Advanced MR Imaging for Lung Diseases

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Lung is most challenging solid organ to image

- 1. Large and moves with respiration (motion artifacts)
- 2. Low density ($\rho = 0.2 \text{ g/cm}^3 \text{ at TLC}$)
- 3. Multiple air-tissue interfaces (alveoli) cause fast MRI decay of signal



X-ray (not tomographic)

X-ray CT (fairly high ionizing radiation)

MRI (no radiation, but historically bad for parenchyma)





neonate

Challenges have caused innovation

1. UTE MRI sequences (echo time < 0.2 ms) Early CF Lung disease Neonates in NICU (with "self"-respiratory gating)

2. A scaled-down, neonatal MRI scanner 1.5T ONI / GE hybrid

3. Hyperpolarized-gas MRI (³He or ¹²⁹Xe) Realtime ventilation (breath hold for 10-15 s) Measure of alveolar-airspace size Measures of gas exchange







Density quantification via MRI is now possible (validation by CT)





NS Higano, et al., *J Magn Reson Imag* 2017 (in press) doi 10.1002/jmri.25643 JM Stein, et al., *Pediatric Radiology* 2016; 46: 1804

Can UTE MRI quantify abnormalities like CT?: 1-3 y.o.

- Score both MRI and CT via Brody Score
- Lung Abnormalities
 - Bronchiectasis (BR)
 - Ground glass opacity (GGO)
 - Bronchial wall thickening (BWT)
 - Mucus Plugging (MP)
 - Consolidation (Con)
 - Air trapping (AT)





Techniques with hyperpolarized ¹²⁹Xe



¹²⁹Xe ventilation MRI: Detection of early obstruction

14 y.o. male control subject, $FEV_1 = 103\%$ (normal lung function)



All control subjects: uniform ¹²⁹Xe ventilation and low ¹²⁹Xe ventilation defect percentage (VDP)

15 y.o. female CF subject, $FEV_1 = 73\%$



CF: ¹²⁹Xe Ventilation Defect Percentage (VDP) in CF

Control FEV₁ = 115%

CFEV₁ = 81%

CF FEV₁ = 102% Much more sensitive than FEV1 Provides spatial heterogentiy complementary to time-heterogeneity with LCI



RL Thomen et al, J Cyst Fibrosis 2016 Jul 28. Kanhere et al., Am J Respir Crit Care Med 2017, 28 Feb

Combining structure and function

UTE MRI Hyperpolarized Xenon MRI function LU LL RU RM RL

structure

Use for chILDs

A post-infectious BOS case...

UTE MRI

¹²⁹Xe ventilation MRI





Threshold & quantify ventilation deficits (blue)





¹²⁹Xe VDP = 40.7%

IRC186H-**38**: 12 y.o. male BOS patient (post-infectious) FEV₁%-pred = 34%

A milder BOS case...(hyperpolarized ¹²⁹Xe MRI)



10 y.o. male BOS patient FEV₁%-pred = 60% 129 Xe VDP = 22.3 %



Conclusions

Pulmonary MRI is feasible, practical

- UTE MRI can depict lung structural abnormalities
- Hyperpolarized-gas MRI can depict and quantify ventilation abnormalities
 - High sensitivity compared to FEV1 (even higher than LCI)

Early results in BOS, NEHI indicate structure-function MRI may be used to quantify earliest forms of disease

Potential to monitor therapeutic efficacy



Core Faculty

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Pulmonology (adult)





Supplemental slides

Rare-lung diseases: bronchiolitis obliterans syndrome (BOS)

Regional structure (UTE MRI) and function (¹²⁹Xe ventilation MRI) Potential applications in lung- and bone-marrow transplantation

10 y.o. post-infectious BOS: 34% FEV₁









¹²⁹Xe VDP: 40.7% Near absence of ventilation in left lung!

Predicting short-term outcomes via MRI

Respiratory support at discharge in 27 patients: 16 discharged on room air, 4 on O_2 , 4 on a ventilator, 3 died before discharge.

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Term (no BPD) Discharge: Room air MRI score = 0		Mild BPD Room air MRI score = 1		Moderate BPD Oxygen MRI score = 8		Severe BF Ventilato MRI score =	2D r = 13	Severe BPD Death MRI score = 13
	Respiratory so discharge (or	upport at death)	Room air (N=16)	O ₂ (N=4)	Ventilator (N=4)	Death (N=3)		
	MRI Ochiai so	ore	1.2 ± 2.2	4.8 ± 2.1	11.5 ± 1.7	12.7 ± 0.6		

Scores correlated significantly with length of hospital stay (slope = 0.06 [score]/day, P<0.0001).

NS Higano, et al., "Early-life MRI of bronchopulmonary dysplasia predicts short-term outcomes", manuscript in preparation

HP Gas Compatible Ventilator

3D Printed Cradle

Teflon Pneumatic Valve



Pressure Transducer

Exhale

