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**Chronic otitis media in indigenous (N643)****ID: 643.2****Management of Indigenous Chronic Middle Ear Disease in North Queensland, Australia**Presenting Author: **Shane Anderson**

Shane Anderson

*The Townsville Hospital*

*Learning Objectives:* The author will describe his experience with dealing with Indigenous Chronic Middle Ear disease in Northern Queensland, Australia. The management of this condition is co-ordinated with the collaboration of the “Deadly Ears” Program. “Deadly Ears” is an outreach program where otolaryngologists and supporting staff travel to rural and remote communities to provide primary intervention and screening. Most of the surgical management in the community involves adenoidectomy and myringotomy with or without ventilation tube insertion. Some simple tympanoplasties are performed in the community. A recent audit has found the repair rates are only 50%. The Authors practice involves providing a dedicated tertiary referral center and subspecialist Otologist support for the more at risk cases that are not treated in the community. By developing a non traditional public health frame work for referral and management, we have been able to achieve a 90% attendance rate for surgical management. Considering the amount of chronic middle ear disease that we see, cholesteatoma is relatively rare. The majority of cholesteatoma disease is mesotympanic in nature with adhesive otitis media a rare finding. The majority of disease that is treated in this setting is Chronic Suppurative Otitis Media with dense granulation. This may be due to the active management that the “Deadly Ears” program provides that may change the nature of pathology. The Author treats all these cases aggressively with cartilage techniques as the mainstay of treatment. Age, air travel back to remote communities and active discharge in this setting has not made a difference to anatomical closure of disease. The author has developed a method of closure and packing that is simple for the local health care workers to manage in their remote community and allows for water exposure.

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**Chronic otitis media in indigenous (N643)****ID: 643.3****Innovation, Excellence and Pragmatism: The Challenges in the management of Chronic Suppurative Otitis Media in Aboriginal people of Northern Australia**Presenting Author: **Hemi Patel**

Hemi Patel, Graeme Crossland, Rohana O’Connell

*Royal Darwin Hospital**Learning Objectives:*

- The prevalence of Otitis Media in Aboriginal people, and the reasons for it.
- The Surgical and non Surgical challenges of Managing Otitis Media in Aboriginal people.
- Historical Data on Surgical interventions in Aboriginal Ear disease.

*Introduction:* Aboriginal people of northern Australia have the highest rates of Chronic Suppurative Otitis Media globally, and of pandemic proportions, yet good evidence to guide our treatment in this population is limited. We present the challenges, some innovative solutions, and our experience.

*Methods:* Prospective cohort study.

*Results:* We present 5 years of prospectively collected data pertaining to Tympanoplasty, Mastoid surgery and TeleOtology in Aboriginal patients of the Northern Territory of Australia.

*Conclusion:* Tympano-mastoid surgery in Aboriginal Australians can achieve similar outcomes (intact graft, recurrence rates, hearing) to traditionally studied cohorts. We emphasise the importance of close pre and post operative follow up, and a standardised intraoperative technique applicable to the unique challenges of the Aboriginal Ear.

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**CI in chronic ears (R644)****ID: 644.1****Cochlear implantation in chronic otitis**Presenting Author: **Milan Profant**Milan Profant<sup>1</sup>, Milan Profant<sup>2</sup>, Miguel Aristegui<sup>3</sup>, Bernard Frayssé<sup>4</sup>, Joachim Mueller<sup>5</sup><sup>1</sup>*Comenius University Medical Faculty,*<sup>2</sup>*Comenius University, Medical Faculty, Dept**ORL HNS, <sup>3</sup>Univ. Madrid, <sup>4</sup>Univ Toulouse,*<sup>5</sup>*Univ Munich*

*Learning Objectives:* This panel will deal with different situations in chronic otitis patients requiring cochlear implantation. Authors will discuss the principles of implantation in cholesteatoma patients doing staging or non-staging procedure. Principles of subtotal petrosectomy with ear canal closure will be discussed as another possibility how to manage these patients. Group of patients with wide open cavity is another problem to be solved in the discussion. Authors will present also changing opinion in surgery in the only hearing ear with possibility to manage unexpected deafness by cochlear implantation. Special case reports will be the subject of discussion after short communication presented by the panel members

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## CI in chronic ears (R644)

**ID: 644.2**

### Subtotal Petrosectomy for cochlear implantation in cases of Chronic Otitis Media

Presenting Author: **Miguel Arístegui**

Miguel Arístegui

*Hospital General Universitario Gregorio Marañón Madrid Spain*

*Learning Objectives:* We will show the safety of subtotal petrosectomy applied to cochlear implantation in cases chronic otitis media, to prevent future infections that might compromise the implant

Expanding indications for cochlear implantation require adaptation of surgical techniques in special cases.

