Sonographic Appearance of Steatocystoma: An Analysis of 14 Pathologically Confirmed Lesions
지선낭종의 초음파 소견: 조직학적으로 진단된 14개 병변의 분석

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Purpose To evaluate the ultrasonographic characteristics of steatocystomas focusing on the features that aid in differentiating them from epidermal inclusion cysts and lipomas.

Materials and Methods The ultrasonographic findings of 14 histologically proven steatocystomas in 10 patients were retrospectively reviewed. The following features were assessed: the layer of involvement, shape, margin, echogenicity, posterior acoustic features, and the presence of a visible wall or intralesional striations. The findings were compared with those of subcutaneous lipomas and epidermal inclusion cysts to identify those findings that aid in the differential diagnosis of steatocystomas.

Results The majority of steatocystomas appeared as a subcutaneous mass (n = 6, 42.9%) or a mass involving both the dermal and subcutaneous layers (n = 6, 42.9%). Steatocystomas exhibited a well-defined smooth margin (n = 12, 85.7%) and homogeneous echogenicity (n = 9, 64.3%), and showed no specific posterior acoustic features (n = 9, 64.3%). The most important features that differentiated steatocystomas from epidermal inclusion cysts were a homogeneous internal echotexture (p = 0.009) and absent or less prominent posterior acoustic enhancement (p < 0.001). The features that distinguished steatocystomas from lipomas were the margin (p < 0.001), echogenicity (p = 0.034), internal echotexture (p = 0.004), and the absence of intralesional striations (p < 0.001).

Conclusion Steatocystomas appeared as well-defined homogeneous masses with mild or absent posterior acoustic enhancement.

Index terms Steatocystoma Multiplex; Soft Tissue Neoplasm; Ultrasonography
INTRODUCTION

Steatocystoma is a lesion that results from a hamartomatous malformation of the pilosebaceous duct, leading to an epithelium-lined cystic lesion containing sebaceous lobules (1). Steatocystomas may occur anywhere in the body, but are more commonly reported where pilosebaceous units are well-developed, including the trunk, neck, axillae, groin, scalp, and proximal extremities. The incidence of steatocystoma is unknown, but is considered a rare lesion that equally affects both sexes (2). They are known to appear during adolescence suggesting a possible hormonal trigger (3, 4).

Steatocystomas can be classified into steatocystoma multiplex and steatocystoma simplex, according to the multiplicity. Steatocystoma multiplex is known to be an autosomal dominant disorder, but sporadic cases have been reported. Recent studies have also revealed that steatocystoma multiplex is associated with mutations in the keratin 17 (K17) gene (5, 6).

Steatocystomas usually manifest as solitary or multiple palpable nodules that are skin-colored and asymptomatic. The diagnosis of steatocystoma multiplex may be performed clinically when intradermal lesions occur in typical locations with multiplicity, when there is a family history (7). However, steatocystoma simplex and sporadic cases of steatocystoma multiplex that manifest as subcutaneous palpable lesions may pose a diagnostic challenge. Clinically, the presentation of these lesions may be similar to those of other more common benign subcutaneous masses such as epidermal inclusion cysts (EICs), lipomas, or lipoma-variants.

The ultrasonographic findings of steatocystomas have been reported in a limited number of studies (7-11), three of which reported mammographic and ultrasonographic findings of steatocystoma multiplex occurring in the breast. Park et al. (10) reported the ultrasound findings of steatocystoma multiplex in four patients; sonographic findings revealed multiple well-circumscribed homogeneous lesions that either confined to the dermal layer or involved the subcutaneous fat.

Based on our clinical experience, we noted that steatocystomas have been erroneously assessed as other subcutaneous masses both clinically and sonographically. Thus, the purpose of this study was to evaluate the ultrasonographic characteristics of steatocystomas, focusing on the features that aid in the differential diagnosis of steatocystomas from EICs and lipomas.

MATERIALS AND METHODS

This retrospective study was approved by our Institutional Review Board, and informed consent was waived (IRB No. B-1904/532-108).

PATIENT SELECTION

A search of the electronic medical records revealed 17 patients that visited our hospital between May 2005 and June 2017, who 1) had been clinically diagnosed with either steatocystoma multiplex or simplex, and 2) had undergone ultrasound examination. Among these patients, we selected those who had undergone surgical excision of the lesion and were
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histologically proven to have steatocystoma. A total of 14 pathologically proven steatocystomas from 10 patients were included in our study. Among the 10 patients, six were diagnosed with steatocystoma multiplex and four with steatocystoma simplex. There were five female patients (mean age, 39.2 years, range 26–51 years) and five male patients (mean age, 53.8 years, range 46–64 years), with an overall mean age of 46.5 years (range, 26–64 years).

