



Sainsbury Wellcome Centre

March 2, 2018

Cerebellar learning

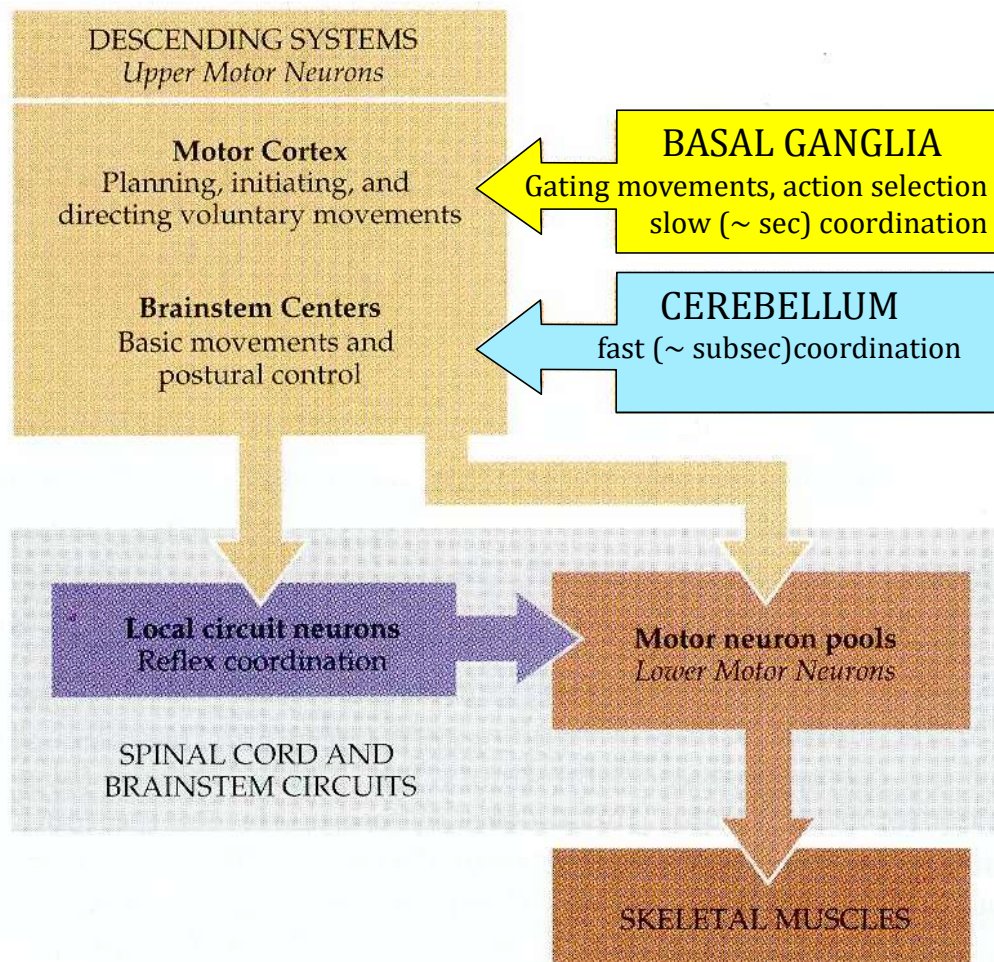
Prof. Tom Otis
t.otis@ucl.ac.uk



- Brief overview of cerebellum
- Behavioural aspects of cerebellar associative learning
- A circuit mechanism and theoretical model
- Cellular mechanisms

