Monika Sakowicz-Burkiewicz¹,A,C,F, Jerzy Kuczkowski²,B,E, Tomasz Przybyła¹,B,C,E, Marzena Grdeń¹,C,E, Anna Starzyńska³,B,E, Tadeusz Pawełczyk¹,A,E,F

Gene expression profile of collagen types, osteopontin in the tympanic membrane of patients with tympanosclerosis

¹Department of Molecular Medicine, Medical University of Gdansk, Poland
²Department of Otolaryngology, Medical University of Gdansk, Poland
³Department of Orofacial and Dental Surgery, Medical University of Gdansk, Poland

A - Research concept and design, B - Collection and/or assembly of data, C - Data analysis and interpretation, D - Writing the article, E - Critical revision of the article, F - Final approval of article

DOI: 10.17219/acem/68984

Article type: ORIGINAL PAPER
Submitted: 18.01.2016
Accepted: 14.02.2017
Published online: 21.09.2017
Abstract

Background. Tympanosclerosis is a pathological process involving the middle ear. The hallmark of this disease is the formation of calcium deposits. In the submucosal layer, as well as in the right layer of the tympanic membrane, the calcium deposits result in a significant increase in the activity of fibroblasts and deposition of collagen fibers.

Objectives. The aim of our study was to examine the expression level of genes encoding collagen type I, II, III and IV (COL1A1, COL2A1, COL3A1, COL4A1) and osteopontin (SPP1) in the tympanic membrane of patients with tympanosclerosis.

Material and methods. The total RNA was isolated from middle ear tissues with tympanosclerosis received from 25 patients and from 19 normal tympanic membranes. The gene expression level was determined by real-time RT-PCR. The gene expression levels were correlated with clinical Tos classification of tympanosclerosis.

Results. We observed that in the tympanic membrane of patients with tympanosclerosis the expression of type I collagen is decreased, while the expression of type II and IV collagen and osteopontin is increased. Moreover, mRNA levels of the investigated genes strongly correlated with the clinical stages of tympanosclerosis.

Conclusions. The strong correlations between the expression of type I, II, IV collagen and osteopontin and the clinical stage of tympanosclerosis indicate involvement of these proteins in excessive fibrosis and pathological remodeling of the tympanic membrane. In the future, a treatment aiming to modulate these gene expressions and/or regulation of the degradation of their protein products could be used as a new medical approach for patients with tympanosclerosis.

Key words: osteopontin, tympanosclerosis, collagen types

External funds
This study was supported by the Poland National Center of Science grant no. NN403211839 to TP.

Conflict of Interest
None declared

Acknowledgment
None declared

Address for correspondence
Monika Sakowicz-Burkiewicz

Department of Molecular Medicine
Medical University of Gdansk
ul. Debinki 7, paw. 27
80-211 Gdansk
Poland

Tel.: +48 58 349 2759
fax: +48 58 349 2797
e-mail address: ssak@gumed.edu.pl

Short title
Collagen and osteopontin in tympanosclerosis

Authors (short)
M. Sakowicz-Burkiewicz et al.
1. Introduction

Tympanosclerosis is a chronic disease of the tympanic membrane and middle ear manifested by the accumulation of collagen in the layer elastic and fibrous lamina propria of the tympanic membrane, submucosal membrane of the tympanic cavity, auditory ossicles and mastoid cavity. The disease process most often refers to the eardrum [1]. Tympanosclerosis limited to the tympanic membrane is called myringosclerosis and occurs in 24-82% of patients with tympanosclerotic lesions [1, 2]. The initiation of the immune response leads to the formation of deposits of hyaline with subsequent calcification and sometimes ossification of tympanosclerotic foci [3, 4]. These changes often lead to restricted mobility of the eardrum and ossicles, resulting in hearing loss. The tympanosclerotic plates formed from deposits of calcium, collagen fibers and hyaline masses result in conductive and mixed hearing impairment, by reducing the mobility of hearing elements, or in rare cases through the perforation of the eardrum, which exacerbates the hearing loss. Tympanosclerosis is characterized by dynamic ischemia and stiffening, and even the disappearance of the middle ear tissue [1, 5].

