

Clinical and hemodynamic outcomes and mortality risk factors in patients undergoing pulmonary thromboendarterectomy

Resultados clínicos y hemodinámicos y factores de riesgo de mortalidad en pacientes sometidos a tromboendarterectomía pulmonar

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Abstract

Background: Pulmonary thromboendarterectomy is the current treatment of choice in patients with chronic thromboembolic pulmonary hypertension. The objective of the present study was to analyze the clinical and hemodynamic outcomes and the risk factors for mortality in a cardiovascular center in Colombia. **Methods:** Cohort study, conducted between 2001 and 2019. All operated patients were included in the study. Risk factors associated with mortality were established by means of a multivariate regression using the COX method and survival was established using the Kaplan–Meier method. $p < 0.05$ was considered statistically significant. **Results:** Seventy-three patients were operated. Median age was 51 years, 55% of females, 79% had functional Class III and IV. The mean pulmonary arterial pressure was 50 mmHg and 640 dyn.s.cm⁻⁵ for pulmonary vascular resistance (PVR). After the intervention, there was a decrease in mean pulmonary artery pressure ($p \leq 0.001$) and in PVR ($p = 0.357$); 21% had evidence of residual pulmonary hypertension. Only 8% and 6% continued with functional Class III and IV at 6 and 12 months, respectively. There were 15 deaths (19.1%; 12% at 30 days). The factors associated with mortality were the diastolic diameter of the right ventricle measured postoperatively (hazard ratio [HR] 10.88 95% confidence interval [CI] 1.97-62, $p = 0.007$), time of invasive mechanical ventilation (HR 1.06 95% CI 1.02-1.09 $p = 0.004$), and the presence of complications during the surgical procedure (HR 5.62 95% CI 1.94-16.22 $p = 0.001$). **Conclusions:** Pulmonary thromboendarterectomy is associated with excellent clinical and hemodynamic outcomes. The mortality risk factors found are not those usually described in the literature.

Keywords: Pulmonary hypertension. Pulmonary embolism. Pulmonary thromboendarterectomy.

Resumen

Antecedentes: La tromboendarterectomía pulmonar es el tratamiento de elección actual en pacientes con hipertensión pulmonar tromboembólica crónica. El objetivo del presente estudio fue analizar los resultados clínicos y hemodinámicos y los factores de riesgo de mortalidad en un centro cardiovascular de Colombia. **Métodos:** Estudio de cohorte entre 2001 y

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2019. Se incluyeron todos los pacientes operados. Los factores de riesgo asociados a la mortalidad se establecieron mediante una regresión multivariante mediante el método COX y la supervivencia se estableció mediante el método de Kaplan-Meier. Los valores de $p < 0.05$ se consideraron estadísticamente significativos. **Resultados:** se operaron 73 pacientes. La mediana de edad fue de 51 años, 55% mujeres, 79% tenían clase funcional III y IV. La presión arterial pulmonar media fue de 50 mmHg y 640 dyn.s.cm⁻⁵ para la resistencia vascular pulmonar. Después de la intervención, hubo una disminución en la presión arterial pulmonar media ($p \leq 0.001$) y en la resistencia vascular pulmonar ($p = 0.357$). El 21% tenía evidencia de hipertensión pulmonar residual. Solo el 8% y el 6% continuaron con clase funcional III y IV a los 6 y 12 meses respectivamente. Hubo 15 muertes (19.1%; 12% a los 30 días). Los factores asociados con la mortalidad fueron el diámetro diastólico del ventrículo derecho medido en el postoperatorio (HR 10.88 IC 95% 1.97-62, $p = 0.007$), el tiempo de ventilación mecánica invasiva (HR 1.06 IC 95% 1.02-1.09 $p = 0.004$) y el presencia de complicaciones durante el procedimiento quirúrgico (HR 5.62 IC 95% 1.94-16.22 $p = 0.001$). **Conclusiones:** La tromboendarterectomía pulmonar se asocia con excelentes resultados clínicos y hemodinámicos. Los factores de riesgo de mortalidad encontrados no son los habitualmente descritos en la literatura.

Palabras clave: Hipertensión pulmonar. Embolia pulmonar. Tromboendarterectomía pulmonar.

