Imaging of otosclerosis
Bildgebung der Otosklerose

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ABSTRACT

Background Otosclerosis is an important cause of hearing loss and a widespread pathologic in ENT medicine. Although a high diagnostic value of CT with impact on therapy is emphasized in the literature, the disease is seldom seen in the radiological routine diagnostics due to ENT findings often being diagnostic. Radiologists detect it rather more often in cases of unclear hearing loss or cochlear implant candidates. The findings may be very subtle. They require a target search and an optimal imaging technique.

Methods This review article is based on a selective search of the literature in PubMed without any time frame restrictions as well as on the long clinical experience of the authors.

Results The paper focuses on imaging aspects of otosclerosis: current role, advice for imaging technique, characteristic imaging signs, radiological differential diagnoses and findings after stapedoplasty. Pathology, clinical signs and therapeutic options are summarized briefly.

Conclusions With an optimal technique and sufficient radiological experience otosclerosis can be correctly diagnosed by imaging in a high percentage of cases. Radiology plays the key role in the diagnosis of retrofenestral otosclerosis. It can provide valuable information for the management of complications after stapedotomy.

Key points:
- X-ray cross-sectional imaging continues to be the method of first choice in the radiological diagnostics of otosclerosis
- With an optimal imaging technique and sufficient experience otosclerosis can be radiologically detected
- In complications after stapedoplasty, causes can be identified supporting the indication for a reoperation

Citation Format

ZUSAMMENFASSUNG


Ergebnisse Dieser Beitrag hat die Otosklerose unter verschiedenen Gesichtspunkten zum Inhalt: Pathologie, klinische Zeichen und Therapieoptionen werden überblickmäßig dargestellt; das Hauptaugenmerk liegt auf der Bildgebung mit heutigem Stellenwert, Untersuchungstechnischen Hinweisen, charakteristischen Bildbefunden, radiologischen Differentialdiagnosen und bildgebenden Befunden nach Stapesplastik.

Schlussfolgerung Bei optimaler Untersuchungstechnik und ausreichender Erfahrung kann die Otosklerose in einem hohen Prozentsatz radiologisch diagnostiziert werden. Eine diagnoseentscheidende Rolle hat der Radiologe bei der retrofenestralen Form der Otosklerose. Im postoperativen Komplikationsmanagement kann er dem HNO-Arzt wertvolle Informationen liefern.
Introduction

Otosclerosis is the most common osteodystrophy of the temporal bone, and only this disease exclusively affects the petrous bone. It is a significant cause of hearing loss. The histological prevalence is about 2.5 % (up to 10 %), clinically it is significantly lower at 0.3 %–0.4 % in Caucasians and only 0.03 %–0.1 % in the African and Asian populations [1]. The disease usually manifests itself in the 3rd–4th decade of life and occurs bilaterally in 70 %–85 % of cases [1], and appears rarely in children [2]. Tending to run in families, twice as many females are affected compared to males [1].

Although otosclerosis is a common disease in ENT medicine, it is rarely encountered by radiologists, most frequently in clarification of unexplained hearing loss or in preoperative cochlear implant (CI) diagnostics or postoperatively in the event of complications. Less frequently there is a specific request for evidence of otosclerosis. X-ray cross-sectional imaging is the method of choice for the detection of otosclerotic foci. The findings can be very subtle and require a targeted search and as well as an optimal examination technique.

This article is intended to inform the reader about otosclerosis with main focus on the current status of imaging, image morphological signs and imaging findings after stapedoplasty.

Etiology and Histopathogenesis

The etiology of otosclerosis is largely not understood. The following models are discussed: Genetic causes (most likely autosomal dominant inheritance), autoimmune, cytokines of bone metabolism, measles infection, hormones, environmental influences [3, 4].

Histological examinations have demonstrated that otosclerosis is characterized by focal changes in the bony labyrinth that occur in stages. Four stages have been distinguished:

- Stage I: Inflammatory resorption of woven bone of the otic capsule by osteoclasts.
- Stage II: Replacement by spongy vascular bone (otospongiotic phase).
- Stage III: Osteoclastic degradation of the spongy bone, replacement by compact lamellar bone (otosclerotic phase).
- Stage IV: Suspension of remodeling processes, in which active resorption and recalcification frequently coexist [5].

Symptoms and clinical ENT Findings

The main symptom of otosclerosis is a unilateral or bilateral progressive conductive hearing loss (CHL) caused by foci on the oval window leading to stapes fixation with mechanical malfunction of the stiffened ring band [1, 3, 4]. Associated with this are typical results in the tuning fork tests (Rinne negative, Weber lateralized into the ear with stronger CHL) and absent stapedius reflex in the diseased ear. The manubrium is easily moved (test with pneumatic ear magnifier), and the middle ear is well-ventilated (test with tympanogram). In general, history of inflammation of the middle ear is absent.

