Advances in Clinical Medical Research and Healthcare Delivery

Volume 2 | Issue 2

Article 5

2022

A case report and literature review of an intrathyroid epidermoid cyst

Satvik R. Hadigal *Rochester Regional Health*, satvikhadigal@gmail.com

Bradley Frate Rochester Regional Health, brad.Frate@rochesterregional.org

Samea Lone *Rochester Regional Health*, snlone@gmail.com

Frank Salamone Rochester Otolaryngology Group, fns99@yahoo.com

Cheris Sachica Rochester Regional Health, Sachica.Cheris@rochesterregional.org

Raj Pyne Rochester Regional Health, Raj.Pyne@rochesterregional.org

Follow this and additional works at: https://scholar.rochesterregional.org/advances



This work is licensed under a Creative Commons Attribution-NonCommercial 4.0 International License

Recommended Citation

Hadigal SR, Frate B, Lone S, Salamone F, Sachica C, Pyne R. A case report and literature review of an intrathyroid epidermoid cyst. *Advances in Clinical Medical Research and Healthcare Delivery*. 2022; 2(2). doi: 10.53785/2769-2779.1092.

ISSN: 2769-2779

This Case Report is brought to you for free and open access by RocScholar. It has been accepted for inclusion in Advances in Clinical Medical Research and Healthcare Delivery by an authorized editor of RocScholar. For more information, please contact Advances@rochesterregional.org.

A case report and literature review of an intrathyroid epidermoid cyst

Abstract

Epidermoid cysts are the most common cutaneous cysts, and infrequently develop within mucosal or glandular tissue. Rarely, epidermoid cysts have been described as arising within the thyroid gland in the form of a nodule. In this paper, we describe a case of a 72-year-old female with a suspicious-appearing thyroid nodule on ultrasound, which was eventually found to be an epidermoid cyst. To our knowledge, this is the first surgical biopsy-proven case that demonstrates an intrathyroid epidermoid cyst in multiple radiographic modalities including ultrasound, nuclear medicine I-123 thyroid scan, and MRI. In addition, we present a concise review of previously described cases of intrathyroid epidermoid cysts, including patient clinical manifestations, initial sonographic appearance, and management of lesions. This case reinforces the importance of considering rare diagnoses in the workup of a thyroid nodule based on its sonographic appearance.

Keywords

epidermoid cyst, epidermoid inclusion cyst, epidermal cyst, intrathyroid epidermoid cyst

Creative Commons License



This work is licensed under a Creative Commons Attribution-NonCommercial 4.0 International License

Conflict of Interest Statement

Authors have no conflict of interest

CASE REPORT

A Case Report and Literature Review of an Intrathyroid Epidermoid Cyst

Satvik R. Hadigal*, Bradley Frate, Samea Lone, Frank Salamone, Cheris Sachica, Raj Pyne

Rochester Regional Health, USA

Abstract

Epidermoid cysts are the most common cutaneous cysts, and infrequently develop within mucosal or glandular tissue. Rarely, epidermoid cysts have been described as arising within the thyroid gland in the form of a nodule. In this paper, we describe a case of a 72-year-old female with a suspicious-appearing thyroid nodule on ultrasound, which was eventually found to be an epidermoid cyst. To our knowledge, this is the first surgical biopsy-proven case that demonstrates an intrathyroid epidermoid cyst in multiple radiographic modalities including ultrasound, nuclear medicine I-123 thyroid scan, and MRI. In addition, we present a concise review of previously described cases of intrathyroid epidermoid cysts, including patient clinical manifestations, initial sonographic appearance, and management of lesions. This case reinforces the importance of considering rare diagnoses in the workup of a thyroid nodule based on its sonographic appearance.

