

Vocal Cord Paralysis After Lung Transplantation Does Not Influence Early Outcome

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BACKGROUND

- Vocal Cord Paralysis (**VCP**) is frequent after cardiothoracic surgery
- Incidence and effect on outcome in lung transplantation (**LTx**) is unknown
- VCP may cause chronic aspiration, which is linked to development of chronic lung allograft dysfunction (**CLAD**)

AIM / METHODS

- AIM:** 1 - to assess incidence of VCP after bilateral LTx
2 - to assess impact on clinical outcomes
- VCP** = inability to achieve vocal cord apposition at 2 occasions assessed by flexible bronchoscopy
- Retrospective assessment of bronchoscopy records 01/10-06/15
- Clinical endpoints: graft survival; CLAD onset; hospitalizations; lower respiratory tract infections (LRTI) in VCP vs comparators (using 5:1 propensity matching).

RESULTS

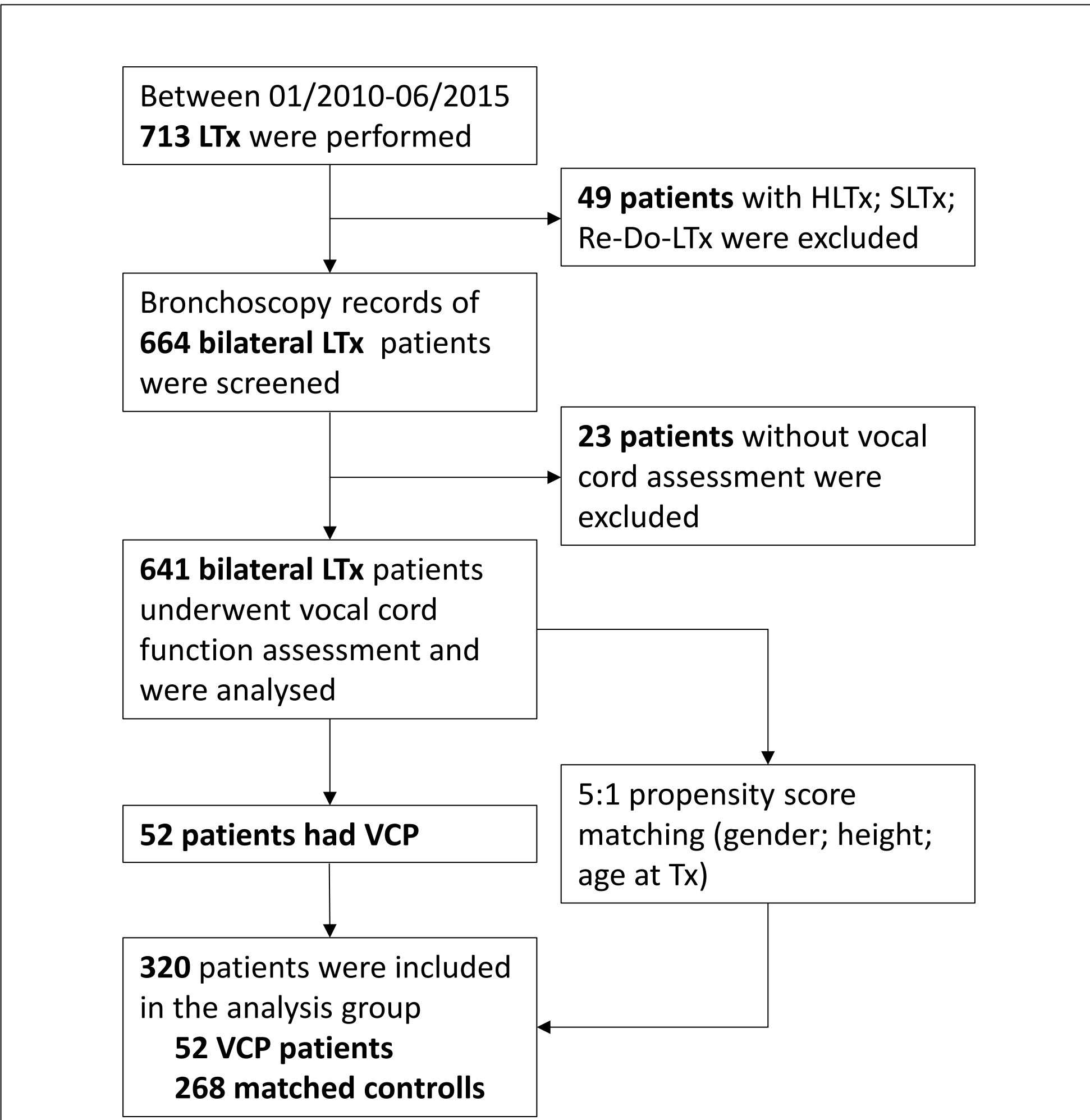


Figure 1. Study population and matching

Table 1. VCP characteristics	
Finding, n (%)	All Patients with VCP (n=52)
Affected side	
Left	40 (77)
Right	9 (17)
Bilateral	3 (6)
Recovery	34 (65)
Time to recovery, months (IQR)	6 (2-12)
Persistent VCP	18 (35)
Symptomatic	15 (29)
Treatment received	15 (29)

