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The impact of patient choice on survival in chronic thromboembolic pulmonary hypertension

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Outcomes for patients undergoing pulmonary endarterectomy are excellent and superior to patients declining surgery http://ow.ly/9UZw30kA28m

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ABSTRACT  Pulmonary endarterectomy (PEA) is the gold standard treatment for operable chronic thromboembolic pulmonary hypertension (CTEPH). However, a proportion of patients with operable disease decline surgery. There are currently no published data on this patient group. The aim of this study was to identify outcomes and prognostic factors in a large cohort of consecutive patients with CTEPH.

Data were collected for consecutive, treatment-naive CTEPH patients at the Pulmonary Vascular Disease Unit of the Royal Hallamshire Hospital (Sheffield, UK) between 2001 and 2014.

Of 550 CTEPH patients (mean±SD age 63±15 years, follow-up ±3 years), 49% underwent surgery, 32% had technically operable disease and did not undergo surgery (including patient choice n=72 and unfit for surgery n=63), and 19% had inoperable disease due to disease distribution. The 5-year survival was superior in patients undergoing PEA (83%) versus technically operable disease who did not undergo surgery (59%) (p<0.001). Survival was superior in patients following PEA compared with those offered but declining surgery (55%) (p<0.001). In patients offered PEA, independent prognostic factors included mixed venous oxygen saturation, gas transfer and patient decision to proceed to surgery.

Outcomes in CTEPH following PEA are excellent and superior to patients declining surgery, and strongly favour consideration of a surgical intervention in eligible patients.

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Introduction
Chronic thromboembolic pulmonary hypertension (CTEPH) is a potentially curable form of pulmonary hypertension [1]. It may follow an acute episode of pulmonary embolism [2, 3], but can present as unexplained pulmonary hypertension with no previous history of venous thromboembolism (VTE) [4–6]. CTEPH occurs as a consequence of failure of clot resolution and secondary pulmonary arterial vasculopathy, leading to the development of pulmonary hypertension, right ventricular dysfunction and ultimately death [5, 7]. Over the past 20 years the treatment of CTEPH has evolved to include pulmonary endarterectomy (PEA) in operable cases [8] and pulmonary hypertension-specific therapy for inoperable disease [9–14]. Balloon pulmonary angioplasty is emerging as a potential treatment option in selected patients with inoperable disease [1, 15, 16].

PEA is currently considered the treatment of choice in patients with operable CTEPH, and is associated with excellent symptomatic benefit and long-term survival [17–19], with 10-year survival of 72–75% [20, 21]. Historical studies in patients treated with anticoagulation alone reported 5-year survival as low as 10% in patients with a mean pulmonary arterial pressure (mPAP) >50 mmHg [22, 23]. However, subsequent large registries have shown significant improvements in outcome in a heterogeneous group of nonoperated patients [24, 25]. Patients with CTEPH may be deemed inoperable where the pulmonary vascular resistance (PVR) is considered to be out of proportion to the degree of surgically accessible obstruction in the pulmonary vasculature [24, 26]. A significant proportion of patients who have technically operable disease do not undergo surgery for a variety of reasons, including comorbidities and patient choice [24–27]. To date, there are only limited data on the clinical course and rationale for treatment decisions in this patient group. In particular, there are very limited data on patients with technically operable disease who have declined surgery.

The aim of the current study was to provide data to help inform patient choice by identifying outcomes and prognostic factors in a large cohort of consecutive patients with CTEPH. Some of the results have been reported previously in abstract form [28, 29].

Methods
Consecutive, treatment-naive patients newly diagnosed with CTEPH at the Pulmonary Vascular Disease Unit of the Royal Hallamshire Hospital (Sheffield, UK) between January 1, 2001 and November 30, 2014 were prospectively recorded in hospital databases, as part of the Sheffield ASPIRE registry (ClinicalTrials.gov identifier NCT02565030). Baseline characteristics, treatment and follow-up data, using a census date of November 30, 2015, were subsequently collected from hospital records and departmental databases. Baseline characteristics, demographics, medical history, pulmonary function, exercise testing, right heart catheter metrics, imaging and treatment were included.

