

Inference of Synaptic Weights in the Integrate-and-Fire Neuron Model

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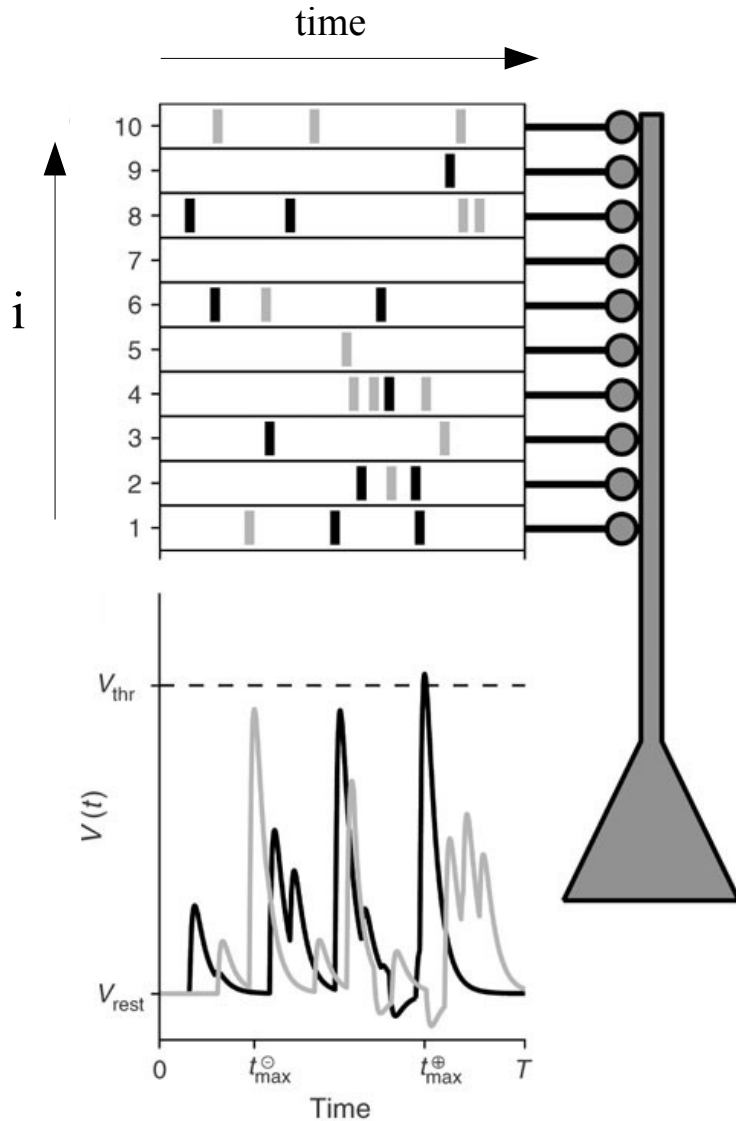
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A « Realistic » Neuron: the Integrate-&-Fire model



