

The role of functional near-infra-red spectroscopy (fNIRS) in early intervention for infants with hearing impairment

Colette McKay

...and EarGenie team



**Bionics
Institute**

The EarGenie Team

BI TEAM

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Dr Demi Gao (Research Engineer)

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DEVELOPMENT/COMMERCIALISATION

NIRGenie company: CEO Dr Jeffry Ng

Prototyping partner: Design + Industry

Our aim...closing the gaps

...between **identification** of hearing loss in an infant and the **accurate fitting of the optimal hearing instrument**

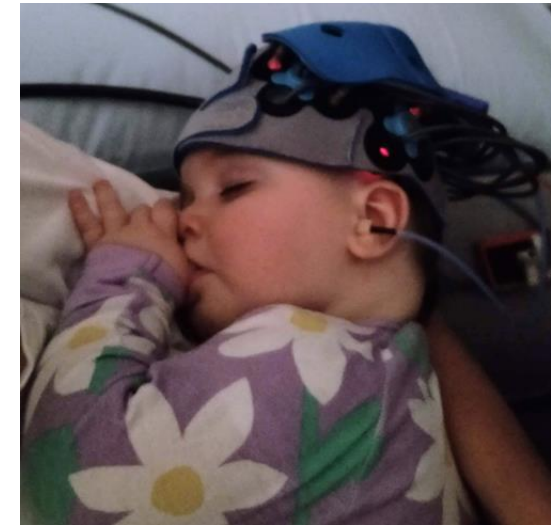
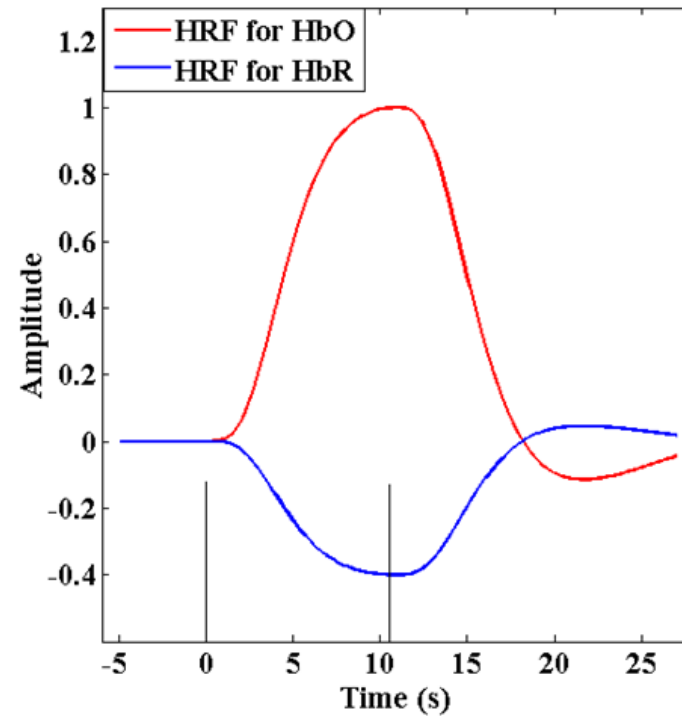
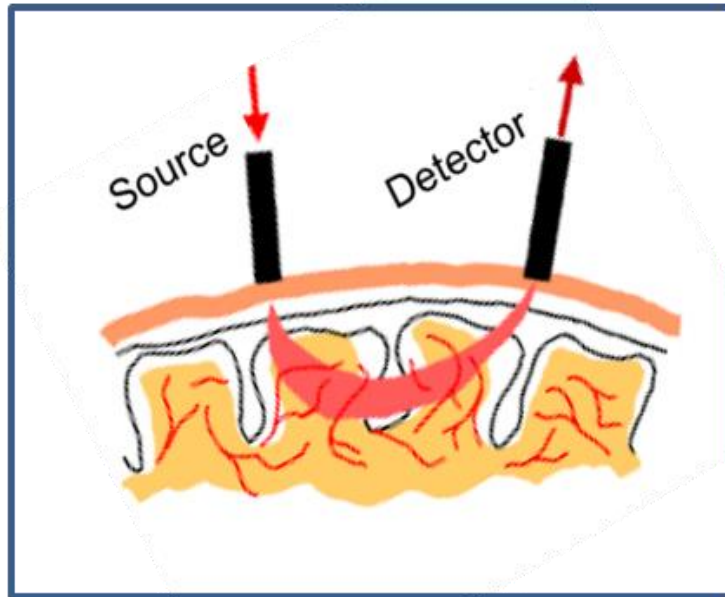
- **Auditory neuropathy**...what is my functional hearing like?
 - Sensitivity and discrimination
- **All infants**...
 - is my hearing aid programmed optimally (sound detection)?
 - and would a cochlear implant be a better option (discrimination)?
- **Cochlear implants**...objective programming and evaluating early programs

What do we need to **fast track** intervention...?