Introduction

Chronic thromboembolic pulmonary hypertension is a rare yet underdiagnosed pulmonary vascular disease, which is a sequela of pulmonary thromboembolism¹ and corresponds to Group 4 of the classification of the World Health Organization².

Although many of the patients who survive a pulmonary embolism present improvement overtime with resolution of the thrombotic load, there is a small group that develops chronic thromboembolic pulmonary hypertension³ due to a different natural history where resolution is limited, with organization and recanalization of the thrombus¹. This is associated with changes in the small pulmonary arteries, which generates an increase in vascular resistance and pulmonary hypertension⁴.

From the epidemiological point of view, there are conflicting data regarding incidence and prevalence. However, recent reviews of hospital databases and clinical studies show that pulmonary hypertension occurs in 3.8-4% of patients who survive a pulmonary thromboembolism event^{3,5,6}. With the above, an incidence of 3-5 cases per 100,000 cases in the United States and Europe can be estimated, with less than one-third of the cases being diagnosed³. Increased pressure and pulmonary vascular resistance (PVR) can lead to the development of the right heart failure and premature mortality⁷.

In selected patients with disease that is susceptible to surgical management, this is the treatment of choice, given the high probability of cure that such management achieves. Data from the different surgical groups show that patients present immediate and sustained improvement in pulmonary hemodynamics. The studies

are consistent in the different series with improvement in PVR from values >800 dyn.s.cm⁻⁵ to less than 400 dyn.s.cm⁻⁵ 8-10. Overtime, not only a decrease in pulmonary resistance has been noted but also an improvement in functional class demonstrated by walking for 6 min¹¹. Data from Italy have shown similar results with sustained improvement in functional class in 97% of patients who were in the WHO functional Class III and IV before the procedure and reaching functional Class I in 74% at 4 years¹².

The objective of this study was to analyze the clinical, hemodynamic results, and the risk factors for mortality in a cardiovascular referral center in Colombia.

Materials and Methods

A retrospective cohort study was conducted between January 2001 and November 2019 of patients undergoing pulmonary thromboendarterectomy in a cardiovascular care referral center. The patients were chosen for the procedure according to a multidisciplinary evaluation with the intervention of cardiovascular surgeons, cardiologists, specialists in heart failure, and pulmonologists from the pulmonary hypertension clinic, taking into account: functional class, accessibility of thrombotic lesions, hemodynamic status, and comorbidities. The pre-surgical evaluation including a detailed medical evaluation, with collection of clinical-demographic data, imaging studies (chest computed tomography, ventilation perfusion scan, echocardiography, and 6 min walk), and hemodynamic study. During the index hospitalization, data related to the surgical procedure and its complications, including mortality, were collected. Patient follow-up was carried out through visits between 1 and 6 months after the procedure, with clinical

evaluation, echocardiography, catheterization (at the discretion of the treating physician), and a 6 min walk. Subsequent follow-ups were carried out depending on the clinical condition of the patient, as well as aspects related to their health insurance system. A telephone contact was made with the patients for the present work to obtain data on the clinical condition as variables of interest until November 2019.

The data included in the analysis were demographic, clinical, hemodynamic (pre-surgical, postsurgical, and follow-up), echocardiographic variables, as well as the characteristics related to the procedure.

Primary outcome: mortality at 30 days and 6 months. Secondary outcomes: change in hemodynamics during follow-up; echocardiographic change in follow-up; complications of the procedure. It was defined to carry out an analysis of mortality before and after the year 2015, year during which pulmonary hypertension group received training at the University of San Diego.

Continuous variables are presented as mean with standard deviation (DS) or as median and interquartile range (IQR) according to their distribution. For the analysis of the change in clinical, echocardiographic, and hemodynamic parameters, the Wilcoxon test was used for non-parametric continuous variables and the Chi-square test for categorical measures, $p \leq 0.05$ was considered statistically significant. A multivariate logistic regression analysis of COX was performed to establish factors associated with mortality. The survival curve was calculated using the Kaplan–Meier method, applying the log-rank test to assess differences. All analyzes were carried out using the SPSS version 22.0 statistical program.

The present study was approved by the Institutional Ethics Committee and the Universidad Pontificia Bolivariana.

