ATRIAL MEASUREMENTS:

CAN THEY PREDICT ADVERSE EVENTS IN PATIENTS WITH ACUTE PE?

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Disclosures

None

Background: Existing data

- Reduced left atrial volume (<62 mls) and increased atrial volume ratio (>1.2) is
 associated with a significantly higher 30-day all-cause mortality⁽¹⁾
- Increased risk of adverse events (30-day PE-related mortality or the need for advanced therapy) when there is evidence of right heart strain on ECHO, ECG and CTA together⁽²⁾
- Septal bowing and increasing pulmonary artery diameter are associated with 5-day adverse outcomes (all-cause mortality, acute decompensation, or need for emergent treatment)⁽³⁾

^{1.} G Aviram et al (2016); ^{2.} B Carroll et al (2018); ^{3.} M Lyhne et al (2019)

Aims

- Determine which atrial measurement(s) (volume, area, diameter) is/are associated with
 30-day PE-related adverse events in patients with acute PE
- Determine if any atrial measurement(s) is/are a better predictor of 30-day PE-related
 adverse events compared to the combination of ECHO, ECG, and CTA
- Determine whether increasing pulmonary artery diameter, septal bowing, or reflux of contrast into the IVC are associated with 30-day PE-related adverse events

Methods

501 patients diagnosed with acute PE between 2007 – 2014

Clinical records:

Adverse events recorded: 30-day PE-related mortality or the need for advanced therapy i.e. thrombolysis, thrombectomy, vasopressors

2. Tests:

- Tricuspid annular plane systolic excursion (TAPSE) measurement from the echo report
 - performed 24hrs before or 48hrs after the diagnostic CT scan
 - classed as abnormal if <16mm
- ECG result recorded where available

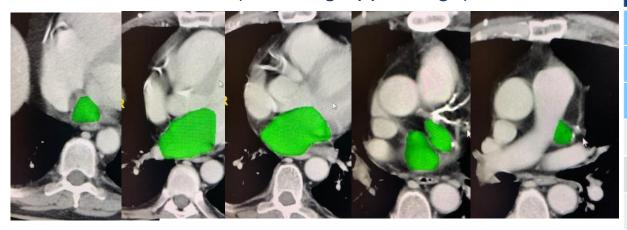
Methods

3. CT measurements (Gated and non-gated chest CTAs) using TeraRecon:

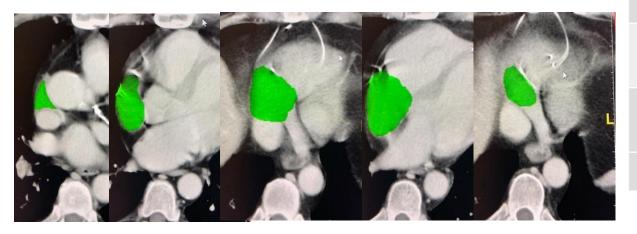
- Right and left atrial volume, area, and diameters
- Main PA diameter
- Ascending aorta diameter
- RV/LV diameter ratio
- Ventricular septal bowing
- Reflux of contrast into the IVC (into hepatic veins)

Methods: Atrial Volume

Left atrium (including appendage):



Right atrium (including appendage):



