

CASE REPORT

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# Thyroid abscess presenting as a discharging sinus: a case report

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## Abstract

**Background** It is unusual to get an isolated thyroid abscess. Thyroid abscess or acute suppurative thyroiditis accounts for 0.7–1% of all instances of thyroid diseases. The thyroid gland typically has a high level of iodine, a well-enveloped capsule, and resistance to infections. The diagnosis and treatment of this unusual entity are covered in this paper.

Case presentation.

An adult male patient presented with anterior neck swelling for 15 days and pus discharge for 2 days. The neck ultrasound revealed characteristics that could indicate a thyroid abscess. The thyroid function test and other laboratory data were within normal bounds. A neck contrast-enhanced computed tomography scan was performed, and it only revealed a single thyroid abscess and no other abnormalities. After starting the patient on intravenous antibiotics, the abscess was incised and drained. The patient's symptoms got better.

**Conclusion** In the era of advanced antibiotics incidence of thyroid abscess is extremely rare. Early diagnosis and intervention may improve the outcome of this condition. Knowledge of disease and its potential complications can prevent morbidity.

**Keywords** Thyroid abscess, Acute suppurative thyroiditis, Sinus

## Background

Due to its capsule, healthy circulatory and lymphatic circulation, and iodine content, the thyroid gland is the least likely gland to become infected. Acute suppurative thyroiditis is a life-threatening endocrine emergency that presents as fever, neck pain, and swelling. Even among those with impaired immune systems, acute suppurative thyroiditis (AST) is a rare disease with the incidence being less than 1% of all thyroid diseases [1]. It is most commonly caused by gram-positive organisms like *Staphylococcus aureus*, haemolytic *Streptococcus*, or

*Streptococcus pneumoniae*, but occasionally it is caused by *Haemophilus* and *Fusobacterium*. Infection caused by *E. coli* in the thyroid has never been reported in the literature. Infection is usually a result of trauma, hematologic seeding from a distant infected site, or direct infection from a deep cervical region. The infection starts in one thyroid lobe, which if left untreated turns into an abscess cavity that breaks through the capsule and extends into the mediastinum or deep neck spaces via fascial planes. On workup usually the patient is euthyroid with normal Erythrocyte sedimentation rate and raised leukocyte count. Diagnosis is usually clinical and management is incision and drainage followed by culture-directed antibiotics.

## Case presentation

A 65-year-old male with no co-morbidities presented to the outpatient department with complaints of anterior neck swelling for 15 days and pus discharge from

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the swelling for 2 days. There was no history of any injectable drug intake. On examination, the patient was afebrile and normotensive with no sign of respiratory distress. Systemic examination was normal. There was no evidence of an immunocompromised state. On head and neck examination a 6×5 cm midline neck swelling was present just above the suprasternal notch with multiple pus points. The swelling was tender with a raised temperature.

Thyroid profile was normal with serum thyroid stimulating hormone (TSH) of 1.46 IU/ml. Viral markers were negative. Fine needle aspiration (FNA) cytology was suggestive of an infected colloid cyst, ruling out the possibility of anaplastic carcinoma. Pus aspirated was sent for culture. Figure 1 shows the contrast-enhanced computed tomography (CECT) neck, which was suggestive of a well-defined lobulated complex hypodense lesion of size 63×60 mm with internal hyperdensities suggestive of debris, cystic areas, and no calcification. The trachea was deviated to the right and retrosternal extension was noted. Routine investigations were sent. Figure 2 shows the chest X-ray with a homogenous soft-tissue density anterior to the trachea, displacing it to the right. The hemoglobin level on admission was 9.5 g/dl with a total leucocyte count (TLC) of 20,500/ $\mu$ L.

Since there was no evident immunocompromised status or other source of infection, such as a urinary tract infection, the underlying etiology of the infection remained unknown. Neither direct laryngoscopy nor a review of neck imaging revealed the presence of a pyri-form sinus fistula.



**Fig. 2** Chest X-ray showing the inferior extent of pus and tracheal shifting

The patient underwent incision and drainage under local anesthesia and 75 ml of yellowish-green pus was suctioned out (Fig. 3).

