Carmine Dario Vizza

List of Publications by Year in descending order

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181 papers 13,556 citations

50170 46 h-index 22102 113 g-index

192 all docs

192 docs citations

192 times ranked 9899 citing authors

#	Article	IF	CITATIONS
1	2019 ESC Guidelines for the diagnosis and management of acute pulmonary embolism developed in collaboration with the European Respiratory Society (ERS). European Heart Journal, 2020, 41, 543-603.	1.0	2,426
2	Macitentan and Morbidity and Mortality in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 809-818.	13.9	1,168
3	Riociguat for the Treatment of Pulmonary Arterial Hypertension. New England Journal of Medicine, 2013, 369, 330-340.	13.9	1,120
4	Initial Use of Ambrisentan plus Tadalafil in Pulmonary Arterial Hypertension. New England Journal of Medicine, 2015, 373, 834-844.	13.9	906
5	2019 ESC Guidelines for the diagnosis and management of acute pulmonary embolism developed in collaboration with the European Respiratory Society (ERS). European Respiratory Journal, 2019, 54, 1901647.	3.1	806
6	Effects of beraprost sodium, an oral prostacyclin analogue, in patients with pulmonary arterial hypertension: a randomized, double-blind, placebo-controlled trial. Journal of the American College of Cardiology, 2002, 39, 1496-1502.	1.2	584
7	Mortality in pulmonary arterial hypertension: prediction by the 2015 European pulmonary hypertension guidelines risk stratification model. European Respiratory Journal, 2017, 50, 1700740.	3.1	489
8	Elderly patients diagnosed with idiopathic pulmonary arterial hypertension: Results from the COMPERA registry. International Journal of Cardiology, 2013, 168, 871-880.	0.8	357
9	Anticoagulation and Survival in Pulmonary Arterial Hypertension. Circulation, 2014, 129, 57-65.	1.6	317
10	Right and Left Ventricular Dysfunction in Patients With Severe Pulmonary Disease. Chest, 1998, 113, 576-583.	0.4	290
11	Pre-Capillary, Combined, and Post-Capillary Pulmonary Hypertension. Journal of the American College of Cardiology, 2016, 68, 368-378.	1.2	244
12	An official European Respiratory Society statement: pulmonary haemodynamics during exercise. European Respiratory Journal, 2017, 50, 1700578.	3.1	222
13	Pregnancy outcomes in pulmonary arterial hypertension in the modern management era. European Respiratory Journal, 2012, 40, 881-885.	3.1	221
14	Prostanoid therapy for pulmonary arterial hypertension. Journal of the American College of Cardiology, 2004, 43, S56-S61.	1.2	184
15	Chronic Inhibition of cGMP Phosphodiesterase 5A Improves Diabetic Cardiomyopathy. Circulation, 2012, 125, 2323-2333.	1.6	171
16	Changes in Right Ventricular Function Measured by Cardiac Magnetic Resonance Imaging in Patients Receiving Pulmonary Arterial Hypertension–Targeted Therapy. Circulation: Cardiovascular Imaging, 2014, 7, 107-114.	1.3	139
17	Outcomes of noncardiac, nonobstetric surgery in patients with PAH: an international prospective survey. European Respiratory Journal, 2013, 41, 1302-1307.	3.1	131
18	COMPERA 2.0: a refined four-stratum risk assessment model for pulmonary arterial hypertension. European Respiratory Journal, 2022, 60, 2102311.	3.1	124