The presence of chronic otitis media (relapsing acute otitis media, chronic suppurative otitis media or cholesteatoma) require special protection in cases of cochlear implantation.

Subtotal petrosectomy offers the best protection option against future infection in these cases.

Cul di sac closure of the external auditory canal, sealing of the Eustachian tube orifice and elimination of middle ear mucosa provides a secure scenario to avoid infections and risk cochlear implant explantation in the future.

Out of 41 cases in which we have used this technique we have 17 cases that were applied to chronic otitis media of the above mentioned categories. We will report on rationale, technique and complications.

Follow up is made with MRI techniques adapted to the type of implant.

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## New diagnostic method in otology (N645)

**ID: 645.1**

### Trends in genetic diagnostics of hereditary hearing loss

Presenting Author: **Ronald Pennings**

Ronald Pennings<sup>1</sup>, Celia Zazo Seco<sup>2</sup>, Mieke Wesdorp<sup>2</sup>, Ilse Feenstra<sup>2</sup>, Hannie Kremer<sup>2</sup>, Lies Hoefsloot<sup>2</sup>, Margit Schradlers<sup>2</sup>, Helger G. Yntema<sup>2</sup>

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*Introduction:* Over the past decades, many deafness genes have been identified to cause hereditary hearing impairment (HI). It therefore, has become possible to screen for these

genes in the out-patient clinic. The importance of genetic screening of HI is that patients can be counseled about the cause and prognosis of their hearing loss and effects of rehabilitation.

Hearing impairment is genetically heterogeneous and testing of several single HI-related genes is laborious and expensive. This study evaluates the diagnostic utility of whole exome sequencing (WES) targeting a panel of HI-related genes.

*Methods:* Two hundred index patients, mostly of Dutch origin, with presumed hereditary HI underwent WES followed by targeted analysis of an HI gene panel of approximately 100 genes. 206 additional patients underwent single gene testing guided by phenotype analyses.

*Results:* We found causative variants underlying the HI in 67 of 200 patients (33.5%). Eight of these patients have a large homozygous deletion involving a known HI gene, which could only be identified by copy number variation detection. Variants of uncertain significance were found in 11 patients (5.5%). In the remaining 122 cases no potentially causative variants were detected (61%). The diagnostic yield of single gene testing in the 206 additional patients was 7.6%.

*Conclusion:* The diagnostic yield for HI using WES targeting a HI gene panel is higher (33.5%) than targeted sequencing of single genes (7.6%). In our patient cohort, causative variants in *GJB2*, *USH2A*, *MYO15A*, *STRC*, and in *MYO6* were the leading causes for autosomal recessive and dominant HI, respectively. Segregation analysis of variants of uncertain significance will further increase the diagnostic yield of WES. A practical workflow for genetic testing of hereditary HI for screening in the out-patient clinic will be presented.

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## New diagnostic method in otology (N645)

**ID: 645.2**

### N645 session : A review of automated audiometry devices and portable smartphone or tablet-based hearing testing systems in otology

Presenting Author: **Allan Ho**

Allan Ho

*University of Alberta*

*Learning Objectives:* Recent advances in portable and automated hearing testing systems has enabled testing to occur outside the traditional sound treated booths. This has far reaching implications for otologists and the patients they treat. It expands the utility of these devices in the community and in the developing world where diagnostic audiology services are scarce. We aim to review automated hearing testing systems which do not require testing in traditional sound treated booths. We will discuss the evidence supporting portable automated hearing testing systems which are available on the web and those that are independent applications for smartphones or tablet computers.

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