

'Silent' olfactory bulb mitral cells emerge from common feature subtraction.

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Background

1. Recently [1] performed blind whole-cell patch clamp of MCs.
2. Spontaneous and odour-evoked activity were **inversely related**.
3. Cells with high baseline activity were often inhibited by odours.
4. 'Silent cells' had low baselines while highly excitable by odours.

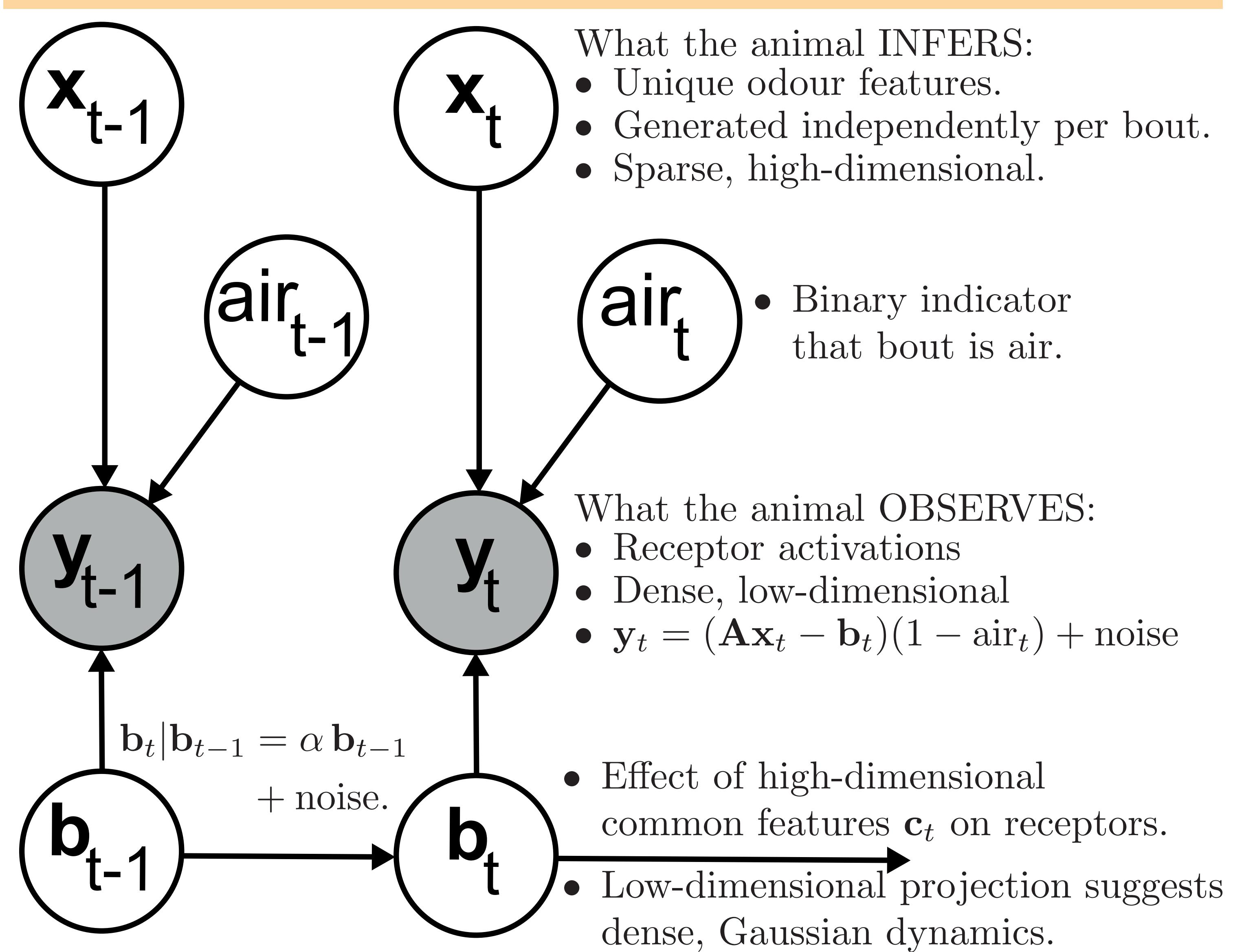
Model

1. Time discretized into $O(1 \text{ sec.})$ bouts $t \in 1 \dots N$.
2. Each bout contains either odour or air.
3. Odours are sparse, high-dimensional.
4. Odour_t = unique features \mathbf{x}_t + common features \mathbf{c}_t .
5. Unique features are generated independently per bout.
6. Common features change slowly over bouts.
7. Animal observes dense, low-D receptor activations \mathbf{y}_t .
8. Animal MAP infers unique features from receptor history:
 $\underset{\mathbf{x}_t}{\operatorname{argmax}} p(\mathbf{x}_t | \mathbf{y}_1, \dots, \mathbf{y}_t)$.