Keywords: Epidermoid cyst, Epidermoid inclusion cyst, Epidermal cyst, Intrathyroid epidermoid cyst

1. Introduction

pidermoid cysts, also known as epidermal cysts, epidermal inclusion cysts, infundibular cysts, and keratin cysts, are the most common benign lesions of the skin, ovaries, and testicles. They represent approximately 85%-90% of all excised cysts. Epidermoid cysts are derived from ectoderm and consist of an epithelial-lined wall that may be partly keratinized; this contrasts with dermoid cysts, which contain mesodermal tissue. Epidermoid cysts seldom occur in mucosal or glandular tissue, especially in the thyroid gland. These can present as solid thyroid nodules on ultrasound (US) and can be misinterpreted as aggressive nodules. In this case, a benign epidermoid cyst initially presented as a highly suspicious nodule classified as TI-RADS 5 (based on the White Paper of ACR TI-RADS committee), underwent pathological analysis and was subsequently surgically removed.¹

2. Case presentation

A 72-year-old female presents with complaint of a palpable midline neck lesion. The patient was found to have subclinical hyperthyroidism with a low TSH of 0.26 (normal range: 0.55-4.78) and a normal T3 of 133 (normal range: 60-181) and normal free T4 of 1.3 (normal range: 0.9-1.8). The patient denied any symptoms of hyper or hypothyroidism.

On ultrasound, the right thyroid lobe measured $4.3 \times 2.2 \times 1.7$ cm and the left thyroid lobe measured $4.1 \times 1.6 \times 1.4$ cm. A well-circumscribed, unilocular, avascular, homogeneous, hypoechoic mass measuring $3.0 \times 2.1 \times 1.3$ cm was found in the right mid-inferior thyroid lobe extending into the isthmus (Fig. 1a and b). Within this mass, there were scattered hyperechoic foci, some of which exhibited posterior acoustic enhancement. These hyperechoic foci were similar in appearance to calcifications and are highly suspicious for cancer. These types of suspicious calcification typically exhibit posterior acoustic shadowing instead of enhancement. Based



* Corresponding author.

E-mail addresses: satvikhadigal@gmail.com (S.R. Hadigal), brad.Frate@rochesterregional.org (B. Frate), snlone@gmail.com (S. Lone), fns99@yahoo.com (F. Salamone), Sachica.Cheris@rochesterregional.org (C. Sachica).

Accepted 28 April 2022. Available online 31 May 2022



Fig. 1. (A) Sagittal view of right thyroid lobe. Hypoechoic lesion with multiple punctate hyperechogenic foci (white arrowheads) which exhibit posterior acoustic enhancement. (B) Transverse view of the right thyroid lobe with hyperechogenic foci (white arrowheads). Same lesion from Fig. 1A shown extending from the right lobe into the isthmus. (C) I-123 thyroid of the thyroid gland shows enlarged right thyroid lobe with mildly elevated 24-h uptake (26.5%; normal 8%–25%) of radioactive iodine. There is a photopenic region (white arrowhead) in the inferior right thyroid lobe on anterior (ANT), left anterior oblique (LAO) and right anterior oblique (RAO).

on its characteristics, the nodule was classified as TI-RADS 5 for solid composition (2 points), hypoechogenicity (2 points), and punctate echogenic foci (3 points).¹ For the sake of comprehensiveness, another smaller nodule that measured $1.0 \times 0.9 \times 0.6$ cm³ was identified incidentally in the left lobe, biopsied, and found to be a benign nodule.

On nuclear medicine I-123 thyroid scan, the right thyroid gland was enlarged with mildly elevated 24h radioactive iodine uptake. There was photopenia in the right inferior thyroid lobe in the region of the US nodule. The above radiographic findings, along with the size of the nodule, warranted an US-guided

https://scholar.rochesterregional.org/advances/vol2/iss2/5 DOI: 10.53785/2769-2779.1092 biopsy based on the suggested diagnostic and treatment approach for thyroid nodules and TI-RADS criteria (Fig. 1C).^{1,2} US-guided fine needle aspiration (FNA) biopsy was performed, which demonstrated nucleated and anucleated squamous cells with few interspersed keratinized squamous cells without evidence of colloid or other thyroid cellular elements (Fig. 2A and B). At this point the differential diagnosis was epidermoid cyst, dermoid cyst, and metastatic squamous cell carcinoma. A repeat FNA biopsy was performed to decrease the chances of missing any underlying and undersampled squamous cell carcinoma especially in older adults such as in this case. The results were unchanged when compared to the first FNA. Given the rarity of this nodule in the thyroid, MRI with contrast was performed to further characterize lesion and for surgical planning. On MRI, the nodule exhibited high T1 intensity, low to intermediate T2 intensity with no significant fat suppression or contrast enhancement (Fig. 3). High T1, suggests that cyst contents are mostly proteinaceous and is





Fig. 2. Fine needle aspiration biopsy of the intra-thyroid lesion. (A) Anucleated (black arrowheads) and nucleated (white arrowheads) squamous cells seen (10x cytospin prep PAP stain). (B) Keratinizing squamous cells (pink colored squamous cells) interspersed with anucleated and nucleated squamous cells (10x PAP stain).