Measurements

Left atrial volume

Right atrial volume

RA/LA volume ratio

Left atrial area

Right atrial area

RA/LA area ratio

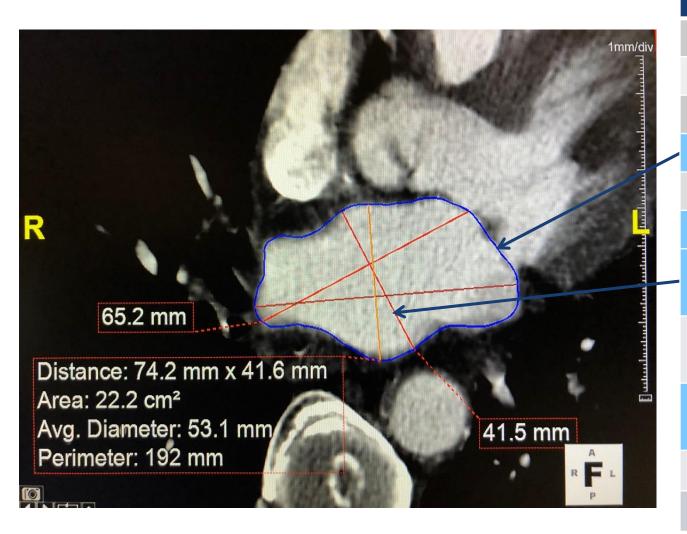
Left atrial short-axis diameter

Right atrial short-axis diameter

RA/LA short-axis diameter ratio

PA diameter

Methods: Left atrial area and diameter



Measurements

Left atrial volume

Right atrial volume

RA/LA volume ratio

Left atrial area

Right atrial area

RA/LA area ratio

Left atrial short-axis diameter

Right atrial short-axis diameter

RA/LA short-axis diameter ratio

PA diameter

Methods: Right atrial area and diameter



Measurements

Left atrial volume

Right atrial volume

RA/LA volume ratio

Left atrial area

Right atrial area

RA/LA area ratio

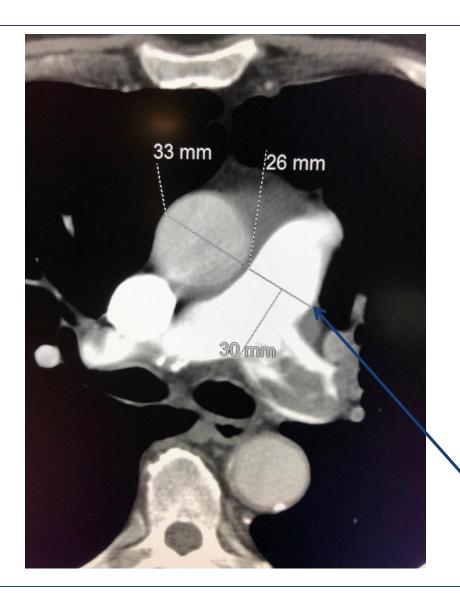
Left atrial short-axis diameter

Right atrial short-axis diameter

RA/LA short-axis diameter ratio

PA diameter

Methods: Pulmonary artery



Measurements

Left atrial volume

Right atrial volume

RA/LA volume ratio

Left atrial area

Right atrial area

RA/LA area ratio

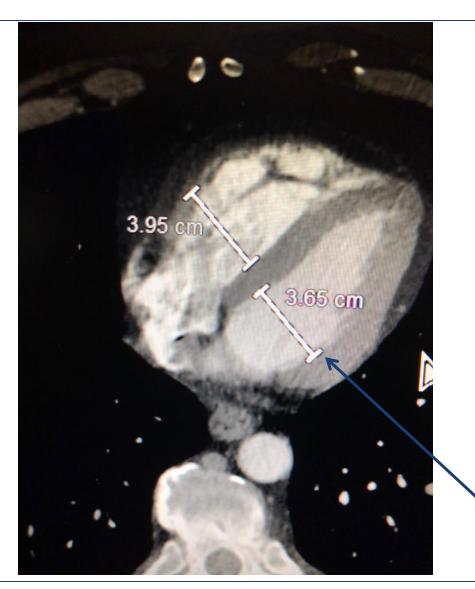
Left atrial short-axis diameter

Right atrial short-axis diameter

RA/LA short-axis diameter ratio

PA diameter

Methods: Ventricular diameters



Measurements

Left atrial volume

Right atrial volume

RA/LA volume ratio

Left atrial area

Right atrial area

RA/LA area ratio

Left atrial short-axis diameter

Right atrial short-axis diameter

RA/LA short-axis diameter ratio

PA diameter

Statistical Analysis

Comparison of subsets (with and without adverse events) was done using **Mann-**Whitney U and Chi-squared tests

- Prediction analysis was done by calculating the **area under the curve (AUC)** for receiver operating characteristic (ROC) curves, and Delong's test was used to compare AUC for paired ROC curves
- Multivariate logistic regression was performed to determine the best predictors of adverse outcomes

