



CSE/NEUBEH 528

Lecture 9: Computation by Networks
(Chapter 7)

Image from <http://clasdean.la.asu.edu/news/images/ubep2001/neuron3.jpg>

Lecture figures are from Dayan & Abbott's book
<http://people.brandeis.edu/~abbott/book/index.html>

Course Summary (thus far)

- ◆ Neural Encoding
 - ⇒ What makes a neuron fire? (STA, covariance analysis)
 - ⇒ Poisson model
- ◆ Neural Decoding
 - ⇒ Stimulus Discrimination based on firing rate
 - ⇒ Spike-train based decoding of stimulus
 - ⇒ Population decoding (Bayesian estimation)
- ◆ Single Neuron Models
 - ⇒ RC circuit model of membrane
 - ⇒ Integrate-and-fire model
 - ⇒ Conductance-based and Compartmental Models

Today's Agenda

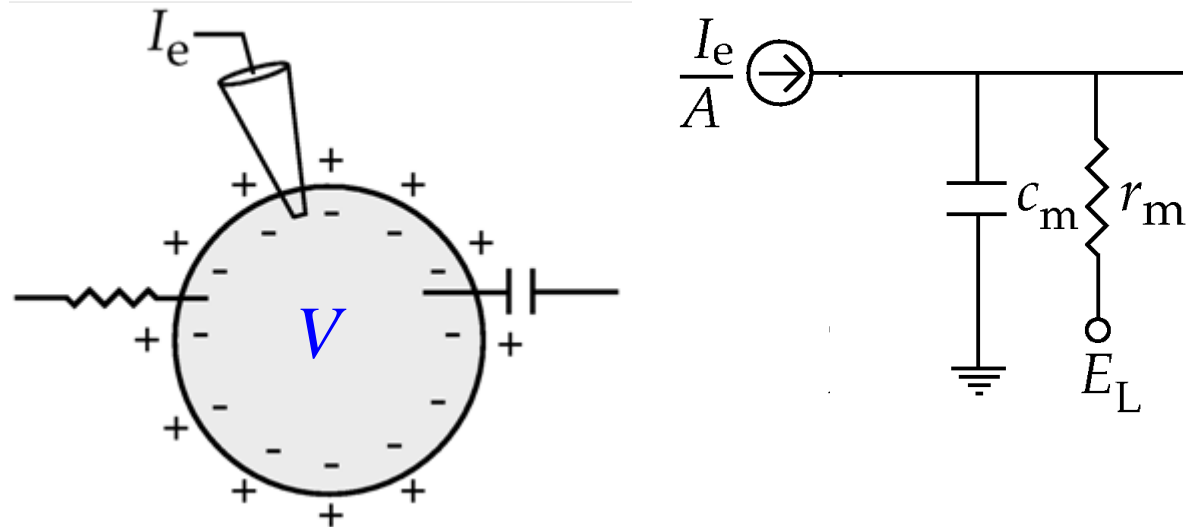
- ◆ Computation in Networks of Neurons
 - ⇒ From spiking to firing-rate based networks
 - ⇒ Feedforward Networks
 - ◆ E.g. Coordinate transformations in the brain
 - ⇒ Linear Recurrent Networks
 - ◆ Can amplify inputs
 - ◆ Can integrate inputs
 - ◆ Can function as short-term memory

Modeling Networks of Neurons

- ◆ Option 1: Use *spiking* neurons (e.g. I & F neurons)
 - ⇒ *Advantages*: Allows computation and learning based on:
 - ◆ Spike Timing
 - ◆ Spike Correlations/Synchrony between neurons
 - ⇒ *Disadvantages*: Computationally expensive
- ◆ Option 2: Use neurons with *firing-rate outputs*
 - ⇒ *Advantages*: Greater efficiency, scales well to large networks
 - ⇒ *Disadvantages*: Ignores spike timing issues
- ◆ Question: How are these two approaches related?

Flashback

1-Compartment Membrane Model



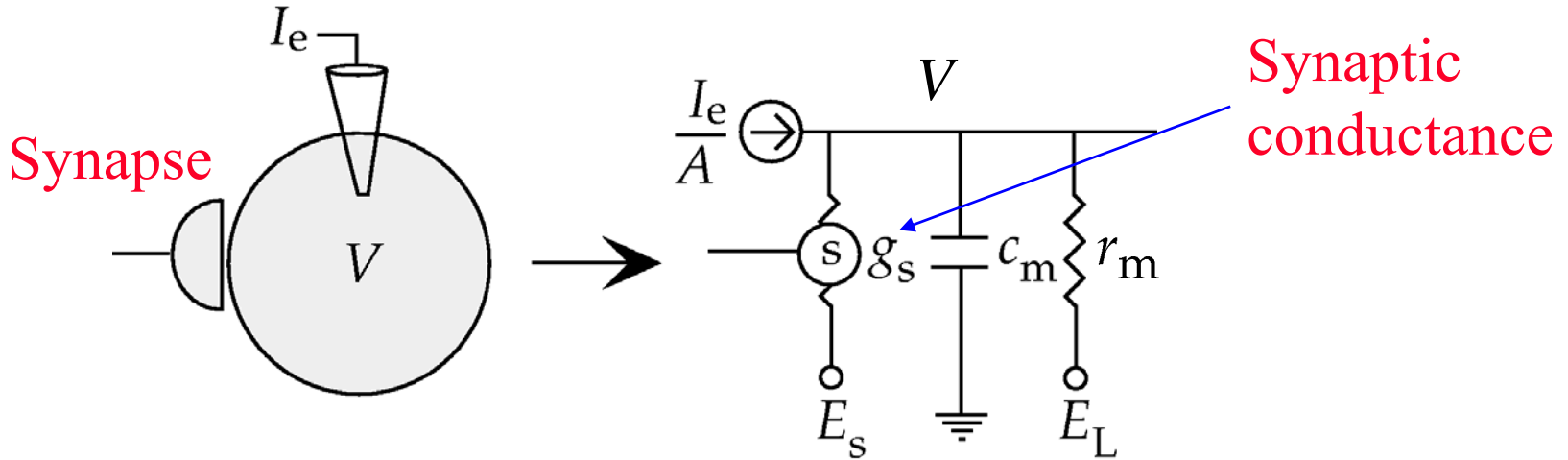
$$C_m \frac{dV}{dt} = -\frac{(V - E_L)}{r_m} + \frac{I_e}{A}, \text{ or equivalently}$$

