

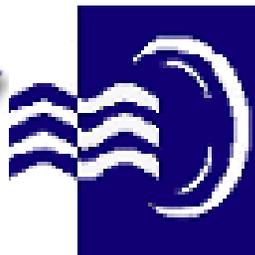
NCRAR Workshop

Ototoxicity Early Identification & Monitoring

VA Rehabilitation Research & Development
National Center for Rehabilitative Auditory Research



Site by [CDR Marketing Group](#)



Outline

- I. Learner Outcomes
- II. Overview: Basic Principles
- III. Tinnitus Monitoring
- IV. Ototoxicity Monitoring in Adults
- V. Objective Monitoring
- VI. Ototoxicity Monitoring in Children
- VII. Establishing Program

VII. Establishing Ototoxicity Monitoring Program

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Enlisting Support

- **Health Care Providers**
 - Primary Care Physicians
 - Oncology and Infectious Disease Physicians
 - Nurses (!!!)
- **Hospital Administrators**
 - Medical Policy Committee
 - Other

Enlisting Support

- Program is worthwhile, feasible, affordable
- Should be standard of healthcare b/c fits WHO definition for screening test
- Support of National Organizations
 - ASHA, AAA
- Support of healthcare providers
 - oncologists, nurses
- Other hospitals do it

Key Questions

- Purpose for identifying Ototoxic change
 - Early Identification, Hearing Loss prevention
- Target patient population
- Tests you will use
 - Sensitivity/Specificity
 - Test-retest reliability
 - Speed of test & its analysis
 - Cost of equipment
- Level of training needed
 - Audiologist
 - Nurse
 - Technician
- Program management costs

*Prevention is
the BEST form of
Rehabilitation*

Major Components

- Patient identification
- Patient status
- Equipment
- Patient Communication
- Serial Monitoring
- Behavioral/Objective measures change criteria
- Test protocol adapted to the patient (flowchart)
- Patient counseling
- Report to primary care provider (PCP)

Patient Identification

- Coordinated effort between the audiologist and health care team
- Medical staff
 - Oncologist / PCP
 - Consults
 - Nurse Practitioners
 - Pharmacist
 - Drugs dispensed and hospital formulary
- Computer generated pharmacy lists
- At Risk Patients:
 - Drugs with high ototoxicity incidence rates (see table)
 - Children
 - Elderly
 - Co-morbidities
 - Multiple ototoxic agents (including radiation)

Ototoxic Medications

<i>Antineoplastic Drugs</i>	<i>Aminoglycosides</i>	<i>Loop Diuretics</i>
Cisplatin Carboplatin Oxaliplatin Nitrogen mustard Methotrexate* Vincristine Dactinomycin Bleomycin	Gentamicin* Neomycin* Kanamycin Amikacin Streptomycin* Tobramycin* Netilmicin	Furosemide* Ethacrynic acid* Bumetanide*
<i>Other Antibiotics</i>	<i>Antimalarial Drugs</i>	<i>Salicylates</i>
Vancomycin See handout	See handout	See handout

**vestibulotoxic*

Patient Status

- Outpatient vs Inpatient
- Responsive vs non-responsive
- Booth vs Ward
- Good vs poor hearing

Patient Status

- Responsive:
 - Alert and Oriented x 4 (person, place, time, purpose)
 - Does not fatigue easily
 - Adults and older children
- Limited Responsive:
 - Alert and Oriented x 3-4
 - Fatigues easily
 - Adults and small children
- Non-responsive:
 - Alert and Oriented x 0
 - Pt sedate or Confused
 - Adults and infants
- Can oscillate between categories

Patient Status

Ward Testing Considerations

- Patient Safety, Mobility issues
 - Patient in ICU
 - Sedated
 - Isolation
 - Monitoring: Telemetry, Sugars, etc.
 - Physical limitations: Bedrest, Non-ambulatory, etc.
- Discuss with responsible attending physician or nurse
- Sound level meter recordings
- Maintain consistent test conditions

Patient Status

Good vs. Poor Hearing

- OAE particularly useful if hearing is good
 - Puretone Thresholds < 50 dB HL
- ABR if hearing loss precludes OAE use
 - Puretone Thresholds >50 dB HL at all test frequencies between 1 kHz and 10,000 Hz
 - Use if patient presents with fluctuating middle ear pathology

Test Equipment

- Audiometer / high-frequency headphones
- Immittance system
- OAE system &/or ABR system
- Maintain consistent conditions / document
- Calibration

Patient Communication

- Reasons for ototoxicity monitoring
- Prevalence of and risk factors for ototoxic hearing loss
- Possibility of late or progressive onset of hearing loss
- Realistic expectations

Serial Monitoring

- Baseline evaluation
 - ~platinum patients 1 week before or within 24 hours
 - antibiotic patients within 72 hours
 - 24 hour recheck evaluation
- Monitor evaluations
 - Performed periodically throughout treatment
 - ~platinum patients: *prior to each dose*
 - Antibiotic patients: 1-2 times per week
 - Re-test if significant changes are noted
- Post-treatment evaluations
 - Immediate post-drug evaluation
 - 1, 3, & 6 month follow-up evaluation
 - Continue to test if ASHA-significant changes are noted until hearing stabilizes

