

Tuberculosis in Children

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> TB Intensive September 13 – 15, 2023 Richmond, TX

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Andrea Cruz, MD, MPH has the following disclosures to make:

- No conflict of interests
- No relevant financial relationships with any commercial companies pertaining to this educational activity



Tuberculosis in Children



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Disclosures

- •Associate editor, *Pediatrics*
- •Levofloxacin is off-label for tuberculosis in children
- •Pictures shared with permission of families, who wished to increase TB awareness among clinicians and policymakers





- •14-month-old girl, no medical history, presented to ED with seizures, no return to baseline
- •Progressively more altered → intubated
- •Fever x 2 weeks, vomiting → prior diagnoses of gastroenteritis
- •CT brain (uncontrasted): prominent ventricles, hypoattenuation of R basal ganglia
- •CSF: 450 WBC, 2 RBC, protein 800, glucose < 20, Gram stain: no organisms seen

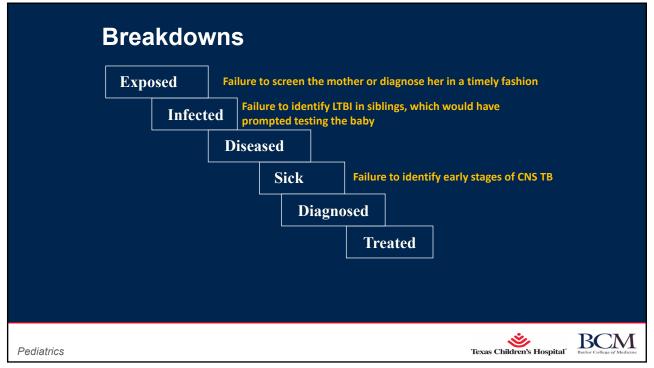
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Objectives

- •To identify when to think about TB
- •To recognize potential complications of TB and what forms of TB can result in rapid decompensation
- •To formulate a diagnostic and therapeutic treatment plan for children with TB infection

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TB Definitions

Class	Sx	Exam	TST/ IGRA	CXR	Contagious	Treatment
Infection	-	-	+	-	Never	Usually 1-2 drugs, given 3-9 months (given by family or health department)
Disease	+	-/+	+/-	+/-	Rarely	Multiple drugs (3-4), given 6-12 months (always given by health department)

TST: tuberculin skin test IGRA: interferon gamma release assay

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US TB Epidemiology: 2022

- •8,300 cases; incidence: 2.2/100,000
- •73% in the foreign-born
 - -Top 5 countries of birth: Mexico (19% of non-US-born persons), Philippines (12%), India (10%), Vietnam (8%), China (6%)
- ·Largest increases:
 - -Children <=4y: up by 29%
 - -15-24y: up by 24%
- •4 states (CA, FL, TX, NY): 50% of US cases

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MMWR 2021;70:409





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Clinical Manifestations



Risk of Progression from TB Infection to Disease by Age

Age at infection (y)	No disease (%)	Pulmonary TB (%)	CNS TB (%)
<1	50	30-40	10-20
1-2	75-80	10-20	2.5
2-5	95	5	0.5
5-10	98	2	<0.5
>10	80-90	10-20	<0.5

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Peds in Review 2010;31:13





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Childhood TB Disease Sites

Site*	% of cases	Median Age (years)
Pulmonary	77.5	6
Lymphatic	13.3	5
Pleural	3.1	16
Meningeal	1.9	2
Bone/joint	1.2	8
Miliary	0.9	1
GU	0.8	16
Peritoneal	0.3	13

*: United States (almost all are immunocompetent)





Signs & Symptoms of Pulmonary TB

Feature		Infant	Child	Adolescent
Symptom	Fever	+++	+	+++
	Cough	+++	+++	+++
	Night sweats	-	-	+
	Hemoptysis	-	-	+
Sign	Rales	+++	-	+
	Wheezing	+++	-	-
	Decreased breath sounds	+++	-	+
Site	Intrathoracic	+++	+++	+++
	Intra + extrathoracic	+++	-	-

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When should I suspect TB disease?

