

CITATION REPORT

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Hydroxychloroquine and surfactant protein C deficiency

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#	Paper	IF	Citations
70	Defects in surfactant synthesis: clinical implications. <i>Pediatric Clinics of North America</i> , 2006 , 53, 911-27, ix	3.6	8
69	Chloroquine normalizes aberrant transforming growth factor beta activity in cystic fibrosis bronchial epithelial cells. <i>Pediatric Pulmonology</i> , 2006 , 41, 771-8	3.5	14
68	Neonatal respiratory failure associated with mutation in the surfactant protein C gene. <i>Journal of Perinatology</i> , 2006 , 26, 67-70	3.1	37
67	Inherited surfactant protein-B deficiency and surfactant protein-C associated disease: clinical features and evaluation. <i>Seminars in Perinatology</i> , 2006 , 30, 316-26	3.3	60
66	Genetically engineered mice in understanding the basis of neonatal lung disease. <i>Seminars in Perinatology</i> , 2006 , 30, 341-9	3.3	6
65	Genetic disorders of surfactant proteins. <i>Neonatology</i> , 2007 , 91, 311-7	4	98
64	What's new in surfactant? A clinical view on recent developments in neonatology and paediatrics. <i>European Journal of Pediatrics</i> , 2007 , 166, 889-99	4.1	21
63	[Lung diseases associated with inherited disorders of surfactant metabolism]. <i>Archives De Pediatrie</i> , 2008 , 15, 1560-7	1.8	2
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61	Inherited Surfactant Disorders. <i>NeoReviews</i> , 2008 , 9, e458-e467	1.1	9
60	Chronic lung disease after premature birth. <i>New England Journal of Medicine</i> , 2008 , 358, 743-5; author reply 745-6	59.2	45
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57	Characteristics of disorders associated with genetic mutations of surfactant protein C. <i>Archives of Disease in Childhood</i> , 2010 , 95, 449-54	2.2	88
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54	Infants and Young Children with Children's Interstitial Lung Disease. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2010 , 23, 25-31	0.8	55

53	Surfactant dysfunction. <i>Paediatric Respiratory Reviews</i> , 2011 , 12, 223-9	4.8	61
52	Diagnosis and management of diffuse lung disease in children. <i>Paediatric Respiratory Reviews</i> , 2011 , 12, 238-42	4.8	27
51	Surfactant protein C G100S mutation causes familial pulmonary fibrosis in Japanese kindred. <i>European Respiratory Journal</i> , 2011 , 38, 861-9	13.6	63
50	Interstitial Lung Disease in Infants and Children: New Classification System with Emphasis on Clinical, Imaging, and Pathological Correlation. 2012 , 99-154		2
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48	Neumopatías intersticiales. <i>Anales De Pediatría Continuada</i> , 2012 , 10, 87-94		
47	[Genetic disorders of surfactant]. <i>Archives De Pédiatrie</i> , 2012 , 19, 212-9	1.8	3
46	The surfactant protein C mutation A116D alters cellular processing, stress tolerance, surfactant lipid composition, and immune cell activation. <i>BMC Pulmonary Medicine</i> , 2012 , 12, 15	3.5	19
45	Imaging in Pediatric Pulmonology. 2012 ,		3
44	Diffuse Lung Disease. 2012 ,		2
43	Lung Diseases Associated with Disruption of Pulmonary Surfactant Homeostasis. 2012 , 810-821		
42	A novel mutation in surfactant protein-B gene resulting in fatal neonatal respiratory distress in an Indian family. <i>Journal of Neonatal-Perinatal Medicine</i> , 2012 , 5, 183-187	1.3	
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17	Recurrent diffuse lung disease due to surfactant protein C deficiency. <i>Respiratory Medicine Case Reports</i> , 2018 , 25, 91-95	1.2	10
16	Lung Diseases Associated With Disruption of Pulmonary Surfactant Homeostasis. 2019 , 836-849.e5		
15	Surfactant protein C dysfunction with new clinical insights for diffuse alveolar hemorrhage and autoimmunity. <i>Pediatric Investigation</i> , 2019 , 3, 201-206	1.3	0
14	Alveolar Dynamics and Beyond - The Importance of Surfactant Protein C and Cholesterol in Lung Homeostasis and Fibrosis. <i>Frontiers in Physiology</i> , 2020 , 11, 386	4.6	4
13	Methylprednisolone pulse treatment improves ProSP-C trafficking in twins with SFTPC mutation: An isoform story?. <i>British Journal of Clinical Pharmacology</i> , 2021 , 87, 2361-2373	3.8	1
12	Hydroxychloroquine, a successful treatment for lung disease in ABCA3 deficiency gene mutation: a case report. <i>Journal of Medical Case Reports</i> , 2021 , 15, 54	1.2	4
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