Pneumoscrotum after ERCP-related duodenal perforation

The incidence of duodenal perforation after endoscopic retrograde cholangiopancreatography (ERCP) is lower than 1%. When it does occur, intestinal gas usually tracks upward through the retroperitoneal space and mediastinum to present as surgical emphysema in the neck [1].

We describe a rare case of a young man developing pneumoscrotum following an ERCP-related duodenal perforation. A 29-year-old man underwent ERCP for investigation and treatment of jaundice due to choledocholithiasis. The papilla was difficult to cannulate. Endoscopic sphincterotomy was performed. Successful biliary drainage was achieved. As stone clearance was incomplete, a temporary plastic stent was placed [2]. The patient immediately complained of a swollen scrotum; otherwise he was completely asymptomatic. There was no abdominal tenderness or guarding. His scrotum was enlarged 2–3 times the normal size (Fig. 1), and was tense and tympanitic. Crepitus was elicited in the scrotum, but not in the abdomen, chest, or neck. Needle aspiration of the scrotum produced gas but no fluid. Abdominal computed tomography scan (Fig. 2) confirmed the diagnosis of pneumoscrotum related to retroperitoneal perforation. Type IV perforation was identified and was treated successfully with conservative management.

Few cases of pneumoscrotum have been reported. A systematic review by Cochetti et al. [3] identified 59 cases of pneumoscrotum. Although the endoscopic procedures accounted for 32% of all iatrogenic cases, only three cases of pneumoscrotum after ERCP have been described previously [3–5]. In cases of pneumoscrotum, the gas could reach the scrotum by tracking along the transversalis fascia, which forms the innermost covering layer of the spermatic cord. This would account for the lack of palpable subcutaneous emphysema over the abdominal wall. Consequently, pneumoscrotum could be the first manifestation of a duodenal perforation, as was the case in our patient. As duodenal perforation is a potentially life-threatening condition, clinicians involved in ERCP management must be aware of this clinical manifestation in order to establish prompt diagnosis and treatment.

Competing interests: None

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References

Bibliography
DOI http://dx.doi.org/10.1055/s-0042-116428
Endoscopy 2016; 48: E295
© Georg Thieme Verlag KG Stuttgart · New York ISSN 0013-726X

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