- •Epidemiology:
 - -Child has contact to a person with suspected TB
- •Symptoms:
 - -Prolonged symptom duration, failed therapy for PNA or adenitis
 - -Pneumonia + weight loss
 - -Recurrent or persistent pneumonia
- •Lab/Radiology findings:
 - -Meningitis + abnormal CXR
 - -CSF pleocytosis with ↑ protein, ↓ glucose
 - -Altered mentation with strokes and/or new-onset hydrocephalus
 - -Certain radiographic patterns

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What will kill a child?

- •TB meningitis: increased ICP
- •Miliary TB: tension pneumothorax (early), bronchiolitis obliterans (late)
- •Lymphadenopathy: airway compression
- Pericarditis
- •Myocarditis: predilection for the conducting system
- •Hemorrhage: erosion of cavity into vessel
- Congenital TB



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CXR Findings in Pediatric TB

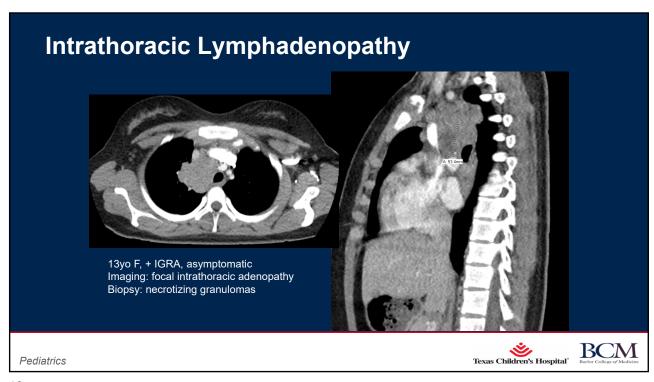
- •Hilar or mediastinal adenopathy
- Segmental/lobar infiltrates
- •Calcifications (seen in 75-80% of children with pulmonary TB)
- Miliary disease
- Pleural effusions

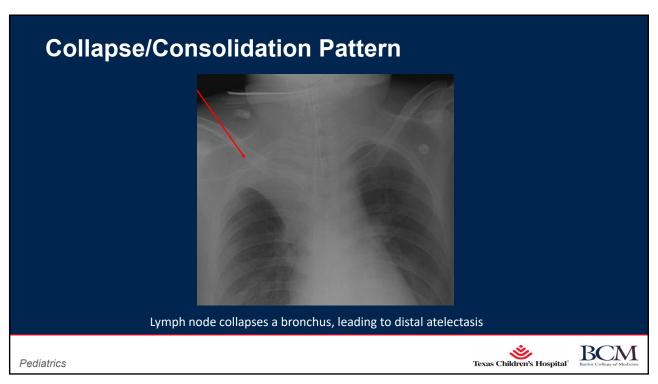
Except for CNS disease, the CXR often looks sicker than the patient

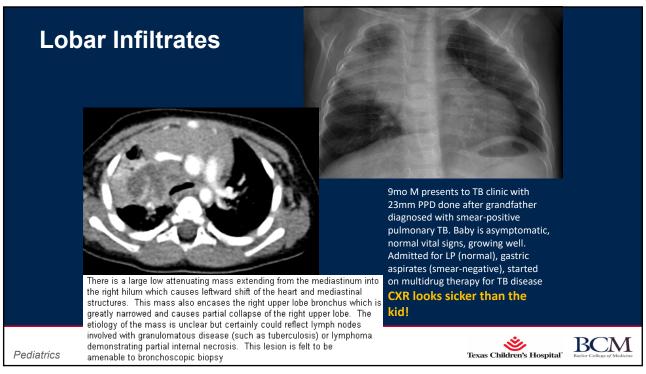
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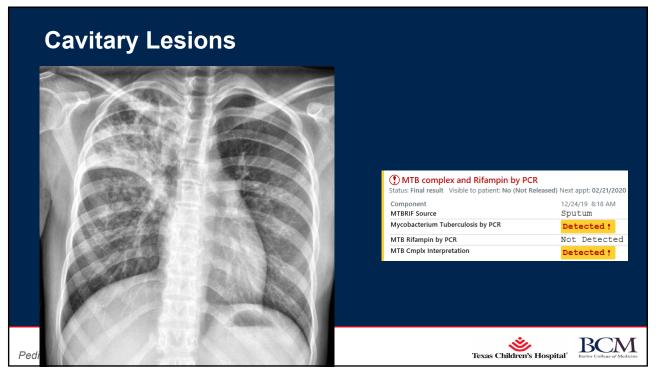


