Dysphagia Lusoria presenting with Pill-induced Oesophagitis - A case report with review of literature

Malhotra A, Kottam RD, Spira RS

Abstract
Extrinsic oesophageal compression from vascular structures usually remains asymptomatic. Occurrence of dysphagia is an uncommon problem. We present a rare case of dysphagia lusoria from right sided aortic arch presenting as pill-induced oesophagitis caused by a testosterone pill. A thorough literature review on both the conditions is then presented to address pathogenesis, diagnosis, and management.

Keywords
Dysphagia Lusoria, Pill-induced oesophagitis, drug-induced oesophagitis

Introduction
Oesophageal compression by a vascular structure resulting in dysphagia is uncommon. There have been multiple reports in the medical literature of almost every major vascular structure in the chest causing some degree of oesophageal compression and subsequent dysphagia[1, 2]. In 1794, David Bayford, described a case of a 62 year-old lady having dysphagia due to an aberrant right subclavian artery. He coined the term “dysphagia lusus naturae”, which means "Freak of nature"[3].

Pill-induced esophagitis is associated with the ingestion of many pills and presents an uncommon cause of erosive oesophagitis[4]. Multiple different classes of drugs have been described to be hazardous to the oesophageal mucosa and cause pill-induced oesophagitis[5, 6]. Although uncommon in itself, dysphagia lusoria has not been described in literature to present as pill-induced oesophagitis. We present the first case of dysphagia lusoria causing pill-induced oesophagitis by testosterone pills in a young healthy man. Pubmed review of the English medical literature has been conducted to discuss the epidemiology, pathogenesis and management of this uncommon disorder.

Case Presentation:
A 26 year-old man with no significant past medical history presented with 5 days of dysphagia (for both solids and liquids), odynophagia and retrosternal chest discomfort. He admitted to having occasional difficulty swallowing for past 2-3 months for solids only. He denied any heartburn, cough, regurgitation, loss of appetite, weight loss, fever, chills, haematemesis or melaena. He denied tobacco or alcohol use. 2 weeks prior to presentation he had started taking testosterone pills for bodybuilding. Physical examination was completely unremarkable.

A barium oesophagram showed extrinsic oblique compression of the oesophagus at the level of the carina as it passes through the aorta. CT scan and MRI of the chest revealed a right-sided aortic arch crossing posteriorly to the oesophagus with proximal oesophageal dilatation consistent with Dysphagia Lusoria. Endoscopy noted erosive oesophagitis/distinct ulceration extending from 18cm into a pulsatile area of narrowing at 20 cm with normal mucosa visualized distally.

Biopsies revealed oesophageal squamous mucosa with marked acute inflammation, reactive changes and no evidence of viral inclusions. Surgical management was discussed with the patient, but given the short duration of symptoms and the patient’s stable weight, providing symptomatic relief with lifestyle changes, together with a trial of medications such as proton pump inhibitors were considered. At 2 weeks follow-up whilst taking proton pump inhibitors and having discontinued the testosterone pills, the patient experienced complete resolution of symptoms.

Epidemiology:
Dysphagia Lusoria: Moltz et al, found that lusonian artery has a prevalence of about 0.7% in the general population, based on the post-mortem findings[7]. Also, out of 1629 patients who underwent endoscopy for various reasons, 0.4% had a finding of a lusorian artery in a report from Fockens et al[8]. It has also been concluded based on the autopsy results and retrospective analysis of patients’ symptoms during life that about 60-70% of these patients remain asymptomatic[7]. Coughing, dysphagia, thoracic pain, syncope and Horner’s syndrome may develop, but usually present in old-age[9].

Pill-Induced Oesophagitis: The data on pill-induced oesophagitis is rather limited. A Swedish study found an incidence of 4 cases per 100,000 population/year[10]. Wright found the incidence of drug-induced oesophageal injury to be 3.9/100,000[11]. This may be underestimated and does not include subclinical or...
misdiagnosed cases. Also, cases are reported selectively, due to clustering of cases, newly implicated pills or rare complications. The incidence today is probably much higher due to increased use of prescription medications, widespread use of endoscopy and an ageing population. All these factors limit our ability to correctly assess the true epidemiology of this iatrogenic disorder.

Pathogenesis

**Dysphagia Lusoria:** During embryological development, the aortic sac gives rise to six aortic arches and with further development the arterial pattern is modified and the fourth arch persists on both sides and some vessels regress. In right arch anomaly, the left arch atrophies and disappears whereas the right arch persists. If both arches persist, they form a double arch or a vascular ring encircling the trachea and oesophagus.

**Pill-induced oesophagitis:** Several lines of evidence confirm that oesophageal mucosal injury is caused by prolonged contact with the drug contents. On clinical grounds, patients frequently report a sensation of a pill stuck in the oesophagus before the development of symptoms and the frequent occurrence of symptoms after improper pill ingestion. Endoscopically, the evidence includes occasional observation of pill fragments at the site of injury, sharp demarcation of the injury site from the normal tissue and the frequent localization of the injury to the areas of oesophageal hypomotility or anatomic narrowing. Therefore factors predisposing to the drug-induced oesophageal injury can be divided into two main categories: patient or oesophageal factors and drug or pharmaceutical factors as shown in tables 1 and 2. It is important to note that most patients who experience pill-induced damage have no antecedent oesophageal disorder, neither obstructive nor neuromuscular. It is the combination of anatomic narrowing coupled with the caustic effects of the implicated drug that caused the oesophageal injury in our case. Although testosterone pills have never been reported to cause pill-induced oesophagitis, 6 cases of corticosteroid-induced oesophagitis have been described in the literature.