Otoscopy often reveals a circumscribed reddening on the promontory (Schwartz's sign), caused by hypervascularized otosclerotic lesions [3].

Tinnitus is evident in the majority of cases [5]. Vestibular symptoms are less frequent and are described as undirected short-term vertigo [3, 5].

Likewise, combined hearing loss resulting from cochlear involvement or up to 10 % purely sensorineural hearing loss (SNHL) in isolated capsular otosclerosis are also observed [6, 7]. However, isolated capsular sclerosis without stapes fixation has been found very rarely in temporal bone preparations [8]. A typical phenomenon is the Carhart depression, a drop in bone conduction threshold of 15 dB in the pure tone audiogram at 1.5–2 kHz [3, 6, 9].

Various hypotheses explain the presence of SNHL in cases of otosclerosis. Histologically, it has been shown that involvement of the cochlear wall in active otosclerotic foci was associated with collagen deposits in the spiral ligament (hyalinization) and atrophy of the adjacent stria vascularis [10–14]; SNHL can be explained by both changes. Other hypotheses assume damage to hair cells by cytotoxic enzymes, inflammatory cytokine mediators, oxidative stress and reactive oxygen species. It is possible that these molecules enter the endolymph, impair hair cell function and cause SNHL [4, 15]. This hypothesis is supported by the detection of antioxidants in the serum of otosclerosis patients [16].

The symptoms and examination findings are not clearly pathognomonic. Clinical differential diagnoses include post-inflammatory ossicular fixation ( tympanosclerosis) or ossicular resorption, traumatic ossicular dislocation, congenital cholesteatoma, middle ear malformations, Paget's disease, brittle bone disease (the latter two are very rare), pseudo CHL in dehiscence syndrome of the superior semicircular canal or in a mass in the inner ear, and apparent CHL in cases of insufficient masking in pure tone audiometry (detailed overview in [3]).

Therapy

To date there is no causal therapy for otosclerosis. Therapy options range from audiological follow-ups, conventional hearing aids, standard surgical therapy in the form of stapedoplasty to implantable hearing aids and cochlear implants [17].

In stapedoplasty, the stapes superstructure is removed, the stapes footplate is perforated (stapedotomy) and a prosthesis is inserted into this perforation, which is attached to the long process of the incus by a loop or clip [17]. Although hearing improvement can be expected in about 90 % of operations [18, 19], and existing tinnitus is reduced after the operation [20], an approximately 1 % risk of deafness must be taken into account [18, 19].

Drug therapy approaches may consist of the use of 3rd generation bisphosphonates to prevent the progression of otosclerosis [21]. This, however, is not considered standard therapy. Active middle ear implants may be indicated in special cases, especially for moderate to profound hearing loss. Advanced otosclerosis with profound hearing loss bordering on deafness is an indication for a cochlear implant [3, 22, 23].
Imaging Diagnostics

Current status

Although the detection of otosclerosis-related lesions has already been achieved by polytomographic examinations [24, 25], and today even very small lesions can be depicted much more precisely by modern X-ray cross-sectional imaging [26–29], patients with a typical clinical otosclerosis constellation are often treated surgically without imaging. In the case of a combined or sensorineural hearing loss, a request for imaging is often made under the indication “unexplained hearing loss”. False negative findings regarding otosclerosis may result if MRI is performed as the first or sole procedure for this issue. The imaging diagnosis of otosclerosis requires a cross-sectional X-ray [26–29], which can also reliably detect other middle ear pathologies causing unexplained hearing loss [26–29]. After a retrospective analysis of 259 CT scans performed on patients with clinically suspected otosclerosis, Dudau et al. [29] conclude that CT is a valuable aid in the management of otosclerosis patients. They determined a positive predictive value of 100% for CT for the diagnosis otosclerosis, a relevant alternative diagnosis in 33% of the cases and an influence of CT findings in the indication for stapedoplasty in 45% of the cases. In this study, image analysis was performed by radiologists experienced in temporal bone reading. The CT examination technology was optimal. In the authors’ point of view, very subtle foci in the cochlear cleft are often overlooked in routine diagnostics, not least due to a suboptimal examination technique.

Hearing success can be proved immediately after stapes surgery. If employed, a postoperative documentation of the prophylaxis by imaging is not performed in Germany. If a progressive decline of inner ear function occurs during the hospitalization phase, revision is frequent regardless of the outcome of imaging, if it is performed at all [30, 31]. In the event of a later recurrence of hearing loss or vestibular symptoms, imaging can disclose the cause in a high percentage of cases, thus supporting patient counseling and indication for re-operation [30, 32–34], where there is a higher risk of deafness than in the initial operation.