The etiology of tympanosclerosis is still not fully understood. A widely-accepted consensus recognizes the tympanosclerotic changes as a complication of acute inflammation or injury within the middle ear [2, 6-10]. A special role is attributed to otitis media with effusion, particularly if patient treatment involved the insertion of a catheter vent [11, 12]. Other factors that are taken into account in the etiology of tympanosclerosis include immunological processes [13, 14], genetic predispositions [15], hypertension and hypercalcemia [16].

Under the influence of cytokines and other regulatory factors in the lamina propria of the tympanic membrane, fibrosis is initiated, which involves degradation and vacuolization of fibrocytes. This results in the disintegration of cells and accumulation of deposits in the spaces between the collagen fibers. Endoplasmic reticulum released from the destroyed cells is equipped with calcium-binding receptors. This leads to an accumulation of calcium deposits and eventual calcification of tympanosclerotic foci [17]. The activated immune cells as well as activated fibroblasts promote tissue remodeling. Our previous in vitro study on fibroblasts isolated from tympanosclerotic lesions demonstrated an up-regulated sensitivity of these cells to mast cell stimulation, which could significantly contribute to the ongoing fibrosis and pathological remodeling of the tympanic membrane [18].

Our present study aimed to evaluation the expression level of genes encoding type I, II, III and IV collagen, and osteopontin in the tympanic membrane of patients with tympanosclerosis. Moreover,
we determined the correlations between the gene expression levels and clinical stages of tympanosclerosis.

2. Material and Methods

2.1 Material

Tympanic membranes were obtained from 25 patients with tympanosclerosis who underwent tympanoplasty in the local Department of Otolaryngology of the Medical University. Patients with accompanied inflammatory disease, with immune deficiency or those taking antibiotics or glucocorticosteroids within the last two weeks were excluded. The clinical stage of the tympanosclerotic lesion was assigned based on the intraoperative evaluation of tympanosclerotic changes according to a modified Tos classification as described previously [19]. Morphologically normal tympanic membranes were dissected from 19 persons who had died suddenly. The institutional review board at the Medical University previously approved all procedures (NKEBN/432/2009), and written consent was obtained from all patients. Immediately after resection, tissues were placed in RNALater stabilization solution (Thermo Fisher Scientific, Walther, Massachusetts, USA) and stored at -20 °C until isolation of RNA.

2.2 Isolation of total RNA

Isolation of total RNA was carried out in accordance with the Chomczyński procedure [20], with our own modifications. A TRI reagent and suspended material were vortexed briefly and then left standing for 10 min at 4 °C. Next, chloroform (250 μl) was added, and the samples were vigorously shaken, incubated at 4 °C for 15 min and centrifuged (10,000×g for 15 min at 4 °C). The upper aqueous phase was removed into a new Eppendorf tube, an equal volume of isopropanol was added and RNA was precipitated by overnight incubation at −20 °C, followed by centrifugation (10,000×g for 15 min at 4 °C). RNA pellets were washed first with 96% and then with 70% (v/v) ethanol, air-dried and resolved in diethyl-pyrocarbonate-treated thermo-sterilized water and stored at −20 °C until further analysis.

2.3 mRNA level determination

The gene expression level was determined by real-time PCR performed in a Light Cycler 480 II (Roche Diagnostic GmbH, Manheim, Germany) using Path-IDTM Multiplex One-
Step RT-PCR Kit and appropriate Universal ProbeLibrary Set, Human (Roche Applied Science). Transcript levels were normalized to that of the β-actin gene (ACTB). The primer sequences, TaqMan probes and cycling conditions used are listed in Table 1.

2.4 Statistical analysis

Statistical analysis was performed using Statistica v.12.0 (StatSoft, Inc., Tulsa, OK, USA). The level of gene expression in a normal tympanic membrane (control) and the tympanosclerotic samples were analyzed with a nonparametric Mann-Whitney U test. The correlations between the level of gene expression and clinical classification of tympanosclerosis were analyzed using Spearman R ratio.

