

# Sleeve lobectomy may provide better outcomes than pneumonectomy for non-small cell lung cancer. A decade in a nationwide study

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#### ▶ To cite this version:

Pierre-Benoit Pagès, Pierre Mordant, Stéphane Renaud, Laurent Brouchet, Pascal-Alexandre Thomas, et al.. Sleeve lobectomy may provide better outcomes than pneumonectomy for non-small cell lung cancer. A decade in a nationwide study. Journal of Thoracic and Cardiovascular Surgery, 2017, 153 (1), pp.184+. 10.1016/j.jtcvs.2016.09.060. hal-01489668

# HAL Id: hal-01489668 https://hal.science/hal-01489668v1

Submitted on 26 Mar 2019

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## **Accepted Manuscript**

Sleeve lobectomy may provide better outcomes than pneumonectomy for non-small cell lung cancer. A decade in a nationwide study

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PII: S0022-5223(16)31275-2

DOI: 10.1016/j.jtcvs.2016.09.060

Reference: YMTC 10927

To appear in: The Journal of Thoracic and Cardiovascular Surgery

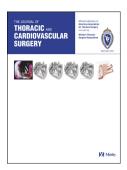
Received Date: 26 January 2016

Revised Date: 15 September 2016

Accepted Date: 23 September 2016

Please cite this article as: Pagès P-B, Mordant P, Renaud S, Brouchet L, Thomas P-A, Dahan M, Bernard A, for the EPITHOR project (French Society of Thoracic and Cardiovascular Surgery), Sleeve lobectomy may provide better outcomes than pneumonectomy for non-small cell lung cancer. A decade in a nationwide study, *The Journal of Thoracic and Cardiovascular Surgery* (2016), doi: 10.1016/j.jtcvs.2016.09.060.

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- **Word count:** 3688 Words
- 18 **Abstract:** 244 words

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#### Conflicts of interest and source of funding

29 The other authors have no disclosure to declare for this work.



30	Abstrac	t

database.

Introduction: Whenever feasible, sleeve lobectomy is recommended to avoid pneumonectomy for lung cancer, but these guidelines are based on limited retrospective series. The aim of our study was to compare outcomes following sleeve lobectomy and pneumonectomy using data from a national

**Methods:** From 2005 to 2014, 941 sleeve lobectomy and 5318 pneumonectomy patients were recorded in the French database Epithor. Propensity score was generated with 15 pretreatment variables and used to create balanced groups with matching (794 matches) and inverse probability of treatment weighting (standardized difference was 0 for matching, and 0.0025 after weighting). Odds ratio of postoperative complications and mortality, hazard ratio for overall survival and disease-free survival were calculated using propensity adjustment techniques and a sensitivity analysis.

**Results:** Postoperative mortality after sleeve resection was similar to that after pneumonectomy (Matching OR 1.24, p=0.4; weighting OR 0.77, p=0.4) despite significantly lower odds of pulmonary complications with pneumonectomy (matching OR 0.4, p<0.0001; weighting OR 0.12, p<0.001). The adjusted hazard for death after pneumonectomy was significantly higher when analyzed using matched analysis but not with weighting (Matching HR 1.63, p=0.002; weighting HR 0.97, p=0.92). The same was true for disease-free survival (Matching HR 1.49, p=0.01; weighting HR 1.03, p=0.84).

**Conclusions:** Despite early differences in perioperative pulmonary outcomes favoring pneumonectomy, early overall and disease-free survival were in favor of sleeve lobectomy in the matched analysis but not the weighted analysis. In our opinion, when it is technically feasible, sleeve lobectomy should be the preferred technique.

**Key-words:** Pneumonectomy, sleeve lobectomy, survival, outcomes

56 Central messag	ge
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- 57 There were no differences in postoperative mortality between the two techniques. Sleeve lobectomy
- increases the risk of pulmonary complications.

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## Perspective statement

- 61 Sleeve lobectomy leads to higher rate of overall and disease-free survival despite an increased risk of
- 62 postoperative pulmonary complications. We think that when it is technically possible, sleeve
- 63 lobectomy has to be the type of resection to favor for central tumor.

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## **Central Picture**

Weighting analysis did not highlight any benefit in overall survival for sleeve lobectomy.

## 68 Abbreviations

- 69 ARDS: Acute Respiratory Distress Syndrome
- 70 ASA: American Society of Anesthesiologists
- 71 BMI: Body Mass Index
- 72 BPF: Bronchopleural Fistula
- 73 CI: Confidence Interval
- 74 DFS: Disease-Free Survival
- 75 FEV1: Forced Expiratory Volume in one second
- 76 HR: Hazard Ratio
- 77 IPTW: Inverse Probability for Treatment Weighting
- 78 NSCLC: Non-small cell lung cancer
- 79 OR: Odds Ratio
- 80 OS: Overall Survival
- 81 PN: Pneumonectomy
- 82 PS: Propensity Score
- 83 SL: Sleeve Lobectomy
- 84 RCT: Randomized Control Trial
- 85 WHO: World Health Organization

#### Manuscript

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#### Introduction

Complete surgical resection is the cornerstone of the management of localized non-small cell lung cancer (NSCLC). For years, surgical resection was restricted to segmentectomy, lobectomy, and pneumonectomy (PN), the last being associated with the highest postoperative morbidity and mortality [1]. Sleeve Lobectomy (SL) was first described for carcinoma by Allison, in 1959, in order to avoid the burden of PN in frail patients with compromised lung function [2]. Given the morbidity and mortality associated with PN, sleeve lobectomy (SL) was initially restricted to patients with compromised lung function who would not tolerate PN. However, the indication for SL has progressively expanded to any tumor that may be completely resected using this technique, particularly on the right side [3]. As a result of this progressive expansion of SL over PN, recent guidelines from the American College of Chest Physicians recommend SL rather than PN in patients with clinical stage I or II central NSCLC in whom complete resection can be achieved [4]. However, these guidelines are based on short surgical series comparing recent SL to historical PN, or meta-analyses that included patients operated on over many decades [5]. No database analysis has been published to date even though every lung cancer surgeon is now able to perform SL and PN, and recent publications have highlighted the impact of surgery improvements on the prognosis of NSCLC during recent decades [6]. We therefore sought to compare short and long-term outcomes following SL and PN for NSCLC during the last decade in France. For this purpose, we used the French national database Epithor and two PS methods: matching and the inverse probability of treatment weighting (IPTW), combined with a

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#### **Materials and Methods**

sensitivity analysis.