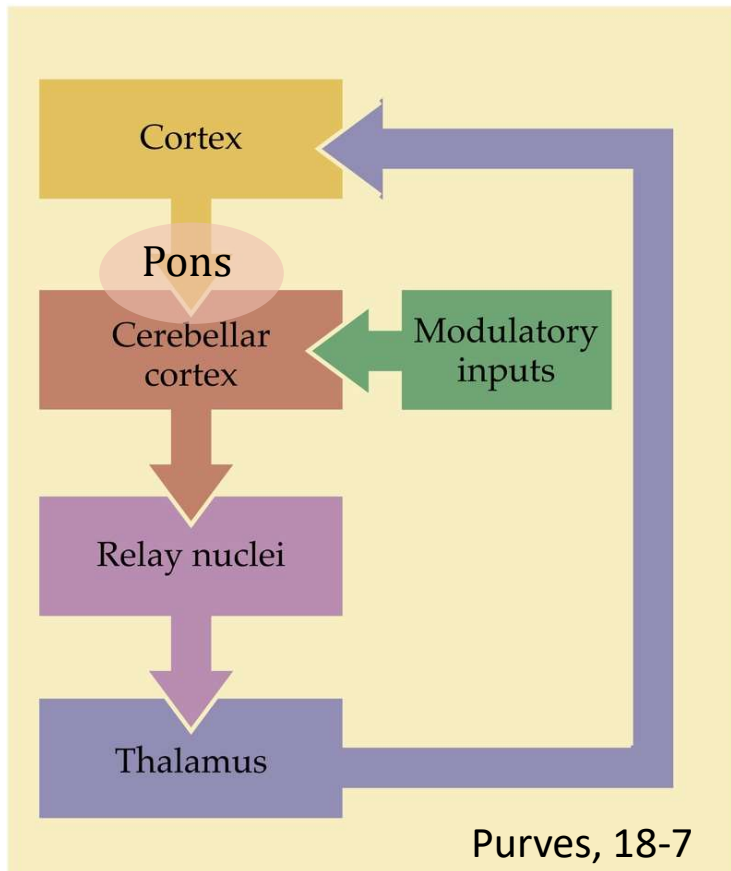
A simplified view of motor system output

The cerebellum functions as a rapid, corrective feedback loop, smoothing and coordinating movements.



from Fig. 15-1, Purves

Fast feedback loops for coordinating movement



Cerebellar lesions cause:

nystagmus

ataxia

dysdiadochokinesia

dysmetria

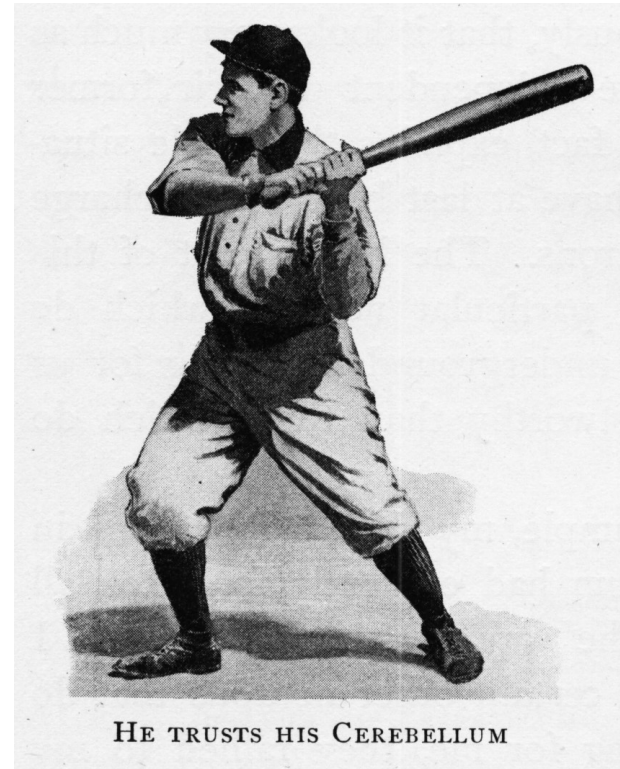
intention tremor

also, deficits in

motor learning

What kinds of information does the cerebellum receive?