CONTROL GROUP SELECTION

On retrospective review of the initial radiologic reports of ultrasound examinations in our institution, we found that steatocystomas had been erroneously assessed as either EICs or lipomas in a majority of cases. In order to identify ultrasonographic findings that aid in the differential diagnosis of steatocystomas from EICs and lipomas, we selected cases of EICs and lipomas as the control group. We searched our electronic medical record between May 2005 and June 2017 to identify patients who had undergone ultrasound examination and was confirmed with either lipoma or EIC on subsequent histologic examination. There were 148 histologically confirmed lipomas and 104 histologically confirmed EICs. Only lesions with size equal to or less than 20mm were selected, considering the small size of the included steatocystomas. Among the lesions fulfilling the size criterion (44 lipomas and 69 EICs), 14 lipomas and 14 EICs were randomly selected based on a random number generated with Microsoft Excel. The lipoma group included 6 females and 8 males with a mean age of 51.5 years (range, 29–70 years), whereas the EIC group included 7 females and 7 males with a mean age of 44.7 years (range, 23–80 years).

SONOGRAPHIC EXAMINATIONS AND IMAGE ANALYSIS

Ultrasoundographic examinations were performed by 23 radiologists with 1–9 years of experience in musculoskeletal ultrasonography. Imaging was performed with either an iU22 scanner (n = 11) or an Epic7 scanner (n = 3) using a 5–12 MHz, 5–17 MHz, or 5–18 MHz linear array transducer (iU22 and EPIQ 7, Philips Healthcare, Best, the Netherlands).

Three readers who did not perform the ultrasonographic examinations retrospectively reviewed the acquired images. First, two readers (Y.K. and H.Y. with 1 and 9-year experience in ultrasonography) independently reviewed the images, blinded to the pathologic report. Discrepancies between the two readers were resolved by a third reader (H.P. with 5-year experience in ultrasonography).

The following features were evaluated; 1) location of lesion (intradermal, subcutaneous layer, involving both dermal and subcutaneous layers), 2) shape of lesion (round or oval, spindle-shaped, irregular), 3) margin of lesion (ill-defined, well-defined smooth, well-defined irregular), 4) echogenicity of lesion, 5) internal echotexture, 6) presence of intraluesional striations, and 7) posterior acoustic features. A lesion with well-defined smooth margin shows an abrupt transition between lesion and surrounding tissue, with an even, gradually curving interface. A well-defined irregular margin refers to lesions with an abrupt transition between lesion and surrounding tissue but shows lobulation, angulation or speculation. Lesions without clear demarcation between the mass and its surrounding tissue were categorized as having an “ill-defined margin.” The echogenicity of the lesions was analyzed with reference to the echogenicity of the dermal and subcutaneous layers, and was classified into 4 levels: hy-
poechoic to fat, isoechoic to fat, hyperechoic to fat but hypoechoic to the dermis, and isoecho-
ic or hyperechoic to the dermis. The internal echotexture of the lesion was assessed as ho-
mogeneous or heterogeneous, and the presence of intralosomal hyperechoic striations was
also recorded. The posterior acoustic features of the lesions were divided into four catego-
ries: no posterior acoustic feature (no posterior acoustic enhancement or shadowing), mild
posterior acoustic enhancement (a slight increase in echogenicity of the area posterior to the
mass, compared to adjacent tissue at the same depth), strong posterior acoustic enhance-
ment (marked increase in echogenicity of the area posterior to the mass, compared to adja-
cent tissue at the same depth), and posterior acoustic shadowing (the area posterior to the
mass appears hypoechoic compared with the adjacent tissue).