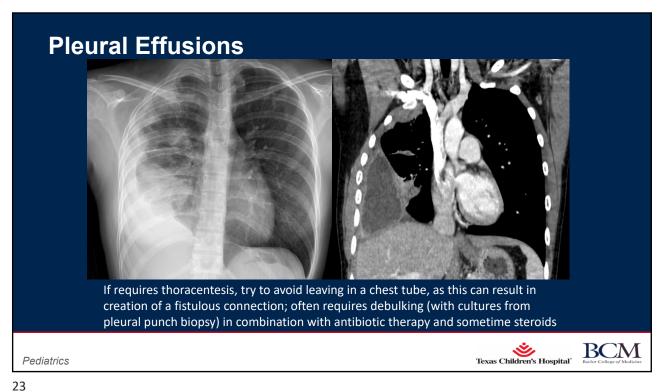












TB Meningitis: Imaging Findings

CT Feature	Sensitivity	Specificity	PPV	NPV	LR+
Basilar enhancement	89%	94%	94%	88%	14.3
Hydrocephalus	68%	72%	74%	66%	2.4
Infarcts	62%	78%	77%	64%	2.8
Granuloma	14%	94%	71%	48%	2.2
"Full House"	41%	100%	100%	59%	N/A

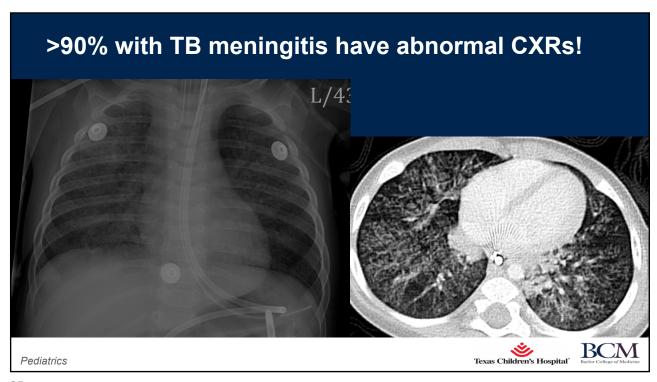
"Full house" – combination of basal enhancement, infarction, and hydrocephalus

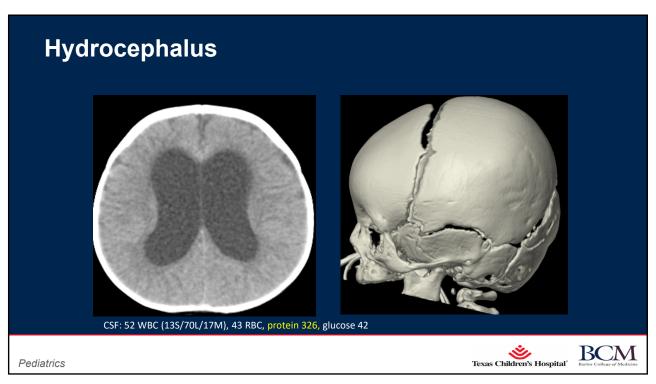
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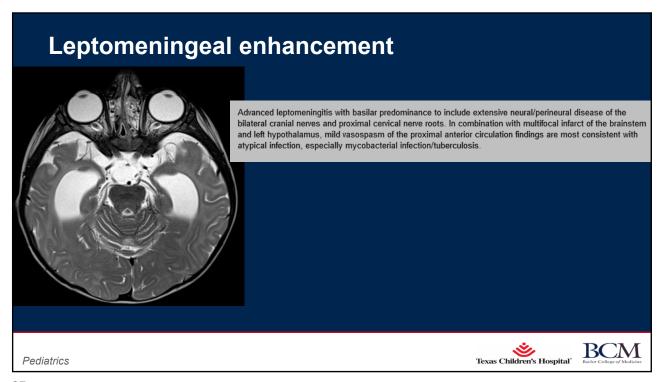
Pediatr Radiol 2004;34:876-885

