



# Intra-Aortic Balloon Pump Placement in the Axillary Artery: Where are We?

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The prevalence of patients with heart failure (HF) in Brazil is high, with an increasing number of hospitalizations for advanced HF in tertiary care services. This suggests that patients' conditions are persistently more severe, with recurrent episodes of pulmonary congestion or low cardiac output requiring frequent hospitalization. Multidisciplinary clinical support and optimized medical therapy are fundamental in the treatment of these patients. However, in refractory cases, bridge or destination therapies such as circulatory or ventricular assist devices (VADs) and heart transplantation may be indicated.<sup>1,2</sup>

Patients with decompensated INTERMACS 2 or 1 HF may have an indication for mechanical circulatory support (MCS) during hospital stay. Devices currently available in Brazil include intra-aortic balloon pump (IABP), Impella CP, venoarterial extracorporeal membrane oxygenation, and Centrimag.<sup>2</sup> The IABP is the most used device in Brazil and worldwide due to easy access, cost-effectiveness, and simple implant procedure and management. Despite having a modest effect on cardiac output, the IABP has a significant impact on circulatory hemodynamics, is simpler to use, and has an equal or superior safety profile compared with more modern devices.<sup>3-5</sup>

Complete or partial patient immobilization is inherent in critical HF and in the use of MCS and thus may be required. The development of additional complications due to immobilization is a risk factor for worse in-hospital outcomes, and complications such as sarcopenia and cachexia are more frequent and often progressive.<sup>6,7</sup>

Aiming at reducing immobility and its consequences while still providing the necessary hemodynamic support, McBride et al. first described in 1989 a technique for surgical placement of an IABP through the axillary artery.<sup>8</sup> In the early 2000s, two case series were published. The first series described the outcomes of 13 patients over a 3-year period who had received IABP support for a mean duration of 37 days. Of these, 10 underwent a heart transplant.<sup>9</sup> The second series reported the outcomes of

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4 patients with ischemic cardiomyopathy on the heart transplant waiting list. Support duration ranged from 12 to 70 days, and all patients underwent successful transplants.<sup>10</sup>

In 2012, a series of 18 patients surgically implanted with an axillary IABP between 2007 and 2010 was published. Median support duration was 19 days, and 72% of patients underwent a heart transplant. Three patients developed device-related complications, ie, IABP displacement, rupture, or kinking. These complications were not associated with worse outcomes. There were no vascular complications or stroke.<sup>11</sup>

The first series of patients treated with an axillary IABP using only percutaneous access was published in 2013. Fifty patients referred for heart transplantation or VAD evaluation received a left axillary IABP between 2007 and 2012. Mean support duration was 18 days, and 84% of patients underwent a heart transplant. Complications requiring intervention included one case of significant bleeding and two cases of left upper extremity ischemia. IABP repositioning was required in 44% of patients, whereas 20% of patients required IABP replacement due to malfunction. There were no IABP-related deaths, strokes, or infections.<sup>12</sup>

In 2020, the same group of researchers expanded on previous experience and published a series of 195 patients who had received an axillary IABP between 2007 and 2018. Patients were divided into two groups according to therapeutic success, which was defined as destination therapy. Success rate was 68%; 120 patients underwent a heart transplant, and 13 patients received a long-term VAD. Among the remaining 62 patients (31.8%), 16 (8%) died, 18 (9.2%) required support escalation, and 28 (14%) underwent IABP removal (22 due to complications and 6 due to contraindications to destination therapy). The 1-year survival rate was 87% for heart transplantation and 62% for VAD implantation. Median support duration was 19 days. IABP replacement or repositioning was common (37%), with a mean number of IABP exchanges per patient of 0.68. Left upper extremity ischemia occurred in 3.5% of patients, but no patient suffered limb loss. Stroke, mesenteric ischemia, and bacteremia rates were 2.5%, 3%, and 9.2%, respectively. Among patients who developed bacteremia, 16.6% required IABP removal due to infection. Implant site-related bleeding occurred in 2.5% of patients, whereas 96 (49%) patients required IABP repositioning at least once.13

More recently, another study described 38 patients treated percutaneously between 2017 and 2020. IABP failure or migration requiring replacement occurred in 21.4% of patients. There were no major complications, and 81.6% of patients

# Viewpoint

received the intended therapy.<sup>14</sup> Nishida et al. reported their experience with 241 patients implanted with an IABP, of whom 58.9% underwent axillary insertion. Ambulation was possible in 90% of patients, and 86.7% received the intended therapy.<sup>15</sup> Vascular complications occurred in 3% of patients who underwent percutaneous IABP placement, and one third of these patients required surgical treatment.<sup>16</sup>

Some Brazilian hospitals perform percutaneous IABP placement in the left upper extremity, but data on MCS implantation and advanced HF treatment are scarce (Figures 1 and 2). Knowledge is essential to better understand the risk factors involved in complications and unfavorable outcomes, as well as to precisely define the role of axillary IABP in the current setting of MCS. Although these approaches have not been directly compared, the positive impact on adequate physical therapy and motor rehabilitation favors IABP placement via the axillary artery compared with the femoral artery. By allowing ambulation and greater mobility, the processes of sarcopenia and cachexia are also likely to be attenuated.

According to the available data, percutaneous axillary IABP placement is a viable and safe alternative for the implantation of an IABP in patients who require long-term support. The data suggest that placement via the axillary artery requires careful attention for correct device positioning, with increased rates of IABP repositioning and exchange compared with the femoral artery. Prospective and randomized clinical trials involving multidisciplinary teams are needed to provide hemodynamic

support and comprehensive care according to the demands and risk profile of each patient in this complex setting of advanced HE.

# **Author Contributions**

Conception and design of the research and Critical revision of the manuscript for intellectual content: Boros GAB, Bernoche CYSM, Nicz PFG; Writing of the manuscript: Boros GAB, Nicz PFG.

#### Potential Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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This study is not associated with any thesis or dissertation work.

### Ethics approval and consent to participate

This article does not contain any studies with human participants or animals performed by any of the authors.



Figure 1 – Arteriography performed with a 5F introducer to confirm puncture positioning

# Viewpoint



Figure 2 – Final position of the intra-aortic balloon pump after percutaneous implantation.

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