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19	RESPITE: switching to riociguat in pulmonary arterial hypertension patients with inadequate response to phosphodiesterase-5 inhibitors. European Respiratory Journal, 2017, 50, 1602425.	3.1	113
20	Riociguat for the treatment of pulmonary arterial hypertension associated with connective tissue disease: results from PATENT-1 and PATENT-2. Annals of the Rheumatic Diseases, 2017, 76, 422-426.	0.5	108
21	Idiopathic pulmonary arterial hypertension phenotypes determined by cluster analysis from the COMPERA registry. Journal of Heart and Lung Transplantation, 2020, 39, 1435-1444.	0.3	104
22	Long-term safety and efficacy of imatinib in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2015, 34, 1366-1375.	0.3	103
23	Risk Reduction and Right Heart Reverse Remodeling by Upfront Triple Combination Therapy in Pulmonary ArterialÂHypertension. Chest, 2020, 157, 376-383.	0.4	97
24	Sildenafil in severe pulmonary hypertension associated with chronic obstructive pulmonary disease: A randomized controlled multicenter clinical trial. Journal of Heart and Lung Transplantation, 2017, 36, 166-174.	0.3	89
25	Combination Therapy with Oral Treprostinil for Pulmonary Arterial Hypertension. A Double-Blind Placebo-controlled Clinical Trial. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 707-717.	2.5	89
26	Locoregional versus general anesthesia in carotid surgery: Is there an impact on perioperative myocardial ischemia? Results of a prospective monocentric randomized trial. Journal of Vascular Surgery, 1999, 30, 131-138.	0.6	87
27	Outcome of Patients with Cystic Fibrosis Awaiting Lung Transplantation. American Journal of Respiratory and Critical Care Medicine, 2000, 162, 819-825.	2.5	86
28	Switching to riociguat versus maintenance therapy with phosphodiesterase-5 inhibitors in patients with pulmonary arterial hypertension (REPLACE): a multicentre, open-label, randomised controlled trial. Lancet Respiratory Medicine, the, 2021, 9, 573-584.	5.2	85
29	Right Intraventricular Dyssynchrony in Idiopathic, Heritable, and Anorexigen-Induced Pulmonary Arterial Hypertension. JACC: Cardiovascular Imaging, 2015, 8, 642-652.	2.3	83
30	Pulmonary Hypertension in Patients with Chronic Fibrosing Idiopathic Interstitial Pneumonias. PLoS ONE, 2015, 10, e0141911.	1.1	80
31	Long term effects of bosentan treatment in adult patients with pulmonary arterial hypertension related to congenital heart disease (Eisenmenger physiology): safety, tolerability, clinical, and haemodynamic effect. Heart, 2007, 93, 621-625.	1.2	75
32	Acute hemodynamic effects of inhaled nitric oxide, dobutamine and a combination of the two in patients with mild to moderate secondary pulmonary hypertension. Critical Care, 2001, 5, 355.	2.5	67
33	Systemic sclerosis patients with and without pulmonary arterial hypertension: a nailfold capillaroscopy study. Rheumatology, 2013, 52, 1525-1528.	0.9	67
34	Treatment of pulmonary hypertension in patients undergoing cardiac surgery with cardiopulmonary bypass: a randomized, prospective, double-blind study. Journal of Cardiovascular Medicine, 2006, 7, 119-123.	0.6	66
35	Right ventricular remodeling in idiopathic pulmonary arterial hypertension: adaptive versus maladaptive morphology. Journal of Heart and Lung Transplantation, 2015, 34, 395-403.	0.3	66
36	Transesophageal dipyridamole echocardiography for diagnosis of coronary artery disease. Journal of the American College of Cardiology, 1992, 19, 765-770.	1.2	62