- somatosensory
- visual
- auditory
- vestibular
- proprioceptive
- *efferent copy*



From *Control of Body and Mind*,
Gulick Hygiene Series, 1908

Movement is fast & nerves are slow coordination requires *prediction*

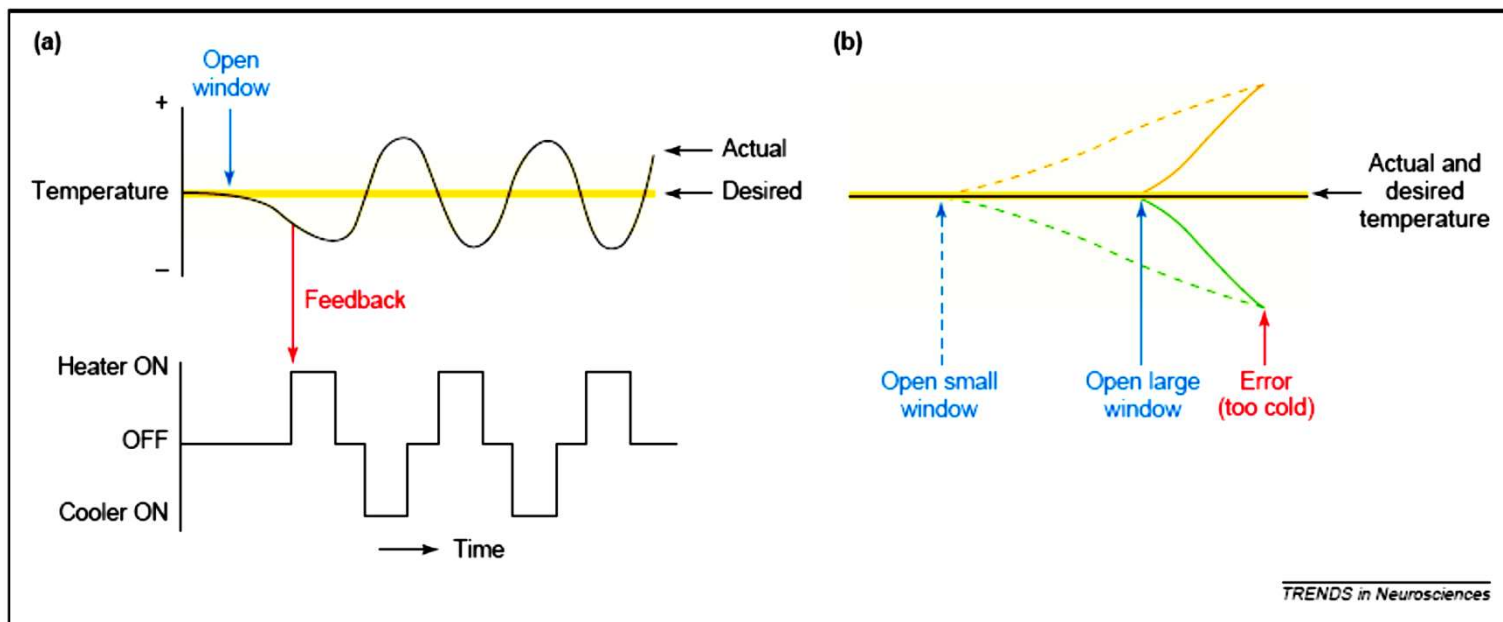


Usain Bolt, 100 m WR: 9.58 s

conduction velocity of most nerve fibers is ~ 10 m/s

some humans run at ~ 10 m/s

For it to be adaptive, control must be “feedforward”



