

Emergent intensity invariance in a physiologically inspired model of the grasshopper auditory system

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1 The sensory world of a grasshopper

Strong dependence on acoustic signals for ranged communication

- Diverse species-specific sound repertoires and production mechanisms
- Different contexts/ranges: Stridulatory, mandibular, wings, walking sounds
- Mate attraction/evaluation, rival deterrence, loss-of-signal predator alarm
- Elaborate acoustic behaviors co-depend on reliable auditory perception

Songs = Amplitude-modulated (AM) broad-band acoustic signals

- Generated by stridulatory movement of hindlegs against forewings
- Shorter time scales: Characteristic temporal waveform pattern
- Longer time scales: High degree of periodicity (pattern repetition)
- Sound propagation: Signal intensity varies strongly with distance to sender
- Ectothermy: Temporal structure warps with temperature
- Sensory constraints imposed by properties of the acoustic signal itself

Multi-species, multi-individual communally inhabited environments

- Temporal overlap: Simultaneous singing across individuals/species common
- Frequency overlap: No/hardly any niche speciation into frequency bands
- "Biotic noise": Hetero-/conspecifics ("Another one's songs are my noise")
- "Abiotic noise": Wind, water, vegetation, anthropogenic
- Effects of habitat structure on sound propagation (landscape - soundscape)
- Sensory constraints imposed by the (acoustic) environment

Cluster of auditory challenges (interlocking constraints → tight coupling):

From continuous acoustic input, generate neuronal representations that...

- 1)...allow for the separation of relevant (song) events from ambient noise floor
- 2)...compensate for behaviorally non-informative song variability (invariances)

3)...carry sufficient information to characterize different song patterns, recognize the ones produced by conspecifics, and make appropriate behavioral decisions based on context (sender identity, song type, mate/rival quality)

How can the auditory system of grasshoppers meet these challenges?

- What are the minimum functional processing steps required?
- Which known neuronal mechanisms can implement these steps?
- Which and how many stages along the auditory pathway contribute?
- What are the limitations of the system as a whole?

How can a human observer conceive a grasshopper's auditory percepts?

- How to investigate the workings of the auditory pathway as a whole?
- How to systematically test effects and interactions of processing parameters?
- How to integrate the available knowledge on anatomy, physiology, ethology?
- Abstract, simplify, formalize → Functional model framework

2 Developing a functional model of the grasshopper auditory pathway

2.1 Population-driven signal pre-processing

"Pre-split portion" of the auditory pathway:

Tympanal membrane → Receptor neurons → Local interneurons

Similar response/filter properties within receptor/interneuron populations (Jan Clemens 2011)

→ One population-wide response trace per stage (no "single-cell resolution")

Stage-specific processing steps and functional approximations:

Initial: Continuous acoustic input signal $x(t)$

Filtering of behaviorally relevant frequencies by tympanal membrane

→ Bandpass filter 5-30 kHz

$$x_{\text{filt}}(t) = x(t) * h_{\text{BP}}(t), \quad f_{\text{cut}} = 5 \text{ kHz}, 30 \text{ kHz} \quad (1)$$

Extraction of signal envelope (AM encoding) by receptor population

→ Full-wave rectification, then lowpass filter 500 Hz

$$x_{\text{env}}(t) = |x_{\text{filt}}(t)| * h_{\text{LP}}(t), \quad f_{\text{cut}} = 500 \text{ Hz} \quad (2)$$

Logarithmically compressed intensity tuning curve of receptors

→ Decibel transformation

$$x_{\text{dB}}(t) = 10 \cdot \log_{10} \frac{x_{\text{env}}(t)}{x_{\text{ref}}}, \quad x_{\text{ref}} = \max[x_{\text{env}}(t)] \quad (3)$$

Spike-frequency adaptation in receptor and interneuron populations

→ Highpass filter 10 Hz

$$x_{\text{adapt}}(t) = x_{\text{dB}}(t) * h_{\text{HP}}(t), \quad f_{\text{cut}} = 10 \text{ Hz} \quad (4)$$

2.2 Feature extraction by individual neurons

”Post-split portion” of the auditory pathway:

Ascending neurons (AN) → Central brain neurons

Diverse response/filter properties within AN population (Jan Clemens 2011)

- Pathway splitting into several parallel branches
- Expansion into a decorrelated higher-dimensional sound representation
- Individual neuron-specific response traces from this stage onwards

Stage-specific processing steps and functional approximations:

Template matching by individual ANs

- Filter base (STA approximations): Set of Gabor kernels
- Gabor parameters: σ, ϕ, f → Determines kernel sign and lobe number

$$k(t) = e^{-\frac{t^2}{2\sigma^2}} \cdot \sin(2\pi ft + \phi) \quad (5)$$

→ Separate convolution with each member of the kernel set

$$c_i(t) = x_{\text{adapt}}(t) * k_i(t) = \int_{-\infty}^{+\infty} x_{\text{adapt}}(\tau) \cdot k_i(t - \tau) d\tau \quad (6)$$