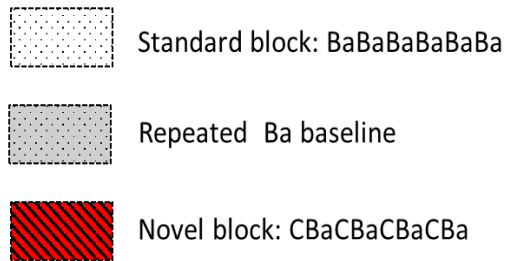
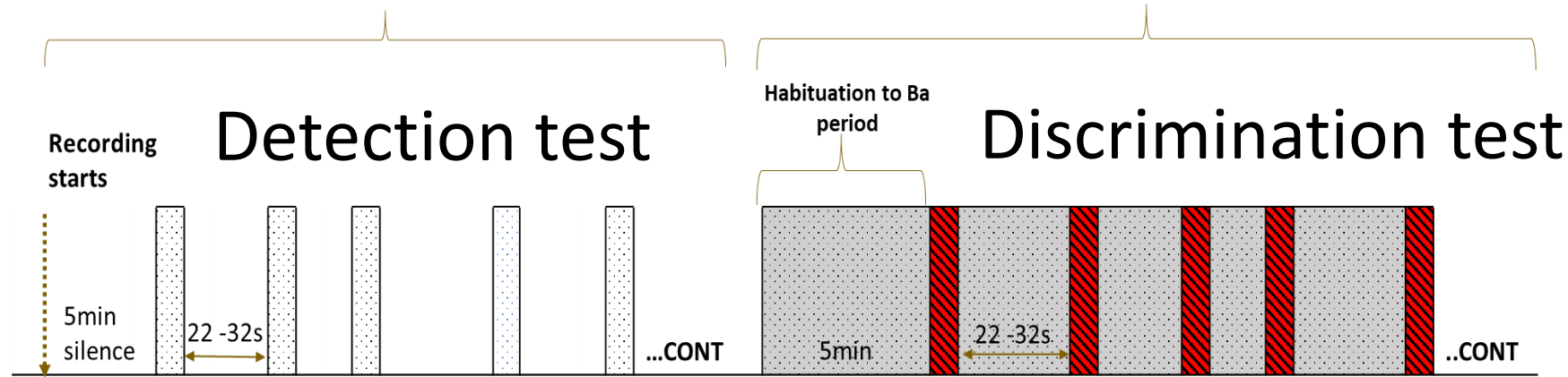
- A reliable and accurate objective test of speech sound detection and discrimination that is
 - Reliable for infants with **auditory neuropathy**
 - Suitable for **very young infants in the natural sleep state**
 - Suitable for **use in cochlear implants** (not affected by electrical artifacts or interference)

Introducing EarGenie

The hearing assessment system that uses fNIRS



Speech Detection and Discrimination

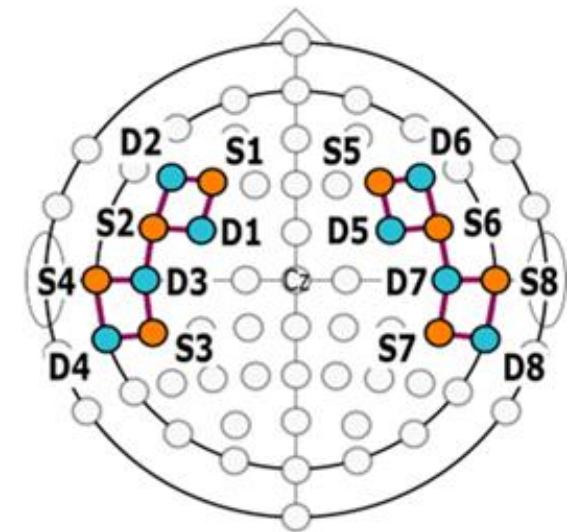


C = Contrast speech sound to be tested

In each test run, C is pseudorandomised within a set of 3 contrasts.

Set 1 contrast sounds:
Tea/Ba, Bee/Ba, Ga/Ba

Set 2 contrast sounds:
Tea/Ba, Pa/Ba, Ma/Ba

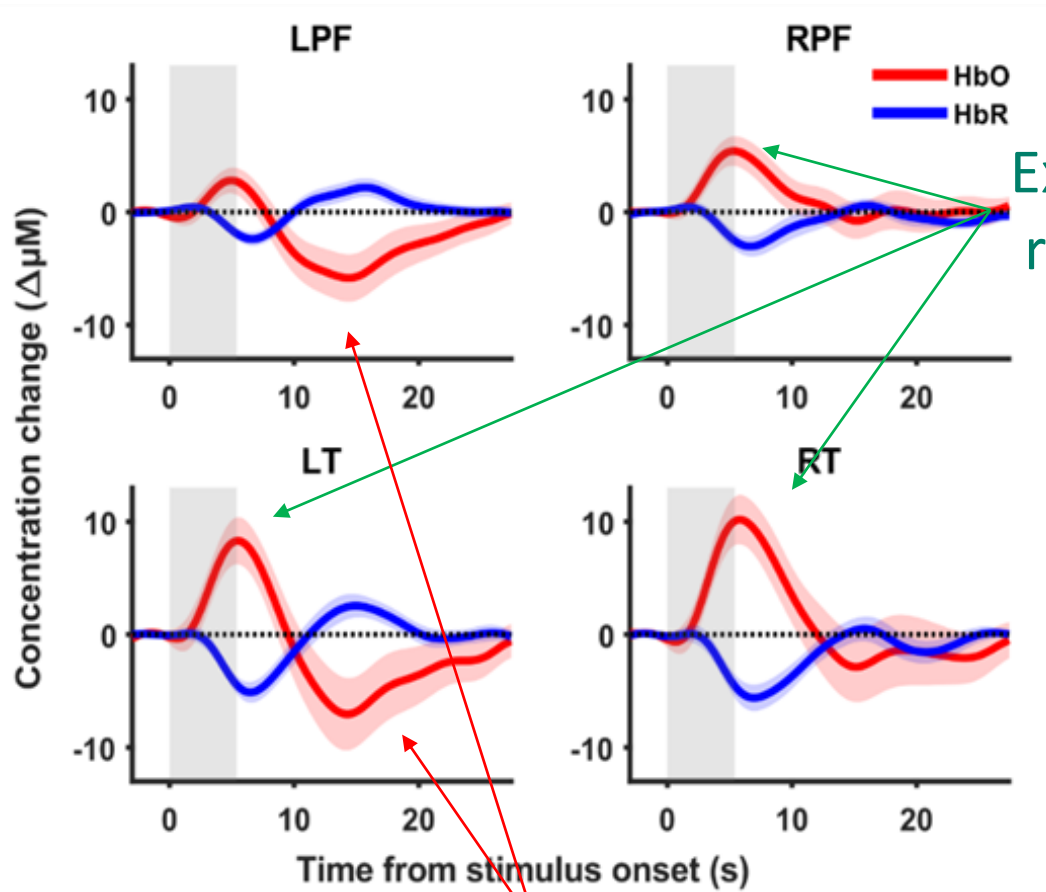


Detection response 65 dB SPL

(n = 16, normal hearing)