STATISTICAL ANALYSIS
Demographic data and imaging findings were summarized using descriptive statistics, in-
cluding means, ranges, and frequencies, as appropriate. Fisher’s exact test was performed to
assess the ultrasonographic findings that aid differentiation between steatocystomas and
EICs and steatocystomas and lipomas. To evaluate the interobserver agreement between the
two radiologists that assessed the sonographic features of the soft tissue masses, Cohen’s kapp-
a statistics were used and the weighted kappa value was obtained to determine the echo-
genicity, which was considered an ordinal variable. Kappa values were evaluated according to
the following classification: 0–0.20, poor agreement; 0.21–0.40, fair agreement; 0.41–0.60,
moderate agreement; 0.61–0.80, good agreement; 0.81–1.0, excellent agreement. All statistical
analyses were performed with SPSS (version 24.0, IBM Corp., Armonk, NY, USA) and Med-
Calc (version 18.10.2, Ostend, Belgium). A p value of < 0.05 was considered statistically signif-
icant.

RESULTS
A summary of the clinical findings is shown in Table 1. Lesions were located in the trunk
(n = 6), upper extremity (n = 4), axilla (n = 2), and the head and neck region (n = 2). Using ul-
trasound examination, the first diagnostic consideration was lipoma or a lipoma-variant in
six lesions, EIC in four lesions, neurofibromas in two lesions, and other benign masses in the
remaining two lesions.

SONOGRAPHIC FINDINGS OF THE STEATOCYSTOMAS
The mean size of the steatocystoma was 7.9 mm; range: 4 mm to 14 mm. Six lesions
(42.9%) were located within the subcutaneous layer (Fig. 1), two lesions (14.3%) were limited
to the dermal layer (Fig. 2A), and six lesions (42.9%) involved both dermal and subcutaneous
layers (Fig. 3). Lesions were round or oval (n = 12), spindle-shaped (n = 2), and irregular-
shaped (n = 1). The majority of lesions (n = 12, 85.7%) had a well-defined smooth margin,
whereas one lesion had a well-defined irregular margin, and another exhibited an ill-defined
margin. Three lesions (21.4%) were hypoechoic to subcutaneous fat (Fig. 4), six lesions
(42.9%) were isoechoic to subcutaneous fat (Fig. 3), three lesions (21.4%) were hyperechoic to
subcutaneous fat (Fig. 4), and five (35.7%) were hyperechoic to subcutaneous fat, but hy-
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The ultrasonographic features of steatocystomas in comparison with EICs and lipomas are summarized in Table 2. Steatocystomas mostly showed no specific posterior acoustic feature or mild posterior acoustic enhancement, whereas EICs mostly showed strong posterior acoustic enhancement (Fig. 5), and the difference was statistically significant (p < 0.001). In addition, steatocystomas mostly showed a homogeneous echotexture, while epidermoid in-
clusion cysts mostly showed a heterogeneous echotexture ($p < 0.009$). Other features, including the location, shape, margin, and echogenicity of lesions and intralesional striation did not show statistically significant differences between the two groups.

A larger proportion of the lipomas showed the following features, compared to steatocystomas, with statistically significant differences: 1) ill-defined margin ($p < 0.001$), 2) hyperechogenicity to subcutaneous fat ($p = 0.0034$), 3) heterogeneous internal echotexture ($p = 0.004$),
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Table 2. Ultrasound Features of Steatocystomas Compared to EICs and Lipomas

|                      | Steatocystoma (n = 14) | EIC (n = 14) | Lipoma (n = 14) | p-Value | p-Value |
|----------------------|------------------------|--------------|-----------------|---------|---------|
| **Location**         |                        |              |                 |         |         |
| Intradermal          | 2                      | 2            | 0               | 0.871   | 0.059   |
| Subcutaneous         | 6                      | 4            | 12              |         |         |
| Involving both layers| 6                      | 8            | 2               |         |         |
| **Shape**            |                        |              |                 |         |         |
| Round or oval        | 11                     | 13           | 5               |         |         |
| Spindle-shaped       | 2                      | 0            | 7               |         |         |
| Irregular            | 1                      | 1            | 2               |         |         |
| **Margin**           |                        |              |                 |         |         |
| Ill-defined          | 1                      | 0            | 11              |         | <0.001  |
| Well-defined smooth  | 11                     | 13           | 2               |         |         |
| Well-defined irregular| 2                      | 1            | 1               |         |         |
| **Echogenicity of the lesion** |              |              |                 |         |         |
| Hypoechoic to fat    | 3                      | 7            | 0               |         |         |
| Isoechoic to fat     | 6                      | 3            | 3               |         |         |
| Hyperechoic to fat, Hypoechoic to dermis | 5 | 3 | 7 |         |         |
| Isoechoic or hyperechoic to dermis | 0 | 1 | 4 |         |         |
| **Internal echotexture** |                      |              |                 |         |         |
| Homogeneous          | 9                      | 2            | 1               | 0.009   | 0.004   |
| Heterogeneous        | 5                      | 12           | 13              |         |         |
| **Intralesional striations** |                |              |                 | 1.000   | <0.001  |
| Absent               | 13                     | 14           | 3               |         |         |
| Present              | 1                      | 0            | 11              |         |         |
| **Posterior acoustic feature** |                |              |                 | <0.001  | 0.678   |
| Absent               | 9                      | 1            | 11              |         |         |
| Mild posterior acoustic enhancement | 5 | 3 | 3 |         |         |
| Strong posterior acoustic enhancement | 0 | 10 | 0 |         |         |
| Posterior acoustic shadowing | 0 | 0 | 0 |         |         |

