## CITATION REPORT List of articles citing

Initial combination therapy with ambrisentan and tadalafil in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH): subgroup analysis from the AMBITION trial

DOI: 10.1136/annrheumdis-2016-210236 Annals of the Rheumatic Diseases, 2017, 76, 1219-1227.

Source: https://exaly.com/paper-pdf/66044731/citation-report.pdf

Version: 2024-04-23

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
120	Ambrisentan: a review of its use in pulmonary arterial hypertension. <b>2017</b> , 11, 233-244		12
119	Pulmonary arterial hypertension: screening challenges in systemic sclerosis and future directions. <b>2017</b> , 49,		3
118	Epidemiology and treatment of pulmonary arterial hypertension. <b>2017</b> , 14, 603-614		182
117	Pulmonary hypertension in systemic sclerosis: different phenotypes. <b>2017</b> , 26,		59
116	New insights into the recognition, classification and management of systemic sclerosis-associated pulmonary hypertension. <b>2017</b> , 29, 561-567		3
115	Selexipag for the treatment of connective tissue disease-associated pulmonary arterial hypertension. <b>2017</b> , 50,		58
114	Early intervention in the management of pulmonary arterial hypertension: clinical and economic outcomes. <b>2017</b> , 9, 731-739		13
113	Can cardiovascular magnetic resonance prompt early cardiovascular/rheumatic treatment in autoimmune rheumatic diseases? Current practice and future perspectives. <b>2018</b> , 38, 949-958		15
112	An Update on Systemic Sclerosis-Associated Pulmonary Arterial Hypertension: a Review of the Current Literature. <b>2018</b> , 20, 10		15
111	[Pulmonary hypertension in connective tissue disease]. <b>2018</b> , 77, 219-230		
110	Safety and feasibility audit of a home-based drug-transitioning approach for patients with pulmonary arterial hypertension: an observational study. <b>2018</b> , 17, 612-618		3
109	Pulmonary arterial hypertension associated with connective tissue diseases: A review focusing on distinctive clinical aspects. <b>2018</b> , 48, e12876		17
108	Initial tadalafil and ambrisentan combination therapy in pulmonary arterial hypertension: cLinical and haemodYnamic long-term efficacy (ITALY study). <b>2018</b> , 19, 12-17		15
107	Predictors of Favorable Responses to Immunosuppressive Treatment in Pulmonary Arterial Hypertension Associated With Connective Tissue Disease. <b>2018</b> , 82, 546-554		23
106	Pulmonary hypertension in patients with interstitial lung disease. <b>2018</b> , 50, 38-46		13
105	Improvement in Right Ventricular Strain with Ambrisentan and Tadalafil Upfront Therapy in Scleroderma-associated Pulmonary Arterial Hypertension. <b>2018</b> , 197, 388-391		24
104	Prevalence of pulmonary hypertension in patients with systemic sclerosis and mixed connective tissue disease. <b>2018</b> , 97, e11437		8

