Coding and computation by neural ensembles in the primate retina

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— with J. Pillow (UT Austin), G. Field, J. Gauthier, J. Shlens (Salk), A. Litke (UCSC),
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Machado, D. Pfau, X. Pitkow, M. Vidne (Columbia).

Retinal ganglion neuronal data

Preparation: dissociated macaque retina

— extracellularly-recorded responses of populations of RGCs



Stimulus: random spatiotemporal visual stimuli (Pillow et al., 2008)

Receptive fields tile visual space



Multineuronal point-process model



$$\lambda_i(t) = f\left(b_i + \vec{k}_i \cdot \vec{x}(t) + \sum_{i',j} h_{i',j} n_{i'}(t-j)\right),$$

— likelihood is easy to compute and to maximize (concave optimization)
(Paninski, 2004; Paninski et al., 2007; Pillow et al., 2008)

— close connections to noisy integrate-and-fire model

Optimal Bayesian decoding

 $E(\vec{x}|spikes) \approx \arg\max_{\vec{x}} \log P(\vec{x}|spikes) = \arg\max_{\vec{x}} \left[\log P(spikes|\vec{x}) + \log P(\vec{x})\right]$

(Loading yashar-decode.mp4)

- Computational points:
 - $\log P(spikes|\vec{x})$ is concave in \vec{x} : concave optimization again.
 - Decoding can be done in linear time via standard Newton-Raphson methods, since Hessian of log P(x|spikes) w.r.t. x is banded (Pillow et al., 2010; Ahmadian et al., 2010).

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- Biological point: paying attention to correlations improves decoding accuracy.

Application: how important is timing?



— further applications: decoding velocity signals (Lalor et al., 2009), tracking images perturbed by eye jitter (Pfau et al., 2009)

Next steps: reconsidering the model

$$\lambda_i(t) = \exp\left(k_i \cdot x(t) + h_i \cdot y_i(t) + \sum_{i \neq j} l_{i,j} \cdot y_j(t)\right)$$



Pros:

- Tractable model-fitting and optimal decoding
- Captures response statistics

Cons:

- Instantaneous coupling filters
- No explicit Common Input

Considering common input effects



— universal problem in network analysis: can't observe all neurons!

Intracellular findings:

 RGCs receive strongly correlated synaptic input in the absence of modulated light stimuli



Khuc Trong & Rieke Nature Neuro 2008

F RGCs



• ON RGCs are weakly electrically coupled

• No electrical coupling seen between OFF RGCs





Direct state-space optimization methods

To fit parameters, optimize approximate marginal likelihood:

$$\log p(spikes|\theta) = \log \int p(Q|\theta)p(spikes|\theta, Q)dQ$$

$$\approx \log p(\hat{Q}_{\theta}|\theta) + \log p(spikes|\hat{Q}_{\theta}) - \frac{1}{2}\log|J_{\hat{Q}_{\theta}}|$$

$$\hat{Q}_{\theta} = \arg \max_{Q} \{\log p(Q|\theta) + \log p(spikes|Q)\}$$

-Q is a very high-dimensional latent (unobserved) "common input" term. Taken to be a Gaussian process here with autocorrelation time ≈ 5 ms (Khuc-Trong and Rieke, 2008).

— correlation strength specified by one parameter per cell pair. — all terms can be computed in O(T) via banded matrix methods (Paninski et al., 2010).

Inferred common input effects are strong



— note that inferred direct coupling effects are now relatively small.

Common-input-only model captures x-corrs



— single and triple-cell activities captured well, too (Vidne et al., 2009)

Decoding the stimulus and hidden input



 $\arg\max_{\vec{x}} p(\vec{x}|y,\theta) = \arg\max_{\vec{x}} \int p(\vec{x},Q|y,\theta) dQ \approx \arg\max_{\vec{x},Q} p(\vec{x},Q|y,\theta)$

Models lead to similar decoding performance



...but CI model is more robust to spike jitter and deletions (Vidne et al., 2009).

Next steps: inferring cones



— cone locations and color identity can be inferred accurately with high spatial-resolution stimuli via maximum a posteriori estimates (Field et al., 2010).

Next steps: inferring circuitry?



OFF parasol



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