2009, 18, 82-96.	2.9	146
25	Insoluble Mutant SOD1 Is Partly Oligoubiquitinated in Amyotrophic Lateral Sclerosis Mice. Journal of Biological Chemistry, 2006, 281, 33325-33335.	3.4	86
26	Protein Nitration in a Mouse Model of Familial Amyotrophic Lateral Sclerosis. Journal of Biological Chemistry, 2005, 280, 16295-16304.	3.4	168