Primary sclerosing lipogranuloma: an unusual scrotal mass

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Sclerosing lipogranuloma (SLG) of the male external genitalia is a rare benign condition presenting as subcutaneous masses. The underlying pathological process is a granulomatous reaction of fatty tissue in this area. The cause of this is unclear but hypothesis stems around the pathogenesis of exogenous lipid degeneration from injection of foreign bodies such as paraffin for penile augmentation. However, endogenous lipid degeneration from other various causes such as infection, trauma, and allergic mechanisms has also been reported. We present the case of a 40-year-old man with primary SLG of the external genitalia. Literature review on the treatment strategies are addressed and discussed.

Keywords: sclerosing lipogranuloma; male genitalia; scrotal mass

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Case report

A 40-year-old man presented with bilateral scrotal, penile, and suprapubic swelling, which was intermittently tender and had progressively enlarged over 4-month period. He denied injecting foreign material into his external genitalia. There was no trauma to the groin and no symptoms of urinary tract infection. Past history revealed an episode of penile discharge 2 years ago which grew Klebsiella species treated with antibiotics. On examination, the penis and scrotum were swollen and tense. The swelling extended to the suprapubic region. There was bilateral inguinal lymphadenopathy. The testes were not palpable due to the scrotal oedema. There were multiple non-tender, irregular masses on the penile shaft dorsally without pus discharge per meatus (Fig. 1).

His white cell counts were 11.8 x 10⁹ and erythrocyte sedimentation rate (ESR) was 79 mm/h. Urine full examination and microscopic examination (FEME) was normal. Ultrasonography showed thickened scrotum due to oedema, minimal hydrocele with normal testes. Blood screen for sexually transmitted diseases were negative. Serology test for tuberculosis, fungal, cytomegalovirus, and filariasis from blood and tissue cytology were negative. Computer tomography of abdomen and pelvis showed bilateral scrotal enlargement with presence of a fat component within, extending superiorly to anterior abdominal wall (Fig. 2). Initial fine needle aspiration of enlarged lymph node yield a mixed population of lymphoid cells, predominantly of small lymphocytes, suggestive of a reactive process. As this diagnosis was inadequate, an excision of lymph node was performed.

Lymph node excision biopsy revealed lymphoid follicles of various sizes with prominent germinal centre and tangible body macrophages. There was
lipo-granulomatous reaction in the sinuses (Fig. 3). He was treated with broad spectrum antibiotics for 8 weeks. He had initially been on intravenous Ciprofloxacin 400 mg bd for 10 days, followed by oral Augmentin (Amoxicillin/Clavulanic Acid) 625 mg bd for 14 days. Later a combination of Doxycycline 100 mg daily and metronidazole 500 mg tds was given for 2 weeks in view of possible sexually transmitted microorganism involvement. Intravenous Ceftriaxone 1 gm daily was added for a total of 10 days. Following a bout of fever, intravenous Meropenem 1 gm tds was commenced for 1 week. Partial resolution of the genitalia and suprapubic swelling with complete resolution of pain was achieved. No surgical excision was required. The bilateral inguinal lymphadenopathy had shown partial resolution prior to discharge but later completely resolved on follow up one month later.

Discussion

Primary SLG of the male genitalia is rare and is believed to result from breakdown of endogenous lipids. This occurs following an allergic pathway induced by heat, extreme cold, infection, and trauma such as bicycle saddle injury and vigorous sexual intercourse. Some authors have reported presence of esonophilia in peripheral blood which supports the hypothesis of an allergic mechanism in these patients (3).

Patients experience painful erections and penile deformity and usually have tender subcutaneous nodules from penile base, along its shaft and in the scrotum. Primary lesions have also been reported in the testis, presenting as a firm mass and requiring a radical orchidectomy to rule out possibility of malignancy (4).

Our patient most likely has a primary cause of this disease as he denied injecting foreign material into the external genitalia. He additionally had a diffuse and tender suprapubic mass that extended to the margins of the inguinal ligament bilaterally that has never been reported thus far. This had made us look for filariasis as a possible cause as Wuchereria bancrofti is known to cause scrotal and suprapubic swellings and is common in Asian countries (5). Recurrent infection with Klebsiella species was ruled out as a possible etiology of primary SLG and bilateral lymphadenopathy based on the normal urine analysis results. This patient had a muscular build and it could be the strenuous muscle exercises resulting in primary lipid degeneration and lipogranuloma formation around his suprapubic and external genitalia region.

However as in most cases, it is almost impossible to pin point the exact cause of primary SLG. Some authors consider most primary cases are actually secondary because patients are not forthcoming about the use of foreign material injections (6). Does this matter? It probably does not as treatment strategies vary and not
influenced by its underlying cause. Treatment is generally divided into conservative and surgical approach. Surgical treatment includes partial or total excision of nodules with or without reconstructive skin flaps depending on extent of deformity. Partial excision may result in spontaneous regression or sometimes, quite the opposite, early recurrence which mimics a malignant disease process. This can cause unwanted anxiety among patients and treating surgeons. Péter et al. (7) advocated complete excision combined with penoplasty for best outcomes.

Conservative management varies from close observation, symptomatic treatment, antibiotic treatment to oral corticosteroids. Recent reports have shown good response to oral steroids over a 6-week period. Steroids also had resolved the associated eosinophilia seen in primary SLG (8). As there are very little data on the use of other drugs in its treatment, systemic steroids thus should be the first-line treatment for primary SLG while surgery is reserved for patients with refractory disease or with recurrences. Our patient benefited from a prolonged treatment of broad spectrum antibiotics as his main concern of pain over the suprapubic mass and scrotum was alleviated. However, he remains under close follow up to detect recurrences for which the oral steroids may be used.

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**References**


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