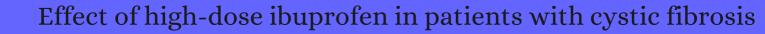
CITATION REPORT List of articles citing



DOI: 10.1056/nejm199503303321303 New England Journal of Medicine, 1995, 332, 848-54.

Source: https://exaly.com/paper-pdf/26408619/citation-report.pdf

Version: 2024-04-28

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
705	SaturdayN children. 1995 , 152, 1-2		7
704	Anti-inflammatory and antipyretic analgesics and drugs used in gout. 1995 , 19, 92-103		
703	News for the practitioner. 1995 , 154, 684-685		
702	Correspondence. 1995 , 19, 320-322		
701	Selected abstracts. 1995 , 19, 320-322		
700	Symposium session summaries. 1995 , 20, 69-177		
699	Cystic fibrosis: pathogenesis, pulmonary infection, and treatment. 1995 , 21, 839-49; quiz 850-1		52
698	Clearance of HIVlessons from newborns. New England Journal of Medicine, 1995, 332, 883-4	59.2	6
69 7	Colon-cancer genes and brain tumors. New England Journal of Medicine, 1995, 332, 884-5	59.2	21
696	Airway inflammation in cystic fibrosis. New England Journal of Medicine, 1995, 332, 886-7	59.2	6
695	Bone marrow transplantation versus chemotherapy in non-HodgkinN lymphoma. <i>New England Journal of Medicine</i> , 1995 , 333, 727; author reply 730	59.2	
694	Transplantation of fetal mesencephalic tissue in ParkinsonN disease. <i>New England Journal of Medicine</i> , 1995 , 333, 730-1	59.2	15
693	Ibuprofen in patients with cystic fibrosis. <i>New England Journal of Medicine</i> , 1995 , 333, 731; author reply 732	59.2	1
692	Serum antibodies to the deleted dystrophin sequence after cardiac transplantation in a patient with BeckerN muscular dystrophy. <i>New England Journal of Medicine</i> , 1995 , 333, 732-3	59.2	18
691	Cost sharing in health insurance. New England Journal of Medicine, 1995, 333, 733-4	59.2	14
690	More on smoking superheroes. New England Journal of Medicine, 1995, 333, 734	59.2	1
689	Drug management of noninfective complications of cystic fibrosis. 1995 , 50, 626-35		5

(1996-1995)

688	Clinical denouement and mutation analysis of patients with cystic fibrosis undergoing liver transplantation for biliary cirrhosis. 1995 , 127, 881-7		62
68 7	Infections in cystic fibrosis. 1995 , 6, 174-181		1
686	Animal studies of cystic fibrosis. 1995 , 1, 336-42		3
685	TREATMENT OF STEROID-RESISTANT ASTHMA. 1996 , 16, 777-796		4
684	Management of pulmonary disease in patients with cystic fibrosis. <i>New England Journal of Medicine</i> , 1996 , 335, 179-88	59.2	412
683	[Mucoviscidosis: anti-inflammatory therapeutic strategies]. 1996 , 3 Suppl 1, 112s-113s		
682	Ibuprofen inhibits pyrogen-dependent expression of VCAM-1 and ICAM-1 on human endothelial cells. 1996 , 58, 2167-81		50
681	Cystic fibrosis-drug therapy. 1996 , 10, 127-34		1
680	Cystic fibrosis. 1996 , 154, 1229-56		778
679	Inflammation in cystic fibrosis. 1996 , 5, 121-43		
678	Linhflammation dans la Mucoviscidose. 1996 , 5, 144-69		1
677	Supplementation with carotenoids corrects increased lipid peroxidation in children with cystic fibrosis. 1996 , 64, 87-93		72
676	Antineutrophil Cytoplasmic Autoantibodies with Specificity Other Than PR3 and MPO (X-ANCA). 1996 , 68-73		2
675	Autoantibodies against bactericidal/permeability-increasing protein in patients with cystic fibrosis. 1996 , 89, 259-65		99
674	Neutrophil proteinases and rhDNase therapy in cystic fibrosis. 1996 , 9, 2193-5		11
673	Liposomal DNA vectors for cystic fibrosis gene therapy. Current applications, limitations, and future directions. 1996 , 19, 73-87		24
672	Bronchiectasis and related disorders. 1996 , 1, 107-14		6
671	Bronchiectasis: a neglected cause of respiratory morbidity and mortality. 1996 , 1, 221-5		37

670	Potential of preventing Pseudomonas aeruginosa lung infections in cystic fibrosis patients: experimental studies in animals. 1996 , 63, 5-42	16
669	Denmark to the rescue. 1996 , 21, 151-2	9
668	Analysis of ibuprofen in serum by capillary electrophoresis. 1996 , 683, 115-8	34
667	Mechanisms of airway inflammation in cystic fibrosis. 1996 , 7, 63-6	4
666	Anti-inflammatory therapy in cystic fibrosis. 1996 , 7, 70-3	1
665	Management of the Child With Cystic Fibrosis. 1996 , 9, 75-90	
664	Relationship between disease severity and inflammatory markers in cystic fibrosis. 1996 , 75, 498-501	29
663	Reduced upper airway nitric oxide in cystic fibrosis. 1996 , 75, 319-22	198
662	Nonsteroidal anti-inflammatory drugs and severe psychiatric side effects. 1996 , 26, 25-34	35
661	Opportunities for the use of aerosolized alpha 1-antitrypsin for the treatment of cystic fibrosis. 1996 , 110, 256S-260S	14
660	Anti-inflammatory and antipyretic analgesics and drugs used in gout. 1997 , 20, 86-102	
659	Controlled trial of inhaled budesonide in patients with cystic fibrosis and chronic bronchopulmonary Psuedomonas aeruginosa infection. 1997 , 156, 1190-6	65
658	Randomised controlled trial of inhaled corticosteroids (fluticasone propionate) in cystic fibrosis. 1997 , 77, 124-30	76
657	Nasal and bronchoalveolar lavage fluid cytokines in early cystic fibrosis. 1997 , 175, 638-47	212
656	Immunologic Aspects of Lung Diseases and Cystic Fibrosis. 1997 , 278, 1924	22
655	New treatments in adult cystic fibrosis. 1997 , 90 Suppl 31, 2-5	O
654	Early inflammation and the development of pulmonary disease in cystic fibrosis. 1997 , 16, 267-8	14
653	[New prospects in cystic fibrosis treatment]. 1997 , 33, 190-5	

652 Concern over ibuprofen/pancreatic enzyme interaction. **1997**, &NA;, 2

651	The biogenesis, traffic, and function of the cystic fibrosis transmembrane conductance regulator. 1997 , 172, 193-241	36
650	The decline and fall of pulmonary function in cystic fibrosis: new models, new lessons. 1997 , 131, 789-90	19
649	Longitudinal analysis of pulmonary function decline in patients with cystic fibrosis. 1997 , 131, 809-14	281
648	Cystic fibrosis gene mutation (deltaF508) is associated with an intrinsic abnormality in Ca2+-induced arachidonic acid release by epithelial cells. 1997 , 16, 749-59	53
647	Cystic fibrosis: a changing clinical perspective. 1997 , 27, 6-11	4
646	Applied pharmacology in pediatric pain management. 1997 , 1, 296-309	1
645	Measurement of plasma ibuprofen by gas chromatography-mass spectrometry. 1997 , 11, 336-9	19
644	Current understanding of the inflammatory process in cystic fibrosis: onset and etiology. 1997 , 24, 137-42; discussion 159-61	242
643	Implications of early inflammation and infection in cystic fibrosis: a review of new and potential interventions. 1997 , 24, 143-5; discussion 159-61	11
642	Indomethacin and pancreatic enzymes synergistically damage intestine of rats. 1998, 43, 2322-32	12
641	Prospects for gene therapy for cystic fibrosis. 1998 , 4, 292-9	16
640	Evidence-based pediatric pulmonary medicine: how can it help?. 1998 , 25, 118-27	6
639	Evidence-based medicine. 1998 , 26, 231-3	
638	Response to evidence-based medicine. 1998 , 26, 232-232	
637	Response to Dr. Marino. 1998 , 26, 232-233	
636	Serum eosinophil cationic protein, eosinophil protein X and eosinophil peroxidase in relation to pulmonary function in cystic fibrosis. 1998 , 28, 241-8	23
635	Cystic fibrosis. 1998 , 351, 277-82	155

634	[Mucoviscidosis: therapeutic strategies are multiplying]. 1998 , 5, 1246-52		3
633	Les explorations fonctionnelles respiratoires dans la mucoviscidose. 1998 , 5, 132s-134s		
632	Relationship between socioeconomic status and disease severity in cystic fibrosis. 1998 , 132, 260-4		76
631	Destruction and loss of bronchial cartilage in cystic fibrosis. 1998 , 29, 65-73		29
630	Basic therapies in cystic fibrosis. Does standard therapy work?. 1998 , 19, 487-504, vi		23
629	Ongoing planned clinical trials investigating the pulmonary management of cystic fibrosis. 1998 , 7, 91-	8	
628	Carrier screening for cystic fibrosis: costs and clinical outcomes. 1998 , 18, 202-12		285
627	The management of young adults with cystic fibrosis: Ngenes, jeans and geniesN1998, 20, 217-25		12
626	Cystic fibrosis: a multiple exocrinopathy caused by dysfunctions in a multifunctional transport protein. 1998 , 104, 576-90		33
625	Clinical pharmacokinetics of ibuprofen. The first 30 years. 1998 , 34, 101-54		304
625 624	Clinical pharmacokinetics of ibuprofen. The first 30 years. 1998 , 34, 101-54 Therapies aimed at airway inflammation in cystic fibrosis. 1998 , 19, 505-13, vi		304
		59.2	28
624	Therapies aimed at airway inflammation in cystic fibrosis. 1998 , 19, 505-13, vi	59.2	28
624	Therapies aimed at airway inflammation in cystic fibrosis. 1998 , 19, 505-13, vi Obesity. <i>New England Journal of Medicine</i> , 1998 , 338, 64; author reply 65 Increasing burden of cardiovascular disease: current knowledge and future directions for research	59.2	28
624 623 622	Therapies aimed at airway inflammation in cystic fibrosis. 1998 , 19, 505-13, vi Obesity. <i>New England Journal of Medicine</i> , 1998 , 338, 64; author reply 65 Increasing burden of cardiovascular disease: current knowledge and future directions for research on risk factors. 1998 , 97, 1095-102 Meta-analyses and large randomized, controlled trials. <i>New England Journal of Medicine</i> , 1998 , 338,		28 3 205
624 623 622	Therapies aimed at airway inflammation in cystic fibrosis. 1998, 19, 505-13, vi Obesity. New England Journal of Medicine, 1998, 338, 64; author reply 65 Increasing burden of cardiovascular disease: current knowledge and future directions for research on risk factors. 1998, 97, 1095-102 Meta-analyses and large randomized, controlled trials. New England Journal of Medicine, 1998, 338, 59; author reply 61-2 Hormone-replacement therapy compared with simvastatin for postmenopausal women with	59.2	28 3 205 28
624623622621620	Therapies aimed at airway inflammation in cystic fibrosis. 1998, 19, 505-13, vi Obesity. New England Journal of Medicine, 1998, 338, 64; author reply 65 Increasing burden of cardiovascular disease: current knowledge and future directions for research on risk factors. 1998, 97, 1095-102 Meta-analyses and large randomized, controlled trials. New England Journal of Medicine, 1998, 338, 59; author reply 61-2 Hormone-replacement therapy compared with simvastatin for postmenopausal women with hypercholesterolemia. New England Journal of Medicine, 1998, 338, 63; author reply 64 Transient renal failure due to simultaneous ibuprofen and aminoglycoside therapy in children with	59.2 59.2	28 3 205 28 4

