

CITATION REPORT

List of articles citing

Initial combination therapy with ambrisentan and tadalafil and mortality in patients with pulmonary arterial hypertension: a secondary analysis of the results from the randomised, controlled AMBITION study

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Lancet Respiratory Medicine, the, 2016, 4, 894-901.

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#	Paper	IF	Citations
52	Time to consider death in clinical trials for PAH. <i>Lancet Respiratory Medicine</i> , 2016 , 4, 849-850	35.1	1
51	Ambrisentan: a review of its use in pulmonary arterial hypertension. <i>Therapeutic Advances in Respiratory Disease</i> , 2017 , 11, 233-244	4.9	12
50	Pulmonary Hypertension-Back to the Future. <i>Revista Espanola De Cardiologia (English Ed)</i> , 2017 , 70, 901-904	0.4	1
49	Outcomes of pulmonary arterial hypertension therapy in Australia: is monotherapy adequate?. <i>Internal Medicine Journal</i> , 2017 , 47, 1124-1128	1.6	
48	Combination therapy in pulmonary arterial hypertension: recent accomplishments and future challenges. <i>Pulmonary Circulation</i> , 2017 , 7, 312-325	2.7	35
47	Hipertensi3 pulmonar. Regreso al futuro. <i>Revista Espanola De Cardiologia</i> , 2017 , 70, 901-904	1.5	2
46	Early intervention in the management of pulmonary arterial hypertension: clinical and economic outcomes. <i>ClinicoEconomics and Outcomes Research</i> , 2017 , 9, 731-739	1.7	13
45	The Low-Risk Profile in Pulmonary Arterial Hypertension. Time for a Paradigm Shift to Goal-oriented Clinical Trial Endpoints?. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 197, 860-868	10.2	27
44	Survival of Idiopathic Pulmonary Arterial Hypertension Patients in the Modern Era in Australia and New Zealand. <i>Heart Lung and Circulation</i> , 2018 , 27, 1368-1375	1.8	17
43	Update on pulmonary arterial hypertension research: proceedings from a meeting of experts. <i>Current Medical Research and Opinion</i> , 2018 , 34, 263-273	2.5	5
42	Room With a View. <i>Circulation: Cardiovascular Imaging</i> , 2018 , 11, e008148	3.9	1
41	Endothelin-receptor antagonists in the management of pulmonary arterial hypertension: where do we stand?. <i>Vascular Health and Risk Management</i> , 2018 , 14, 253-264	4.4	20
40	Targeted therapy of pulmonary arterial hypertension: Updated recommendations from the Cologne Consensus Conference 2018. <i>International Journal of Cardiology</i> , 2018 , 272S, 37-45	3.2	33
39	Ambrisentan + tadalafil in WHO functional class II/III pulmonary arterial hypertension: a guide to its use in the EU. <i>Drugs and Therapy Perspectives</i> , 2018 , 34, 289-299	1.5	
38	Choice of Initial Oral Therapy for Pulmonary Arterial Hypertension: Age and Long-Term Survival. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018 , 198, 1090-1093	10.2	2
37	Modern treatment to reduce pulmonary arterial pressure in pulmonary arterial hypertension. <i>Journal of Cardiology</i> , 2018 , 72, 466-472	3	24
36	Treatment of pediatric pulmonary arterial hypertension: A focus on the NO-sGC-cGMP pathway. <i>Pediatric Pulmonology</i> , 2019 , 54, 1516-1526	3.5	11

