# Pulmonary Function Improvement after Lung Volume Reduction Surgery: A Twenty-year Experience

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#### Disclosures

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Reddy, RM-Intuitive Surgical, Auris Health, Medtronic, Genentech, On Target Labs

Lin, J-Intuitive Surgical

Non relevant to this talk



### National Emphysema Treatment Trial (NETT) Findings

- NETT distinguished favorable versus unfavorable characteristics of candidates for LVRS
  - Recommended for upper-lobepredominant (Fishman, et al. 2003)
- LVRS, supplemental oxygen and smoking cessation improve survival in selected patients (Criner, et al. 2011)
- LVRS remains an underused therapy



## Objective

To evaluate post-operative outcomes of lung volume reduction surgery (LVRS) at a high-volume institution that participated in the National Emphysema Treatment Trial (NETT) and continues to follow NETT criteria.



#### Criteria for LVRS

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#### Inclusion

History and physical examination consistent with emphysema CT Scan evidence of bilateral upper-lobe-dominant emphysema Pre-rehabilitation post-bronchodilator TLC≥100% and RV≥150%

FEV1 ≤45%

PaCO<sub>2</sub>≤60mmHg

PaO<sub>2</sub>≥45mmHg

Cotinine ≤ 13.7ng/ml

BMI  $\leq$ 31.1 (males) or  $\leq$ 32.3 (females)

Quit smoking at least 4 months before surgery Completion of pre-rehabilitation assessments

Completion of NETT rehabilitation program

#### Exclusion

Daily use of prednisone >20mg

No other major disease

Previous heart or lung surgery

## Criteria for study based on NETT



#### Methods

#### Patients who underwent LVRS identified between July 1998 and April 2019

Chart review- demographics, pre-operative diagnostic tests (PFTs, 600m walk, oxygen supplementation), procedural information, and post-operative outcomes recorded

Analyzed whole cohort data

Divided cohort based on procedure type (MS vs. VATS)

Analyzed data based on procedure type



## Cohort Demographics

Average age was 62±7.6 years

Total of 151 patients

Average smoking history of 54±24.3 pack year

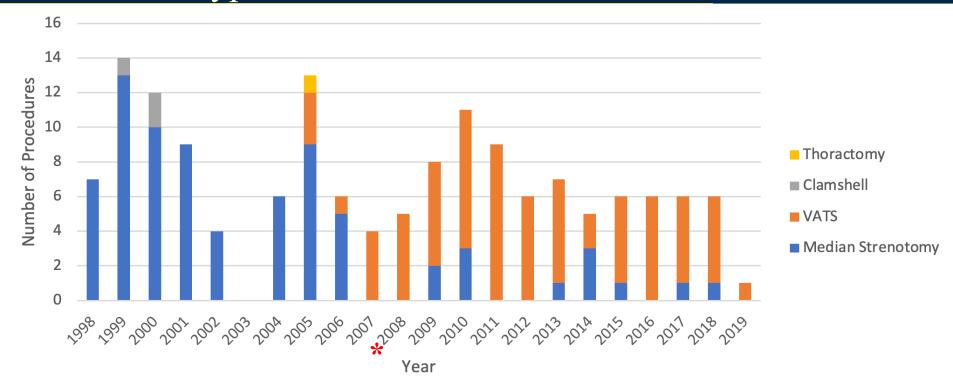
#### Cohort Pre-operative Pulmonary Function Tests

Pre-Op Pulmonary Function Tests	Cohort
FVC (L) (n=150)	2.4±0.76
FEV1 (L) (n=150)	0.72±0.2
FEV1 % of Predicted Value (n=150)	27.4±6.9
FEV1:FVC % (n=150)	30.9±5.98
Carbon Monoxide Diffusing Capacity (mL/min/mmHg) (n=122)	8.97±4.2
Carbon Monoxide Diffusing Capacity % of Predicted Value (n=122)	35.6±13.4
Total Lung Capacity (L) (n=140)	8.5+10.7

All subjects met NETT criteria for pre-operative PFTs



#### Procedure Type



#### MS vs. VATS

	Median Sternotomy (n=75)	VATS (n=72)
Characteristics		
Age at Surgery (yr)*	63±7	60.6±7.9
Sex- no. (%)		
Female	31 (41)	38 (53)
Male	44 (59)	34(47)
Body Mass Index	25.4±4.14	25.9±4.03
Smoking History (Pack-years) *	59.5±26.4	49.2±21.6
No. of years quit before surgery	7.26±6.63	6.1±6.54
Pre-Op Supplemental Oxygen (L)	1.8±1.5	2.4±1.5
Pre-Op Exercise Capacity (watt)	37.5±16.8	35.7±19.6
Distance walked in 6 min (ft).	886±468.5	813.1±437.9
PaO2- mmHg*	65.8±13.2	78.9±20.4
PaCO2- mmHg	41.7±4.04	42.9±5.38
Pre-Op Pulmonary Function Tests		
FVC (L)	2.47±0.76	2.31±0.75
FEV1 (L)	0.72±0.2	0.72±0.2
FEV1 % of Predicted Value	27.7±7.45	27.9±6.43
Carbon Monoxide Diffusing Capacity (mL/min/mmHg)	9.1±3.4	8.9±4.7
Carbon Monoxide Diffusing Capacity % of Predicted	25.2.2	25.0.45.5
Value	35.3±9.7	35.8±15.6
Total Lung Capacity (L)	7.99±4.2	9.2±15.4