3. Results

To evaluate the contribution of collagens and osteopontin to the pathogenesis of sclerotic changes in the tympanic membrane, we determined the gene expression profile of COL1A1, COL2A1, COL3A1, COL4A1 and SPP1 in the tympanic membrane of patients with tympanosclerosis. The real-time PCR analysis performed showed the presence of COL1A1, COL2A1, COL3A1, COL4A1 and SPP1 in normal and sclerotic tympanic membranes, but the level of COL2A1 mRNA was barely detectable in the normal tympanic membrane. The level of gene profile expression of the collagen types and osteopontin in the selected structures of the middle ear with tympanosclerosis differed significantly compared to the healthy tympanic membranes (control group) (Figure 1). We observed that the level of COL1A1 transcript was significantly decreased in tympanic membrane subjects with tympanosclerosis (Figure 1A), whereas the expression level of COL2A1 was higher in sclerotic lesions of the tympanic membrane (Figure 1B). The transcript levels of COL3A1 and COL4A1 were not altered in sclerotic tympanic membranes compared to controls (Figures 1C, D). The expression level of the osteopontin gene (SPP1) was 7-fold higher in tympanosclerotic membranes compared to that determined in normal tympanic membranes (Figure 1E).

Figure 1

Altered expression of genes (COL1A1, COL2A1, COL3A1, COL4A1) encoding collagens; type I (A), type II (B), type III (C), type IV (D), and gene (SPP-1) encoding osteopontin (E) in sclerotic lesion of tympanic membranes (TTM) and normal tympanic membranes (NTM). The data is means from at least three independent measurements performed on isolated total RNA from tympanic membranes. *p < 0.0007; * * p < 0.002; * * * p < 0.00004
Moreover, we compared the mRNA levels of the investigated genes within the patient groups arranged based on Tos’s clinical classification of tympanosclerosis with our own modification. Our modification of the Tos classification concerns the study group division into two (instead of three), i.e. group I – myringosclerosis with/or without perforation; group II – tympanosclerosis with/or without perforation. We observed a much lower mRNA level of the COL1A1 gene in patients with stage II of tympanosclerosis. Expression of the COL1A1 gene was inversely correlated with the degree of tympanosclerotic changes (Spearman’s rank correlation (R) = -0.82; p<0.05). Moreover, increased expression of COL2A1, and SPP1 genes strictly correlated with the severity of the disease (R=0.8 and R=0.82 p<0.05, respectively). Interestingly, the mRNA level of the COL4A1 gene, which was not different in the study group compared to the control group, was correlated with the degree of tympanosclerotic changes (R=0.48 p<0.05, Figure 2C).

Figure 2

Relation of the tympanosclerotic stage and the expression level of genes (COL1A1, COL2A1, COL4A1) encoding collagens; type I (A), type II (B), type IV (C), and gene (SPP-1) encoding osteopontin (D). The data is means from at least three independent determinations performed on isolated total RNA from tympanic membranes. *p < 0.0007; ** p < 0.0002; *** p < 0.02; **** p < 0.05

4. Discussion

Tympanosclerosis (TS) is a chronic disease that is encountered at any age, but usually occurs between 30-50 years. However, as many as 87% of patients are over 40 years old [21, 22]. The pathogenesis of TS is not clear. It usually develops following middle ear infection during the resolution phase of chronic otitis media. The clinical observations indicate that there are some differences in the disease procession between old and young patients. Tympanosclerosis in children is associated with secretory otitis media and the lesions are mostly limited to the tympanic membrane, whereas the tympanosclerotic changes in the elderly are also observed in other middle ear sites: the ossicular chain or the mastoid cavity. These changes are often accompanied by perforation of the tympanic membrane, varying degrees of destruction of the ossicles and accompanied diseases, such as hypercalcemia, hyperlipidemia or hypertension and atherosclerosis [1, 9]. The risk of TS especially increases in children which had ventilation tubes inserted [12, 23]. The clinical observations indicate that the frequency of myringosclerosis is much higher in tympanic membranes with tympanostomy tube insertion than in tympanic membranes with no tympanostomy tube insertion (23-70% and 0-13%, respectively) [5, 11, 23-30]. It was also observed that the myringosclerosis rate increased with a larger size of tube, several tube insertions and time of tympanostomy tube stay [25, 29, 31, 32]. Based on
computer modeling, it has been shown that ventilation tube insertion induces shear stress in the structure of the tympanic membrane. The areas of maximal shear stresses have been found in the same positions as tympanosclerosis. It has been proposed that such stresses could damage the fibrils that connect the fibrous layer of the lamina propria and lead to TS [33]. Furthermore, hyperoxic conditions, foreign body reaction, inflammation, fibrous hyperplasia, hemorrhage and the release of free hemoglobin between the layers of the tympanic membrane are considered as factors involved in the development of myringosclerosis following tympanostomy tube insertion [25, 29, 34].