Results

Over an 18-year period, 73 patients underwent pulmonary thromboendarterectomy. The median age was 51 years, 55% of the patients were women. About 93% of the population had a history of pulmonary thromboembolism and 32% a history of thrombophilia, with antiphospholipid syndrome the most important condition (23%). The most frequent presenting symptoms were dyspnea and fatigue; 20% presented syncopal episodes, with a time between the onset of symptoms and their diagnosis of 12 months. The majority of the patients had marked symptoms, 79% being in functional Class III and IV (Table 1). The initial echocardiogram

Table 1. Baseline and surgical characteristics of the population

Demographic and anthropometric	
Age, years [IQR]	51 [39-61]
Male, n (%)	33 (45)
BMI [IQR]	24.6 [22.05-27.15]
Medical record	
Pulmonary embolism, n (%)	68 (93)
Thrombophilia, n (%)	23 (32)
Collagen disease, n (%)	12 (16)
Symptoms	
Dyspnea, n (%)	72 (99)
Fatigue, n (%)	70 (96)
Edema, n (%)	29 (40)
Chest pain, n (%)	30 (41)
Hemoptysis, n (%)	4 (6)
Syncope, n (%)	14 (19)
Initial WHO functional class, n (%)	
– I	0 (0)
– II	15 (21)
– III	44 (60)
– IV	14 (19)
Symptom time at diagnosis, months [IQR]	12 [6-48]
Diagnostic time to surgery, months [IQR]	12 [4-48]
Symptom time to surgery, months [IQR]	24 [8-64]
Pre-operative	
Anticoagulation, n (%)	45 (60.8%)
– Warfarin	24 (32.4%)
– Rivaroxaban	11 (14.8)
– Apixaban	9 (12.1%)
– Dabigatran	1
Inferior vena cava filter, n (%)	50 (69)
Perfusion time, minutes [IQR]	292 [244-326]
Aortic clamp time, minutes [IQR]	120 [96-142]
Circulatory arrest time, minutes [IQR]	64 [42-75]
Additional surgeries, n (%)	17 (23.2%)
– Tricuspid plasty	7 (41.1%)
– ASD closure	4 (23.5%)
– Pulmonary artery plasty	2 (11.7%)
– Coronary artery bypass graft surgery	2 (11.7%)
Intraoperative complications, n (%)	8 (10.9%)
– Vascular injuries	7 (87.5%)
– Bronchial injuries	1 (12.5%)
Mechanical ventilation time, days [IQR]	2 [1-6]
Reperfusion edema, n (%)	31 (43)
Post-operative vasoactive support	
– Dobutamine, n (%)	39 (53)
– Norepinephrine, n (%)	32 (43)
– Nitric oxide, n (%)	25 (34)
– Vasopressin, n (%)	5 (6.8)
– Milrinone, n (%)	21 (29)
– Levosimendan, n (%)	4 (6)
– Nitroprusside, n (%)	3 (4)
Hospital stay	
– ICU, days [IQR]	6 [3-13]
– Hospitalization, days [IQR]	20 [14-29]

showed right ventricular dilation in 63%, with a median pulmonary artery pressure of 79 mmHg, tricuspid annular plane systolic excursion (TAPSE) of 16, and a quarter of the population had moderate-to-severe tricuspid regurgitation (Table 1 supplementary appendix). The mean pressure of the pulmonary artery was 50 mmHg in the right catheterization, with PVR of 640 dyn.s.cm⁻⁵ and systemic resistance of 1600 dyn.s.cm⁻⁵. About 46.5% of the population received specific treatment for pulmonary hypertension, sildenafil (n: 25; 34.2%), and bosentan (n: 11; 15%) being the most frequently used (14 in combination therapy and of these 6 received the sildenafil-bosentan combination). A little more than half were in anticoagulation management (60%), half were taking warfarin, and the remaining were direct anticoagulants (rivaroxaban and apixaban the most frequent). Before the procedure, two-thirds of the population received an inferior vena cava filter implant (Table 1).

Regarding of the surgical procedure, the median perfusion time was 292 min, the aortic clamp time was 120 min, and the circulatory arrest time was 64 min. Additional procedures were performed in 17 patients (23%), with tricuspid plasty and the correction of an atrial septal defect being the most frequent. Complications related to the procedure occurred in a minority of cases (10%), of which vascular lesions were the main ones; no patient died during the procedure (Table 1).