VATS group was younger (p=0.045) and presented with fewer smoking pack-years (p=0.013)



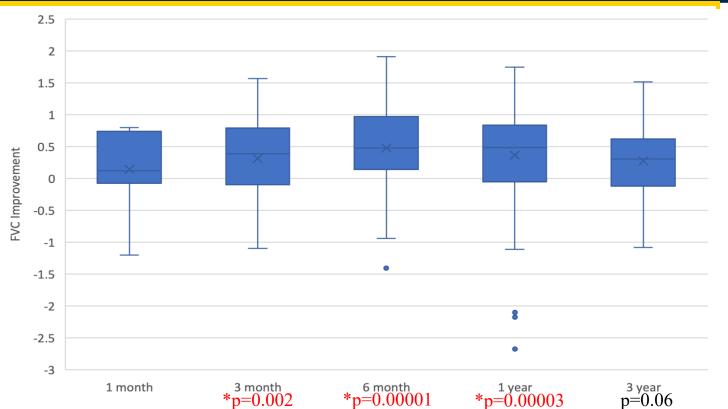
## Perioperative Outcomes

	Median Sternotomy (n=75)	VATS (n=72)
Complications	No. (%)	No. (%)
New CT placed	6 (8)	7 (10)
Respiratory distress	13 (17)	22 (31)
Respiratory failure	7 (9)	10 (14)
Pneumonia	20 (27)	14 (19)
Reintubation	10 (13)	10 (13)
Tracheostomy	2 (3)	1 (1)
Afib_SVT	16 (21)	25 (35)
PE	0 (0)	1 (1)
MI	0 (0)	2 (3)
AKI	8 (11)	3 (4)
Wound Infection	3 (4)	1 (1)
Re-op	8 (11)	5 (7)
Re-admit to ICU	8 (11)	10 (14)
Hospital Mortality	9 (12)	5 (7)
SQ emphysema* (p=0.03)	6 (8)	15 (21)
Initial Air leak* (p=0.03)	45 (60)	55 (76)
Prolong air leak	37 (49)	45 (63)
	Mean ± STD	Mean ± STD
Hospital LOS	11.37±9.56	11.97±10.68
ICU LOS	3.02±6.54	4.52±8.24
Chest tube days	15.63±18.3	15.52±17.29
Ventilator time	0.027±0.23	1.04±4.75
Reintubation Days	2.44±3.84	6.58±8.5
EBL (cc)* (p=0.03)	133±145.2	88.4±130.3
Surgery Duration(min)	192.28±44.7	232.3±155.6

- Decrease in EBL over time\*
- VATS associated with increased risk of initial air leaks and subcutaneous emphysema
- No other significant differences between MS and VATS



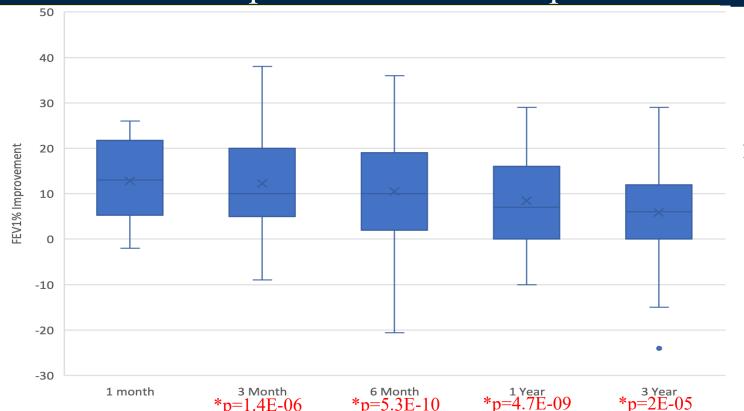
## Cohort Post-operative FVC Improvement



PFTs taken at 3 months, 6 months and 1 year showed significant improvement in FVC compared to pre-operation