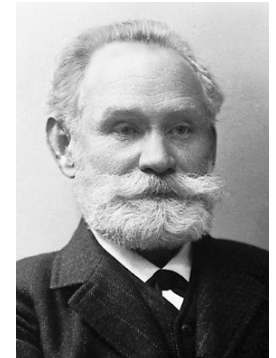
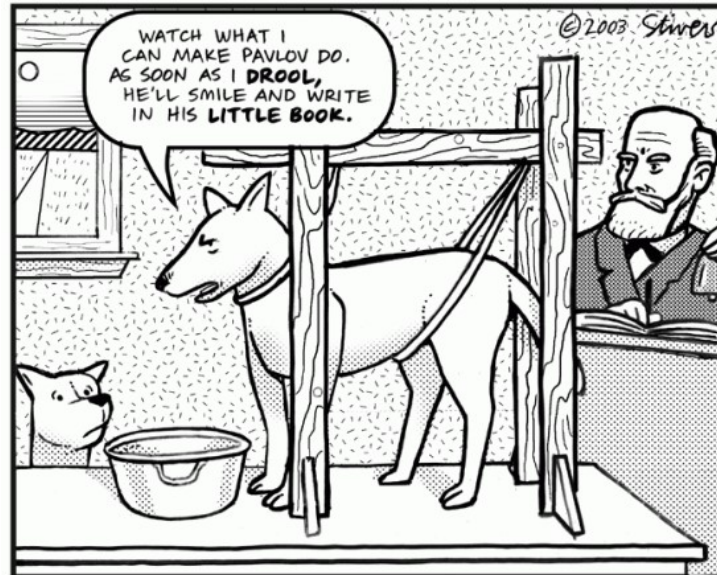
Ohyama et al., 2003

Behavioural aspects of cerebellar associative learning

Classical or Pavlovian conditioning

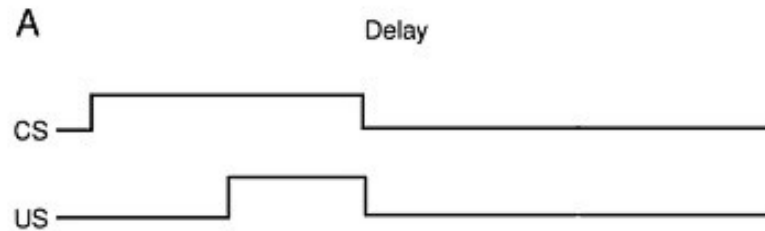
A form of associative learning in which a conditioned stimulus (CS) is linked to an unconditioned stimulus/response (US/UR).

After learning the CS elicits a conditioned response (CR) when delivered by itself.

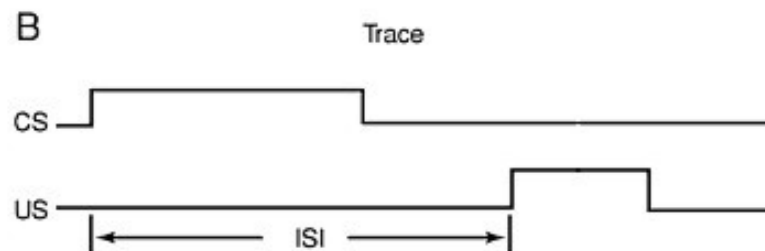


Ivan Pavlov
Nobel Prize, 1904

Paradigms for classical conditioning:

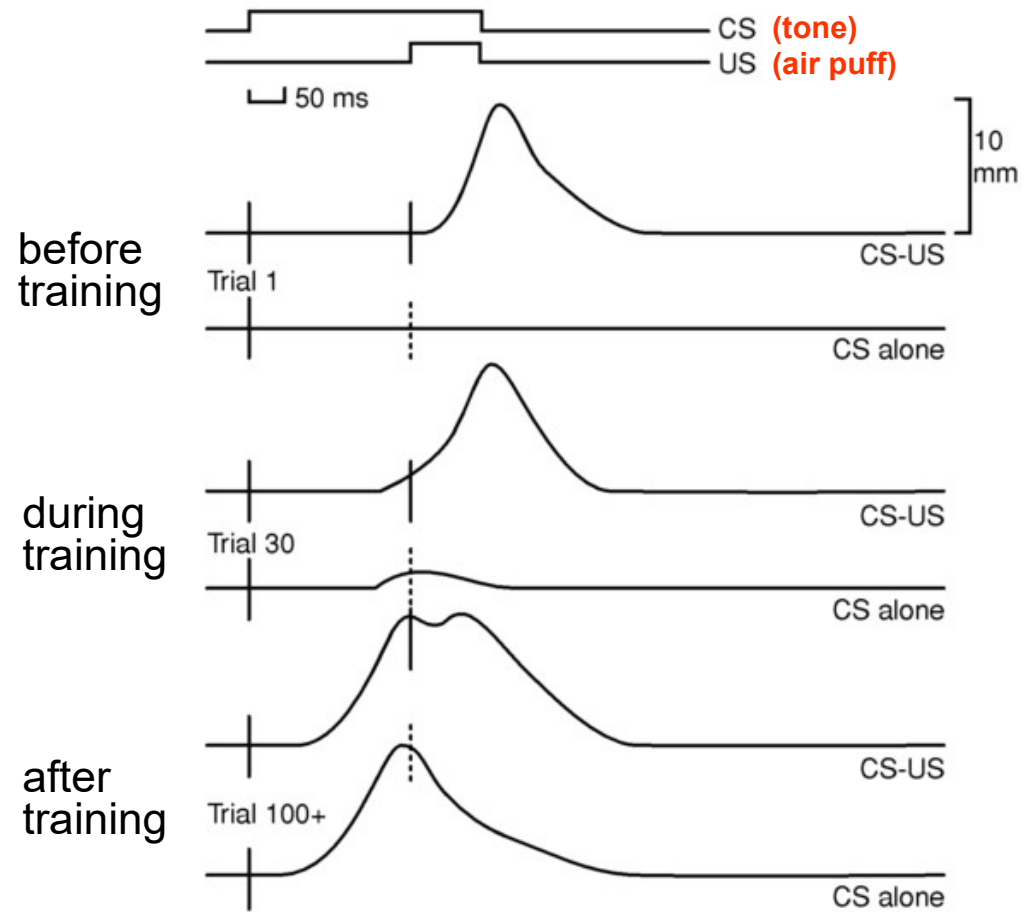


Cerebellar lesions disrupt delay conditioning



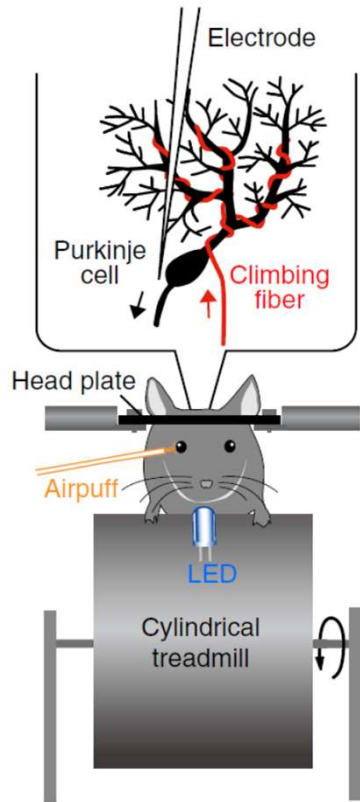
Both cerebellar and hippocampal lesions disrupt trace conditioning

Eyelid movements during a classical conditioning experiment

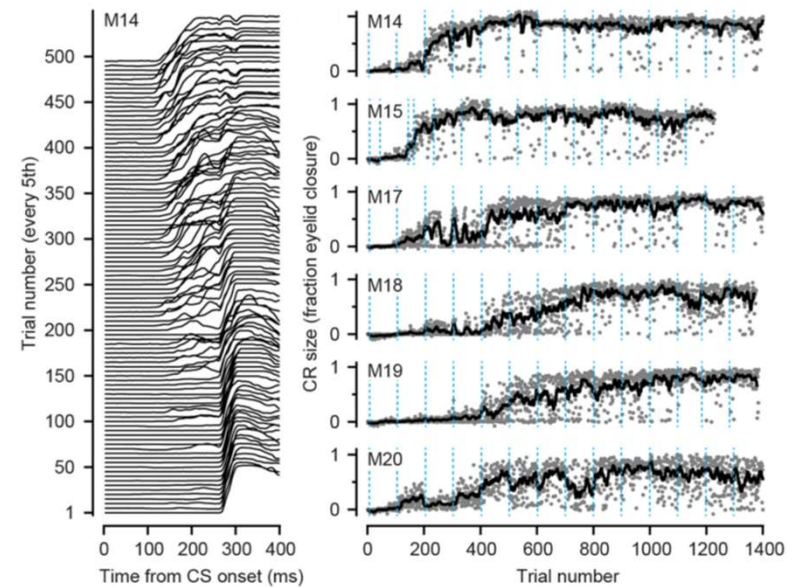
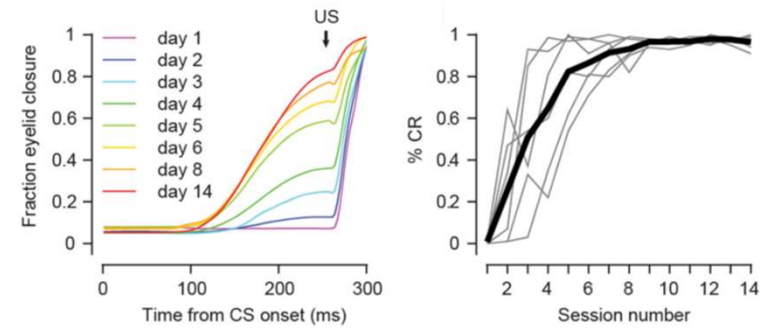


