

Infant assessment using the auditory brainstem response (ABR) and auditory steady-state-response (ASSR): Updates for clinical practice



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# **Overview of brainstem ABR & ASSRs**

**EEG parameters** 



**Estimation of infant hearing thresholds** 

**Isolation of test cochlea** 

**Clinical implications/ Future research needed** 



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# Overview

Clinical goal for infant ABR & ASSR testing?

- Identification of hearing loss
  - -- Air-conduction (AC) thresholds within normal limits?
  - -- AC thresholds elevated?
- If AC thresholds elevated, estimate bone-conduction (BC) thresholds
  - -- type of hearing loss
  - -- degree of conductive loss if present
- When hearing loss is identified, <u>frequency- & ear-specific</u> <u>thresholds</u> estimated to plan intervention services



What are ABRs?

Evoked potentials elicited to transient stimuli analyzed in the time domain: amplitude & latency characteristics of peaks/troughs in waveform

#### For threshold estimation:

- Wave V tracks close to threshold levels
- Subjectively interpret presence/absence of response using visual replicability with some objective tools (e.g., residual noise, SNR)
- $\circ~$  Assess one ear and one frequency at a time
- Most commonly used clinical method to identify hearing loss in infants



#### What are ASSRs?

- Evoked potential that is <u>repetitive</u> in nature & is analyzed in terms of its <u>frequency components</u> rather than its waveform
- For high enough rates, a "sinusoidal" response is elicited with a frequency that matches the presentation or "modulation" rate

#### Amplitude maxima in adults (reviewed in Picton et al., 2003)

- 70-110 Hz modulation rate: 1<sup>o</sup> brainstem response (Picton et al., 2003)
- ~40 Hz modulation rate: 1<sup>0</sup> cortical & brainstem (Herdman et al, 2002)
- > Most research and clinical applications for <u>infants</u>
  - -- 40-Hz smaller in sleep in infants versus adults (Picton et al., 2003)
  - -- 80-Hz or "brainstem" most of research & today's focus!

Single- & multiple-ASSRs presented to two ears simultaneously
 -- depends on equipment available (focus on multiple ASSRs)



- Why consider ASSRs for the clinic when we have brief-tone ABRs?
- -- brief-tone ABRs require considerable training & skill to interpret:
- Visual replicability of wave V? Absence of response? Waveform too noisy to interpret? Amplitude & latency features across test conditions?





Large pediatric centres: skilled, experience clinicians are available for ABR testing and do an excellent job!

**Practical challenges:** 

- (i) New clinicians
- (ii) Clinicians with low infant-ABR case loads
- (iii) Countries or regions within countries with fewer resources for training
  - -- face difficulties conducting/interpreting AC & BC ABRs
- Solutions:
- (i) Method that requires less training & skill- ASSR?
- (ii) Telehealth ABR (emerging but still requires skilled clinician)



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- > Why ASSRs?
- (i) frequency-specific stimuli
  - growing # of choices (advantage or disadvantage?)
- (ii) response presence/absence is statistically determined
  - objective rather than subjective interpretation of waveforms
- (iii) multiple stimuli can be presented to both ears simultaneously
  - efficient use of clinical time (2/3 time of ABR)

[van Maanen & Stapells, 2009]

#### **One example of ASSR analysis**

Comparison of response amplitude @ modulation rate to surrounding noise frequencies: F statistic (p < .05) (for review see Picton et al., 2003)





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EEG parameters - very briefly





Can record EEG ipsilateral & contralateral to mastoid stimulated to assist with isolation of the test ear (more later in presentation)







- can avoid post-auricular muscle response







Can record EEG ipsilateral & contralateral to mastoid stimulated to assist with isolation of the test ear (more later in presentation)



# AC methodology

-- nothing new



preferred- better for setting hearing aid targets







# BC methodology: ASSR threshold data (Small et al., 2007)

Bone oscillator coupling method in infants



No significant differences (with training)

**Recommend: <u>Either</u>** 

#### **Bone oscillator placement**

Recommend: <u>"T" position</u>



#### Occlusion effect (OE): earphones in or out during infant BC testing?





Young infants (< 12 months) - negligible OE

Older infants (1-2 years) - emerging occlusion effect

(Small et al., 2007, Small & Hu, 2011)

Recommend: <u>0-1 year: leave earphones in</u> <u>1-2 years +: remove earphones (conservative)</u>