#### Data Collection

113 Epithor is a government-recognized clinical database, financially supported by the National Cancer

Institute (Institut National du Cancer) for data-quality monitoring. Epithor is accredited by the French

Health Authorities (Haute Autorité de Santé), a governmental agency dedicated to improving the quality of patient care and to guaranteeing equality within the health care system, as a methodologically appropriate tool to assess professional surgical practices. Participating in Epithor is now a requirement for medical accreditation and thoracic surgery unit certification in France [7,8]. The accuracy of data collection is checked in regular external on-site audits initiated in 2010 [7]. Data are sent by Internet to the national database; surgeons and patients are anonymous. Surgeons can check the quality of the way they enter the data by comparing their data with national data through a quality score ranging from 0% to 100%. Moreover, participants have to check the quality of the local database for missing values by comparing its completeness with that of the national database. This comparison is expressed through a quality score ranging from 0 to 100%. A score exceeding 80% is mandatory to have the local data incorporated in the national database and to benefit from the accreditation. Every surgeon receives a personal quality score, thus inciting them to update their data. This induces a virtuous cycle. Almost all of the teams that participate in Epithor have a score above 80% for data entry. The Institutional Review Board of the French Society of Thoracic and Cardiovascular Surgery approved the electronic prospective database used for this study and the study itself. Patients' consent was obtained, and patients were aware that the data collected would be used for clinical research purposes. The institutional review board of the French Society of Thoracic and Cardiovascular Surgery certifies that this study respects the current regulations that govern clinical research in France, referenced as: "CERC-SFCTCV-2015-8-14-16-5-39-PAPi.

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#### Study population

From January 2005 to December 2014, 6259 patients underwent SL or PN for NSCLC in 103 centers in France. The baseline demographic and clinical characteristics included age, sex, American Society of Anesthesiologists (ASA) score, World Health Organization (WHO) performance status, body mass index (BMI), the Forced Expiratory Volume in 1 second (FEV1) as a percentage and the dyspnea score according to the Medical Research Council [9]. The number of comorbid diseases per patient

was considered a categorical variable because recent data from Epithor consistently suggested that this variable was superior to individual comorbidities in a predictive model for operative mortality [10]. Systematic nodal dissection included node sampling or radical lymphadenectomy. NSCLC histology was classified according to the most recent WHO classification [11]. Tumor and Nodal stages were classified postoperatively according to the pathologic examination and the most recent IASLC classification [12].

#### Outcome measurements

The primary endpoint was postoperative complications, which included cardiopulmonary morbidity, bronchopleural fistula (BPF), empyema, and hemorrhage. Cardiopulmonary morbidity was reported as proposed by the European Society of Thoracic Surgery and included postoperative pulmonary complications (pneumonia, atelectasis requiring bronchoscopy, adult respiratory distress syndrome (ARDS), mechanical ventilation for more than 2 days and arrhythmia [13].

The secondary endpoints were (i) postoperative mortality, defined as any patient who died within the first 30 days following surgery, or the initial hospitalization if longer; (ii) length of hospital stay, (iii)

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visit; and (iiii) disease-free survival (DFS), defined as the time from surgery until disease recurrence

overall survival (OS), defined as the time from surgery until death from any cause or the last follow-up

or the last follow-up visit.

## Variables used for PS analysis

Variables used to estimate the PS were age, sex, performance status, number of comorbidities, dyspnea score, FEV1, BMI, induction chemotherapy, side, histology, T status, N status, year of surgery, type of center and hospital volume.

#### Missing data

The proportion of missing FEV1 for this study was 7%. We assumed that the missing data were missing at random. We applied a multiple imputation framework (20 imputations) to compensate for missing prognostic factor data for FEV1.

For missing data for tumor pathological features, lymph nodes and histology, we created a variable category to include in the analysis. For each hospital, the hospital volume was calculated for the period from 2005 to 2014. Centers were ranked by number of procedures performed per year. We created a categorical variable detailing five categories of hospital volume.

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#### Statistical analysis

The PS is the conditional probability of assignment to a particular treatment given a vector of observed covariates [14]. PS techniques were used to balance the distributions of measured potentially confounding covariates for patients treated by SL or PN. A mirrored histogram was used to measure the discriminatory ability of PS matching, and the standardized difference for the IPTW analysis. Matching and IPTW tend to eliminate systematic differences between experimental and control subjects to a greater degree than does stratification or covariate adjustment [15,16]. Matching used a search algorithm to find a set of weights for each covariate such that the version of optimal balance is achieved after matching [17]. We used 1:1 matching without replacement in descending order with a caliper of 0.01. With IPTW, each individual is weighted by the inverse probability of receiving the treatment that they actually received. In this way, each group is weighted to represent the full population sample, thus revealing treatment effects. We evaluated two PS techniques in their ability to balance the measured covariates between SL and PN by reducing the standardized difference [16]. The standardized difference is the difference between sample means in the SL and PN group divided by the standard deviation in the treatment group overall [17]. Finally, odds ratios (OR) were used for dichotomous variables such as postoperative mortality, atelectasis, pneumonia, arrhythmia, BPF, empyema and hemorrhage, ventilation > 2 days and ARDS. Logistic models were used. For the length of hospital stay, the difference of means was used. Logistic models and linear regression were used. For OS and DFS, we used the adjusted Kaplan-Meier estimators for curves using IPTW data and the log-rank test. Cox proportional hazards models were used to estimate Hazard ratios (HR). Hospital level clustering was used to estimate the robust standard error for each coefficient of the logistic model, linear regression and Cox model.

199	Sensitivity analysis
200	The Q Mantel-Haenszel test statistic was used; if $\acute{\Gamma}=1$ , the statistic test is significant and the study is
201	free of hidden bias [19]. And if the value of the Q Mantel-Haenszel test is greater, then the study is
202	insensitive to bias [20].
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204	Results
205	Study cohort
206	From 2005 to 2014, 941 SL and 5318 PN were included in the Epithor database and make up the study
207	cohort. As compared to the PN group, patients in the SL group were characterized by a younger age
208	(median 62 years old for both groups), higher BMI (median 25 for both groups), higher FEV1, lower
209	ASA score, lower WHO status, and less frequent induction therapy (Table 1). As compared to the PN
210	group, the SL group was characterized by the predominance of right-sided surgery and squamous cell
211	carcinoma. T and N stages in the SL group were lower than those in the PN group (Table 1).
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213	Surgery
214	More than half of the SL involved the right upper lobe, followed by the left upper, left lower and right
215	lower respectively. Middle lobectomy and bilobectomy SL occurred in less than 5% of patients each
216	(Table 1). Both SL and PN were predominantly performed in teaching hospitals. SL was more
217	frequently performed in high-volume centers, whereas PN was more frequently performed in low-
218	volume centers. The number of SL increased in the last decade whereas the number of PN decreased
219	steadily (Table 1).
220	
221	PS estimation
222	The mirrored histogram shows the good distribution of the covariates after PS matching (Figure 1).
223	The median distribution of standardized biases was 0.024 for the matching approach (1st and 3rd
224	quartile: 0.013-0.0355), and 0.04 for IPTW (1st and 3rd quartile: 0.0145-0.0654). The standardized
225	difference never reached the value of 10%, which highlighted that the two groups were well balanced
226	for covariates by matching and IPTW [19] (Table 2)