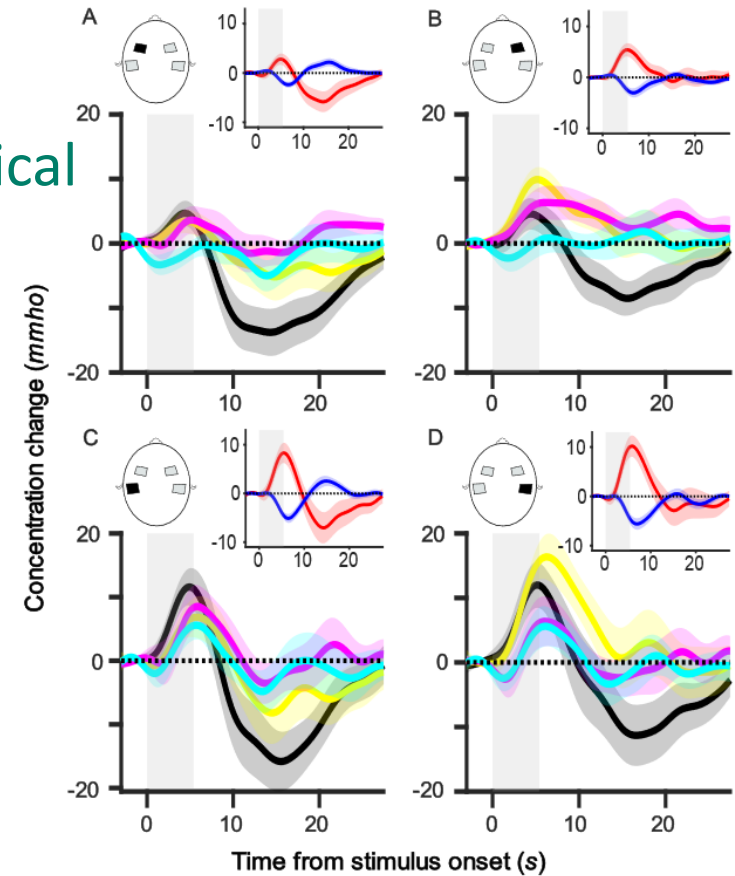
Grand averages 20 epochs per ROI

Grand averages every 5 epochs per ROI



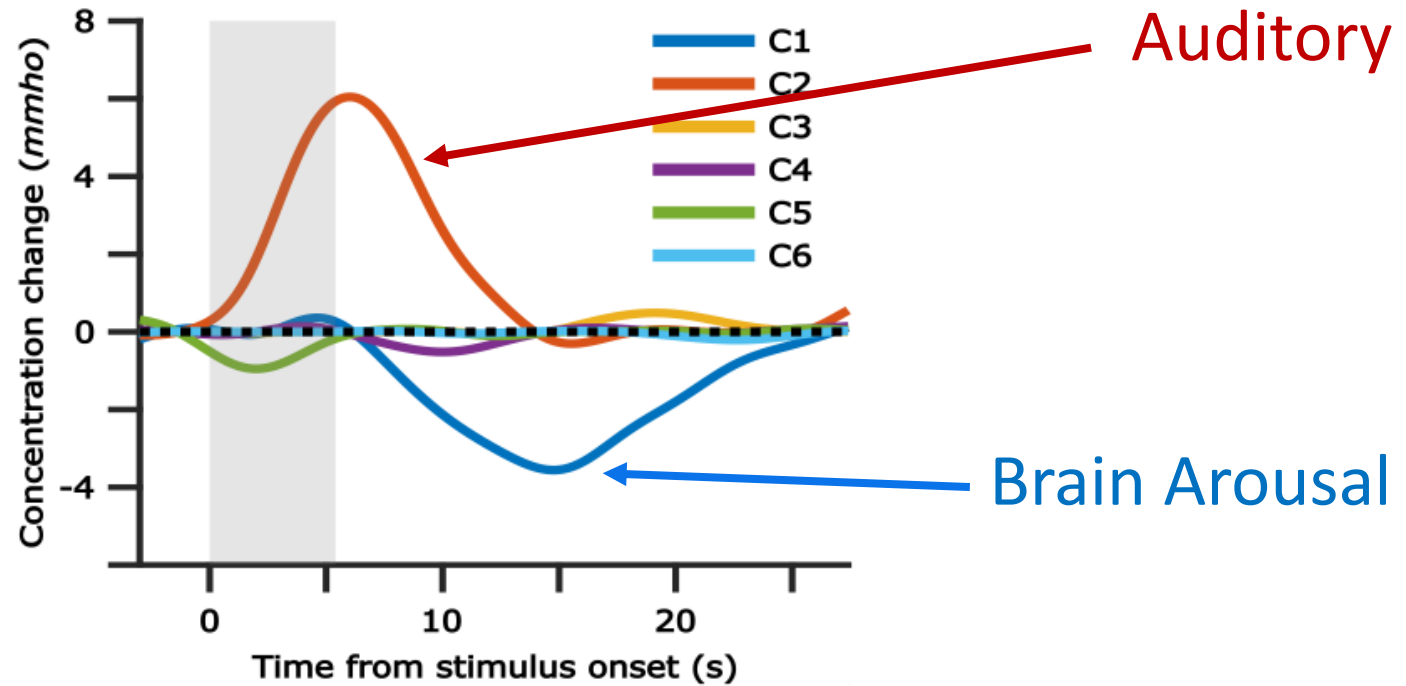
Expected canonical response

Unexpected negative component



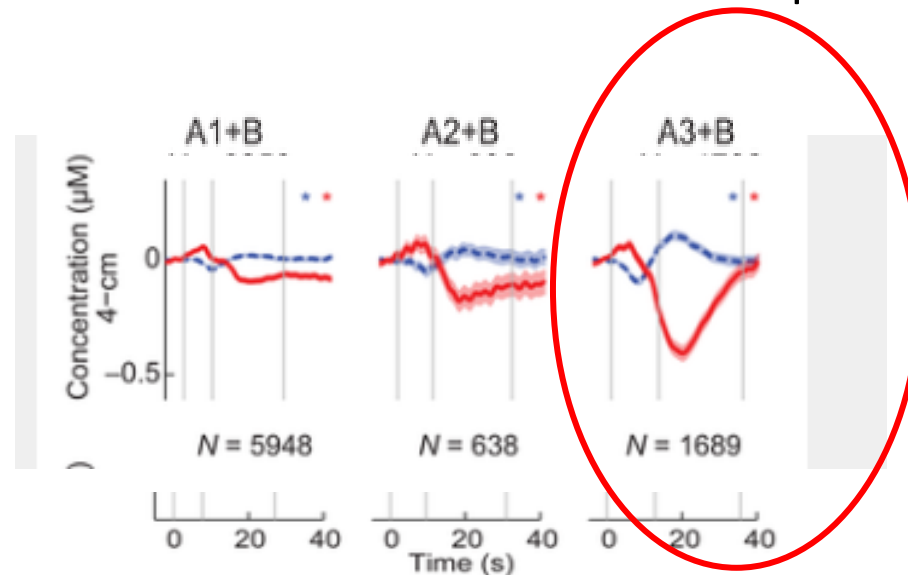