$$\frac{dV}{dt}(t) = -g V(t) + \sum_i J_i \sum_n C(t-t_{i,n}) + \eta(t)$$

Time-scale $\ll 1/g$

active

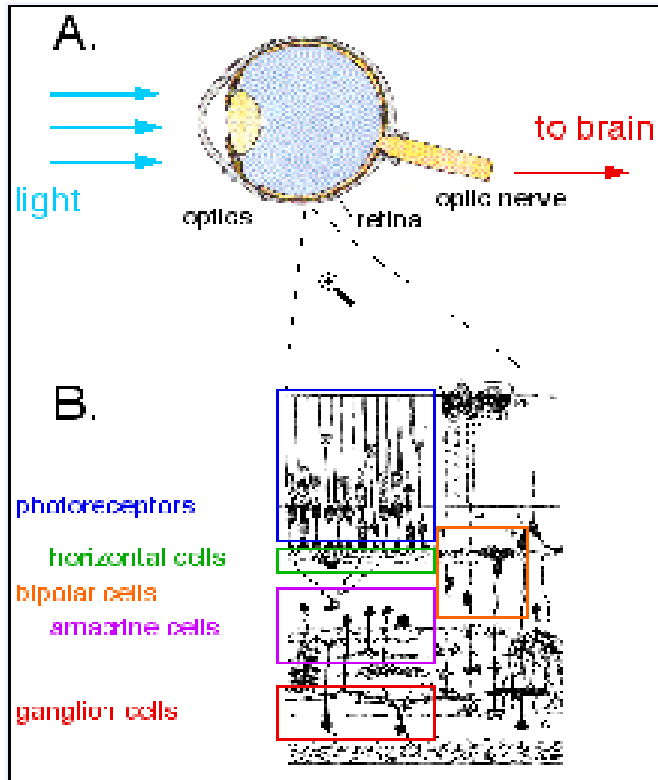


or

silent

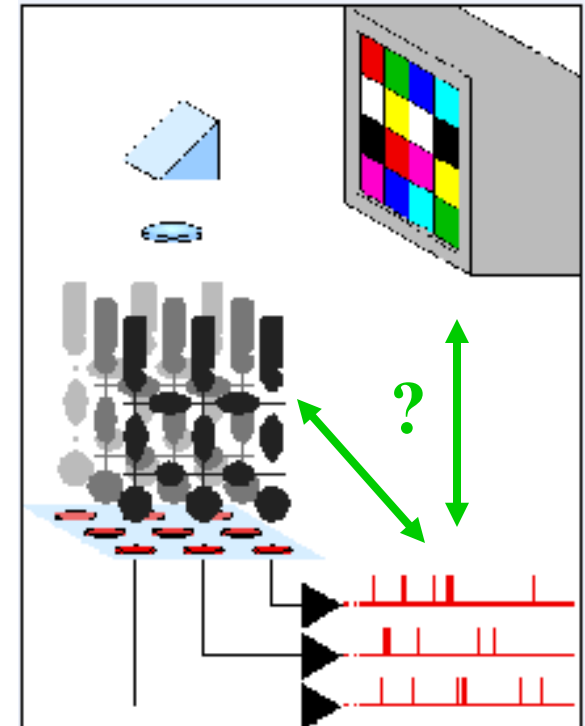


Multi-electrode recording from retina



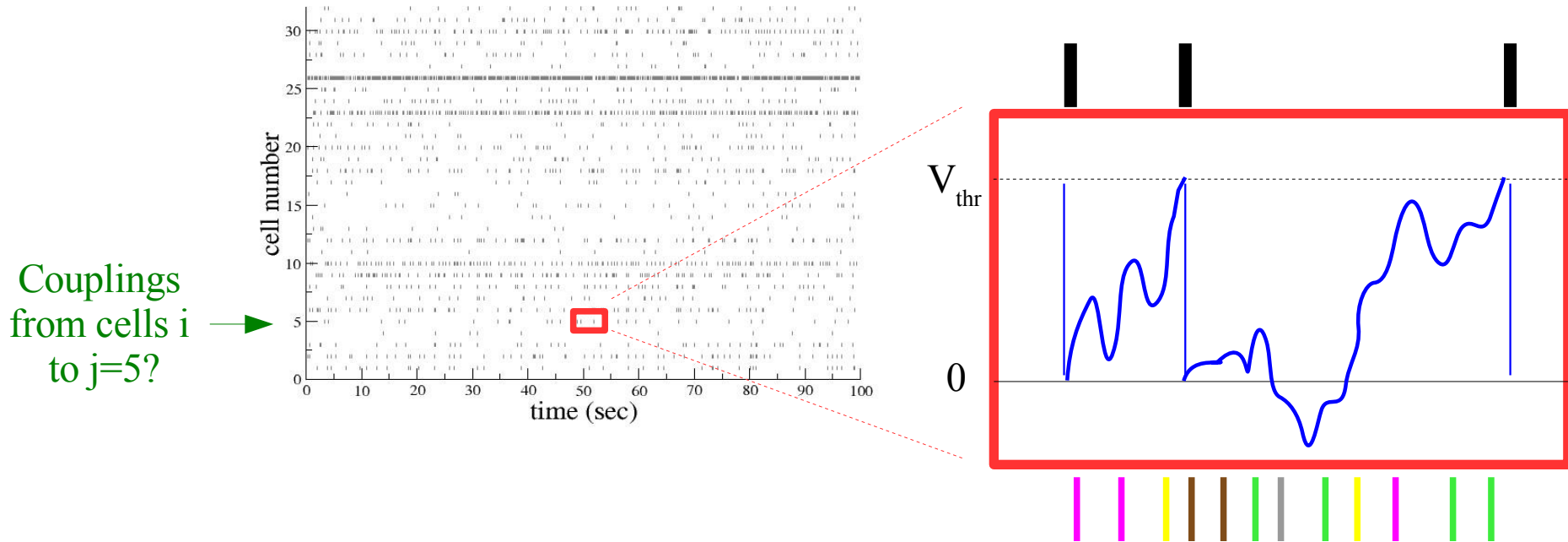
Berry et al. (2006): 40 ganglion cells, natural movies