$\tau_m = r_m C_m =$
membrane time
constant

$$\tau_m \frac{dV}{dt} = -(V - E_L) + I_e R_m$$

Flashback

Modeling Synaptic Inputs from other Neurons



$$\tau_m \frac{dV}{dt} = -(V - E_L) - r_m g_s (V - E_s) + I_e R_m$$

$$g_s = g_{s,\max} P_{rel} P_s$$

← Probability of postsynaptic channel opening
(= fraction of channels opened)

← Probability of transmitter release given an input spike

Basic Synapse Model

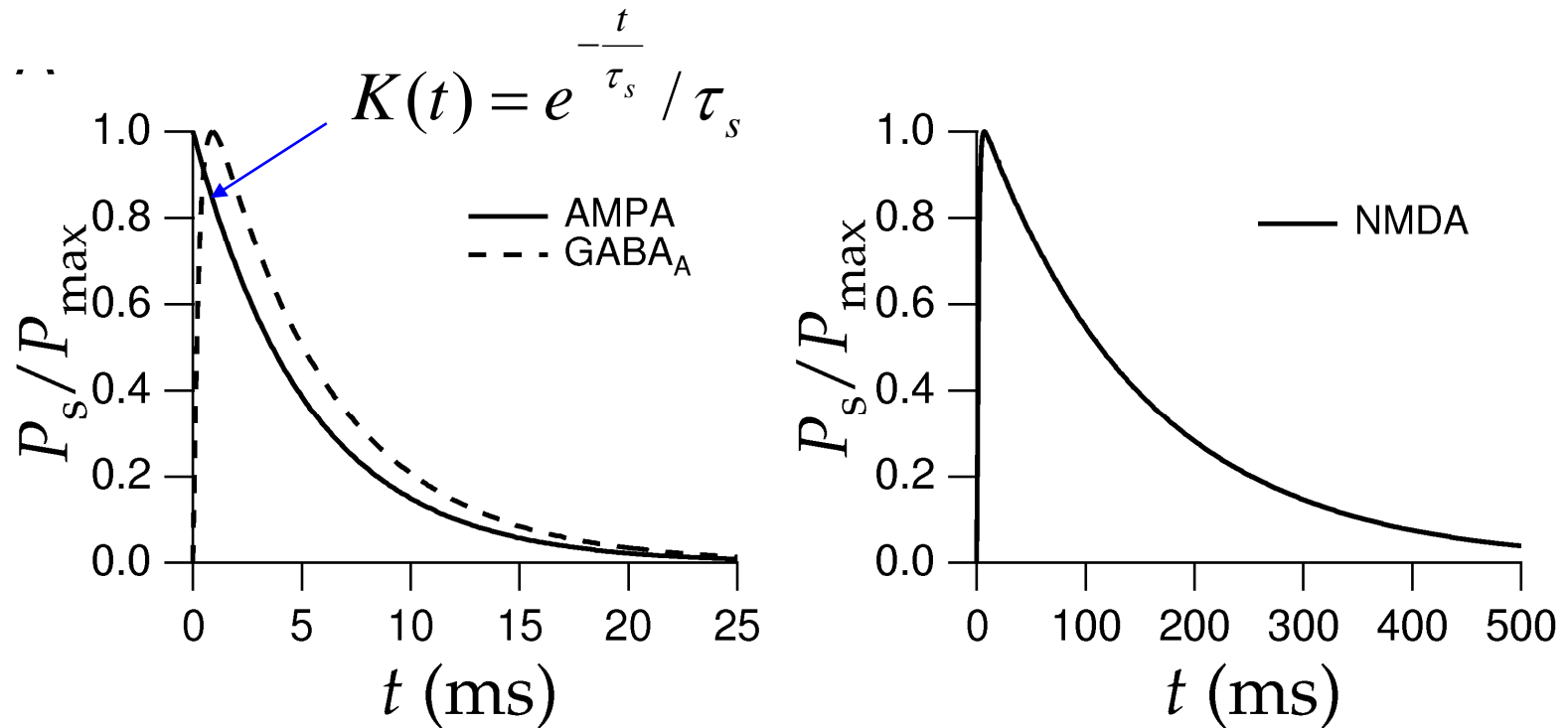
- ◆ Assume $P_{\text{rel}} = 1$
- ◆ Model the effect of a single spike input on P_s
- ◆ Kinetic Model: $\text{closed} \xrightarrow{\alpha_s} \text{open}$
 $\text{open} \xrightarrow{\beta_s} \text{closed}$

$$\frac{dP_s}{dt} = \alpha_s (1 - P_s) - \beta_s P_s$$

Opening rate Closing rate Fraction of channels open

Fraction of channels closed

Postsynaptic Data



Exponential function $K(t)$ gives reasonable fit to biological data (other options: difference of exponentials, “alpha” function)

