Lung MRI in Children: The Road Less Travelled

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Abstract

Keywords

- children
- lung
- magnetic resonance imaging
- pediatric

Magnetic resonance imaging (MRI) of the lungs is one of the most underutilized imaging modality when it comes to imaging of thoracic diseases in children. This is largely due to less-than-optimal image quality and multiple technical challenges involved with MRI of the lungs. Advances in MRI technology along with increased awareness about optimization of MR protocol have led to it being viewed as a feasible option for evaluation of various chest diseases in children. This short review article takes the reader to the road less travelled to explore newer horizons for applications of this rapidly evolving magnetic resonance technique in the field of thoracic diseases in children.

Introduction

Chest radiograph is the most common radiological investigation performed worldwide in children. It is considered appropriate and adequate in most of the clinical conditions involving the respiratory system. However, computed tomography (CT) of the chest is often performed wherever detailed anatomical information is required and in some specific clinical indications. It provides additional information but involves radiation exposure, which can have detrimental effects on children.¹ Magnetic resonance imaging (MRI) of the lungs has been underutilized in daily clinical radiological practice. Advances in MRI technology in the past decade have changed the way we look at the lungs and offer comparative radiological information as CT in some specific clinical indications.^{2–4} It has the inherent advantage of being nonionizing and can be repeated in children without any radiation risks. This article aims to provide a short review of indications, protocols, advantages, and limitations of thoracic MRI in children.

Indications

Thoracic MRI can now be performed in virtually all pathologies where CT is performed. Indications can be broadly divided into lung parenchymal, mediastinal, and chest wall pathologies (► **Figs 1–5**). Thoracic MRI has been reported to be of high sensitivity and specificity, nearly at par with CT, in mediastinal masses and cysts, mediastinal lymph nodes, pulmonary embolism, pulmonary nodules larger than 4 mm, bronchiectasis, congenital entities, pleural collections, chest wall tumors, and infections.^{5–9} However, as of now, thoracic MRI is considered inferior to CT in imaging evaluation of interstitial lung diseases, air trapping, pulmonary nodules less than 3 to 4 mm, and calcifications.^{10,11} This is largely due to lower signal resolution in the lungs and susceptibility differences at the interface.

Specifically, thoracic MRI has demonstrated great diagnostic utility as a radiation-free imaging modality in children with infections (tubercular/nontubercular/hydatid), more so in the setting of immunocompromised children and cystic

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Fig. 1 (a) Normal TRUFI, (b) T2W, and (c) T1 VIBE images at the level of the aortic arch in a 7-year-old boy.

Fig. 2 Necrotic nodes (arrows) on (a) BLADE and (b) T2-weighted images in an 8-year-old boy with febrile neutropenia.

fibrosis.¹²⁻¹⁶ Thoracic MRI is also reported to be useful in the evaluation of large airways in children.^{17,18}

►**Table 1** provides a complete list of broad categories of indications of thoracic MRI in children.

MRI Protocol

Typical MR scans in children for lung MRI are performed with either respiratory- or nonrespiratory-gated and nonelectrocardiogram (non-ECG)-gated MRI sequences. Respiratory and ECG gating does offer the advantage of reduced image artifacts and better image quality; however, these gated sequences would entail much higher scan time and this would risk overall compliance of the child for such a study.

In children, time is the most crucial component. MR sequences should be faster and offer great contrast and higher signal-to-noise ratio. Further, to avoid respiratory artifacts, all sequences can be acquired with the breath-holding technique. All children with the help of their parents/guardians should be first taught appropriate

Fig. 3 Left-sided empyema and subsegmental collapse on (a) CT, (b) T2, and (c) BLADE in a 9-year-old girl with fever and chest pain for 4 weeks.

Fig. 4 Consolidation in lingular segments with breakdown on (a) CT, (b) T2, and (c) T1 VIBE in a 12-year-old boy with acute myelogenous leukemia. Mild bronchial wall thickening (arrows) is also noted in the right lower lobe.

Fig. 5 Small pulmonary nodule in the superior lingula on (a) CT, (b) T2, and (c) T1 VIBE in a 12-year-old boy with acute lymphoblastic leukemia and febrile neutropenia.

minimum breath-holding (10–15 seconds). Children who are sick or cannot hold the breath for short intervals cannot undergo breath-hold MRI sequences, and this is one of the limitations of MRI. However, in children, who cannot hold breath, respiratory triggered free-breathing, T2-weighted sequences can then be utilized. It has to be deployed differently in each child and one size cannot fit all.

Table 1 Typical indications of thoracic MRI in children

Lung parenchymal	Mediastinal	Chest wall
Congenital causes	Lymph nodes	Tumors
Infective causes	Masses	Abscesses/infections
Interstitial diseases	Pulmonary embolism	
Airway diseases		
Masses/metastasis		
Vascular abnormalities		
Pleural diseases (associated with parenchymal disease or only pleural)		

Sedation for smaller children who cannot hold breath or cannot lie on their own for an MRI study would be required. This can be one of the limiting factors for MRI of the lungs in smaller children. Hence, its utility is best advocated for children older than 5 years who can be taught breath-holding and who do not require sedation.

Typical MR sequences include (► **Fig. 1**):

- HASTE T2: Half-Fourier-acquisition single-shot turbo spin-echo.
- BLADE T2: Proprietary name for periodically rotated overlapping parallel lines with enhanced reconstruction (PROPELLER) in MR systems.
- TRUFI T2: True fast imaging with steady-state free precession sequence.
- VIBE T1: Volumetric interpolated breath-hold examination.