Half of the patients required inotropic support (n: 39; 53%) after the procedure and in a lesser percentage vasopressor support (n: 32; 43%). About 43% of the patients had reperfusion edema. The stay in the intensive care unit was 6 days, with a median of 2 days of mechanical ventilation and a total time of hospitalization of 20 days (Table 1).

When analyzing the echocardiographic behavior (Table 1 supplementary appendix), there are three moments in which the patients were evaluated: pre-surgery, immediately after the procedure during hospitalization and in outpatient follow-up. The parameters that showed variation were the diastolic diameter of the right ventricle, which decreased from 4.7 cm to 4.1 cm and finally 3.6 cm ($p \leq 0.001$). Of 63 patients with right ventricular dilation in the initial study, only 21 patients continued with dilatation criteria in follow-up; something similar was noted in the systolic pressure of the pulmonary artery, which decreased from 76 mmHg to 45 mmHg after surgery, statistically significant change ($p \leq 0.001$) and to 43 mmHg in the last available study. However, only 46 patients (63% of the total population) had an echocardiogram at follow-up, and of

these 25 patients had pressures > 40 mmHg, which corresponds to 54%. The TAPSE showed a decrease in the immediate post-operative period to return to its initial value in the follow-up (16 mm; $p = 0.002$). Of the 18 patients with moderate-to-severe tricuspid regurgitation, only three remained with this criterion in the postsurgical echocardiogram and five in the outpatient echocardiogram ($p = 0.004$). No other significant changes were noted at the echocardiographic level.

Regarding the hemodynamic parameters (Table 2) after the surgical intervention, there was a significant decrease in parameters such as pulmonary artery systolic pressure (change of 46%; $p = 0.008$), mean pulmonary artery pressure (change of 40%; $p \leq 0.001$), pulmonary artery diastolic pressure (43.8% change; $p \leq 0.001$), systemic vascular resistance (SVR) (42.8% change; $p \leq 0.005$), and PVR (79% change; $p = 0.357$); the cardiac index also increased (24% increase; $p = 0.003$). Only 22 patients had right catheterization during follow-up, since only those who remained symptomatic or had functional impairment were subjected to this study. Mean pulmonary pressures > 25 mmHg were found in 16 patients in this group, of which only seven had values >40 mmHg. When analyzing PVR, 13 had > 240 dyn.s.cm⁻⁵ (equivalent to > 3 Wood units), 4 had values > 500 dyn.s.cm⁻⁵ (equivalent to > 6.2 Wood units), and only 1 presented a value dyn.s.cm⁻⁵ (equivalent to > 12 Wood units). With these data, 21% of the patients had invasive evidence of residual pulmonary hypertension.

At discharge, 20.5% of the patients continued with the specific treatment for pulmonary hypertension, with sildenafil monotherapy being the strategy most used in these patients.

In the 6-month follow-up, there were four hospitalizations: two for cardiovascular causes (heart failure) and two non-cardiovascular (abdominal pain and infection of the surgical site).

The change in the clinical condition of the patients was evaluated by functional class and walking at 6 min at 6 and 12 months (data presented in Table 3). The baseline functional class of the patients was III and IV in 79% of the population; at 6 months, only 7% were in functional Class III and one patient reported a functional Class IV; 54% of the patients reported functional Class I and II. At 6 months, the patients increased their 6 min walk by 15% (approximately 70 m), something that was statistically significant ($p = 0.04$)

Regarding mortality, there were 14 events (19.17%) in the study period (Fig. 1 and table 4). Most of these (n: 9; 12.32%) occurred at 30 days (Fig. 2), septic shock

Table 2. Hemodynamic variables

Parameter	Initial, (n = 72*)	Post-operative, (n = 68*)	Follow-up, (n = 22)	p
RAP mmHg	10 [6-16]	8 [4-10]	6 [3-8]	0.29
PASP mmHg	87 [80-100]	47 [36-57]	43 [32-53]	0.008
PADP mmHg	32 [25-39]	18 [15-23]	18 [12-21]	< 0.001
mPAP mmHg	50 [44-59]	30 [22-35]	33 [27-40]	< 0.001
PVR dyn.seg.cm ⁻⁵	640 [12-876]	136 [94-305]	352 [186-473]	0.357
SVR dyn.seg.cm ⁻⁵	1600 [1264-1865]	916 [694-1161]	1452 [955-1689]	0.005
Wedge mmHg	13 [9-16]	14 [12-28]	11 [7-14]	0.14
CI, L/min/m ²	2.4 [2.0-2.7]	3.2 [2.6-3.7]	3.2 [2.8-3.2]	0.003