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# Stimuli



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#### Many types of "frequency-specific" ASSR stimuli



# **Estimation of infant hearing thresholds**



### **Definition of terms currently used for ABR** (BCEHP, 2012) *Normal behavioural threshold*:

• 25 dB HL

#### Normal ABR maximum level:

• ABR presentation level at which the majority of normalhearing infants have a response present

normal?

response must be present at normal ABR (dB nHL) max

#### eHL correction:

• Correction factor used to estimate behavioural hearing threshold (dB HL) from the ABR threshold

ABR threshold \_\_\_\_\_ eHL correction \_\_\_\_\_ = (dB nHL)

estimated behavioural threshold (dB HL)

# Normal ABR maximum levels & eHL correction for infants Air- and bone-conduction ABR

	500	Hz	1000 Hz		2000 Hz		4000 Hz	
	AC	BC	AC	BC	AC	BC	AC	BC
BC EHP								
Normal ABR Max	35	20	35	na	30	30	25	na
(dB nHL)								
Range in literature	30-35	20	30-35	na	20-30	30	20-25	na
BC EHP	10	5	10	na	5	5	0	na
eHL correction (dB)	10	5	Ĩ	na	5	5	Ŭ	na
Range in literature	10-15	-5	5-10	na	0-5	5	-5-0	na

(BC-EHP 2012, 2015; Small & Stapells, Ch. 21, 2017)

#### Mean AC & BC ASSR thresholds across 11 infant & 10 adult studies AM; AM/FM; AM<sup>2;</sup> COS<sup>3</sup>



AC: low > high frequencies
BC: low < high frequencies</li>
Maturational air-bone gap

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(Lins et al, 1996; Cone-Wesson et al., 2002; John et al., 2004; Rance et al., 2005; Swanepoel & Steyn, 2005; Luts et al., 2006; Rance & Tomlin, 2006; van Maanen & Stapells, 2009; Ribeiro et al., 2010; Casey & Small, 2014; Valeriote & Small, 2015)

AC & BC: similar across frequency -- tendency for BC 500 Hz to be greater than other frequencies

(reviewed in Tlumak et al., 2007)



# How well do AC ASSRs predict the audiogram in infants?

#### AC multiple ASSR versus AC behavioural thresholds/brief-tone ABR

- **Correlation coefficients:**
- <u>Adult</u>
- ➤ .70-.85 for 500 Hz
- > .80-.95 for 1000-4000 Hz (for review see Tlumak et al., 2007)

#### <u>Infant</u>

- > .97 @ 500-4000 Hz (includes profound loss with "no response")
- .77-.89 @ 500-4000 Hz (excludes "no responses")
  - (Van Maanen & Stapells, 2010)

# Normal ASSR maximum levels & eHL correction for infants *Air-conduction ASSR*

#### Preliminary & conservative!

 $\Delta M/FM$ 

$COS^3$	500	Hz	1000	Hz	2000 Hz		4000 Hz	
AM <sup>2</sup> (Ages:0-79 ms)	AC		AC		AC		AC	
10 studies								
Normal ASSR Max	40-50		40-45		40		40	
(dB HL)								
Range in literature	40-52		30 to >50		30-50		28-44	
6 studies**								
eHL correction (dB)	10-20		10-15		10-15		5-15	
Range in literature	-3 to 20		0-17		0 - 6		-3 - 14	

(reviewed in Small & Stapells, Ch. 21, 2017: \*Lins et al, 1996; John et al., 2004; Rance et al., 2005; Swanepoel & Steyn, 2005; Luts et al., 2006; Rance & Tomlin, 2006; van Maanen & Stapells, 2009; Ribeiro et al., 2010; Casey & Small, 2014; Valeriote & Small, 2015;\*\*Rance & Briggs, 2002; Hanh et al., 2006; Luts et al, 2006; wan Maanen & Stapells, 2010; Rodrigues & Lewis, 2010; Chou et Al., 2012)



# How well do BC ASSRs predict the audiogram in infants?

### BC multiple ASSR versus AC behavioural thresholds/brief-tone ABR

- **Correlation coefficients:**
- Adult (sensorineural & simulated)
- ➤ .71 for 500 Hz
- .84-.94 for 1000-4000 Hz (Ishida, Cuthbert & Stapells, 2011)
- Adult BC-ASSR data is promising

#### <u>Infant</u>

> No correlational data available

Valeriote & Small (in prep): Infant: normal hearing versus mild conductive loss at 500 Hz