Two independent and concurrent responses to auditory stimuli (N = 16, normal hearing)

Grand average of reconstructed ICA components
(All epochs and channels used)

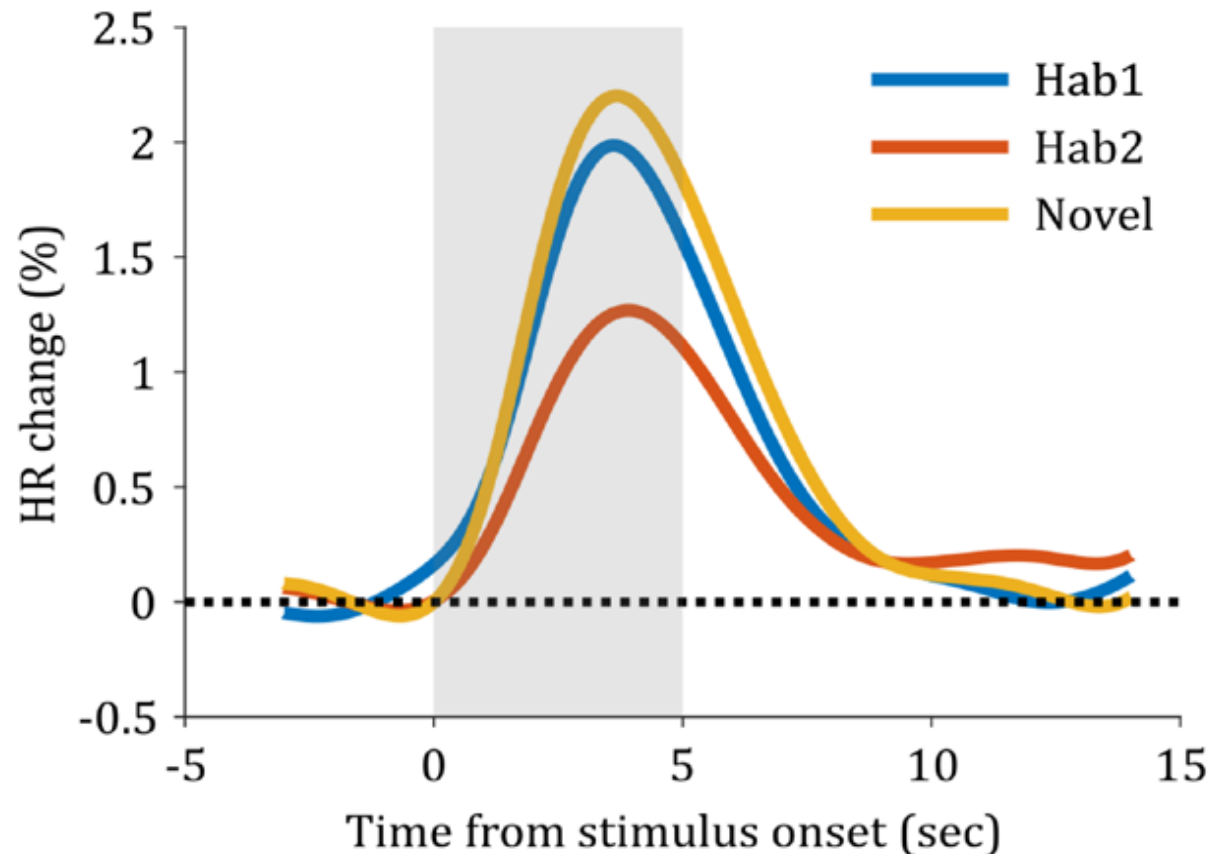


Why do we think it is a brain arousal response?

