NT journal review

Eustachian Tube Dysfunction



INTRODUCTION

- The middle ear is an air-filled chamber within the skull that is periodically vented when the eustachian tube opens.
- No well-accepted definition of eustachian tube dysfunction
- More accurately defined as failure of the functional valve of the eustachian tube to open and/or close properly



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Normal function of E-tube

- Three functional roles
- Equalization of pressure across the tympanic membrane
- Protecting the middle ear from infection and reflux of nasopharyngeal contents
- Clearance of middle ear secretions
- Open when swallowing and yawning
- <u>84 times per hours</u> in the daytime, less during sleep
- The valve is roughly 5 to 8 mm long in the middle of the cartilaginous portion of the tube.

Pathophysiology ~ Pressure dysregulation

- Fail to open adequately to allow for ventilation of the middle ear space
- Functional obstruction
- Mucosal inflammation with edema and secretions limiting the ability of the valve to open
- Failure of muscular actions
- Negative pressure make tubal muscles harder to dilate the lumen
- Floppy tube (young children with cleft palate, Down's syndrome, and other disorders with craniofacial anomalies)
- True anatomic obstruction: infrequently
- Severe inflammation of mucosa, adenoid mass, congenital or acquired stenosis, or nasopharyngeal neoplasm

Pathophysiology ~ Impaired protective function

- Reflux into the eustachian tube of nasopharyngeal pathogens, allergy-inducing proteins, gastric secretions, and sounds from one's own breathing or speaking (autophony).
- Reflux occurs in the following conditions:
- Abnormal patent E-tube due to congenital or acquired factors
- Short, floppy E-tube, creaniofacial anomalies (risk: Bottle-fed>breast-fed)
- Abnormal positive pressure in the nasopharynx (obstruction, blowing the nose, and crying)
- Loss of immune protection due to copious or viscous secretions, failre to expel pathogens
- Loss of mucosal protection from gastric enzymes and acid

Pathophysiology ~ Impaired clearance

- Loss of mucociliary function can result in the inability to clear viscous material and pathogens from the middle ear
- Primary loss of mucociliary function occurs in cystic fibrosis and other ciliary dysmotility disorders.
- Ciliary function may be secondarily impaired in inflammatory conditions.
- Impaired clearance may also be due to an obstructive process or inadequacy of tubal pump function.

Etiology

- Dilatory dysfunction:
- Failure of tubal dilatory action
- responsible for a large proportion of otitis media and chronic ear disease
- much more common than patulous dysfunction
- Patulous dysfunction
- valve incompetency leading to chronic patency (stuck open)
- bothersome symptoms
- Not lead to significant acute illness or chronic disease

Etiology~ Dilatory dysfunction

- Any cause of inflammation
- Allergies
- Irritants such as tobacco smoke, wood burning stoves, and pollution
- Laryngopharyngeal and gastroesophageal reflux
- Hormonal changes
- Primary mucosal disease (eg, granulomatous disease, Samter's triad)
- Ciliary disorders (eg, primary ciliary dyskinesia)
- Pressure dysregulation during scuba diving or descent during air travel
- Acquired anatomic abnormalities
- Nasopharyngeal masses
- Trauma (injury to palatopterygoid bone, trigeminal nerve or its mandibular branch)
- Congenital abnormalities (cholesteatoma, dermoid cysts, etc)
- Degenerative and metabolic diseases affecting the tubal musculature(ex MG)

Etiology~ Patulous dysfunction

- Weight loss, as little as a six pound reduction in weight (chronic illnesses, especially rheumatologic diseases)
- Scarring of mucosa as a result of previous procedures, inflammation, or radiation
- Neuromuscular disorders (eg, from cerebrovascular accidents, multiple sclerosis, poliomyelitis, traumatic injury to the trigeminal nerve)
- Allergy and chronic reflux of gastric contents, ultimately leading to mucosal atrophy.
- Hormonal factors (high estrogen levels in pregnancy or high dose oral contraceptives in women, estrogen therapy for prostate cancer in men)
- Others nasal decongestants or cocaine, craniofacial abnormalities, palatal myoclonus, chronic gum chewing

Clinical evaluation

Dilatory dysfunction

- Ear pain, ear fullness or pressure, hearing loss, tinnitus, "plugged"
- Popping and snapping noises
- <u>Vertigo and disequilibrium</u> (negative pressure or MEE)
- Delayed speech or language development

Patulous tube dysfunction

- Autophony!
- Hearing his/her own voice and breathing sounds, "talking into a barrel"
- Relieved with head in dependant position
- Worsed by exercise and prolonged speaking
- Any retractions, effusions, atelectasis,
 cholesteatoma, perforations or tympanosclerotic plaques
- Excursions of tympanic membrane accompany the patient's breathing, stop after lying

Conductive hearing loss

- nil
- Nasal endoscopy, Audiometry, and Radiographic imaging.