#### Postoperative Mortality

Postoperative mortality was 4.99% (n=47) in the SL group and 5.89% (n=313) in the PN group (p=0.279). There were no significant differences in post-operative mortality between SL and PN, according to the 2 PS methods, with an OR associated with PN of 1.24 (95%CI) [0.74-2.1] for matching, and 0.77 (95%CI) [0.4-1.5] for IPTW (OR=1 for SL, **Table 3**).

#### Postoperative complications and Length of Hospital stay

Postoperative pulmonary complications overall (pneumonia, atelectasis requiring bronchoscopy, ARDS, mechanical ventilation for more than 2 days) were significantly less frequent in the PN group than in the SL group according to the two PS methods (**Table 3**). In detail, only atelectasis and pneumonia were significantly less frequent following PN whatever the PS analysis performed. The two PS methods showed a trend towards a higher rate of postoperative arrhythmia, and a significantly higher rate of BPF and empyema in the PN group than in the SL group (**Table 3**). The incidence of hemorrhage was significantly higher in the PN group but only by matching (**Table 3**). BPF occurred in 2.6% of patients in the PN group (n=138) and 1.59% in SL group (n=15) (p=0.067). In the PN group, there was no significant difference in the occurrence of BPF whether the patient receive preoperative irradiation (3.17%, n=4) or not (2.34%, n=122) (p=0.679). Length of hospital stay was significantly shorter in the SL group than in the PN group by matching but not by IPTW analysis (**Table 3**).

## Overall Survival and Disease-free Survival

The median follow-up time was 10.89 months (1st and 3rd quartile: 1.66-15) for OS and 9.6 months (1st and 3rd quartile: 1-14) for DFS. From the 941 patients of the SL group, 463 were evaluable at 1 month, 174 at 12 months, 39 at 36 months and 10 at 60 months. From the 5318 patients of the PN group, 2222 were evaluable at 1 month, 719 at 12 months, 154 at 36 months and 20 at 60 months. Three-year OS was  $71.86 \pm 3.29\%$  for the SL group and  $60.76 \pm 1.69\%$  for the PN group. As compared with SL, there was an increased risk of death in the PN group by matching, with HR of 1.63 [1.19-2.21], but not by the IPTW method (Table 3, Figure 2).

Three-year DFS was  $46.41 \pm 4.01\%$  for the SL group and  $31.63 \pm 1.59\%$  for the PN group. As compared with SL, there was an increased risk of recurrence in the PN group by matching, with HR of

1.49 [1.1-2], but not by the IPTW method (Table 3 & Figure 3).

#### Sensitivity analysis

For the matching analysis, the Q Mantel-Haenszel (Qmh) statistic test was used and showed a potential hidden bias (Qmh value close to 1) only for postoperative mortality (**Supplementary Table 4**). For postoperative complications, the Qmh value was high, indicating the study was insensitive to a bias that would the odds [19]. These results confirmed that compared to SL, PN was associated with a significant decrease in the rate of pulmonary complications overall (pneumonia, atelectasis requiring bronchoscopy, ARDS, mechanical ventilation for more than 2 days), a significant decrease in the rate of atelectasis and pneumonia considered separately, a non-significant increase in the rate of arrhythmias, and a significant increase in the risk of BPF, empyema and hemorrhage. For the IPTW method, the sensitivity analysis confirmed these data, except for arrhythmias and hemorrhages (**Supplementary Table 5**).

#### Discussion

#### Reminder of the main results

In studying outcomes after PN and SL for NSCLC using data from a nationwide database, two PS methods concluded that SL was not associated with any significant difference in postoperative mortality, but with a significant increase in the rate of pulmonary complications (atelectasis, pneumonia), and a significant decrease in the rate of bronchopleural fistulae and empyema as compared with PN. Matching analysis also found that SL was associated with improved 3-year OS as compared with PN, but not IPTW analysis.

#### Treatment allocation

Initially developed for patients with insufficient pulmonary reserve, which did not allow PN, SL has now become widely accepted as a reliable and safe procedure to allow the complete resection of

NSCLC [22]. Indeed, SL makes it possible to spare lung parenchyma, and thus provides better postoperative FEV1 than that achieved with PN [23–25]. Quality of life after SL is also better than that after PN, as highlighted by Balduyck et al., who reported a high burden of dyspnea, general pain, thoracic pain and shoulder dysfunction after PN [26]. This is the first time, however, that the impact of SL on postoperative complications, mortality, and long-term survival had been reported in the context of a large study based on a national database, which allows the analysis of real-life events. In this respect, one can assume that over the last decade all French thoracic surgeons were able to perform both procedures. The choice to perform one procedure rather than the other was therefore based on disease severity and the patient's clinical status rather than technical preferences. This probable difference between groups in baseline characteristics of patients and tumors constitutes the strongest argument in favor of adequate statistical analysis, including PS analysis, of this large real-life dataset.

#### Postoperative mortality and postoperative complications

We found no significant difference regarding postoperative mortality between SL and PN groups, whatever the statistical method used. In a recent paper from the Epithor Group, Morgant et al. reported PN and bronchial SL to be major prognostic factors associated with postoperative death following lung cancer surgery, with OR ranging from 4.4 to 8.2 for PN and 2.4 to 2.9 for bronchial SL [6]. In a recent meta-analysis that included more than 3800 patients, Shi et al. showed a significant difference in postoperative mortality favoring SL over PN with an OR of 0.5 [0.34-0.72] [5]. Similarly, in a matching analysis comparing SL and PN, Park et al. found postoperative mortality to be significantly lower in the SL group (1%) than in the PN group (8.6%, p<10-4) [27]. In contrast, we found no difference between SL and PN for postoperative mortality, despite a significantly greater incidence of BPF and empyema after PN. We found no impact of preoperative irradiation on the occurrence of BPF after PN, as already highlighted in a recent paper establishing a predictive score for BPF [28]. This absence of difference can be attributed either to the low mortality associated with PN, or to the high mortality associated with SL in our study.