Meister et al. (1997): 60 ganglion cells in the dark,
random flickering stimuli



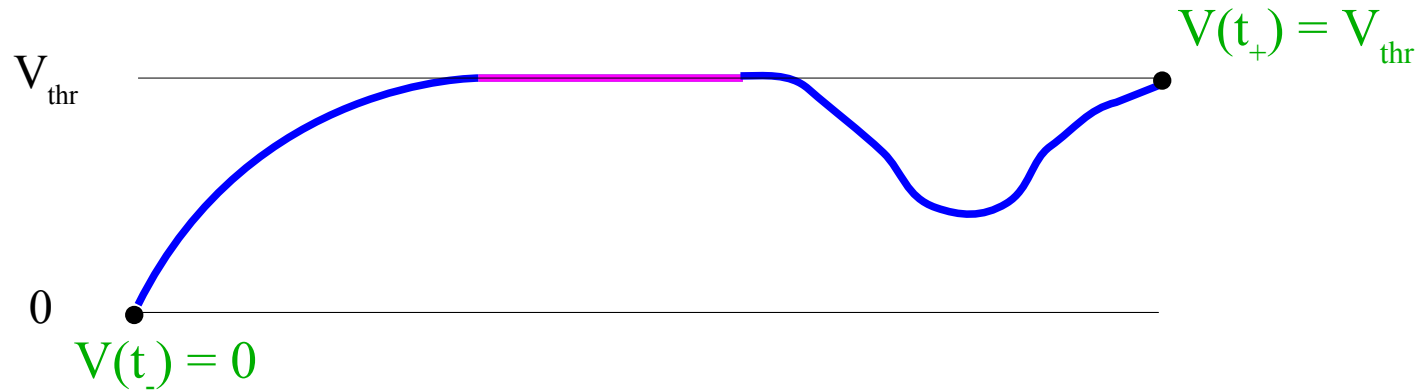
Experiments typically run for 1 hour

Inference of Synaptic Weights from Raster Plots



- $P[\{t_{i,n}\}|\{J_i\}] = \text{product of FPT probabilities}$
- $\{J_i\}$ maximizing $P[\{J_i\}|\{t_{i,n}\}]$?
- Instanton calculation (variance of noise $\rightarrow 0$):
optimal path for $V(t)$ conditioned by $\{t_{i,n}\}$