616	Lung infections. 3. Pseudomonas aeruginosa and other related species. 1998, 53, 213-9	69
615	Allograft colonization and infections with pseudomonas in cystic fibrosis lung transplant recipients. 1998 , 113, 1235-43	77
614	Effect of dirithromycin on Haemophilus influenzae infection of the respiratory mucosa. 1998, 42, 772-8	15
613	Update on clinical trials in the treatment of pulmonary disease in patients with cystic fibrosis. 1999 , 8, 1917-1927	
612	IL-10 attenuates excessive inflammation in chronic Pseudomonas infection in mice. 1999 , 160, 2040-7	108
611	Interest of colchicine for the treatment of cystic fibrosis patients. Preliminary report. 1999 , 8, 13-5	15
610	Inflammation and CFTR: might neutrophils be the key in cystic fibrosis?. 1999, 8, 7-11	33
609	Aspirin and some other nonsteroidal anti-inflammatory drugs inhibit cystic fibrosis transmembrane conductance regulator protein gene expression in T-84 cells. 1999 , 8, 219-27	10
608	Priming of blood neutrophils in children with cystic fibrosis: correlation between functional and phenotypic expression of opsonin receptors before and after platelet-activating factor priming. 1999 , 179, 151-62	26
607	Oral non-steroidal anti-inflammatory drug therapy for cystic fibrosis. 2000 , CD001505	10
606	Quantitation of inflammatory responses to bacteria in young cystic fibrosis and control patients. 1999 , 160, 186-91	310
605	Quantitative and qualitative differences in bronchoalveolar inflammatory cells in Pseudomonas aeruginosa-resistant and -susceptible mice. 1999 , 115, 103-9	34
604	rhDNase therapy for the treatment of cystic fibrosis patients with mild to moderate lung disease. 1999 , 24, 415-26	20
603	Update and Review: Cystic Fibrosis. 1999 , 8, 137-62	4
602	Pyloric channel stricture secondary to high-dose ibuprofen therapy in a patient with cystic fibrosis. 1999 , 33, 693-6	12
601	High-resolution computed tomography of the chest in children with cystic fibrosis: support for use as an outcome surrogate. 1999 , 29, 731-5	113
600	Pharmacokinetics of ibuprofen in patients with cystic fibrosis. 1999 , 19, 340-5	13
599	Ibuprofen: new explanation for an old phenomenon. 1999 , 57, 313-20	63

598	Rheumatic disease and cystic fibrosis. 1999 , 42, 1563-71	24
597	Patterns of medical practice in cystic fibrosis: part II. Use of therapies. Investigators and Coordinators of the Epidemiologic Study of Cystic Fibrosis. 1999 , 28, 248-54	72
596	Direct injection HPLC analysis of some non-steroidal anti-inflammatory drugs on restricted access media columns. 1999 , 13, 51-6	22
595	Abstruse comparisons: the problems of numerical contrasts of two groups. 1999 , 52, 13-8	10
594	The pharmacokinetics of ibuprofen suspension, chewable tablets, and tablets in children with cystic fibrosis. 1999 , 134, 58-63	32
593	Goals of therapy in antibody deficiency syndromes. 1999 , 104, 911-3	10
592	Pulmonary abnormalities in patients with primary hypogammaglobulinemia. 1999 , 104, 1031-6	150
591	Dornase alpha and survival of patients with cystic fibrosis. 1999 , 60, 736-9	2
590	The use of anti-inflammatory medications in cystic fibrosis: trends and physician attitudes. 1999 , 115, 1053-8	79
589	Emerging therapies for cystic fibrosis lung disease. 1999 , 115, 1120-6	38
588	Lymphocyte glutathione levels in children with cystic fibrosis. 1999 , 116, 201-5	40
587	Pharmacokinetics of ibuprofen enantiomers in children with cystic fibrosis. 2000 , 40, 861-8	10
586	Cartilaginous airway wall dimensions and airway resistance in cystic fibrosis lungs. 2000 , 15, 735-42	60
585	Selectivity of cyclo-oxygenase inhibitors in human pulmonary epithelial and smooth muscle cells. 2000 , 15, 751-6	25
584	Prenatal Smoking Increases Risk of Cleft Lip/Palate. 2000 , 3, 61-62	
583	Diagnosis and treatment of cystic fibrosis. 2000 , 67, 239-47	74
582	Re: Rost et al. describing the effects of neck position on endotracheal tube (ETT) location in low birth weight infants. 2000 , 29, 242-4	1
581	The authors respond. 2000 , 29, 243-244	

(2000-2000)

580	2000, 29, 244	13
579	Ibuprofen therapy in cystic fibrosis. 2000 , 29, 244-5	2
578	Response by author. 2000 , 29, 245	
577	Modern statistical techniques for the analysis of longitudinal data in biomedical research. 2000 , 30, 330-44	135
576	Mouse models of chronic lung infection with Pseudomonas aeruginosa: models for the study of cystic fibrosis. 2000 , 30, 413-24	47
575	Activation of human CIC-2 Cl- channels: implications for cystic fibrosis. 2000 , 27, 896-900	10
574	Ibuprofen use to reduce the incidence and severity of bronchopulmonary dysplasia: a pilot study. 2000 , 20, 13-6	21
573	Neutrophils: molecules, functions and pathophysiological aspects. 2000 , 80, 617-53	792
572	PKA and arachidonic acid activation of human recombinant ClC-2 chloride channels. 2000 , 279, C40-50	63
571	Mycobacterium abscessus infection in cystic fibrosis. Colonization or infection?. 2000 , 161, 641-5	135
570	Chest pain in women with normal coronary angiograms. <i>New England Journal of Medicine</i> , 2000 , 342, 885-7	28
569	Growth in children with chronic lung disease. <i>New England Journal of Medicine</i> , 2000 , 342, 887-8 59.2	14
568	Clinical implications of inflammation in young children. 2000 , 162, S11-4	12
567	In vivo lipid peroxidation and platelet activation in cystic fibrosis. 2000 , 162, 1195-201	77
566	Transport of bifunctional proteins across respiratory epithelial cells via the polymeric immunoglobulin receptor. 2000 , 161, 944-51	19
565	Effect of Pseudomonas infection on weight loss, lung mechanics, and cytokines in mice. 2000 , 161, 271-9	91
564	Diagnosis of lung cancer: FOB before CT or CT before FOB?. 2000 , 55, 350-1	2
563	CXC chemokine receptor CXCR2 is essential for protective innate host response in murine Pseudomonas aeruginosa pneumonia. 2000 , 68, 4289-96	233

562	Maintenance treatment of chronic pseudomonas aeruginosa infection in cystic fibrosis. 2000 , 55, 349-50	22
561	Superantigens and cystic fibrosis: resistance of presenting cells to dexamethasone. 2000 , 7, 553-6	7
560	Maternal Anticonvulsant Therapy and Neonatal Blood Coagulation. 2000 , 3, 62-63	
559	Role of bronchial responsiveness testing in asthma prevalence surveys. 2000 , 55, 352-4	30
558	Inflammation in cystic fibrosis and its management. 2000 , 1, 101-6	26
557	Newer therapies for cystic fibrosis. 2000 , 1, 107-13	1
556	A randomized controlled trial of a 3-year home exercise program in cystic fibrosis. 2000 , 136, 304-10	206
555	[Correction of CFTR anomalies: pharmacological approach]. 2000 , 7 Suppl 2, 352s-354s	
554	Statistical analysis of the long-term effects of recombinant human deoxyribonuclease on pulmonary function in cystic fibrosis patients. 2000 , 10, 287-97	1
553	Prospects for the prevention and control of pseudomonal infection in children with cystic fibrosis. 2000 , 2, 451-63	35
552	New and emerging therapies for pulmonary complications of cystic fibrosis. 2001 , 61, 1379-85	8
551	[SEPAR (Spanish Society of Pneumology and Thoracic Surgery) Guidelines. Guideline for diagnosis and treatment of respiratory involvements in cystic fibrosis]. 2001 , 37, 316-24	25
550	Utilizacifi de la alternancia de antipirficos en el tratamiento de la fiebre en Espa li . 2001 , 55, 503-510	6
549	Antipyretics: mechanisms of action and clinical use in fever suppression. 2001 , 111, 304-15	184
548	Defining a pulmonary exacerbation in cystic fibrosis. 2001 , 139, 359-65	219
547	A two-year randomized, placebo-controlled trial of dornase alfa in young patients with cystic fibrosis with mild lung function abnormalities. 2001 , 139, 813-20	235
546	What is the cystic fibrosis clinician supposed to do with human recombinant dornase alfa?. 2001 , 139, 768-70	3
545	[Management of pulmonary involvement in mucoviscidosis in the child]. 2001 , 8 Suppl 5, 856s-883s	1

(2001-2001)

544	Cystic fibrosis and the use of pharmacogenomics to determine surrogate endpoints for drug discovery. 2001 , 1, 223-38	8
543	Risks and benefits of nonsteroidal anti-inflammatory drugs in children: a comparison with paracetamol. 2001 , 3, 817-58	101
542	CFTR and pseudomonas infections in cystic fibrosis. 2001 , 6, d890-897	6
541	Proinflammatory cytokine responses to P. aeruginosa infection in human airway epithelial cell lines. 2001 , 280, L493-502	147
540	CFTR and pseudomonas infections in cystic fibrosis. 2001 , 6, D890-7	17
539	Bacterial infections and inflammation in the lungs of cystic fibrosis patients. 2001 , 20, 207-13	27
538	Inpatient care of the adult with an exacerbation of cystic fibrosis. 2001, 12, 293-304	3
537	Bronchiectasis. 2001 , 14, 193-7	17
536	Inflammation and infection in cystic fibrosishen or egg?. 2001 , 17, 4-5	6
535	Pulmonale Manifestation der Zystischen Fibrose. 2001 , 149, 222-238	4
534	Leukotrienes and cystic fibrosis. 2001 , 1, 175-177	
533	Pseudomonas aeruginosa and the in vitro and in vivo biofilm mode of growth. 2001 , 3, 23-35	303
532	Effect of choice of reference equation on analysis of pulmonary function in cystic fibrosis patients. 2001 , 31, 227-37	47
531	Renal failure and vestibular toxicity in an adolescent with cystic fibrosis receiving gentamicin and standard-dose ibuprofen. 2001 , 31, 314-6	30
530	Gene therapy for cystic fibrosis. 2001 , 3, 409-17	51
529	Inflammation and cystic fibrosis pulmonary disease. 2001 , 21, 593-603	12
528	Interleukin-8 up-regulation by neutrophil elastase is mediated by MyD88/IRAK/TRAF-6 in human bronchial epithelium. 2001 , 276, 35494-9	130
527	P-113D, an antimicrobial peptide active against Pseudomonas aeruginosa, retains activity in the presence of sputum from cystic fibrosis patients. 2001 , 45, 3437-44	79