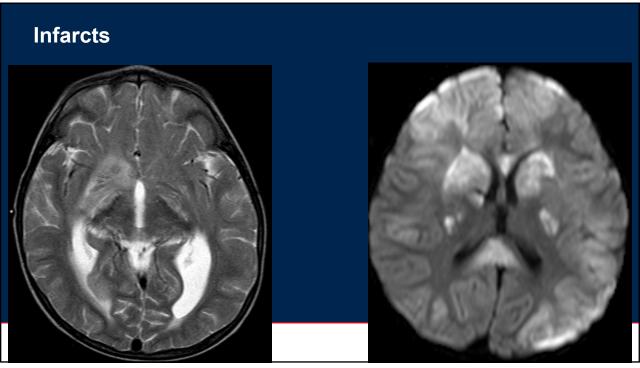


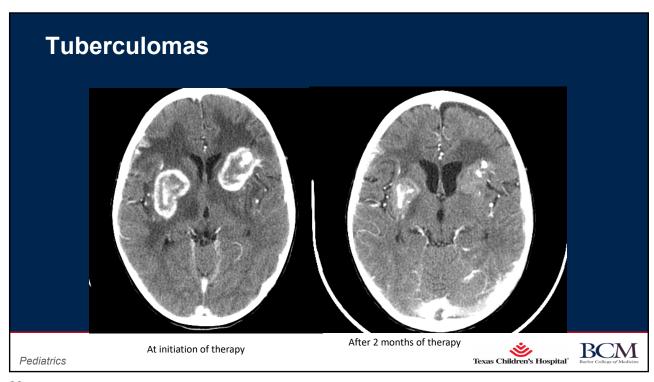












Who do I tap?

- •Assuming it is safe to tap them from a herniation standpoint; would obtain CNS imaging first
- •Any child with symptoms concerning for TB meningitis
- •Any infant in first year of life with suspected pulmonary TB
- Consider in any child with miliary pattern on CXR

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Who do I image, preferably with MRI?

- •All children with pleocytosis
- Children too unstable to do lumbar puncture
- •All infants with miliary TB, even if they do not have a CSF pleocytosis
- •Why do I image?
 - Can see tuberculomas in absence of meningitis
 - Young children can have unreliable physical examinations
 - Exam findings can be subtle early on
 - Changes management

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Why do we insist on LP (and MRI)?

- Changes drugs used
 - -Ethambutol penetrates blood/brain barrier poorly, so we instead use ethionamide
- Changes decision to use systemic steroids
 - -Usually 1 month with slow taper
- Changes duration of therapy
 - -Minimum 9-12 months for CNS TB
- Changes prognostic conversations
 - -Most children with TB meningitis suffer sequelae





TB adenitis

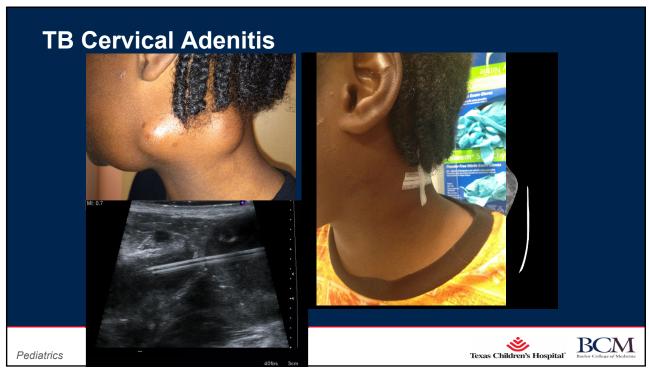
- •Painless, can be come tethered to overlying skin, central fluctuance
- •Reddish or violaceous discoloration
- •Can form draining sinus tracts
- •AVOID incision and drainage; prefer FNA or needle aspiration

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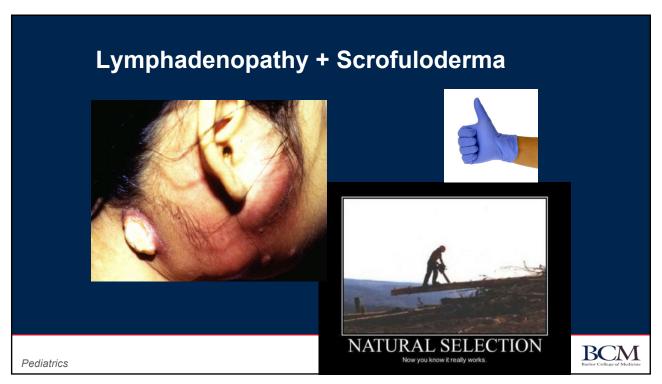