Zigmond et al., 1999

Mouse eyeblink data

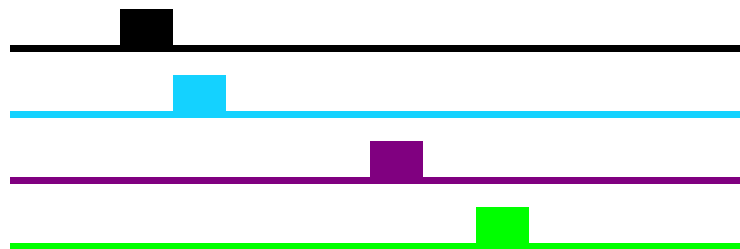
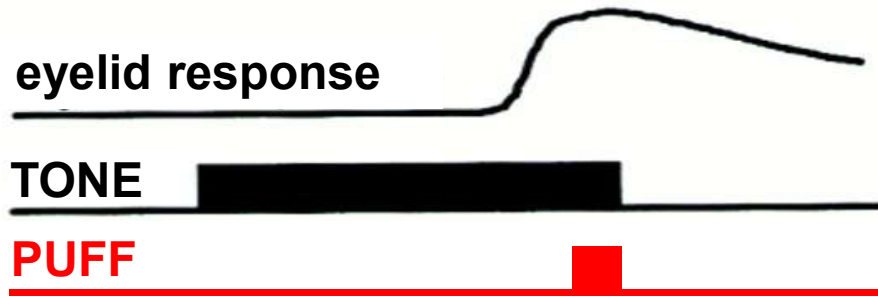