526	Ursodeoxycholic acid ameliorates ibuprofen-induced enteropathy in the rat. 2001 , 32, 270-3	9
525	Administration of aerosolized antibiotics in cystic fibrosis patients. 2001 , 120, 107S-113S	71
524	Cystic fibrosis. 2001 , 56, 237-41	46
523	Management of Pulmonary Disease in Patients with Cystic Fibrosis. 2001 , 14, 207-227	2
522	Antibacterial activity of apical surface fluid from the human airway cell line Calu-3: pharmacologic alteration by corticosteroids and beta(2)-agonists. 2001 , 25, 196-202	28
521	The association of socioeconomic status with outcomes in cystic fibrosis patients in the United States. 2001 , 163, 1331-7	242
520	Fitness, acute exercise, and anabolic and catabolic mediators in cystic fibrosis. 2001 , 164, 1432-7	64
519	The human ATP-binding cassette transporter genes: from the bench to the bedside. 2001 , 1, 45-65	121
518	The pharmacokinetics of colistin in patients with cystic fibrosis. 2001 , 41, 645-54	110
517	The Pseudomonas autoinducer N-(3-oxododecanoyl) homoserine lactone induces cyclooxygenase-2 and prostaglandin E2 production in human lung fibroblasts: implications for inflammation. 2002 , 169, 2636-42	135
516	Nutrition and growth in cystic fibrosis. 2002 , 58 Suppl 1, 16-20	17
515	Inflammatory response in airway epithelial cells isolated from patients with cystic fibrosis. 2002 , 166, 1248-56	104
514	Pulmonary function, body composition, and protein catabolism in adults with cystic fibrosis. 2002 , 165, 495-500	74
513	Murine models of CF airway infection and inflammation. 2002 , 70, 495-515	7
512	Global genomic analysis of AlgU (sigma(E))-dependent promoters (sigmulon) in Pseudomonas aeruginosa and implications for inflammatory processes in cystic fibrosis. 2002 , 184, 1057-64	90
511	Asthma morbidity after the short-term use of ibuprofen in children. 2002 , 109, E20	121
510	Development of population pharmacokinetic models and optimal sampling times for ibuprofen tablet and suspension formulations in children with cystic fibrosis. 2002 , 24, 315-21	10
509	Potential role of macrolide antibiotics in the management of cystic fibrosis lung disease. 2002 , 8, 521-8	23

(2002-2002)

5	08	Bradykinin increases IL-8 generation in airway epithelial cells via COX-2-derived prostanoids. 2002 , 283, L612-8	31
5	07	Lung infections associated with cystic fibrosis. 2002 , 15, 194-222	1158
5	06	Macrolides in cystic fibrosis: is there a role?. 2002 , 1, 235-41	14
5	05	Fibrosis quatica. Fisiopatologa, gentica, aspectos claicos y terapliticos. 2002 , 37, 1-22	
5	04	An epidemiological investigation of a sustained high rate of pediatric parapneumonic empyema: risk factors and microbiological associations. 2002 , 34, 434-40	292
5	03	Treatment of severe small airways disease in children with cystic fibrosis: alternatives to corticosteroids. 2002 , 4, 381-9	10
5	02	Cystic Fibrosis Methods and Protocols. 2002 ,	1
5	01	Pilot safety study of liposomal prostaglandin (PGE1) in respiratory exacerbations in cystic fibrosis. 2002 , 1, 90-3	3
5	00	Understanding bacterial biofilms in patients with cystic fibrosis: current and innovative approaches to potential therapies. 2002 , 1, 249-54	93
4	.99	Cystic fibrosis: a 2002 update. 2002 , 140, 156-64	76
4	.98	Effects of megestrol acetate on weight gain, body composition, and pulmonary function in patients with cystic fibrosis. 2002 , 140, 439-44	50
4	97	Hyperacidification in cystic fibrosis: links with lung disease and new prospects for treatment. 2002 , 8, 512-9	62
4	.96	A(2) adenosine receptors regulate CFTR through PKA and PLA(2). 2002, 282, L12-25	53
4	.95	Pharmacological approaches for the discovery and development of new anti-inflammatory agents for the treatment of cystic fibrosis. 2002 , 54, 1409-23	64
4	.94	Transplantation for cystic fibrosis: outcome following early liver transplantation. 2002, 17, 208-13	80
4	.93	Jointly modelling the relationship between survival and pulmonary function in cystic fibrosis patients. 2002 , 21, 1271-87	86
4	.92	Evidence-based medicine in cystic fibrosis: how should practice change?. 2002 , 34, 242-7	2
4	.91	Symposium summaries S13.4B18.4. 2002 , 34, 153-181	

490	The role of inflammation in the pathophysiology of CF lung disease. 2002 , 23, 5-27	180
489	Use of modulators of airways inflammation in patients with CF. 2002 , 23, 29-39	3
488	Cystic fibrosis-related diabetes. 2003 , 20, 425-36	67
487	Dornase alfa in the treatment of cystic fibrosis in Europe: a report from the Epidemiologic Registry of Cystic Fibrosis. 2003 , 36, 427-32	36
486	Pathophysiology and management of pulmonary infections in cystic fibrosis. 2003, 168, 918-51	1230
485	Improved glutathione status in young adult patients with cystic fibrosis supplemented with whey protein. 2003 , 2, 195-8	72
484	Cystic fibrosis. 2003 , 361, 681-9	800
483	Low-dose methotrexate for advanced pulmonary disease in patients with cystic fibrosis. 2003 , 97, 498-500	15
482	Growth and nutritional indexes in early life predict pulmonary function in cystic fibrosis. 2003, 142, 624-30	292
481	[Question 1. What is the influence of nutritional status on the progression of cystic fibrosis? Physiopathologic aspects of nutritional disorders in the course of cystic fibrosis]. 2003 , 10 Suppl 3, 413s-420s	1
480	[Anti-inflammatory therapy in cystic fibrosis]. 2003, 10 Suppl 2, 370s-375s	
479	Newer therapies for cystic fibrosis. 2003 , 13, 259-263	3
478	[Complementary therapies in cystic fibrosis: evidence of therapeutic benefits and treatment recommendations]. 2003 , 58, 39-44	3
477	Azithromycin in patients with cystic fibrosis chronically infected with Pseudomonas aeruginosa: a randomized controlled trial. 2003 , 290, 1749-56	668
476	Microarray analysis of global gene expression in mucoid Pseudomonas aeruginosa. 2003, 185, 1071-81	135
475	Population pharmacokinetics of high dose ibuprofen in cystic fibrosis. 2003 , 88, 1128-30	9
474	Altered tissue distribution in adults with cystic fibrosis. 2003, 58, 885-9	27
473	Effect of ibuprofen on neutrophil migration in vivo in cystic fibrosis and healthy subjects. 2003 , 306, 1086-91	71

472	Hidden depletion of fat-free mass and bone mineral density in adults with cystic fibrosis. 2003 , 124, 2220-8	57
471	Overview of cystic fibrosis and non-CF bronchiectasis. 2003 , 24, 619-28	6
470	Cor pulmonale in cystic fibrosis. 2003 , 24, 323-30	13
469	Inflammatory and microbiologic markers in induced sputum after intravenous antibiotics in cystic fibrosis. 2003 , 168, 1471-5	141
468	Emerging drug treatments for cystic fibrosis. 2003 , 8, 523-35	7
467	Factors influencing outcomes in cystic fibrosis: a center-based analysis. 2003 , 123, 20-7	169
466	Early aggressive intervention in cystic fibrosis: is it time to redefine our "best practice" strategies?. 2003 , 123, 1-3	26
465	Current treatment of chronic bronchial suppuration. 2003 , 403-427	
464	Current and future treatment of cystic fibrosis. 2003, 428-450	
463	. 2003,	15
463 462	. 2003, Pseudomonas aeruginosa Biofilms in Lung Infections. 2003, 287-310	15
		15 2
462	Pseudomonas aeruginosa Biofilms in Lung Infections. 2003 , 287-310	
462 461	Pseudomonas aeruginosa Biofilms in Lung Infections. 2003, 287-310 Mucoactive Agents for the Treatment of Cough. 269-281	2
462 461 460	Pseudomonas aeruginosa Biofilms in Lung Infections. 2003, 287-310 Mucoactive Agents for the Treatment of Cough. 269-281 Cystic fibrosis: bench to bedside 2003. 2003, 10, 161-4	3
462 461 460 459	Pseudomonas aeruginosa Biofilms in Lung Infections. 2003, 287-310 Mucoactive Agents for the Treatment of Cough. 269-281 Cystic fibrosis: bench to bedside 2003. 2003, 10, 161-4 Cytokine secretion by cystic fibrosis airway epithelial cells. 2004, 169, 645-53	2 3 124
462 461 460 459	Pseudomonas aeruginosa Biofilms in Lung Infections. 2003, 287-310 Mucoactive Agents for the Treatment of Cough. 269-281 Cystic fibrosis: bench to bedside 2003. 2003, 10, 161-4 Cytokine secretion by cystic fibrosis airway epithelial cells. 2004, 169, 645-53 Lung inflammation as a therapeutic target in cystic fibrosis. 2004, 31, 377-81	2 3 124 86

454	Ibuprofen inhibits survival of bladder cancer cells by induced expression of the p75NTR tumor suppressor protein. 2004 , 64, 6207-13	64
453	Ibuprofen and increased morbidity in children with asthma: fact or fiction?. 2004, 6, 267-72	8
452	New therapeutic key for cystic fibrosis: a role for lipoxins. 2004 , 5, 357-8	15
451	High-dose ibuprofen therapy associated with esophageal ulceration after pneumonectomy in a patient with cystic fibrosis: a case report. 2004 , 4, 19	11
450	Pulmonary exacerbations in cystic fibrosis. 2004 , 37, 400-6	141
449	Molecular imaging for pediatric lung diseases. 2004 , 37, 286-96	7
448	CF: an X-ray database to assess effect of aerosolized tobramycin. 2004 , 38, 23-30	14
447	Symposium summaries. 2004 , 38, 92-188	78
446	Early interventions in CF. 2004 , 26, 88-90	4
445	Comparing the efficacy of NSAIDs and paracetamol in children. 2004 , 14, 201-17	30
444	Early intervention and prevention of lung disease in cystic fibrosis: a European consensus. 2004, 3, 67-91	213
443	Cytokines and inflammatory mediators in cystic fibrosis. 2004 , 3, 223-31	80
442	Pharmacokinetics of Ibuprofen in children with cystic fibrosis. 2004 , 43, 145-56	4
441	NSAID-related psychiatric adverse events: who is at risk?. 2004 , 64, 2619-27	40
440	Cystic fibrosis lung disease: when does it start, and how can it be prevented?. 2004 , 145, 6-7	1
439	Effects of CFTR, interleukin-10, and Pseudomonas aeruginosa on gene expression profiles in a CF bronchial epithelial cell Line. 2004 , 10, 562-73	43
438	Pulmonary Effects of Low-Dose Methotrexate Therapy. 2004 , 11, 307-317	1
437	Lung function decline in cystic fibrosis patients and timing for lung transplantation referral. 2004 , 126, 412-9	112