AC & BC ASSR data fall within ABR normal maximum levels

 AC: trend for elevated ASSR thresholds
 but overlap for NH and mild CHL for ASSR

BC: CHL and NH did not differ significantly as expected

Valeriote & Small (in prep)



# Case 1: Adult with asymmetric conductive loss (stapes fixation bilaterally, poor surgical outcome left)

Small, unpublished

# Normal ASSR maximum levels & eHL correction for infants Bone-conduction ASSR Preliminary & conservative!

	500	Hz	100	0 Hz	200	0 Hz	400	0 Hz
AM/FM AM <sup>2</sup>		BC		BC		BC		BC
8 studies (0-24 mos)								
Normal ASSR Max		30		20		40		30
(dB HL)								
Range in literature		30-40		10-30		30-40		10-40
BC EHP								
eHL correction (dB)		lid		lld		lld		IId
Range in literature		na		na		na		na

(Small & Stapells, Ch. 21, 2017)

## More recently...chirp stimuli

Why chirps?

• <u>Clicks/brief-tones</u> do not compensate for temporal delays in frequency contributions when basilar membrane is activated

**Consequences**:

- (i) superimposition of responses from individual nerves is suboptimal
- some destructive interference-- reduces ABR/ASSR amplitudes

- <u>Chirps</u> are designed to compensate for basilar membrane travelling-wave delays (> compensation for low frequencies) (Shore & Nutall, 1985; Dau et al., 2000; Elberling et al., 2007)
  - e.g., \*CE-chirp (Broadband) (Elberling et al., 2007; Elberling, Don et al., 2007)
  - -- low-frequencies precede high-frequencies
  - -- maximizes synchrony of neural firing across frequencies
  - -- larger response amplitudes result

\* registered trade name



#### For stimuli < 60 dB nHL:

> ABR amplitude CE-chirp versus click: 1.5-2 X larger

(e.g., Dau et al., 2000; Elberling & Don, 2008)

Advantage for screening purposes in infants (e.g., Cebulla et al., 2014)

#### For CE-Chirps @ higher intensities (> 60 dB nHL):

- there is upward spread of excitation by the individual frequency components of the chirp
- as level increases, area of cochlear excitation broadens & desynchronization occurs
- chirps can be modelled with different delays most efficient design is to increase the delay for decreasing stimulus levels
  - -- stacked ABR data was used to model CE-Chirps

(Fobel & Dau, 2004; Elberling & Don, 2008; Elberling et al., 2010)

<u>Note</u>: refer to papers on Level Specific CE-Chirps for more information on optimizing CE-Chirps (e.g., Kristensen & Elberling., 2012)

 Refer to the literature for more on "broad-band" CE-Chirp thresholds in infants & adults with normal hearing and hearing loss (ABR & ASSR)- presenting only more frequencyspecific data today Chirps for frequency-specific threshold estimation? Specifically in infants?

**Optimize response amplitudes?** 

**Increase detectability?** 

**Reduce testing time?** 

#### Accurately predict behavioural thresholds?

narrowband chirp stimuli were designed (NB CE-Chirps)
 -- same compensation for travelling-wave delay but for much narrower range of frequencies

<u>Question</u>: Is the NB CE-Chirp better than brief-tones for <u>ABR</u> or other stimuli for <u>ASSRs</u> @ estimating threshold? Is the amplitude advantage also present for the NB CE-Chirps? For all test levels?

# <u>Note</u>: Today's focus is frequency-specific stimuli & brainstem ABR/ (80-Hz) ASSR

# **NB CE-Chirps**

(Elberling, 2011)



Stimulus timing NB-Chirps temporally adjusted

If stimulus onset is not adjusted, wave V latencies for the low frequencies are shorter for NB CE-Chirps than brief tones due to stimulus design (e.g., Rodrigues et al., 2013)

Stimulus onset is adjusted to expected response latencies (same delay model for all levels) (e.g., Interacoustics, 2012)

# <u>ABR</u> NB CE-Chirps versus brief tones NH infants

#### ABR studies in NH infants: NB CE-Chirps versus brief tones

Study	Age (wks)	f (Hz)	Optimal stimulus level (dB nHL)	Level (dB nHL)	Ear phone	Greater Amp	% with >AMP	> AMP with increase in level
Ferm et al. 2013/15	0-12	500	≤ 30	10-50*	TDH-39	<b>1.3 x</b>	71	Νο
N=30-39								