Key Ideas

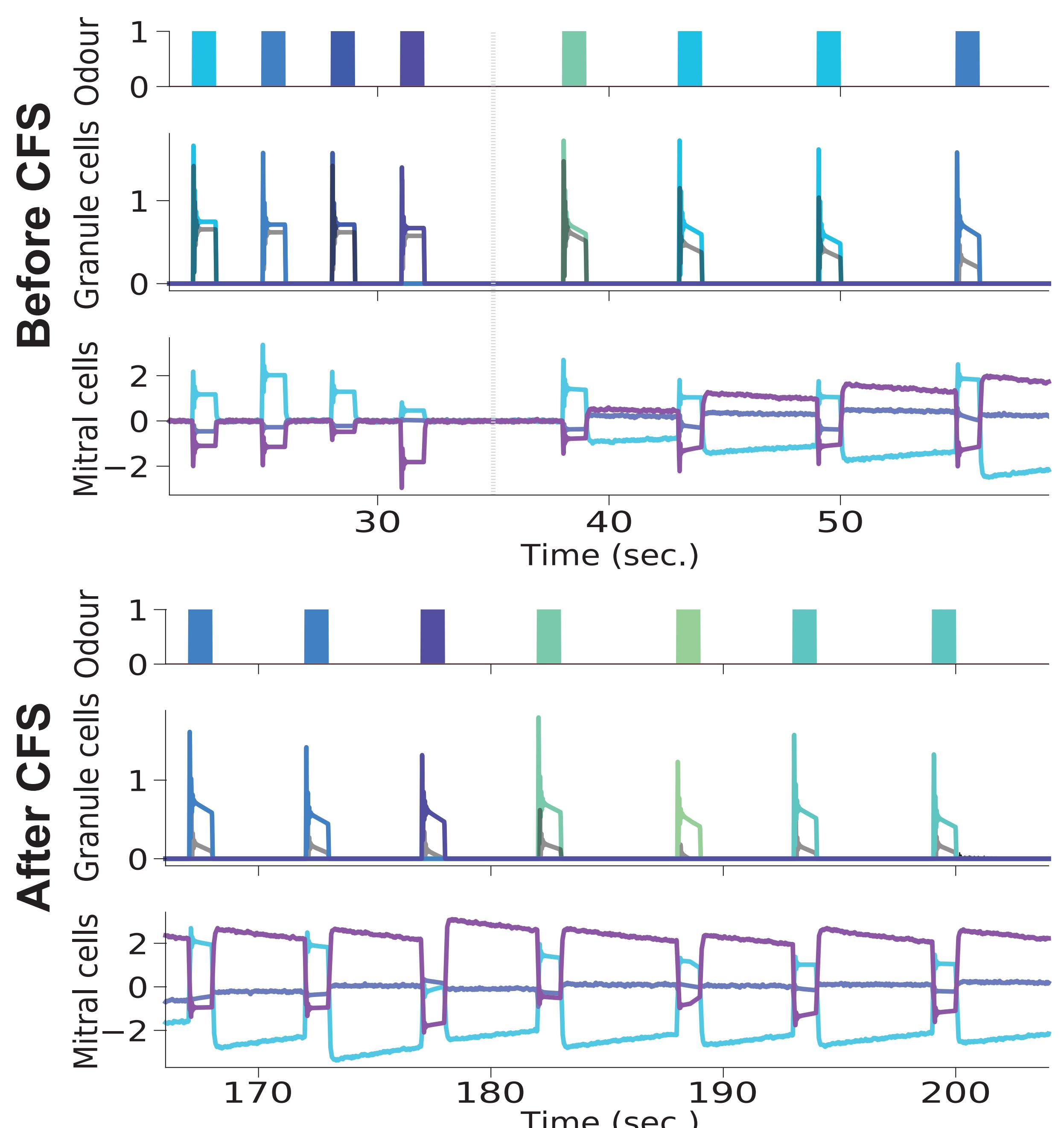
1. Common features \mathbf{c}_t are high-D but are *not inferred*.
2. They are accounted for through their effect \mathbf{b}_t on receptors.
3. Projection to low-D receptor space suggests Gaussian dynamics.
4. MC baselines reflect **negative** of expected receptor activations.
5. In odour, baseline removes component due to common features.
6. Unique features remain, inferred by fast MC/GC dynamics à la [2].

Probabilistic Model



Neural Dynamics

$$\begin{aligned} \tau_\lambda \dot{\lambda} &= -\lambda + (\mathbf{y} - \mathbf{Ax} + \alpha \mathbf{b}) / \sigma_{\text{eff}}^2 & (\text{MC voltage}) \\ \tau_b \dot{\mathbf{b}} &= -\mathbf{b} - (1 - \text{air}(t)) \kappa \lambda & (\text{MC baseline}) \\ \tau_v \dot{\mathbf{v}} &= -\mathbf{v} + \mathbf{A}^T \lambda & (\text{GC voltage}) \\ \mathbf{x} &= [\mathbf{v} - \beta]_+ / \gamma & (\text{GC firing rate}) \end{aligned}$$



Data vs. Model