436	Cystic fibrosis adult care: consensus conference report. 2004 , 125, 1S-39S	394
435	Mucoviscidose. 2004 , 1, 1-14	
434	Effects of azithromycin on clinical isolates of Pseudomonas aeruginosa from cystic fibrosis patients. 2005 , 128, 912-9	70
433	Cystic fibrosis: an overview. 2005 , 39, 307-17	30
432	Platelet activation in cystic fibrosis. 2005 , 105, 4635-41	93
431	Perioperative pharmacokinetics of ibuprofen enantiomers after rectal administration. 2005 , 15, 566-73	12
430	Effects of liver transplantation on the nutritional status of patients with cystic fibrosis. 2005 , 18, 246-55	42
429	Amphiphilic pyridinium salts block TNF alpha/NF kappa B signaling and constitutive hypersecretion of interleukin-8 (IL-8) from cystic fibrosis lung epithelial cells. 2005 , 70, 381-93	18
428	Antiinflammatory therapies for cystic fibrosis: past, present, and future. 2005 , 25, 555-73	33
427	Cystic fibrosis lung disease: genetic influences, microbial interactions, and radiological assessment. 2005 , 35, 739-57	28
426	DNA concentrations in BAL fluid of cystic fibrosis patients with early lung disease: influence of treatment with dornase alpha. 2005 , 39, 1-4	67
425	CFTR: more than just a chloride channel. 2005 , 39, 292-8	68
424	Cytokine levels in sputum of cystic fibrosis patients before and after antibiotic therapy. 2005 , 40, 15-21	69
423	Anti-Inflammatory Agents. 2005 , 34, 187-194	
422	Surfactant replacement does not reduce duration of ventilatory support in paediatric acute lung injury. 2005 , 60, 325-325	78
421	The association of acetaminophen, aspirin, and ibuprofen with respiratory disease and lung function. 2005 , 171, 966-71	92
420	Computed tomography correlates with pulmonary exacerbations in children with cystic fibrosis. 2005 , 172, 1128-32	123
419	Progression of lung disease on computed tomography and pulmonary function tests in children and adults with cystic fibrosis. 2006 , 61, 80-5	148

418	Association of tumour necrosis factor alpha variants with the CF pulmonary phenotype. 2005 , 60, 320-5	35
417	Interleukin-1beta differentially regulates beta2 adrenoreceptor and prostaglandin E2-mediated cAMP accumulation and chloride efflux from Calu-3 bronchial epithelial cells. Role of receptor changes, adenylyl cyclase, cyclo-oxygenase 2, and protein kinase A. 2005 , 280, 23451-63	22
416	Assessment of hypoxia in children with cystic fibrosis. 2005 , 90, 1138-43	37
415	Cystic fibrosis, disease severity, and a macrophage migration inhibitory factor polymorphism. 2005 , 172, 1412-5	73
414	Diabetes: a major co-morbidity of cystic fibrosis. 2005 , 31, 221-32	80
413	Use of macrolides and tetracyclines for chronic inflammatory diseases. 2005 , 39, 86-94	37
412	Development of medicines for children in Europe: ethical implications. 2005 , 6, 45-51	23
411	Relation of exaggerated cytokine responses of CF airway epithelial cells to PAO1 adherence. 2005 , 6, 69	12
410	Docosahexaenoic acid trials in cystic fibrosis: a review of the rationale behind the clinical trials. 2005 , 4, 27-34	15
409	Disease-specific reference equations for lung function in patients with cystic fibrosis. 2005 , 172, 885-91	56
408	Anti-inflammatory medications for cystic fibrosis lung disease: selecting the most appropriate agent. 2005 , 4, 255-73	27
407	Pseudomonas aeruginosa tolerance to tobramycin, hydrogen peroxide and polymorphonuclear leukocytes is quorum-sensing dependent. 2005 , 151, 373-383	389
406	Cystic fibrosis pulmonary exacerbations. 2006 , 148, 259-64	109
405	WhatN new in CF airway inflammation: an update. 2006 , 7 Suppl 1, S70-2	24
404	CF-Emerging therapies: Modulation inflammation. 2006 , 7 Suppl 1, S170-4	10
403	Mucoviscidose´: physiopathologie, gfilique, aspects cliniques et thlapeutiques. 2006 , 1, 1-23	
402	Advances in cystic fibrosis therapies. 2006 , 18, 604-13	37
401	Mucoviscidosis (Cystic Fibrosis), Molecular Cell Biology of. 2006 ,	

(2007-2006)

400	evaluating the effect of tobramycin solution for inhalation in cystic fibrosis patients. 2006 , 41, 1129-37	18
399	Interaction of ibuprofen and other structurally related NSAIDs with the sodium-coupled monocarboxylate transporter SMCT1 (SLC5A8). 2006 , 23, 1209-16	36
398	Zystische Fibrose. 2006 , 3, 325-339	O
397	Biological effects of a dietary omega-3 polyunsaturated fatty acids supplementation in cystic fibrosis patients: a randomized, crossover placebo-controlled trial. 2006 , 25, 418-27	67
396	A pilot trial on safety and efficacy of erythrocyte-mediated steroid treatment in CF patients. 2006 , 6, 17	14
395	Absence of cochleotoxicity measured by standard and high-frequency pure tone audiometry in a trial of once- versus three-times-daily tobramycin in cystic fibrosis patients. 2006 , 50, 2293-9	34
394	Quantifying pulmonary inflammation in cystic fibrosis with positron emission tomography. 2006 , 173, 1363-9	85
393	Adenosine regulation of cystic fibrosis transmembrane conductance regulator through prostenoids in airway epithelia. 2006 , 34, 600-8	41
392	Inhaled heparin in cystic fibrosis. 2006 , 27, 354-8	32
391	Inflammation is a modulator of the insulin-like growth factor (IGF)/IGF-binding protein system inducing reduced bioactivity of IGFs in cystic fibrosis. 2006 , 154, 47-52	54
390	COX-2: a link between airway inflammation and disordered chloride secretion in cystic fibrosis?. 2006 , 61, 552-3	6
389	Talniflumate increases survival in a cystic fibrosis mouse model of distal intestinal obstructive syndrome. 2006 , 317, 275-83	31
388	Macrolide resistance of Staphylococcus aureus and Haemophilus species associated with long-term azithromycin use in cystic fibrosis. 2006 , 57, 741-6	79
387	Upregulation of COX-1 and COX-2 in nasal polyps in cystic fibrosis. 2006 , 61, 592-6	29
386	FDG-PET imaging of pulmonary inflammation in healthy volunteers after airway instillation of endotoxin. 2006 , 100, 1602-9	68
385	The inflammatory role of platelets in cystic fibrosis. 2006 , 173, 483-90	64
384	Improving rate of decline of FEV1 in young adults with cystic fibrosis. 2006, 61, 155-7	82
383	Novel pharmaceutical approaches for treating patients with cystic fibrosis. 2007 , 13, 3252-63	7

382	Oral non-steroidal anti-inflammatory drug therapy for cystic fibrosis. 2007 , CD001505	32
381	Parthenolide inhibits IkappaB kinase, NF-kappaB activation, and inflammatory response in cystic fibrosis cells and mice. 2007 , 36, 728-36	109
380	Ibuprofen therapy and nasal polyposis in cystic fibrosis patients. 2007 , 36, 309-14	23
379	Advancing outcome measures for the new era of drug development in cystic fibrosis. 2007, 4, 370-7	43
378	Oral corticosteroid therapy in cystic fibrosis patients hospitalized for pulmonary exacerbation: a pilot study. 2007 , 132, 1212-8	44
377	IL-23 mediates inflammatory responses to mucoid Pseudomonas aeruginosa lung infection in mice. 2007 , 292, L519-28	154
376	Inflammation in cystic fibrosiswhen and why? Friend or foe?. 2007, 28, 286-94	7
375	Clinical use of Ibuprofen is associated with slower FEV1 decline in children with cystic fibrosis. 2007 , 176, 1084-9	144
374	Heritability of lung disease severity in cystic fibrosis. 2007 , 175, 1036-43	147
373	Pharmacotherapy for bronchiectasis. 2007 , 8, 3183-93	5
3,3		
372	Exacerbations in cystic fibrosis: 2 . prevention. 2007 , 62, 723-32	33
372	Exacerbations in cystic fibrosis: 2 . prevention. 2007 , 62, 723-32	33
37 ² 37 ¹	Exacerbations in cystic fibrosis: 2 . prevention. 2007 , 62, 723-32 Pulmonary exacerbations in cystic fibrosis and bronchiectasis. 2007 , 62, 288-90	33 17
372 371 370	Exacerbations in cystic fibrosis: 2 . prevention. 2007 , 62, 723-32 Pulmonary exacerbations in cystic fibrosis and bronchiectasis. 2007 , 62, 288-90 Dysphonia. 2007 , 46, 283-5	33 17 1
37 ² 37 ¹ 37 ⁰ 369	Exacerbations in cystic fibrosis: 2 . prevention. 2007, 62, 723-32 Pulmonary exacerbations in cystic fibrosis and bronchiectasis. 2007, 62, 288-90 Dysphonia. 2007, 46, 283-5 Azithromycin protects against hyperoxic lung injury in neonatal rats. 2007, 55, 299-305 The aryl propionic acid R-flurbiprofen selectively induces p75NTR-dependent decreased survival of	33 17 1
372 371 370 369 368	Exacerbations in cystic fibrosis: 2 . prevention. 2007, 62, 723-32 Pulmonary exacerbations in cystic fibrosis and bronchiectasis. 2007, 62, 288-90 Dysphonia. 2007, 46, 283-5 Azithromycin protects against hyperoxic lung injury in neonatal rats. 2007, 55, 299-305 The aryl propionic acid R-flurbiprofen selectively induces p75NTR-dependent decreased survival of prostate tumor cells. 2007, 67, 3254-62	33 17 1 19 48

(2007-2007)