On the one hand, recent publications tend to disfavor PN, which, for example, was associated with a

mortality rate of 7.8% over the last decade in the French database [28]. On the other hand, in our

study, mortality following SL remained far higher than that reported in the most recent studies [5,22,24], and markedly higher than the 2.7% reported after regular lobectomy in the same database [7]. Indeed, in our study, postoperative atelectasis and pneumonia were significantly more frequent after SL than after PN, whatever the statistical method considered. This high rate of parenchymal complications may explain the increased postoperative mortality after SL than after regular lobectomy [7]. Altogether, these data suggest that the improvement in patient selection and postoperative care developed in PN should be applied to SL in order to decrease its postoperative morbidity and mortality.

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## Overall Survival and Disease-free Survival

We found that OS was lower in the PN group than in the SL group only by matching. Similar results were reported in the meta-analysis by Shi et al, with a combined HR of 0.63 [0.56-0.71], and in the matching analysis by Park et al. (5-year OS for PN vs SL, 32.1% vs 58.4%, respectively, p=0.0002) [5,27]. In our study, patients undergoing PN were significantly more likely to have T3 and T4 tumors, and N2 lymph-node involvement than were patients undergoing SL. The TNM classification alone might lack precision: T stages have recently been separated according to tumor size [29], the N stage might need to be deciphered according to the number of lymphatic chains and stations involved [30], and the number of molecular biomarkers is constantly increasing [31]. However, as in our study, the meta-analysis of Shi et al. showed significantly more advanced disease in the PN group (47.96% of Stage III in the PN group vs. 38.32% in the SL group; p<0.001), which could explain by itself the worse prognosis in patients who underwent pneumonectomy [5]. Therefore, PS methods tend to balance the distributions of potentially confounding covariates, but the lack of precision of the covariates measured might limit the impact of PS in this setting. As compared with the SL group, the PN group had significantly more recurrences according to matching. These findings were different from those of Park et al. whose sample size was too small to draw any definitive conclusions [27], or by Shi et al., whose matching analysis allowed a more powerful comparison of OS and DFS following SL and PN [5].

However, the two PS analyses did not provide the same results, and the lower OS and DFS in the PN group were not significant in the IPTW analysis. It is important to bear in mind that, unlike randomized assignments of treatments, PS typically does not balance covariates that were not observed [32]. Matching reduces the sample size since matches may not be found for some patients, whereas weighting allows the comparison of expectations and distributions between treated and untreated subjects [16,33]. The combination of matching and IPTW tends to eliminate systematic differences between experimental and control subjects to a greater degree than does stratification or covariate adjustment [15,16]. Even if these two techniques well balanced the distribution of the covariates, they do not permit to conclude which one provide the "true results" and also raised the question of the reality of the results published in studies using only matching analysis. Given the differences in the results of matching and IPTW, the results for long-term survival and recurrence should be interpreted with caution.

#### Strengths and limitations

To the best of our knowledge, this is the first study to compare SL and PN using a large dataset, a national database, two PS methods and a sensitivity analysis. The main strengths of this study are the use of a national database, which provided a large number of patients, a homogeneous population. The large number of patients in both groups allowed powerful comparisons. The homogeneous population reduced the sample size needed for the matching comparison and the reasonable length of study period tended to decrease historical bias.

However, any study involving a large database raises the question of the quality and exhaustiveness of the prospectively entered data, such as comorbidities, and observational studies are notoriously full of no responses and missing values [28]. Few details were available about the surgical technique, especially concerning bronchial stump coverage and pulmonary artery reconstruction. We used PS to create comparable cohorts; however, we cannot be certain that the PS perfectly neutralized all of the confounding variables, as suggested by the differences in the results for the two methods used.

The main limitation of this study is the dramatic loss of follow-up data (50% of missing data after only 1 month of follow-up), which could be explained by various factors: at the visit 1 month after the

surgery, some patients are not seen by the thoracic surgeon but by the medical oncologist or the 367 pneumologist; and some of the surgeons do not always enter follow-up data into the Epithor database, 368 369 for example, when they hear news about the patients or correspondence announcing their death. Moreover, we must underline that PS cannot replace randomized control trials (RCTs). In RCTs, 370 random allocation of patients to either an experimental or a control arm guarantees that treatment 371 allocation is unrelated to measured and unmeasured patients' characteristics. It enables researchers to 372 373 draw unbiased conclusions about a treatment effect, provided that the number of randomized patients is large enough to minimize random variation [33]. Even though this study has a high level of 374 evidence, the conclusions that can be drawn will never be as strong as those from prospective 375 multicenter RCT, which seem to be impossible to conduct nowadays. 376

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#### Conclusion

Despite the limits of our study, especially the dramatic loss of follow-up data, we highlighted the interest of SL, which could lead to better 3-year OS and DFS as compared to PN for NSCLC patients.

Whenever it is technically possible, surgeons must perform SL in order to provide more long-term

survival benefits to patients even with the risk of more postoperative pulmonary complications.

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#### Acknowledgments

This work was performed on behalf of the French Society of Thoracic and Cardiovascular surgery

(Société Française de Chirurgie Thoracique et CardioVasculaire - SFCTCV). We are grateful to all

the French thoracic surgeons who participated in this study and have listed them in e-appendix 1.

Pierre-Benoit Pagès is the recipient of research grants allocated by the Nuovo-Soldati foundation for

Cancer Research (Fondation Nuovo-Soldati) and the French Society of Thoracic and Cardiovascular

Surgery (SFCTCV).

The authors would like to thank Philip Bastable (Dijon University Hospital) for reviewing the

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Table 1
 Characteristics of patients undergoing sleeve lobectomy and pneumonectomy.