Differential diagnosis

- Temporomandibular joint dysfunction
- •Meniere's disease
- •Superior semicircular dehiscence syndrome
- •Ear blockage, disequilibrium, vertigo with loud noise, conductive hearing loss, and autophony
- •MRI or high resolution CT scan

Treatment for Dilatory dysfunction

- Most commonly caused by inflammation
- If not bacterial infection, systemic decongestants (pseudoephedrine and phenylephrine), antihistamines, or nasal steroid sprays for presumptive viral rhinosinusitis or allergic rhinitis.
- Direct toward the suspected etiology of tubal dysfunction

Medical management for Dilatory dysfunction: Decongestants

- Systemic decongestants helpful for congestive symptoms (ear fullness or pressure)
- Systemic or topic decongestants are not effective in cases of OME

Johnson D et al. Intranasal phenylephrine-surfactant treatment is not beneficial in otitis media with effusion. Int J Pediatr Otorhinolaryngol 2008; 72:1085

- Topical nasal decongestant sprays or drops may be used for symptomatic relief of nasal congestion or rhinitis with/without ear blockage symptoms
- Topical nasal decongestants may be helpful for difficulty clearing the ears during flights and scuba diving

Shapiro GG et al. *Treatment of persistent eustachian tube dysfunction in children with aerosolized nasal dexamethasone phosphate versus placebo*. Ann Allergy 1982; 49:81

Medical management for Dilatory dysfunction: Glucocortisoids

• Animal studies have found oral methylprednisolone may be effective in relieving effusion and improving acute E-tube dysfunction.

Aynali G, et al. *The effects of methylprednisolone, montelukast and indomethacine in experimental otitis media with effusion*. Int J Pediatr Otorhinolaryngol 2011; 75:15.

• Topical nasal steroids failure to show benefit in the absence of sinonasal inflammation.

Gluth MB, et al. *Management of eustachian tube dysfunction with nasal steroid spray: a prospective, randomized, placebo-controlled trial*. Arch Otolaryngol Head Neck Surg 2011; 137:449

- No evidence of that 6 wks nasal steroids was effective in patients with OME +/negative ME pressure.
- No evidence of changing tympanogram from B/C to type A.

Norman, G., et al. Systematic review of the limited evidence base for treatments of Eustachian tube dysfunction: a health technology assessment. Clinical Otolaryngology 2014; 39, 6-21

Medical management for Dilatory dysfunction: Politzer maneuver

• A systematic review of six randomized trials found that Politzer devices are effective for treatment of otitis media with effusion in children.

Silman S, Arick D. Efficacy of a modified politzer apparatus in management of eustachian tube dysfunction in adults. J Am Acad Audiol 1999; 10:496 Perera R, et al. Autoinflation for hearing loss associated with otitis media with effusion Cochrane Database Syst Rev 2006

 It is uncertain which patients with E-tube dysfunction would be most likely to benefit from Politzer devices.



Surgical management for Dilatory dysfunction

- Surgery is indicated when medical management fails.
- Tympanostomy tubes are indicated for persistence over 90 days despite medical therapy in children with OME secondary to E-tube dysfunction.
- Longer-term tympanostomy tubes may have a higher risk for subsequent tympanic membrane perforation

van Heerbeek N, et al. *Therapeutic improvement of Eustachian tube function: a review*. Clin Otolaryngol Allied Sci 2002; 27:50

• Eustachian tuboplasty: reduces the thickness of the mucosa and submucosa of the more bulky posteriormedial wall of the tubal orifice and lumen in order to facilitate tubal dilation.



Abstract

FOUNDATION

Balloon Eustachian Tuboplasty: A Systematic Review

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OBJECTIVE: A systematic review and meta-analysis of the evidence on balloon Eustachian tuboplasty (BET) as a treatment modality for Eustachian tube dysfunction (ETD). We followed the PRISMA guideline and registered with PROSPERO No. CRD42014009461.

DATA SOURCES: We searched 12 databases including PubMed and Embase from <u>January 1, 2010, to April 7, 2014</u>, for studies of BET. Endpoints: change in symptoms, middle ear pathology, eardrum status, Eustachian tube function tests, hearing, adverse events, complications, and health-related quality of life.

REVIEW METHODS: Study quality was assessed using the modified Delphi technique quality appraisal tool for case series studies. Risk of bias was assessed using the Cochrane Collaboration's tool for assessing risk of bias.

RESULTS: Nine case series studies with 443 patients (642 tubes) were included. Population size n = 4 (7 tubes) to n = 210 (320 tubes). All studies were of poor quality and featured a high risk of bias. We found reduction of patient symptoms in ETD questionnaire (P < .001), postoperative normalization of the tympanic membrane, conversion of type B or type C into type A tympanograms, reduced mucosal inflammation, increased number of positive Valsalva test and Swallowing tests, improvement in Eustachian tube score, reduction in Sino-Nasal Outcome Test (SNOT)-22 score (P = .001), and increased quality of life (P = .001). No serious adverse events were found.

CONCLUSION: The evidence of BET is poor and biased. No firm conclusions can be made to identify patients who will benefit from the procedure or to accurately predict surgical results. Randomized controlled trials or case-control trials are needed.