Examination technology principles

CT, Cone Beam CT (CB-CT) and 3D rotational angiography are suitable for the detection of typical otosclerosis changes and postoperative complications. Currently, CT is regarded as the method of choice, as it is the only method that is widely available or can be used for otosclerosis diagnostics with respect to capacity. Due to the very small temporal bone structures, it is important to achieve optimal spatial resolution in all 3 procedures. High-quality axial and coronary multiplanar reconstructions (MPR) with a slice thickness of 0.3–0.6 mm should be created from a thin-layer isotropic axial data set; for CT, the spiral technique with the narrowest possible collimation should be selected [35]. The region of the oval window including the stapes plays a special role in pre- and postoperative imaging of otosclerosis patients. It can best be assessed by means of paraxial MPR [36], prepared with high magnification parallel to the longitudinal axis of the stapes by planning the sectional position on the coronal images (Fig. 1). Due to spatial resolution and radiation hygiene, it is advisable to examine both temporal bones separately in CB-CT [37] and to blend exactly on temporal bone borders in 3D rotational angiography.

Signs of otosclerosis on X-ray cross-sectional imaging

Depending on the localization of the bone remodeling foci, a distinction can be made between fenestral and retrofenestral forms, whereby retrofenestral otosclerosis rarely occurs in isolation without fenestral involvement and is also assumed to be a continuation of the fenestral form [26, 38]. Of the histologically described stages, the otospongiotic phase is characterized by areas of reduced bone density (demineralized zones) and is the easiest to detect. In the otosclerotic stage and when remodeling processes are suspended, the foci are again more dense and more difficult to differentiate from normal surrounding bone. A concomitance of otospongiotic and sclerotic components is also frequently observed on imaging. Determination of density values in the CT analysis of otosclerosis lesions [39] has not become generally applied.

In fenestral otosclerosis the bone remodeling foci are located along the medial tympanic cavity wall. They are most commonly found in the fissula ante fenestram (Fig. 2), a gap between the middle and inner ear located in front of the oval window and filled with fibrocartilaginous tissue in the embryonic period. Less frequent are plaques on the promontory, the round window (Fig. 3) and the tympanic facial canal.

As the disease progresses, the entire footplate may be affected, which is sometimes very thickened after many years (Fig. 2d). Familiarity with a thickened stapes footplate is helpful for stapedoplasty. During the recalcification phase, bony protrusions into the vestibule can occur (Fig. 3c), which then constrict it. The windows, very rarely also the stapes, can also be completely ossified (Fig. 3b, c). If a CI should be inserted, a bony closed round window provides an obstacle to the insertion of the electrode array which must be removed or bypassed.

A little-known phenomenon is incudomalleolar separation as a result of otosclerosis (Fig. 3d) [26]. Due to the close positional relationship, plaques on the promontory often contact the basal turn of the cochlea, so that strictly speaking a mixed fenestral-retrofenestral form is thus present (Fig. 3a, d).

Retrofenestral involvement (Fig. 4) occurs much more frequently due to the progression of fenestral otosclerosis with encroachment on the bone surrounding the membranous labyrinth than as an isolated manifestation (Fig. 5). Depending on the stage of the disease, more or less dense, spotty or band-shaped plaques are found in the os petrosum. The cochlea may have a double-ring appearance (Valvassori sign) (Fig. 4a). The otosclerotic lesions can reach all structures of the membranous labyrinth (cochlea, vestibule, semicircular canals, endo- and peri-lymphatic duct), the facial or internal auditory canal, but they can also lie isolated in the bone.

Despite the current high level of X-ray cross-sectional imaging technology, false negative findings in the sclerotic stage of fenestral otosclerosis cannot be completely excluded if there are no contour irregularities [40].
In addition to the classification into fenestral and retrofenstral otosclerosis, various grading systems have been proposed in imaging studies depending on focal location, density and extent [41–43], none of which has been generally accepted to date.

**Signs of otosclerosis on the MRI image**
During the active phase, the otosclerotic foci accumulate contrast agent [44, 45] (Fig. 6), which is only noticeable on thin-sliced (≤ 2 mm) contrast-supported T1-weighted or FLAIR sequences. The foci are even more difficult to identify and only partially visi-
ble as intermediate foci in the bony labyrinth on native T1-w sequences. If the findings are recognized, an X-ray cross-sectional image procedure should be performed to confirm the diagnosis. If the contrast enhancement is low or if the plaques are very small, in the experience of the authors, a false negative result is usually obtained in routine diagnostics. In the sclerotic stage, the diagnosis cannot be made by MRI.

