

Anna Tamanini

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5949726/publications.pdf>

Version: 2024-02-01

49
papers

1,675
citations

236925

25
h-index

289244

40
g-index

50
all docs

50
docs citations

50
times ranked

2471
citing authors

#	ARTICLE	IF	CITATIONS
1	Heparan Sulfate Glycosaminoglycans Are Involved in Adenovirus Type 5 and 2-Host Cell Interactions. <i>Virology</i> , 2000, 268, 382-390.	2.4	236
2	A microRNA signature from serum exosomes of patients with glioma as complementary diagnostic biomarker. <i>Journal of Neuro-Oncology</i> , 2018, 136, 51-62.	2.9	125
3	Pyrogallol, an active compound from the medicinal plant <i>Emblica officinalis</i> , regulates expression of pro-inflammatory genes in bronchial epithelial cells. <i>International Immunopharmacology</i> , 2008, 8, 1672-1680.	3.8	87
4	Mapping the Transcriptional Machinery of the IL-8 Gene in Human Bronchial Epithelial Cells. <i>Journal of Immunology</i> , 2011, 187, 6069-6081.	0.8	84
5	Association Between Carrier Screening and Incidence of Cystic Fibrosis. <i>JAMA - Journal of the American Medical Association</i> , 2009, 302, 2573.	7.4	82
6	Interaction of Adenovirus Type 5 Fiber with the Coxsackievirus and Adenovirus Receptor Activates Inflammatory Response in Human Respiratory Cells. <i>Journal of Virology</i> , 2006, 80, 11241-11254.	3.4	77
7	Activation of NF- κ B mediates ICAM-1 induction in respiratory cells exposed to an adenovirus-derived vector. <i>Gene Therapy</i> , 2001, 8, 1436-1442.	4.5	71
8	A Peptide Nucleic Acid against MicroRNA miR-145-5p Enhances the Expression of the Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) in Calu-3 Cells. <i>Molecules</i> , 2018, 23, 71.	3.8	43
9	MAP kinases and NF- κ B collaborate to induce ICAM-1 gene expression in the early phase of adenovirus infection. <i>Virology</i> , 2003, 307, 228-242.	2.4	41
10	Decoy oligodeoxyribonucleotides and peptide nucleic acids as DNA chimeras targeting nuclear factor kappa-B: Inhibition of IL-8 gene expression in cystic fibrosis cells infected with <i>Pseudomonas aeruginosa</i> . <i>Biochemical Pharmacology</i> , 2010, 80, 1887-1894.	4.4	41
11	Molecular basis of cystic fibrosis: from bench to bedside. <i>Annals of Translational Medicine</i> , 2018, 6, 334-334.	1.7	36
12	Early diagnosis from newborn screening maximises survival in severe cystic fibrosis. <i>ERJ Open Research</i> , 2018, 4, 00109-2017.	2.6	35
13	Phospholipase C- β 3 Is a Key Modulator of IL-8 Expression in Cystic Fibrosis Bronchial Epithelial Cells. <i>Journal of Immunology</i> , 2011, 186, 4946-4958.	0.8	34
14	Trimethylangelicin reduces IL-8 transcription and potentiates CFTR function. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2011, 300, L380-L390.	2.9	34
15	Transient Receptor Potential Ankyrin 1 Channels Modulate Inflammatory Response in Respiratory Cells from Patients with Cystic Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2016, 55, 645-656.	2.9	34
16	Targeting Transcription Factor Activity as a Strategy to Inhibit Pro-Inflammatory Genes Involved in Cystic Fibrosis: Decoy Oligonucleotides and Low-Molecular Weight Compounds. <i>Current Medicinal Chemistry</i> , 2010, 17, 4392-4404.	2.4	32
17	β -Sitosterol Reduces the Expression of Chemotactic Cytokine Genes in Cystic Fibrosis Bronchial Epithelial Cells. <i>Frontiers in Pharmacology</i> , 2017, 8, 236.	3.5	32
18	Cystic fibrosis carrier screening effects on birth prevalence and newborn screening. <i>Genetics in Medicine</i> , 2016, 18, 145-151.	2.4	31