*Variables presented as median [IQR]. P value obtained from the Wilcoxon test for non-parametric continuous variables and X2 for categorical variables. PASP: pulmonary artery diastolic pressure; PADP: pulmonary artery systolic pressure; mPAP: mean pulmonary artery pressure; RAP: right atrial pressure; CI: cardiac index; SVR: systemic vascular resistance; PVR: pulmonary vascular resistance.

Table 3. Clinical variables

Variable n (%)	Initial, (n = 72*)	6 months, (n = 45*)	12 months, (n = 25)	p
Class functional				0.34
I	0 (0)	23 (32)	19 (26)	
II	15 (21)	16 (22)	2 (3)	
III	44 (60)	5 (7)	4 (6)	
IV	14 (19)	1 (1)	0 (0)	
6 min walk distance, mts [IQR]	407 [321-493]	475 [420-565]	432 [366-484]	0.04

*Variables presented as n (%) or median [IQR]. P value obtained from the Wilcoxon test for non-parametric continuous variables and Chi-square X2 for categorical variables.

Table 4. Aspects related to mortality

Deaths n (%)	14 (19.17%)
30-day mortality, n (%)	9 (12.32%)
Septic shock	6
Ventilator-associated pneumonia	3
Nosocomial pneumonia	1
Intestinal necrosis	1
Cardiogenic shock	3
Mortality at 6 months, n (%)	5 (6.85%)

being the main cause followed by cardiogenic etiology. The deaths during the first 6 months were 5; of these three due to pulmonary infectious complications and in two of these events, the cause could not be established. An analysis was performed in two periods: between 2001 and 2014 and 2015 until 2019. In the first

period, 18 procedures were performed with 6 deaths (33%). In the second period, 55 procedures were performed with eight deaths for a mortality rate of 14.54% with a tendency to a better result in patients operated in survival at 6 months, however, without differences statistically neither at 30 days nor at 6 months (Fig. 3).

To establish the risk factors for mortality in our population, a multivariate logistic regression analysis of COX was performed. Demographic, clinical, surgical, and echocardiographic hemodynamic parameters were included (Table 2 supplementary appendix). The factors found associated with mortality were the diastolic diameter of the right ventricle measured postoperatively (hazard ratio [HR] 10.88 95% confidence interval [CI] 1.97-62, p = 0.007), time of invasive mechanical ventilation (HR 1.06 95% CI 1.02-1.09, p = 0.004), and the presence of complications during the surgical

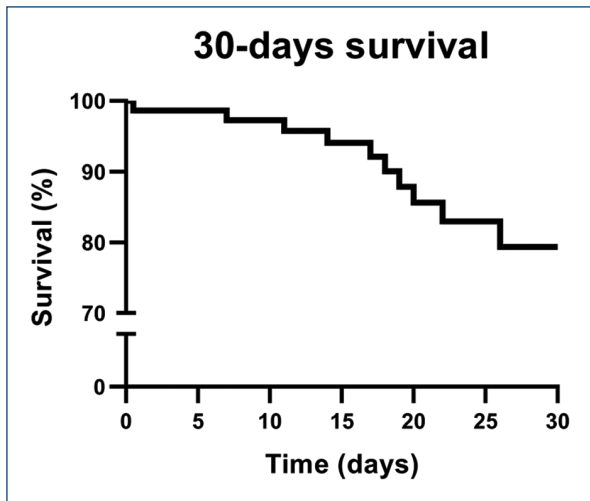


Figure 1. Survival curves (Kaplan–Meier) at 30 days.