Results

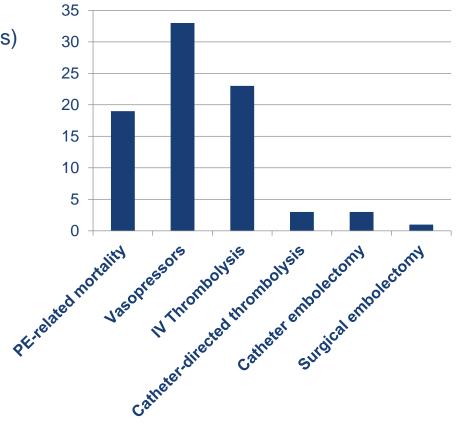
Demographics

• N = 493 pts

(8 excluded - incomplete/unavailable images)

- Adverse events = 62/493 pts (12.6%)
- Mean age = 63±16 years (p=0.2)
- Male 48%; Female 52% (p=0.6)

Number of patients with 30-day adverse events

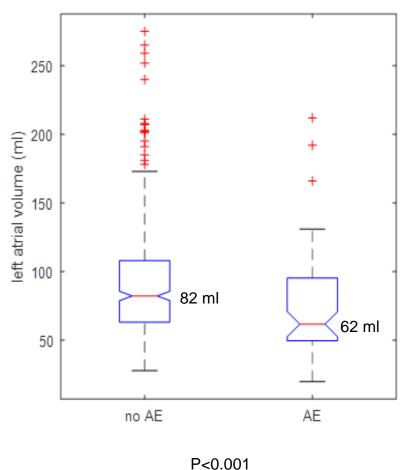


Results: Association between measurements and adverse events

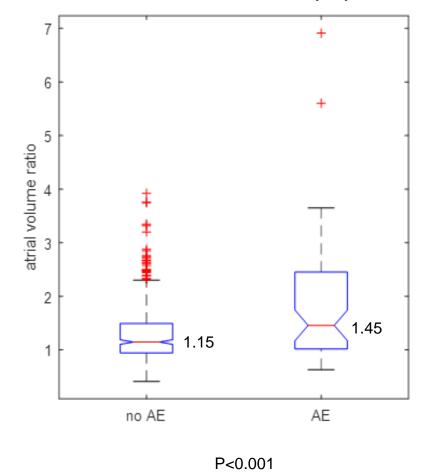
Measurements	No Adverse Event (Median)	Adverse Event (Median)	p-value
Left atrial volume	82 mls	62 mls	<0.001
RA/LA volume ratio	1.15	1.46	<0.001
Left atrial area	21 cm ²	18 cm ²	0.001
RA/LA area ratio	1.19	1.45	<0.001
Left atrial short-axis diameter	42 mm	37 mm	0.001
RA/LA short-axis diameter ratio	1.41	1.62	<0.001
RV/LV diameter ratio	1.0	1.21	<0.001

Results: Left atrial volume

LA volume in patients with and without adverse events (AE)



RA/LA volume ratio in patients with and without adverse events (AE)



Results: Association between measurements and adverse events

		Adverse Event Median (N = 62)	P-value
TAPSE	18 mm	14.5 mm	<0.001
PA diameter	30 mm	30 mm	0.88

Measurements	No Adverse Event (% of group)	Adverse Event (% of group)	P-value
ECG	40%	55%	0.02
Septal Bowing	3%	10%	0.01
Reflux of contrast into IVC	26%	50%	<0.001

Results: Prediction of adverse events

Measurements	AUC
Left atrial volume	0.67
TAPSE+ECG+(RV/LV)	0.71

No sig difference between them (P>0.07)

Limitations

Retrospective study

- **Selection bias:**
 - Acute PE only
 - Original database only included patients with an echo performed
- Slightly **limited precision** of measurements due to:
 - Motion artefact
 - Varying quality of contrast opacification in atria
 - Streak artefact from contrast/lines in the right atrium
- Mostly Non-ECG gated studies therefore unable to account for systole/diastole

Conclusions

 LA measurements are associated with 30-day PE-related adverse events in patients with acute PE

- LA volume is an equivalent predictor of 30-day PE-related adverse events compared to the combination of modalities (TAPSE, ECG, and RV/LV diameter ratio)
- Reflux of contrast into the IVC and septal bowing are associated with 30-day PE-related adverse events, however PA diameter is not

Ongoing and Future Work

Assess inter-rater variability with a randomly selected subset

- Assess the association of atrial measurements with 90-day and 1-year mortality
- Repeat the measurements using the same automatic software as Aviram et al (2016) for our cohort to see if our results are reproducible

References

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