Thresholding nonlinearity in ascending neurons (or further downstream)

- Binarization of AN response traces into ”relevant” vs. ”irrelevant”
- Shifted Heaviside step-function $H(c_i - \Theta_i)$ (or steep sigmoid threshold?)

$$b_{i,\Theta}(t) = \begin{cases} 1, & c_i(t) > \Theta_i \\ 0, & c_i(t) \leq \Theta_i \end{cases} \quad (7)$$

Temporal averaging by neurons of the central brain

- Finalized set of slowly changing kernel-specific features (one per AN)
- Different species-specific song patterns are characterized by a distinct combination of feature values → Clusters in high-dimensional feature space
- Lowpass filter 1 Hz

$$f_{i,\Theta}(t) = b_{i,\Theta}(t) * h_{\text{LP}}(t), \quad f_{\text{cut}} = 1 \text{ Hz} \quad (8)$$

3 Two mechanisms driving the emergence of intensity-invariant song representation

3.1 Logarithmic scaling & spike-frequency adaptation

Envelope $x_{\text{env}}(t) \xrightarrow{\text{dB}}$ Logarithmic $x_{\text{dB}}(t) \xrightarrow{h_{\text{HP}}(t)}$ Adapted $x_{\text{adapt}}(t)$

- Rewrite signal envelope $x_{\text{env}}(t)$ (Eq. 2) as a synthetic mixture:
 - 1) Song signal $s(t)$ ($\sigma_s^2 = 1$) with variable multiplicative scale $\alpha \geq 0$
 - 2) Fixed-scale additive noise $\eta(t)$ ($\sigma_\eta^2 = 1$)

$$x_{\text{env}}(t) = \alpha \cdot s(t) + \eta(t), \quad x_{\text{env}}(t) > 0 \quad \forall t \in \mathbb{R} \quad (9)$$

- Signal-to-noise ratio (SNR): Ratio of variances of synthetic mixture $x_{\text{env}}(t)$ with ($\alpha > 0$) and without ($\alpha = 0$) song signal $s(t)$, assuming $s(t) \perp \eta(t)$

$$\text{SNR} = \frac{\sigma_{s+\eta}^2}{\sigma_\eta^2} = \frac{\alpha^2 \cdot \sigma_s^2 + \sigma_\eta^2}{\sigma_\eta^2} = \alpha^2 + 1 \quad (10)$$

Logarithmic component:

- Simplify decibel transformation (Eq. 3) and apply to synthetic $x_{\text{env}}(t)$
- Isolate scale α and reference x_{ref} using logarithm product/quotient laws

$$\begin{aligned} x_{\text{dB}}(t) &= \log \frac{\alpha \cdot s(t) + \eta(t)}{x_{\text{ref}}} \\ &= \log \frac{\alpha}{x_{\text{ref}}} + \log \left[s(t) + \frac{\eta(t)}{\alpha} \right] \end{aligned} \quad (11)$$

- In log-space, a multiplicative scaling factor becomes additive
- Allows for the separation of song signal $s(t)$ and its scale α
- Introduces scaling of noise term $\eta(t)$ by the inverse of α
- Normalization by x_{ref} applies equally to all terms (no individual effects)

Adaptation component:

- Highpass filter over $x_{\text{dB}}(t)$ (Eq. 4) can be approximated as subtraction of the local signal offset within a suitable time interval T_{HP} ($0 \ll T_{\text{HP}} < \frac{1}{f_{\text{cut}}}$)

$$x_{\text{adapt}}(t) \approx x_{\text{dB}}(t) - \log \frac{\alpha}{x_{\text{ref}}} = \log \left[s(t) + \frac{\eta(t)}{\alpha} \right] \quad (12)$$

Implication for intensity invariance:

- Logarithmic scaling is essential for equalizing different song intensities
- Intensity information can be manipulated more easily when in form of a signal offset in log-space than a multiplicative scale in linear space

- Scale α can only be redistributed, not entirely eliminated from $x_{\text{adapt}}(t)$
 \rightarrow Turn initial scaling of song $s(t)$ by α into scaling of noise $\eta(t)$ by $\frac{1}{\alpha}$
- Capability to compensate for intensity variations, i.e. selective amplification of output $x_{\text{adapt}}(t)$ relative to input $x_{\text{env}}(t)$, is limited by input SNR (Eq. 10):
 $\alpha \gg 1$: Attenuation of $\eta(t)$ term $\rightarrow s(t)$ dominates $x_{\text{adapt}}(t)$
 $\alpha \approx 1$ Negligible effect on $\eta(t)$ term $\rightarrow x_{\text{adapt}}(t) = \log[s(t) + \eta(t)]$
 $\alpha \ll 1$: Amplification of $\eta(t)$ term $\rightarrow \eta(t)$ dominates $x_{\text{adapt}}(t)$
 \rightarrow Ability to equalize between different sufficiently large scales of $s(t)$
 \rightarrow Inability to recover $s(t)$ when initially masked by noise floor $\eta(t)$
- Logarithmic scaling emphasizes small amplitudes (song onsets, noise floor)
 \rightarrow Recurring trade-off: Equalizing signal intensity vs preserving initial SNR