	Full Sa	Full Sample			
	Sleeve Lobectomy	Pneumonectomy	P Value		
	Group	Group	1 , 411010		
	(n=941)	(n=5318)			
<u>Demographics</u>					
Sex Male	716 (76.1%)	4216 (79.3%)	0.027		
Female	225 (23.9%)	1102 (20.7%)			
Age (Years)	$60.9 \pm 12.6$	$61.9 \pm 10.2$	0.014		
Body Mass Index (Kg/m²)	$25.6 \pm 4.5$	$25.1 \pm 4.1$	0.0009		
ASA 1	181 (19.2%)	817 (15.3%)	0.006		
2	522 (54.5%)	2951 (55.5%)			
3	224 (23.8%)	1434 (27%)			
4	14 (1.5%)	116 (2.2%)			
4	14 (1.5%)	110 (2.270)			
WHO Performance Status 0	442 (47%)	2124 (40%)	< 0.0001		
1	420 (44.6%)	2574 (48.4%)			
2	72 (7.6%)	523 (9.8%)			
3	7 (0.8%)	97 (1.8%)			
3	7 (0.8%)	97 (1.8%)			
<b>FEV</b> (%)	$74.1 \pm 17.6$	$62.9 \pm 20.5$	< 0.00001		
Dyspnea score 0	434 (46.1%)	2596 (48.8%)	0.37		
1	328 (34.9%)	1831 (34.4%)	0.57		
		, , ,			
2	148 (15.8%)	715 (13.5%)			
3	21 (2.2%)	121 (2.3%)			
4	10 (1%)	55 (1%)			
Number of comorbidities per patient 0	131 (13.9%)	702 (13.2%)	0.003		
1	226 (24%)	1471 (27.6%)	0.003		
	271 (28.8%)	1543 (29%)			
$\frac{2}{3}$	252(26.8%)	1393 (26.2%)			
3	· · · · · · · · · · · · · · · · · · ·				
47	61 (6.5%)	209 (4%)			
Perioperative management					
Neoadjuvant Chemotherapy	167 (17.7%)	1282 (24.2%)	< 0.0001		
Neoadjuvant Radiotherapy	11 (1.2%)	126 (2.4%)	0.02		
Adjuvant Chemotherapy for N1	202 (78%)	2378 (64%)	0.466		
N2	135 (90%)	1138 (83%)	0.4		
<b>Tumor characteristics</b>					
Side Right	652 (69.6%)	2172 (41%)	< 0.0001		
Left	285 (30.4%)	3122 (59%)	10.0001		
	(= (= (= (= (= (= (= (= (= (= (= (= (= (	(52.73)			
Tumor T1	190 (20.2%)	347 (6.5%)	< 0.0001		
T2	408 (43.3%)	1813 (34.1%)			
T3	169 (18%)	1482 (27.9%)			
T4	32 (3.4%)	690 (13%)			
Missing	142 (15.1%)	986 (18.5%)			
wiissing	142 (13.170)	700 (10.370)			
Lymph nodes N0	385 (40.9%)	1394 (26.2%)	< 0.0001		
N1	259 (27.5%)	1522 (28.6%)			
1/1	239 (21.3%)	1322 (28.0%)			

N2		150 (16%)	1372 (25.8%)	
N3		1 (0.1%)	22 (0.4%)	
Missing		146 (15.5%)	1008 (19%)	
Postoperative Histology	Squamous	514 (54.6%)	2570 (48.3%)	< 0.0001
	Adenocarcinoma	160 (17%)	1602 (30.1%)	
	Carcinoid	138 (14.7%)	69 (1.3%)	
	Others	57 (6%)	477 (9%)	
	Missing	72 (7.7%)	600 (11.3%)	
Resection Margins R0		797 (84.7%)	4388 (82.5%)	< 0.0001
R1		66 (6.4%)	232 (4.4%)	
R2		3 (0.3%)	47 (0.9%)	
Missing		81 (8.6%)	651 (12.2%)	
Characteristic	cs of Center			
<b>Type of center</b> Non-Ac	ademic	53 (5.6%)	669 (12.6%)	< 0.0001
Private		265 (28.2%)	1556 (29.3%)	
Acaden	nic	623 (66.2%)	3087 (58.1%)	
Hospital volume of activity	ty (number of		)	
procedures/year) <48		136 (14.5%)	1235 (23.2%)	< 0.0001
49-84		193 (20.5%)	1023 (19.2%)	
85-13		163 (17.3%)	903 (17%)	
134-1		213 (22.6%)	1214 (22.8%)	
>17	1	236 (25.1%)	943 (17.8%)	
Year of treatment 2005-	2006	175 (18.6%)	1003 (18.9%)	0.005
2007-	2008	164 (17.4%)	1189 (22.4%)	
2009-	2010	207 (22%)	1174 (22%)	
2011-	2012	189 (20%)	938 (17.6%)	
2013-	2014	206 (22%)	1014 (19.1%)	
		4		

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ASA: American Society of Anesthesiologists; FEV: Forced Expiratory Volume; WHO: World Health

488 Organization

PB. Pagès et al. Sleeve lobectomy vs Pneumonectomy

Patients' baseline characteristics for sleeve lobectomy and pneumonectomy (in parentheses percentage or mean) with their standardized difference. A standardized difference greater than 0.1 (10%) represents meaningful imbalance in a given variable between treatment groups.

Table 2

		Full sampl	le	Matched			Weighted	l	
Variables	Sleeve (n=941)	PN (n=5318)	standardized difference	Sleeve (n=794)	PN (n=794)	standardized difference	Sleeve (n=941)	PN (n=5318)	standardized difference
Demographics					)				
Sex Male	715 (76%)	4201 (79%)	0.078	635 (80%)	627 (79%)	-0.015	734 (78%)	4201 (79%)	0.017
BMI	25	25.6	-0.107	25.5	25.5	0.001	25.3	25.2	-0.028
WHO status 1	423 (45%)	2552 (48%)	0.075	381 (48%)	381 (48%)	0.013	489 (52%)	2552 (48%)	-0.064
2	75 (8%)	532 (10%)	0.081	63 (8%)	63 (8%)	0.00001	84 (9%)	532 (10%)	0.006
3	9 (1%)	106 (2%)	0.094	8 (1%)	8 (1%)	-0.0022	28 (3%)	106 (2%)	-0.082
FEV	74%	63%	-0.583	73%	73%	-0.011	66.5%	64.7%	-0.098
Dyspnea score 1	329 (35%)	1808 (34%)	-0.009	119 (15%)	301 (38%)	0.029	320 (34%)	1861 (35%)	0.009
2	150 (16%)	691 (13%)	-0.067	119 (15%)	119 (15%)	-0.021	132 (14%)	744 (14%)	-0.001
3	18 (2%)	106 (2%)	0.003	24 (3%)	16 (2%)	-0.034	37 (4%)	53 (1%)	-0.124
4	9 (1%)	53 (1%)	-0.003	8 (1%)	8 (1%)	-0.012	19 (2%)	53 (1%)	-0.083
Number of comorbidities per									
patients 1	226 (24%)	1489 (28%)	0.083	190 (24%)	182 (23%)	-0.02	263 (28%)	1435 (27%)	-0.017
2	273 (29%)	1542 (29%)	0.003	238 (30%)	246 (31%)	0.011	244 (26%)	1542 (29%)	0.065
3	254 (27%)	1382 (26%)	-0.01	222 (28%)	222 (28%)	0.014	291 (31%)	1382 (26%)	-0.112
4	66 (7%)	212 (4%)	-0.115	47 (6%)	47 (6%)	-0.04	47 (5%)	212 (4%)	-0.01
Perioperative management		7							
Neoadjuvant chemotherapy	169 (18%)	1276 (24%)	0.156	159 (20%)	167 (21%)	0.009	216 (23%)	1276 (24%)	0.013