364	A review of ibuprofen and acetaminophen use in febrile children and the occurrence of asthma-related symptoms. 2007 , 29, 2716-23	37
363	Actualit the the peutiques 2007 dans la mucoviscidose. 2007 , 24, 109-111	
362	Inflammation and anti-inflammatory therapies for cystic fibrosis. 2007, 28, 331-46	78
361	Cystic fibrosis pulmonary guidelines: chronic medications for maintenance of lung health. 2007 , 176, 957-69	502
360	Pseudomonas aeruginosa and the host pulmonary immune response. 2007 , 1, 121-37	5
359	Use of high-dose ibuprofen in a pediatric cystic fibrosis center. 2007 , 6, 153-8	27
358	Bioelectric effects of quinine on polarized airway epithelial cells. 2007 , 6, 351-9	4
357	Clinical trials in cystic fibrosis. 2007 , 6, 85-99	37
356	Sputum biomarkers of inflammation in cystic fibrosis lung disease. 2007 , 4, 406-17	122
355	Pharmacologic, pharmacodynamic, and pharmacokinetic considerations with intravenous Ibuprofen lysine. 2007 , 12, 158-70	2
354	Abstracts of the 21st Annual North American Cystic Fibrosis Conference, October 3-6, 2007, Anaheim, California, USA. 2007 , 30, 99-412	
353	Recurrent Pseudomonas bronchopneumonia and other symptoms as in cystic fibrosis in a child with type I pseudohypoaldosteronism. 2007 , 87, 472-474	28
352	50 Years Ago in The Journal of Pediatrics. 2007 , 151, 139	
351	Risk factors for rate of decline in forced expiratory volume in one second in children and adolescents with cystic fibrosis. 2007 , 151, 134-9, 139.e1	328
350	Pacing the marathon: rate of decline of pulmonary function in cystic fibrosis. 2007, 151, 111-3	4
349	High-dose ibuprofen in cystic fibrosis: Canadian safety and effectiveness trial. 2007 , 151, 249-54	114
348	Non! to non-steroidal anti-inflammatory therapy for inflammatory lung disease in cystic fibrosis (at least at the moment). 2007 , 151, 228-30	15
347	Do inhaled corticosteroids impair long-term growth in prepubertal cystic fibrosis patients?. 2007 , 166, 23-8	28

346	Chronic inflammation in the cystic fibrosis lung: alterations in inter- and intracellular signaling. 2008 , 34, 146-62	86
345	Anti-inflammatory therapies for cystic fibrosis-related lung disease. 2008 , 35, 135-53	30
344	WhatN new in cystic fibrosis? From treating symptoms to correction of the basic defect. 2008, 167, 839-49	57
343	Dornase alfa and progression of lung disease in cystic fibrosis. 2008 , 43, S24-S28	5
342	A role for aerosolized antibiotics. 2008 , 43, S29-S34	5
341	Relationship between inhaled corticosteroid therapy and rate of lung function decline in children with cystic fibrosis. 2008 , 153, 746-51	65
340	Association between practice patterns and body mass index percentile in infants and young children with cystic fibrosis. 2008 , 7, 385-90	3
339	Recent advances in cystic fibrosis. 2008 , 9, 144-8	31
338	Clinical trials in infants with cystic fibrosis detected following newborn screening. 2008, 9, 176-80	3
337	Oral DHA supplementation in DeltaF508 homozygous cystic fibrosis patients. 2008 , 78, 109-15	47
337 336	Oral DHA supplementation in DeltaF508 homozygous cystic fibrosis patients. 2008 , 78, 109-15 Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008 , 55, 99-121	47
336	Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008 , 55, 99-121	7
336 335	Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008 , 55, 99-121 Proteases and cystic fibrosis. 2008 , 40, 1238-45	7
336 335 334	Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008, 55, 99-121 Proteases and cystic fibrosis. 2008, 40, 1238-45 Neutrophils in cystic fibrosis. 2009, 64, 81-8	7 144 142
336335334333	Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008, 55, 99-121 Proteases and cystic fibrosis. 2008, 40, 1238-45 Neutrophils in cystic fibrosis. 2009, 64, 81-8 Respiratory Manifestations. 2008, 871-887	7 144 142
336335334333332	Cystic fibrosis: a review of pulmonary and nutritional therapies. 2008, 55, 99-121 Proteases and cystic fibrosis. 2008, 40, 1238-45 Neutrophils in cystic fibrosis. 2009, 64, 81-8 Respiratory Manifestations. 2008, 871-887 Intensive care management of the patient with cystic fibrosis. 2008, 23, 159-77	7 144 142 1

(2009-2008)

328	Risk factors for increased need for intravenous antibiotics for pulmonary exacerbations in adult patients with cystic fibrosis. 2008 , 5, 29-33	19
327	Peroxisome proliferator-activated receptor-gamma in cystic fibrosis lung epithelium. 2008 , 295, L303-13	48
326	Persistent methicillin-resistant Staphylococcus aureus and rate of FEV1 decline in cystic fibrosis. 2008 , 178, 814-21	243
325	Nonsteroidal anti-inflammatory drugs upregulate function of wild-type and mutant CFTR. 2008 , 32, 334-43	18
324	Cystic fibrosis papers of the year 2007. 2008 , 101 Suppl 1, S10-4	1
323	Innate immunity mediated by TLR5 as a novel antiinflammatory target for cystic fibrosis lung disease. 2008 , 180, 7764-73	71
322	Cystic fibrosis lung disease in adult patients. 2008 , 120, 64-74	6
321	CHOP transcription factor mediates IL-8 signaling in cystic fibrosis bronchial epithelial cells. 2008 , 38, 176-84	51
320	Ibuprofen therapy for cystic fibrosis lung disease: revisited. 2008, 14, 567-73	49
319	End of life care in chronic obstructive pulmonary disease: in search of a good death. 2008 , 3, 11-29	46
318	Multicenter cross-sectional study of nontuberculous mycobacterial infections among cystic fibrosis patients, Israel. 2008 , 14, 378-84	148
317	NS-398, ibuprofen, and cyclooxygenase-2 RNA interference produce significantly different gene expression profiles in prostate cancer cells. 2009 , 8, 261-73	36
316	A pipeline of therapies for cystic fibrosis. 2009 , 30, 611-26	22
315	The triterpenoid CDDO limits inflammation in preclinical models of cystic fibrosis lung disease. 2009 , 297, L828-36	36
314	The clinical approach to lung disease in patients with cystic fibrosis. 2009, 30, 505-13	28
313	Stabilization of lung function and clinical symptoms in a patient with cystic fibrosis (CF) after institution of infliximab: a monoclonal antibody that binds tumor necrosis factor alpha. 2009 , 187, 149-52	9
312	Ibuprofen: pharmacology, efficacy and safety. 2009 , 17, 275-342	302
311	Symposium Summaries. 2009 , 44, 109-212	5

310	Highlights of a workshop to discuss targeting inflammation in cystic fibrosis. 2009, 8, 1-8	14
309	Inhaled medication and inhalation devices for lung disease in patients with cystic fibrosis: A European consensus. 2009 , 8, 295-315	193
308	[Protocol for the diagnosis and follow up of patients with cystic fibrosis]. 2009 , 71, 250-64	11
307	Bronchiectasis in children. 2009 , 56, 157-71, xi	20
306	The role of inhaled corticosteroids in the management of cystic fibrosis. 2009, 11, 101-13	31
305	Characterizing aggressiveness and predicting future progression of CF lung disease. 2009 , 8 Suppl 1, S15-9	41
304	Cystic fibrosis. 2009 , 373, 1891-904	972
303	A phase II study on safety and efficacy of high-dose N-acetylcysteine in patients with cystic fibrosis. 2009 , 14, 352-8	34
302	Inflammation, Hyperinflammation & Cystic Fibrosis Lung Disease 🖪 Paradigm Shift?. 2009 , 5, 136-148	
301	New Genetic and Pharmacological Treatments for Cystic Fibrosis. 2009 , 5, 8-27	2
300	Update on new pulmonary therapies. 2009 , 15, 604-10	4
299	Prostaglandin-endoperoxide synthase genes COX1 and COX2 - novel modifiers of disease severity in cystic fibrosis patients. 2010 , 51, 323-30	13
298	Protracted bacterial bronchitis (PBB). 2010 , 11, S71-S73	
297	Cystic fibrosis: exploiting its genetic basis in the hunt for new therapies. 2010 , 125, 219-29	45
296	Ibuprofen for neuroprotection after cerebral ischemia. 2010 , 139, 489-93	14
295	Co-spray-dried mannitol-ciprofloxacin dry powder inhaler formulation for cystic fibrosis and chronic obstructive pulmonary disease. 2010 , 40, 239-47	78
294	Cystic fibrosis lung disease starts in the small airways: can we treat it more effectively?. 2010 , 45, 107-17	126
293	Real-time, once-daily monitoring of symptoms and FEV in cystic fibrosis patientsa feasibility study using a novel device. 2010 , 4, 74-82	19

292	Potential of anti-inflammatory treatment for cystic fibrosis lung disease. 2010 , 3, 61-74	11
291	Pediatric Respiratory Assembly. Mini symposium on lung inflammation. 2010 , 17, e35-41	
290	High-Dose Ibuprofen in Cystic Fibrosis. 2010 , 3, 2213-2224	18
289	A disease-relevant high-content screening assay to identify anti-inflammatory compounds for use in cystic fibrosis. 2010 , 15, 1204-10	7
288	Molecular Basis of Pulmonary Disease. 2010 ,	3
287	Improving evidence-based care in cystic fibrosis through quality improvement. 2010 , 164, 957-60	22
286	TLR5 as an anti-inflammatory target and modifier gene in cystic fibrosis. 2010 , 185, 7731-8	56
285	In cystic fibrosis homozygotes and heterozygotes, neutrophil apoptosis is delayed and modulated by diamide or roscovitine: evidence for an innate neutrophil disturbance. 2010 , 2, 260-6	71
284	Platelet proinflammatory activity in clinically stable patients with CF starts in early childhood. 2010 , 9, 179-86	10
283	Design and powering of cystic fibrosis clinical trials using rate of FEV(1) decline as an efficacy endpoint. 2010 , 9, 332-8	35
282	Dietary supplementation with pressurized whey in patients with cystic fibrosis. 2010 , 13, 77-82	31
281	Emerging therapies in cystic fibrosis. 2010 , 4, 177-85	31
280	Pseudomonas aeruginosa biofilm infections in cystic fibrosis: insights into pathogenic processes and treatment strategies. 2010 , 14, 117-30	95
279	Consequences of airway neutrophilia in children. 2010 , 11, S67-S71	
278	Ibuprofen modulates NF-kB activity but not IL-8 production in cystic fibrosis respiratory epithelial cells. 2010 , 79, 234-42	28
277	Clinical significance of microbial infection and adaptation in cystic fibrosis. 2011 , 24, 29-70	271
276	Targeting airway inflammation in cystic fibrosis in children: past, present, and future. 2011 , 13, 141-7	18
275	Managing cystic fibrosis: strategies that increase life expectancy and improve quality of life. 2011 , 183, 1463-71	207

