Oophoritis with Xanthogranulomatous Change – A Case Report.

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Short Communication

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ABSTRACT

Xanthogranulomatous oophoritis is an unusual chronic inflammation and clinically it form a mass-like lesion in the pelvic cavity and invades the surrounding tissue which mimic the tumour. Histologically, it is characterized by the presence of lipid filled macrophages with admixed lymphocytes, plasma cells and neutrophils. Granulomatous change occurs in other organs like gall bladder, kidney etc and etiology is unknown.

INTRODUCTION

Xanthogranulomatous inflammation of the female genital tract is an uncommon lesion, in the pelvic cavity that invades the surrounding tissues, which may mimic the tumour clinically and by imaging [1]. It is destructive to normal tissue of affected organs [2].

Xanthogranulomatous inflammation has been reported. Only a few cases involving the ovary have been reported to date in India. We describe a rare case of Xanthogranulomatous oophoritis.

CASE REPORT

A 65 yr old multiparous, postmenopausal female presented to Gynaecology OPD, of our institution with complaints of bloating, retching and pain in the lower abdomen for one year. She has off and on fever for few days. On clinical examination, patient had a firm, non tender, oval mass in right iliac, lumbar and umbilical area. Laboratory tests showed elevated ESR and white blood cell count. Ultrasound showed right adnexal mass. CT guided FNAC revealed purulent material, polymorphonuclear cells and foamy macrophages along with lymphocytes in background of necrotic debris.

Gross findings

A flap like structure measuring 7.5x6.5x2 cm in size (figure A). Outer surface was yellowish with small nodule.

Microscopic findings

Examined multiple pieces show ovarian tissue infiltrated by acute as well as chronic inflammatory infiltrate consisting of many foamy macrophages with abundant cytoplasm and hypochromatic central nuclei, polymorphonuclear cells, some lymphoid cells and
fibrosis (figure B&C). These foamy macrophages are scattered in various parts of lesion. The fibrosis is marked at places with hyalinisation. Many congested and thick walled blood vessels and hemorrhages are present (figure D).

**DISCUSSION**

Xanthogranulomatous oopheritis is often misdiagnosed by pathologists if they do not keep this entity in mind this may be due to the rarity of the condition. If the lesion is mainly focal scattered lymphocytes, it may be misdiagnosed as secondary lymphoma or leukemia. If the lymphocytes are scattering diffusely and foam cell are seldom, a diagnosis of malignant small cell tumour with stromal luteinzation may be rendered. If small amount of obvious fibrosis fibrosis and foam cells , a diagnosis of sclerosing stromal tumor may be made [3].

Correct diagnosis is made chiefly by histology, a suggestive preoperative diagnosis of xanthogranulomatous oopheritis could be lead to less radical surgery. The treatment of choice for Xanthogranulomatous oopheritis is oophrectomy. Patient with Xanthogranulomatous oopheritis should be monitor closely as it is associated with pelvic inflammatory diseases, endometriosis and intrauterine death [4,5,7].

The average age of patients of Xanthogranulomatous Oopheritis is 31 years. Although the pathogenesis of ovarian lesions is not fully understood, but proposed causes are infection, ineffective antibiotic therapy, abnormality in lipid metabolism, endometriosis or ineffective clearance of bacteria by phagocytes [8], rarely does a chronic ovarian abscess result in a solid tumor like mass, which is called either an ovarian Xanthogranuloma or Xanthogranulomatous oopheritis. This lesion occurs in patients with recurrent pelvic inflammatory disease. The involved ovary in each of the previously reported cases was replaced by a solid, yellow, lobulated mass that was well circumscribed and consisted of Xanthogranulomatous inflammation [6].

Hence, a diagnosis of Acute or Chronic Oophritis with Abundant Xanthogranulomatous Change was made. The case is being presented for its rarity.

**REFERENCES**