### Table 1: Patient/Esophageal factors for pill-induced esophagitis

<table>
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<tr>
<th>Factor</th>
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<tr>
<td>Old Age</td>
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<td>Decreased Salivation</td>
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<td>Pill intake in recumbent position</td>
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<tr>
<td>Lack of adequate fluid intake with the drug</td>
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<td>Structural abnormalities of esophagus</td>
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<td>Hypomobility of the esophagus</td>
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### Table 2: Drug related factors for pill-induced esophagitis

<table>
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<th>Factor</th>
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<tr>
<td>Chemical structure (sustained release pills, gelatinous surface)</td>
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<tr>
<td>Formal structure (capsule increases risk over the tablet)</td>
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<tr>
<td>Solubility</td>
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<td>Simultaneous administration of multiple medications</td>
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Clinical Presentation

**Dysphagia Lusoria:** As previously mentioned the disorder remains asymptomatic in majority of the patients. Symptomatic adults usually present with dysphagia for solids, (91%), chest pain (20% or less). Less commonly, patients may have cough, thoracic pain or Horner’s syndrome. In infants, respiratory symptoms are the most predominant mode of clinical presentation. This is believed to be due to absence of tracheal rigidity, allowing for its compression with resulting stridor, wheezing, cyanosis etc.

**Pill-induced oesophagitis:** Patients with pill-induced injury usually present with odynophagia, dysphagia and/or retrosternal chest pain. Symptoms can occur after several days after starting a drug, but frequently occur after the first dose. Fever and haematemesis signifying a possible mediastinal extension can occur without chest pain. Pharyngitis due to the pill lodged in the hypopharynx has been reported.

Our case presents a typical example of asymptomatic Dysphagia Lusoria, who developed acute dysphagia, odynophagia and retrosternal chest discomfort immediately after the initiation of the offending agent; which is very typical of pill-induced oesophageal injury.

Diagnostic approach

**Dysphagia Lusoria:** The best method to diagnose an aberrant right subclavian artery presenting with difficulty swallowing is initially with a barium oesophagram followed by a CT or MRI scan. Angiography although considered gold standard for the diagnosis of vascular abnormalities is now largely supplanted by newer less invasive techniques such as CT or MR angiography. Upper endoscopy may reveal a pulsating compression of the posterior wall of the oesophagus as in our case.

Figure 1 - Barium esophagram depicting the extrinsic indentation of the esophagus as it crosses the aorta.
Endoscopic ultrasound, especially with Doppler technology may be helpful to confirm the vascular nature of the abnormality\(^{(27)}\). Oesophageal manometry usually shows non-specific findings. High peristaltic pressures have been reported in the proximal oesophagus above the level of the compression \(^{(9,35)}\).

*Pill-induced oesophagitis*: Barium studies can be normal, and slowing of barium column may be the only abnormality seen\(^{(31)}\). Double contrast studies may however, increase the yield of a positive result\(^{(36)}\). Kikendall et al. reported that endoscopy revealed the evidence of injury in all the patients \(^{(5)}\). Endoscopy most commonly reveals one or more discrete well demarcated ulcers with normal surrounding mucosa. Ulcers may range from pin-point to several centimetres in diameter\(^{(5)}\). Biopsies, if performed, help to distinguish the condition from infection and neoplasia.

Our case shows a distinct oblique compression in the posterior wall of oesophagus on the barium study (fig 1) and classic findings on MR/CT with contrast which also excluded any other thoracic vascular abnormalities (fig 2-5). Endoscopic images of a shallow ulcer are shown in fig 6, 7.
Treatment

**Dysphagia Lusoria:** The treatment of patients with DL primarily depends upon the severity of symptoms. Mild to moderate cases are managed by lifestyle and dietary changes such as eating slower, chewing well, sipping liquids, weight reduction and reassurance as in our case.\(^{(9)}\) Jansen et al also reported in a series of 6 patients that 3/6 improved with proton-pump inhibitor alone or in combination with the prokinetic drug cisapride \(^{(9)}\). Severe symptoms and failure of medical therapy may need surgical evaluation and treatment. Richter et al. reported 14/24 patients who underwent surgical repair of the aberrant vessel for DL \(^{(27)}\). Bogliolo et al proposed endoscopic dilation as a temporary alternative to relieve symptoms in patients who are poor surgical candidates \(^{(9)}\).

**Pill-induced oesophagitis:** Most uncomplicated cases of pill-induced oesophagitis may heal spontaneously, with resolution of symptoms in a few days to a few weeks. Withdrawal of the offending drug and avoidance of topicaly irritating foods such as citrus fruits, alcohol is imperative to aid healing.\(^{(29)}\). Sucralfate, topical anaesthetics, and acid suppression are often used to aid in relief of pain \(^{(14, 15)}\). Rarely, in severe cases, parenteral nutrition or endoscopic dilation of chronic strictures may be required. \(^{(15, 25)}\)

**Conclusion**

Our case demonstrates a typical case of DL presenting with pill-induced oesophagitis who responded to conservative and acid suppressive therapy. Identifying the risk factors and adequate patient education is the key to prevention.

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