- McNamara et al J Appl Physiol (1998) (sleeping infants) tactile stimulus induced cortical arousal
- Cortical arousal defined as a sudden change in alpha activity
- Cortical arousal rapidly habituated with repeated stimuli
- Nasi et al, PLoS ONE 2012 (sleeping adults)
- Cyclic Alternating Patterns A-B occur in non-REM sleep.
- A1: Cluster of high amplitude alpha activity
- A2: transient high amplitude activity
- A3: transient high amplitude activity with desynchronization
- B: following A
- fNIRS epoched at starts of A phases



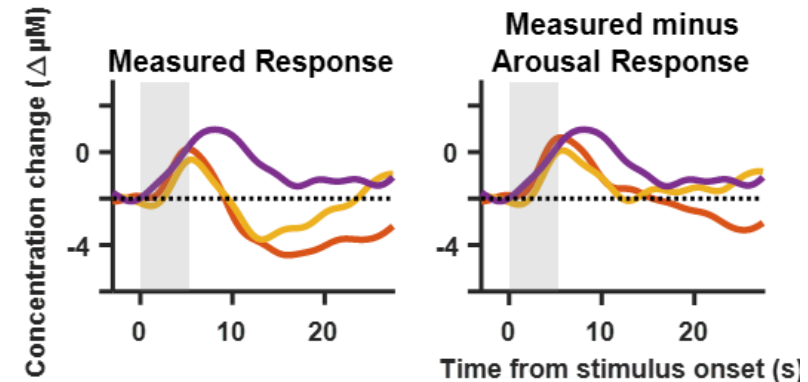
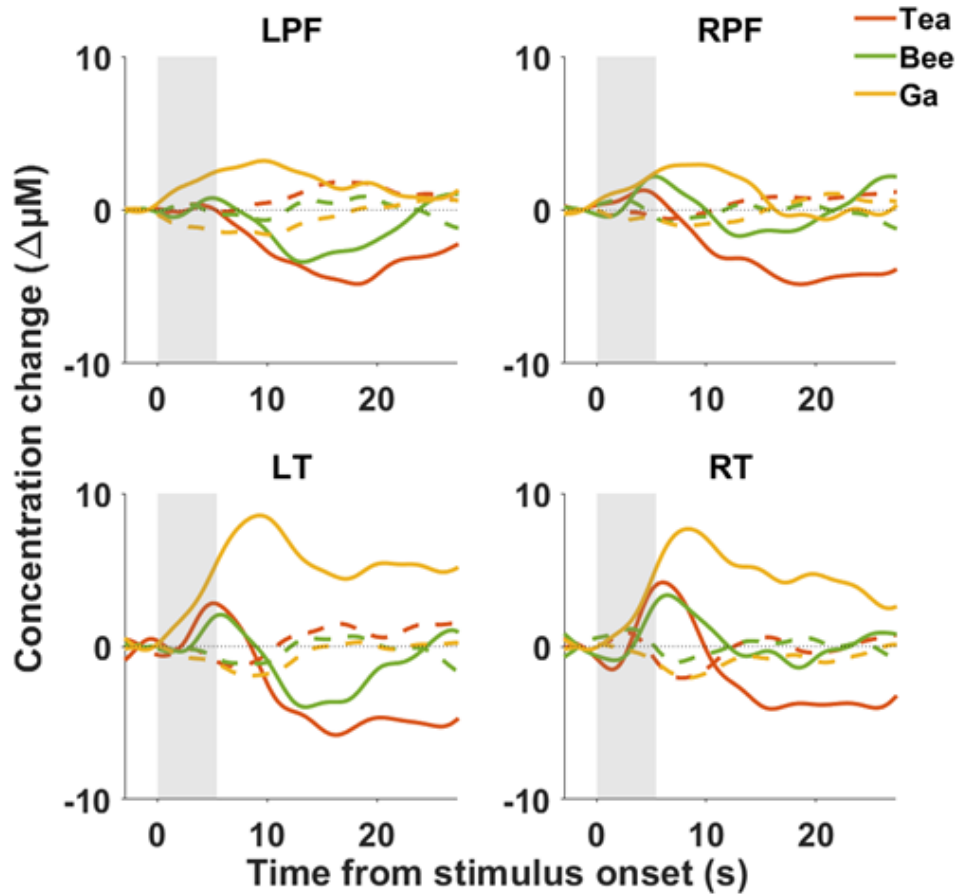
..and another physiological response to hearing sounds



- Rapidly habituates (similarly to arousal response)
- Is only reliably measurable at higher intensities in normal hearing babies (≥ 65 dB SPL)