103	Risk assessment in scleroderma patients with newly diagnosed pulmonary arterial hypertension: application of the ESC/ERS risk prediction model. <b>2018</b> , 52,	17
102	Endothelin-receptor antagonists in the management of pulmonary arterial hypertension: where do we stand?. <b>2018</b> , 14, 253-264	20
101	Treatment Algorithms for Systemic Sclerosis According to Experts. <b>2018</b> , 70, 1820-1828	120
100	Long-Term Outcomes in Systemic Sclerosis-Associated Pulmonary Arterial Hypertension From the Pulmonary Hypertension Assessment and Recognition of Outcomes in Scleroderma Registry (PHAROS). <b>2018</b> , 154, 862-871	47
99	Can treprostinil-induced early gastrointestinal side effects serve as predictors of pulmonary arterial hypertension prognosis?. <b>2018</b> , 264, 187	
98	Ambrisentan – tadalafil in WHO functional class II/III pulmonary arterial hypertension: a guide to its use in the EU. <b>2018</b> , 34, 289-299	
97	Aggressive combination therapy for treatment of systemic sclerosis-associated pulmonary hypertension <b>2018</b> , 3, 30-38	2
96	Lung Involvements in Rheumatic Diseases: Update on the Epidemiology, Pathogenesis, Clinical Features, and Treatment. <b>2018</b> , 2018, 6930297	22
95	Pulmonary Circulation on the Crossroads Between the Left and Right Heart in Systemic Sclerosis: A Clinical Challenge for Cardiologists and Rheumatologists. <b>2018</b> , 14, 271-281	8
94	[What rheumatologists can learn from pneumologists]. 2018, 77, 477-483	1
93	Hypertension pulmonaire et connectivites. 2018, 85, 210-220	
92	Major lung complications of systemic sclerosis. <b>2018</b> , 14, 511-527	43
91	Pulmonary Manifestations of Systemic Sclerosis and Mixed Connective Tissue Disease. <b>2019</b> , 40, 501-518	13
90	Screening for pulmonary arterial hypertension in systemic sclerosis. <b>2019</b> , 28,	21
89	A multicenter randomized, double-blind, placebo-controlled pilot study to assess the efficacy and safety of riociguat in systemic sclerosis-associated digital ulcers. <b>2019</b> , 21, 202	10
88	Advances in Management of Pulmonary Hypertension Associated with Systemic Sclerosis. 2019,	1
87	Pulmonary arterial hypertension in connective tissue disorders: Pathophysiology and treatment. <b>2019</b> , 244, 120-131	16
86	Pulmonary Hypertension Associated with Connective Tissue Disease. <b>2019</b> , 40, 173-183	15

85	Validation of the REVEAL Prognostic Equation and Risk Score Calculator in Incident Systemic Sclerosis-Associated Pulmonary Arterial Hypertension. <b>2019</b> , 71, 1691-1700	7
84	Guidelines for the Treatment of Pulmonary Hypertension (JCS 2017/JPCPHS 2017). <b>2019</b> , 83, 842-945	67
83	OP0067 UTILITY OF RISK STRATIFICATION IN PREDICTING OUTCOMES OF INITIAL MONOTHERAPY VERSUS COMBINATION THERAPY IN PULMONARY ARTERIAL HYPERTENSION ASSOCIATED WITH CONNECTIVE TISSUE DISEASE: A POST-HOC ANALYSIS OF THE AMBITION STUDY. <b>2019</b> ,	
82	OP0068 ABATACEPT IN EARLY DIFFUSE CUTANEOUS SYSTEMIC SCLEROSISIRESULTS OF A PHASE 2 INVESTIGATOR-INITIATED, MULTICENTER, DOUBLE-BLIND RANDOMIZED PLACEBO-CONTROLLED TRIAL. <b>2019</b> ,	1
81	Therapeutic Challenges And Advances In The Management Of Systemic Sclerosis-Related Pulmonary Arterial Hypertension (SSc-PAH). <b>2019</b> , 15, 1427-1442	6
80	Pulmonary Arterial Hypertension In Systemic Sclerosis: Challenges In Diagnosis, Screening And Treatment. <b>2019</b> , 11, 323-333	8
79	Interstitial Lung Disease and Other Pulmonary Manifestations in Connective Tissue Diseases. <b>2019</b> , 94, 309-325	37
78	Pulmonary arterial hypertension: In Asia, as elsewhere, still a lethal disease despite modern treatment. <b>2019</b> , 24, 99-100	1
77	Risk Factors for Mortality and Cardiopulmonary Hospitalization in Systemic Sclerosis Patients At Risk for Pulmonary Hypertension, in the PHAROS Registry. <b>2019</b> , 46, 176-183	14
76	Longterm Efficacy and Safety of Monotherapy versus Combination Therapy in Systemic Sclerosis-associated Pulmonary Arterial Hypertension: A Retrospective RESCLE Registry Study. <b>2020</b> , 47, 89-98	7
75	Systemic sclerosis: Recent insight in clinical management. <b>2020</b> , 87, 293-299	18
74	Combination therapy with bosentan and sildenafil for refractory digital ulcers and Raynaud's phenomenon in a 30-year-old woman with systemic sclerosis: Case report and literature review <b>2020</b> , 5, 159-164	
73	Survival Improved in Patients Aged 170 Years With Systemic Sclerosis-Associated Pulmonary Arterial Hypertension During the Period 2006 to 2017 in France. <b>2020</b> , 157, 945-954	5
72	Mortality and hospitalization outcomes of interstitial lung disease and pulmonary hypertension in the Singapore systemic sclerosis cohort. <b>2020</b> , 50, 473-479	8
71	Patient-reported outcome instruments in clinical trials of systemic sclerosis <b>2020</b> , 5, 90-102	8
70	Hemodynamics and risk assessment 2 years after the initiation of upfront ambrisentan-tadalafil in pulmonary arterial hypertension. <b>2020</b> , 39, 1389-1397	3
69	Effectiveness and safety of endothelin receptor antagonists, alone and in combination therapy, in the pulmonary arterial hypertension-connective tissue disease subtype: A systematic review and meta-analysis. <b>2020</b> , 23, 1276-1287	1
68	Efficacy and safety of ambrisentan in Chinese patients with connective tissue disease-pulmonary arterial hypertension: a post-hoc analysis. <b>2020</b> , 20, 339	O