The diagnosis of CTEPH was based on findings obtained after at least 3 months of anticoagulation, and required the patient to have undergone right heart catheterisation and to have mPAP \( \geq 25 \) mmHg at rest and at least one segmental perfusion defect detected by perfusion lung scan or pulmonary artery obstruction seen by multidetector computed tomography angiography or conventional pulmonary angiography with other causes of pulmonary hypertension excluded [18, 19]. Patients were further classified into CTEPH: 1) operated group, 2) technically-operable-not-operated group and 3) patients with nonsurgical disease distribution. The technically-operable-not-operated group was further subclassified into four subgroups based on the reasons for the surgery not being performed: 1) declined surgery (patient choice), 2) being unfit for surgery following multidisciplinary team (MDT) assessment, 3) CTEPH with other contributors to symptoms in addition to clot burden and 4) reason for decision not clear. Suitability for PEA surgery was assessed by a review of clinical and radiological information by the surgical MDT at the national PEA centre at Papworth Hospital (Cambridge, UK).

The date of diagnosis of CTEPH was recorded as the date of the first right heart catheterisation. All patients were followed up until death or the census date of November 30, 2015. Ethical approval was granted for this study (REC 06/Q2308/8).

Statistical analysis
Descriptive data are presented using mean and standard deviation. Comparisons between groups were made using the t-test (for two groups) and ANOVA (with Bonferroni corrections for three groups) for continuous
data, and the Chi-squared test for categorical data. Survival was assessed by the Kaplan–Meier method. Comparisons between two groups were performed using the log-rank test. Prognostic variables were assessed using univariate and multivariate Cox regression analysis for survival. 72 variables in the CTEPH-surgical-operated group and 71 variables in the CTEPH-surgical-not-operated and CTEPH-nonsurgical groups were identified based on previous literature [24, 25, 30] and entered into univariate Cox regression analysis. Variables with a p-value of <0.20 at univariate analysis and <10% missing data were included for the multivariate analysis using the forward logistic regression method. To allow comparison between continuous variables, hazard ratios (HRs) were scaled using the standard deviation of the variable. A p-value of <0.05 was considered statistically significant. Accuracy of variables for prediction of mortality at 3 years was assessed using receiver operator characteristic (ROC) curve analysis. Statistical analysis was performed using SPSS version 25 (IBM, Armonk, NY, USA) and Prism software (GraphPad, La Jolla, CA, USA).

550 patients with CTEPH (mean±SD age 63±15 years, 50% female) were identified. A flowchart giving the classification with a detailed breakdown of patients is shown in figure 1.

Results
Patient characteristics
CTEPH (whole cohort)
Baseline characteristics for the major groups are summarised in table 1 and for technically-operable-not-operated subgroups in supplementary table S1. There was excellent data completeness, with data on lung function testing, exercise testing and imaging available in 94%, 94% and 93% of patients, respectively. Right heart catheterisation was performed in all patients. Survival data to the census date were available for all patients. The most common presenting symptoms were breathlessness (98%), ankle swelling (38%), pre-syncpe (27%) and chest pain (19%). There was no significant difference in symptoms or duration of symptoms prior to diagnosis between patients with technically operable disease undergoing and not undergoing surgery and those with a nonsurgical disease distribution. In terms of ethnicity, 95% of patients were White, with no significant difference in ethnicity between the CTEPH groups. With respect to

![Patient cohort flowchart showing group/subgroup classification. CTEPH: chronic thromboembolic pulmonary hypertension; CTED: chronic thromboembolic disease; RHC: right heart catheterisation.](https://doi.org/10.1183/13993003.00589-2018)
anticoagulation, 90% of the patients received a vitamin K antagonist, 4.5% received anti-Xa oral anticoagulants and 5% received low-molecular-weight heparin, while 0.5% were not anticoagulated. 76% of patients received pulmonary hypertension therapy, of whom 315 received phosphodiesterase type 5 inhibitor (PDE-5i) monotherapy, 49 received endothelin receptor antagonist monotherapy, 22 received prostanoid monotherapy, 18 received combination therapy with a PDE-5i and endothelin receptor antagonist, and 12 received combination therapy with a PDE-5i and prostanoid. No patients received riociguat.

CTEPH-operated group

Patients with CTEPH who underwent surgery were younger with a male predominance, and with better exercise capacity and gas transfer compared with patients in the other groups (table 1). The mean waiting time from date of diagnosis to surgery at the national PEA centre was 290±175 days. 74% received bridging therapy, of whom 74% received monotherapy with PDE-5i. Bridging therapy was commenced at diagnosis and did not lead to delay in referral for surgical consideration or time to surgery (supplementary table S2).