#	ARTICLE	IF	CITATIONS
19	Circulating microRNAs as emerging non-invasive biomarkers for gliomas. <i>Annals of Translational Medicine</i> , 2017, 5, 277-277.	1.7	31
20	Pancreatic phenotype in infants with cystic fibrosis identified by mutation screening. <i>Archives of Disease in Childhood</i> , 2007, 92, 842-846.	1.9	30
21	Use of a Membrane Potential-Sensitive Probe to Assess Biological Expression of the Cystic Fibrosis Transmembrane Conductance Regulator. <i>Human Gene Therapy</i> , 1995, 6, 1275-1283.	2.7	28
22	Molecular Mechanism of Action of Trimethylangelicin Derivatives as CFTR Modulators. <i>Frontiers in Pharmacology</i> , 2018, 9, 719.	3.5	28
23	Effect of modulation of protein kinase C on the cAMP-dependent chloride conductance in T84 cells. <i>FEBS Letters</i> , 1992, 311, 25-28.	2.8	26
24	Unravelling the role of sphingolipids in cystic fibrosis lung disease. <i>Chemistry and Physics of Lipids</i> , 2016, 200, 94-103.	3.2	26
25	Sphingolipids role in the regulation of inflammatory response: From leukocyte biology to bacterial infection. <i>Journal of Leukocyte Biology</i> , 2018, 103, 445-456.	3.3	26
26	Modulation of expression of IL-8 gene in bronchial epithelial cells by 5-methoxypsoralen. <i>International Immunopharmacology</i> , 2009, 9, 1411-1422.	3.8	25
27	Time Trends in Birth Incidence of Cystic Fibrosis in Two European Areas: Data from Newborn Screening Programs. <i>Journal of Pediatrics</i> , 2008, 152, 25-32.	1.8	22
28	Inconclusive Cystic Fibrosis neonatal screening results: long-term psychosocial effects on parents. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2009, 98, 1927-1934.	1.5	21
29	<i>Pseudomonas aeruginosa</i> reduces the expression of CFTR via post-translational modification of NHERF1. <i>Pflügers Archiv European Journal of Physiology</i> , 2014, 466, 2269-2278.	2.8	21
30	cAMP-Dependent protein kinase inhibits the chloride conductance in apical membrane vesicles of human placenta. <i>Journal of Membrane Biology</i> , 1991, 119, 25-32.	2.1	20
31	Synthesis and Therapeutic Applications of Iminosugars in Cystic Fibrosis. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3353.	4.1	20
32	Treatment of human airway epithelial Calu-3 cells with a peptide-nucleic acid (PNA) targeting the microRNA miR-101-3p is associated with increased expression of the cystic fibrosis Transmembrane Conductance Regulator (CFTR) gene. <i>European Journal of Medicinal Chemistry</i> , 2021, 209, 112876.	5.5	18
33	ICAM-1 induction in respiratory cells exposed to a replication-deficient recombinant adenovirus in vitro and in vivo. <i>Gene Therapy</i> , 1998, 5, 131-136.	4.5	17
34	Evidence for the Involvement of Lipid Rafts and Plasma Membrane Sphingolipid Hydrolases in <i>Pseudomonas aeruginosa</i> Infection of Cystic Fibrosis Bronchial Epithelial Cells. <i>Mediators of Inflammation</i> , 2017, 2017, 1-16.	3.0	16
35	Exploring the effect of chirality on the therapeutic potential of N-alkyl-deoxyiminosugars: anti-inflammatory response to <i>Pseudomonas aeruginosa</i> infections for application in CF lung disease. <i>European Journal of Medicinal Chemistry</i> , 2019, 175, 63-71.	5.5	16
36	PLCB3 Loss of Function Reduces <i>Pseudomonas aeruginosa</i> -Dependent IL-8 Release in Cystic Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2018, 59, 428-436.	2.9	15

#	ARTICLE	IF	CITATIONS
37	Protein a, hydroxyapatite and diethylaminoethyl: Evaluation of three procedures for the preparative purification of monoclonal antibodies by high-performance liquid chromatography. <i>Journal of Chromatography A</i> , 1989, 465, 101-111.	3.7	12
38	Protracted neonatal hypertrypsinogenaemia, normal sweat chloride, and cystic fibrosis. <i>Archives of Disease in Childhood</i> , 2000, 82, 481-482.	1.9	12
39	GM1 as Adjuvant of Innovative Therapies for Cystic Fibrosis Disease. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4486.	4.1	11
40	Sphingolipids and plasma membrane hydrolases in human primary bronchial cells during differentiation and their altered patterns in cystic fibrosis. <i>Glycoconjugate Journal</i> , 2020, 37, 623-633.	2.7	10
41	Enhancing the Expression of CFTR Using Antisense Molecules against MicroRNA miR-145-5p. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 1443-1444.	5.6	9
42	A Peptide-Nucleic Acid Targeting miR-335-5p Enhances Expression of Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) Gene with the Possible Involvement of the CFTR Scaffolding Protein NHERF1. <i>Biomedicines</i> , 2021, 9, 117.	3.2	9
43	Chemical conjugation of 125 I-F508-CFTR corrector deoxyspergualin to transporter human serum albumin enhances its ability to rescue Cl $^{-}$ channel functions. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2008, 295, L336-L347.	2.9	8
44	miRNAs in Serum Exosomes for Differential Diagnosis of Brain Metastases. <i>Cancers</i> , 2022, 14, 3493.	3.7	8
45	Alternative splicing of a previously unidentified CFTR exon introduces an in-frame stop codon 5' of the R region. <i>FEBS Letters</i> , 1993, 329, 159-162.	2.8	7
46	Late generation lentiviral vectors: Evaluation of inflammatory potential in human airway epithelial cells. <i>Virus Research</i> , 2009, 144, 8-17.	2.2	7
47	Peptide Nucleic Acids for MicroRNA Targeting. <i>Methods in Molecular Biology</i> , 2020, 2105, 199-215.	0.9	7
48	Sweat chloride and immunoreactive trypsinogen in infants carrying twoCFTRmutations and not affected by cystic fibrosis. <i>Archives of Disease in Childhood</i> , 2017, 102, 644-646.	1.9	6
49	Adenosine 3':5'-Monophosphate-Dependent Protein Kinase from Human Placenta: Characterization of the Catalytic Subunit. <i>Enzyme</i> , 1991, 45, 97-108.	0.7	5