D



likely due to keratin given the FNA results showed keratinized squamous cells in Fig. 3B.

The decision to perform a right hemithyroid ectomy was made based on the nodule's suspicious appearance and high TI-RADS score for thyroid cancer. On gross pathology after resection, the cyst contained thick, dark brown fluid with a smooth and tan-pink cyst lining. The surrounding parenchyma was rubbery and dark-red, suggestive of a fibrous layer (Fig. 4A). The final diagnosis of epidermoid inclusion cyst was made based on histology. The cyst contained abundant keratin and the cyst-wall lining was composed of stratified squamous epithelium with a granular layer which are characteristic features of an epidermoid cyst. No papillary areas, columnar cells, cuboid cells, cilia, glands, or hair follicles were found within the cyst wall (Fig. 4B and C). Initially, the possibility of an intrathyroid thyroglossal duct cyst was entertained. However, due to the absence of columnar or cuboid epithelium and glands in the cyst wall, as well as the presence of keratinizing squamous epithelium, this diagnosis was unlikely.

3. Discussion

The pathophysiology of intrathyroid epidermoid cysts is not well understood. Classified as congenital or acquired, many theories have been proposed regarding their pathophysiology in the neck. Congenital cysts can arise from reactivation of the ectodermic elements entrapped during the midline fusion of the first and second branchial arches. Acquired cysts are theorized to be derived from traumatic or iatrogenic inclusion of epithelial cells.³

Squamous cells and keratin are principal components of epidermoid cysts. Histologically, epidermoid cysts are unilocular, well circumscribed, and lined with mature superficial squamous cells with intact nuclei, anucleate squames, and clusters of neutrophils, lymphocytes, and macrophages in a background of amorphous debris. Keratin debris is usually found within the cyst.

A brief review of 18 cases from literature search along with this case report is presented in Table 1. Based on a review of these 19 case studies describing intrathyroid epidermoid cysts, the typical age of presentation ranged from 4 to 78 years of age

Fig. 3. MRI of the neck. (A) Axial T1 sequence shows a high T1 intensity nodule in the right thyroid lobe (white arrows), (B) Axial T1 fat saturation sequence shows no significant fat suppression of the contents (white arrows), (C) Axial T1 fat saturation sequence with intravenous ProHance contrast shows no significant enhancement, (D) Axial T2 sequence shows a low to intermediate signal.

Α





Fig. 4. (A) Right hemithyroidectomy gross pathology showing right lobe of thyroid measuring $5.0 \times 4.0 \times 2.5$ cm with a nodule (*) measuring $3.5 \times 3.0 \times 1.5$ cm. (B) Histology of the surgical specimen shows the lining of the intrathyroid cyst with stratified squamous epithelium with granular layer; note the surrounding thyroid follicles and the lack of columnar cells, cuboidal cells, sebaceous glands, or eccrine glands within the wall; the cyst (Cy) contents are composed of some keratin flakes shown by arrows (10x H&E stain). (C) Abundant keratin flakes (black arrows) within the cyst (4x H&E stain).

with a bimodal age distribution of 20's and 50's with a mean age of 39.9 years (Table 1). A majority of these are unilocular and well-circumscribed with a maximum lesion size ranging from 2.0 to 6.5 cm. There is no definite predilection for sex or location of the lesion within the thyroid gland (Table 1).

https://scholar.rochesterregional.org/advances/vol2/iss2/5 DOI: 10.53785/2769-2779.1092

The most common presenting symptom is a painless, non-tender thyroid nodule (11 out of 19). Of the 19 cases, 12 are biochemically euthyroid given the benign nature of this lesion. Pain is reported in only 4 cases. Other presenting symptoms are neck mass, globus pharyngeus, dysphonia, dysphagia, and left recurrent nerve palsy. In our case, patient presented with a swelling in the neck with no other associated symptoms even though biochemically patient was subclinically hyperthyroid. The pathogenesis of this nodule associated with subclinical hyperthyroidism is anomalous given that this epidermoid nodule was cold on the nuclear medicine thyroid scan. Likely, this patient's subclinical hyperthyroidism is an incidental discovery. There are no surrounding inflammatory signs to suggest leakage of thyroid hormone like thyroiditis based on radiographic imaging, gross pathology, and histology.