Modeling Synaptic Input Current

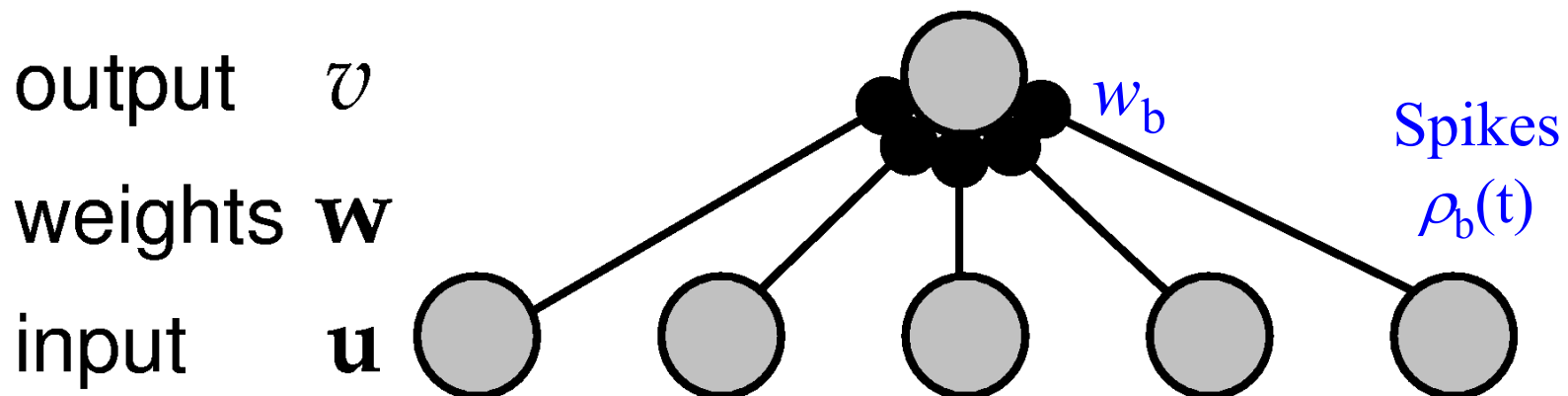
Synaptic kernel: $K(t) = e^{-\frac{t}{\tau_s}} / \tau_s$

Synaptic current: $I_s(t) = w_s \int_{-\infty}^t K(t - \tau) \rho_s(\tau) d\tau$

where $\rho_s(t)$ is the input spike train:

$$\rho_s(\tau) = \sum_i \delta(\tau - t_i) \quad (t_i \text{ are the spike times})$$

From Spiking to Firing Rate Models



Current at synapse b

$$I_b(t) = w_b \int_{-\infty}^t K(t - \tau) \rho_b(\tau) d\tau$$

Spike train $\rho_b(t)$

$$\approx w_b \int_{-\infty}^t K(t - \tau) u_b(\tau) d\tau$$

Firing rate $u_b(t)$

Total synaptic current

$$I_s(t) = \sum_b I_b(t)$$

Synaptic Current Dynamics

- ◆ If synaptic kernel K is an exponential function: $K(t) = e^{-t/\tau_s} / \tau_s$

Differentiating
$$I_s(t) = \sum_b w_b \int_{-\infty}^t K(t - \tau) u_b(\tau) d\tau$$

We get
$$\begin{aligned} \tau_s \frac{dI_s}{dt} &= -I_s + \sum_b w_b u_b \\ &= -I_s + \mathbf{w} \cdot \mathbf{u} \end{aligned}$$