## (2021-2020)

67	Diagnostic and prognostic markers and treatment of connective tissue disease-associated pulmonary arterial hypertension: current recommendations and recent advances. <b>2020</b> , 16, 993-1004	4
66	Effect of ambrisentan on echocardiographic and Doppler measures from patients in China with pulmonary arterial hypertension. <b>2020</b> , 18, 643-649	O
65	Pharmacological treatment of systemic sclerosis-associated pulmonary hypertension: A systematic literature review. <b>2020</b> , 27, 135-145	
64	Screening for pulmonary arterial hypertension in systemic sclerosis: A systematic literature review. <b>2020</b> , 78, 17-25	13
63	Randomised controlled trials in systemic sclerosis: patient selection and endpoints for next generation trials. <b>2020</b> , 2, e173-e184	8
62	Management of systemic sclerosis: the first five years. <b>2020</b> , 32, 228-237	15
61	Pulmonary Hypertension Phenotypes in Systemic Sclerosis: The Right Diagnosis for the Right Treatment. <b>2020</b> , 21,	8
60	A case report, a case who developed limited cutaneous scleroderma and pulmonary hypertension 8 years after diagnosis of anti-centromere antibody-positive Sjören syndrome. <b>2020</b> , 4, 248-252	1
59	The future of treatment in systemic sclerosis: can we design better trials?. <b>2020</b> , 2, e185-e194	8
58	Challenges in evidence-based therapy for systemic sclerosis associated interstitial lung disease. <b>2020</b> , 8, 226-227	O
57	The Effect of Grapefruit Juice on the Pharmacokinetics of Tadalafil in Rats. 2020, 2020, 1631735	5
56	Systemic sclerosis: Advances towards stratified medicine. <b>2020</b> , 34, 101469	3
55	Management of Pulmonary Arterial Hypertension in Patients with Systemic Sclerosis. <b>2020</b> , 13, 15-29	13
54	Initial combination therapy of ambrisentan and tadalafil in connective tissue disease-associated pulmonary arterial hypertension (CTD-PAH) in the modified intention-to-treat population of the 2.4 AMBITION study: post hoc analysis. <i>Annals of the Rheumatic Diseases</i> , <b>2020</b> , 79, 626-634	11
53	Donnês rêentes sur la prise en charge clinique de la sclêodermie systênique. <b>2021</b> , 88, 24-31	
52	Pulmonary hypertension in connective tissue diseases, new evidence and challenges. <b>2021</b> , 51, e13453	9
51	Hemodynamic Response to Treatment and Outcomes in Pulmonary Hypertension Associated With Interstitial Lung Disease Versus Pulmonary Arterial Hypertension in Systemic Sclerosis: Data From a Study Identifying Prognostic Factors in Pulmonary Hypertension Associated With Interstitial Lung	10
50	Disease. <b>2021</b> , 73, 295-304 Risk Reduction and Hemodynamics with Initial Combination Therapy in Pulmonary Arterial Hypertension. <b>2021</b> , 203, 484-492	12