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37	Acute Hemodynamic Effects of Singleâ€Dose Sildenafil When Added to Established Bosentan Therapy in Patients With Pulmonary Arterial Hypertension: Results of the COMPASSâ€1 Study. Journal of Clinical Pharmacology, 2009, 49, 1343-1352.	1.0	57
38	Temporal trends in pulmonary arterial hypertension: results from the COMPERA registry. European Respiratory Journal, 2022, 59, 2102024.	3.1	57
39	Phenotyping of idiopathic pulmonary arterial hypertension: a registry analysis. Lancet Respiratory Medicine,the, 2022, 10, 937-948.	5.2	57
40	Therapy for pulmonary arterial hypertension due to congenital heart disease and Down's syndrome. International Journal of Cardiology, 2013, 164, 323-326.	0.8	55
41	Pulmonary Hypertension in Patients With COPD. Chest, 2021, 160, 678-689.	0.4	55
42	Right ventricular dyssynchrony in idiopathic pulmonary arterial hypertension: Determinants and impact on pump function. Journal of Heart and Lung Transplantation, 2015, 34, 381-389.	0.3	54
43	Echocardiography Combined With Cardiopulmonary Exercise Testing for the Prediction of Outcome in Idiopathic Pulmonary Arterial Hypertension. Chest, 2016, 150, 1313-1322.	0.4	51
44	Pulmonary hemodynamics contribute to indicate priority for lung transplantation in patients with cystic fibrosis. Journal of Thoracic and Cardiovascular Surgery, 2000, 119, 682-689.	0.4	50
45	Long term treatment of pulmonary arterial hypertension with beraprost, an oral prostacyclin analogue. British Heart Journal, 2001, 86, 661-665.	2.2	50
46	Prognostic factors in severe pulmonary hypertension patients who need parenteral prostanoid therapy: The impact of late referral. Journal of Heart and Lung Transplantation, 2012, 31, 364-372.	0.3	50
47	Circulating biomarkers in pulmonary arterial hypertension: Update and future direction. Journal of Heart and Lung Transplantation, 2015, 34, 282-305.	0.3	49
48	Influence of various therapeutic strategies on right ventricular morphology, function and hemodynamics in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2018, 37, 365-375.	0.3	49
49	Prognostic relevance of right heart reverse remodeling in idiopathic pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2018, 37, 195-205.	0.3	46
50	Assessment of severity of coronary narrowings by quantitative exercise echocardiography and comparison with quantitative arteriography. American Journal of Cardiology, 1991, 67, 1201-1207.	0.7	41
51	Tissue-Type Plasminogen Activator Therapy Versus Primary Coronary Angioplasty: Impact on Myocardial Tissue Perfusion and Regional Function 1 Month After Uncomplicated Myocardial Infarction. Journal of the American College of Cardiology, 1998, 31, 338-343.	1.2	41
52	Risk Reduction and Hemodynamics with Initial Combination Therapy in Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 484-492.	2.5	41
53	The importance of right ventricular evaluation in risk assessment and therapeutic strategies: Raising the bar in pulmonary arterial hypertension. International Journal of Cardiology, 2020, 301, 183-189.	0.8	40
54	Reconstruction of the pulmonary artery by a conduit of autologous pericardium. Journal of Thoracic and Cardiovascular Surgery, 1995, 110, 867-868.	0.4	38