**Differential diagnoses on imaging**

Radiology-based differential diagnoses primarily concern other osteodystrophies, which, however, are not isolated diseases of the temporal bone, but which can also affect it. Due to similar bone remodeling processes, otosclerosis is also known as isolated Paget’s disease; image morphology is correspondingly similar (▶ Fig. 7a). No differentiation to retrofenestral otosclerosis can be made if only a cross-sectional X-ray examination of the temporal bone is available, and the onset of the disease or the age of manifestation, which in the case of Paget’s disease is beyond the age of 40, is unknown. Involvement of the temporal bone in brittle bone disease is a rarity. Here the images likewise resemble those of retrofenestral otosclerosis [46]. However, it is unlikely that the diagnosis is not known at the time of imaging because patients exhibit typical clinical findings. Other bone remodeling diseases such as osteopetrosis and fibrous dysplasia are not considered as differential diagnoses due to their different appearance on X-Ray cross-sectional imaging.

Relevant differential imaging diagnoses are globuli ossei (▶ Fig. 7b), norm variants that the authors have so far only observed in younger children. These are remnants of embryonic cartilage, which, when located in the fissula ante fenestram fissula, are also referred to as the cochlear cleft [47]. They apparently ossify later (▶ Fig. 7c), and are seen between 32–40 % in the pediatric population under 10 years of age [47, 48].

Congenital stapes fixation cannot be differentiated from fenestral otosclerosis on the basis of clinical findings and formerly could not be detected by CT. Advances in X-ray cross-sectional imaging have increasingly made it possible to visualize a thickened stapes base plate, sometimes also an ossified stapes superstructure, as an imaging morphological correlate when making paraxial reconstructions (▶ Fig. 7d). It is important that otosclerosis-typical foci with reduced density are absent.

Diseases that cause osteolysis, hydroptic ear disease and malformations are characterized by a different image morphology and therefore they are not a radiological differential diagnosis. A recent study has been shown that endolymphatic hydrops is common in advanced otosclerosis [49].

**Imaging after stapedoplasty**

Prostheses used today are usually made of titanium and have significantly fewer metal extinction artifacts than previously-used gold prostheses. Simplified, stapes prostheses consist of a piston and a loop or clip [17]. The different types differ mainly in the loop
design, which cannot be differentiated on imaging. From the radiological point of view, the main task is to assess the position of the prosthesis on the basis of cross-sectional X-ray imaging. For an exact assessment, axis-aligned reconstructions are necessary, sometimes thin-sliced MIPs can be helpful.

When properly positioned, the stapes prosthesis is located in the posterior third of the oval window and does not protrude significantly beyond the window plane in the direction of the vestibule (▶ Fig. 8a, b) [30–34]. In the middle ear it is usually attached to the long process of the incus.

Prosthesis dislocation is the most frequent complication after stapedoplasty, which is usually accompanied by an acute functional disorder. If there is dislocation into the tympanic cavity (▶ Fig. 8c, d), then a more or less large gap arises between the end of the prosthesis and the stapes footplate, which prevents sound transmission to the inner ear fluid, and the patient suffers from renewed hearing loss. In dislocations into the vestibule (▶ Fig. 9) usually vertigo is to the fore. In some cases, triggering factors for prosthesis dislocation can also be found, such as a loosening of the anchorage on the long process of the incus (▶ Fig. 9c, d), a prosthesis fracture (▶ Fig. 10a, b) or a necrosis of the incus as a late complication (▶ Fig. 10c); in the latter case, there is a wafer-thin long incus process. Scarring (▶ Fig. 10d) can be assumed if the prosthesis is surrounded by flat or strand-like opacifications that draw towards the eardrum, which appears retracted.

Rare complications include perilymph fistula and granulations. Indirect signs of a perilymph fistula include air in the vestibule (this sign is useless shortly after stapedotomy, as air may have penetrated intraoperatively and has not yet been resorbed [31]) and/or a fluid level in the tympanic sinus.

If none of the described changes are present on X-Ray cross-sectional images in suspected complications, it is recommended to perform an MRI of the temporal bone, since granulation can only be classified reliably in this way [30]. They are characterized by strong contrast enhancement. In extremely rare cases, they can extend intracranially via the inner ear/inner auditory canal.
Summary

Unlike the ENT physician, the radiologist rarely encounters otosclerosis. The much more common fenestral form is associated with a typical clinical constellation where imaging is not considered necessary to confirm the diagnosis. In this form the image morphological changes can be very discreet. On X-ray cross-sectional imaging technique and targeted search are necessary for their detection. The situation is different in mixed forms and the very rare isolated retrofenstral otosclerosis. Here the radiologist plays a crucial role in establishing the correct diagnosis. If complications after stapedoplasty are suspected, the radiologist can provide information regarding the cause and can support the indication for repeated surgery.
Conflict of Interest

The authors declare that they have no conflict of interest.

References


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