Typically, tympanosclerotic changes proceed via the destruction of connective tissue followed by fibrosis resulting in elevated deposition of collagens and subsequent calcification of tissues in the middle ear. A recent histological study demonstrated that a healthy human tympanic membrane consists of collagen type I, II, III and IV [35]. All these collagen types have different mechanical properties. The type I collagen fibers are resistance to force, and type II fibers are resistance to deformation. The type III collagen fibers are flexible and elastic, whereas type IV collagen provides support and transport [36].

Earlier studies showed that an autoimmune reaction (both cellular and humoral) to type II collagen, an essential component of ear tissue, leads to, among other things, sensorineural hearing loss, vestibular dysfunction, endolymphatic hydrops, Eustachian tube inflammation and otitis media with effusion (not infectious). The tympanosclerotic membrane also has C3 and Ig deposits, which may suggest that tympanosclerosis is also induced by type II collagen immunization, especially in patients undergoing surgical incision of the tympanic membrane [37, 38]. The mechanisms of this type II collagen autoimmune-mediated middle ear disease are not clear. In our study we observed that a normal tympanic membrane exhibits a very low level of COL2A1 mRNA, which tremendously increased (4.9-fold) in tympanosclerotic foci.

Our study has not shown any significant changes in COL4A1 expression in the whole group of tympanosclerotic lesions, but we did observe an increase in type IV collagen expression in a subgroup of patients with stage II of tympanosclerosis compared to patients with stage I. This might indicate that the expression of COL4A1 increases with the propagation of the disease.

Osteopontin (OPN), also known as bone sialoprotein I, is a universal regulator of inflammation, biomineralization and tissue remodeling. Osteopontin is expressed by a variety of cell types including fibroblasts, osteoblasts, osteocytes, odontoblasts, hypertrophic chondrocytes, dendritic cells and macrophages. The elevation of OPN level accompanies the exposure of cells to pro-inflammatory cytokines (e.g. TNFα, IL-1β, TGFβ). Several studies indicate that OPN is also up-regulated at sites of
pathologic, ectopic calcification [29]. A study by Makiishi-Shimobayashi et al. [40] suggested that macrophage-derived increased expression of SPP1 in inflammatory tissues of the middle ear is involved in the development of tympanosclerosis. Our present study showed that the expression of SPP1 is significantly higher in tympanosclerotic foci and positively correlated with the degree of tympanosclerosis changes. Therefore, manipulation of local OPN levels may be useful in the treatment of tympanosclerosis.

5. Conclusions

Tympanosclerosis is a result of post-inflammatory fibrosis characterized by elevated deposition of collagens, and calcification. The present study, with a detailed analysis of the expression of collagen types and osteopontin during the tympanosclerotic process, shows that in the tympanosclerotic membrane the expression of type I collagen is decreased, and the expression of type II and IV collagen and osteopontin is increased.

The altered secretory phenotype of cells from the middle ear induces histological remodeling of the tympanic membrane and correlates with the progression of tympanosclerosis.
References


[23] Vlastarakos PV, Nikolopoulos TP, Korres S, Tavouli E, Tzagaroulakis A, Ferekidis E:


Table 1. Primers and TaqMan probes and cycling conditions used for RT-PCR

<table>
<thead>
<tr>
<th>Gene transcript</th>
<th>Primers</th>
<th>TaqMan probe</th>
</tr>
</thead>
</table>
| COL1A1          | (F) gggattccttgacctaaag  
                 | (R) ggaacacctctgccttca  | Universal ProbeLibrary Probe # 67 (Roche) |
| COL2A1          | (F) tggtgctaagggcagaag  
                 | (R) cccagctctccacgtccac | Universal ProbeLibrary Probe # 4 (Roche) |
| COL3A1          | (F) agctggaaagattggtgcag  
                 | (R) ccttgaggacggagcag  | Universal ProbeLibrary Probe # 18 (Roche) |
| COL4A1          | (F) tggtgacaagagacagcag  
                 | (R) ggttcacccctttcagctg | Universal ProbeLibrary Probe # 81 (Roche) |
| SPP1            | (F) cccacagacctcctcaagta  
                 | (R) acactatcctggcagcatc | Universal ProbeLibrary Probe # 18 (Roche) |

Reverse transcription: 48°C (10 min), 95°C (10 min). Amplification: 95°C (10 s), 60°C (45 s)