#### ABR studies in NH infants: NB CE-Chirps versus brief tones

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Ferm et al. 2013/15	0-12	500	≤ 30	10-50*	TDH-39	<b>1.3 x</b>	71	Νο
N=30-39		1000	≤ 40	10-45	TDH-39	1.6 x	98	Yes
		2000	≤ 40	0-40	TDH-39	1.5 x	93	Yes
		4000	≤ 45	0-40	TDH-39	1.6 x	100	Yes

NB CE-Chirp responses present at levels 10 dB < brief tones for 52-62 % of infants tested— less so for 500 Hz\*

#### ABR studies in NH infants: NB CE-Chirps (NBCh) versus brief tones (BT)

Study	Age (wks)	f (Hz)	Optimal stimulus level TDH39 (dB nHL)	Level (dB nHL)	Ear phone	Greater Amp?
Rodrigues et al., 2013	4-12	500	≤ 30	80	ER3-A	No BT>NBCh*
				60	ER3-A	No NBCh~BT*
				20-40	ER3-A	Yes

500 Hz: only advantage seen is for low levels; no advantage for mid levels and <u>disadvantage</u> for high levels\*

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Rodrigues et al., 2013	4-12	500	≤ 30	80	ER3-A	No BT>NBCh*
				60	ER3-A	No NBCh~BT*
				20-40	ER3-A	Yes
		1000 - 4000	≤ 40-45	80	ER3-A	No NBCh~BT*
		1000 - 4000	≤ 40-45	20-60	ER3-A	Yes

- 500 Hz: only advantage seen is for low levels; no advantage for mid levels and <u>disadvantage</u> for high levels\*
- > 1000-4000 Hz: advantage for low/mid levels but not for high level\*

#### ABR studies in NH infants: NB CE-Chirps versus brief tones

Study	Age (hours)	f (Hz)	Optimal stimulus level TDH39 (dB nHL)	Level (dB nHL)	Ear phone	Greater Amp?
Cobb & Stuart, 2016	11-104 N=20-22	500	≤ 30	30-60	ER3-A	No*



#### ABR studies in NH infants: NB CE-Chirps versus brief tones

Study	Age (hours)	f (Hz)	Optimal stimulus level TDH39 (dB nHL)	Level (dB nHL)	Ear phone	Greater Amp?
Cobb & Stuart, 2016	11-104 N=20-22	500	≤ 30	30-60	ER3-A	No*
		1000 2000	≤ 40	30-60	ER3-A	Yes
		4000	≤ 40	30-45	ER3-A	Yes
		4000	≤ 45	60	ER3-A	No*

- 500 Hz: no advantage for low-mid levels\*
- > 4000 Hz: no advantage for mid levels\*
- > 4000 Hz: advantage for low levels
- > 1000-2000 Hz: advantage for low & mid levels

#### Across NH infant <u>ABR</u> studies: NB CE-Chirps versus brief tones for frequency-specific threshold estimation

## <u>AC stimuli</u>

**Optimize response amplitudes?** 

- NH : Yes, on average, amplitudes larger; greatest advantage 1000-4000 Hz for levels < 60-80 dB nHL</li>
- However @ 500 Hz: amplitude disadvantage at 80 dB nHL, no advantage at 45-60 dB nHL, & may or may not show an amplitude advantage at 20-30 dB nHL

**Increase detectability?** 

- NH infants: Yes, ~ 52-62% of cases
- Hearing loss: No data but expect less benefit with increase in stimulus level

## AC stimuli cont'd

#### **Reduce testing time?**

- NH infants: Yes
- Milder losses: No ABR data but probably (more so at 1-4 kHz)
- Moderate/severe losses: No ABR data: maybe a disadvantage?