Output Firing-Rate Dynamics

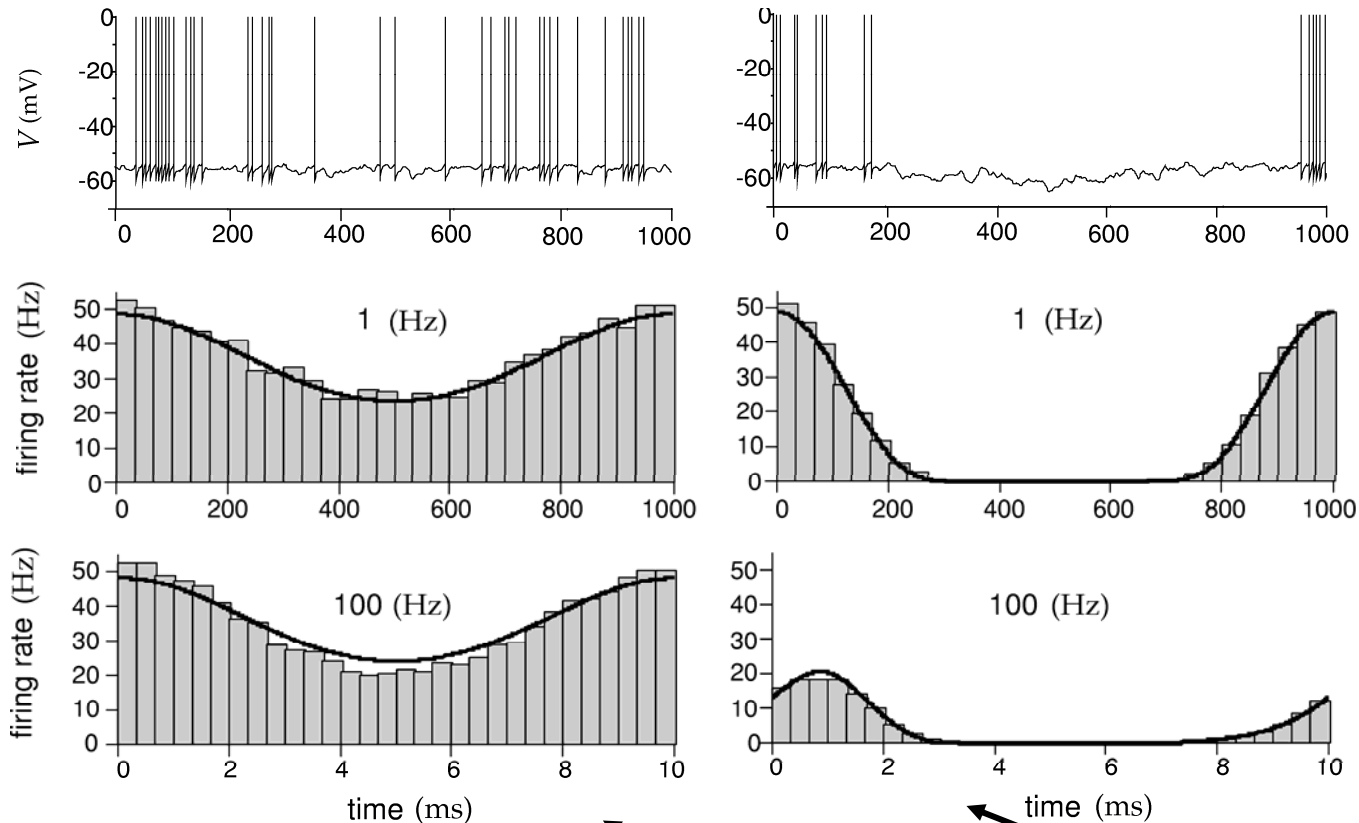
- ◆ How is the output firing rate ν related to synaptic inputs?

$$\tau_r \frac{d\nu}{dt} = -\nu + F(I_s(t))$$

- ◆ On-board derivations of special cases obtained from comparing τ_r and $\tau_s \dots$

(see also pages 234-236 in the text)

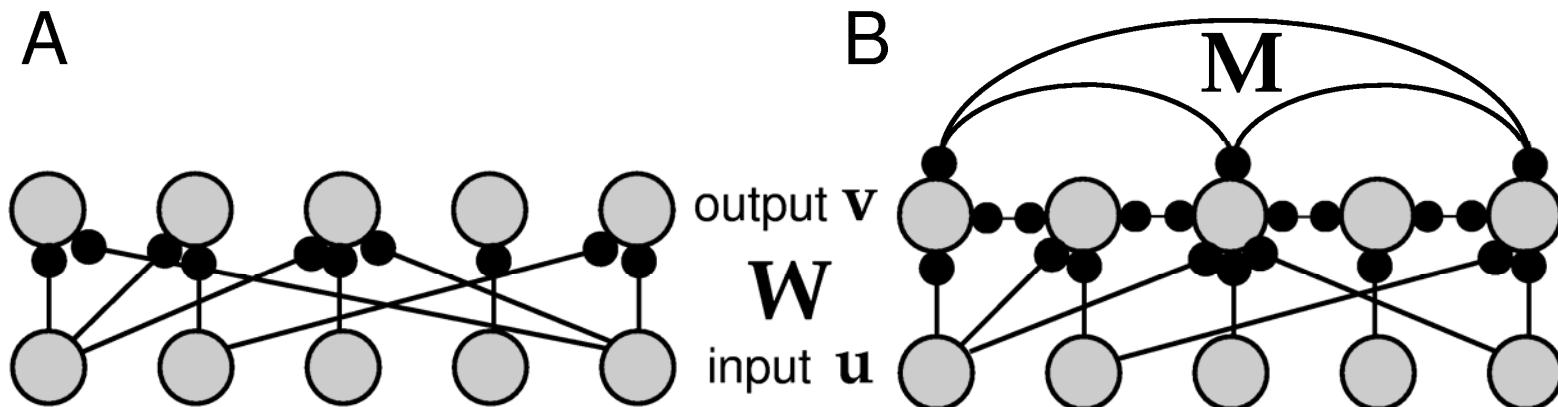
How good are the Firing Rate Models?



