ACUTE APPENDICITIS IN AN INGUINAL HERNIA SAC: A VARIANT OF AMYAND'S HERNIA.

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ABSTRACT

Background: Amyand's hernia; a rare form of inguinal hernia is the presence of a vermiform appendix in inguinal hernia sac. It is often misdiagnosed as an incarcerated or strangulated hernia. The rarity of this condition is to be emphasized as well as review of literature.

Aim: This is to report a case of an inflamed appendix in an inguinal hernia sac.

Summary: A 29 year old male student, who was referred from a peripheral centre with right inguinoscrotal swelling of 14 years duration and severe persistent pain of four hours duration prior to presentation. He had repeated vomiting of recently ingested meal. There was constipation and no abdominal distension. The working diagnosis was dynamic intestinal obstruction from strangulated right inguinoscrotal hernia. He had emergency inguinoscrotal exploratory following resuscitation. The findings were: Inflamed vermiform appendix measuring about 15 cm in length and part of viable terminal ileum in the hernia sac. Appendicectomy and herniorrhaphy were done. His out-patient follow-up was been uneventful.

Conclusion: Acute appendicitis could have a variety of presentation, likewise inguinoscrotal hernia. In an extremely rare form, inflamed appendix could be a content of an inguinal hernia sac.

Key words: Hernia, Acute Appendicitis, Amyands.

INTRODUCTION:

Inguinal hernia is one of the most common operations done. About 1% of these operative findings contain appendix in a hernia sac. Amyand's hernia; a rare form of inguinal hernia is the presence of a vermiform appendix in inguinal hernia sac. It is often misdiagnosed as an incarcerated or strangulated hernia and delayed mesh repair is a better surgical option in perforated appendix cases.

CASE REPORT:

A 29 year old male student, who was referred from a peripheral centre with right inguinoscrotal swelling of 14 years duration and severe constant pain of four hours duration prior to presentation. He had repeated vomiting of recently ingested meal. There was constipation and no abdominal distension. Patient had similar occurence of pain 3 days prior to presentation but symptoms resolved spontaneously after hours of lying down. Swelling of the right groin started 14 years prior to presentation. Swelling was intermittent at first, spontaneously reducing, then sometimes with manual reduction by patient, associated with mild pain which relieves at rest. No history of chronic cough, drenching

night sweats, fever, weight loss, difficulty in passing urine or other urinary symptoms. There is a positive history of lifting heavy objects. No history of swelling on any other part of the body. At onset of symptoms he presented to a medical centre from where he was reffered to University of Port Harcourt Teaching Hospital (UPTH). Past medical and surgical history were not contributory. No history of drug allergy. Second in a family of 8 children. 3 boys and 5 girls in a monogamous setting. Takes alcohol(about 2 bottles of star a day, but does not take any tobacco products).

Clinical examination revealed he was a youngman in mild painful distress, drowsy, not pale, afebrile, anicteric, acyanosed, not dehydrated, no pedal oedema. Pulserate was 78 beats per minute, blood pressurewas 130/70 mmHg, only first and second heart sounds were heard, respiratory rate was 24 cycles/minute, and breath sounds were vesicular. Abdomen was Flat, soft, moved with respiration, the liver and spleen were not palpable and kidneys was non ballotable. Bowel sounds werehyperactive. Right groin swelling (huge) was tender, soft, not fluctuant. The Mass is separate from testes andone could not get above it, nor could it be reduced.

Rectal examination revealed normal findings.

Diagnosis of Obstucted right inguinoscrotal hernia possible strangulation was made. He was placed onnil per os, intravenous fluids, antibiotics and analgesics. The investigation and results were as follows: Packed cell volume was 36%, serology wasnegative for HIV I and II, Na was 137 mmol/L, K was 3.4 mmol/L, HCO₃was 24mmol/L, Urea was 1.9mmol/L and Creatinine was1.5mg/dl.Urinalysis showed glucose was 2+.

He had emergency inguinoscrotal exploration following resuscitation. The findings were: serosanguinous exudate, inflamed vermiform appendix(acute appendicitis complicating hernia) measuring about 15 cm in length and part of viable terminal ileum in the hernia sac(Fig 1-5). Appendicectomy and herniorrhaphy were done. Posterior wall repair was done using nylon darn. Histology result showed inflamed appendix. His out-patient follow-up was uneventful.

DISCUSSION:

Amyand's hernia is difficult to diagnose pre-operatively. The management is also not straightforward in most cases. This condition has been reported by some authors. Bhide in United Kingdom reported a case of Amyand's hernia in a 2 years old boy who presented with septic shock. Logan et al in Columbia, United Sates of America (USA), reported a case of incarcerated and perforated appendix within an inguinal hernia. They also recommended that repair should be undertaken without the use of synthetic mesh through a preperitoneal incision that gives access to the inguinal regions. In our case, the repair was done using nylon darn through a groin crease incision. Kaymakci et al in Instanbul, Turkey, reported a retrospective analysis of 30 patients aged between 19 days and 8 years with an Amyand's hernia operated in their institution from 1998 to 2009. In their study, hernia repair without an appendicetomy was



Fig 1 Inflamed appendix in an inguinal hernia sac.

performed in patients with normal appendix. Also, emergency appendicectomy through herniotomy was performed in cases of inflamed and perforated appendices.

Cankorkmaz et al reported also in a retrospective analysis study in Sivas Turkey, between 1998 and 2006, showed the management of 564 pateints with acute appendicitis, 1090 patients with inguinal hernia, 33 patients with incarcerated inguinal hernia and 12 patients with Amyand's hernia in paediatric surgery service. They found that all patients with Amyand's hernia were boys with a median age of 40 days. Findings intraoperatively included 2 normal appendices, 6 inflamed appendices, and 4 appendices with external signs of serosal inflammation of uncertain significance in the inguinal hernia sac. Two patients with a normal appendix had hernia repair without an appendicectomy.

The other 10 patients with an abnormal appendix underwent an emergency open appendicectomy with repair of the inguinal hernia. They also found none of the patients developed recurrence within a median postoperative follow-up of 2.5 years. In a retrospective study done by Sharma et al in Merseyside, UK, out of 18 consecutive patients with Amyand's hernia from 1991 to 2005, 17 were men only 1 was a woman. Their median age was 42 years. None was diagnosed preoperatively.

They concluded that inflammatory status of the appendix determines the type of hernia repair and the surgical approach. Incidental appendicectomy is not favoured. Park et al in New York, USA reported incarcerated Amyand's hernia in a premature infant associated with circumcision. Zaharie et al in Romania, reported perforated appendix with peri-appendicular abscess in an inguinal sac in an 81 year old patient. Bassini' procdure was the method for repair of the hernia.



Fig 2 Inflamed appendix in an inguinal hernia sac.

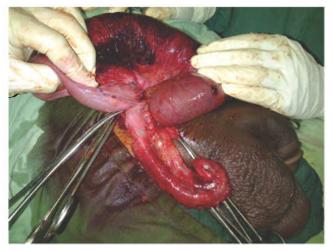


Fig 3.Inflamed appendix in an inguinal hernia sac.



Fig 5 Inflamed appendix in an inguinal hernia sac.

CONCLUSION:

Acute appendicitis could have a variety of presentation, likewise inguinoscrotal hernia. In an extremely rare form, inflamed appendix could be a content of an inguinal hernia sac as seen in index case.

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Fig 4. Inflamed appendix in an inguinal hernia sac

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