### TABLE 1 Baseline characteristics of patients with chronic thromboembolic pulmonary hypertension (CTEPH)

|                      | CTEPH (whole cohort) | CTEPH-operated group | CTEPH-technically-operable-not-operated group | CTEPH-nonsurgical-disease-distribution group |
|----------------------|----------------------|----------------------|----------------------------------------------|---------------------------------------------|
| Subjects             | 550 (100)            | 272 (49)             | 176 (32)                                     | 102 (19)                                    |
| Age years            | 63±15                | 58±14*               | 69±14*                                       | 65±15*                                      |
| Female               | 50                   | 45%                  | 50%                                          | 42%                                         |
| BMI kg·m⁻²           | 29±7                 | 30±7                 | 29±8                                         | 29±6                                        |
| Duration of symptoms |                      |                      |                                              |                                              |
| <1 year              | 15                   | 14                   | 17                                           | 12                                          |
| 1–2 years            | 38                   | 40                   | 35                                           | 33                                          |
| 2–5 years            | 24                   | 27                   | 19                                           | 28                                          |
| >5 years             | 16                   | 16                   | 14                                           | 18                                          |
| Not clear            | 8                    | 3                    | 14                                           | 9                                           |
| WHO FC I/II versus III/IV | 11/89             | 13/87*               | 11/89                                        | 3/97*                                       |
| ISWD m               | 189±177              | 232±185%             | 142±157%                                     | 155±160%                                    |
| RAP mmHg             | 11±5                 | 11±5                 | 11±6                                         | 12±5                                        |
| mPAP mmHg            | 46±11                | 47±11*               | 43±11*                                       | 48±12*                                      |
| Cardiac index L-min⁻¹·m⁻² | 2.5±0.8            | 2.5±0.7              | 2.6±0.8                                      | 2.5±0.8                                     |
| PCWP mmHg            | 12±5                 | 12±4                 | 12±6                                         | 11±6                                        |
| PVR WU               | 7.7±6                | 7.7±6                | 7.4±5.6                                      | 8.7±5.6                                     |
| SvO₂, %              | 61±8                 | 61±8                 | 61±9                                         | 61±9                                        |
| FEV₁ % pred          | 80±35                | 83±43†               | 75±23*                                       | 80±24                                       |
| FVC % pred           | 92±24                | 95±21                | 89±25                                        | 90±27                                       |
| Dłco % pred          | 61±17                | 65±15†               | 55±19*                                       | 61±17†                                      |
| History of acute VTE | 71                   | 74*                  | 74*                                          | 57*†                                        |
| IVC filter           | 3                    | 3                    | 5                                            | 1                                           |
| Thrombophilia        | 5                    | 6                    | 3                                            | 3                                           |
| History of cancer    | 11                   | 6*                   | 15*                                          | 15*                                         |
| Smoking              | 38                   | 38                   | 43                                           | 30                                          |
| Obesity              | 36                   | 39                   | 32                                           | 32                                          |
| Spleenectomy         | 5                    | 3*                   | 3*                                           | 12*†                                        |
| Thyroid disorder     | 12                   | 8*                   | 14                                           | 18*                                         |
| VA shunt/PPM infection | 2                  | 2                    | 0                                            | 4                                           |
| IBD                  | 1                    | 1                    | 2                                            | 2                                           |
| CAD                  | 11                   | 10                   | 14                                           | 7                                           |
| LV dysfunction       | 4                    | 2*                   | 8*†                                          | 2†                                          |
| Valvular heart disease | 2                | 2                    | 2                                            | 2                                           |
| CKD                  | 7                    | 3*                   | 13*                                          | 6                                           |
| COPD                 | 19                   | 13†                  | 30*                                          | 19                                          |
| PH therapy following diagnosis | 76 | 74*           | 72*                                          | 86*†                                        |

Data are presented as n (%), mean±SD or %. BMI: body mass index; WHO FC: World Health Organization Functional Class; ISWD: Incremental shuttle walk distance; RAP: right atrial pressure; mPAP: mean pulmonary arterial pressure; PCWP: pulmonary capillary wedge pressure; PVR: pulmonary vascular resistance; WU: Wood units; SvO₂: mixed venous oxygen saturation; FEV₁: forced expiratory volume in 1 s; FVC: forced vital capacity; Dłco: diffusing capacity of the lung for carbon monoxide; VTE: venous thromboembolism; IVC: inferior vena cava; VA: ventriculo-atrial; PPM: permanent pacemaker; IBD: inflammatory bowel disease; CAD: coronary artery disease; LV: left ventricular; CKD: chronic kidney disease; COPD: chronic obstructive pulmonary disease; PH: pulmonary hypertension. *: p<0.05 compared with CTEPH-operated; †: p<0.05 compared with CTEPH-technically-operable-not-operated; ‡: p<0.05 compared with CTEPH-nonsurgical-disease-distribution.