Balloon dilation of eustachian tube



(A) Preoperative 30 degree angled rigid endoscopic view of left eustachian tube nasopharyngeal orifice showing inflamed, edematous mucosa circumferentially and especially at the posterior cushion. The lumen is swollen closed.

(B) Intraoperative inflation of a 7 mm diameter by 16 mm long balloon placed within the cartilaginous eustachian tube.

(C) Immediate postdilation view of the eustachian tube lumen showing some minor mucosal laceration and significant persistent widening of the lumen.

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Table 1. Inclusion and Exclusion Criteria.

| Participants | Persons eligible for balloon Eustachian tuboplasty with a clinical diagnosis of Eustachian tube dysfunction |
|--------------|--|
| | Exclusion: diagnosis of adenoid tissue, rhinopharyngeal tumors, patoulous tube, cleft palate |
| Intervention | Balloon dilation Eustachian tuboplasty |
| Comparators | Any or none |
| Outcomes | Change in symptoms (severity or frequency), middle ear pathology, eardrum status, Eustachian tube function tests, hearing, adverse events, complications, health-related quality of life |
| Study design | Any |
| | Exclusion: abstracts, publications without peer review |

Table 2. Resources Searched.

| | No. Hit |
|---|---------|
| PubMed | 36 |
| Embase | 36 |
| Web of Science (Science Citation Index Expanded & Conference Proceedings Citation Index-Science) | 41 |
| Scopus | 40 |
| Cochrane Database of Systematic Reviews (SDSR) | 1 |
| Cochrane Database of Abstracts of Reviews of Effects (DARE) | 0 |
| Cochrane Central Register of Controlled Trials (CENTRAL) | 0 |
| Cochrane Methodology Register | 0 |
| Health Technology Database | 0 |
| NHS Economic Evaluation Database | 0 |
| Latin America and Caribbean Health Sciences (LILACS) | 0 |
| Others: | |
| Clinicatrials.gov | 0 |
| Controlled-trials.com | I |
| Clinicaltrialsregister.eu | 0 |
| MHRA.gov.uk | 0 |
| Fda.gov | 0 |
| Who.int/ictrp/en | 0 |

ts

- Balloon dilatation Eustachian tuboplasty(BET)
- was first described by Ockermann et al in 2010
- The search was limited to results after Jan.2010
- No attempt was made to pool the data for a meta-analysis
- Using the modified **Delphi technique** quality
- appraisal tool for case series studies



Figure 1. Flow chart of the study selection.

*Reasons for exclusion: records not on subject, conference abstracts. **Reason for exclusion: no isolated balloon Eustachian tuboplasty results could be extracted.

• Dx: endoscopy, tubomanometry, Swallowing test,

CT, ENT exam, audiometry, tympanometry...

| Study | Design | n | Diagnostics |
|-------------------|--|---------------------------|--|
| Ockermann (2010) | Case series | 8 (13 tubes) | Rhinopharyngeal endoscopy, otomicroscopy, tubomanometry, Valsalva, Swallowing test, HR-CT of the temporal bone |
| Poe (2011) | Case series | (tubes) | Rhinopharyngeal endoscopy, otomicroscopy, tympanometry, HR-CT of the temporal bone, Valsalva |
| McCoul (2012) | Case series | 22 (35 tubes) | ENT exam, pneumatic otoscopy, tympanometry, pure-tone audiometry, CT of the temporal bone |
| Schröder (2012) | Case series | I group of 66 (II5 tubes) | ENT exam, audiometry, tympanometry, tubomanometry, Valsalva, Swallowing test, CT of the temporal bone |
| | | I group of I2 (20 tubes) | - |
| Catalano (2012) | Case series | 70 (100 tubes) | Tympanometry |
| Jurkiewicz (2012) | Case series | 4 (7 tubes) | ENT exam, rhinopharyngeal endoscopy, tympanometry, pure-tone audiometry, pressure- swallow test, Valsalva, CT angiography of the carotid artery |
| Tisch (2013) | Case series | 210 (320 tubes) | Valsalva, Toynbee test, subjective evaluation, tympanometry, otomicroscopy |
| Bast (2014) | Retrospective quality of life questionnaire | 30 | ENT exam, audiometry, tympanometry, CT of the petrosal bone |
| Silvola (2014) | Prospective series Some of the patients are follow-up from a pilot study (Poe 2011) | 37 (41 tubes) | Rhinopharyngeal endoscopy with systematic mucosal inflammation score, otomicroscopy, tympanometry, HR-CT of the temporal bone, Valsalva |

Abbreviation: HR-CT, high-resolution computed tomography.

Table 3. Overview of Studies.