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55	Comparative Effects of Irbesartan Versus Amlodipine on Left Ventricular Mass Index in Hypertensive Patients with Left Ventricular Hypertrophy. Journal of Cardiovascular Pharmacology, 2003, 42, 622-628.	0.8	37
56	Right ventricular dyssynchrony and exercise capacity in idiopathic pulmonary arterial hypertension. European Respiratory Journal, 2017, 49, 1601419.	3.1	37
57	Pulmonary Arterial Dilatation in Pulmonary Hypertension: Prevalence and Prognostic Relevance. Cardiology, 2012, 121, 76-82.	0.6	36
58	Preoperative Evaluation of Patients Undergoing Lung Resection Surgery: Defining the Role of the Anesthesiologist on a Multidisciplinary Team. Journal of Cardiothoracic and Vascular Anesthesia, 2016, 30, 530-538.	0.6	35
59	Pathophysiological adaptations to walking and cycling in primary pulmonary hypertension. European Journal of Applied Physiology, 2008, 102, 417-424.	1.2	32
60	Prognostic relevance of pulmonary arterial compliance after therapy initiation or escalation in patients with pulmonary arterial hypertension. International Journal of Cardiology, 2017, 230, 53-58.	0.8	32
61	Sildenafil dosed concomitantly with bosentan for adult pulmonary arterial hypertension in a randomized controlled trial. BMC Cardiovascular Disorders, 2017, 17, 239.	0.7	32
62	The added value of cardiopulmonary exercise testing in the follow-up of pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2019, 38, 306-314.	0.3	32
63	Usefulness of Tricuspid Annular Velocity in Identifying Global RV Dysfunction in Patients with Primary Pulmonary Hypertension: A Comparison with 3D Echoâ€Derived Right Ventricular Ejection Fraction. Echocardiography, 2008, 25, 289-293.	0.3	31
64	Clinical implications of idiopathic pulmonary arterial hypertension phenotypes defined by cluster analysis. Journal of Heart and Lung Transplantation, 2020, 39, 310-320.	0.3	31
65	Usefulness of the dipyridamole-Doppler test for diagnosis of coronary artery disease. American Journal of Cardiology, 1990, 65, 829-834.	0.7	30
66	Human Herpesvirus 8 and Pulmonary Hypertension. Emerging Infectious Diseases, 2005, 11, 1480-1482.	2.0	30
67	Right Ventricular Strain Curve Morphology and Outcome in IdiopathicÂPulmonary Arterial Hypertension. JACC: Cardiovascular Imaging, 2021, 14, 162-172.	2.3	29
68	Occult lung cancer in patients with bullous emphysema. Thorax, 1997, 52, 289-290.	2.7	29
69	Right ventricular concentric hypertrophy and clinical worsening in idiopathic pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2016, 35, 1321-1329.	0.3	28
70	Increased plasma levels of adrenomedullin, a vasoactive peptide, in patients with end-stage pulmonary disease. Regulatory Peptides, 2005, 124, 187-193.	1.9	27
71	Analysis of endothelin-1 and endothelin-1 receptor A gene polymorphisms in patients with pulmonary arterial hypertension. Internal and Emergency Medicine, 2012, 7, 425-430.	1.0	27
72	Haemodynamic effects of an acute vasodilator challenge in heart failure patients with reduced ejection fraction and different forms of postâ€capillary pulmonary hypertension. European Journal of Heart Failure, 2018, 20, 725-734.	2.9	27

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73	Aggressive Afterload Lowering to Improve the Right Ventricle: A New Target for Medical Therapy in Pulmonary Arterial Hypertension?. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 751-760.	2.5	27
74	HIV-related pulmonary hypertension. Acta Cardiologica, 2004, 59, 323-330.	0.3	26
75	Inhaled areosolized prostacyclin and pulmonary hypertension during anesthesia for lung transplantation. Transplantation Proceedings, 2001, 33, 1634-1636.	0.3	25
76	Right Heart and Pulmonary Vessels Structure and Function. Echocardiography, 2015, 32, S3-10.	0.3	25
77	Effects of Glucose-Insulin-Potassium Infusion on Myocardial Perfusion and Left Ventricular Remodeling in Patients Treated With Primary Angioplasty for ST-Elevation Acute Myocardial Infarction. American Journal of Cardiology, 2006, 98, 1349-1353.	0.7	24
78	Hemodynamics and risk assessment 2 years after the initiation of upfront ambrisentanâ€'tadalafil in pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2020, 39, 1389-1397.	0.3	24
79	Pulmonary hypertension in systemic sclerosis: prevalence, incidence and predictive factors in a large multicentric Italian cohort. Clinical and Experimental Rheumatology, 2013, 31, 31-6.	0.4	24
80	Usefulness of Adding Echocardiography of the Right Heart to Risk-Assessment Scores in Prostanoid-Treated Pulmonary Arterial Ahypertension. JACC: Cardiovascular Imaging, 2020, 13, 2054-2056.	2.3	23
81	Mid-Term Efficacy of Beraprost, an Oral Prostacyclin Analog, in the Treatment of Distal CTEPH: A Case Control Study. Cardiology, 2006, 106, 168-173.	0.6	21
82	Pulmonary Hypertension in Adults with Congenital Heart Disease: Real-World Data from the International COMPERA-CHD Registry. Journal of Clinical Medicine, 2020, 9, 1456.	1.0	21
83	Demographics, clinical characteristics, health resource utilization and cost of chronic thromboembolic pulmonary hypertension patients: retrospective results from six European countries. BMC Health Services Research, 2014, 14, 246.	0.9	20
84	Safety and efficacy evaluation of ambrisentan in pulmonary hypertension. Expert Opinion on Drug Safety, 2012, 11, 1003-1011.	1.0	19
85	Myocardial and microvascular inflammation/infection in patients with HIV-associated pulmonary artery hypertension. Aids, 2014, 28, 2541-2549.	1.0	18
86	Interleukin-32 in systemic sclerosis, a potential new biomarker for pulmonary arterial hypertension. Arthritis Research and Therapy, 2020, 22, 127.	1.6	18
87	Venous endotelin-1 (ET-1) and brain natriuretic peptide (BNP) plasma levels during 6-month bosentan treatment for pulmonary arterial hypertension. Regulatory Peptides, 2008, 151, 48-53.	1.9	17
88	Elevated serum levels of macrophage migration inhibitory factor and stem cell growth factor \hat{l}^2 in patients with idiopathic and systemic sclerosis associated pulmonary arterial hypertension. Reumatismo, 2014, 66, 270-276.	0.4	17
89	Efficacy of 1, 5, and 20Âmg oral sildenafil in the treatment of adults with pulmonary arterial hypertension: a randomized, double-blind study with open-label extension. BMC Pulmonary Medicine, 2017, 17, 44.	0.8	17
90	ISHLT consensus statement: Perioperative management of patients with pulmonary hypertension and right heart failure undergoing surgery. Journal of Heart and Lung Transplantation, 2022, 41, 1135-1194.	0.3	17