### CTEPH-operated group

Patients with CTEPH who underwent surgery were younger with a male predominance, and with better exercise capacity and gas transfer compared with patients in the other groups (table 1). The mean waiting time from date of diagnosis to surgery at the national PEA centre was 290±175 days. 74% received bridging therapy, of whom 74% received monotherapy with PDE-5i. Bridging therapy was commenced at diagnosis and did not lead to delay in referral for surgical consideration or time to surgery (supplementary table S2).
CTEPH-technically-operable-not-operated subgroup: declined surgery (patient choice)
Compared with patients undergoing PEA, patients who declined surgery were older, more likely to be female, and had more comorbidities and lower exercise capacity (all p<0.05) (table 2).

CTEPH-technically-operable-not-operated subgroup: not offered surgery (unfit for surgery, other contributors to symptoms in addition to clot burden)
Patients who were unfit for surgery had worse exercise capacity, lower diffusing capacity of the lung for carbon monoxide (DLCO), were more likely to be current smokers and had a higher prevalence of chronic obstructive pulmonary disease (COPD) than other technically operable patients not undergoing surgery. Patients with other contributors to symptoms in addition to clot burden (where the risk of surgery was felt to outweigh the benefit) had milder pulmonary haemodynamics and a better exercise capacity than the other two subgroups. The baseline characteristics of these patients are described in further detail in supplementary table S1.

CTEPH-nonsurgical-disease-distribution group
History of VTE was less common in patients with nonsurgical disease. A history of thyroid disease and splenectomy was also significantly more common in this subgroup compared with patients undergoing PEA (table 1).

### TABLE 2 Baseline characteristics of chronic thromboembolic pulmonary hypertension (CTEPH)-operated group versus patients who declined surgery (patient choice)

| Subjects n | CTEPH-operated | Declined surgery (patient choice) | p-value |
|------------|----------------|-----------------------------------|---------|
| Age years  | 58±14          | 68±16                             | <0.001  |
| Female     | 45             | 63                                | 0.007   |
| BMI kg·m⁻² | 30±7           | 29±7                              | 0.485   |
| WHO FC I/II versus II/IV | 13/87       | 17/83                             | 0.462   |
| Duration of symptoms |         |                                   | 0.009   |
| <1 year    | 14             | 22                                |         |
| 1–2 years  | 40             | 38                                |         |
| 2–5 years  | 27             | 17                                |         |
| >5 years   | 16             | 11                                |         |
| Not clear  | 3              | 13                                |         |
| ISWD m     | 232±185        | 169±177                           | 0.009   |
| RAP mmHg   | 11±5           | 12±6                              | 0.192   |
| mPAP mmHg  | 47±11          | 46±10                             | 0.360   |
| Cardiac index L·min⁻¹·m⁻² | 2.5±0.8     | 2.6±0.8                           | 0.305   |
| PCWP mmHg  | 12±4           | 12±5                              | 0.667   |
| PVR WU     | 7.7±4          | 8±4                               | 0.767   |
| SO₂, %     | 61±8           | 61±9                              | 0.610   |
| FEV₁ % pred| 83±43          | 82±21                             | 0.714   |
| FVC % pred | 95±21          | 96±23                             | 0.702   |
| DLCO % pred| 65±15          | 61±17                             | 0.084   |
| History of VTE | 74       | 69                                | 0.489   |
| History of cancer | 6        | 6                                 | 0.827   |
| Smoking    | 38             | 35                                | 0.664   |
| Obesity    | 39             | 26                                | 0.048   |
| CAD        | 11             | 8                                 | 0.620   |
| LV dysfunction | 2        | 6                                 | 0.131   |
| CKD        | 3              | 14                                | <0.001  |
| COPD       | 13             | 17                                | 0.356   |
| PH therapy following diagnosis | 74       | 75                                | 0.849   |

Data are presented as mean±sd or %, unless otherwise stated. BMI: body mass index; WHO FC: World Health Organization Functional Class; ISWD: incremental shuttle walk distance; RAP: right atrial pressure; mPAP: mean pulmonary arterial pressure; PCWP: pulmonary capillary wedge pressure; PVR: pulmonary vascular resistance; WU: Wood units; SO₂: mixed venous oxygen saturation; FEV₁: forced expiratory volume in 1 s; FVC: forced vital capacity; DLCO: diffusing capacity of the lung for carbon monoxide; VTE: venous thromboembolism; CAD: coronary artery disease; LV: left ventricular; CKD: chronic kidney disease; COPD: chronic obstructive pulmonary disease; PH: pulmonary hypertension. Comparisons between continuous variables and categorical variables were made by the t-test and the Chi-squared test, respectively.
Outcomes
CTEPH (whole cohort)
During a mean±SD follow-up of 4±3 years, 182 (32%) patients died: 51 in the CTEPH-surgical-operated group, 76 in the CTEPH-technically-operable-not-operated group and 49 in the CTEPH-nonsurgical-disease-distribution group. Estimated 1-, 3- and 5-year survival from date of right heart catheterisation was superior in the CTEPH-operated group (97%, 87% and 83%, respectively) compared with both the CTEPH-technically-operable-not-operated group (87%, 63% and 53%, respectively) and CTEPH-nonsurgical-disease-distribution group (92%, 75% and 59%, respectively) (p<0.001) (figure 2).