Table 4. Overview of Studies, Inclusions and Exclusions.

| Study | Inclusion (OME, score, | Exclusion | | | |
|-------------------|---|---|--|--|--|
| Ockermann (2010) | history) Symptoms of chronic obstructive ETD or recent tympanoplasties caused by acute relapsing OM or chronic OME as a consequence of ETD | N/a (TM joint, tumor, structure | | | |
| Poe (2011) | Unilateral or bilateral OME for ≥5 consecutive years, broken only by grommet insertion or TM perforation | N/a | | | |
| McCoul (2012) | Age >18 years, abnormal tympanogram (non-A- curve with admittance <0.2 mmho or resting pressure <-100 daPa, abnormal otosopic examination, symptoms of unilateral or bilateral ETD, failure to improve symptoms by medical therapy over 2 months | Any head and neck surgery within 3 months, history of RT to the head and neck, sinonasal malignancy, acute upper respiratory infections, nasal polyposis, cleft palate, craniofacial syndrome, cystic fibrosis or other immunodeficiency | | | |
| Schröder (2012) | Tuba score \leq 5, anamnestic and clinical signs of ETD | Age < 18 years, tumor of the rhinopharynx, dehiscent bony canal of the carotid artery, severe septal deviation, severe hypertrophy of the nasal turbinates | | | |
| Catalano (2012) | Adults with a history of chronic ETD symptoms | Temperomandibular joint disease, early hydrops | | | |
| Jurkiewicz (2012) | Lasting or periodic uni- or bilateral partial hearing loss, feeling of obstruction, clicking noises | N/a | | | |
| Tisch (2013) | Adult patients with symptoms of ETD and prior failed attempts to treat with medicine or surgery | N/a | | | |
| Bast (2014) | Chronic ETD | Age <18 years, unidentifiable opening of the ET in the nasopharynx, dehiscence of the bony canal of the carotid artery, septal deviation, hyperplastic turbinates | | | |
| Silvola (2014) | Unilateral or bilateral persistent OME or significant nonadherent TM ≥5 years and follow-up ≥1, 5 years | Dehiscence of the bony canal of the carotid artery | | | |

Abbreviations: ET, Eustachian tube; ETD, Eustachian tube dysfunction; OM, otitis media; OME, otitis media with effusion; RT, radiotherapy; TM, tympanic membrane.

Table 5. Overview of Studies, Methods and Follow-up.

| dies, Flechous and Follow-up. | |
|---|--|
| Methods | Follow-up |
| Bielefeld balloon system. The catheter was introduced into the cartilaginous and the bony canal of the ET. Inflation to 20 mm length and 3 mm width at 10 bar for 2 minutes | I, 2, and 8 weeks |
| Reliva Solo Sinus Balloon Dilation System, catheter introduced into the cartilaginous part of the ET. Inflation to 12 atm for 1 minute in 7 cases and 8-10 atm for 1 minute in 5 cases 5 patients had adjunctive procedures (2 grommet insertions, 3 | I and 6 months |
| ÷ , | 3, 6, 12 weeks and 6 months |
| | 5, 6, 12 weeks and 6 months |
| cartilaginous part of the ET. Adjunctive surgery: all patients had partial inferior turbinectomy; 15 submucosal resection of the nasal septum, 12 sphenoethmoidectomy with maxillary sinostomy, 2 revision ethmoidectomy, 1 grommet inserted, 1 | Three different instruments are in use: 1. Bielefeld Balloon System (Spiggle & Theiss, Overath, Germany) |
| Bielefeld Balloon System, dilation of the cartilaginous part of the ET; 6 patients had adjunctive surgery of paranasal sinuses, nasal | (length 20 mm, diameter 3.28 mm) 2. Reliva Solo Sinus Balloon Dilation System |
| Relieva Vortex Sinus Irrigation Catheter introduced into the | (length 16 mm, diameter 7 mm) |
| 44 patients had BET under GA (no specified pressure or time of dilation) 5 had adjunctive otologic surgery, 39 sinonasal surgery 26 patients had BET under LA in office setting. n = 5 tolerated only 6 atm dilation, n = 20 8 atm for 10 seconds, n = 22 8 atm | Reliva Vortex Sinus Irrigation Catheter (length 16 mm diameter 5 mm) (Acclarent, Inc, Menlo Park, California, USA). |
| Bielefeld Balloon System, dilation of the cartilaginous part of the ET | 6 weeks |
| Bielefeld Balloon System, dilation of the cartilaginous part of the ET | N/a |
| Bielefeld Balloon System, dilation of the cartilaginous part of the ET | 6-18 months |
| Reliva Solo Sinus Balloon Dilation System, catheter introduced into the cartilaginous part of the ET, 12 atm for 1 minute (15 patients from the pilot) | 1,5 years |
| | |
| All patients instructed to do Valsalva ≥2 times a day for 1 week after surgery | |
| | Methods Bielefeld balloon system. The catheter was introduced into the cartilaginous and the bony canal of the ET. Inflation to 20 mm length and 3 mm width at 10 bar for 2 minutes Reliva Solo Sinus Balloon Dilation System, catheter introduced into the cartilaginous part of the ET. Inflation to 12 atm for 1 minute in 7 cases and 8-10 atm for 1 minute in 5 cases S patients had adjunctive procedures (2 grommet insertions, 3 grommet removals) Reliva Solo Sinus Balloon Dilation System, catheter (5 × 24 mm² [n = 17] or 7 × 24mm² [n = 18]) introduced into the cartilaginous part of the ET. Adjunctive surgery: all patients had partial inferior turbinectomy; 15 submucosal resection of the nasal septum, 12 sphenoethmoidectomy with maxillary sinostomy, 2 revision ethmoidectomy if grommet inserted, 1 myringoplasty Bielefeld Balloon System, dilation of the cartilaginous part of the ET; 6 patients had adjunctive surgery of paranasal sinuses, nasal septum, or turbinates Relieva Vortex Sinus Irrigation Catheter introduced into the cartilaginous part of the ET 44 patients had BET under GA (no specified pressure or time of dilation) 5 had adjunctive otologic surgery, 39 sinonasal surgery 26 patients had BET under LA in office setting, n = 5 tolerated only 6 atm dilation, n = 20 8 atm for 10 seconds, n = 22 8 atm for 30 seconds Bielefeld Balloon System, dilation of the cartilaginous part of the ET Bielefeld Balloon System, dilation of the cartilaginous part of the ET Reliva Solo Sinus Balloon Dilation System, catheter introduced into the ET Bielefeld Balloon System, dilation of the cartilaginous part of the ET Bielefeld Balloon System, dilation of the cartilaginous part of the ET Bielefeld Balloon System, dilation of the cartilaginous part of the ET Reliva Solo Sinus Balloon Dilation System, catheter introduced into the cartilaginous part of the ET Reliva Solo Sinus |