Firing rate $v(t) = F(I(t))$ describes this well but **not this case**

R. Rao, CSE528: Lecture 9 Input $I(t) = I_0 + I_1 \cos(\omega t)$

Feedforward versus Recurrent Networks

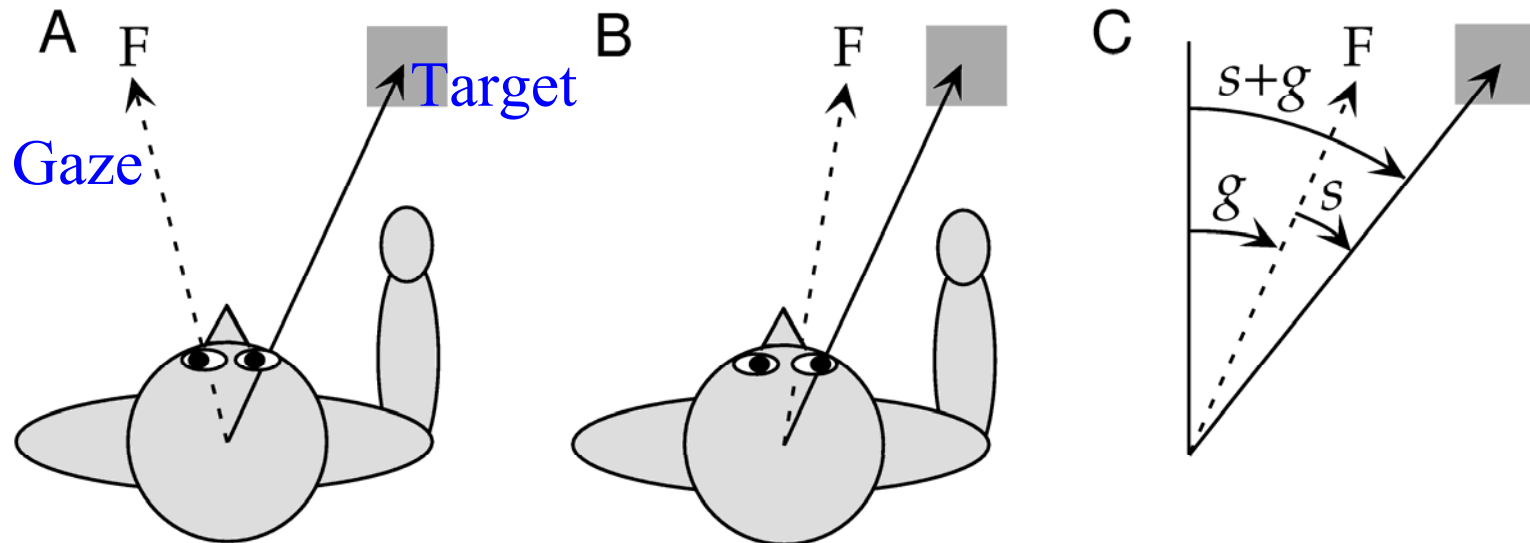


$$\tau \frac{d\mathbf{v}}{dt} = -\mathbf{v} + F(\mathbf{W}\mathbf{u} + \mathbf{M}\mathbf{v})$$

Output Decay Input Feedback

(For feedforward networks, matrix $\mathbf{M} = 0$)

The Problem of Coordinate Transformations



g = gaze angle *relative to body*

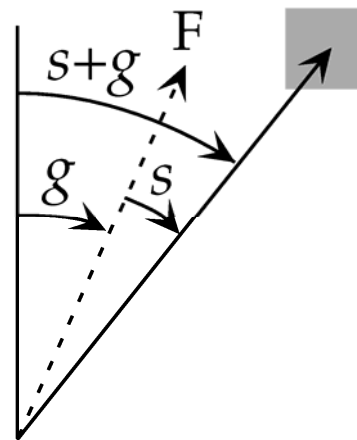
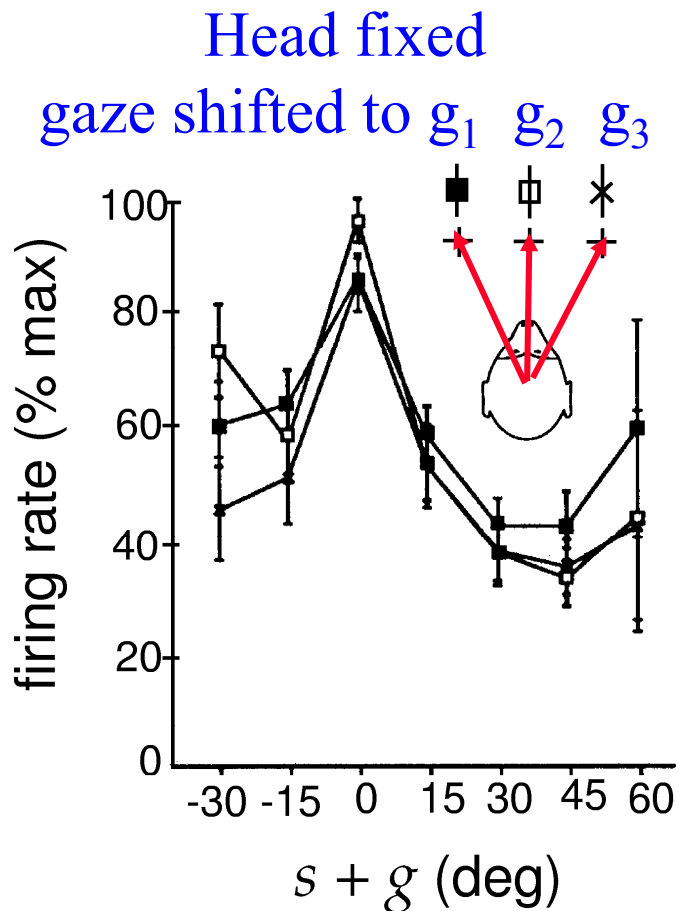
s = stimulus or target angle *relative to gaze (retinal coordinates)*

$s+g$ = stimulus relative to body

Same arm movement required in A and B but s and g are different

How does the brain solve this problem?

