

# Lofgren's syndrome – A Case Report

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## ABSTRACT

Lofgren's syndrome is an acute form of sarcoidosis, characterized by the triad of polyarthrits, hilar adenopathy and erythema nodosum.

Here, we report a 41-year-old male who presented with polyarthrits and reddish rash all over the body.

Skin biopsy revealed the reddish rash to be erythema nodosum.

Chest imaging showed enlarged multiple mediastinal lymph nodes.

Endoscopy fine needle aspiration of mediastinal lymph node showed noncaseating granuloma consistent with sarcoidosis.

## INTRODUCTION

1) Lofgren's syndrome is characterized by the triad of acute polyarthrits, hilar adenopathy and erythema nodosum.

2) It is an acute form of sarcoidosis<sup>1</sup>

3) It is a self-limiting disease with a very good prognosis<sup>1</sup>

4) Non steroidal anti-inflammatory drugs (NSAIDs) and steroids along with supportive care and close monitoring, resolve the symptoms for treatment<sup>1</sup>

## CASE HISTORY

A 41-year-old male presented with multiple joint pain and swelling of 3 weeks duration.

Ankles, knees, wrist, proximal interphalangeal and metacarpophalangeal joints were involved.

There was associated fever with reddish colored rashes all over the body.

He gives history of heaviness in the chest and weight loss.

There was no past history of tuberculosis and bronchial asthma.

## EXAMINATION

1) Afebrile and hemodynamically stable.

2) Significant synovitis around both knees and ankle joints.

3) ESR was high (50mm/hr).

4) RA, ANA and ANCA serology were all negative

5) Serum calcium was 9.2 mg/dl.

6) Skin biopsy of the reddish rash was consistent with erythema nodosum.

7) Chest radiograph showed bilateral hilar prominence.

8) CT chest revealed multiple enlarged mediastinal lymphnodes in paratracheal, pretracheal, subcarinal, prevascular right and left hilar regions (Figure: 1,2).

9) Angiotensin converting enzyme level was within the normal range (38 IU/L).

10) Endoscopy fine needle aspiration of mediastinal lymphnode showed non caseating granulomas consistent with sarcoidosis (Figure: 3,4).

## DIAGNOSIS

The symptoms, clinical findings, imaging and biopsy results in our patient fulfilled the triad of acute polyarthrits, hilar adenopathy and erythema nodosum.

The patient was managed with low dose steroids and NSAIDs.

## DISCUSSION

1) Sven Lofgren first described in 1955.

2) Biopsy: definitive diagnosis of sarcoidosis.

3) Differentials: fungal infection, tuberculosis, lymphomas, and bronchogenic carcinoma.

4) Histopathology of the lymph nodes reveals non-caseating granulomas.

5) Closely related to HLA-B8 and DR3 in Caucasians<sup>2</sup>

6) Chest radiographic findings are seen in approximately 90 % of patients with sarcoidosis.

7) Our case can be categorized as stage 1, according to Stoltzsch classification, based on the presence of bilateral hilar with mediastinal adenopathy, without pulmonary infiltrates.

8) Treatment: NSAIDs + Bed rest

9) Steroids can be used in serious arthritis, hypercalcaemia, and granulomatous skin lesions<sup>3</sup>

## CONCLUSIONS

1) Lofgren's syndrome can present as fever with generalized rash and arthralgia.

2) Lofgren's syndrome mimics viral exanthematous fever.

3) Lofgren's syndrome has a very good prognosis.

## REFERENCES

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- 3) Ohta H, Nakase H, Nakamura A, et al. Acute-onset sarcoidosis with erythema nodosum and polyarthralgia (Lofgren's syndrome): a report of a case report and a review of the literature. Intern Med. 2005; 44(5):559-562.



Figure 1: Axial CT



Figure 2: Coronal CT

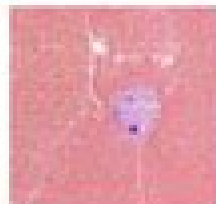


Figure 3: x 400

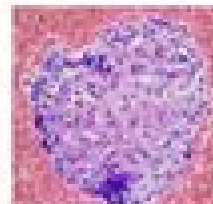


Figure 4: x 400

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## **Table of Contents Poster Presentation Case Report**

1. Understanding the eBook Poster Presentation Case Report
  - The Rise of Digital Reading Poster Presentation Case Report
  - Advantages of eBooks Over Traditional Books
2. Identifying Poster Presentation Case Report
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an Poster Presentation Case Report
  - User-Friendly Interface
4. Exploring eBook Recommendations from Poster Presentation Case Report
  - Personalized Recommendations



- Poster Presentation Case Report User Reviews and Ratings
- Poster Presentation Case Report and Bestseller Lists
- 5. Accessing Poster Presentation Case Report Free and Paid eBooks
  - Poster Presentation Case Report Public Domain eBooks
  - Poster Presentation Case Report eBook Subscription Services
  - Poster Presentation Case Report Budget-Friendly Options
- 6. Navigating Poster Presentation Case Report eBook Formats
  - ePub, PDF, MOBI, and More
  - Poster Presentation Case Report Compatibility with Devices
  - Poster Presentation Case Report Enhanced eBook Features
- 7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of Poster Presentation Case Report
  - Highlighting and Note-Taking Poster Presentation Case Report
  - Interactive Elements Poster Presentation Case Report
- 8. Staying Engaged with Poster Presentation Case Report
  - Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - Following Authors and Publishers Poster Presentation Case Report
- 9. Balancing eBooks and Physical Books Poster Presentation Case Report
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection Poster Presentation Case Report
- 10. Overcoming Reading Challenges
  - Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
- 11. Cultivating a Reading Routine Poster Presentation Case Report
  - Setting Reading Goals Poster Presentation Case Report
  - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Poster Presentation Case Report
  - Fact-Checking eBook Content of Poster Presentation Case Report

- Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
  - Utilizing eBooks for Skill Development
  - Exploring Educational eBooks
- 14. Embracing eBook Trends
  - Integration of Multimedia Elements
  - Interactive and Gamified eBooks

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