3.2 Threshold nonlinearity & temporal averaging

Convolved $c_i(t) \xrightarrow{H(c_i - \Theta_i)}$ Binary $b_{i,\Theta}(t) \xrightarrow{h_{\text{LP}}(t)}$ Feature $f_{i,\Theta}(t)$

Thresholding component:

- Within an observed time interval T , $c_i(t)$ follows probability density $p(c_i, T)$
- Within T , $c_i(t)$ exceeds threshold value Θ_i for time T_1 ($T_1 + T_0 = T$)
- Threshold $H(c_i - \Theta_i)$ splits $p(c_i, T)$ around Θ_i in two complementary parts

$$\int_{\Theta_i}^{+\infty} p(c_i, T) dc_i = 1 - \int_{-\infty}^{\Theta_i} p(c_i, T) dc_i = \frac{T_1}{T} \quad (13)$$

\rightarrow Semi-definite integral over right-sided portion of split $p(c_i, T)$ gives ratio of time T_1 where $c_i(t) > \Theta_i$ to total time T due to normalization of $p(c_i, T)$

$$\int_{-\infty}^{+\infty} p(c_i, T) dc_i = 1 \quad (14)$$

Averaging component:

- Lowpass filter over binary response $b_{i,\Theta}(t)$ (Eq. 8) can be approximated as temporal averaging over a suitable time interval T_{LP} ($T_{\text{LP}} > \frac{1}{f_{\text{cut}}}$)
- Within T_{LP} , $b_{i,\Theta}(t)$ takes a value of 1 ($c_i(t) > \Theta_i$) for time T_1 ($T_1 + T_0 = T_{\text{LP}}$)

$$f_{i,\Theta}(t) \approx \frac{1}{T_{\text{LP}}} \int_t^{t+T_{\text{LP}}} b_{i,\Theta}(\tau) d\tau = \frac{T_1}{T_{\text{LP}}} \quad (15)$$

\rightarrow Temporal averaging over $b_{i,\Theta}(t) \in [0, 1]$ (Eq. 7) gives ratio of time T_1 where $c_i(t) > \Theta_i$ to total averaging interval T_{LP}

\rightarrow Feature $f_{i,\Theta}(t)$ approximately represents supra-threshold fraction of T_{LP}

Combined result:

- Feature $f_{i,\Theta}(t)$ can be linked to the distribution of $c_i(t)$ using Eqs. 13 & 15

$$f_{i,\Theta}(t) \approx \int_{\Theta_i}^{+\infty} p(c_i, T_{LP}) dc_i = P(c_i > \Theta_i, T_{LP}) \quad (16)$$

→ Because the integral over a probability density is a cumulative probability, the value of feature $f_{i,\Theta}(t)$ (temporal compression of $b_{i,\Theta}(t)$) at every time point t signifies the probability that convolution output $c_i(t)$ exceeds the threshold value Θ_i during the corresponding averaging interval T_{LP}

Implication for intensity invariance:

- Convolution output $c_i(t)$ quantifies temporal similarity between amplitudes of template waveform $k_i(t)$ and signal $x_{\text{adapt}}(t)$ centered at time point t
 - Based on amplitudes on a graded scale
- Feature $f_{i,\Theta}(t)$ quantifies the probability that amplitudes of $c_i(t)$ exceed threshold value Θ_i within interval T_{LP} around time point t
 - Based on binned amplitudes corresponding to one of two categorical states
 - Deliberate loss of precise amplitude information
 - Emphasis on temporal structure (ratio of T_1 over T_{LP})
- Thresholding of $c_i(t)$ and subsequent temporal averaging of $b_{i,\Theta}(t)$ to obtain $f_{i,\Theta}(t)$ constitutes a remapping of an amplitude-encoding quantity into a duty cycle-encoding quantity, mediated by threshold function $H(c_i - \Theta_i)$
- Different scales of $c_i(t)$ can result in similar T_1 segments depending on the magnitude of the derivative of $c_i(t)$ in temporal proximity to time points at which $c_i(t)$ crosses threshold value Θ_i
 - The steeper the slope of $c_i(t)$, the less T_1 changes with scale variations
 - Extreme amplitudes of $c_i(t)$ (peaks/troughs)
 - Only amplitudes of
 - Absolute amplitudes of peaks/troughs of $c_i(t)$
 - Acuity of peaks/troughs in $c_i(t)$ matters, not their absolute amplitude
- From graded stimulus to categorical behavioral decision:

4 Discriminating species-specific song patterns in feature space

5 Conclusions & outlook