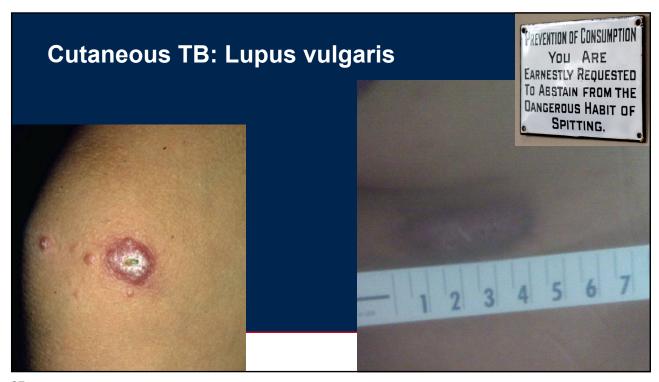


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Baseline testing

- •CBC, HIV, LFTs
- •Place TST, order TSPOT and QuantiFERON interferon gamma release assays
- •Xpert TB PCR (also detects RIF resistance)
- •CXR, other appropriate imaging studies
- •LP if indicated
- •CXRs on 2 caregivers (done at TCH expense)
- •Respiratory samples: gastrics vs sputum induction vs expectorated sputa

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Acid-Fast Culture Yield

Specimen	Culture Yield	
Sputum/gastric aspirate	30-40%	
Lymphatic tissue	75%	
Pleural fluid	20-40%	
Cerebrospinal fluid	20-50%	
Pericardial fluid	0-42%	
Ascitic fluid	30%	
Skin biopsy	20-50%	
Skeletal biopsy	75%	

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Paed Resp Rev 2007;8:107





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Diagnosis of pediatric TB

- •Given poor culture yield, the diagnosis of childhood TB is usually based on the following triad:
 - -Positive PPD or IGRA
 - -Compatible radiographic and/or clinical findings
 - -Epidemiologic link to a person with known or suspected TB

If you wait upon microbiologic confirmation for children with suspected TB, you will be grossly under-diagnosing pediatric TB and kids will die

Public health contact tracing invaluable

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Initial regimens (not discussing MDR)

- •Non-CNS:
 - Rifampin, INH, pyrazinamide, ethambutol (RIPE)
 - SHINE Trial results: 4m course of therapy
- •CNS: Start minimum 4-drug therapy
 - INH, rifampin, pyrazinamide, ethionamide, and often add another drug (e.g., levofloxacin)
 - -Steroids
 - -Consider admission to the ICU

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Who gets steroids?

- •Always:
 - -CNS disease (meningitis, tuberculomas)
 - -Pericarditis
- •Sometimes:
 - -Pleural disease
 - -Bulky lymphatic disease
 - -Endobronchial disease
 - -Immune reconstitution / paradoxical worsening





Acute decompensation

- •Neurologic:
 - -Hydrocephalus
 - -Shunt malfunction
 - -Worsening of tuberculomas
- •Respiratory:
 - -IRIS with worsening of adenopathy around the airway
 - -Pneumothorax (tension or otherwise)
 - -Erosion into vessel
- •Cardiac(ish):
 - -PEA in child with miliary disease: needle the chest

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Notes on TB Drugs

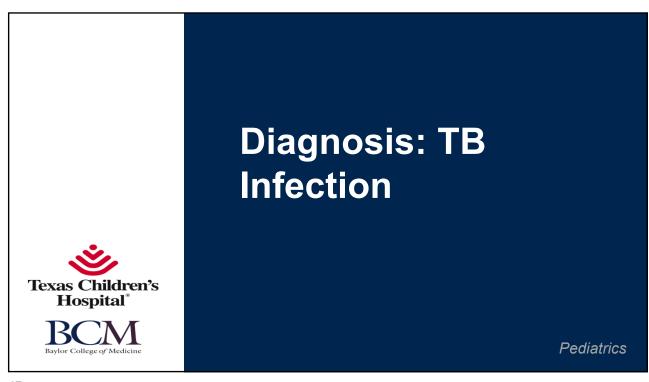
Drug	Side Effects	Other notes
INH	Peripheral neuropathy; seizures in overdose	B6 helps prevent neuropathy and is only treatment for INH seizures, but doesn't prevent hepatotoxicity
RIF	Orange discoloration of secretions; inactivates oral contraceptives; many drug interactions	Please warn of Astros-orange urine!
PZA	Can increase uric acid → gout symptoms; rash	Of 1 st -line drugs, greatest association with hepatotoxicity
EMB	Optic neuritis, red-green color blindness	Despite side effects, has very poor CNS penetrance and not used for meningitis