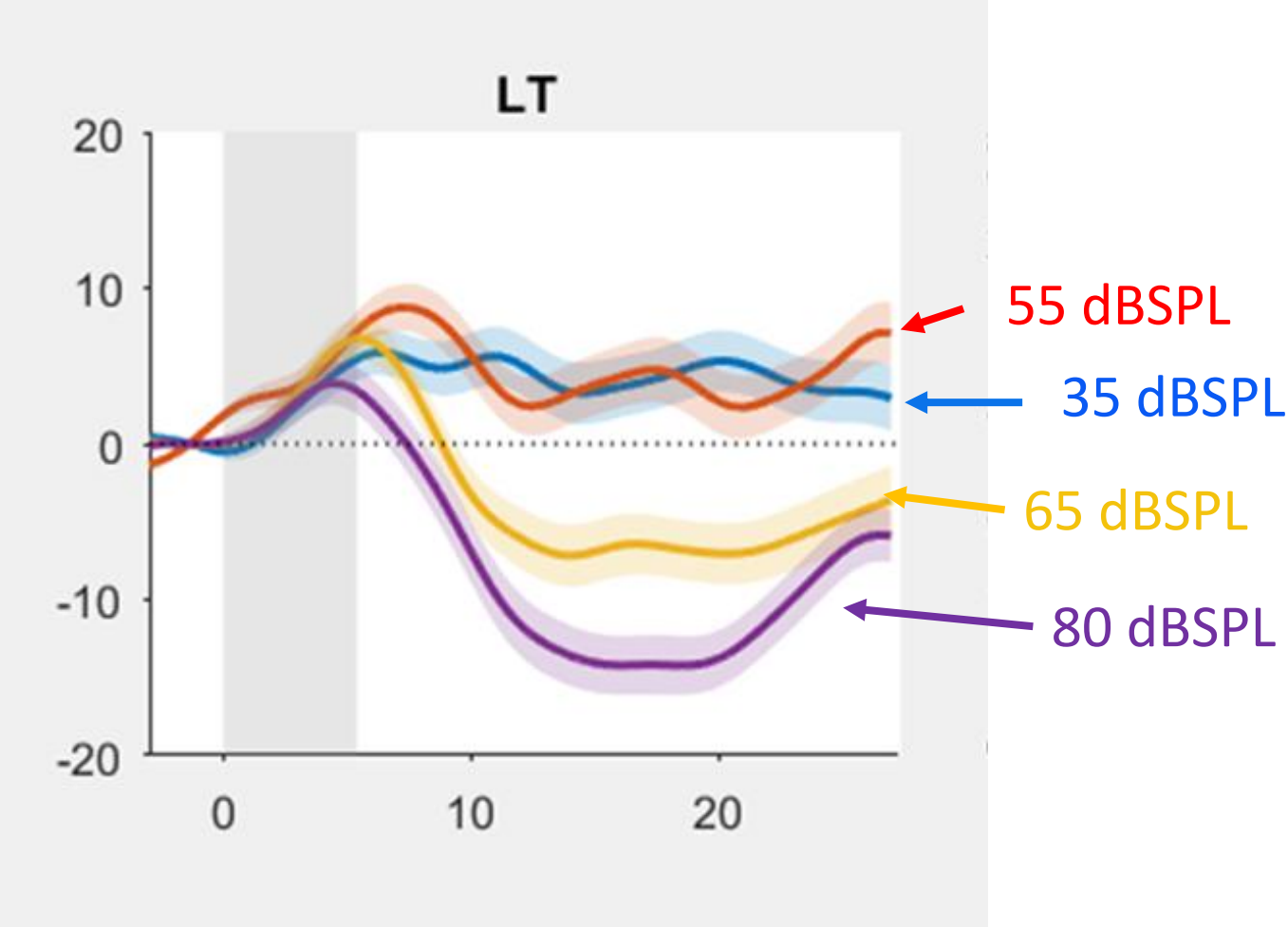
Discrimination response “Ba” versus “Tea”, “Bee”, and “Ga” at 65 dB SPL

Grand averages per ROI

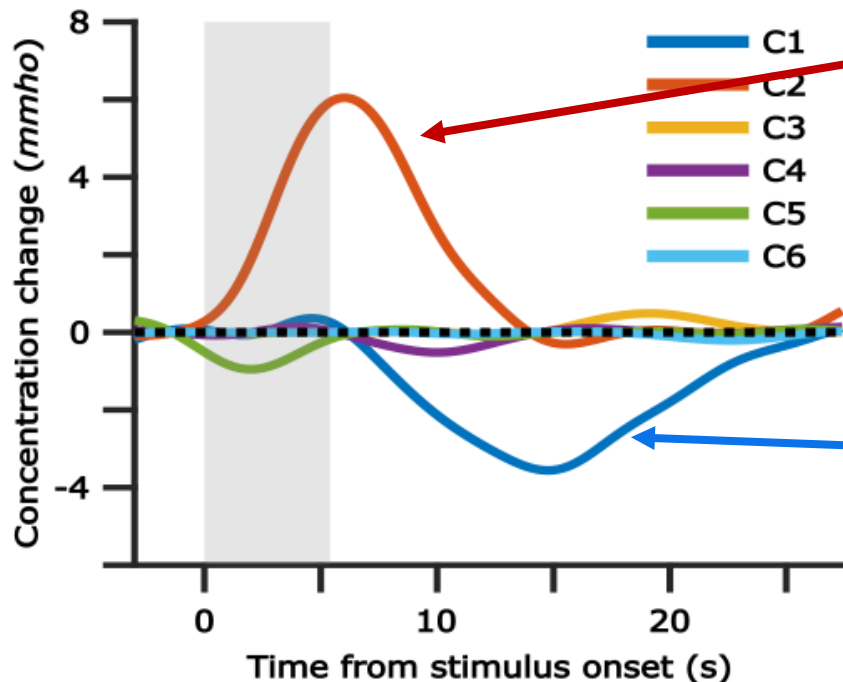


- Same negative arousal component present
- Arousal **sensitive to degree of contrast**
- Positive acoustic component is broader and has longer peak latency than the detection response

Effect of stimulus level on fNIRS response



Summary of fNIRS response characteristics in sleeping infants



Auditory: latency and duration are affected by sound intensity or discrimination contrast

Brain Arousal: rapidly habituates
Size is affected by sound intensity and discrimination contrast

