Short Editorial



Coronary Compression in Pulmonary Hypertension: A Treatable Threat Hidden in Plain Sight

Allan K. N. Alencar¹

Department of Biomedical Engineering, Tulane University,¹ New Orleans – USA

Short Editorial related to the article: Left Main Coronary Artery Angioplasty for the Treatment of Extrinsic Compression in Patients with Pulmonary Hypertension

Pulmonary hypertension (PH) represents a severe, progressive condition associated with right ventricular failure, impaired functional capacity, and increased mortality. 1.2 Although angina is not a hallmark symptom of PH, it can affect up to 30% of patients and is frequently overlooked in clinical practice. 3 The study published in this issue of *Arquivos Brasileiros de Cardiologia* 4 addresses a rare yet critical cause of chest pain in PH: extrinsic compression of the left main coronary artery (LMCA) by a dilated pulmonary artery. The authors present an observational case series of 12 patients undergoing percutaneous coronary intervention (PCI) with stenting for LMCA compression, reporting both feasibility and sustained symptom relief.

While LMCA atherosclerosis is well recognized for its prognostic impact, compression of this vessel due to pulmonary artery dilation in PH remains underdiagnosed. The authors highlight that up to 76% of patients with advanced PH exhibit pulmonary artery enlargement, which may result in significant extrinsic compression of the LMCA, even in the absence of coronary atherosclerosis. ^{4,5} Importantly, LMCA compression has been associated with sudden cardiac death in PH cohorts, emphasizing the need for high clinical suspicion. ⁶⁻⁸

The current study reinforces findings from previous reports, such as the Italian series by Saia et al.⁹ which demonstrated successful outcomes following stenting in 53 patients. In this Brazilian series, all 12 patients reported symptom improvement, with angiographic success and no immediate complications. Over a 33-month follow-up, only four deaths occurred, all attributed to the progression of underlying PH, with no stent-related events reported.

This case series is valuable for several reasons. First, it provides real-world evidence from a high-volume PH center in Latin America, a region underrepresented in coronary-vascular PH research. Second, it demonstrates

Keywords

Pulmonary Hypertension; Percutaneous Coronary Intervention; Angioplasty.

Mailing Address: Allan K. N. Alencar •

Department of Biomedical Engineering, Tulane University, 500 Lindy Boggs Center, New Orleans, LA, 70118 - USA E-mail: aalencar@tulane.edu

Manuscript received May 14, 2025, revised manuscript May 21, 2025, accepted

DOI: https://doi.org/10.36660/abc.20250352i

a structured diagnostic and therapeutic algorithm involving multidisciplinary evaluation, advanced imaging with coronary computed tomography angiography, and individualized decision-making — a model worth replicating in specialized centers.

The study also aligns with growing awareness of nonatherosclerotic causes of myocardial ischemia in systemic diseases. The distinct pathophysiology of PH contributes to coronary underperfusion due to increased right ventricular wall stress and altered coronary reserve. The addition of mechanical compression by the pulmonary trunk creates a 'double-hit' mechanism that may go unrecognized unless coronary imaging is proactively pursued in symptomatic patients.

Importantly, the authors describe technical nuances of stent selection and deployment. In most cases, a single stent sufficed; however, in 25% of cases, a second stent was required to optimize radial force and lumen gain. Although intravascular imaging was used sparingly, its selective application underscores the real-world constraints and the need for flexible, pragmatic approaches in resource-limited settings.

One of the key messages is that LMCA compression should be systematically considered in PH patients with angina or unexplained ventricular dysfunction. While revascularization strategies for atherosclerotic LMCA disease are well established, 10 the role of PCI in extrinsic compression is still evolving. The long-term prognostic impact of stenting in this context is not fully known, and the current study adds to a growing body of literature suggesting symptomatic benefit, though randomized studies are unlikely due to the rarity of the condition.

Future research should explore the utility of routine screening in high-risk PH patients, especially those with pulmonary artery diameters >40 mm or anatomic proximity between the pulmonary artery and LMCA on imaging.¹¹ Additionally, assessing the impact of PCI on right ventricular function and long-term survival could yield insights into the integrative management of PH.¹²

In conclusion, the authors provide compelling evidence that LMCA compression in PH is both clinically significant and therapeutically actionable. Their experience supports the incorporation of coronary evaluation into PH care algorithms and positions PCI as a safe and effective intervention in selected cases. In an era of precision cardiology, recognizing and treating such anatomical culprits is not only feasible — it may also prove to be life-saving.

