



Approach to cervical lymphadenopathy: A case-based scenario

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Case introduction

- A 6-year-old boy was brought to your clinic with unilateral cervical LAP since 10 days earlier.
- The LN is 1.5*1.5 cm in the submandibular area
- LN features: soft, non-tender, mobile
- General PE is normal

Definition

Lymphadenopathy refers to enlargement of the lymph node.

➤ **Size:**

Larger than 1cm in cervical, axillary

1.5 cm in inguinal

0.5 cm in epitrochlear area

any size in supraclavicular areas

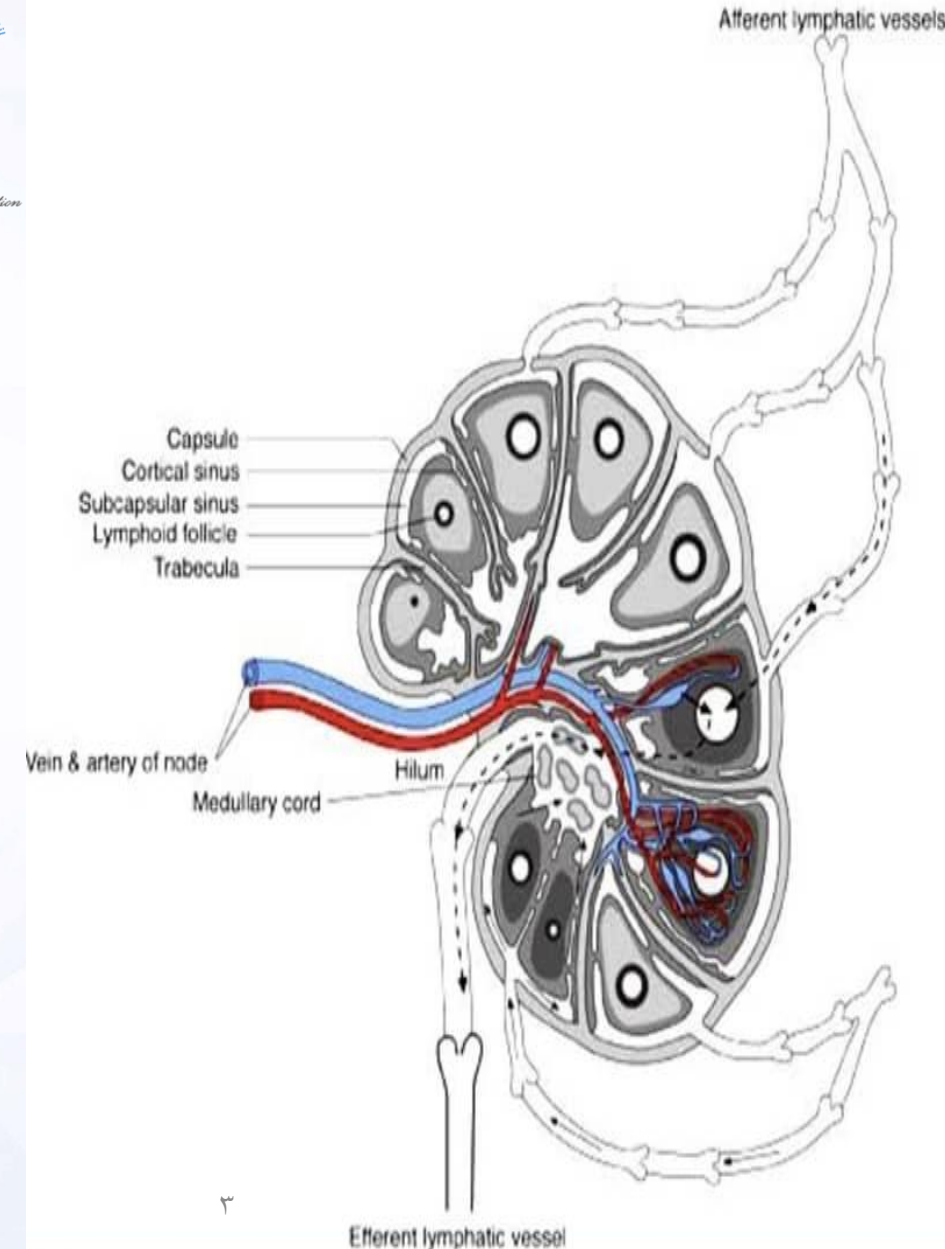
➤ **Duration:**

Acute LAP if less than 2 weeks duration

Subacute if 2-6 weeks duration

Chronic if not resolved by 6 weeks

➤ **Localized vs generalized**



Differential Dx of cervical LAP

Table III. Common congenital neck masses.

Neck mass	Characteristics	Common age of presentation
Thyroglossal duct cysts	<ul style="list-style-type: none">• Most common congenital neck mass• Moves up with tongue protrusion	2-10 years
Branchial cleft anomalies	<ul style="list-style-type: none">• Second most common congenital neck mass• Includes cysts, sinuses and fistulas• Manifests as presence of pit, dimple, sinus or mass along anterior margin of sternocleidomastoid muscle	Fistulas: infancy Cysts: older children > 10 years but can present at any age
Cystic hygroma	<ul style="list-style-type: none">• Soft, painless, fluctuant masses with violaceous appearance• Transilluminable	Birth to 2 years
Dermoid cysts	<ul style="list-style-type: none">• Midline lesions that do not move up on swallowing or tongue protrusion• Rubbery consistency, attached to overlying skin	Aged < 5 years
Congenital muscular torticollis	<ul style="list-style-type: none">• Firm, fibrous mass within belly of sternocleidomastoid muscle• Causes torticollis and positional plagiocephaly	Birth to first few weeks of life
Infantile haemangioma	<ul style="list-style-type: none">• Flat purplish lesions at birth• Enlarge in first year of life – soft, compressible, red or purple masses, not transilluminable• Spontaneous involution thereafter	Birth

Causes of generalized LAP