Complications Study Results Ockermann (2010) Significant improvement in ETS at all follow-up examinations None Pre-BET mean ETS = 1.077, week 1 = 4.154, week 2 = 5.846, week 8 = 7.539 Pre-BET swallowing test = 12 never, 1 infrequent; week 8 = 1 never, 6 infrequent, 5 always Pre-BET Valsalva = 9 never, 2 infrequent, 1 always; week 8 = 1 never, 6 infrequent, 5 always Poe (2011) TM status recovered to normal in 2/4 with preoperative intact TM, the 2 Mucosal tear n = 5, C6-7 contralateral radiculopathy due remaining had grommets inserted when seen at follow-up. Tympanometry measured, but results are inconclusive due to perforations and to neck extension in I patient grommets. Mucosal inflammation was reduced from a mean of 2.91 (0.83) to 1.73 (no SD available) no P value reported. Preoperative Valsalva = 11 never; 6 month = 11 always McCoul (2012) Pre-BET TM status: 33 TM showed retraction (4 with OME, 1 with early attic n = I bleeding from the cholesteatoma), I had a grommet, and I had a stable TM perforation turbinectomy site on day 3 Post-BET TM status: 35 free of retraction, I ear had a resolving perforation resulting in bilateral from grommet insertion haematotympanon, Pre-BET tympanometry: type C = 20, type B = 5, type A = 10. myringotomy was required Post-BET tympanometry: type A = 34, type B = 1 (perforated) Mean ETDQ-7 scores improved significantly at all follow-up visits. Mean SNOT-22 scores improved significantly at all follow-up visits. Schröder (2012) Significant improvement after 2 months in tuba score by ≥ 2 points in 60%, 2 reports of self-limiting 1-2 points in 19%, and no change in 21% from a subgroup of 66 patients enhancement of known (115 tubes) tinnitus, I self-limiting epistaxis The first 20 patients were followed 12 months; 12 were available for follow-up; in this subgroup the tubas core was improved by ≥ 2 points in 10 patients and 1-2 points in 2 patients Catalano (2012) Pre-BET tympanometry: abnormal (type B or C) in 28 ears, post-BET I preauricular emphysema, improvement (type A) in 25 of the 28 resolved spontaneously Overall 71% patients showed notable improvement or reduction in symptoms (no information of specific endpoints included), Jurkiewicz (2012) Pre-BET tympanometry: abnormal (type B or C) in 7 ears, post-BET None improvement (type A) in 6 of the 7 Pre-BET Valsalva positive in 1/7 ears, post-BET Valsalva positive in 6/7 ears Pre-BET PST negative in 7/7 ears, post-BET PST positive in 7/7 ears Otoscopic examination pre-BET 7/7 TM retraction, post-BET otoscopic examination 5/7 TM normal, 2/7 no change Tisch (2013) Post-BET patient report of symptoms: 150 a lot better or completely 10 cases of minor epistaxis, 1 resolved, 35 better, 25 no change case of emphysema in the face, Valsalva pre-BET was negative in 92%, post-BET in 10% neck and mediastinum Significant improvement in total score (P = .001) and subscores general health Bast (2014) N/a (P = .001) and physical health (P = .039) in GBI TM status: pre-BET 41/41 abnormal, post-BET 37/41 normal Silvola (2014) None OME: pre-BET 38/41, post-BET 1 Retraction/atelectasis: pre-BET 3/41, post-BET 3/41 Tubal inflammation was reduced from 2.8 to 1.4 (P < .001) Tympanometry: pre-BET type A = 1, type B or C = 16 perforation or grommet 24; post-BET: type A = 23, type B = 0, type C = 6 perforation or grommet 12

Valsalva pre-BET was negative in 100%, post-BET in 20%

Table 6. Overview of Studies, Results and Complications.