► **Table 2** summarizes the various intrinsic parameters of the sequences used.

These sequences provide most of the required pathological and anatomical information.^{2–4} Additional sequences, such as STIR (short tau inversion recovery) sequence, diffusion-weighted imaging (DWI), and postcontrast imaging, can be added to a given study, on an individual case basis.

Table 2 Intrinsic parameters of our typical MRI sequences

Parameters	T2 HASTE	T1 VIBE	TRUFI/SSFP
FoV (mm)	275	400	300
FoV phase%	78.1	75	85.9
Base resolution	320	384	256
Phase resolution (%)	80	75	100
Slice thickness (mm)	4	4	4
Phase partial Fourier	4/8	7/8	Off
Pixel size (mm)	0.7 × 0.7 mm	0.7 × 0.7 mm	0.6 × 0.6 mm
Distance factor	20	20	60
TR (ms)	500	4.06	580.23
TE (ms)	36	2.07	1.48
Flip-angle (degree)	146	5	40
Bandwidth (Hz/pixel)	781	540	1,028
iPAT (no. of ref. lines)	2(1)	2(24)	2(26)

Abbreviations: FoV, field of view; HASTE, half-fourier-acquisition single-shot turbo spin-echo; iPAT, integrated parallel acquisition techniques; SSFP, steady-state free precession sequence; TR, repetition time; TE, echo time; TRUFI, true fast imaging with steady-state free precession sequence; VIBE, volumetric interpolated breath-hold examination.

Contrast Administration in Lung MRI

Contrast is not required to be administered in all routine thoracic MRI cases. However, as per indication, and on a case basis, contrast can be given, for example, in mediastinal masses or suspected abscess formation. Few MR sequences (steady-state free precession [SSFP]/TRUFI) depict inherent vascular contrast in mediastinal vessels without the need for giving any contrast. Anatomy of mediastinal vessels and even pulmonary artery embolism can be detected even without contrast administration.

Lungs are scanned in the axial plane in the craniocaudal direction in all children. The total MRI scanning time can vary from 2 to 14 minutes for all sequences combined. Few authors have reported faster (2 minutes) but limited protocol for use in lung MRI in children.^{3,7}

Advances in MRI

Advances in MR technology including newer MRI lung sequences, e.g., ultrashort TE sequence and zero echo time, offer great potential and show promise of yielding very high resolution images at par with high-resolution CT.¹⁹⁻²² Both these newer sequences deploy very short TEs to capture magnetic resonance signals from very short intrinsic T2/T2* tissues. Thus, these provide much superior lung parenchymal images.

Diffusion imaging of the lungs is another promising tool, which has as yet not been explored in the pediatric population. DWI with functional MRI has been reported in adults to provide structural as well as functional information in patients with primary antibody deficiencies.²³ Diffusion imaging of lungs has also been found useful for the detection of inflammatory changes of lung during respiratory tract exacerbations in cystic fibrosis patients.²⁴

Functional assessment of the lung parenchyma is another exciting potential application of lung MRI. Functional MRI studies have demonstrated expiratory–inspiratory MR signal-intensity differences, which can differentiate ventilation defects of different types in cystic fibrosis lung disease, and this kind of functional impairment is reported to be directly related to the diseased structural abnormalities.²⁵ Dynamic ventilation and perfusion studies are performed and perfusion maps are generated, deploying postprocessing software and algorithms, which can guide patient monitoring in these patients.²⁶

Advantages and Limitations of MRI

Thoracic MRI has its inherent set of advantages and limitations, which have been summarized in tabulated form in **Table 3**. It offers a radiation-free alternative to CT scan in most of the common pathologies and provides both anatomic and functional information in one setting.^{15,26,27} It is a particularly more attractive option in children in whom repeated follow-up imaging (e.g., cystic fibrosis) is required as a part of the management protocol. In smaller children, where it

Table 3 Advantages and limitations of lung MRI

Advantages	Limitations
Lack of ionizing radiations	Higher cost factor
Can be repeated: no safety issues of radiations for follow-up in younger children	Not widely available
Higher soft-tissue contrast	Smaller children would require sedation
Vascular information can be obtained without administration of contrast	More image artifacts and poor image quality if child not compliant
Provides both anatomical and functional information in one setting	Less spatial resolution overall
	Lower sensitivity in smaller nodules (< 4 mm), cysts, air trapping, peripheral bronchiectasis
	Lower sensitivity in small calcifications

is not considered safe to administer MR contrast, MRI provides vascular information even without administration of gadolinium-based contrast and specific MR sequences (SSFP) can be performed in these clinical settings.

Conclusion

Lung MRI in children is difficult and challenging due to low signal generated from lungs, relatively longer imaging times, motion artifacts, and nonuniform cooperation by children. However, it offers a radiation-free alternative to CT, and can provide both anatomical and functional assessment of chest in the same sitting. Lung MRI can be utilized in many specific clinical indications, e.g., pulmonary infections in immunocompromised children, lymph node evaluation, cystic fibrosis, and follow-up imaging in children. CT, however, still remains the investigation of choice in evaluation of interstitial lung disease, smaller airways, and pulmonary micronodules. It is important to optimize all cases of lung MRI in children to reduce MRI scan time with judicious planning and adapting MR protocols and get MR images of high quality.

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Conflict of Interest

The authors declare no conflict of interest.

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