49	The effects of oral treatment for systemic sclerosis related pulmonary arterial hypertension: A systematic review and meta-analysis. <b>2021</b> , 31, 151-161	2
48	Clinical Treatment Options in Scleroderma: Recommendations and Comprehensive Review. <b>2021</b> , 1	9
47	The multifaceted problem of pulmonary arterial hypertension in systemic sclerosis. 2021, 3, e149-e159	4
46	Performance Under Pressure: The Relevance of Pulmonary Vascular Response to Exercise Challenge in Scleroderma. <b>2021</b> , 159, 481-483	1
45	Recent advances in the management of pulmonary arterial hypertension: lessons from the upfront combination of ambrisentan and tadalafil. <b>2021</b> , 15, 493-504	2
44	Group 3 Pulmonary Hypertension: A Review of Diagnostics and Clinical Trials. <b>2021</b> , 42, 59-70	3
43	Pharmacological Interventions for Pulmonary Involvement in Rheumatic Diseases. <b>2021</b> , 14,	2
42	Long-Term Outcomes in Patients With Connective Tissue Disease-Associated Pulmonary Arterial Hypertension in the Modern Treatment Era: Meta-Analyses of Randomized, Controlled Trials and Observational Registries. <b>2021</b> , 73, 837-847	9
41	Combining Data Sets as Well as Therapies Shows Improved Outcome in Connective Tissue Disease-Associated Pulmonary Hypertension. <b>2021</b> , 73, 725-727	1
40	Pulmonary arterial hypertension in systemic sclerosis. <b>2021</b> , 50, 104062	2
39	Clinical characteristics and survival of patients with three major connective tissue diseases associated with pulmonary hypertension: A study from China. <b>2021</b> , 22, 925	2
38	An update on targeted therapies in systemic sclerosis based on a systematic review from the last 3 years. <b>2021</b> , 23, 155	8
37	Recognition and Management of Cutaneous Connective Tissue Diseases. <b>2021</b> , 105, 757-782	O
36	The Role of Exercise Doppler Echocardiography to Unmask Pulmonary Arterial Hypertension in Selected Patients with Systemic Sclerosis and Equivocal Baseline Echocardiographic Values for Pulmonary Hypertension. <b>2021</b> , 11,	3
35	[Characteristics of patients with connective tissue disease-associated pulmonary arterial hypertension treated with prostanoids: A multicenter retrospective study]. <b>2021</b> , 42, 825-831	
34	Systemic sclerosis-associated pulmonary arterial hypertension in children. <b>2021</b> , 11, 1137-1143	Ο
33	Pulmonary hypertension phenotypes in patients with systemic sclerosis. <b>2021</b> , 30,	5
32	Low-Dose Dobutamine Stress Echocardiography for the Early Detection of Pulmonary Arterial Hypertension in Selected Patients with Systemic Sclerosis Whose Resting Echocardiography Is 5.2 Non-Diagnostic for Pulmonary Hypertension. <i>Journal of Clinical Medicine</i> , <b>2021</b> , 10,	0 1