Outcome and follow up

- Tisch et al and Catalano et al. : Increased satisfaction of subjective outcome
- McCoul and Anand: positive change in mean score of ETDQ-7(4.5->2.8) at 6 months
- Tympanometry show a high rate of conversion of type B or C into type A when follow-up.(6wk-1.5y)
- Mucosal inflammation at the tubal orifice improved at 6 months and 1.5 years
- Bast et al: QoL improved

| | Ockermann | Poe | McCoul | Schröder | Catalano | Jurkiewicz | Tisch | Bast | Silviola |
|--|-----------------|-----------------|---------|----------|----------|-----------------|-----------------|-----------------|----------|
| Hypothesis stated? | Yes | Yes | Yes | Unclear | Yes | Unclear | Yes | Yes | Yes |
| Participants described? | Yes | Yes | Yes | Yes | Yes | Yes | Yes | Yes | Yes |
| Inclusion and exclusion criteria? | Partial | Partial | Yes | Yes | Yes | Partial | Partial | Yes | Yes |
| Consecutive recruitment? | No | Unclear | Yes | Unclear | Unclear | Unclear | Unclear | Yes | Yes |
| Inclusion at similar point in the disease? | No | No | Unclear | No | Unclear | No | No | No | Yes |
| Procedure described? | Yes | Yes | Yes | Yes | Yes | Yes | Yes | Yes | Yes |
| Co-interventions reported? | No | Unclear | Unclear | No | Unclear | No | No | No | Unclear |
| Outcomes clearly defined? | Yes | Yes | Yes | Yes | No | Yes | Yes | Yes | Yes |
| Outcomes appropriately measured? | Yes | Yes | Yes | Yes | No | Yes | No | Yes | Yes |
| Outcomes measured before and after? | Yes | Yes | Yes | Yes | No | Yes | Yes | Not relevant | Yes |
| Appropriate statistical tests? | Not relevant | Yes | Yes | Yes | Unclear | Not relevant | No | Yes | Yes |
| Follow-up reported? | Yes | Yes | Yes | Yes | Yes | Yes | Not relevant | Yes | Yes |
| Loss to follow-up reported? | Not relevant | Not relevant | Yes | Yes | Yes | Not relevant | Not relevant | Yes | Yes |
| Variability in outcomes reported? | Yes | Yes | Yes | Yes | No | Not relevant | No | Yes | Yes |
| Adverse events reported? | Yes | Yes | Yes | Yes | Yes | Yes | Yes | No | No |
| Conclusion supported by results? | Yes | Yes | Yes | No | No | Yes | Yes | Yes | No |
| Conflict of interest reported? | Yes | Yes | Partial | Yes | Partial | Partial | Unclear | Yes | Yes |
| Prospective study? | Yes | Unclear | Yes | Unclear | Unclear | Unclear | Unclear | No | Yes |

Table 7. Study Quality.

The assessment of quality guided by modified Delphi technique quality tool for case series studies is listed in this table.

The overall quality of all the included studies is pool.

Discussion

- The underlying mechanism has not yet been identified, but it is hypothesized that submucosal microhemorrhages from the applied pressure cause fibrosis and expansion of the internal ET diameter during healing.
- Preopera-tive CT scan of the temporal bone and the exclusion of patients with a bony dehiscence or petrous aneurysms.
- Use EDTQ-7(>2.1), Valsalva's test and Tubomanometry for evaluation of outcome.

Validating the clinical assessment of eustachian tube dysfunction: The Eustachian Tube Dysfunction Questionnaire (ETDQ-7).

McCoul ED¹, Anand VK, Christos PJ.

Author information

Abstract

OBJECTIVES/HYPOTHESIS: Eustachian tube dysfunction (ETD) is a common condition that is associated with otologic and rhinologic symptoms. The complete assessment of ETD is limited without a valid symptom score. We developed and conducted initial validation of the seven-item Eustachian Tube Dysfunction Questionnaire (ETDQ-7), a disease-specific instrument to assess symptoms with respect to ETD.

STUDY DESIGN: Validation study.

METHODS: The ETDQ-7 was developed using standard survey methodology. The ETDQ-7 was completed by a group of 50 consecutive adult patients diagnosed with ETD and 25 non-ETD patients who served as a control group. Tympanometry was used as a criterion standard to distinguish the two groups. A subset of respondents repeated the ETDQ-7 at a time point 4 weeks later.

RESULTS: Content validity for the ETDQ-7 was established by focus group and review of the literature. Reliability testing indicated acceptable internal consistency for the entire instrument (Cronbach α = .71). The test-retest reliability indicated good correlation between the two questionnaires completed by the same patient 4 weeks apart (r = 0.78). The ETDQ-7 was able to discriminate between patients with ETD and those without (P < .001), indicating excellent discriminant validity.