Data are number of patients, unless indicated otherwise. p values are for comparison between groups with Fischer exact test. EIC = epidermal inclusion cyst

and 4) intralesional striations (p < 0.001) (Fig. 6).

**INTEROBSERVER AGREEMENT BETWEEN THE TWO READERS IN EVALUATING THE SONOGRAPHIC FEATURES OF THE SOFT TISSUE MASSES**

The interobserver agreement between the two readers was fair to moderate. The kappa coefficient for each variable was calculated as follows: 1) location (κ = 0.52), 2) shape (κ = 0.30), 3) margin (κ = 0.55), 4) echogenicity of the lesion (weighted κ = 0.51), 5) internal echotexture (κ = 0.46), 6) intralesional striations (κ = 0.50), and 7) posterior acoustic feature (κ = 0.57).
DISCUSSION

In our study, steatocystomas showed a well-defined smooth margin, homogeneous internal echotexture, and no specific posterior acoustic feature or mild posterior acoustic enhancement. The majority of cases in our study were initially diagnosed as lipomas, EICs, and neurofibromas. This aligns with previous reports that have shown that steatocystomas may be clinically confused with EICs, lipomas, xanthomatosis, neurofibromatosis, and syringomas (12, 13). Many benign subcutaneous masses are considered as nonspecific solid masses and require biopsy or excision for definitive diagnosis (14). The results of our study may aid in the differential diagnosis of steatocystomas from other benign subcutaneous masses based on ultrasonographic findings.

Ultrasonography can play an important role because subcutaneous lesions can be difficult to differentiate based on clinical findings. Park et al. (10) reported the ultrasound findings of 18 steatocystomas in four patients: sonographic findings revealed multiple well-circumscribed oval hypoechoic lesions with posterior enhancement. They reported that all 18 lesions appeared homogeneous internally, which is consistent with the results of our study. In their study, the lesions were either confined to the dermal layer (44.4%) or showed involvement of both dermal and subcutaneous fat layers (55.6%). However, in our study, only 2 lesions were confined to the dermal layer and the others were appeared as a subcutaneous mass or a mass involving both dermal and subcutaneous layers. This discrepancy may have resulted from a
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Selection bias in the inclusion of patients for our study, as typical multiple intradermal steatocystomas were not referred for ultrasound examination. A recent report by Zussino et al. (11) showed that steatocystomas may appear as hypoechoic nodules with well-defined hyperechoic borders with absence of Color Doppler signal. Furthermore, posterior acoustic enhancement could be seen when the internal content was liquid. This is consistent with the results of our study: 92.9% of the steatocystomas showed well-defined borders and 35% showed mild posterior acoustic enhancement. The well-defined borders of steatocystomas are attributable to the histologic feature of steatocystomas: the cyst walls are composed of three to five layers of squamous epithelium and are lined with flattened sebaceous glands (15, 16).

In our study, steatocystomas mostly showed a homogeneous echotexture and absent or mild posterior acoustic enhancement, whereas EICs showed a heterogeneous internal echotexture with strong posterior acoustic enhancement. The ultrasonographic features of EICs found in the present study are consistent with previously reported findings (17-20). Huang et al. (19) reported a pseudotestis appearance of EICs, with intralesional bright echogenic reflectors and filiform anechoic areas. This coincides with the heterogeneous internal echogenicity noted in EICs in our study. EICs are known to show a strong posterior acoustic enhancement (17, 18). This was found to be an important feature that differentiate EICs from steatocystomas, in our study. The internal content of the steatocystomas have been reported to be yellowish-dirty fluid that has acellular, granular debris, rare anucleated squamous cells and rare cholesterol crystals on histologic examination (21). The heterogeneity of the internal content of steatocystomas may have possibly resulted in a less prominent posterior acoustic enhancement compared to EICs. On encountering a subcutaneous mass with homogeneous echotexture and absent or mild posterior acoustic enhancements on ultrasound, one should consider the possibility of a steatocystoma rather than an EIC.