Table 2. Secondary clinical endpoints				
Endpoint	VCP (n=52)	Control (n=268)	Hazard Ratio (95% CI) *	p
ICU re-intubation	8 (15)	38 (14.2)	1.1 (0.48-2.52)	0.821
Bronchial stenosis	10 (19)	48 (18)	1.06 (0.50-2.27)	0.872
Pulmonary function test, % of predicted (IQR)				
FEV ₁	93 (82-104)	92 (75-109)		0.835
FVC	98.5 (85-116)	102 (89-117)		0.353
MEF 25-75%	90.5 (78-118)	89 (63-117)		0.243
Bronchoscopies*	7 (6-9)	8 (6-10)		0.181

*within first 24 months following LTx after initial hospital discharge

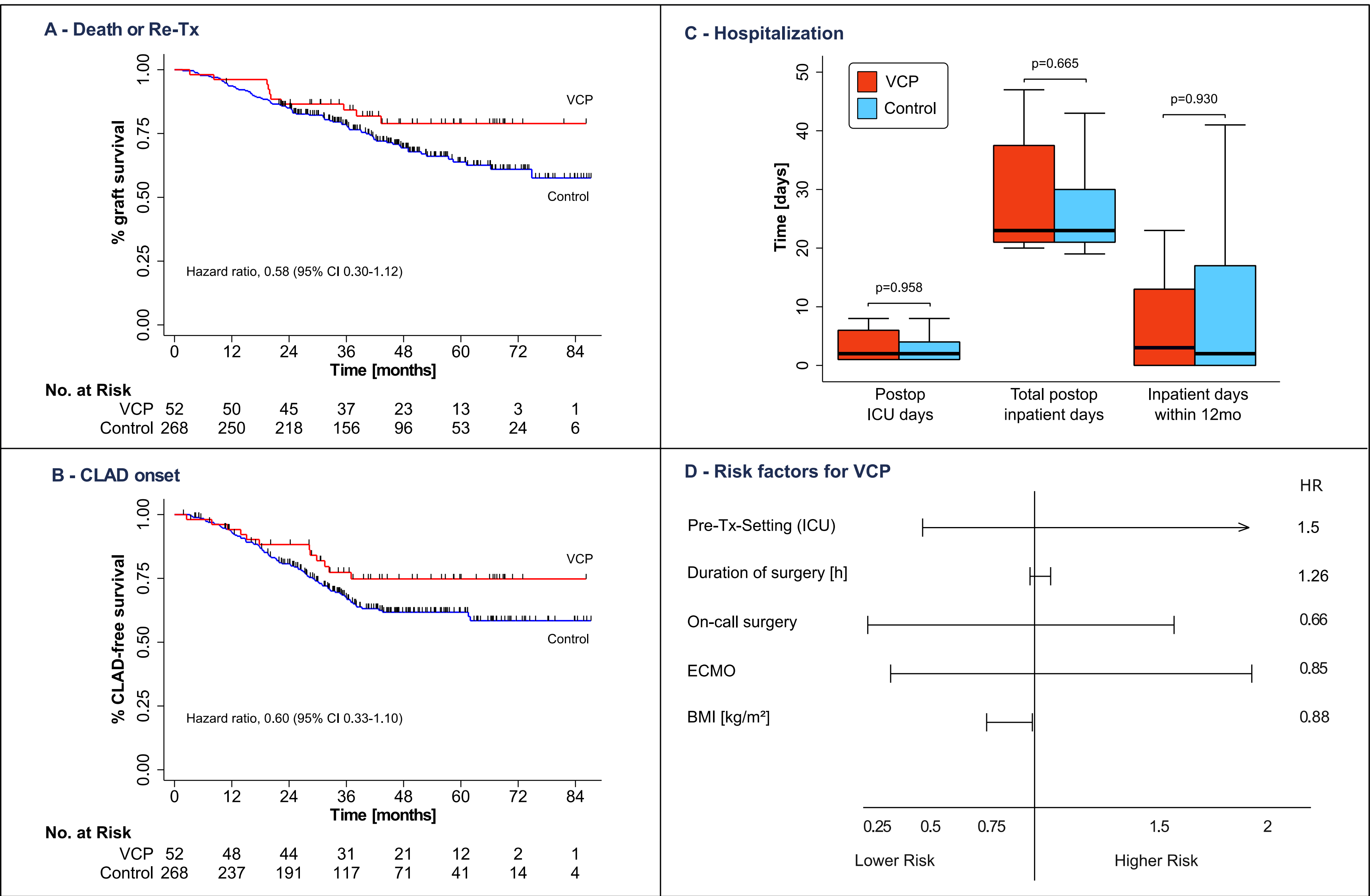


Figure 2. A) Graft survival (Death or Re-Tx) in VCP vs controls. B) Chronic allograft dysfunction (CLAD) in VCP vs controls. C) Median post-op ICU stay, median post-op hospital stay in day and subsequent inpatient days within first year after Tx. D) Hazard ratios for potential risk factors for occurrence of VPC following Tx (with 95% CI).

CONCLUSIONS

- VCP was present in 52/641 bilateral LTx recipients (8.1%) (**Figure 1**).
- VCP was transient in most patients with full recovery in 65% (**Table 1**).
- Using 5:1 propensity matching, VCP was not associated with increased graft –loss or CLAD-onset (**Figure 2A+B**).
- Duration of hospitalization including ICU-stay was not increased in the VCP-group (**Figure 2C**).
- Lower BMI is a potential risk factor for VCP (**Figure 2D**).
- There was no increased rate of re-intubation or development of bronchial stenosis in the VCP group (**Table 2**).
- Overall, VCP did not adversely impact clinical outcome in LTx-recipients.**