

Mucoid *Pseudomonas aeruginosa* is a marker of poor su

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Nosocomially Acquired <i>Pseudomonas cepacia</i> Infection in Patients with Cystic Fibrosis. <i>Infection Control and Hospital Epidemiology</i> , 1993, 14, 124-126.	1.0	8
2	Nosocomially Acquired <i>Pseudomonas cepacia</i> Infection in Patients with Cystic Fibrosis. <i>Infection Control and Hospital Epidemiology</i> , 1993, 14, 124-126.	1.0	15
3	Possible Nosocomial Transmission of <i>Pseudomonas cepacia</i> in Patients With Cystic Fibrosis. <i>JAMA Pediatrics</i> , 1994, 148, 805.	3.6	34
4	Mechanism of Action of Antibiotics in Chronic Pulmonary <i>Pseudomonas</i> Infection. <i>Advances in Pharmacology</i> , 1994, 30, 53-84.	1.2	5
5	Gender differences in cystic fibrosis: <i>Pseudomonas aeruginosa</i> infection. <i>Journal of Clinical Epidemiology</i> , 1995, 48, 1041-1049.	2.4	263
6	Cystic fibrosis.. <i>American Journal of Respiratory and Critical Care Medicine</i> , 1996, 154, 1229-1256.	2.5	859
7	The prognosis of cystic fibrosis in the Western Cape region of South Africa. <i>Journal of Paediatrics and Child Health</i> , 1996, 32, 323-326.	0.4	7
8	Potential of preventing <i>Pseudomonas aeruginosa</i> lung infections in cystic fibrosis patients: Experimental studies in animals. <i>Apmis</i> , 1996, 104, 5-42.	0.9	18
9	A prognostic model for the prediction of survival in cystic fibrosis. <i>Thorax</i> , 1997, 52, 313-317.	2.7	72
10	<i>Pseudomonas aeruginosa</i> : Assessment of Risk from Drinking Water. <i>Critical Reviews in Microbiology</i> , 1997, 23, 47-75.	2.7	177
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16	Antipseudomonal antibiotics and cystic fibrosis. <i>Australian and New Zealand Journal of Medicine</i> , 1999, 29, 5-8.	0.5	1
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18	Changing Epidemiology of <i>Pseudomonas aeruginosa</i> Infection in Danish Cystic Fibrosis Patients (1974-1995)., 1999, 28, 159-166.		161
19	Management of Patients with Cystic Fibrosis. <i>Disease Management and Health Outcomes</i> , 1999, 6, 93-108.	0.3	11

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21	Establishment of <i>Pseudomonas aeruginosa</i> infection: lessons from a versatile opportunist ¹ *Address for correspondence: Channing Laboratory, 181 Longwood Avenue, Boston, MA 02115, USA. <i>Microbes and Infection</i> , 2000, 2, 1051-1060.	1.0	1,191
22	Secondary genetic factors in cystic fibrosis lung disease. <i>Thorax</i> , 2000, 55, 446-446.	2.7	9
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32	A randomised clinical trial of nebulised tobramycin or colistin in cystic fibrosis. <i>European Respiratory Journal</i> , 2002, 20, 658-664.	3.1	251
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40	The Role of Inflammation in the Pathophysiology of CF Lung Disease. <i>Clinical Reviews in Allergy and Immunology</i> , 2002, 23, 005-028.	2.9	214
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71	Inhaled Tobramycin in Non-”Cystic Fibrosis Patients with Bronchiectasis and Chronic Bronchial Infection with <i>Pseudomonas Aeruginosa</i> . <i>Annals of Pharmacotherapy</i> , 2005, 39, 39-44.	0.9	201
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134	Measuring and improving respiratory outcomes in cystic fibrosis lung disease: Opportunities and challenges to therapy. <i>Journal of Cystic Fibrosis</i> , 2010, 9, 1-16.	0.3	93
135	An aerobiological model of aerosol survival of different strains of <i>Pseudomonas aeruginosa</i> isolated from people with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2010, 9, 64-68.	0.3	21
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