CTEPH-operated group
Persistent pulmonary hypertension (defined by mPAP ⩾ 25 mmHg) post-PEA surgery was found in 108 (40%) patients in our study. There was no significant difference in long-term survival between the patients who developed pulmonary hypertension post-PEA versus the patients who did not develop pulmonary hypertension post-PEA (p=0.288). However, those with a post-operative PVR above the median (3 Wood units (WU)) had a worse prognosis following surgery (p=0.013). There was no significant difference in survival between patients who received bridging therapy versus patients who did not receive bridging therapy prior to surgery (3-year survival 86% versus 90%; p=0.447), although those who received bridging therapy had more severe pulmonary haemodynamics at diagnosis (mPAP 50 versus 40 mmHg and PVR 8.8 versus 5 WU; both <p<0.001).

CTEPH-technically-operable-not-operated subgroup: declined surgery (patient choice)
The estimated 5-year survival of patients declining surgery was significantly worse than those undergoing PEA (55% versus 83%; p<0.001) (figure 3a). The impact of age on long-term outcome in patients offered surgery is shown in figure 3b–d. A survival benefit was seen in patients aged <60 years (p=0.036) and ⩾60 years (p<0.001) (figure 3b and c, respectively). A trend in favour of surgery was observed in more elderly patients aged ⩾70 years (p=0.056) (figure 3d).

CTEPH-technically-operable-not-operated subgroup: not offered surgery (unfit for surgery, other contributors to symptoms in addition to clot burden)
Patients deemed unfit for surgery had a significantly worse survival than patients offered surgery who declined, who had a significantly worse survival than patients with other contributors to symptoms in addition to clot burden (p<0.05) (supplementary figure S1).

FIGURE 2 Kaplan–Meier estimates of survival from date of diagnosis in the chronic thromboembolic pulmonary hypertension [CTEPH]-operated, CTEPH-technically-operable-not-operated and CTEPH-nonsurgical patient groups.

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**Prognostic indicators**

**CTEPH (whole cohort)**

Univariate analysis of the whole cohort identified a number of predictors of outcome (table 3). Independent predictors of mortality identified from multivariate analysis were: PEA (HR 0.38, 95% CI 0.23–0.63), DLCO (HR 0.59, 95% CI 0.46–0.74), mixed venous oxygen saturation ($SvO_2$) (HR 0.71, 95% CI 0.57–0.87), history of cancer (HR 2.24, 95% CI 1.28–3.95), chronic kidney disease (HR 2.20, 95% CI 1.22–4.71) and age (HR 1.39, 95% CI 1.06–1.80).

**CTEPH-offered-surgery**

Four independent predictors of survival were identified in the combined group of patients offered surgery: patient choice (HR 3.64, 95% CI 1.95–6.81), $SvO_2$ (HR 0.66, 95% CI 0.49–0.89), DLCO (HR 0.67, 95% CI 0.47–0.95) and the presence of coronary artery disease (HR 2.34, 95% CI 1.11–4.96) (table 4).

**CTEPH-technically-operable-not-operated subgroup: declined surgery (patient choice)**

ROC curve analysis for prediction of 3-year mortality was performed for the three continuous prognostic variables identified at univariate analysis (supplementary table S3): DLCO (area under the curve (AUC) 0.87), RAP (AUC 0.81) and $SvO_2$ (AUC 0.85). Using median thresholds of DLCO 62%, RAP 11 mmHg and $SvO_2$ 62%, the sensitivity, specificity, positive predictive value and negative predictive value for predicting...
TABLE 3 Cox regression survival analysis for chronic thromboembolic pulmonary hypertension (CTEPH) (whole cohort)