Body-Based Representation in the Monkey



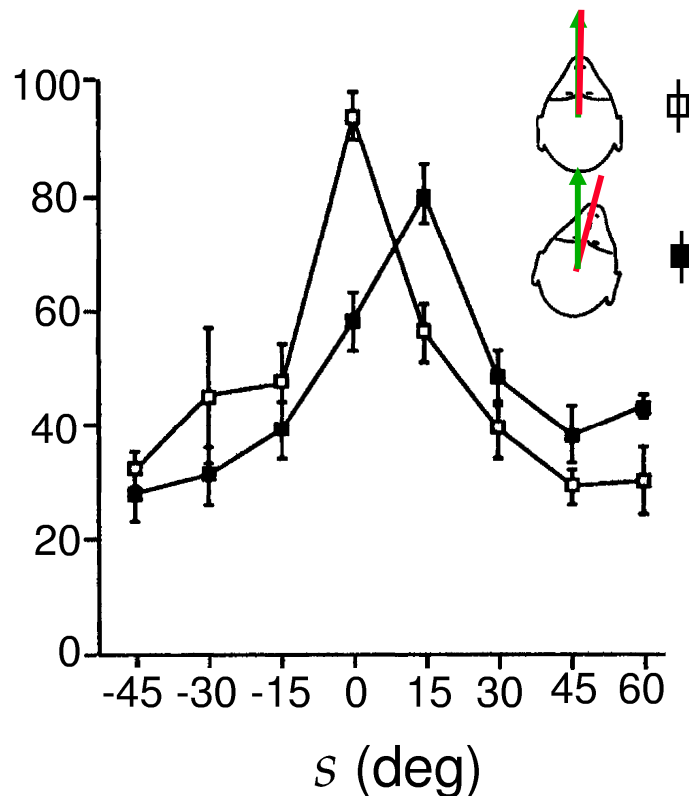
Objects approaching
at different angles

← Same tuning curve
regardless of gaze angle

Premotor cortex neuron responds
to stimulus location *relative to*
body, not retinal image location

Body-Based Representation in the Monkey

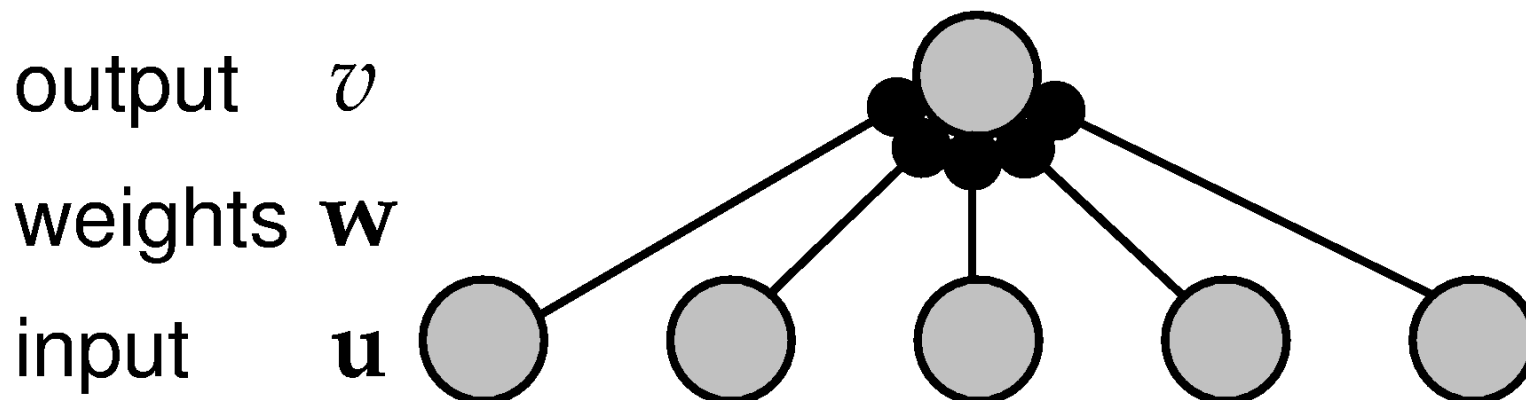
When head is moved but
gaze remains unchanged:



After head is moved 15° ,
objects approaching at 15°
in retinal image now elicit
the highest response \rightarrow
Tuning curve in retinal
coordinates has shifted