Optimal Paths for Potential and Noise



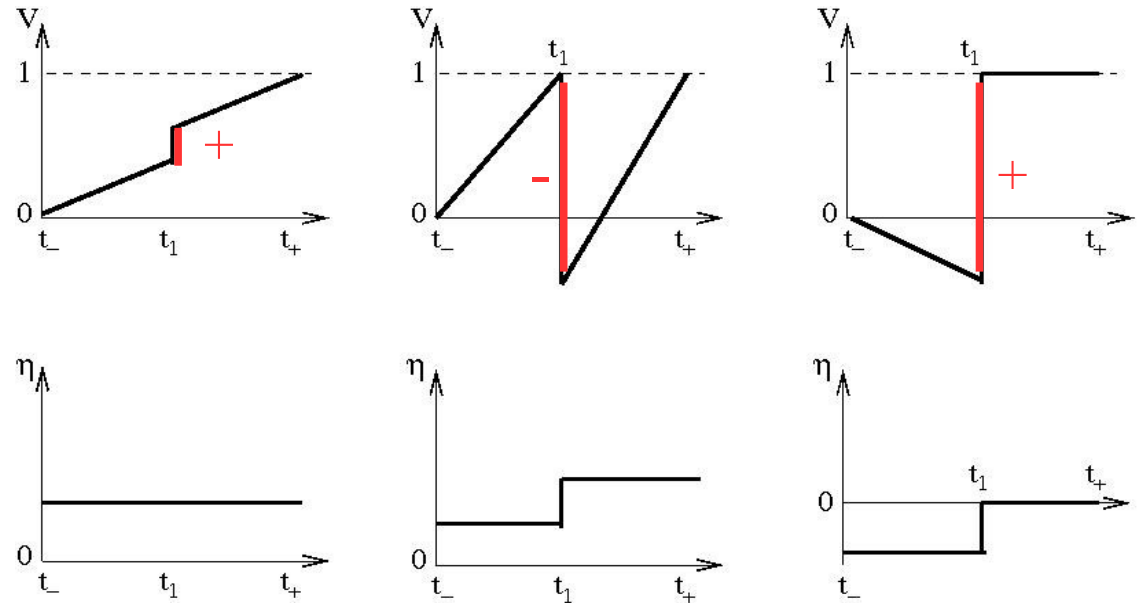
$$\left. \begin{aligned} \frac{dV}{dt}(t) &= -g V(t) + \sum_i J_i \sum_n C(t-t_{i,n}) + \eta(t) \\ \frac{d\eta}{dt}(t) &= +g \eta(t) \end{aligned} \right\} \text{or} \left\{ \begin{aligned} V(t) &= V_{thr} \\ \eta(t) &= g V_{thr} - \sum_i J_i \sum_n C(t-t_{i,n}) \end{aligned} \right.$$

- Where are contact intervals? ... not easy to find out ...
- Limit-case of instantaneous synapses: $C(t)=\delta(t)$
- $g = 0$ (not essential): noise is constant, except over contact intervals

- Intervals reduce to isolated points coinciding with (some of the) entering spikes
- Noise^{after} can take any value $>$ Noise^{before}

Looking for Contact Points

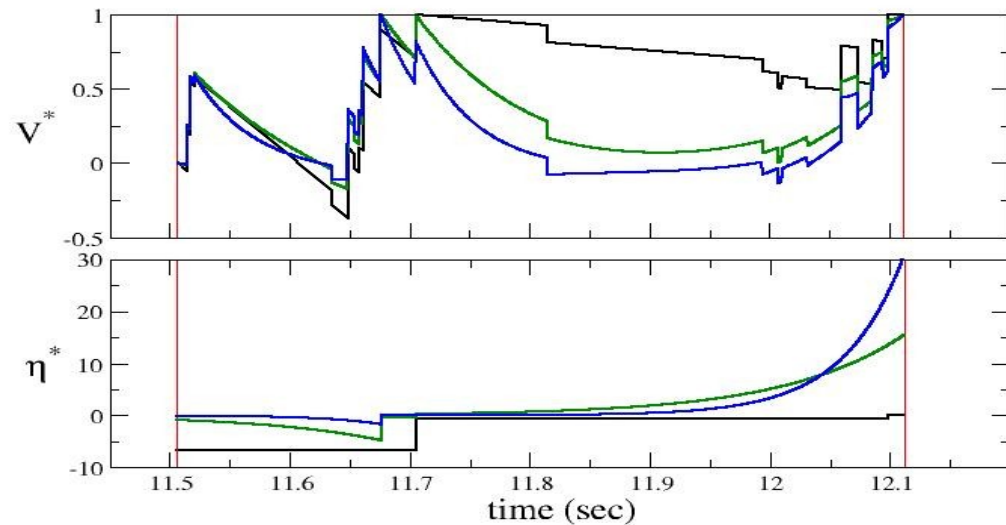
1 incoming spike



More incoming spikes

(20 spikes from 31 cells)