274	Applying clinical outcome variables to appropriate aerosolized antibiotics for the treatment of patients with cystic fibrosis. 2011 , 105 Suppl 2, S18-23	14
273	Pain is a common problem affecting clinical outcomes in adults with cystic fibrosis. 2011 , 140, 1598-1603	40
272	Inverse relation between vitamin D and serum total immunoglobulin G in the Scandinavian Cystic Fibrosis Nutritional Study. 2011 , 65, 102-9	61
271	Transcript profiling identifies novel key players mediating the growth inhibitory effect of NS-398 on human pancreatic cancer cells. 2011 , 650, 170-7	20
270	The cystic fibrosis neutrophil: a specialized yet potentially defective cell. 2011 , 59, 97-112	61
269	Clinical use of dornase alpha is associated with a slower rate of FEV1 decline in cystic fibrosis. 2011 , 46, 545-53	66
268	Update in cystic fibrosis 2010. 2011 , 183, 1620-4	16
267	Inhaled corticosteroids and lower lung function decline in young children with cystic fibrosis. 2011 , 37, 1091-5	23
266	Down-regulation of cytokine-induced interleukin-8 requires inhibition of p38 mitogen-activated protein kinase (MAPK) via MAPK phosphatase 1-dependent and -independent mechanisms. 2011 , 286, 15998-6007	24
265	Platelets in pulmonary vascular physiology and pathology. 2012 , 2, 291-308	33
264	Effect of pulmonary exacerbations on long-term lung function decline in cystic fibrosis. 2012, 40, 61-6	146
263	Integrative Therapies in Lung Health and Sleep. 2012 ,	
262	Progressive flow-to-volume dysanapsis in cystic fibrosis: a predictor for lung transplantation?. 2012 , 186, 82-7	16
261	Outcome measures for clinical trials assessing treatment of cystic fibrosis lung disease. 2012 , 2, 163-175	28
2 60	Atypical activation of the unfolded protein response in cystic fibrosis airway cells contributes to p38 MAPK-mediated innate immune responses. 2012 , 189, 5467-75	51
259	New therapies in cystic fibrosis. 2012 , 18, 614-27	19
258	Progress in cystic fibrosis and the CF Therapeutics Development Network. 2012 , 67, 882-90	49
257	Effect of dornase alfa on inflammation and lung function: potential role in the early treatment of cystic fibrosis. 2012 , 11, 78-83	52

256	State of progress in treating cystic fibrosis respiratory disease. 2012 , 10, 88	46
255	Pulmonary Disease in Cystic Fibrosis. 2012 , 770-780	6
254	Ibuprofen: Pharmacology, Therapeutics and Side Effects. 2012 ,	18
253	The Prognosis of Cystic Fibrosis - A ClinicianN Perspective. 2012 ,	
252	Channel Replacement Therapy for Cystic Fibrosis. 2012,	О
251	Liver transplantation in chronic cholestatic conditions. 2012 , 17, 959-69	12
250	Cystic fibrosis: a mucosal immunodeficiency syndrome. 2012 , 18, 509-19	336
249	Pathogenesis and management of nasal polyposis in cystic fibrosis. 2012 , 12, 163-74	52
248	Metabolomic profiling of regulatory lipid mediators in sputum from adult cystic fibrosis patients. 2012 , 53, 160-71	106
247	Enhancement of anti-inflammatory drug activity by multivalent adamantane-based dendrons. 2012 , 33, 5610-7	26
246	CFTR negatively regulates cyclooxygenase-2-PGE(2) positive feedback loop in inflammation. 2012 , 227, 2759-66	36
245	R-flurbiprofen, a novel nonsteroidal anti-inflammatory drug, decreases cell proliferation and induces apoptosis in pituitary adenoma cells in vitro. 2012 , 106, 561-9	14
244	Mechanism and Causality in Biology and Economics. 2013,	3
243	New advances in the therapy of non-cystic fibrosis bronchiectasis. 2013 , 19, 266-275	5
242	Is adolescentsNeligious coping with cystic fibrosis associated with the rate of decline in pulmonary function?-A preliminary study. 2013 , 19, 33-42	12
241	Cystic fibrosis therapeutics: the road ahead. 2013 , 143, 207-213	65
240	Safety and early treatment effects of the CXCR2 antagonist SB-656933 in patients with cystic fibrosis. 2013 , 12, 241-8	84
239	FDG PET imaging in cystic fibrosis. 2013 , 43, 412-9	12

238	New advances in the therapy of non-cystic fibrosis bronchiectasis. 2013 , 19, 266-75		7
237	Early lung disease in cystic fibrosis. 2013 , 1, 148-57		58
236	Cystic fibrosis pulmonary guidelines. Chronic medications for maintenance of lung health. 2013 , 187, 680-9		432
235	Risk factors for bronchiectasis in children with cystic fibrosis. <i>New England Journal of Medicine</i> , 2013 , 368, 1963-70	59.2	369
234	Synthesis and Spectrophotometric Determination Ibuprofen Charge Transfer Complexes with P-Chloranil, 7,7,8,8-Tetracyanoquinodimethane, Bromothymol Blue, Methyl Orange and Picric Acid. 2013 , 05,		
233	Antibiotic and anti-inflammatory therapies for cystic fibrosis. 2013 , 3, a009779		40
232	Oral contraceptives do not appear to affect cystic fibrosis disease severity. 2013 , 41, 67-73		17
231	The role of endoscopy and biopsy in the management of severe gastrointestinal disease in cystic fibrosis patients. 2013 , 48, 1181-9		8
230	Induction of cyclooxygenase-2 signaling by Stomatococcus mucilaginosus highlights the pathogenic potential of an oral commensal. 2013 , 191, 3810-7		13
229	Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. 2013 , CD001505		23
228	Anti-inflammatory therapies for cystic fibrosis lung disease. 2013 , 82-92		
227	Important role of platelets in modulating endotoxin-induced lung inflammation in CFTR-deficient mice. 2013 , 8, e82683		12
226	Deciphering miRNA transcription factor feed-forward loops to identify drug repurposing candidates for cystic fibrosis. 2014 , 6, 94		16
225	Histatins: salivary peptides with copper(II)- and zinc(II)-binding motifs: perspectives for biomedical applications. 2014 , 281, 657-72		76
224	What is the optimal management of chronic rhinosinusitis in cystic fibrosis?. 2014 , 22, 42-6		11
223	Lung inflammation in cystic fibrosis: pathogenesis and novel therapies. 2014 , 47, 539-46		40
222	High-dose ibuprofen is not associated with increased biomarkers of kidney injury in patients with cystic fibrosis. 2014 , 49, 148-53		21
221	Symposium Session Summaries. 2014 , 49, S116-S215		1

(2015-2014)

220	Dissecting the reaction of Phase II metabolites of ibuprofen and other NSAIDS with human plasma protein. 2014 , 5, 3789-3794	14
219	Pediatric Chest Imaging. 2014 ,	1
218	Risk factors for the progression of cystic fibrosis lung disease throughout childhood. 2014 , 11, 63-72	47
217	A randomized double blind, placebo controlled phase 2 trial of BIIL 284 BS (an LTB4 receptor antagonist) for the treatment of lung disease in children and adults with cystic fibrosis. 2014 , 13, 148-55	93
216	Advances in the diagnosis and treatment of cystic fibrosis. 2014 , 61, 225-43	10
215	Thoracic Manifestations of Systemic Diseases. 2014 , 395-429	1
214	Improving complex medical care while awaiting next-generation CFTR potentiators and correctors: The current pipeline of therapeutics. 2015 , 50 Suppl 40, S66-73	6
213	Symposium Summaries. 2015 , 50 Suppl 41, S108-92	1
212	Pharmacology and Toxicology of Ibuprofen. 2015 , 132-236	2
211	Essential oils in the treatment of respiratory tract diseases highlighting their role in bacterial infections and their anti-inflammatory action: a review. 2015 , 30, 331-341	44
210	Decline in lung function does not predict future decline in lung function in cystic fibrosis patients. 2015 , 50, 856-62	13
209	Risk factors for lung function decline in a large cohort of young cystic fibrosis patients. 2015 , 50, 763-70	71
208	Gene expression profile of ABC transporters and cytotoxic effect of ibuprofen and acetaminophen in an epithelial ovarian cancer cell line in vitro. 2015 , 37, 283-90	8
207	Genetic Deletion and Pharmacological Inhibition of PI3K (Reduces Neutrophilic Airway Inflammation and Lung Damage in Mice with Cystic Fibrosis-Like Lung Disease. 2015 , 2015, 545417	13
206	Use of ibuprofen to assess inflammatory biomarkers in induced sputum: Implications for clinical trials in cystic fibrosis. 2015 , 14, 720-6	43
205	The Evolution of Cystic Fibrosis Care. 2015 , 148, 533-542	32
204	Spanish consensus on the prevention and treatment of Pseudomonas aeruginosa bronchial infections in cystic fibrosis patients. 2015 , 51, 140-50	29
203	The challenges of developing effective anti-inflammatory agents in cystic fibrosis. 2015 , 14, 164-6	6

202	[Cystic fibrosis and associated complications]. 2015 , 56, 263-74	2
201	Spanish Consensus on the Prevention and Treatment of Pseudomonas aeruginosa Bronchial Infections in Cystic Fibrosis Patients. 2015 , 51, 140-150	4
200	Pharmacokinetics and tolerability of oral sildenafil in adults with cystic fibrosis lung disease. 2015 , 14, 228-36	29
199	Cystic fibrosis. 2015 , 1, 15010	258
198	Considerations for the Conduct of Clinical Trials with Antiinflammatory Agents in Cystic Fibrosis. A Cystic Fibrosis Foundation Workshop Report. 2015 , 12, 1398-406	28
197	Long-term treatment with oral N-acetylcysteine: affects lung function but not sputum inflammation in cystic fibrosis subjects. A phase II randomized placebo-controlled trial. 2015 , 14, 219-27	73
196	42-year-old man with asthma symptoms and recurrent bronchitis. 2015 , 90, 527-31	
195	Pulmonary function outcomes for assessing cystic fibrosis care. 2015 , 14, 376-83	11
194	Chiral liquid membrane for enantioselective separation of racemic ibuprofen by L-tartaric acid derivatives. 2015 , 5, 41729-41735	12
193	Inflammation in cystic fibrosis lung disease: Pathogenesis and therapy. 2015 , 14, 419-30	276
193 192	Inflammation in cystic fibrosis lung disease: Pathogenesis and therapy. 2015 , 14, 419-30 Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015 , 14, 763-9	276 30
	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic	
192	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015 , 14, 763-9	30
192 191	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015 , 14, 763-9 Public health evolutionary biology of antimicrobial resistance: priorities for intervention. 2015 , 8, 223-39 Lumacaftor and ivacaftor in the management of patients with cystic fibrosis: current evidence and	30
192 191 190	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015, 14, 763-9 Public health evolutionary biology of antimicrobial resistance: priorities for intervention. 2015, 8, 223-39 Lumacaftor and ivacaftor in the management of patients with cystic fibrosis: current evidence and future prospects. 2015, 9, 313-26	30
192 191 190	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015, 14, 763-9 Public health evolutionary biology of antimicrobial resistance: priorities for intervention. 2015, 8, 223-39 Lumacaftor and ivacaftor in the management of patients with cystic fibrosis: current evidence and future prospects. 2015, 9, 313-26 Ibuprofen. 2016, 5-12	30 37 52
192 191 190 189	Treatment and demographic factors affecting time to next pulmonary exacerbation in cystic fibrosis. 2015, 14, 763-9 Public health evolutionary biology of antimicrobial resistance: priorities for intervention. 2015, 8, 223-39 Lumacaftor and ivacaftor in the management of patients with cystic fibrosis: current evidence and future prospects. 2015, 9, 313-26 Ibuprofen. 2016, 5-12 Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. 2016, 4, CD001505	30 37 52 20