| Covariate               | Univariate analysis | Multivariate analysis |
|-------------------------|---------------------|-----------------------|
|                         | HR (95% CI)         | p-value               | HR (95% CI)         | p-value               |
| Age                     |                     |                       |                       |                       |
| Per 15 years            | 1.68 [1.40–2.02]    | <0.001                | 1.39 [1.06–1.80]     | 0.016                 |
| BMI                     |                     |                       |                       |                       |
| Per 7 kg·m⁻²             | 0.82 [0.68–0.99]    | 0.046                 |                       |                       |
| WHO FC                  |                     |                       |                       |                       |
| I/II or III/IV; Ref.=I/II| 2.20 [1.12–4.30]   | 0.021                 |                       |                       |
| ISWD                    |                     |                       |                       |                       |
| Per 177 m               | 0.51 [0.41–0.63]    | <0.001                |                       |                       |
| RAP                     |                     |                       |                       |                       |
| Per 5 mmHg              | 1.30 [1.14–1.49]    | <0.001                |                       |                       |
| Cardiac index           |                     |                       |                       |                       |
| Per 0.8 L·min⁻¹·m⁻²     | 0.75 [0.64–0.88]    | <0.001                |                       |                       |
| PVR                     |                     |                       |                       |                       |
| Per 4.3 WU              | 1.36 [1.16–1.60]    | <0.001                |                       |                       |
| SvO₂                    |                     |                       |                       |                       |
| Per 8%                  | 0.66 [0.57–0.76]    | <0.001                | 0.71 [0.57–0.87]     | 0.001                 |
| FEV, #                  |                     |                       |                       |                       |
| Per 35%                 | 0.65 [0.49–0.86]    | 0.002                 |                       |                       |
| DLCO, #                 |                     |                       |                       |                       |
| Per 17%                 | 0.52 [0.44–0.62]    | <0.001                | 0.59 [0.46–0.74]     | <0.001                |
| VTE                     | Ref.=absent         |                       |                       |                       |
| Cancer                  | Ref.=absent         |                       |                       |                       |
| Obesity                 | Ref.=absent         |                       |                       |                       |
| Thyroid disorders       | Ref.=absent         |                       |                       |                       |
| CAD                     | Ref.=absent         |                       |                       |                       |
| LV dysfunction          | Ref.=absent         |                       |                       |                       |
| CKD                     | Ref.=absent         | 0.001                 | 2.20 [1.22–4.71]     | 0.021                 |
| PEA                     | Ref.=not performed  | 0.31 [0.22–0.43]     | 0.38 [0.23–0.63]     | <0.001                |

Data shown for univariate analysis where p<0.20; 72 variables were imported into univariate analysis. HR: hazard ratio; Ref.: reference parameter; BMI: body mass index; WHO FC: World Health Organization Functional Class; ISWD: incremental shuttle walk distance; RAP: right atrial pressure; mPAP: mean pulmonary arterial pressure; PVR: pulmonary vascular resistance; WU: Wood units; SvO₂: mixed venous oxygen saturation; FEV₁: forced expiratory volume in 1 s; DLCO: diffusing capacity of the lung for carbon monoxide; VTE: venous thromboembolism; CAD: coronary artery disease; LV: left ventricular; CKD: chronic kidney disease; PEA: pulmonary endarterectomy. # these variables are scaled so that the hazard ratio is the change by 1 SD.

3-year mortality were 100%, 63%, 31% and 100%, 80%, 70%, 32% and 95%, and 90%, 60%, 30% and 97%, respectively.

CTEPh-technically-operable-not-operated group

For the whole group, univariate analysis identified a number of predictors of outcome, including age, World Health Organization Functional Class, exercise capacity, pulmonary haemodynamics and comorbidities. Univariate and multivariate analysis for each of the three subgroups is shown in supplementary table S3. Independent predictors of outcome were SvO₂ (HR 0.53, 95% CI 0.38–0.76), DLCO (HR 0.54, 95% CI 0.38–0.75) and cancer (HR 4.10, 95% CI 2.02–8.37). In those who declined surgery, SvO₂ (HR 0.24, 95% CI 0.12–0.51) was an independent predictor of survival; in those unfit for surgery, exercise capacity (HR 0.51, 95% CI 0.26–0.96), PVR (HR 1.56, 95% CI 1.02–2.38) and comorbidities (cancer (HR 11.46, 95% CI 3.46–37.99) and chronic kidney disease (HR 8.87, 95% CI 2.45–32.17)) independently predicted outcome, while in patients with other contributors to symptoms in addition to clot burden, cancer (HR 9.93, 95% CI 1.98–49.85) was an independent predictor of outcome.

Discussion

To the best of our knowledge, this is the first study primarily focusing on patients with technically operable CTEPH who did not undergo PEA. Patient choice, lack of fitness for surgery and the presence of other contributors to symptoms in addition to clot burden were the commonest reasons for patients not undergoing surgery, while pulmonary haemodynamic severity, DLCO and comorbidities were independent predictors of survival. In addition, in a large cohort of consecutive patients with CTEPH we have shown that long-term survival of patients undergoing PEA is excellent and superior to patients declining surgery, strongly favouring consideration of a surgical intervention in eligible patients.

