Pediatric High-Resolution Computed Tomography

Alan S. Brody, MD





 Alan Brody: alan.brody@cchmc.org

Slides at:
<u>http://www.cincinnatichildrens.org/</u>
<u>research/div/radiology/present.htm</u>

Contributors

Eric Crotty
 Robin Deterding
 Eric Effmann
 Gail Deutsch

Claire Langston
 Fred Long
 Javier Lucaya
 Robert Wood

Pediatric HRCT

What do I do with a request for an HRCT in a child?
How can I get better images?
What do I say about the images once I've got them?

High-Resolution CT

 HRCT provides a low dose sampling technique for diffuse or widespread abnormalities

 HRCT does not evaluate the mediastinum or central airways
 HRCT is very unlikely to be useful if conventional CT is normal

High-Resolution CT

- High-resolution CT is a misleading term
- "Limited sample parenchymal evaluation CT" is more accurate
- HRCT is more technically demanding than conventional CT
- HRCT is only useful if high quality images are obtained

HRCT Technique



HRCT Technique

Weight/size/age based mAs ♦ kVp 80-120 - Increased kVp markedly increases dose ♦ 1mm sections \diamond 7mm (infants) to 20mm (limited survey) intervals for inspiratory images Maximum 1:2 inspiratory to expiratory ratio, minimum of 4 Fastest scan speed

Dose Considerations

Perform HRCT first if unsure whether to do conventional or HRCT
Scan limited area when appropriate
Evaluate the relative dose of available CT scanners
Use breast shields

Cassese, et al. J Thorac Imaging. 2003;18:242-5 Fricke BL, et al. AJR 2003;180:407-11

Breast Shields





30% lower breast dose, no increase in noise

Patient Preparation for HRCT

HRCT requires cooperation or control
Inspiratory images 4 to 6 years old
Expiratory images 6 to 8 years old
Coach in room helpful until 10-12 years old

Patient Preparation

Explain and practice the procedure before entering the scan room
Practice again on the scanner table
Talk your patient through the entire procedure
It helps if an adult stays in the room

during scanning

New Imaging Techniques

- Imaging quietly breathing young children is not adequate for HRCT
 - Motion degrades images
 - Lung volumes are variable, and level of inspiration is unknown
 - Comparable inspiratory and expiratory images cannot be obtained

Controlling Lung Volume

Decubitus imaging
 Controlled ventilation CT
 General anesthesia

Decubitus Imaging



Decubitus Imaging

 Perform initial HRCT
 Place child in lateral decubitus position
 Down side is expiratory, up side is

well inflated

Lucaya, et al. AJR 2000 174:235-41

2 Year Old

Normal appearance



Multiple Bronchial Atresias





5 Year Old, ? Bronchiectasis



5 year old

Bronchi abut the mediastinal pleura indicating bronchiectasis



Controlled Ventilation CT



Controlled Ventilation CT (CVCT)

 Mask ventilate sedated child
 CO₂ and chest stretch receptors produce 10-15 seconds of apnea
 Obtain inspiratory and expiratory images during apneic period

Long et al. Radiology, Aug 1999; 588-93





2 Year Old with CF



Courtesy Frederick R. Long, MD

2 yo with CF, Inspiratory CVCT

[m:1

R

)FOV 22.0cm 30NE

Courtesy Frederick R. Long, MD

ļ

2 yo with CF, CVCT





Courtesy Frederick R. Long, MD

Controlled-Ventilation CT

 Safe; technique used for infant PFTs on thousands of children
 Effective; success rate > 90%

 Requires a trained RT or other HCP
 Must have a well-established sedation program in place

General Anesthesia



General Anesthesia Inspiratory and Expiratory Images



General Anesthesia

The greatest technical problem is atelectasis
Begin scanning as soon as possible
Maintain 30 cm water inspiratory pressure with frequent sighs
Insist on prone images if posterior opacities are present

Interpreting Pediatric HRCT



Interpreting Pediatric HRCT

 "Evaluation of the lung parenchyma is not straightforward in neonates and infants"