An automatic analysis method that is reliable and accurate in individual infants

- Needs to be able to automatically **detect non-stationary responses with differing morphology**
- A brief overview (patent protected)
 - Standard pre-processing steps to clean the signal and convert light intensities to HbO and HbR
 - A stochastic process captures the unique statistical properties of neural responses, illustrating neighborhood covariance relative to stimulus onset across a series of post-stimulus responses spanning the expected response lengths
 - Statistical significance is established by comparing the derived neural stochastic process against arbitrary baseline signals (from silence period or non-silence baseline).
 - To assess the probability of a false positive for that infant/ROI, control baseline signals are compared to each other using the same statistical process

Accuracy analysis in individual infants (normal hearing)

Specificity 95% (calculated across multiple control versus control tests in around 40 infants i.e. false positive estimate = 5%)

Secondary false positive check: control versus “0 dB SPL stimulus” 14 trials, no false positives

Sensitivity

Detection:

65 dB SPL 97-100% (28/29 infant tests)

Other levels 35 dB SPL to 90 dB SPL 100% (38 tests in all 5-8 tests per level)

Discrimination:

Ba/Tea 65 dB SPL 15/16 infants (94%)

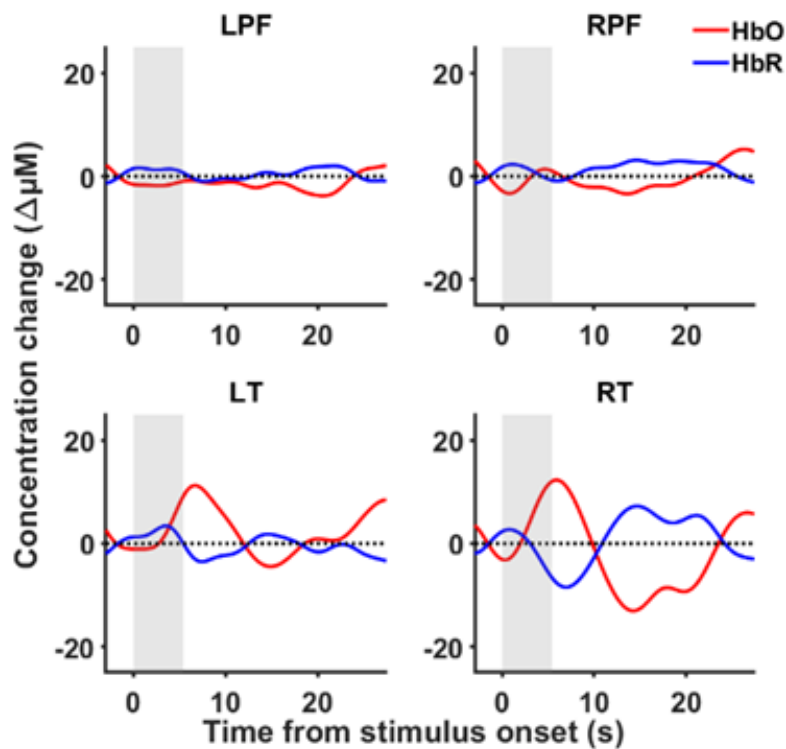
Ba/Bee 65-75 dB SPL 30/33 infants (91%)

Ba/Bee 35-50 dB SPL 20/21 infants (95%)

Ba/Ga 65 dB SPL 14/16 infants (88%)

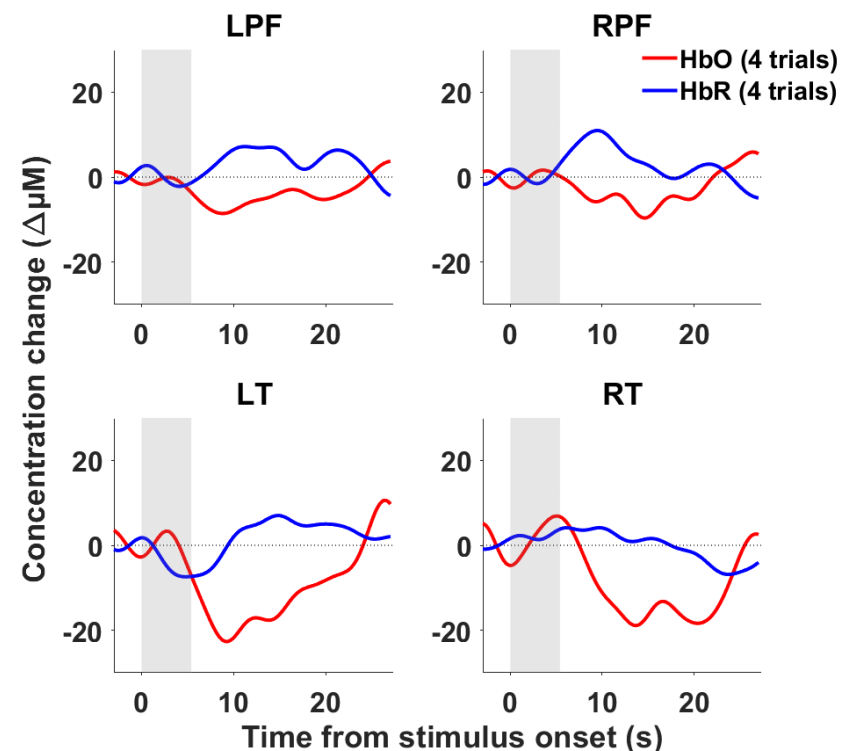
Example: infant (6m) with moderate hearing loss: Aided condition at 65 dB SPL: Detection responses

Average over 20 epochs



Algorithm says "Yes, detected"