On sonographic imaging, an epidermoid cyst typically appears as a unilocular, well-circumscribed, and hypoechoic which is similar to our case. Hypoechogenicity is due to the densely packed proteinaceous/keratinaceous material in the cvst. Cvsts can present with hyperechoic foci as the result of highly compact deposits of cholesterol or fat as was observed in our case which are similar in appearance to microcalcifications and hence can be considered highly suspicious (Fig. 1). In contrast to the typical US characteristics, one case from literature describes internal content of the cyst as isoechoic, which likely is due to variable ratio of proteinaceous to fat content. Another distinct epidermoid cyst consists of a superficial hyperechoic wall with heavy posterior acoustic shadowing. Due to this shadowing artifact, internal contents are not visualized.⁴ This is likely due to a partially calcified epidermal cyst with active surrounding inflammation, which has been reported in epidermal cysts in other regions.⁵

Based on sonographic imaging and FNA in our case, besides thyroid parenchymal neoplasms, differential diagnoses which should be considered include dermoid/epidermoid, laryngocele, thymic cyst, lymphatic malformation, intrathyroid thyroglossal duct cyst and intrathyroid branchial cleft cyst.⁶ Although the risk of malignant transformation is rare, squamous cells within the thyroid gland can present as metaplastic or dysplastic cells. Benign squamous metaplasia can occur in goiter, myxedema, and thyroiditis.⁷ Squamous dysplasia can be seen in papillary carcinoma, squamous carcinoma, mucoepidermoid carcinoma, adenosquamous carcinoma, and teratomas.

In the majority of the cases, including in our case, the lesions are primarily treated with

Table 1. Cases reported to date.

Case	Age	Gender	Location	Size	US	Symptoms, Physical exam	Biochemical	Treatment	Reference (Ref)
1	28	М	Right	$6.5 \times 6.0 \text{ cm}$	_	Painless, firm, swelling for 10 years	Euthyroid	Hemithyroidectomy	Ref ⁹
2	57	М	Right	$3.1 \times 1.8 \times 3.2$ cm	Cystic, hypoechoic with punctate echogenic foci, and smooth margins	Dysphonia and dysphagia	Euthyroid	Hemithyroidectomy	Ref ¹⁰
3	28	М	-	_	_	Painless swelling of neck for 2 years	Euthyroid	Surgical excision	Ref ¹¹
4	15	М	Left	$2.0 \times 2.8 \times 3.2$ cm	Solid and homogeneous hypoechoic lesion macrocalcifications	Smooth, painful, palpable lump for 3 weeks with globus pharyngeus and pain with swallowing	Euthyroid	Drainage with close followup	Ref ⁸
5	54	М	Right	$3.2\times3.0\times2.6~cm$	Hypoechoic mass	Asymptomatic	Euthyroid	Hemithyroidectomy	Ref ¹²
6	46	F	Right	$2 \times 3 \text{ cm}$	Characteristics of a colloid goiter (iso-hypoechoic with possible multiple echogenic foci)	Painless neck swelling for 6 months.	Euthyroid	Hemithyroidectomy	Ref ¹³
7	24	М	Left	$4 \times 3.5 \times 2.0$ cm	Homogenous, isoechoic to thyroid parenchyma with a clear margin	Painless neck swelling for 10 years	Euthyroid	Surgical excision	Ref ¹⁴
8	26	F	-	$2 \times 2 \text{ cm}$	_	Painless, firm mass for 6 months	Euthyroid	Hemithyroidectomy	Ref ¹⁵
9	26	F	Right	-	-	Pain with swallowing and neck swelling for 6–7 months	Euthyroid	Hemithyroidectomy	Ref ¹⁶
10	38	F	Left	$1.2 \times 3.0 \times 1.0~\text{cm}$	Hypoechoic lesion	Nonpalpable	Euthyroid	Hemithyroidectomy	Ref ³
11	27	Μ	Left	3.2 × 1.5 cm	Crescentic, slightly hyper- echoic anterior surface with deep posterior acoustic attenuation, making it impossible to evaluate the internal structure	Palpable lesion slowly growing for 10 years.	Euthyroid	Hemithyroidectomy	Ref ⁴
12	60	М	Left	$3.0 \times 3.0 \text{ cm}$	Hypoechoic mass	Dysphonia, and left recurrent laryngeal nerve palsy	_	Hemithyroidectomy	Ref ¹⁷
13	50	F	Right	4.4 cm	Hypoechoic mass with multiple echogenic foci	Nontender, palpable mass in neck with discomfort on swallowing	Euthyroid	Hemithyroidectomy	Ref ¹⁸
14	9	F	Left	$3.0 \times 2.0 \text{ cm}$	_		_	Surgical excision	Ref ¹⁹
15	78	М	Left	2.0 cm	_	Tender nodule and discom- fort with swallowing	-	Hemithyroiidectomy and isthmusectomy	Ref ²⁰
16	58	М	Right	1.1 cm	Hypoechoic	Asymptomatic, incidentally found	-	Hemithyroiidectomy and isthmusectomy	Ref ²⁰
17	26	F	Left	_	_	Tender, palpable, mobile mass in neck	Euthyroid	Simple excision	Ref ²¹