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91	Initial tadalafil and ambrisentan combination therapy in pulmonary arterial hypertension. Journal of Cardiovascular Medicine, 2018, 19, 12-17.	0.6	16
92	Timing and Priorities for Cystic Fibrosis Patients Candidates to Lung Transplantation. European Journal of Pediatric Surgery, 1998, 8, 274-277.	0.7	14
93	Heart failure in patients with human immunodeficiency virus. Journal of Cardiovascular Medicine, 2015, 16, 383-389.	0.6	14
94	The impact of delayed treatment on 6-minute walk distance test in patients with pulmonary arterial hypertension: A meta-analysis. International Journal of Cardiology, 2018, 254, 299-301.	0.8	14
95	Ambrisentan for the treatment of adults with pulmonary arterial hypertension: a review. Current Medical Research and Opinion, 2015, 31, 1793-1807.	0.9	13
96	The Growing Role of Echocardiography in Pulmonary Arterial Hypertension Risk Stratification: The Missing Piece. Journal of Clinical Medicine, 2021, 10, 619.	1.0	13
97	Riociguat treatment in patients with pulmonary arterial hypertension: Final safety data from the EXPERT registry. Respiratory Medicine, 2021, 177, 106241.	1.3	13
98	Human herpesvirus 8 and pulmonary hypertension. Emerging Infectious Diseases, 2005, 11, 1480-2.	2.0	13
99	Incremental value of cardiopulmonary exercise testing in intermediate-risk pulmonary arterial hypertension. Journal of Heart and Lung Transplantation, 2022, 41, 780-790.	0.3	13
100	Relationship between baseline ET-1 plasma levels and outcome in patients with idiopathic pulmonary hypertension treated with bosentan. International Journal of Cardiology, 2013, 167, 220-224.	0.8	12
101	Clinical impact of echocardiography in prognostic stratification after acute myocardial infarction. American Journal of Cardiology, 1998, 81, 17G-20G.	0.7	10
102	Prognostic significance of the echocardiographic estimate of pulmonary hypertension and of right ventricular dysfunction in acute decompensated heart failure. A pilot study in HFrEF patients. International Journal of Cardiology, 2018, 271, 301-305.	0.8	10
103	Hemodynamics during inhaled nitric oxide in lung transplant candidates. Transplantation Proceedings, 1997, 29, 3367-3370.	0.3	9
104	Improved results with lung transplantation for cystic fibrosis. Transplantation Proceedings, 2001, 33, 1632-1633.	0.3	9
105	ECMO Assistance during Mechanical Ventilation: Effects Induced on Energetic and Haemodynamic Variables. Computer Methods and Programs in Biomedicine, 2021, 202, 106003.	2.6	9
106	Prognostic value of improvement endpoints in pulmonary arterial hypertension trials: A COMPERA analysis. Journal of Heart and Lung Transplantation, 2022, 41, 971-981.	0.3	9
107	Imaging risk in pulmonary arterial hypertension. European Respiratory Journal, 2020, 56, 2002313.	3.1	8
108	Medical treatment of pulmonary hypertension in adults with congenital heart disease: updated and extended results from the International COMPERA-CHD Registry. Cardiovascular Diagnosis and Therapy, 2021, 11, 1255-1268.	0.7	8

