


1-1-2020

Kikuchi-Fujimoto Disease

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Recommended Citation

Karkra, Rohan (2020) "Kikuchi-Fujimoto Disease," *Digital Journal of Clinical Medicine*: Vol. 2: Iss. 1, Article 7.

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Kikuchi-Fujimoto Disease

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CLINICAL HISTORY:

A 25 year male patient came with the complaints fever with chills since 3 months. Fever was high grade, intermittent and showed evening rise of temperature

He gives h/o myalgia, fatigue, loss of appetite and significant weight loss.

No h/o of cough, rashes, abdominal pain, diarrhoea.

Not a k/c/o diabetes mellitus, hypertension, thyroid disorders, cardiac disorders

Past History:

He was a k/c/o of kikuchi fujimoto disease, diagnosed 3 years back at same hospital.

EXAMINATION AND INVESTIGATIONS:

General Physical Examination:

Febrile

Pallor: present

B/L cervical (posterior and anterior triangle), axillary, inguinal, and epitrochlear non-tender lymphadenopathy.

Lymph nodes were soft and non-matted.



Systemic Examination:

ABD: No organomegaly

Normal bowel sounds.

CVS: S₁S₂ heard. No murmurs

RS: B/L NVBS. No added sounds.

Blood Investigations: Hb: 10.8 g% PCV: 32.4% MCHC: 33.3 g/dL TLC: 3510 cells/cumm Platelets: 1.5 lakh/cumm PBS: Microcytic hypochromic anaemia with leukopenia.

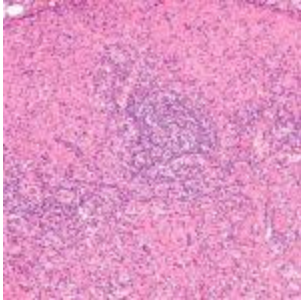
Malaria Card Test: Immunochromatographic test – negative

Tuberculosis Test: Acid fast bacilli (AFB) staining negative.

Paul Bunnell Test: Negative for infectious mononucleosis

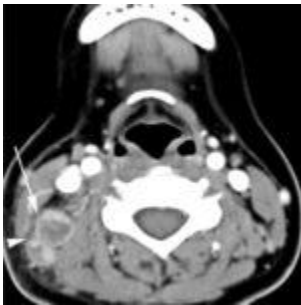
Fine Needle Aspiration Cytology Reactive lymphadenitis with histiocytes, tingible body macrophages.

Biopsy: Biopsy of scalene lymph node show no granulomas but areas of necrosis and signs of lymphadenitis



CT FINDINGS:

2 enlarged lymph nodes showing necrotic foci and some perinodal infiltration (arrow head)



FINAL DIAGNOSIS:

1. Kikuchi Fujimoto Disease
2. Tuberculosis
3. Infectious Mononucleosis
4. Systemic Lupus Erythematosus

Treatment Given:

TAB. PARACETAMOL 650 mg 1-1-1 PO

TAB DEXAMETHASONE 4 mg 1-1-1 PO TILL SYMPTOMS PERSIST, THEN
TAPERED OVER 1 WEEK

TAB PREDNISOLONE 20 mg 1-1-0 PO TILL SYMPTOMS PERSIST, THEN TAPERED OVER 1 WEEK

TAB PANTOPRAZOLE 40 mg 1-0-0 PO

INJ. PARACETAMOL 1gm i.v SOS

DISCUSSION:

Kikuchi Fujimoto is an extremely rare, sporadic and benign condition of unknown etiology which causes fever, necrotizing lymphadenopathy, rashes, headache and occasionally leukopenia^[1]. The symptoms are usually self-limiting. The condition was first described in 1972 in Japan as a new type of lymphadenitis with focal reticulum cell hyperplasia^[2]. This condition is almost always found in young women^[3]. Over a period of 15 years (1986-2000), only 330 cases were reported worldwide^[4], in contrast to the period 1996-2012 in which only 9 cases were reported^[5]. This makes this case a particularly rare diagnosis.

Kikuchi disease is often associated with Systemic Lupus Erythematosus and hence must be evaluated for the same using an antinuclear antibody profile.

Clinically, the disease is often confused for tuberculosis and sarcoidosis due to similar presentation.

Histologically, necrosis, karryorhectic debris, crescent shaped histiocytes and plasmacytoid monocytes can be seen. Treatment is mainly symptomatic and relies heavily on NSAIDs and corticosteroids. Death is unlikely and there is a 3% recurrence .

ACKNOWLEDGEMENTS: None

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