(continued on next page)

1 able	T. (CO1.	аппиеа)							
Case	Age	Gender	Location	Size	ns	Symptoms, Physical exam	Biochemical	Treatment	Reference (Ref)
18	37	М	Right	. 1	Mixed echogenic multi- loculated solid-cystic lesion	Painful palpable neck swelling for 6 weeks followed by sore throat and progres-	Subclinical hypothyroidism with low TSH and normal T3 and T4	Hemithyroidectomy	Ref ²²
19	72	ц	Right	$5.0 \times 4.0 \times 2.5 \text{ cm}$	Hypoechoic mass with scattered hyperechoic foci	sive neck swelling for 2 weeks Asymptomatic painless neck swelling	Subclinical hypothyroidism with normal T3 and T4	Hemithyroidectomy	current case

hemithvroidectomy and among 19 cases, only 2 reported simple excision. A decision to perform a hemithyroidectomy was made based on the high TI-RADS criteria. None of the case reports mention any recurrence. Interestingly, one case reports drainage of the cyst with close ultrasound follow-up for a 15year-old patient.8 The initial size of the lesion measured 2.0 \times 2.8 \times 3.4 cm, which was followed by drainage and reduction in the size of the lesion to $1.1 \times 0.5 \times 2.5$ cm on follow-up with no recurrence. Drainage is a possible alternative, especially if patients would like to avoid surgery due to the potential postoperative morbidity, including recurrent laryngeal nerve injury, bleeding, and infection. However, long-term follow-up of these patients is needed, to ascertain if this is a reliable alternative to simple excision or hemithyroidectomy.

Conflict of interest

The authors have no conflict of interest to declare. All co-authors have seen and agree with the contents of the manuscript and there is no financial interest to report.