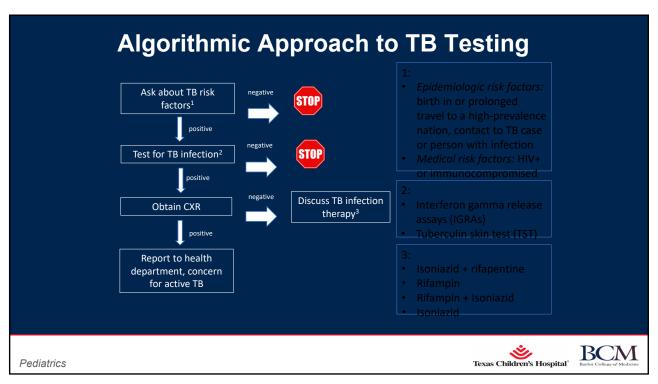
*All primarily hepatically metabolized, except EMB, which is also renally excreted

Peds in Review 2010;31:13 Pediatrics









Comparison of Skin T	est & IGRA

Characteristic	TST	IGRA
Antigens studied	Many -PPD	ESAT-6, CFP-10, (TB-7.7)
Cross-reactivity with BCG	Yes	Unlikely
Cross-reactivity with NTM	Yes	Less Likely
Estimated sensitivity, TB in immunocompetent adults	75-90%	75-95%
Estimated specificity, TB in immunocompetent adults	70-95%	90-100%
Distinguish between TB infection and TB disease	No	No
Boosting	Yes	No
Patient visits required	Two	One

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Positive PPDs

- Generally, skin test conversion occurs within 2 months of contact
- Measure only induration
- Record millimeters of induration (never record "+" or "-")
- Any induration seen only in the first 24 hours should be ignored
- Induration after 72 hours counts
- Blistering also counts



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What is a Positive PPD? ≥ 5mm ≥ 10mm ≥15mm HIV-infected Children < 4 years of age Anyone, even without risk factors Contact to a TB case Children exposed to high-risk adults† Child in whom you suspect TB Immigrants from high-prevalence disease regions* Children with diabetes or other immunocompromising conditions † HIV-infected, incarcerated, IV drug use *Low prevalence regions: US, Canada, Scandinavia, Western Europe, Australia, New Zealand Texas Children's Hospital BCM**Pediatrics**

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PPD Limitations

False positives:

- •Exposure to mycobacteria other than
- BCG vaccine
- False negatives:
- Corticosteroid usage
- Other immunocompromise
- Viral suppression: measles, mumps, influenza

- •Inter-observer variability
- ·Sliding scale for what is considered positive can be confusing
- •Until very recently, lack of any confirmatory tests





Recommendation	2015	2018	2021	2024
Age	≥ 5 years	≥ 2 years	Unchanged, but indicated some experts will use down to 1 year of age	All ages!!
Preferred test for BCG- immunized children	Yes	Unchanged	Unchanged	Unchanged
Use in immunocompromised children (both TST and IGRA)	Cautiously	Unchanged	Unchanged	Unchanged

Rec	2015	2018	2021 (no change for 2024)
Preferred regimen	INH	No specific preference (this is order in Red Book): • 3m INH + Rifapentine* • 4m Rifampin • 9m INH	Several regimens are recommended, depending on circumstance: • 3m INH + Rifapentine • 4m Rifampin • 3m INH + Rifampin (if 3HP or 4R not feasible) • 6-9m INH
RIF role	Limited: INH intolerance or INH resistance in child's contact	Expanded	Unchanged
3НР	Use in ≥ 12 years	Use in ≥ 2 years	Unchanged

- •14yo F from Afghanistan, + IGRA, no symptoms
- ·Massive mediastinal mass, encasing the aorta, internal jugular, subclavian

•What next?