(2017-2016)

184	New and emerging targeted therapies for cystic fibrosis. 2016 , 352, i859	91
183	Current and future treatment options for cystic fibrosis lung disease: latest evidence and clinical implications. 2016 , 7, 170-83	31
182	Key mechanisms governing resolution of lung inflammation. 2016 , 38, 425-48	138
181	Handbook of Cystic Fibrosis. 2016 ,	4
180	Treatment of Cystic Fibrosis. 2016 , 77-99	1
179	Background and Epidemiology. 2016 , 63, 567-84	43
178	Cystic Fibrosis: Microbiology and Host Response. 2016 , 63, 617-36	23
177	Improved Efficiency of Ibuprofen by Cationic Carbosilane Dendritic Conjugates. 2016 , 13, 3427-3438	12
176	Biomarkers for cystic fibrosis drug development. 2016 , 15, 714-723	46
175	Aminoglycoside antibiotics. 2016 , 216-236	O
174	Carrier-free high-dose dry powder inhaler formulation of ibuprofen: Physicochemical characterization and in vitro aerodynamic performance. 2016 , 511, 403-414	23
173	Question 5: What is the role of macrolide antibiotics as anti-inflammatory treatment in Cystic fibrosis?. 2016 , 18, 55-57	
172	CFTR and Cystic Fibrosis. 2016 , 519-552	3
171	Cystic Fibrosis and Its Management Through Established and Emerging Therapies. 2016 , 17, 155-75	28
170	Using Cystic Fibrosis Therapies for Non-Cystic Fibrosis Bronchiectasis. 2016 , 37, 139-46	8
169	Cystic Fibrosis. 2016 , 822-852.e17	4
168	Bronchiectasis. 2016 , 853-876.e9	3
167	Use of FEV in cystic fibrosis epidemiologic studies and clinical trials: A statistical perspective for the clinical researcher. 2017 , 16, 318-326	54

166	Anti-Inflammatory Therapies for Cystic Fibrosis. 2017 , 139-151	3
165	AJRCCM: 100-Year Anniversary. Progress along the Pathway of Discovery Leading to Treatment and Cure of Cystic Fibrosis. 2017 , 195, 1092-1099	13
164	Working Towards an Appropriate Use of Ibuprofen in Children: An Evidence-Based Appraisal. 2017 , 77, 1295-1311	37
163	Efficacy of Rhesus Theta-Defensin-1 in Experimental Models of Pseudomonas aeruginosa Lung Infection and Inflammation. 2017 , 61,	15
162	The treatment of the pulmonary and extrapulmonary manifestations of cystic fibrosis. 2017, 46, e139-e164	9
161	Medical and Surgical Advancements in the Management of Cystic Fibrosis Chronic Rhinosinusitis. 2017 , 5, 24-34	28
160	A little CFTR can change a lot: slowing cystic fibrosis progression. 2017 , 5, 86-87	4
159	Assessment of safety and efficacy of long-term treatment with combination lumacaftor and ivacaftor therapy in patients with cystic fibrosis homozygous for the F508del-CFTR mutation (PROGRESS): a phase 3, extension study. 2017 , 5, 107-118	158
158	Comparison of FEV reference equations for evaluating a cystic fibrosis therapeutic intervention. 2017 , 52, 1013-1019	6
157	Mapping targetable inflammation and outcomes with cystic fibrosis biomarkers. 2017 , 52, S21-S28	7
156	Corticosteroid-resistant inflammatory signalling in -infected bronchial cells. 2017, 3,	3
155	Association of Acetaminophen and Ibuprofen Use With Wheezing in Children With Acute Febrile Illness. 2017 , 51, 239-244	6
154	Phase I Studies of Acebilustat: Biomarker Response and Safety in Patients with Cystic Fibrosis. 2017 , 10, 28-34	33
153	Neutrophils. 2017 , 253-272	
152	Brazilian guidelines for the diagnosis and treatment of cystic fibrosis. 2017 , 43, 219-245	39
151	Effect of inhaled steroids on clinical and inflammatoryparameters in children with cystic fibrosis. 2017 , 47, 1432-1440	2
150	Reference percentiles of FEV1 for the Canadian cystic fibrosis population: comparisons across time and countries. 2018 , 73, 446-450	5
149	Ajulemic acid: potential treatment for chronic inflammation. 2018 , 6, e00394	39

(2018-2018)

148	Association of High-Dose Ibuprofen Use, Lung Function Decline, and Long-Term Survival in Children with Cystic Fibrosis. 2018 , 15, 485-493	38
147	Increased platelet activation occurs in cystic fibrosis patients and correlates to clinical status. 2018 , 162, 32-37	14
146	Antimicrobial Activity of Ibuprofen against Cystic Fibrosis-Associated Gram-Negative Pathogens. 2018 , 62,	26
145	KB001-A, a novel anti-inflammatory, found to be safe and well-tolerated in cystic fibrosis patients infected with Pseudomonas aeruginosa. 2018 , 17, 484-491	41
144	Impact of a cystic fibrosis transmembrane conductance regulator (CFTR) modulator on high-dose ibuprofen therapy in pediatric cystic fibrosis patients. 2018 , 53, 1035-1039	5
143	The Long and the Short of It in Cystic Fibrosis Clinical Research Outcomes. 2018 , 15, 430-431	
142	Formulation and delivery strategies of ibuprofen: challenges and opportunities. 2018, 44, 173-183	52
141	Infectious Complications in Special Hosts. 2018 , 643-651.e3	
140	Is there a role for inhaled anti-inflammatory drugs in cystic fibrosis treatment?. 2018, 6, 69-84	2
139	Rapid therapeutic advances in CFTR modulator science. 2018 , 53, S4-S11	27
139	Rapid therapeutic advances in CFTR modulator science. 2018 , 53, S4-S11 Symposium Summaries. 2018 , 53, S36-S147	27 O
138	Symposium Summaries. 2018 , 53, S36-S147	0
138	Symposium Summaries. 2018, 53, S36-S147 Omega-3 PUFA vs. NSAIDs for Preventing Cardiac Inflammation. 2018, 5, 146	3
138 137 136	Symposium Summaries. 2018, 53, S36-S147 Omega-3 PUFA vs. NSAIDs for Preventing Cardiac Inflammation. 2018, 5, 146 Risks associated with lung transplantation in cystic fibrosis patients. 2018, 12, 893-904	3
138 137 136	Symposium Summaries. 2018, 53, S36-S147 Omega-3 PUFA vs. NSAIDs for Preventing Cardiac Inflammation. 2018, 5, 146 Risks associated with lung transplantation in cystic fibrosis patients. 2018, 12, 893-904 Induced pluripotent stem cells for treating cystic fibrosis: State of the science. 2018, 53, S12-S29 EMPIRE-CF: A phase II randomized placebo-controlled trial of once-daily, oral acebilustat in adult	o 3 2 16
138 137 136 135	Symposium Summaries. 2018, 53, S36-S147 Omega-3 PUFA vs. NSAIDs for Preventing Cardiac Inflammation. 2018, 5, 146 Risks associated with lung transplantation in cystic fibrosis patients. 2018, 12, 893-904 Induced pluripotent stem cells for treating cystic fibrosis: State of the science. 2018, 53, S12-S29 EMPIRE-CF: A phase II randomized placebo-controlled trial of once-daily, oral acebilustat in adult patients with cystic fibrosis - Study design and patient demographics. 2018, 72, 86-94 High Concentrations of Sodium Chloride Improve Microbicidal Activity of Ibuprofen against	o 3 2 16

130	Unmet needs in cystic fibrosis: the next steps in improving outcomes. 2018 , 12, 585-593	15
129	Pulmonary Disease in Cystic Fibrosis. 2019 , 777-787.e4	3
128	A specialized method of sputum collection and processing for therapeutic interventions in cystic fibrosis. 2019 , 18, 203-211	8
127	Cystic Fibrosis: an Update on Disease Pathophysiology, Management, and Novel Modalities of Therapy. 2019 , 6, 226-237	
126	Acute Pulmonary Exacerbations in Cystic Fibrosis. 2019 , 40, 792-803	15
125	Anti-inflammatories and mucociliary clearance therapies in the age of CFTR modulators. 2019 , 54 Suppl 3, S46-S55	14
124	Treating the Airway Consequences of Cystic Fibrosis Transmembrane Conductance Regulator Dysfunction. 2019 , 40, 751-761	
123	Targeting airway inflammation in cystic fibrosis. 2019 , 13, 1041-1055	10
122	Recombinant Human Deoxyribonuclease I. 2019 , 471-488	3
121	Acetyl-CoA carboxylase inhibition regulates microtubule dynamics and intracellular transport in cystic fibrosis epithelial cells. 2019 , 316, L1081-L1093	3
120	Gene therapy-emulating small molecule treatments in cystic fibrosis airway epithelial cells and patients. 2019 , 20, 290	9
119	Oral non-steroidal anti-inflammatory drug therapy for lung disease in cystic fibrosis. 2019 , 9, CD001505	20
118	Stem Cell-Based Therapy for Lung Disease. 2019 ,	3
117	Physiological study of pulmonary involvement in adults with cystic fibrosis through simulated modeling of different clinical scenarios. 2019 , 57, 413-425	O
116	Azithromycin is the answer in paediatric respiratory medicine, but what was the question?. 2020 , 34, 67-74	10
115	Strong toll-like receptor responses in cystic fibrosis patients are associated with higher lung function. 2020 , 19, 608-613	2
114	Cystic Fibrosis and Genotype-Dependent Therapy: Is There a Need for a Sex-Specific Therapy?. 2020 , 4, 247028972093702	О
113	Targeting the Heme Oxygenase 1/Carbon Monoxide Pathway to Resolve Lung Hyper-Inflammation and Restore a Regulated Immune Response in Cystic Fibrosis. 2020 , 11, 1059	11

(2021-2020)