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109	The paradox of pulmonary arterial hypertension in Italy in the COVID-19 era: is risk of disease progression around the corner?. European Respiratory Journal, 2022, 60, 2102276.	3.1	8
110	â€~Real-life' information on pulmonary arterial hypertension: the iPHnet Project. Current Medical Research and Opinion, 2014, 30, 2409-2414.	0.9	7
111	Endogenous opioid system modulation in anginal pain: Demonstration of its central activity. American Heart Journal, 1992, 124, 589-595.	1.2	6
112	Intra-aortic balloon counterpulsation timing: A new numerical model for programming and training in the clinical environment Computer Methods and Programs in Biomedicine, 2020, 194, 105537.	2.6	6
113	Response to Letters Regarding Article, "Anticoagulation and Survival in Pulmonary Arterial Hypertension: Results From the Comparative, Prospective Registry of Newly Initiated Therapies for Pulmonary Hypertension (COMPERA)― Circulation, 2014, 130, e110-2.	1.6	5
114	Identification of coronary artery by-pass grafts: reliability of MRI in clinical practice. International Journal of Cardiovascular Imaging, 1992, 8, 85-94.	0.2	4
115	Usefulness of 2D echo Doppler in the preoperative assessment of cystic fibrosis patients who are candidates for lung transplantation. Transplantation Proceedings, 2001, 33, 1628-1629.	0.3	4
116	Choosing the best first line oral drug agent in patients with pulmonary hypertension: Evidence from a network meta-analysis. International Journal of Cardiology, 2013, 168, 4336-4338.	0.8	4
117	The importance of right ventricular function in patients with pulmonary arterial hypertension. Expert Review of Respiratory Medicine, 2018, 12, 809-815.	1.0	4
118	Pulmonary hypertension due to lung disease – Results from COMPERA. , 2015, , .		4
119	Optimal duration of dual anti-platelet therapy after percutaneous coronary intervention. Journal of Cardiovascular Medicine, 2017, 18, 1-9.	0.6	3
120	Pulmonary hypertension in left heart disease: The need to continue to explore. International Journal of Cardiology, 2019, 288, 132-134.	0.8	3
121	Future perspective in diabetic patients with pre- and post-capillary pulmonary hypertension. Heart Failure Reviews, 2023, 28, 745-755.	1.7	3
122	Plasma adrenomedullin and endothelin-1 concentration during low-dose dobutamine infusion: Relationship between pulmonary uptake and pulmonary vascular pressure/flow characteristics. Regulatory Peptides, 2006, 136, 85-91.	1.9	2
123	SPHERIC-1 (Sildenafil and Pulmonary HypERtension in COPD): Intention-to-Treat (ITT) Analysis of Safety and Efficacy Data. Journal of Heart and Lung Transplantation, 2014, 33, S148-S149.	0.3	2
124	Riociguat in Combination With Prostacyclin Analogs for the Treatment of Pulmonary Arterial Hypertension (PAH): A Subgroup Analysis of the PATENT Studies. Chest, 2015, 148, 922A.	0.4	2
125	Switching from PDE5i to Riociguat in the RESPITE Study: Effect on Right Heart Function. Journal of Heart and Lung Transplantation, 2018, 37, S152.	0.3	2
126	Exercise energy expenditure in patients with idiopathic pulmonary arterial hypertension: Impact on clinical severity and survival. Respiratory Physiology and Neurobiology, 2019, 264, 33-39.	0.7	2