Suggested Feedforward Network

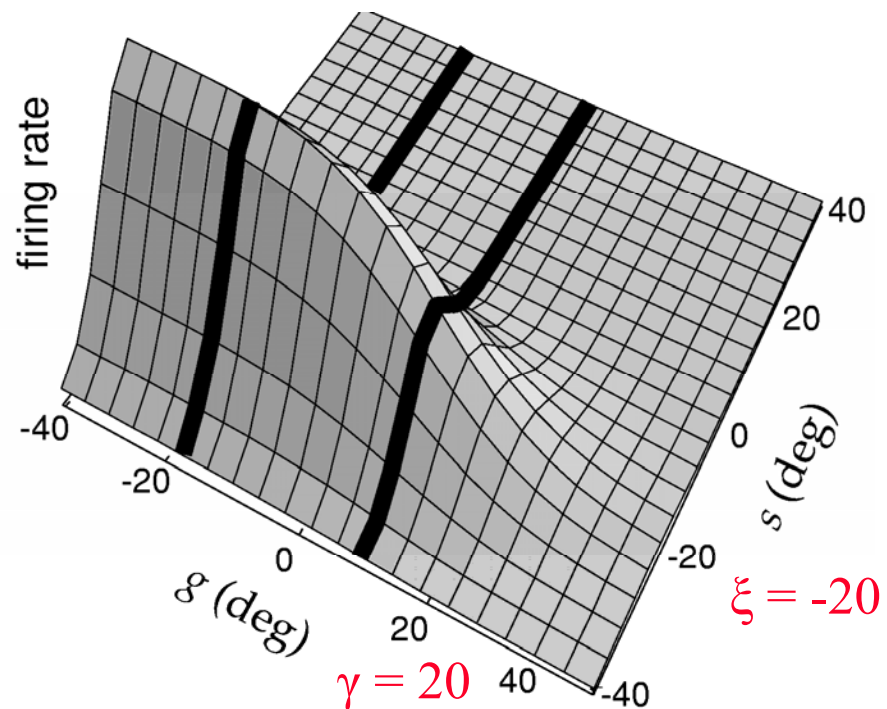
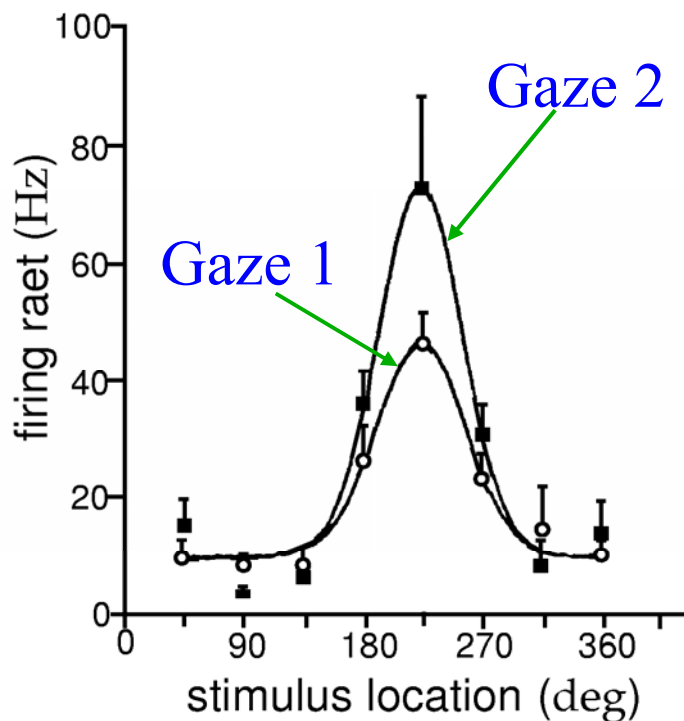
Output: Premotor Cortex Neuron with Body-Based Tuning Curves



Input: Area 7a Neurons with Gaze-Dependent Tuning Curves

Input neurons exhibit *gaze-dependent gain modulation*

Gaze-Dependent Gain Modulation

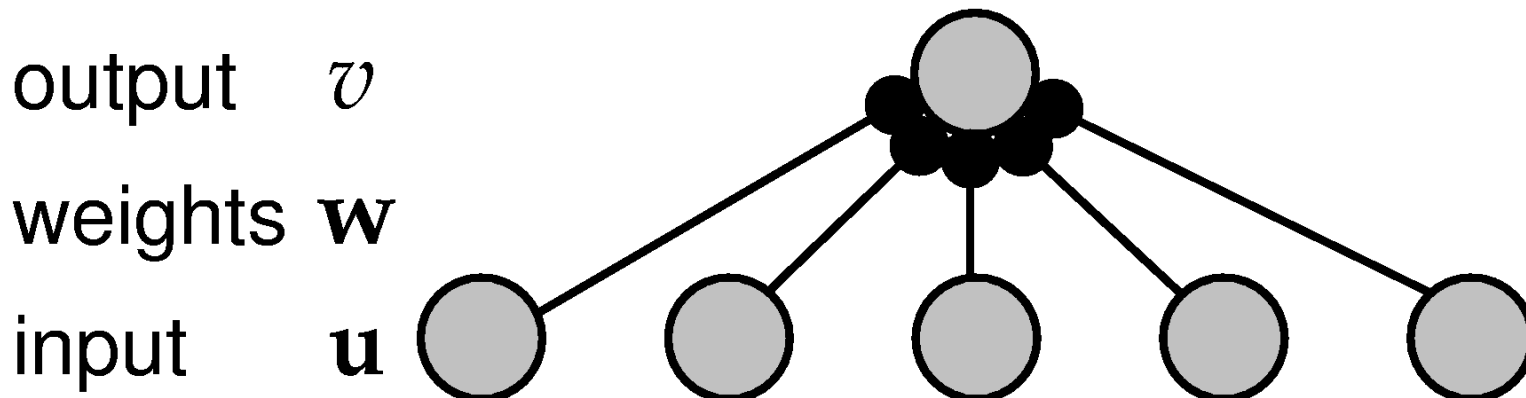


Responses of Area 7a neuron

Example of a gain-modulated tuning curve

What should the weights be?