CTEPh (whole cohort)

PEA is considered the treatment of choice for suitable patients with CTEPH, and is thought to provide the best prospect of improved quality and quantity of life [17–19]. Our study, conducted in a large cohort of consecutive patients with CTEPH, confirms the results of the international CTEPH Registry that PEA is an independent predictor of survival [24]. In operated patients it was associated with an excellent long-term outcome with an estimated 5-year survival of 83%, similar to data from the international [24],
Although 482 patients (82%) had technically operable disease distribution, despite the proven benefits of PEA only 272 (49% of the total cohort) underwent surgery. Previous registries reported similar proportions of patients who underwent PEA, but provided only limited data on reasons for not undergoing surgery and predictors of long-term outcome [24, 34, 35].

CTEPH-technically-operable-not-operated group

176 (39% of patients with technically operable disease) did not undergo surgery due to patient choice (n=72), concerns regarding fitness to undergo surgery (n=63) or having other contributors to symptoms in addition to clot burden (n=31 where the benefits of surgery were felt to be minimal). The 5-year survival in patients with technically accessible disease not undergoing surgery was 53%, significantly better than historical studies of patients with CTEPH treated with anticoagulation alone [22, 23]. Survival was related to the rationale underpinning the treatment decision, with those declining surgery having a superior survival to those who were deemed unfit for surgery. Not only did markers of disease severity such as $SvO_2$ and $DLCO$ independently predict survival, but also the presence of comorbidities, emphasising the impact of conditions apart from the pulmonary vasculature when making treatment decisions.

CTEPH-technically-operable-not-operated subgroup: declined surgery (patient choice)

The proportion of patients with technically operable disease who were offered surgery (n=344) but declined (n=72 (21%)) is larger than previously noted in other registries and may reflect the consecutively enrolled nature of our study [24, 34, 35]. These data highlight the importance of patients being referred for assessment and counselling by a PEA surgeon and experienced MDT. In patients who declined surgery the severity of pulmonary haemodynamics and $DLCO$ predicted outcome, with median thresholds for $DLCO$, right atrial pressure and $SvO_2$ having negative predictive values for 3-year mortality >95%. For a selected cohort of patients who despite counselling decline surgery this information may be useful, although it must also be emphasised that quality of life benefits in the majority of patients are greater with surgery [36, 37]. In contrast, patients with severe haemodynamics assessed to be good surgical candidates may find data highlighting a poor prognosis in the absence of a surgical intervention an aid to decision making.

This study has demonstrated a significantly superior survival in patients with CTEPH who were offered surgery and underwent PEA compared with those who declined. Although patients declining surgery were older, with a poorer exercise capacity and more comorbidities, declining surgery due to patient choice was

### Table 4: Cox regression survival analysis in chronic thromboembolic pulmonary hypertension (CTEPH)-technically-operable who were offered surgery (operated group) and declined surgery (patient choice) subgroups

| Covariate                  | Univariate analysis | Multivariate analysis |
|----------------------------|---------------------|-----------------------|
|                            | HR (95% CI)         | p-value               |
|                            |                     |                       |
| Age: per 15 years          | 1.71 [1.30–2.25]    | <0.001                |
| WHO FC: I/II or II/IV; Ref.=I/II | 3.98 [1.25–12.65]   | 0.019                 |
| Cardiac arrhythmia: Ref.=absent | 2.16 [1.03–4.53]    | 0.043                 |
| ISWD: per 185 m            | 0.56 [0.41–0.75]    | <0.001                |
| RAP: per 6 mmHg            | 1.57 [1.25–1.98]    | <0.001                |
| PVR: per 4 WU              | 1.39 [1.08–1.78]    | 0.009                 |
| $SvO_2$: per 8%            | 0.62 [0.49–0.77]    | <0.001                |
| $DLCO$: per 16%            | 0.56 [0.42–0.75]    | <0.001                |
| Cancer: Ref.=absent        | 0.62 [0.39–0.98]    | 0.045                 |
| Obesity: Ref.=absent       | 1.77 [0.85–3.69]    | 0.127                 |
| Thyroid disorders: Ref.=absent | 0.55 [0.33–10.92]   | 0.024                 |
| CAD: Ref.=absent           | 2.21 [1.24–3.94]    | 0.007                 |
| CKD: Ref.=absent           | 1.90 [0.82–4.38]    | 0.132                 |
| Patient choice: Ref.=surgery | 2.56 [1.57–4.16]    | <0.001                |
|                            | 3.64 [1.95–6.81]    | <0.001                |