#### Accurately predict behavioural thresholds?

- Lack of ABR data for NB CE-Chirps in infants with hearing loss
- Ferm et al. recommended correction factors to dB eHL for NB CE-Chirps as 5 dB < brief tone correction factors</li>
- Not yet verified in a clinical population

# <u>BC stimuli</u>

- No published ABR data for NB CE-Chirp stimuli -- NH infants or infants with hearing loss
- Currently for clinic- no date to support estimation of frequencyspecific BC thresholds using these stimuli
  - -- only one research group has looked at AC and BC ABRs in infants -- only for broadband CE-Chirps (Cobb & Stuart, 2016a, b& c)
  - -- we know there are frequency-dependent differences in infant BC thresholds so need to investigate BC ABRs to NB CE-Chirps

#### <u>ASSRs</u>

# NB CE-Chirps versus brief tones ABR/behavioural measures: NH infants & infants with hearing loss Adults with hearing loss

#### **ADULTS WITH HEARING LOSS:**

#### AC ASSR NB CE-Chirps & AM/FM versus behavioural

Study	Age	C	<b>Correlation coefficient</b>								
		500	1000	2000	4000						
NB CE-Chirps versus behavioural											
Lee et al. (2016)	ADULTS N=19	.67*	.87	.76	.91						
	AM/FM sinusoid versus behavioural										
Lee et al. (2016)	ADULTS N=19	.67*	.81	.68	.80						

Note: all correlation coefficients were statistically significant

Larger amplitudes for NB CE-Chirps for 1000-4000 Hz similar to ABR

- Slightly stronger correlation coefficients compared to AM/FM
   > 500 Hz
- > no advantage at 500 Hz\*

#### **NORMAL HEARING INFANTS**

#### AC ASSR NB CE-Chirps versus brief-tone ABR

Study	Age (mos)	Level at whic PRESENT res (dBnHL)	Testing time (min)			
		500	1000	2000	4000	
		NB CE-Ch	nirps			
Rodrigues & Lewis (2014)	NH neonates N=30	35*	28	13	15	<b>21±5</b>
		Brief-tone	e ABR			
Small & Stapells, 2017 <sup>a</sup> Janssen et al. (2010) <sup>b</sup>	NH (across studies)	<b>30-35</b> ª*	<b>30-35</b> ª	<b>20-30</b> <sup>a</sup>	<b>20-25</b> <sup>a</sup>	<b>49-58</b> b

normal maximum levels: lower for NB CE-Chirps 1000-4000 Hz
 no advantage at 500 Hz\*

#### **INFANTS WITH HEARING LOSS:**

#### AC ASSR NB CE-Chirps versus brief-tone ABR

Study	Age (mos)	Correl	Correlation coefficient*			Difference	dB nHL to dB eHL	Testing in min
		500	1000	2000	4000			
NB CE-Chirp ASSR versus brief-tone ABR								
Michel & 4-22 Foldager N=67	4-22	.90	.90	.96	.95	500-4000 Hz	5 dB <	
	N=67	+ infan	its with h	nearing lo	ss only	Mean: - 4-5 dB	ABR	
(2016) <sup>+</sup>								
		NB	CE-Ch	irp ASSI	R versi	s behavioural		
Venail et al. (2014)	2-12 N=32	.87	.91	s .93	.86	500 & 2000 Hz Mean - 4 dB 1000 & 4000 Hz Mean: -1-2 dB		23±16
		В	rief-to	ne ABR	versus	behavioural		
Stapells et	4	.94ª		<b>.95</b> <sup>a</sup>	.97ª			<b>49-58</b> <sup>b</sup>
al., 1995 <sup>a</sup> , Janssen et al. (2010) <sup>b</sup>	N=188		simil	ar				shorter testing time

Across infant <u>ASSR</u> studies: NB CE-Chirps versus AM/FM tones for frequency-specific threshold estimation

#### <u>AC stimuli</u>

**Optimize response amplitudes? Increase detectability?** 

- No amplitude data provided in the two infant studies available (1 NH and 1 hearing loss)
- NH infants: Normal levels were slightly lower for 1000-4000 Hz but not 500 Hz (Rodrigues et al.; 2014)
- Hearing loss: One study with correlation data showed strong correlations but slightly less compared to brief-tone ABR data

#### **Reduce testing time?**

• NH and hearing loss: Yes, significantly

## Accurately predict frequency-specific behavioural thresholds?

#### Potential issues:

- (i) Limited ASSR data for NB CE-Chirps in infants with hearing loss
- No recommended correction factors to dB eHL

(ii) No place/frequency specificity studies to explore whether the advantage of using NB CE-Chirps is due to compensation for basilar membrane travelling-wave delays OR stimulation of broader portion of the basilar membrane (i.e., less frequency specific)

# <u>BC stimuli</u>

• No published ASSR data for BC NB CE-Chirp stimuli in infants

#### **Spectral analysis: NB CE-Chirps versus 2-1-2 brief tones**

Adjekum, Chan, Stapells & Small, in prep - also Cobb & Stuart (2016)