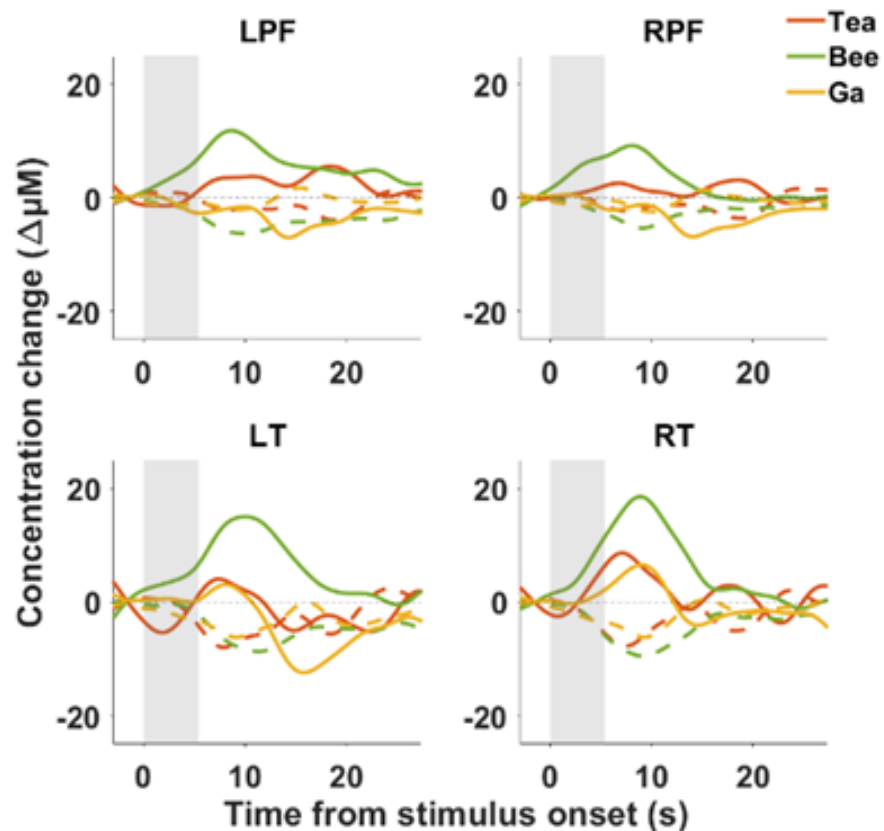
Average over first 4 epochs



Note dominance of arousal response at start

Example: infant (6m) with moderate hearing loss: Aided condition at 65 dB SPL: Discrimination responses

Average over 20 epochs

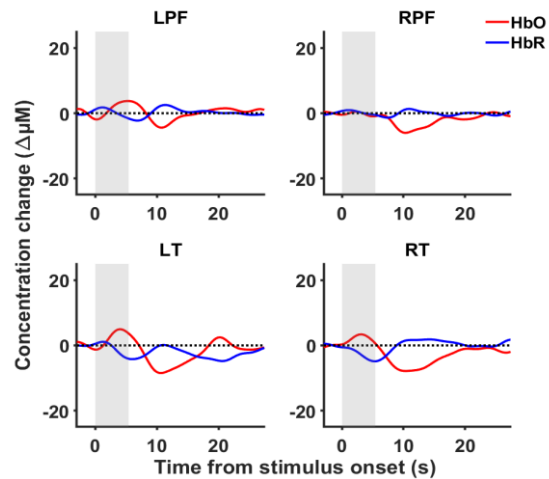


Note delayed positive peaks.

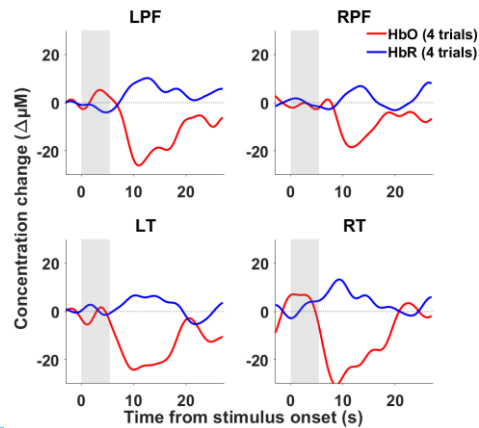
Algorithm confirms discrimination of all contrasts

Example: infant (bilateral severe/profound loss 5m and repeat 9m): aided condition at 65 dB SPL

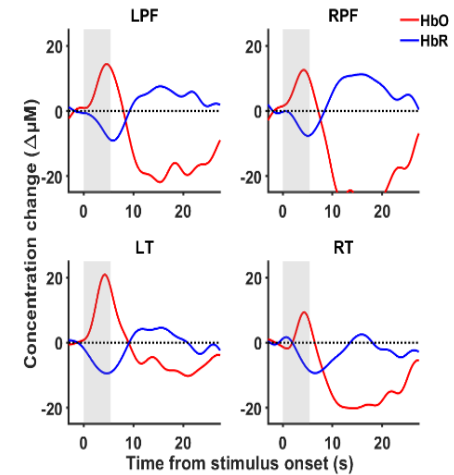
Detection responses at 5 months



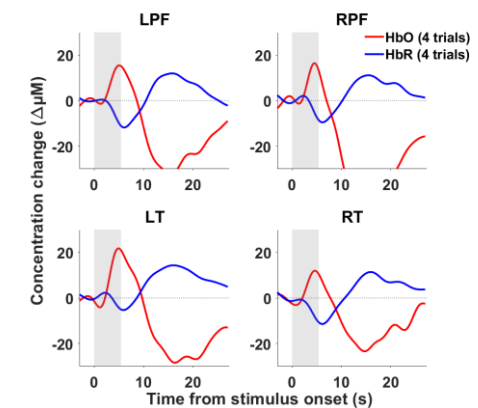
First 4 epochs



Detection responses at 9 months



First 4 epochs



Developing a clinically easy system ...ready for clinical testing September 2023



Baby-friendly head gear with fixed montage

Wireless connection



Clinician friendly user interface

Automatic analysis and report



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