CONCLUSIONS: The ETDQ-7 is a valid and reliable symptom score for use in adult patients with ETD that may facilitate clinical practice by highlighting the impact of ETD. Further testing is needed to determine its usefulness in assessing treatment response.

| The Seven Item Evet | TABLE I. | | Ouestiennei | *0 | | | |
|---|----------|-------------|------------------|-----|---|-------------------|---|
| The Seven-Item Eusta | | Dystunction | Questionnai | re. | | | |
| Over the past 1 month, how much has each of the following been a problem for you? | No Pr | oblem | Moderate Problem | | | Severe Problem | |
| 1. Pressure in the ears? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 2. Pain in the ears? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 3. A feeling that your ears are clogged or "under water"? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 4. Ear symptoms when you have a cold or sinusitis? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 5. Crackling or popping sounds in the ears? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 6. Ringing in the ears? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
| 7. A feeling that your hearing is muffled? | 1 | 2 | 3 | 4 | 5 | 6 | 7 |

Conclusion

- First systemic review to describe the evidence for BET.
- No RCTs, or case-control studies on BET were identified.
- Poor study design: no absolute indication, no accurate prediction of results.
- To establish consensus of diagnostic criteria is important.
- \rightarrow the evidence of BET is poor and biased.
- → No firm conclusions can be made to identify patients who will be benefit from the procedure or to accurately predict surgical results.

Treatment for Patulous dysfunction

- •Treatment should be directed at underlying medical condition, if known
- Weight gain is rarely effective and not recommended unless underweight
- Decongestants or nasal steroid ate not effective and may worsen symptoms
- Reassurance about the benign nature
- •Medical therapy should be initiated if severe s/s lasting for more than 6 weeks

Medical management for Patulous dysfunction

- Good hydration +/- nasal saline drops as needed
- Thickening the mucus can be attempted using potassium iodine eight to ten drops in a glass of juice orally 3 times daily

Dyer RK Jr, McElveen JT Jr et al. *The patulous eustachian tube: management options* Otolaryngol Head Neck Surg. 1991 Dec;105(6):832-5

Surgical management for Patulous dysfunction

- Tympanostomy tube insertion:
- first-line surgical treatment (only effective in relieving the bothersome sensation of movements of the tympanic membrane during breathing, not autophony)
- Intraluminal catheter placement:
- An intravenous catheter sealed with bone wax is a method to block the wide eustachian tube by plugging its lumen
- Cartilage grafting:
- after failure of intraluminal catheters
- Endoscopic approaches are used for reconstruction of the patulous defect while preserving the eustachian tube's function. The technique implants conchal or septal cartilage grafts to augment the concave defect within the anterolateral wall.
- Complete occlusion of the eustachian tube + VT

Surgical Management of Patulous Eustachian Tube: A Systematic Review 1960-2014

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Objectives: Patulous Eustachian tube (PET) is a challenging clinical problem with limited medical and surgical options. The current study systematically reviews the literature to determine the safety and efficacy of surgical treatments for PET. **Data Sources:** Medline, Google Scholar, and Cochrane databases.

Methods: Studies evaluating the surgical management of PET were extracted based on defined inclusion criteria. Data including surgical techniques, outcomes, and complications were extracted and analyzed.

Results: A total o<u>f 1,616 studies</u> were retrieved from the initial search. Of these, 14 studies comprising a total of 226 patients (253 sides) met inclusion criteria and were evaluated for surgical techniques, patient outcomes, and complications. As defined by the Oxford Center for Evidence-Based Medicine (Oxford, UK), all studies were classified as level 4 evidence. The most commonly reported techniques were ET plugging (3 studies), PE tube placement (2 studies), and suture ligation (2 studies). Postoperative follow-up ranged from 2 to 60 months (mean, 20.6 months). Outcome measures varied significantly between individual studies, with overall symptom improvement reported between 22% and 100% (mean 72.4%; 95% CI, 62.5%–81.2%). A low incidence of minor complications was reported in nine of 14 studies.

Conclusions: Current literature evaluating the surgical management of PET is limited and comprised entirely of level 4 studies. Comparisons between techniques were not possible due to the small number of studies and variable outcome measures. Future larger studies evaluating defined outcomes and quality-of-life measures are needed to determine the comparative efficacy of surgical treatments for this challenging condition.

Key Words: Patulous eustachian tube, eustachian tube dysfunction, autophony, eustachian, patulous, outcomes, surgery.

Laryngoscope, 00:000-000, 2015

Introduction

- Standardized treatment algorithms have yet to be established.
- Treatment aim to narrow or close the ET pharyngeal orifice.
- Nonsurgical methods: weight gain, topical estrogen, and insufflation with boric or salicylic acid.
- Surgical method: injection of bulking agents, fat/cartilage plugging, ligation of the orifice, endoluminal cauterization, and hamulotomy.