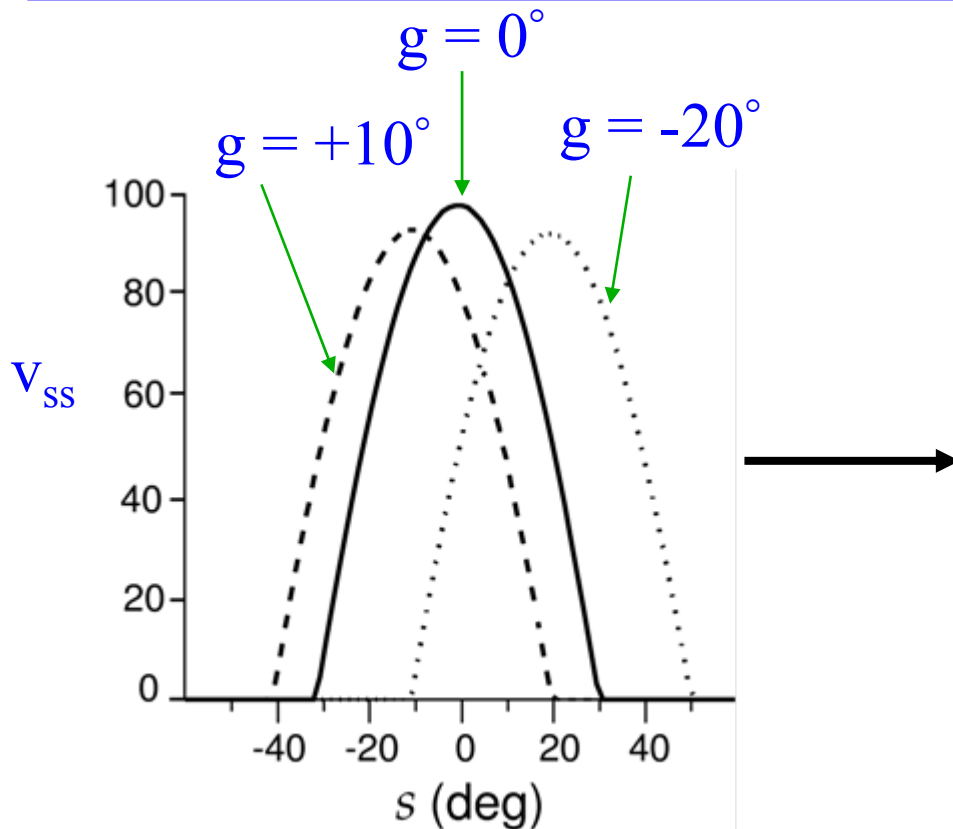
Output: Premotor Cortex Neuron with Body-Based Tuning Curves



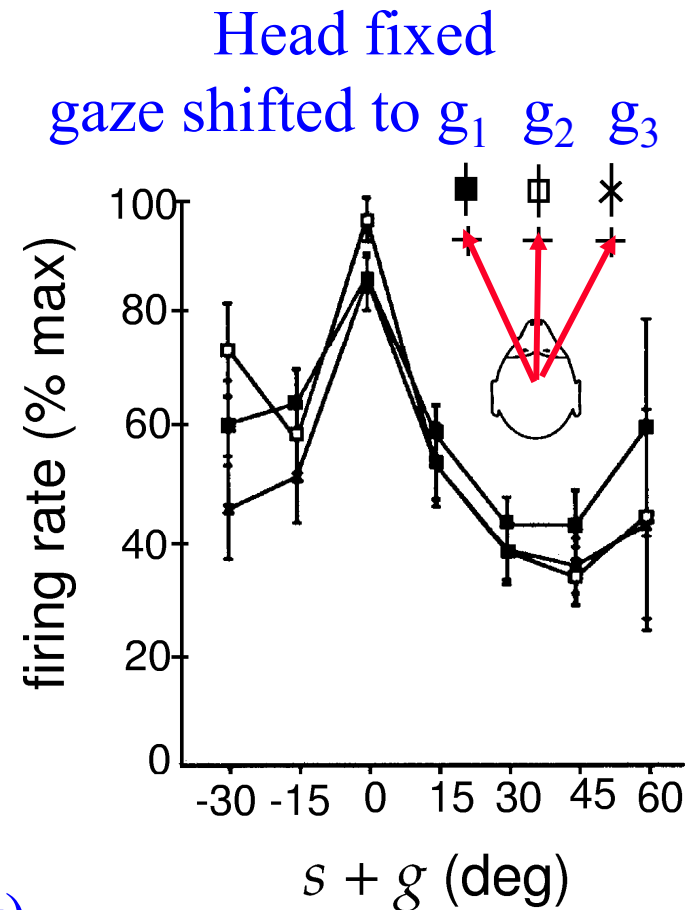
Input: Area 7a Neurons with Gaze-Dependent Tuning Curves

Weights $w(\xi, \gamma)$ need to be a function of $\xi + \gamma$

Output of a Simulated Feedforward Network



Retinal tuning curves shift to compensate for g (i.e. stable for $s + g$)



Next Class: Recurrent Networks

◆ Things to do:

- ⇒ Finish reading Chapter 7
- ⇒ Homework #3 due next Thursday May 14
- ⇒ Start working on mini-project