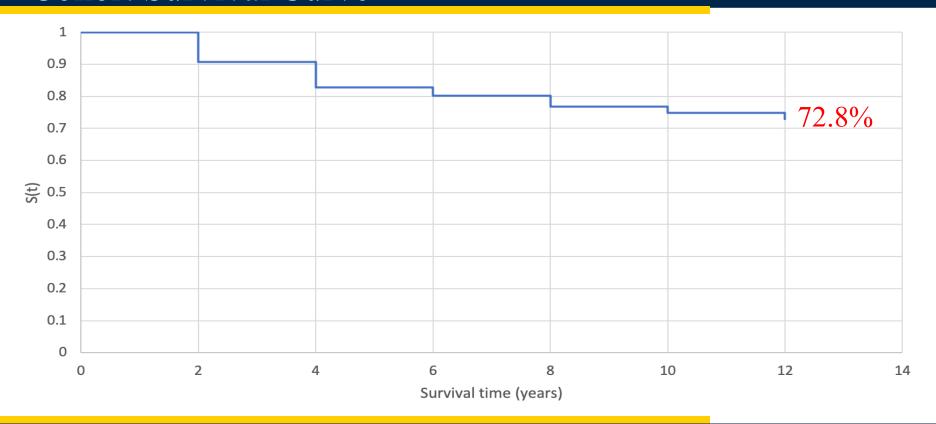
#### Cohort Post-operative FEV1% Improvement



Significant improvement in FEV1% seen at 3 months, 6 months, 1 year, and 3 years

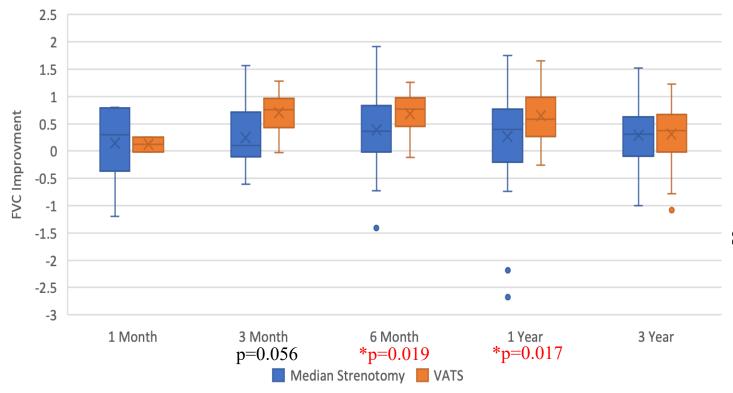


#### Cohort Survival Curve



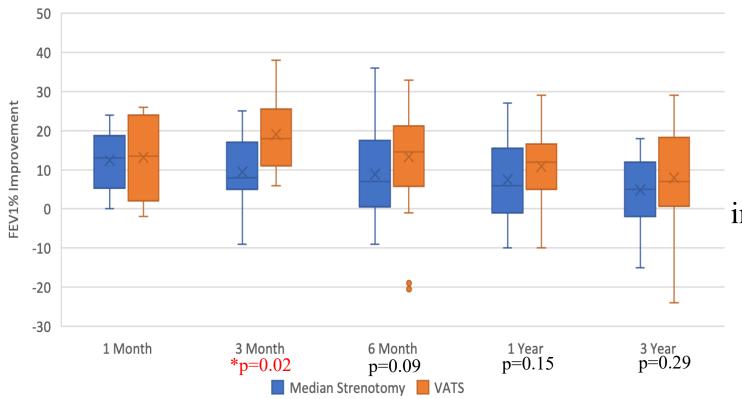


## Post-operative FVC Improvement by Procedure Type



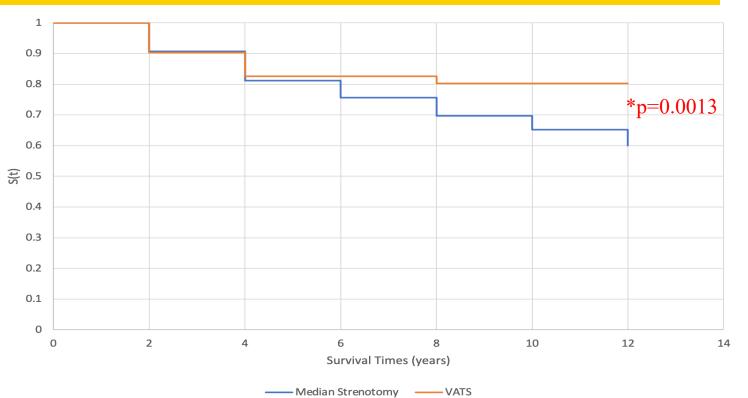
Significant
difference in
FVC
improvement
seen at 6 months,
1 year in VATS
group

### Post-operative FEV1% Improvement by Procedure



Significant difference in FEV1% improvement only seen at 3 months

### Survival Curve by Procedure Type



**VATS** showed improved survival at 12 years post-LVRS when compared to MS

#### Conclusions

VATS showed improvement in post-operative FVC up to 1 year post-operatively when compared to MS

Patients
undergoing LVRS
had significant
increases in postoperative FVC and
FEV1% lasting up
to 3 years postsurgery

VATS
demonstrated
improved survival
rate 12 years postsurgery\*



Thank you! <a href="mailto:scjulia@med.umich.edu">scjulia@med.umich.edu</a>