References

- 1. Tessler FN, Middleton WD, Grant EG, et al. ACR thyroid imaging, reporting and data system (TI-RADS): white paper of the ACR TI-RADS committee. *J Am Coll Radiol*. 2017;14(5): 587–595. https://doi.org/10.1016/j.jacr.2017.01.046.
- 2. Knox MA. Thyroid nodules. Am Fam Phys. 2013;88(3):193-196.
- Palombini L, Cozzolino I, Luigi S, Pifano A. Fine needle cytology of intrathyroid epidermoid cyst. *Diagn Cytopathol.* 2015;43(5):390–391. https://doi.org/10.1002/dc.23202.
- Bekele W, Gerscovich EO, Naderi S, Bishop J, Gandour-Edwards RF, McGahan JP. Sonography of an epidermoid inclusion cyst of the thyroid gland. *J Ultrasound Med.* 2012; 31(1):128–129. https://doi.org/10.7863/jum.2012.31.1.128.
 Smirniotopoulos JG, Chiechi MV. Teratomas, dermoids,
- Smirniotopoulos JG, Chiechi MV. Teratomas, dermoids, and epidermoids of the head and neck. *Radiographics*. 1995;15(6):1437–1455. https://doi.org/10.1148/radiographics. 15.6.8577967.
- Yilmaz M, Haciyev Y, Mamanov M, Cansiz H, Yilmaz R. Epidermal inclusion cyst of the larynx. J Craniofac Surg. 2011; 22(6):e1-e2. https://doi.org/10.1097/SCS.0b013e31822ec818.
- Frank E, Macias D, Hondorp B, Kerstetter J, Inman JC. Incidental squamous cell carcinoma in an epidermal inclusion cyst: a case report and review of the literature. *Case Rep Dermatol.* 2018;10(1):61–68. https://doi.org/10.1159/000487794.
- Lauffer P, van Schuppen J, Mooij CF. Epidermal inclusion cyst of the thyroid: a rare case of a nodule-like structure at ultrasound. *BJR Case Rep.* 2020;6(4), 20200038. https://doi.org/ 10.1259/bjrcr.20200038.
- Mansi JP, Riti TKS. A rare case of epidermal inclusion cyst of thyroid gland. J Med Sci Health. 2017;4(2):22–24. https:// doi.org/10.46347/jmsh.2018.v04i02.004.
- Palacio MN, Gonzalez-Mosquera L, Rosenthal D, et al. Epidermal inclusion cyst in the thyroid gland [published correction appears in Stem Cell Investig. 2021 Aug 17;8:17. *Stem Cell Invest*. 2020;7:18. https://doi.org/10.21037/sci-2020-021.
- Kuduban O. Epidermal inclusion cyst of thyroid gland. *Eur J Med.* 2015;47(1):78. https://doi.org/10.5152/eajm.2014.15.

- 12. Chen KT. Fine-needle aspiration cytology of epidermoid cyst of the thyroid: report of a case and review of seven cases. *Diagn Cytopathol.* 2007;35(2):123–124. https://doi.org/10.1002/dc.20592.
- Dayanand DC, Nichat PD, Agarsal S, Turabi MA, Tayade MB. Epidermal cyst presenting as a solitary thyroid nodule: a rare case report. J Med Sci Clin Res. 2015;3(8). https://doi.org/ 10.18535/jmscr/v3i8.39.
- Kannan S, Akila L, Kuppuswamy M, Hedne N. Epidermal inclusion cyst in the neck masquerading as a thyroid neoplasm. *Thyroid Res Pract*. 2015;12(1):32–34. https://doi.org/ 10.4103/0973-0354.147290.
- Surbhi SK, Vyas AS. Thyroid epidermal cyst– a common cyst, rare site. Natl J Med Res. 2016;6(3):290–291.
- Binayke R, Deshpande KP. Epidermal inclusion cyst of the thyroid gland: an uncommon entity. *Int J Curr Res Med.* 2017; 3(3):72–75. https://doi.org/10.22192/ijcrms.2017.03.03.009.
- 17. Salib RJ, Radcliffe G, Gallimore A. Intra-parenchymal thyroid epidermal cyst presenting with a left recurrent laryngeal

nerve palsy. J Laryngol Otol. 2001;115(3):247-249. https:// doi.org/10.1258/0022215011907127.

- Hatada T, Ichii S, Sagayama K, et al. Intrathyroid thyroglossal duct cyst simulating a thyroid nodule. *Tumori*. 2000;86(3): 250–252. https://doi.org/10.1177/030089160008600313.
- Sonnino RE, Spigland N, Laberge JM, Desjardins J, Guttman FM. Unusual patterns of congenital neck masses in children. J Pediatr Surg. 1989;24(10):966–969. https://doi.org/ 10.1016/S0022-3468(89)80192-7.
- North Jr JH, Foley CAM, Hamill LRL. Intrathyroid cysts of thyroglossal duct origin. *Am Surg.* 1998;64(9):886–888.
 Impieri M, Russo R, Cappagli M, Cucchi I, Poggi P,
- Impieri M, Russo R, Cappagli M, Cucchi I, Poggi P, Calcina GG. Epidermoid cyst of the thyroid: report of a case. *Acta Chir Belg.* 1987;87(6):385–386.
- 22. Chatchomchaun W, Thewjitcharoen Y, Krittadhee K, et al. Epidermoid cyst abscess of the neck masquerading as a thyroid abscess [published online ahead of print, 2020 Jun 4]. Endocrinol Diab Metab Case Rep. 2020;2020:20–47. https:// doi.org/10.1530/EDM-20-0047.