Deciding When to Monitor

- According to drug regimen
 - Chart review
 - Chemotherapeutic: Review hem/onc chart note for specific treatment regimen (i.e. Day 1, 8, 22, 29)
 - Antibiotics: Typically infused every day, sometimes 2-3 times per day (i.e. BID or Q12)
- When patient subjectively reports any difficulties
 - hearing difficulties, tinnitus, fullness, vertigo, etc...
- Audiologist responsibility to track/coordinate

Change Criteria

- ASHA Ototoxic Change Criteria

- ≥ 20 dB shift at one frequency
- ≥ 10 dB shift at 2 consecutive test frequencies
- “Response” shifting to “no response” at 3 consecutive test frequencies

- Change confirmed by retest

- DPOAE Ototoxic Change Criteria

- ≥ 6 dB shift
 - If worried about over referrals, make more stringent by requiring two consecutive test frequencies
- Change confirmed by retest
- Verify YOUR own test-retest reliability

*EACH SUBJECT
SERVES AS THEIR
OWN CONTROL*

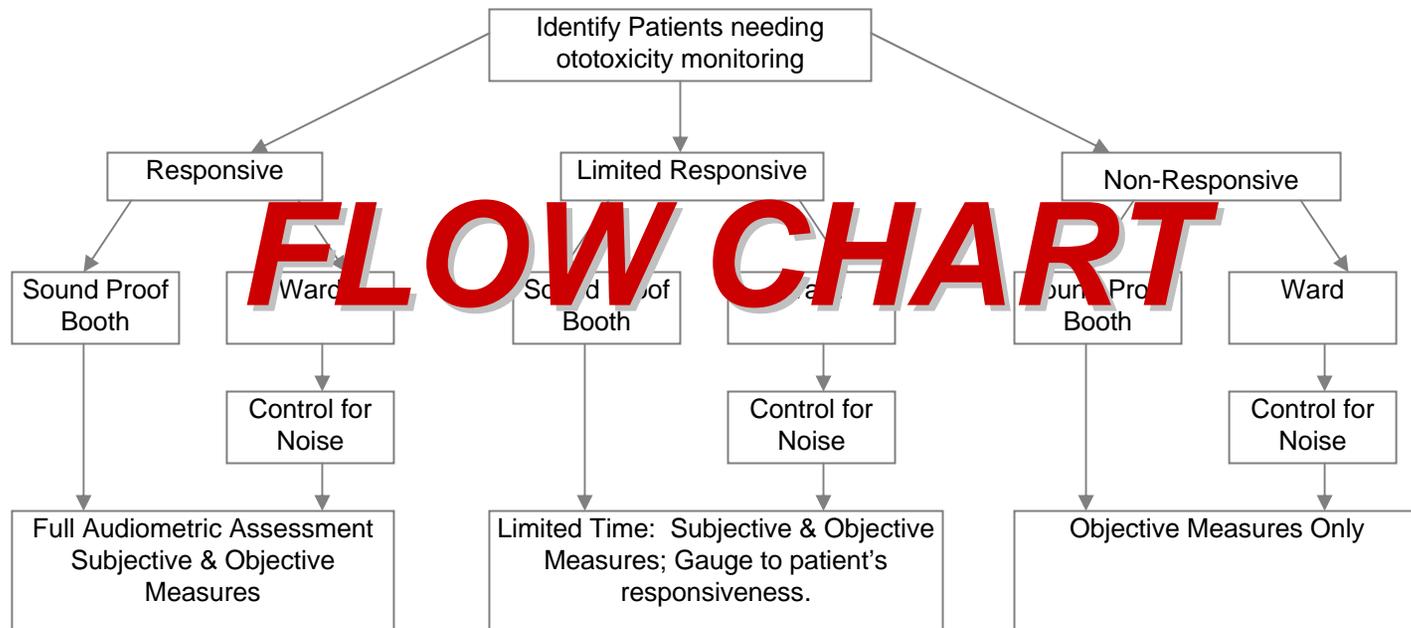
Patient Counseling

- Hearing loss
 - Potential recovery
 - Permanent
 - Realistic expectations
- Other symptoms (tinnitus, dizziness)
- Noise potentiation
 - Use ear protection
 - Up to 6 months
- Amplification
 - Caution against over-amplification
 - Work with dispensing audiologist

Report to PCP

- Test results
 - Type of test
- Behavioral hearing change noted
 - ASHA significant criteria
 - Frequencies demonstrating ototoxic change
 - Confirmed by re-test
- Objective hearing change noted
 - Exceeds established (published) test-retest reliability, which has been validated in your clinic
 - Confirmed by re-test
- Other symptoms
 - Dizziness
 - Tinnitus

Testing Protocol



Conclusion

- Ototoxicity causes high burden disorders
 - Hearing loss, tinnitus, dizziness
 - Incidence may justify cost
 - Individuals at high risk can be targeted
- Test procedures exist that reliably identify ototoxic hearing loss
- Identifying ototoxicity can improve outcome of care



<http://www.ncrar.org/home.htm>