> Work in progress in my lab: - investigating place specificity of NB CE-Chirps

# Table 1: Characteristics of brief tone spectra: NB CE Chirps versus 2-1-2 tones (Interacoustics)

Nominal Frequency	0.5 KHz		1 KHz		2 KHz		4 KHz	
Stimuli	Chirp	2-1-2	Chirp	2-1-2	Chirp	2-1-2	Chirp	2-1-2
Geometric Mean	485	472	1006	973	2076	1945	3618	3764
Bandwidth	560	302	1077	560	1229	1120	3015	2024
Deviation (%)	3.03	5.64	0.63	2.74	3.79	2.74	9.56	5.90

#### Work in progress in my lab...



# Are multiple ASSRs more or less efficient than single ASSRs?



(Hatton & Stapells, 2011 & 2013)

Note: stimuli with broader spectra or higher presentation levels exhibit > interactions (Ishida & Stapells, 2012; Mo & Stapells, 2008, Wood, 2009)

> Recommend: Low-mid intensities – multiple ASSR High intensities – consider single ASSR



# What about simultaneous AC & BC multiple ASSRs?

New study from Cuba (Torres-Fortuny et al., 2016)
 -- compared ASSR amplitudes elicited to AC & BC stimuli at same time in both ears to only one mode at a time in NH infants



No significant reduction in amplitude for simultaneous AC/BC conditions; more data needed but clinical potential ...



# AC & BC ASSRs & severe-to-profound loss

- Caution: can elicit vestibular responses to high-intensity AC & BC stimuli using ABR & ASSRs
- ABR- negative wave at ~ 3 ms at 95 & 110 dB nHL due to activation of the vestibular system- not auditory in nature but easy to identify in the waveform (Stapells, 2011)
- ASSRs can also be elicited from vestibular sources- cannot be differentiated from auditory responses - no time domain waveform available
  - -- spurious responses recorded at 50-60 dB HL for BC ASSRs; 118-120 dB HL for AC ASSRs (Small & Stapells, 2004)



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# Isolation of test cochlea

# **BC ABR: Utilize ipsilateral/contralateral asymmetries**

Expected pattern for normal cochleae up to 1-2 years of age -normal hearing or conductive loss (e.g., aural atresia)

[e.g., Foxe & Stapells, 1993; Stapells & Ruben, 1989; Stapells & Mosseri, 1991]



Amplitude: contra smaller than ipsi

Latency: contra later than ipsi



#### Factors contributing to ipsi/contra asymmetries?

1. Greater IA (10-35 dB) compared to adults due to unfused cranial sutures

(Yang & Stuart 1987; Small & Stapells, 2008; Hansen & Small, 2012)

2. Infant-adult differences in positioning of neural generators



(see for review: Small & Stapells, 2017)

 Infant BC ABR/ASSRs show consistent ipsi/contra asymmetries @ near-threshold levels (adult do not) BC ABR: 500 & 2000 Hz (e.g., Stapells & Ruben, 1989)
 BC ASSR: 500 & 4000 Hz (less consistent @1000 & 2000 Hz) (Small & Stapells, 2008; Small & Love, 2014)

> more research needed for ASSRs to determine accuracy in infants with hearing loss



What if ipsi/contra asymmetries in BC ABR or ASSRs are ambiguous?

#### > MASK!

Main reason masking not routinely used clinically for infant BC ABRs: -- effective masking levels (EMLs) for BC ABR stimuli in young infants have not been measured directly

[these data are currently being collected in my lab (Lau, M.Sc. thesis)]

> We have estimated EMLs for BC ASSRs using binaural AC masking

(Hansen & Small, 2012; Small, Smyth & Leon, 2014)



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#### Recommended EMLs (dB SPL) for BC ASSR stimuli presented at 35 dB HL

	Frequency (Hz)									
	500	1000	2000	4000						
Infant	م 81 <sup>-</sup>	15 68 <b>7</b>	<mark>5</mark> 59	45 T	-10					
Adult	66	* 63	* 59	55 -	*					

\* Significant infant <u>minus</u> adult EML difference (dB)

Frequency-dependent infant-adult differences in EMLs except at 2000 Hz

(Hansen & Small, 2012; Small, Smyth & Leon, 2014)



# **Threshold: Clinical Implications/Future Research**





# **Threshold: Clinical Implications/Future Research**



no BC NB CE-Chirp data



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# **More Future Research needed**