Steatocystomas differed from lipomas in terms of location, internal echotexture, and the presence of intralesional striations. Steatocystomas were present in variable locations and exhibited a homogeneous internal echotexture, but lipomas were almost invariably located in the subcutaneous fat layer and showed a heterogeneous internal echotexture. The typical ultrasound findings of lipoma are well-circumscribed mass with variable echogenicity compared with the surrounding soft tissue (14). Although lipomas usually have a well-defined appearance with an identifiable thin capsule, some may have an ill-defined margin that blends with the adjacent subcutaneous fat (22-24). In our study, lipomas were more frequently assessed to have ill-defined margins compared to steatocystomas. Our study results coincide with the study by Inampudi et al. (23), in which the majority of lipomas had ill-defined borders; 52–60% of the lipomas included in their studies were assessed to have ill-defined borders. Another characteristic feature of lipomas is that curved echogenic lines can be noted within the mass (14). This was one of the key features that differentiated lipomas from steatocystomas in our study.

Several limitations of our study are noteworthy. First, due to the retrospective design of the study, we could not control biases resulting from the use of different ultrasound machines, probes and the differences in the level of experience of the examiners. Second, the study sample was small, and further studies with a larger number of subjects are required. Third, there may be a selection bias in our patient group, as only a number of patients with steatocystoma
in the database underwent ultrasound; those with typical clinical presentations were diagnosed clinically and were not subjected to sonographic evaluation. Fourthly, due to the retrospective nature of the study, advanced ultrasound techniques such as elastography could not be applied. Lastly, the interobserver agreement of the sonographic features were limited in our study. The low agreement could have possibility resulted from the diverse ultrasonographic examination settings, and further analysis with controlled ultrasound examination settings may be necessary.

In conclusion, steatocystomas were noted as well-defined homogeneous masses with mild or absent posterior acoustic enhancement, and were located in the subcutaneous layer or involved both dermal and subcutaneous layers. These findings may aid the differential diagnosis of steatocystomas from other subcutaneous soft tissue masses on ultrasound.

**Author Contributions**

Conceptualization, Y.H., K.Y.; data curation, K.Y.; formal analysis, Y.H., K.Y., P.H.; investigation, Y.H., K.Y.; methodology, K.Y., A.J.M., K.H.S.; project administration, K.Y.; resources, Y.H., K.Y.; supervision, K.Y.; visualization, Y.H.; writing—original draft, Y.H., K.Y.; and writing—review & editing, P.H., L.E., L.J.W., A.J.M., K.H.S.

**Conflicts of Interest**

The authors have no potential conflicts of interest to disclose.

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지선낭종의 초음파 소견: 조직학적으로 진단된
14개 병변의 분석

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목적 지선낭종의 초음파 소견을 분석하고, 표피낭종 및 지방종과의 감별진단에 도움이 되는 소견을 알아보고자 하였다.

대상과 방법
10명의 환자에서 14개의 조직학적으로 확진된 지선낭종의 초음파 소견을 후향적으로 검토하였다. 병변의 위치, 모양, 경계, 에코 발생 정도, 후방 음향 특징 및 테두리 벽 또는 병변 내 줄무늬의 존재 여부를 평가하였다. 지선낭종의 초음파 소견을 분석하고, 표피낭종 및 지방종과의 감별진단에 도움이 되는 소견을 알아보고자 하였다.

결과 지선낭종의 대부분은 피하 종괴(n = 6, 42.9%) 또는 피부층과 피하층을 함께 침범한 종괴로(n = 6, 42.9%) 나타났다. 병변의 위치, 모양, 경계, 에코 발생 정도, 후방 음향 특징 및 테두리 벽 또는 병변 내 줄무늬의 존재 여부를 평가하였다. 지선낭종의 초음파 소견을 분석하고, 표피낭종 및 지방종과의 감별진단에 도움이 되는 소견을 알아보고자 하였다.

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결론 지선낭종은 초음파 검사상 경계가 좋고 균질하며, 후방 음향이 경도로 있거나 없는 종괴로 관찰되었다.