Short Editorial

References

- Humbert M, Kovacs C, Hoeper MM, Badagliacca R, Berger RMF, Brida M, et al. 2022 ESC/ERS Guidelines for the Diagnosis and Treatment of Pulmonary Hypertension. Eur Heart J. 2022;43(38):3618-731. doi: 10.1093/eurheartj/ ebac237
- Simonneau G, Montani D, Celermajer DS, Denton CP, Gatzoulis MA, Krowka M, et al. Haemodynamic Definitions and Updated Clinical Classification of Pulmonary Hypertension. Eur Respir J. 2019;53(1):1801913. doi: 10.1183/13993003.01913-2018.
- Galiè N, Saia F, Palazzini M, Manes A, Russo V, Reggiani MLB, et al. Left Main Coronary Artery Compression in Patients with Pulmonary Arterial Hypertension and Angina. J Am Coll Cardiol. 2017;69(23):2808-17. doi: 10.1016/j.jacc.2017.03.597.
- Bichuette LD, Calderaro D, Lemos Neto PA, Fonseca EKUN, Tatagiba LS, Vieira TM, et al. Angioplastia do Tronco da Artéria Coronária Esquerda no Tratamento de Compressão Extrínseca em Pacientes com Hipertensão Pulmonar. Arq Bras Cardiol. 2025; 122(7):e20250002. DOI: https://doi. org/10.36660/abc.20250002i.
- Badagliacca R, Poscia R, Pezzuto B, Papa S, Nona A, Mancone M, et al. Pulmonary Arterial Dilatation in Pulmonary Hypertension: Prevalence and Prognostic Relevance. Cardiology. 2012;121(2):76-82. doi: 10.1159/000336172.
- Hoette S, Figueiredo C, Dias B, Alves JL Jr, Gavilanes F, Prada LF, et al. Pulmonary Artery Enlargement in Schistosomiasis Associated Pulmonary Arterial Hypertension. BMC Pulm Med. 2015;15:118. doi: 10.1186/ s12890-015-0115-y.

- Żyłkowska J, Kurzyna M, Florczyk M, Burakowska B, Grzegorczyk F, Burakowski J, et al. Pulmonary Artery Dilatation Correlates with the Risk of Unexpected Death in Chronic Arterial or Thromboembolic Pulmonary Hypertension. Chest. 2012;142(6):1406-16. doi: 10.1378/chest.11-2794.
- Lee SE, An HY, Im JH, Sung JM, Cho IJ, Shim CY, et al. Screening of Mechanical Complications of Dilated Pulmonary Artery Related to the Risk for Sudden Cardiac Death in Patients with Pulmonary Arterial Hypertension by Transthoracic Echocardiography. J Am Soc Echocardiogr. 2016;29(6):561-6. doi: 10.1016/j.echo.2016.02.002.
- Saia F, Dall'Ara G, Marzocchi A, Dardi F, Palazzini M, Manes A, et al. Left Main Coronary Artery Extrinsic Compression in Patients with Pulmonary Arterial Hypertension: Technical Insights and Long-Term Clinical Outcomes after Stenting. JACC Cardiovasc Interv. 2019;12(3):319-21. doi: 10.1016/j. jcin.2018.08.002.
- Crystal GJ, Pagel PS. Right Ventricular Perfusion: Physiology and Clinical Implications. Anesthesiology. 2018;128(1):202-18. doi: 10.1097/ ALN.00000000001891.
- Neumann FJ, Sousa-Uva M, Ahlsson A, Alfonso F, Banning AP, Benedetto U, et al. 2018 ESC/EACTS Guidelines on Myocardial Revascularization. Eur Heart J. 2019;40(2):87-165. doi: 10.1093/eurhearti/ehy394.
- Mesquita SM, Castro CR, Ikari NM, Oliveira SA, Lopes AA. Likelihood of Left Main Coronary Artery Compression Based on Pulmonary Trunk Diameter in Patients with Pulmonary Hypertension. Am J Med. 2004;116(6):369-74. doi: 10.1016/j.amjmed.2003.11.015.