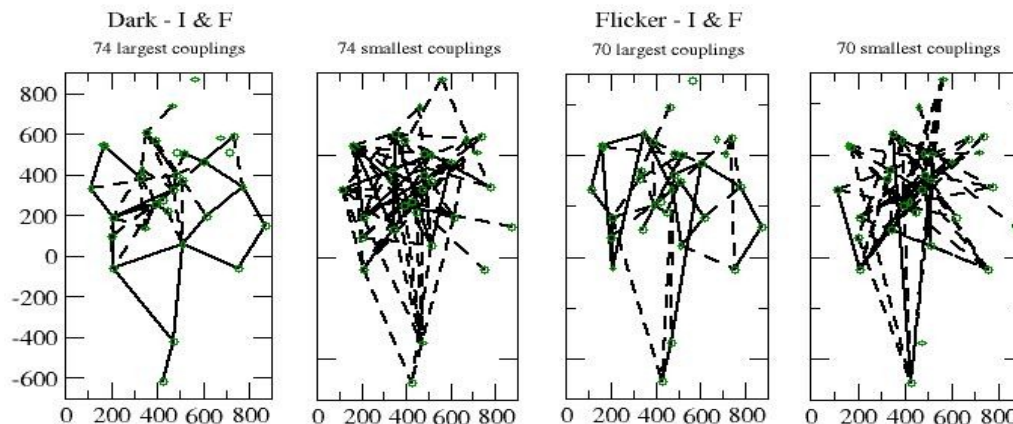
$g=0, 10, 20 \text{ sec}^{-1}$



Searching for Optimal Interactions

- Optimal action (log-likelihood) is a convex function of $\{J_j\}$ \rightarrow gradient descent
- Repeat for all cells \rightarrow matrix of interactions $\{J_{ij}\}$
- Applications to multi-electrode recordings of retinal ganglion cell activity:
(up to 60 cells so far \rightarrow 1770 couplings)

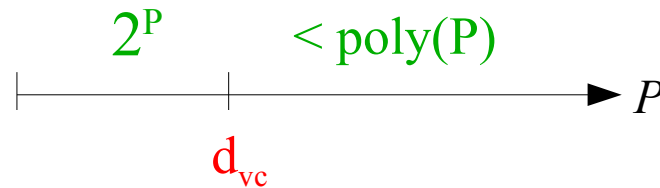
Allow us to study various aspects of interactions of biological interest:
spatial structure, symmetry, dependence on stimulus, on time-scales, ...



Computational Capabilities of the I&F neuron

Input activity $\xi_i(t) \rightarrow$ Output $\sigma = 0$ (silent) or 1 (active)

Set of P inputs $\{\xi_i^\mu(t)\}$ for $\mu = 1, \dots, P$: how many output sets $\{\sigma^\mu\}$ are realizable?



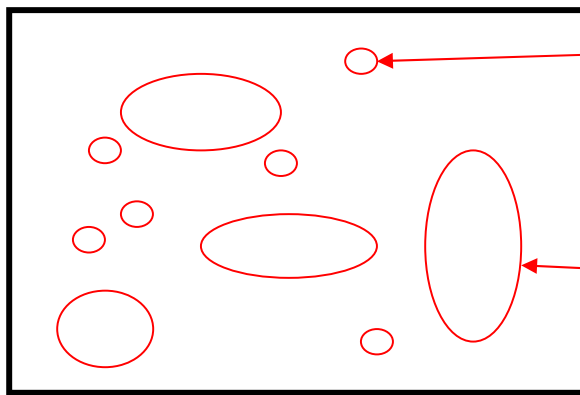
Vapnik
Chervonenkis
1971

Dimensionless time: $K = T \times g$

- $K \ll 1$: only $\xi_i^\mu = \langle \xi_i^\mu(t) \rangle$ matters \rightarrow perceptron
- $K \gg 1$? $1/\log K < d_{vc}/N < \log K \dots$

