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Case Report

The new onset of dysphagia four years after anterior cervical discectomy and fusion: Case report and literature review

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INTRODUCTION

Anterior cervical discectomy and fusions (ACDF) complications, include dysphagia (most common), followed by recurrent laryngeal nerve injury, Horner syndrome, esophageal perforation, infection, and graft extrusion. The prevalence of dysphagia 1–6 weeks postoperatively ranges from 28% to 57%. Although the majority of symptoms resolve spontaneously, some complaints may last up to two postoperative years. Here, we present the new-onset dysphagia in a 49-year-old male, 4 years following a 2-level C3-C4/C5-C6 ACDF. Symptoms were attributed to a left C3 screw extrusion into the posterior pharyngeal soft tissue with the loosening of the right C3 screw. The patient required resection of instrumentation followed by a posterior C2T2 fusion. The point of this study was to highlight that the new-onset dysphagia occurring even years after an ACDF should be carefully
evaluated for instrumentation failure (e.g., perforation of the pharyngeal wall).

**CASE REPORT**

At age 45, the patient with a history of renal insufficiency, had a C3-4 and C5-6 ACDF. Four years later, he prior presented with neck/right upper extremity pain, and worsening dysphagia. The physical exam revealed 3/5 deltoid strength on the right side, but was otherwise unremarkable.

**Diagnostic studies**

Lateral radiographs of the cervical spine revealed an extruded left C3 screw, which had migrated into the prevertebral soft tissues at the C4-5 level, as well as loosening of the right C3 screw [Figure 1]. A barium swallow confirmed screw migration superiorly with swallowing, indicating it was embedded in the posterior soft tissue of the pharyngeal wall [Video 1].

**Surgery**

With the help of an otolaryngologist ear, nose, and throat specialist, the left C3 screw was removed from within the left lateral pharyngeal wall; scar tissue on the posterior aspect of the screw pocket was opened, and the inflammatory capsule surrounding it was excised without evidence of esophageal perforation. Next, the spine surgeons removed the anterior instrumentation (e.g., plate C3-C4/C5-C6), followed by placement of a new interbody structural allograft at C4-5 with a new plate, and posteriorly, and instrumented fusion from C2-T2. The patient was discharged without complaints.

**DISCUSSION**

Following anterior-approach cervical operations, many patients experience some degree of dysphagia, which is almost always self-limiting and may take up to 2 years to resolve.\(^4\)\(^6\)\(^7\) It usually appears 2–7 days after surgery but may begin up to a month after surgery in 8–25% of patients; estimates of the incidence of dysphagia in the literature range from 3.6% to 83%.\(^3\)\(^4\)\(^8\)

To assess dysphagia, dynamic imaging studies include modified barium swallow study and fiberoptic endoscopic evaluation of swallowing.

Delayed-onset dysphagia may be variously attributed to: mass effect exerted on the posterior laryngeal wall by an abscess, delayed instrumentation failure with pharyngeal versus esophageal defect.\(^2\) Fryer et al. recently reported a case of hardware failure post-ACDF associated with dysphagia;\(^3\) our case had clear ventral screw migration.

**Radiographic recommendations to evaluate delayed onset dysphagia**

We recommend preoperative dynamic imaging and intraoperative fluoroscopy for all cases of delayed postoperative dysphagia, especially those with potential for or known screw migration/instrumentation failure. A multidisciplinary approach with an otolaryngologist may be valuable to ensure safe instrumentation mobilization and removal, as well as to prepare for a possible esophageal repair.\(^2\)

**CONCLUSION**

Anterior cervical instrumentation failure should be considered when patients who previously underwent an ACDF newly present with dysphagia months to years later; the preoperative workup should include X-rays, magnetic resonance imaging, computed tomography, and videofluoroscopic swallowing studies.

**Declaration of patient consent**

The authors certify that they have obtained all appropriate patient consent.

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**Conflicts of interest**

There are no conflicts of interest.

**REFERENCES**


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