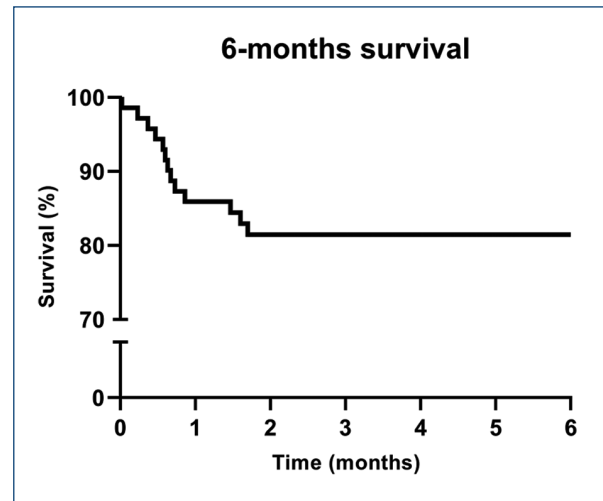


Figure 2. Survival curves (Kaplan–Meier) at 6-month mortality.

procedure (HR 5.62 95% CI 1.94-16.22, $p = 0.001$). None of the other variables included in the model presented a significant association. An exploratory analysis of survival was performed using the Kaplan–Meier method at both 30 days and 6 months for PVR and surgical times. No differences were found according to PVR, nevertheless, it clear that visually, there seems to be a trend toward better survival in patients with PVR $<1000 \text{ dyn.s.cm}^{-5}$ (Figure 1 of the supplementary appendix) and neither according to the surgical times (Figures 2, 3, and 4 of the supplementary appendix).

Discussion

Although the exact prevalence of chronic thromboembolic pulmonary hypertension is unknown, it is believed that 4-5% of patients who present with an episode of pulmonary embolism develop it. Pulmonary thromboendarterectomy remains the management strategy of choice in patients with pulmonary hypertension associated with chronic thromboembolic disease due to the results, it has shown in recent years (low mortality rate, complications, and high possibility of cure)¹³.

This study presents the clinical and hemodynamic results associated with the surgical management of thromboembolic pulmonary hypertension over a period of 18 years; previous reports in Colombia have included a small number of patients¹⁴.

Regarding the baseline characteristics of the population, these accord with those of the group of patients

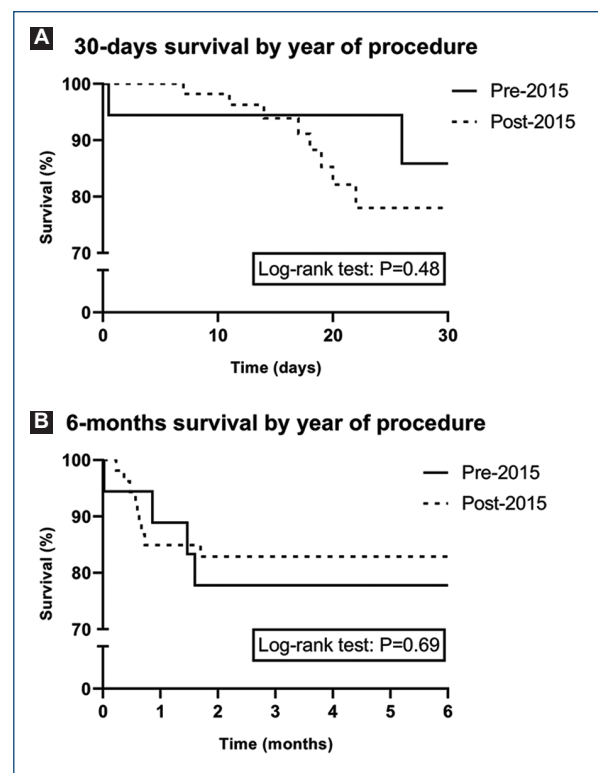


Figure 3. Survival curve of stratified patients before and after 2015. **A:** 30-day survival. **B:** 6-month survival. The difference between curves was evaluated by the log-rank test.