Gardner 1987
Gardner, Derrida 1988

I&F Neuron and Internal Representations

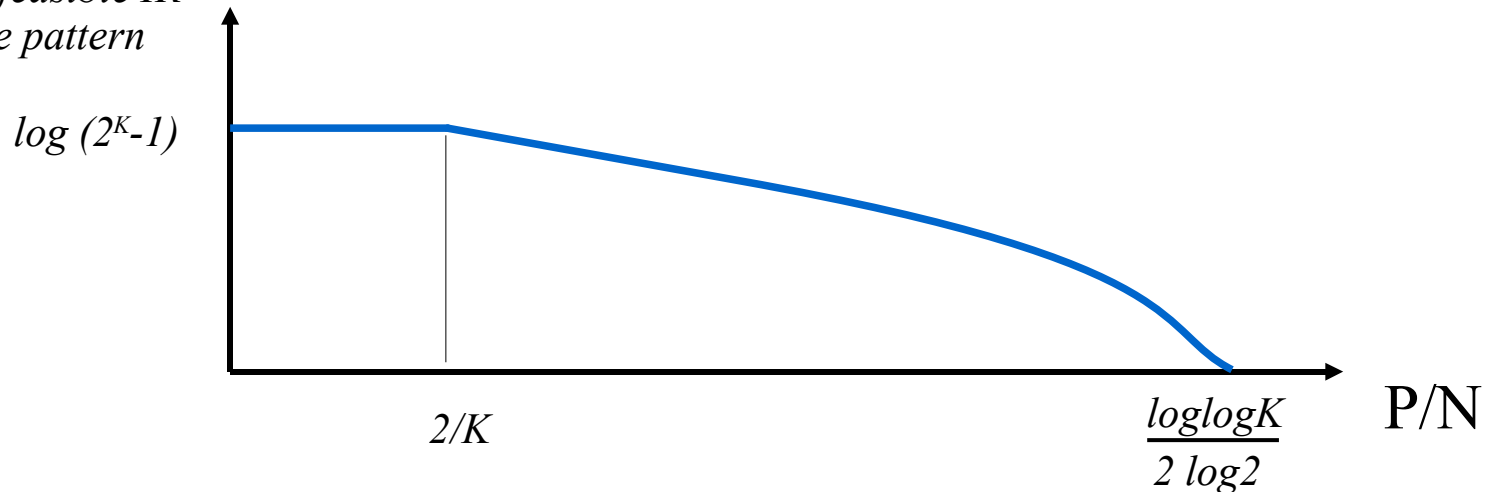


{ active pattern 1: 00011000
 active pattern 2: 01100100

{ active pattern 1: 00101000
 active pattern 2: 00010000

How many domains are not empty i.e. IR are feasible?

*Entropy of feasible IR
per active pattern*

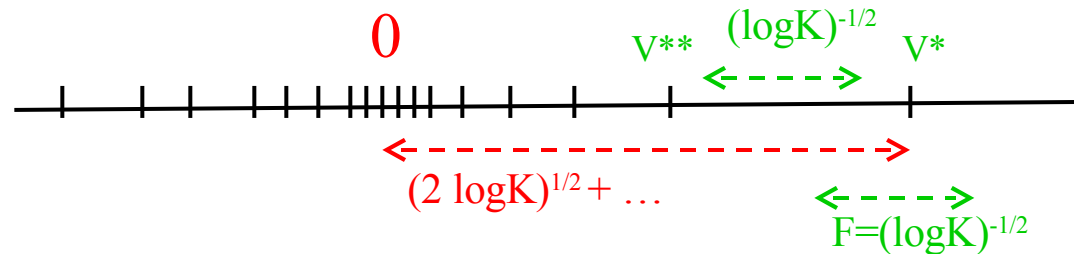


Robustness against Time and Space Correlations

$$\frac{d_{vc}}{N} = \frac{\log \log K}{2 \log 2}$$

obtained for uncorrelated patterns (in time, space) ...

**Extreme
Value
Statistics**



very robust against correlations! (Berman, 1964)