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Tissue is important

- Admitted for oncology evaluation
- •Trans-bronchial biopsy of the mass showed necrotizing granulomas
- •Smear, PCR, culture negative
- •Being treated empirically for TB





- •3-month-old boy, US born, recurrent admissions for fever, no cause identified
- •Concern for miliary TB
- •CSF: 4 wbc, 5 rbc, protein 36, glucose 53, CSF PCR negative



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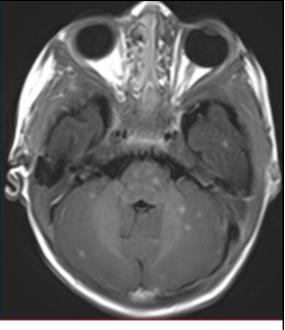
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Neuroimaging is your friend!

What does this change?

- Decision to use steroids
- •Use of 'E' drug

Take-home: you can see tuberculomas in absence of meningitis!



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- •6-month-old presents with fever, vomiting, seizures
- •CT head shows hydrocephalus, CXR shows miliary pattern
- •To OR with neurosurgery → VPS placed

Source	WBC	RBC	Protein	Glucose
EVD	10	500	46	65

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Next steps

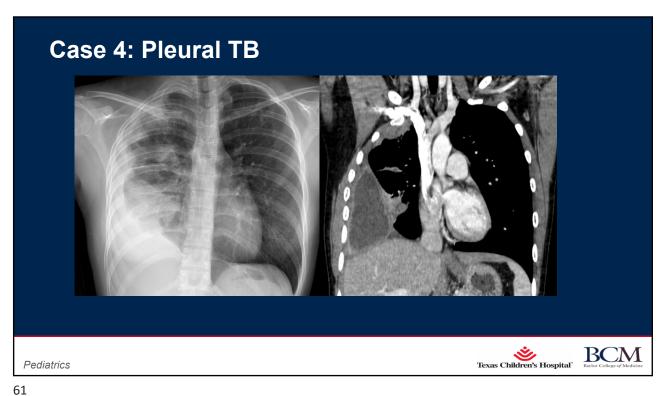
- •You ask the ICU to perform an LP
- •Response: Why, Andrea? We already have CSF from the EVD

Source	WBC	RBC	Protein	Glucose
EVD	10	500	46	65
LP	94	3	173	<20

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When do I consider steroids?

- •Worsening effusion and other factors excluded:
 - -Adherence
 - -Malabsorption / metabolism
 - -Drug resistance
- •Have to consider immune reconstitution inflammatory syndrome (IRIS)
- •If tap the effusion, don't leave chest tube in!!!!

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Case 5: Oy, a rash....

- •2-month-old infant exposed to her mom, who has INH monoresistant TB
- •Baby asymptomatic, normal exam and CXR
- Start rifampin
- •Rash → CBC, LFTs normal, no findings of anaphylaxis or DRESS
- •What now?

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Risk/benefit

- •High risk of progression if infected
- Quinolones are well tolerated
- •Could consider levofloxacin for younger children and moxifloxacin for older children





Case 5 (part deux)

- •Father is in clinic with the baby
- •He wonders why management is different for the baby than for his 2 older children (7, 12 years old)
 - -This comes up all the time with children of different ages
- •What do you tell him?

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Case 5 (part deux)

- •Infant's risk of progression to disease is high
 - -Intrathoracic
 - -Disseminated
- •Less confidence in a negative test in a very young exposed infant





Change in Practice

- Children are often the first in the family to be diagnosed
 - Often, there is no apparent family history at first
- Certain combinations of symptoms and radiographic findings should increase your suspicion for TB
 - Meningitis + pneumonia
 - Pneumonia + weight loss
 - Pneumonia and lymphadenitis refractory to therapy
- Traditional culture techniques are of much lower yield in children as compared with adults
 - Start medications before microbiologic confirmation

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Reference List

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- •Updated guidelines for using IGRAs to detect M. tuberculosis infection. MMWR. 2010;59:1.
- •Guidelines for the treatment of LTBI. MMWR. 2020;69:1
- •Updated recommendation for use of once-weekly isoniazidrifapentine to treat LTBI. MMWR. 2018;67:723.
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