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127	Hemodynamic Evaluation of the Right Heart-Pulmonary Circulation Unit in Patients Candidate to Transjugular Intrahepatic Portosystemic Shunt. Journal of Clinical Medicine, 2022, 11, 461.	1.0	2
128	Computational Simulator Models and Invasive Hemodynamic Monitoring as Tools for Precision Medicine in Pulmonary Arterial Hypertension. Journal of Clinical Medicine, 2022, 11, 82.	1.0	2
129	Comment on TopyÅ,a-Putowska et al. Echocardiography in Pulmonary Arterial Hypertension: Comprehensive Evaluation and Technical Considerations. J. Clin. Med. 2021, 10, 3229. Journal of Clinical Medicine, 2022, 11, 3337.	1.0	2
130	Telehealth: A winning weapon to face the COVID-19 outbreak for patients with pulmonary arterial hypertension. Vascular Pharmacology, 2022, 145, 107024.	1.0	2
131	Efficacy of Amlodipine in the Treatment of Stable Effort Angina. Clinical Drug Investigation, 1997, 13, 108-112.	1.1	1
132	Tissue viability by contrast echocardiography. European Journal of Echocardiography, 2006, 7, S22-S29.	2.3	1
133	Unusual presentation for a patent ductus arteriosus. European Respiratory Review, 2009, 18, 174-176.	3.0	1
134	Mode of Death in Patients with Pulmonary Arterial Hypertension. Journal of Heart and Lung Transplantation, 2013, 32, S17-S18.	0.3	1
135	Right Ventricular Remodeling in Idiopathic Pulmonary Arterial Hypertension: Concentric Versus Eccentric Hypetrophy. Journal of Heart and Lung Transplantation, 2014, 33, S148.	0.3	1
136	FRIO445â€Efficacy and Safety of Riociguat in Patients with Pulmonary Arterial Hypertension (PAH) Associated with Connective Tissue Disease (CTD): Results from Patent-1 and Patent-2. Annals of the Rheumatic Diseases, 2015, 74, 588.2-589.	0.5	1
137	Effects of Riociguat in Treatment-Naive vs Pretreated Patients With Pulmonary Arterial Hypertension: 2-Year Efficacy Results From the PATENT-2 Study. Chest, 2016, 150, 1162A.	0.4	1
138	Letter to the editor about the paper "Right ventricular dyssynchrony predicts clinical outcomes in patients with pulmonary hypertension―by Murata et al International Journal of Cardiology, 2017, 234, 128.	0.8	1
139	Oxygen supplementation for pulmonary arterial hypertension? Clues from the REVEAL registry. Journal of Heart and Lung Transplantation, 2018, 37, 941-942.	0.3	1
140	Longâ€ŧerm study of oral treprostinil to treat pulmonary arterial hypertension: dosing, tolerability, and pharmacokinetics. Pulmonary Circulation, 2020, 10, 1-9.	0.8	1
141	Peripheral Arterial Stiffness in Acute Pulmonary Embolism and Pulmonary Hypertension at Short-Term Follow-Up. Journal of Clinical Medicine, 2021, 10, 3008.	1.0	1
142	Complex connections: A young man presenting with shortness of breath, hypoxemia, right lumbar pain and left limb swelling. Echocardiography, 2022, , .	0.3	1
143	198â€∫Intrapulmonary shunt assessment in pulmonary arterial hypertension. European Heart Journal Supplements, 2021, 23, .	0.0	1
144	BOSENTAN THERAPY IN PATIENTS WITH PULMONARY HYPERTENSION SECONDARY TO CONGENITAL HEART DISEASE (EISENMENGER PHYSIOLOGY). Chest, 2005, 128, 366S.	0.4	0