PB. Pagès et al. Sleeve lobectomy vs Pneumonectomy

Tumor characteristics									
Right side	659 (70%)	2180 (41%)	-0.599	532 (67%)	516 (65%)	-0.042	423 (45%)	2393 (45%)	0.008
Tumor T2	404 (43%)	1808 (34%)	-0.19	349 (44%)	333 (42%)	-0.047	3667 (39%)	1914 (36%)	-0.062
T3 T4	169 (18%) 28 (3%)	1489 (28%) 691 (13%)	0.241 0.354	166 (21%) 32 (4%)	174 (22%) 47 (6%)	0.036 0.075	216 (23%) 84 (9%)	1382(26%) 638 (12%)	0.088 0.081
Missing	28 (3%) 141 (15%)	957 (18%)	0.088	32 (4%) 135 (17%)	127 (16%)	-0.017	188 (20%)	957 (18%)	-0.049
Wiissing	141 (1370)	937 (1070)	0.000	133 (1770)	127 (10%)	-0.017	166 (2070)	937 (1670)	-0.049
Lymph Nodes N1	263 (28%)	1542 (29%)	0.025	238 (30%)	230 (29%)	-0.036	254 (27%)	1542 (29%)	0.037
N2	150 (16%)	1382 (26%)	0.249	143 (18%)	150 (19%)	0.028	225 (24%)	1329 (25%)	0.026
N3	0 (0%)	0 (0%)	0.061	0 (0%)	8 (1%)	0.074	0 (0%)	0 (0%)	0.03
Missing	150 (16%)	1010 (19%)	0.087	135 (17%)	135 (17%)	-0.02	197 (21%)	1010 (19%)	-0.068
Postoperative histology	1.60 (1.70)	1505 (200)	0.217	150 (200)	150 (200)	0.002	244 (260)	1.400 (200()	0.054
Adenocarcinoma	160 (17%)	1595 (30%)	0.317	159 (20%)	159 (20%)	0.003	244 (26%)	1489 (28%)	0.054
Carcinoid	141 (15%)	53 (1%)	-0.51	40 (5%)	47 (6%)	0.019	28 (3%)	159 (3%)	-0.02
Others	56 (6%) 75 (8%)	478 (9%) 584 (11%)	0.111 0.121	55 (7%) 71 (9%)	63 (8%) 63 (8%)	0.038 -0.026	94 (10%) 94 (10%)	478 (9%)	-0.043 0.01
Missing	73 (8%)	364 (11%)	0.121	71 (9%)	03 (8%)	-0.026	94 (10%)	585 (11%)	0.01
<b>Characteristics of Center</b>									
Character issues of Contor									
<b>Type of center</b> Private	263 (28%)	1542 (29%)	0.02	230 (29%)	214 (27%)	-0.031	244 (26%)	1542 (29%)	-0.022
Academic	621 (66%)	3084 (58%)	-0.164	516 (65%)	508 (64%)	-0.013	564 (60%)	3137 (59%)	-0.022
			Y						
Hospital volume of activity			Y						
(number of procedures/year)			$\sim$						
49-84	197 (21%)	1010 (19%)	-0.032	159 (20%)	150 (19%)	-0.025	179 (19%)	1010 (19%)	-0.005
85-133	160 (17%)	904 (17%)	-0.008	135 (17%)	143 (18%)	0.027	179 (19%)	904 (17%)	-0.058
134-171	207 (22%)	1223 (23%)	0.012	190 (24%)	190 (24%)	-0.00001	216 (23%)	1223 (23%)	0.007
>171	235 (25%)	957 (18%)	-0.18	182 (23%)	174 (22%)	-0.028	169 (18%)	1010 (19%)	0.016
V 2007 2000	1.60 (100/)	11(0)(220()	0.122	150 (100/)	125 (170/)	0.025	225 (240()	1160 (220)	0.05
Year 2007-2008 2009-2010	169 (18%)	1169 (22%)	0.122 0.005	150 (19%)	135 (17%)	-0.035 0.04	225 (24%)	1169 (22%)	-0.05
2009-2010 2011-2012	207 (22%) 188 (20%)	1169 (22%) 957 (18%)	0.005 -0.062	174 (22%) 159 (20%)	182 (23%) 174 (22%)	0.04	225 (24%) 150 (16%)	1169 (22%) 957 (18%)	0.046 0.05
2011-2012 2013-2014	207 (22%)	957 (18%) 1010 (19%)	-0.062 -0.071	159 (20%)	174 (22%) 150 (19%)	-0.031	188 (20%)	1010 (19%)	-0.01
Overall propensity score	201 (2270)	1010 (17/0)	0.071	137 (2070)	130 (17/0)	0.031	100 (2070)	1010 (17/0)	0.01
Mean			0.134			0.0258			0.040
Median			0.085			0.0255			0.029

## Table 3

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Effects of pneumonectomy vs sleeve lobectomy according to different statistical analyses (OR for SL=1, if OR <1 means a protective effect of pneumonectomy, OR>1 means a deleterious effect of

496 pneumonectomy).

	Matching 1	:1	Weighting		
		P value		P value	
Postoperative mortality*	1.24 (0.74 – 2.1)	0.4	0.77 (0.4-1.5)	0.4	
			<b>Y</b>		
Postoperative complications*					
Postoperative pulmonary complications	0.4 (0.3-0.6)	< 0.0001	0.12 (0.08-0.2)	< 0.0001	
Atelectasis	0.14 (0.08-0.25)	< 0.0001	0.56 (0.36-0.87)	< 0.0001	
Pneumonia	0.4 (0.24-0.69)	< 0.001	0.43 (0.31-0.61)	0.01	
Ventilation > 2 days	1.4 (0.8-2.5)	0.2	1.25 (0.7-2.3)	0.4	
ARDS	1.4 (0.7-2.5)	0.3	1.22 (0.6-2.5)	0.5	
Arrhythmia	1.6 (0.95-2.6)	0.08	1.67 (0.9-2.9)	0.06	
Bronchopleural fistula	2.9 (1.4-6.2)	0.005	2.5 (1.3-4.7)	0.004	
Empyema	7 (1.9-27)	0.004	15 (4-57)	< 0.0001	
Hemorrhage	4 (1.9-8)	< 0.0001	2 (0.7-6)	0.2	
Length of hospital stay**	-1.31 (-2.540.09)	0.03	-1.18 (-3-0.63)	0.2	
Overall survival***	1.63 (1.19-2.21)	0.002	0.97 (0.63-1.51)	0.92	
Disease-Free survival***	1.49 (1.1-2.01)	0.01	1.03 (0.73-1.45)	0.84	