published by two of the largest registries in the United States¹⁵ and Europe⁸. The history of thrombophilia is in a medium range in relation to the different series, coinciding with the antiphospholipid syndrome being the

main alteration^{16,17}. Most of the patients (79%) presented severe and limiting symptoms measured in their baseline functional class (III and IV) something that is also similar to those documented in large centers^{8,15}. An important and higher proportion than that reported presented chest pain in our population^{8,18}. The time from the onset of symptoms to diagnosis is similar to that reported (12 months), not being the case for the times from diagnosis to surgery (78 days vs. 12 months) and from the onset of symptoms to the surgical procedure (24 months vs. 17 months)^{8,15,16}, surely something related to the Colombian health-care system. The stay in the intensive care unit was shorter than that reported (6 vs. 14) with similar days of mechanical ventilation (2 on average)^{8,15}. Among the complications after the procedure, a higher frequency of reperfusion edema was found compared to other studies^{8,16} (43% vs. 9.6-20%), something that did not increase the need for ventilatory support or stay in the unit intensive care, nor was it a risk factor for mortality in the population.

Regarding the technical aspects of the procedure, the median time of perfusion as for aortic clamp was like the data of large series^{8,15}, not being the case for the time of circulatory arrest, since in our series is double that reported (35 min for the California University in San Diego and the European vs. record 64 min in our cohort). Although this parameter has been related to transient neurological dysfunction, memory alterations, and focal motor deficits in some studies^{19,20}, these abnormalities were not found in the population studied. Complications occurred during surgery in eight patients, most of them arterial injuries, but none died as a cause of these; this parameter was one of the risk factors for death found, something that can be explained by an indirect relationship since these patients required a longer time of ventilatory support and stay in the intensive care unit, which increases the possibility of nosocomial or care-related infections, the main cause of death in the present study.

The hemodynamic and follow-up data found were similar to those reported worldwide with a remarkable improvement immediately after surgery, which in the long term translates into an improvement in dyspnea parameterized by functional class as well as an increase in the 6 min walk distance^{8,10,15,21,22} and the echocardiographic parameters (right ventricular diameter, significant tricuspid regurgitation), reinforcing the benefit of surgical management. The rate of residual pulmonary hypertension (21%) was similar to that reported by the San Diego group (16%).

The reported in-hospital mortality varies worldwide from 4.4 to 16%^{8,10,15} and is related to aspects such as the experience of the surgical group. In the California University in San Diego, mortality for the first 200 procedures was 17%, which decreased to 8.8% in the next 500 patients in 4 years, reaching the current reported rate of 2.2%¹⁵. Global mortality was 19.17% and in-hospital at 20 days 12.32%, what agrees with the aforementioned. In the analyzes carried out for the 2 time periods, in the first 18 procedures, mortality was 30% (2001-2014), decreasing to 14.54% in the following 55 interventions (2015-2019), something that reinforces the observations of improvement in mortality with the increase in the number of patients, procedures, and the experience of the surgical group. The main cause of death was infectious complications, which again denotes the safety of the intervention.

The risk factors for mortality in our population were the diameter of the right ventricle after surgery, the time of mechanical ventilation, and the presence of complications; these last findings are surely related to the fact that the main cause of death was septic shock, due to ventilator-associated pneumonia. No relationship or association was found with parameters related to higher mortality previously described, such as PVR before the procedure, the 6 min walk distance, and the functional class of the patients^{8,15,23}.

The present study has several limitations: the first is that the retrospective nature allows that biases cannot be completely ruled out during the study process. They are data from a single center. Patient follow-up by the medical group was low, with few data from studies such as echocardiography, 6 min walk distance, paraclinical tests such as natriuretic peptides, and right catheterization at 12 months. However, due to the health insurance model in Colombia, it is something that researchers cannot control.

Conclusions

Pulmonary thromboendarterectomy is the medical management of choice in patients with pulmonary hypertension in Group 4, due to the clinical and hemodynamic results and its safety.

In our population, the results obtained in a center with high volume and cardiovascular complexity was similar to those reported worldwide, with mortality that is decreasing as the number of cases increased. This should be considered the first treatment option in candidate patients in our setting.

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Conflicts of interest

None to disclose.

Ethical disclosures

Protection of human and animal subjects. The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

Confidentiality of data. The authors declare that they have followed the protocols of their work center on the publication of patient data.

Right to privacy and informed consent. The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

Supplementary Data

Supplementary data are available at Archivos de Cardiología de México online (<https://www.archivoscardiologia.com>). These data are provided by the corresponding author and published online for the benefit of the reader. The contents of supplementary data are the sole responsibility of the authors.

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