35	Selexipag in the management of pulmonary arterial hypertension: an update. <i>Drug, Healthcare and Patient Safety</i> , 2019 , 11, 55-64	1.6	3
34	Safety and Efficacy of Ambrisentan-Phosphodiesterase Type 5 (PDE5) Inhibitor Combination Therapy for Japanese Pulmonary Arterial Hypertension Patients in Real-World Clinical Practice. <i>Circulation Reports</i> , 2019 , 1, 268-275	0.7	1
33	Improving Survival in Patients with Pulmonary Arterial Hypertension: Focus on Intravenous Epoprostenol. <i>American Journal of Cardiovascular Drugs</i> , 2019 , 19, 99-105	4	5
32	A Phase I Study to Show the Relative Bioavailability and Bioequivalence of Fixed-Dose Combinations of Ambrisentan and Tadalafil in Healthy Subjects. <i>Clinical Therapeutics</i> , 2019 , 41, 1110-1127 ³⁵		1
31	Pulmonary arterial hypertension in systemic sclerosis: Diagnosis and treatment according to the European Society of Cardiology and European Respiratory Society 2015 guidelines.. <i>Journal of Scleroderma and Related Disorders</i> , 2019 , 4, 35-42	2.3	9
30	Survival Improved in Patients Aged \geq 70 Years With Systemic Sclerosis-Associated Pulmonary Arterial Hypertension During the Period 2006 to 2017 in France. <i>Chest</i> , 2020 , 157, 945-954	5.3	5
29	Combination Therapy with Oral Treprostinil for Pulmonary Arterial Hypertension. A Double-Blind Placebo-controlled Clinical Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020 , 201, 707-717	10.2	40
28	Pathological Mechanisms and Potential Therapeutic Targets of Pulmonary Arterial Hypertension: A Review. 2020 , 11, 1623-1639		9
27	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 AMBITION study: post hoc analysis. <i>Annals of the Rheumatic Diseases</i> , 2020 , 79, 626-634	2.4	11
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25	Recent advances in the management of pulmonary arterial hypertension: lessons from the upfront combination of ambrisentan and tadalafil. <i>Expert Review of Respiratory Medicine</i> , 2021 , 15, 493-504	3.8	2
24	Association between Initial Treatment Strategy and Long-Term Survival in Pulmonary Arterial Hypertension. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2021 , 204, 842-854	10.2	13
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21	Advances in the management of pulmonary arterial hypertension. <i>Journal of Investigative Medicine</i> , 2021 , 69, 1270-1280	2.9	4
20	How We Would Treat Our Own Pulmonary Hypertension if We Needed to Undergo Cardiac Surgery. <i>Journal of Cardiothoracic and Vascular Anesthesia</i> , 2021 ,	2.1	0
19	New drugs and emerging therapeutic targets in the endothelin signaling pathway and prospects for personalized precision medicine. <i>Physiological Research</i> , 2018 , 67, S37-S54	2.1	14
18	Temporal trends in pulmonary arterial hypertension: Results from the COMPERA registry. <i>European Respiratory Journal</i> , 2021 ,	13.6	6

17	Mortality trends in pulmonary arterial hypertension in canada: a temporal analysis of survival per ESC/ERS Guideline Era. <i>European Respiratory Journal</i> , 2021 ,	13.6	6
16	Positioning Newer Agents: Riociguat, Selexipag, and Oral Treprostinil in the Current Landscape. <i>Advances in Pulmonary Hypertension</i> , 2017 , 15, 193-197	0.5	
15	Positive Predictors for Response to Ambrisentan Combination Therapy in Pulmonary Arterial Hypertension.. <i>International Heart Journal</i> , 2022 , 63, 99-105	1.8	
14	The Transition From Ambrisentan to Macitentan in Patients With Pulmonary Arterial Hypertension: A Real-world Prospective Study.. <i>Frontiers in Pharmacology</i> , 2021 , 12, 811700	5.6	0
13	Dan-Shen-Yin Granules Prevent Hypoxia-Induced Pulmonary Hypertension STAT3/HIF-1 α /VEGF and FAK/AKT Signaling Pathways.. <i>Frontiers in Pharmacology</i> , 2022 , 13, 844400	5.6	2
12	Transdisciplinary Imagination: Addressing Equity and Mistreatment in Perinatal Care.. <i>Maternal and Child Health Journal</i> , 2022 ,	2.4	
11	To be or not to be treated with initial combination therapy, that is the (PAH) question. <i>European Respiratory Journal</i> , 2022 , 59, 2200390	13.6	
10	Profiles and treatment patterns of patients with pulmonary arterial hypertension on monotherapy at experienced centres. <i>ESC Heart Failure</i> ,	3.7	0
9	Prognostic impact of follow-up pulmonary vascular resistance in pulmonary arterial hypertension. <i>Open Heart</i> , 2022 , 9, e002054	3	
8	2022 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension. 2200879		18
7	2022 ESC/ERS Guidelines for the diagnosis and treatment of pulmonary hypertension.		58
6	Assessment of Clinical Worsening End Points as a Surrogate for Mortality in Pulmonary Arterial Hypertension: A Systematic Review and Meta-Analysis of Randomized Controlled Trials. 2022 , 146, 597-612		0
5	Selexipag-based triple combination therapy improves prognosis in Chinese pulmonary arterial hypertension patients. 9,		0
4	Efficacy and safety of switching from bosentan or ambrisentan to macitentan in pulmonary arterial hypertension: A systematic review and meta-analysis. 9,		0
3	Portopulmonary Hypertension Rethinking Our Current Approach. 2022 , 97, 2189-2191		0
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