- Infections: viral bacterial spirochetal protozoal fungal
- Malignancy
- Lymphoproliferative disorders
- Immunologic
- Endocrine
- Drugs
- Miscellaneous

Clinical approach

Important aspects of History:

- **Characteristics of the lymph node enlargement.**
- **Associated symptoms(local or systemic)**
- **potential exposures**
- **Past medical history.**

Physical examination

- **General appearance, vital signs, growth parameters.**
- **Head & neck**
- **chest**
- **Abdomen**
- **Skin**
- **Lymph nodes**



LN features

Location

Size

Consistency

Fixation

Tenderness

Clinical features to differentiate benign from malignant lymphadenopathy

Features	Malignant	Benign
Size	> 2cm	< 2 cm (<1 cm)
Consistency	Hard, firm or rubbery	Soft
Duration	> 2 weeks	< 2 weeks
Mobility	Fixed, matted	Mobile
Location	Supraclavicular, epitrochlear, generalized	Inguinal, submandibular
Tenderness	Usually non-tender	Usually tender

Q1: Early LN biopsy is mandatory except?

- 1- high ESR
- 2- mediastinal widening in chest X-ray
- 3- Pancytopenia in CBC
- 4- night sweating

Answer: 1 is correct

Clinical features worrisome for malignancy or granulomatous disease in children with peripheral lymphadenopathy



- Systemic symptoms
(fever > 1 week, night sweats, weight loss [$> 10\%$ of body weight])
- Supraclavicular (lower cervical) nodes
- Generalized lymphadenopathy
- Fixed non-tender nodes in the absence of other symptoms; matted nodes
- Nontender lymph nodes > 1 cm with onset in the neonatal period

Clinical features worrisome for malignancy or granulomatous disease in children with peripheral lymphadenopathy



- Nontender lymph nodes ≥ 2 cm in diameter that increase in size from baseline or do not respond to antibiotic therapy
- Abnormal Chest X-ray (particularly mediastinal mass or hilar adenopathy)
- Abnormal CBC (lymphoblasts, cytopenia in more than 1 cell line)
- Absence of symptoms in the ear, nose, and throat regions
- Persistently elevated ESR/CRP or rising ESR/CRP despite antibiotic therapy

No symptoms or signs of infection, <2 cm in diameter

- No worrisome features
- No obvious cause based on the history and examination

Q2: What is the next step?