Data shown for univariate analysis where p<0.20; 72 variables were imported into univariate analysis. HR: hazard ratio; Ref.: reference parameter; WHO FC: World Health Organization Functional Class; ISWD: incremental shuttle walk distance; RAP: right atrial pressure; PVR: pulmonary vascular resistance; WU: Wood units; $SvO_2$: mixed venous oxygen saturation; $DLCO$: diffusing capacity of the lung for carbon monoxide; VTE: venous thromboembolism; CAD: coronary artery disease; CKD: chronic kidney disease. #: these variables are scaled so that the hazard ratio is the change by 1 SD.
an independent predictor of a worse outcome. In patients who declined compared with those who underwent surgery, there was a female predominance [27] and sex-specific factors related to risk-taking may play a role [38]. Given the findings of this study and the benefit of PEA, more work is required to understand the factors underlying decisions to decline surgery.

**Unfit for surgery**
In expert hands PEA has a perioperative mortality of <5% and offers the best chance of longer-term survival, but requires careful assessment of risks versus benefits for individual patients [21, 24]. We found 63 patients (14% of those with technically operable disease) were deemed unfit for surgery by the MDT. These patients had a significantly poorer survival than those declining surgery. Alternative interventions to surgery such as balloon pulmonary angioplasty may appear attractive in these patients, but the presence of significant comorbidities may be primary determinants of survival. A meticulous assessment balancing the potential symptomatic benefit versus the risks of such interventions is therefore paramount where mortality benefit is not clear.

**CTEPH-nonsurgical-disease-distribution group**
19% of all patients were deemed to have nonsurgical disease distribution. This proportion is similar to the international CTEPH Registry (20%) [24], but less than the UK (32%) [35] and Dutch (26%) [34] registries. This may reflect an increasing willingness among surgeons to operate on patients with disease that would previously have been considered too distal to benefit from surgery. Indeed, a number of centres have shown that outcomes in patients with type 3 disease (more distal) in expert hands are now similar to more proximal disease (type 1 and 2) [21, 39]. We noted a female predominance, an increased incidence of thyroid disease and splenectomy, and reduced incidence of VTE in this group (table 1), in keeping with previous reports [33].

**CTEPH-operated group: timing of surgery and pulmonary vasodilator therapy**
The mean±SD time from diagnosis to PEA surgery was 290±175 days, longer than in previously reported studies [24, 32, 35, 40], but in line with waiting times for surgery in the UK during the duration of this study, although UK waiting times are now falling. In keeping with data from the international CTEPH Registry, the duration of delay did not impact on long-term survival [24]. A large number of patients were bridged to PEA with off-label pulmonary arterial hypertension-specific therapies (74%). Although there is no published evidence to support this practice, this may reflect the longer time from referral to surgery in the UK during the study period compared with that reported in the international CTEPH Registry [24]. Importantly, bridging therapy had no effect on time to referral or to surgery. Patients receiving bridging therapy in the present study had similar haemodynamics to those who received bridging therapy in the international CTEPH Registry, while patients who were not bridged to surgery had milder haemodynamics than those in the international CTEPH Registry [24]. Importantly, in our study receiving bridging therapy was not associated with adverse outcome at univariate analysis.

**Limitations**
This study pre-dates the availability of balloon angioplasty and riociguat therapy in the UK, and therefore the impact of these interventions and their potential benefits cannot be assessed. Although patient-specific data were enriched by retrospective case note review and interrogation of databases, this resulted in higher levels of data completeness than in other contemporary registries. Furthermore, the consecutive nature of enrolment in the ASPIRE registry reduces recruitment bias associated with previous nonconsecutively enrolled studies. This study provides no data on the reasons for patients declining surgery. Although the results suggest that surgery improves survival in patients with technically operable disease who were offered surgery, patients judged to be unfit for surgery by the MDT and those in whom there were other contributors to symptoms in addition to clot burden were excluded. For the individual patient, factors including age and comorbidities will influence outcome following surgery. How these factors influence the patient’s decision requires further research.

**Conclusions**
We report results from a large consecutively enrolled registry of patients with CTEPH, and have been able to compare characteristics and have identified predictors of survival in patients who did not undergo surgery despite having technically operable disease. Our data show that survival of patients undergoing PEA is excellent and superior to patients declining surgery, and strongly favours consideration of a surgical intervention in eligible patients. More work is required to understand factors influencing decision making in CTEPH, and to ensure that patients are counselled and supported to make informed decisions.
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