112	Ibuprofen, a traditional drug that may impact the course of COVID-19 new effective formulation in nebulizable solution. 2020 , 144, 110079	8
111	Preparation, characterization and improved release profile of ibuprofen-phospholipid association. 2020 , 60, 101951	10
110	Cystic fibrosis - Ten promising therapeutic approaches in the current era of care. 2020 , 29, 1107-1124	4
109	Selectively targeting key inflammatory pathways in cystic fibrosis. 2020 , 206, 112717	2
108	Role of Cystic Fibrosis Bronchial Epithelium in Neutrophil Chemotaxis. 2020 , 11, 1438	9
107	The Pseudomonas aeruginosa protease LasB directly activates IL-1 2020 , 60, 102984	7
106	Pharmacological and Genomic Approaches in Management of Cystic Fibrosis. 2020, 30, 191-197	0
105	Evaluation of a five-year predicted survival model for cystic fibrosis in later time periods. 2020 , 10, 6602	3
104	Building global development strategies for cf therapeutics during a transitional cftr modulator era. 2020 , 19, 677-687	8
103	Glycosaminoglycans as Multifunctional Anti-Elastase and Anti-Inflammatory Drugs in Cystic Fibrosis Lung Disease. 2020 , 11, 1011	4
102	The Resolution Approach to Cystic Fibrosis Inflammation. 2020 , 11, 1129	2
101	Resveratrol restores intracellular transport in cystic fibrosis epithelial cells. 2020 , 318, L1145-L1157	1
100	Evaluation of airway and circulating inflammatory biomarkers for cystic fibrosis drug development. 2021 , 20, 50-56	Ο
99	Role of the COX2-PGE axis in -induced exacerbation of experimental fibrosis. 2021, 320, L377-L392	2
98	Drug development for cystic fibrosis. 2021 , 56 Suppl 1, S10-S22	1
97	Novel Immunomodulatory Therapies for Respiratory Pathologies. 2021,	
96	Ibuprofen-based advanced therapeutics: breaking the inflammatory link in cancer, neurodegeneration, and diseases. 2021 , 53, 100-121	2
95	Untapped Potential: Therapeutically Targeting Eicosanoids and Endocannabinoids in the Lung. 2021 , 110, 69-81	1

94	Dysfunctional Inflammation in Cystic Fibrosis Airways: From Mechanisms to Novel Therapeutic Approaches. 2021 , 22,	3
93	Cystic Fibrosis Lung Disease in the Aging Population. 2021 , 12, 601438	3
92	Fibrotic expression profile analysis reveals repurposed drugs with potential anti-fibrotic mode of action. 2021 , 16, e0249687	2
91	Treatment of Pulmonary Disease of Cystic Fibrosis: A Comprehensive Review. 2021 , 10,	3
90	Antimicrobial Effects of Ibuprofen Combined with Standard of Care Antimicrobials against Cystic Fibrosis Pathogens.	
89	Tobramycin and Colistin display anti-inflammatory properties in CuFi-1 cystic fibrosis cell line. 2021 , 902, 174098	O
88	The indispensable role of pharmacy services and medication therapy management in cystic fibrosis. 2021 ,	
87	Reversal of SARS-CoV2-Induced Hypoxia by Nebulized Sodium Ibuprofenate in a Compassionate Use Program. 2021 , 10, 2511-2524	2
86	Empire-CF study: A phase 2 clinical trial of leukotriene A4 hydrolase inhibitor acebilustat in adult subjects with cystic fibrosis. 2021 , 20, 1026-1034	2
85	Current and Emerging Therapies to Combat Cystic Fibrosis Lung Infections. 2021 , 9,	2
84	Serious infectious events and ibuprofen administration in pediatrics: a narrative review in the era of COVID-19 pandemic. 2021 , 47, 20	4
83	Platelets in Pulmonary Vascular Physiology and Pathology. 2011 , 371-384	1
82	Recombinant Human Deoxyribonuclease I. 2013 , 321-336	4
81	Cystic Fibrosis. 1998 , 329-338	1
80	Integrative Therapies for People with Cystic Fibrosis. 2012 , 113-126	3
79	Therapeutic Approaches to the Lung Problems in Cystic Fibrosis. 1997 , 341-364	1
78	Nasal Polyps in Cystic Fibrosis. 2010 , 145-152	1
77	Mechanisms Versus Causes in Biology and Medicine. 2013 , 19-34	10

(2021-1997)

76	Therapeutic drug monitoring by capillary electrophoresis. 1997 , 386-408	3
75	Pulmonary Disease in Cystic Fibrosis. 2006 , 873-886	3
74	Cystic Fibrosis. 2010 , 985-1022	6
73	Bronchiectasis. 2010 , 1023-1046	2
72	Cystic Fibrosis. 2015 , 874-885.e3	1
71	Malnutrition in children with cystic fibrosis: the energy-balance equation. 1997 , 25, 127-36	22
70	Diffuse microcystic pancreatic enlargement in a cystic fibrosis patient causing severe gastrointestinal symptoms and successfully treated by total pancreatectomy. 1998 , 26, 454-7	1
69	The effect of intestinal permeability on pancreatic enzyme-induced enteropathy in the rat. 1998 , 26, 489-95	15
68	The effects of high-dose ibuprofen and pancreatic enzymes on the intestine of the rat. 1999 , 29, 178-83	5
67	Role of Inflammasome-independent Activation of IL-1lby the Pseudomonas aeruginosa Protease LasB.	2
66	Pseudomonas aeruginosa Infections. 305-326	5
65	Cystic fibrosis transmembrane conductance regulator dysfunction in platelets drives lung hyperinflammation. 2020 , 130, 2041-2053	20
64	Ibuprofen inhibits cystic fibrosis transmembrane conductance regulator-mediated Cl- secretion. 1998 , 102, 679-87	29
63	Taxonomy of Mucoactive Medications. 2004 , 129-145	5
62	Identifying treatments that halt progression of pulmonary disease in cystic fibrosis. 1997 , 41, 161-5	69
61	Cyclooxygenase-2 promotes ovarian cancer cell migration and cisplatin resistance via regulating epithelial mesenchymal transition. 2020 , 21, 315-326	4
60	New developments in cystic fibrosis. 1998 , 27, 515-20	8
59	Anti-Inflammatory Therapy of Infections. 2021,	

58	Increasing life expectancy in cystic fibrosis: Advances and challenges. 2021,	4
57	Myeloperoxidase Activity of Neutrophils in Cystic Fibrosis. 2000 , 107-113	
56	Serum Drug Monitoring by Capillary Electrophoresis. 2001 , 355-383	1
55	Atemwegserkrankung. 2001 , 265-366	
54	Infektionen der unteren Atemwege. 2004 , 243-300	
53	Zystische Fibrose. 2004 , 966-996	
52	Cystic Fibrosis. 2005 , 116-130	1
51	Cystic Fibrosis. 2007 , 494-501	
50	Infectious Complications in Special Hosts. 2008 , 633-641	
49	Thoracic Manifestations of Systemic Diseases. 2008 , 289-312	
48	CYSTIC FIBROSIS. 2009 , 342-364	
47	Cystic Fibrosis. 2010 , 947-955	1
46	Cystic Fibrosis. 2010 , 339-368	
45	Cystic Fibrosis. 2010 , 1037-1044.e2	
44	The Respiratory System. 2012 , 321-394	
43	Biodisposition in Relation to Actions. 2012, 5-42	
42	Infectious Complications in Special Hosts. 2012 , 633-641.e4	
41	Cystic Fibrosis Needs Attention in Iran. 2013 , 2, 107-8	

40	The many uses of ibuprofen. 1996 , 21, 56	
39	The General Approach to Cystic Fibrosis-Related Pulmonary Infection in the United States. 1996 , 99-110	
38	Mucoactive Agents: Old and New. 1997 , 155-179	2
37	Anti-inflammatory Therapies for Chronic Lung Diseases in Children. 1997 , 113-136	
36	Cystic Fibrosis. 1998 , 223-232	
35	Obstructive Pulmonary Diseases. 1999 , 185-225	
34	Rational treatment of pulmonary infections in patients with cystic fibrosis. 1999 , 12, 257-63	1
33	Human Toxicity of Ibuprofen. 498-517	
32	Clinical Application of Stem/Stromal Cells in Cystic Fibrosis. 2019 , 179-198	
31	CYSTIC FIBROSIS: A BREAKTHROUGH IN 21ST-CENTURY THERAPY. 2020 , 23, 35-41	
30	Drug Repurposing Through a Bioinformatics Pipeline Applied on Fibrotic Diseases.	
29	Inflammation in CF: Key Characteristics and Therapeutic Discovery. 2020 , 161-179	1
28	Targeting cystic fibrosis inflammation in the age of CFTR modulators: focus on macrophages. 2021 , 57,	2
27	Cystic fibrosis. 2020 , 4151-4165	
26	Current evidence on the effect of highly effective CFTR modulation on interleukin-8 in cystic fibrosis. 2021 , 1-14	О
25	Cystic Fibrosis. 2006 , 251-258	
24	Role of corticosteroids in cystic fibrosis lung disease. 1996 , 89 Suppl 27, 8-13	4
23	Liver transplantation for hepatic cirrhosis in cystic fibrosis. 1996 , 89 Suppl 27, 31-7	9

22	Cystic fibrosis in adults. From researcher to practitioner. 1996 , 164, 321-34	6
21	Understanding non-compliance with treatment in adults with cystic fibrosis. 2000 , 93 Suppl 38, 2-8	12
20	New therapeutic approaches for cystic fibrosis lung disease. 2002 , 95 Suppl 41, 58-67	1
19	Recent Progress on Ultra-Small Nanostructure Materials as Fluorescent Nanosensors for Monitoring of Various Molecules in Environmental Samples.	
18	Xylitol as a Hydrophilization Moiety for a Biocatalytically Synthesized Ibuprofen Prodrug 2022, 23,	1
17	Ultrasound-Assisted Slug-Flow Tubular Crystallization for Preparation of Fine Ibuprofen Crystals.	O
16	The Effect of CFTR Modulators on Airway Infection in Cystic Fibrosis 2022, 23,	2
15	Cystic Fibrosis Mucus Model to Design More Efficient Drug Therapies 2021 ,	3
14	Losartan ameliorates TGF-11-induced CFTR dysfunction and improves correction by cystic fibrosis modulator therapies 2022 ,	1
13	Buprofenin FarmasBik Preperatlarda UPLC-MS/MS YBtemiyle Tayini.	
12	TREATMENT OF STEROID-RESISTANT ASTHMA. 1996 , 16, 777-796	
11	Is CF airway inflammation still relevant in the era of highly effective modulators?. 2022,	
10	Systemic Corticosteroids in the Management of Pediatric Cystic Fibrosis Pulmonary Exacerbations.	О
9	Cystic Fibrosis. 2022 ,	1
8	Novel Applications of Biomarkers and Personalized Medicine in Cystic Fibrosis. 2022, 43, 617-630	О
7	Infectious Complications in Special Hosts. 2023 , 665-673.e4	0
6	Advances in Ultra-small Fluorescence Nanoprobes for Detection of Metal Ions, Drugs, Pesticides and Biomarkers.	1
5	Lessons from other fields of medicine, Part 2: Cystic fibrosis. 2023 , 119-130	O

CITATION REPORT

4	Diagnosis and Management of Cystic Fibrosis Exacerbations. 2023 , 44, 225-241	Ο
3	Preparation and Characterization of Ibuprofen Containing Nano-Embedded-Microparticles for Pulmonary Delivery. 2023 , 15, 545	O
2	High efficiency removal of ibuprofen in water using activated carbon derived from Radix Angelica Dahurica residue.	0
1	Inflammation as a Regulator of the Airway Surface Liquid pH in Cystic Fibrosis. 2023, 12, 1104	O