David Hansell, HRCT of Diffuse Lung Disease, Radiol Clin North Am, Nov 2001

Interpreting Pediatric HRCT

Evaluate the large and small airways
 Identify the predominant parenchymal abnormalities

- Ground glass Nodules Cysts

Emphysema Linear / reticular densities
 Adult terms work well for description
 Diagnostic possibilities are often very different

Illustrative Cases

Children are not little adults
Common things occur commonly
Make friends with your pathologist
A pediatric pulmonologist may be more helpful than a radiologist

15 Year Old, Shortness of Breath



?? Idiopathic Pulmonary Fibrosis ??

Appearance in children often associated with autoimmune/ connective tissue disorders
Little fibrosis on biopsy
May respond to steroids or hydroxychloroquine
Often stable for long periods of time

Ideopathic pulmonary fibrosis in infants: good prognosis with conservative management. Hacking, et al. Arch Dis Child 2000;83:152-157

?? Idiopathic Pulmonary Fibrosis ??

 Increasingly restrictive criteria in adults

 "Adult" IPF rarely seen in children
 Should not be diagnosed by imaging appearance without biopsy

11 yo with Frequent Infections











Tree-In-Bud

 Inspissated material in distal bronchioles Frequently ascribed to infection, especially non-tuberculous mycobaterium In children without an underlying condition probably most often seen with chronic aspiration

Three Children with Tachypnea



2 year old, Follicular Bronchiolitis



8 year old, Follicular Bronchiolitis



4 year old, Nonspecific Cellular Infiltrate



Follicular Bronchiolitis





Nonspecific Cellular Infiltrate





Follicular Bronchiolitis

 Commonly associated with HIV, collagen vascular disease, and congenital immune deficiency
 Good prognosis in adults, can be progressive in children

Lymphoid Infiltrative Disorders of the Lung

Lymphocytic bronchiolitis
 Lymphoid interstitial pneumonitis
 Follicular Bronchiolitis
 Lymphocytic alveolitis

 Pathologic diagnosis depends on the predominant cell type, the location, and the degree of follicle formation

Pathology Slide Review

The 2 yo with follicular bronchiolitis and the 4 year old with a nonspecific cellular infiltrate had a nearly identical pathologic appearance Diagnosis was most likely lymphoid bronchiolitis in both cases The 8 year old with follicular bronchiolitis had a very different appearance

Neonate with Cyanosis



Courtesy Robin Deterding, MD

Congenital Pulmonary Alveolar Proteinosis



Pulmonary Alveolar Proteinosis

 Proteinaceous fluid fills alveoli Variable prognosis, usually poor in newborns AP is a specific pulmonary response to insult, but it is not an etiology Associations include surfactant protein abnormalities, inhalation exposure, lipoid aspiration, immune compromise

Pulmonary Alveolar Proteinosis

 Prognosis in children depends on etiology

 Identification of PAP requires a complete evaluation for known associations

Five yo with Chronic Lung Disease



13 yo with Chronic Lung Disease



Surfactant Protein C Deficiency

One of several surfactant protein abnormalities
Increasingly recognized as a cause of chronic lung disease
Variable course and presentation
Infection likely causes lung damage out of proportion to the infection

Sisters with Surfactant Protein C Deficiency



5 year old

13 year old

Niemann-Pick Type B



Niemann-Pick Type B

 Enzyme defect leads to buildup of sphingomyelin

 Type B does not involve the CNS, symptoms are abdominal distension and respiratory

 Abnormal macrophages ("foam cells") accumulate in the lungs producing nodular and ground glass opacities

Niemann-Pick Type B

 Pathophysiology of lung disease and imaging appearance is well understood

 There is no significant relationship between the severity of radiographic abnormality and the severity of respiratory symptoms

Conclusion

 New techniques allow high quality images in patients of all ages
 Identifying the presence and location of disease is an important contribution
 HRCT findings are rarely specific, and lung biopsy will often be needed

Thank You for Your Attention

Alan Brody alan.brody@cchmc.org