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\*Odds Ratio (OR) with 95% confidence interval

\*\*Difference of the mean number of days

\*\*\*Hazard ratio with 95% confidence interval

ARDS: Acute Respiratory Distress Syndrome

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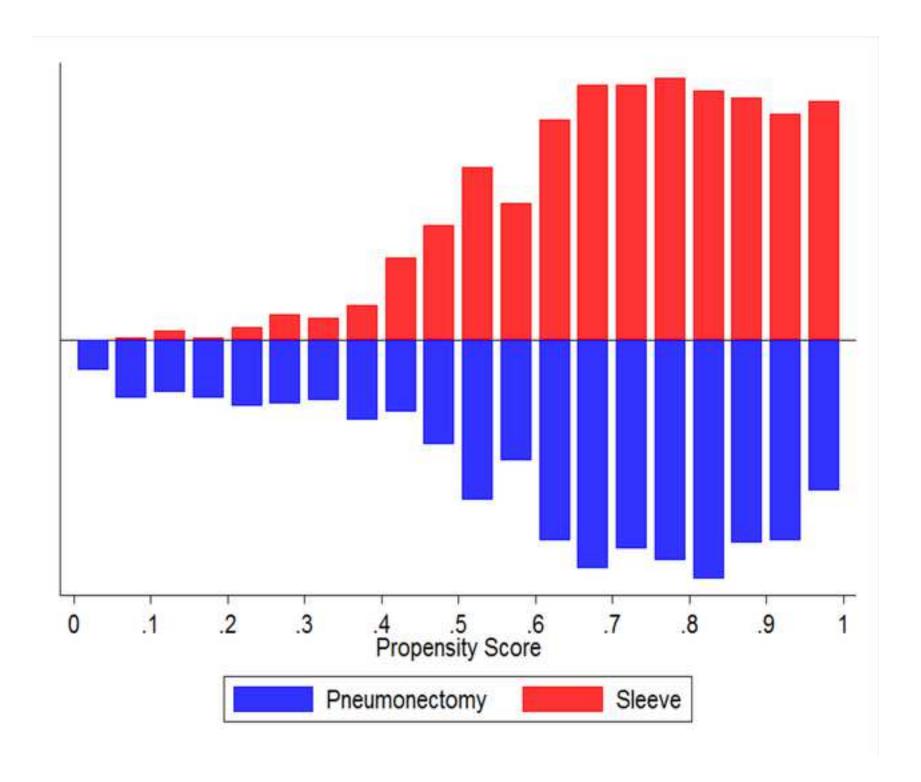
## Figure legends

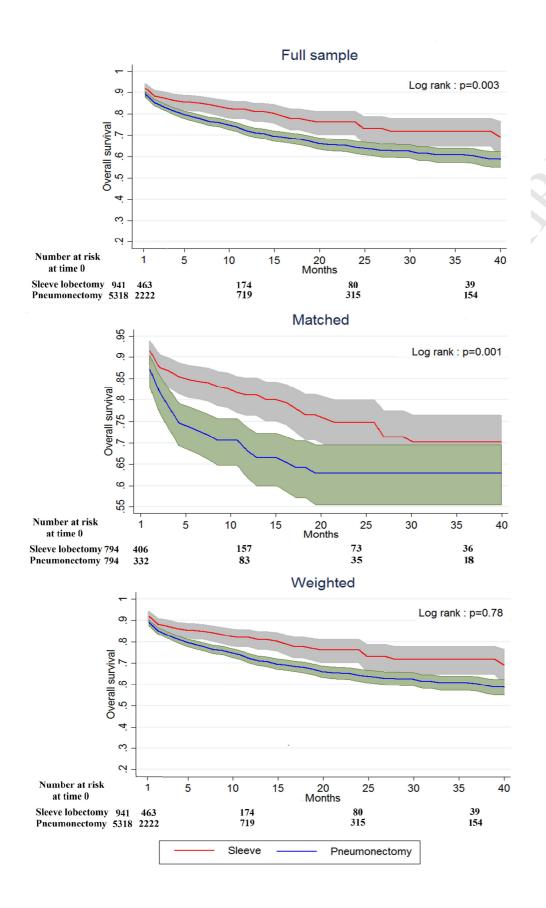
**Figure 1:** Kaplan-Meier plots of overall survival from the unmatched full sample (A), matched (B) and weighted (C) data for PN (continuous red line) and SL (discontinuous blue line).

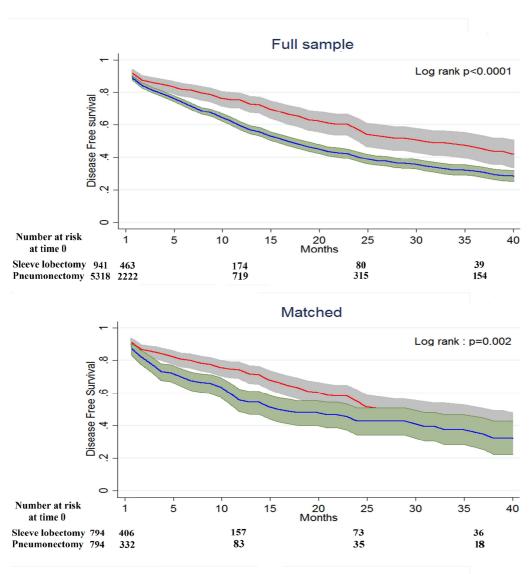
**Figure 2:** Kaplan-Meier plots of disease-free survival from the unmatched full sample (A), matched (B) and weighted (C) data for PN (continuous red line) and SL (discontinuous blue line).

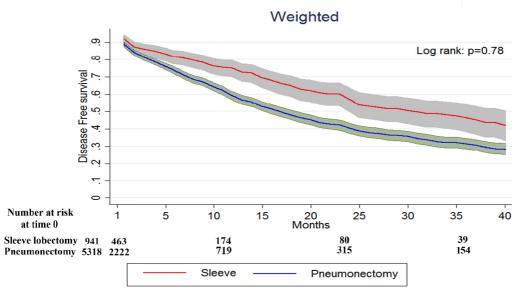
**Figure 3:** Graph of propensity scores in the 2 groups of patients. Each bar represents the number of patients with the same propensity score in both groups.