31	Sudden Cardiac Death in Systemic Sclerosis: Diagnostics to Assess Risk and Inform Management. <b>2021</b> , 11,		О
30	The role of pulmonary arterial hypertension-targeted therapy in systemic sclerosis. <b>2019</b> , 8,		3
29	New Drugs, Therapeutic Strategies, and Future Direction for the Treatment of Pulmonary Arterial Hypertension. <b>2019</b> , 26, 2844-2864		11
28	Systemic Sclerosis and Other Connective Tissue Diseases. <b>2017</b> , 16, 55-60		
27	Gaps and Controversies of New Treatment Recommendations in Recent Pulmonary Hypertension Guidelines: What We Know and What We Don't. <b>2017</b> , 16, 20-25		
26	III. The Particularity of CTD-PH Management. <b>2018</b> , 107, 214-218		
25	Pulmonary involvement in systemic lupus erythematosus, Sjgren syndrome and mixed connective tissue disease. <b>2019</b> , 106-123		1
24	TĒkiyeĒle tersiyer bir merkezde sistemik skleroz hastalar⊞nda pulmoner hipertansiyon taramas⊞; kesitsel orjinal Ēl⊞Ēha.		
23	BOEHRINGER INGELHEIM - Fibroses pulmonaires : maladies pulmonaires ou maladies systEhiques ?. <b>2020</b> , 12, S58-S63		
22	Beyond Scleroderma: Pulmonary Arterial Hypertension in Patients with Other Connective Tissue Diseases. <b>2020</b> , 51-60		
21	Tratamiento farmacolgico de la hipertensifi pulmonar asociada a la esclerosis sistfinica: revisifi sistemfica de la literatura. <b>2020</b> , 27, 135-145		
20	Pulmonary Hypertension Associated with Connective Tissue Disease. <i>Cardiology Clinics</i> , <b>2022</b> , 40, 29-43	2.5	1
19	2020 Clinical practice guidelines for Pulmonary hypertension, including chronic thromboembolic pulmonary hypertension. <i>Russian Journal of Cardiology</i> , <b>2022</b> , 26, 4683	1.3	9
18	Upfront Combination Therapy: Growing the Case to Get Ahead of Pediatric Pulmonary Arterial Hypertension <i>Annals of the American Thoracic Society</i> , <b>2022</b> , 19, 163-165	4.7	
17	Management of Endothelial Dysfunction in Systemic Sclerosis: Current and Developing Strategies <i>Frontiers in Medicine</i> , <b>2021</b> , 8, 788250	4.9	3
16	The treatment strategy of connective tissue disease associated pulmonary arterial hypertension: Evolving into the future <i>Pharmacology &amp; Therapeutics</i> , <b>2022</b> , 108192	13.9	1
15	Changes in the Characteristics and Initial Treatments of Pulmonary Hypertension Between 2008 and 2020 in Japan. <i>JACC Asia</i> , <b>2022</b> , 2, 273-284		0
14	Effect of vasodilator and immunosuppressive therapy on the endothelial dysfunction in patients with systemic sclerosis. <i>Clinical and Experimental Medicine</i> ,	4.9	1

13	Risk Stratification of Patients with Pulmonary Arterial Hypertension: The Role of Echocardiography. <i>Journal of Clinical Medicine</i> , <b>2022</b> , 11, 4034	5.1	O
12	New Era in Systemic Sclerosis Treatment: Recently Approved Therapeutics. <b>2022</b> , 11, 4631		О
11	Pulmonary Hypertension in SclerodermalEvaluation and Management. 2022, 101468		0
10	VIII. Progress in the Management of Pulmonary Arterial Hypertension Associated with Connective Tissue Disease. <b>2021</b> , 110, 2213-2220		O
9	Pulmonary Arterial Hypertension in Connective Tissue Diseases Beyond Systemic Sclerosis. <b>2023</b> , 19, 45-54		0
8	Systemic sclerosis. <b>2022</b> ,		2
7	Treatment of autoimmunity: The impact of disease-modifying therapies in multiple sclerosis and comorbid autoimmune disorders. <b>2023</b> , 22, 103312		О
6	Interstitial Lung Disease. <b>2023</b> , 49, 279-293		O
5	Pulmonary Hypertension. <b>2023</b> , 49, 345-357		О
4	State-of-the-art evidence in the treatment of systemic sclerosis. <b>2023</b> , 19, 212-226		O
3	Hospitalization Among Pulmonary Arterial Hypertension Patients With and Without Connective Tissue Disease Comorbidities Prescribed Oral Selexipag.		О
2	Economic burden of illness among patients with pulmonary arterial hypertension (PAH) associated with connective tissue disorders (CTD). <b>2023</b> , 13,		O
1	Lung Disease in Systemic Lupus Erythematosus, Myositis, Sjgren Disease, and Mixed Connective Tissue Disease. <b>2023</b> , 223-239		0