- 1- A 2-week trial of antibiotic
- 2- CBC, ESR, CRP, LDH
- 3- LN. biopsy
- 4- Observation for 2 weeks

Answer: 4 is correct



- For children with cervical lymphadenopathy <2 cm
- no findings of infection within or distal to the node
- no worrisome features
- no obvious cause based on the history and examination
- we observe for 10 to 14 days

The patient was observed for 2 weeks, but the LN does not regress in size.

Q3: What will you recommend next?

- 1- CBC, ESR, CRP
- 2- Serology for EBV, CMV, HIV
- 3- A trial of empiric antibiotic therapy
- 4- All of the above

Answer: 4 is correct



- If the lymph node does not regress or enlarges, then a course of antibiotic therapy may be indicated.
- We also obtain CBC, ESR/CRP, and serology for EBV, CMV, and HIV
- evaluate for Kawasaki disease and other uncommon causes of cervical lymphadenopathy as indicated by the history and examination.

Q4: What will be the choice of antibiotic?

- 1- Cefixime
- 2- Azithromycin
- 3- Clindamycin
- 4- Cephalexin

Answer: 3 is correct

Empiric antibiotic therapy

- Coverage for common pathogens such as group A *Streptococcus* and *S. aureus*
 - High CA-MRSA prevalence: Clindamycin
 - Low CA-MRSA prevalence – First-generation cephalosporin (eg, cephalexin) or amoxicillin-clavulanate
 - For patients with exposure to cats or kittens, we also include coverage for *B. henselae* (eg, azithromycin) in the initial regimen

If the patient's systemic symptoms (eg. fever) do not improve within **72 hours** or the lymph node increases in size (at any point during treatment):

- For patients initially treated with a first-generation cephalosporin or amoxicillin-clavulanate, we switch to clindamycin to provide coverage for CA-MRSA

- For patients with exposure to cats or kittens, we add coverage for *B. henselae* (eg, [azithromycin](#)) if it was not included in the initial regimen

- For patients initially treated with clindamycin, we may add **coverage for NTM** if there are clinical features suggestive of NTM infection and contraindications to surgical excision (if the suspicion for NTM is high, **excisional biopsy** is more appropriate)

Case scenario (continued)

- The laboratory tests are within the normal range
- Clindamycin was prescribed for 2 weeks
- The next visit revealed a non-tender, mobile, soft, 1.5*1 cm LN in the submandibular area
- General PE is normal
- No worrisome features

Q5: What will you do next?

- 1- add azithromycin to the antibiotic regimen
- 2- Excisional biopsy
- 3- Continue observation
- 4- Both 2 & 3 can be advised

Answer: 4 is correct



- We obtain a **biopsy** after four weeks if the diagnosis remains uncertain and the lymph node has not regressed in size.
- However, **continued observation** may be reasonable if there are no worrisome features.

For children with cervical lymphadenopathy ≥ 2 cm
no findings of infection, and no worrisome features

- Initial Laboratory investigation:
 - CBC, ESR, CRP
 - Chest X-ray
- If features suggestive of malignancy:
 - Early biopsy

LN \geq 2 cm, no sign of infection, no worrisome features; normal initial evaluation



Q6: what will you do next?

- 1- tuberculin skin test
- 2- Empiric antibiotic therapy for 2 weeks
- 3- both 1& 2 are correct
- 4- excisional LN. biopsy

Answer: 3 is correct



- IF TST is negative and no response to the antibiotic trial:
 - Serology for EBV, CMV, HIV
 - Evaluate for Kawasaki disease and other uncommon causes of cervical LAP
 - **LN biopsy** if LN does not regress after **4 weeks**

Thank you