Arch Otolaryngol. 1982 Nov;108(11):735-9.

Surgical treatment of patulous eustachian tube.

Virtanen H, Palva T.

Abstract

Thirteen patients (16 ears) with patulous eustachian tubes were treated with pterygoid hamulotomy combined with transposition (eight ears) or transection (eight ears) of the tendon of the tensor veli palatini muscle. Tubal function was tested preoperatively and postoperatively by sonotubometry. Prior to surgery, only one ear showed normal tubal function on swallowing at the time of the test. The follow-up period ranged from two months to five years. Nine ears yielded normal sonotubometric results during the average follow-up time of two years, while two tubes opened on swallowing, remained open for some time, and closed little by little. The remaining five tubes stayed continuously open as before surgery, and tubal symptoms were unaltered. Transection operation had been performed on two of these ears and transposition on three. The transposition or transection procedure can be used as a routine procedure for correction of patulous tubes. Good results can be expected in about 70% of cases.



Fig. 1. Article selection process for systemic literature search.

TABLE I. Characteristics and Outcomes of Included Studies. Overall symptoms

| | | Characteristics and Outc | omes of Included S | improvement: 22-100% | |
|------------------|------------------|---|--------------------|---|------------|
| Study | Patients (Sides) | Procedure | F/U, Months | Outcome Measures | % Improved |
| Vaezeafshar 2014 | 14 (23) | Endoscopic endonasal injection of hydroxyapatite | 17.5 (9–36) | Subjective improvement (complete/ significant/unchanged/worse) - autophony, fullness, popping/ clicking, pain, vertigo | 59 |
| Rotenberg 2013 | 11 (14) | Endoscopic endonasal multilayer closure (fat plugging, endolu- minal cauterization, suture ligation) | 6 | Autophony score, Postoperative audiometry | 86 |
| Ikeda 2011 | 14 (19) | Ventilation tube placement and/ or myrigotomy with ET plugging | NS | Improvement in habitual sniffing | 89 |
| Poe 2007 | 11 (14) | Endoscopic transnasal/transoral reconstruction of ET with auto- logous cartilage or alloderm | 15.8 (3–30) | Subjective improvement in autoph- ony (complete/significant/ unchanged/worse) | 93 |
| Takano 2007 | 10 (15) | Endoscopic transnasal/transoral ligation of eustachian tube | 13–27 | Symptom resolution and sonotubometry | 60 |
| Sato 2005 | 35 (42) | Trans-tympanic insertion of sili- cone plug | 38.9 (6–68) | Symptom resolution, sonotubometry and tubotympanoaerodynamography | 71 |
| Dyer 1991 | 4 (4) | Trans-tympanic ET occlusion via catheter with ventilation tube | NS | Symptom resolution | 100 |
| Chen 1990 | 46 (60) | Ventilation tube placement | NS | Symptom resolution | 53 |
| Robinson 1989 | 8 (9) | ET diathermy | 15.9 (3–36) | Symptom resolution | 22 |
| O'Connor 1981 | 7 (9) | ET cauterization with silver nitrate | NS | Symptom resolution | 78 |
| Bluestone 1981 | 4 (4) | Trans-tympanic ET occlusion via catheter with ventilation tube | Up to 36 months | Symptom resolution | 100 |
| Virtanen 1982 | 13 (16) | Pterygoid hamulotomy with transposition or transection of tensor veli palatini tendon | 24 | Symptom resolution and sonotubometry | 69 |
| Ogawa 1976 | 16 (22) | Transnasal infusion of gelatin sponge into ET | NS | Symptom resolution | 73 |
| Stroud 1974 | 3 (3) | Transpalatal transposition of tensor veli palatini | NS | Symptom resolution | 100 |

ET = eustachian tube; F/U = followup; N/S = not specified.

outcome

Proportion meta-analysis plot [random effects]



Fig. 2. Forest plot for the success rate (% of patients with improvement in symptoms) of patulous Eustachian tube surgery.

Discussion

- Patulous E-tube is a fairly rare but often frustrating clinical entity for both patient and practitioner.
- Base on this review, most surgical techniques appear to be moderately effective at addressing PE symptoms and improving patient quality of life.
- Narrow margin between symptom relief and pathologic occlusion of the E-tube.
- Limitation: lack of validated metrics and diagnostic criteria in most studies.
- First review of surgical techniques for the management of PET.
- No single technique has been found to be superior in either surgical outcomes or safety.
- To offer interventions based on experience, personal skillset and patient wishes.

Conclusion

- Difficult to treat both medically and surgically.
- Surgical intervention appears to be a safe treatment modality with moderately successful treatment outcomes.

Take home message

- No well-accepted definition of eustachian tube dysfunction.
- Clinical diagnosis, mainly base on history and physical examination.
- Dilatory dysfunction or patulous dysfunction
- Treat directed at the underlying etiology, if known.
- Treat medically at first and then surgically.
- Surgical intervention may relieve the symptoms witohout evidence base.

Thanks for your attention!

