Magali Taulan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7209607/publications.pdf

Version: 2024-02-01

26 papers	775 citations	759233 12 h-index	610901 24 g-index
26 all docs	26 docs citations	26 times ranked	1109 citing authors

#	Article	IF	Citations
1	Oxytocin and Vasopressin V1a and V2 Receptors Form Constitutive Homo- and Heterodimers during Biosynthesis. Molecular Endocrinology, 2003, 17, 677-691.	3.7	296
2	Transcription factors and miRNAs that regulate fetal to adultCFTRexpression change are new targets for cystic fibrosis. European Respiratory Journal, 2015, 45, 116-128.	6.7	65
3	Renal Toxicogenomic Response to Chronic Uranyl Nitrate Insult in Mice. Environmental Health Perspectives, 2004, 112, 1628-1635.	6.0	54
4	Comprehensive analysis of the renal transcriptional response to acute uranyl nitrate exposure. BMC Genomics, 2006, 7, 2.	2.8	49
5	Large genomic rearrangements in the CFTRgene contribute to CBAVD. BMC Medical Genetics, 2007, 8, 22.	2.1	42
6	Small-scale high-throughput sequencingâ€“based identification of new therapeutic tools in cystic fibrosis. Genetics in Medicine, 2015, 17, 796-806.	2.4	31
7	NF-E2-related factor 2, a key inducer of antioxidant defenses, negatively regulates the CFTR transcription. Cellular and Molecular Life Sciences, 2010, 67, 2297-2309.	5.4	27
8	Binding of serum response factor to cystic fibrosis transmembrane conductance regulator CArG-like elements, as a new potential CFTR transcriptional regulation pathway. Nucleic Acids Research, 2005, 33, 5271-5290.	14.5	23
9	A Classification Model Relative to Splicing for Variants of Unknown Clinical Significance: Application to theCFTRGene. Human Mutation, 2013, 34, 774-784.	2.5	23
10	Targeted RNA-Seq profiling of splicing pattern in the DMD gene: exons are mostly constitutively spliced in human skeletal muscle. Scientific Reports, 2017, 7, 39094.	3.3	23
11	Variants in CFTR untranslated regions are associated with congenital bilateral absence of the vas deferens. Journal of Medical Genetics, 2011, 48, 152-159.	3.2	22
12	Should diffuse bronchiectasis still be considered a CFTR-related disorder?. Journal of Cystic Fibrosis, 2015, 14, 646-653.	0.7	20
13	The HDAC inhibitor SAHA does not rescue CFTR membrane expression in Cystic Fibrosis. International Journal of Biochemistry and Cell Biology, 2017, 88, 124-132.	2.8	13
14	CCSP G38A polymorphism environment interactions regulate CCSP levels differentially in COPD. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2016, 311, L696-L703.	2.9	11
15	A novel double deletion underscores the importance of characterizing end points of the CFTR large rearrangements. European Journal of Human Genetics, 2009, 17, 1683-1687.	2.8	10
16	Current and future molecular approaches in the diagnosis of cystic fibrosis. Expert Review of Respiratory Medicine, 2018, 12, 415-426.	2.5	10
17	Identification of a novel duplication CFTRdup2 and functional impact of large rearrangements identified in the CFTR gene. Gene, 2012, 500, 194-198.	2.2	9
18	Phosphorylated C/EBPÎ² Influences a Complex Network Involving YY1 and USF2 in Lung Epithelial Cells. PLoS ONE, 2013, 8, e60211.	2.5	9

#	Article	IF	Citations
19	miRNA repertoires of cystic fibrosis ex vivo models highlight miRâ€181a and miR â€101 that regulate WISP1 expression. Journal of Pathology, 2021, 253, 186-197.	4.5	7
20	Functional analysis of a promoter variant identified in the CFTR gene in cis of a frameshift mutation. European Journal of Human Genetics, 2012, 20, 180-184.	2.8	6
21	The CYSMA web server: An example of integrative tool for in silico analysis of missense variants identified in Mendelian disorders. Human Mutation, 2020, 41, 375-386.	2.5	6
22	Splicing mutations in the CFTR gene as therapeutic targets. Gene Therapy, 2022, 29, 399-406.	4.5	6
23	Lethal factor VII deficiency due to novel mutations in the F7 promoter: Functional analysis reveals disruption of HNF4 binding site. Thrombosis and Haemostasis, 2012, 108, 277-283.	3.4	5
24	Exon identity influences splicing induced by exonic variants and in silico prediction efficacy. Journal of Cystic Fibrosis, 2021, 20, 464-472.	0.7	5
25	Role of Non-coding RNAs in Cystic Fibrosis. , 2015, , .		2
26	New Molecular Diagnosis Approaches â€" From the Identification of Mutations to their Characterization. , 2015, , .		1