The patient was treated empirically with intravenous piperacillin and tazobactam at a dose of 4.5 g thrice a day and metronidazole at a dose of 500 mg three times a day. Cultures of the drained pus yielded *Escherichia coli*. After a week's course of intravenous antibiotics and daily dressing, the patient improved symptomatically and was discharged home 7 days after admission. The TLC on discharge came down to 6310/ $\mu$ L, with a chest X-ray



**Fig. 1** Large collection within the thyroid gland, shifting the trachea



**Fig. 3** Thyroid abscess after drainage



**Fig. 4** No residual swelling on 1 month follow-up

showing the return of the trachea to its normal position. On 1 month follow-up, the patient's wound healed completely with no residual swelling (Fig. 4).

## Discussion

Acute suppurative thyroiditis is an uncommon form of thyroiditis and its progression to thyroid abscess is rare. Thyroid disease, anatomical anomalies, or immunocompromised conditions may predispose patients to developing thyroid abscesses [2, 3]. The most common cause of thyroiditis in children is a pyriform fistula, while in adults it is immunocompromised status. Idiopathic thyroid abscess is a rare occurrence with few reports [4]. In our case the cause of the abscess is unknown.

The most common causative organism for thyroid abscess is *Staphylococcus aureus* and *Streptococci*. Other organisms, though less commonly identified are *Aspergillus*, *Brucella*, *Klebsiella*, *Eikenella*, *Salmonella*, and *Acinetobacter* but often the infection is polymicrobial. Rarely can methicillin-resistant *Staphylococcus aureus* and *E. coli*-related thyroid abscesses develop [5–7]. Thyroid gland infections caused by *Tuberculosis*, *Salmonella*, and *Eikenella* have also been documented [8, 9]. Upon reviewing the literature, a few cases of acute suppurative thyroiditis caused by *E. coli* in immunocompromised patients were identified [10, 11].

A thyroid abscess following fine needle aspiration of the thyroid nodule has been reported [4]. The hematogenous spread of infection from urinary tract infection has also been reported [12]. In this case, no source of infection was found.

Acute suppurative thyroiditis is a rare entity, which progresses rapidly and can be fatal if not identified and treated promptly. For an immunocompromised patient or a child presenting with an acute onset anterior neck swelling with fever with an underlying thyroid disease the possibility of acute suppurative thyroiditis should be considered. Ultrasound is the preferred imaging technique for the diagnosis of thyroid diseases as well as it aids in needle aspiration. The use of computed tomography and magnetic resonance imaging in diagnosis has a limited role. If an iodine scan is done, abscess areas may appear cold [13].

Treatment of thyroid abscess is incision and drainage followed by culture-directed antibiotics. In the case of pyriform sinus fistulas, chemo cauterization can be done [14].

Thyroid abscess secondary to branchial cleft anomalies may require lobectomy [4, 15]. Recurrent and non-responsive cases may require thyroidectomy, though there is an increased risk of parathyroid and recurrent laryngeal nerve injury.

Thyroid abscesses presenting with complications, including mediastinitis, stridor, internal jugular vein thrombosis, sepsis, and death have been reported [4, 16, 17].

## Conclusion

The potential morbidity of thyroid abscesses emphasizes the necessity of timely detection and treatment. Although thyroid abscess is a rare diagnosis, it can have serious side effects. Therefore, in order to implement early aggressive care, practitioners must be informed of the diagnosis.

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## Authors' contributions

VS and AS prepared the manuscript and were involved in patient management. AJ did the editing and was involved in patient management, and AKG was involved in patient management and preparing the final draft for submission. All authors read and approved the final manuscript.

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## Availability of data and materials

This is a case report, and all the data pertaining to the case report have been included in this published article.

## Declarations

### Ethics approval and consent to participate

Approval from the Institute Ethics Committee was sought prior to reporting the case. IEC approval no- 2023-19-0th-exp-5-CR dated 19-9-23. The article was prepared in accordance with the Declaration of Helsinki. Written informed consent was obtained prior to the publication of the case report from the patient.

### Consent for publication

Written informed consent for publication of the patient's clinical details and clinical images was obtained from the patient.

### Competing interests

The authors declare that they have no competing interests.

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