

# Ilaria Musante

## List of Publications by Year in descending order

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Version: 2024-02-01

21  
papers

536  
citations

623734

14  
h-index

713466

21  
g-index

21  
all docs

21  
docs citations

21  
times ranked

946  
citing authors

#	ARTICLE	IF	CITATIONS
1	Loss of Neuron Navigator 2 Impairs Brain and Cerebellar Development. <i>Cerebellum</i> , 2022, , 1.	2.5	5
2	A Phenotypic-Driven Approach for the Diagnosis of WOREE Syndrome. <i>Frontiers in Pediatrics</i> , 2022, 10, 847549.	1.9	3
3	Comprehensive Analysis of Combinatorial Pharmacological Treatments to Correct Nonsense Mutations in the CFTR Gene. <i>International Journal of Molecular Sciences</i> , 2021, 22, 11972.	4.1	21
4	Ionocytes and CFTR Chloride Channel Expression in Normal and Cystic Fibrosis Nasal and Bronchial Epithelial Cells. <i>Cells</i> , 2020, 9, 2090.	4.1	44
5	Targeting Alternative Splicing as a Potential Therapy for Episodic Ataxia Type 2. <i>Biomedicines</i> , 2020, 8, 332.	3.2	13
6	Brain Organoids as Model Systems for Genetic Neurodevelopmental Disorders. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 590119.	3.7	31
7	Furocoumarins as multi-target agents in the treatment of cystic fibrosis. <i>European Journal of Medicinal Chemistry</i> , 2019, 180, 283-290.	5.5	18
8	Two CFTR mutations within codon 970 differently impact on the chloride channel functionality. <i>Human Mutation</i> , 2019, 40, 742-748.	2.5	33
9	Peripheral localization of the epithelial sodium channel in the apical membrane of bronchial epithelial cells. <i>Experimental Physiology</i> , 2019, 104, 866-875.	2.0	11
10	The Autophagy Inhibitor Spautin-1 Antagonizes Rescue of Mutant CFTR Through an Autophagy-Independent and USP13-Mediated Mechanism. <i>Frontiers in Pharmacology</i> , 2018, 9, 1464.	3.5	15
11	Combination potentiator (â€œco-potentiatorâ€™™) therapy for CF caused by CFTR mutants, including N1303K, that are poorly responsive to single potentiators. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 595-606.	0.7	48
12	Genetic inactivation of mGlu5 receptor improves motor coordination in the Grm1 mouse model of SCAR13 ataxia. <i>Neurobiology of Disease</i> , 2018, 109, 44-53.	4.4	15
13	Increased expression of ATP12A proton pump in cystic fibrosis airways. <i>JCI Insight</i> , 2018, 3, .	5.0	43
14	In-vivo effects of knocking-down metabotropic glutamate receptor 5 in the SOD1 mouse model of amyotrophic lateral sclerosis. <i>Neuropharmacology</i> , 2017, 123, 433-445.	4.1	30
15	Phenotypic characterization of Grm1 crv4 mice reveals a functional role for the type 1 metabotropic glutamate receptor in bone mineralization. <i>Bone</i> , 2017, 94, 114-123.	2.9	4
16	Assessment of copy number variations in 120 patients with Poland syndrome. <i>BMC Medical Genetics</i> , 2016, 17, 89.	2.1	20
17	Intermolecular Interactions in the TMEM16A Dimer Controlling Channel Activity. <i>Scientific Reports</i> , 2016, 6, 38788.	3.3	11
18	Goblet Cell Hyperplasia Requires High Bicarbonate Transport To Support Mucin Release. <i>Scientific Reports</i> , 2016, 6, 36016.	3.3	75

#	ARTICLE	IF	CITATIONS
19	De novo deletion of chromosome 11q12.3 in monozygotic twins affected by Poland Syndrome. BMC Medical Genetics, 2014, 15, 63.	2.1	32
20	Knocking down metabotropic glutamate receptor 1 improves survival and disease progression in the SOD1G93A mouse model of amyotrophic lateral sclerosis. Neurobiology of Disease, 2014, 64, 48-59.	4.4	42
21	Compensatory Molecular and Functional Mechanisms in Nervous System of the Grm1crv4 Mouse Lacking the mGlu1 Receptor: A Model for Motor Coordination Deficits. Cerebral Cortex, 2013, 23, 2179-2189.	2.9	22