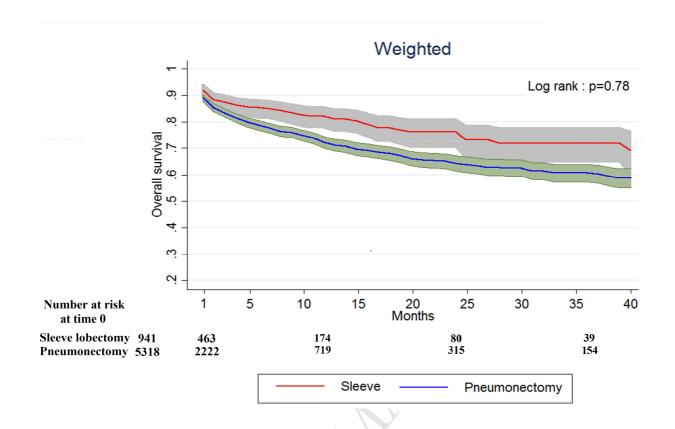
**Video legend:** this is a video of a double (vascular and bronchial) left upper lobectomy by thoracotomy.

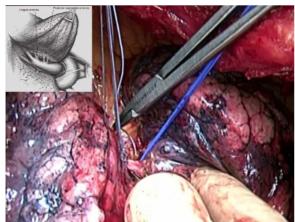












#### Appendix 1

The authors would like to thank all the French thoracic surgeons who participated in Epithor and subsequently in this study in order to improve thoracic surgery quality: Dr Michel Alauzen (Montpellier), Dr Jean-François Andro (Quimper), Dr Maxime Aubert (Grenoble), Dr Jean Philippe Avaro (Marseille), Dr Patrick Bagan (Argenteuil), Dr Francois Bellenot (Cergy Pontoise), Vincent Blin (Vannes), Dr Philippe Boitet (Harfleur), Dr Laurent Bordigoni (Toulon), Professor Jacques Borrelly (Nancy), Professor Pierre-Yves Brichon (Grenoble), Dr Gilles Cardot (Boulogne sur Mer), Dr Jean Michel Carrie (Saint Jean), Dr François Clement (Besançon), Professor Pierre Corbi (Poitiers), Dr Michel Debaert (Lille), Dr Bertrand Debrueres (Ploemeur), Dr Jean Dubrez (Bayonne), Dr Xavier Ducrocq (Strasbourg), Dr Antoine Dujon (Bois Guillaume), Professor Pascal Dumont (Tours), Dr Philippe Fernoux (Chalon sur Saône), Professor Marc Filaire (Clermont-Ferrand), Dr Eric Frassinetti (Chambéry), Dr Gil Frey (Saint Etienne), Dr Dominique Gossot (Paris), Professor Gilles Grosdidier (Nancy), Dr Benoit Guibert (Lyon), Dr Olivier Hagry (Chalon sur Saône), Dr Sophie Jaillard (Lille), Dr Jean-Marc Jarry (Aix en Provence), Dr David Kaczmarek (Saint Etienne), Dr Yves Laborde (Pau), Dr Bernard Lenot (Saint Brieuc), Dr Francis Levy (Bordeaux), Dr Laurent Lombart (Béziers), Dr Eric Marcade (Saint Grégoire), Dr Jean Paul Marcade (La Rochelle), Dr Jean Marzelle (Créteil), Professor Gilbert Massard (Strasbourg), Dr Florence Mazeres (Bayonne), Dr Eric Mensier (Lille), Dr David Metois (Orléans), Dr J. L. Michaud/E Paris (Nantes), Dr Philippe Mondine (Brest), Dr Michel Monteau (Reims), Dr Jean-Michel Moreau (Nantes), Professor Jérôme Mouroux (Nice), Dr Antoine Mugniot (Nantes), Dr Pierre Mulsant (Lyon), Dr Nidal Naffaa (Avignon), Dr Gérard Pavy (Arras), Professor Christophe Peillon (Rouen), Professor Francois Pons (Percy), Professor Henri Porte (Lille), Professor Jean-François Regnard (Paris), Professor Marc Riquet (Paris), Dr Babak Sadeghi Looyeh (Morlaix), Professor Olivier Tiffet (Saint Etienne), Dr Bruno Tremblay (Meaux), Dr Jean Valla (Charenton le Pont), Professor Jean-François Velly (Pessac), Dr Bernard Wack (Metz), Dr Jean-Didier Wagner (Colmar), and Dr Didier Woelffe (Valenciennes).

## Supplementary table 4

Sensitivity analysis to Hidden Bias for matching one-to-one Sensitivity (Mantel-Haenszel test statistic).

	Γ=1 (assumption no hidden bias)		Conclusion
	Qmh	P value	
Postoperative mortality	0.93	0.17	Caution, possible hidden bias
Postoperative complications			
Atelectasis	8.4	0.0001	No hidden bias
Pneumonia	4.4	0.0001	No hidden bias
Postoperative pulmonary complications	6.3	0.0001	No hidden bias
Arrhythmias	2.5	0.006	No hidden bias
Bronchopleural fistula	3.3	0.004	No hidden bias
Empyema	2.7	0.003	No hidden bias
Hemorrhage	4	0.0001	No hidden bias
Ventilation > 2 days	1.02	0.15	Caution, possible hidden bias
ARDS	0.9	0.16	Caution, possible hidden bias

ARDS: Acute Respiratory Distress Syndrome

## **Supplementary Table 5**

Assessment of sensitivity to positivity violations for weighting method (excluding observations in regions of non-overlap).

	Weighting			
	(n=4829)			
		P value		
Postoperative mortality	0.74 (0.4-1.44)	0.4		
Postoperative complications				
Atelectasis	0.13 (0.07-0.22)	0.0001		
Pneumonia	0.55 (0.35-0.88)	0.01		
Postoperative pulmonary complications	0.47 (0.33-0.67)	0.0001		
Arrhythmias	1.86 (1.06-3.23)	0.03		
Bronchopleural fistulae	2.4 (1.23-4.8)	0.01		
Empyema	13 (3.5-48)	0.0001		
Hemorrhage	1.8 (0.57-6)	0.3		
Ventilation > 2 days	1.12 (0.6-2.1)	0.7		
ARDS	1.05 (0.5-2.2)	0.9		
Overall survival	0.99 (0.89-1.11)	0.9		
Disease-Free survival	1.1 (1.01-1.2)	0.03		

ARDS: Acute Respiratory Distress Syndrome