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145	ACUTE HEMODYNAMIC EFFECT OF SILDENAFIL IN COMPARISON WITH INHALED NITRIC OXIDE IN PATIENTS RECEIVING BOSENTAN THERAPY FOR PULMONARY ARTERIAL HYPERTENSION. Chest, 2007, 132, 488A.	0.4	O
146	258: Predictors of Clinical Worsening in Patients with PAH Treated with Bosentan. Journal of Heart and Lung Transplantation, 2009, 28, S156-S157.	0.3	0
147	Right Ventricular Mass/diastolic Ratio: An Expression Of "adequate" RV Adaptation In Pulmonary Arterial Hypertension ?. , 2010, , .		0
148	Outcome Of Pregnancies In Women With Pulmonary Arterial Hypertension In The Modern Management Era. , $2011, \ldots$		0
149	A Health Economic Analysis Of Sitaxentan For The Treatment Of Pulmonary Arterial Hypertension In Europe., 2011,,.		0
150	PCV22 Treatment Patterns and Outcomes in Patients with Pulmonary Arterial Hypertension and Chronic Thromboembolic Pulmonary Hypertension: Results of a Retrospective Chart Review in Six European Countries. Value in Health, 2012, 15, A115.	0.1	0
151	Right Ventricular Dyssynchrony Predicts Clinical Worsening in Idiopathic Pulmonary Arterial Hypertension. Journal of Heart and Lung Transplantation, 2013, 32, S79.	0.3	0
152	Myocardial and micro-vascular inflammation/infection in HIV/HCV associated pulmonary artery hypertension. European Heart Journal, 2013, 34, P228-P228.	1.0	0
153	Right Ventricular Dyssynchrony in Idiopathic Pulmonary Arterial Hypertension: Insights From Echocardiographic and Cardiac Magnetic Resonance Imaging. Journal of Heart and Lung Transplantation, 2014, 33, S229.	0.3	0
154	Concentric Hypertrophy Protects Against Clinical Worsening in Idiopathic Pulmonary Arterial: Hypertension: Insights From Magnetic Resonance Imaging. Journal of Heart and Lung Transplantation, 2015, 34, S117.	0.3	0
155	Determinats and Prognostic Significance of Right Ventricular Reverse Remodeling in Idiopathic Pulmonary Arterial Hypertension Receiving Specific Medical Treatment. Journal of Heart and Lung Transplantation, 2015, 34, S116-S117.	0.3	0
156	Muscular Efficiency in Patients With Idiopathic Pulmonary Arterial Hypertension (iPAH): Impact on Clinical Severity and Survival. Journal of Heart and Lung Transplantation, 2015, 34, S339.	0.3	0
157	Right Ventricular Dyssynchrony and Exercise Capacity in Idiopathic Pulmonary Arterial Hypertension: Insights Form Echocardiography and Cardiopulmonary Exercise Test. Journal of Heart and Lung Transplantation, 2016, 35, S149.	0.3	0
158	Incremental Benefit of Echocardiographic Imaging and Cardiopulmonary Exercise Test in Prognostic Evaluation of Idiopathic Pulmonary Arterial Hypertension. Journal of Heart and Lung Transplantation, 2016, 35, S149.	0.3	0
159	Incremental Benefit of Cardiopulmonary Exercise Testing for the Prediction of Outcome in Stable Prevalent Pulmonary Arterial Hypertension Patients. Journal of Heart and Lung Transplantation, 2017, 36, S76.	0.3	0
160	Echocardiography Predicts the Outcome in Pulmonary Arterial Hypertension Patients Treted With Parenteral Prostanoids. Journal of Heart and Lung Transplantation, 2018, 37, S206.	0.3	0
161	Initial combination therapy for patients with pulmonary arterial hypertension (PAH): a budget impact analysis from the perspective of the Italian national healthcare system. Expert Opinion on Orphan Drugs, 2018, 6, 1-7.	0.5	0
162	Clinical Impact of Right Ventricular Diastolic Patterns in Idiopathic Pulmonary Arterial Hypertension by Speckle Traiking. Journal of Heart and Lung Transplantation, 2019, 38, S487.	0.3	0

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164	Suppressor of Tumorigenicity 2 as a Biomarker in Pulmonary Arterial Hypertension and its Association with REVEAL Risk Score in Riociguat-Treated Patients in the RESPITE Study. Journal of Heart and Lung Transplantation, 2019, 38, S96.	0.3	0
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