250 ms CS: LED US: Airpuff

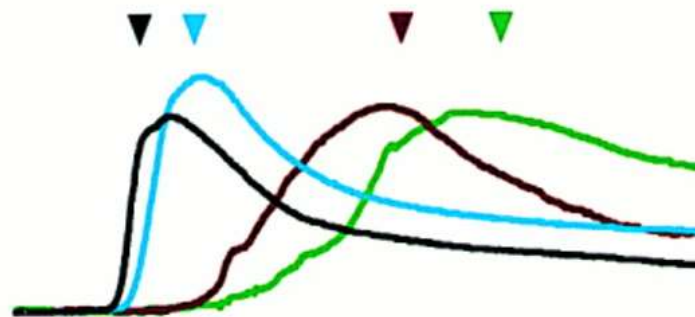


Heiney et al, *J. Neurosci.*, 2014

Timing of learned responses dictated by CS-US timing during training



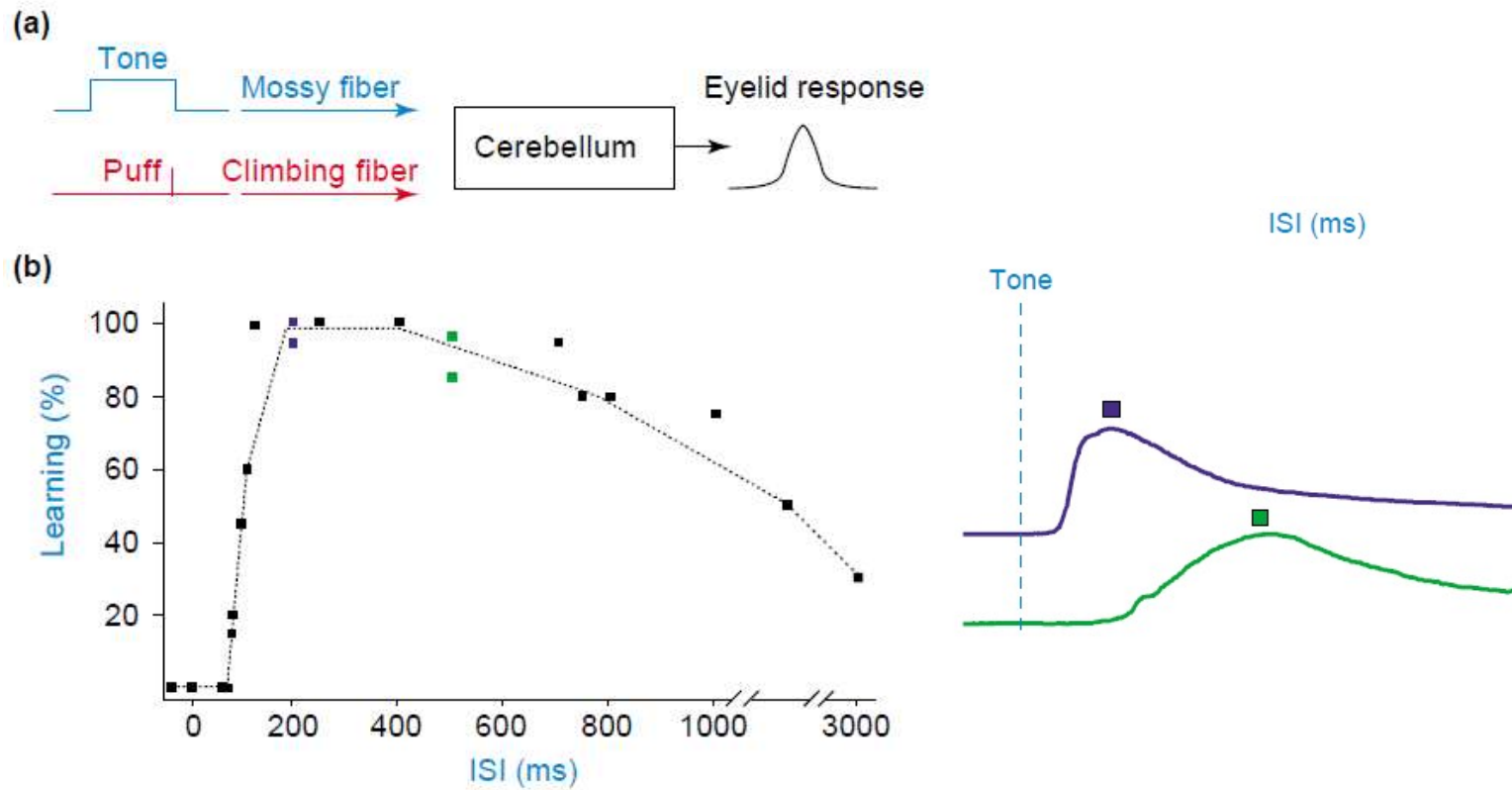
differently timed
puffs during
training



responses after
training

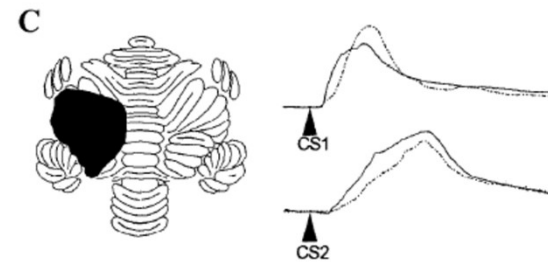
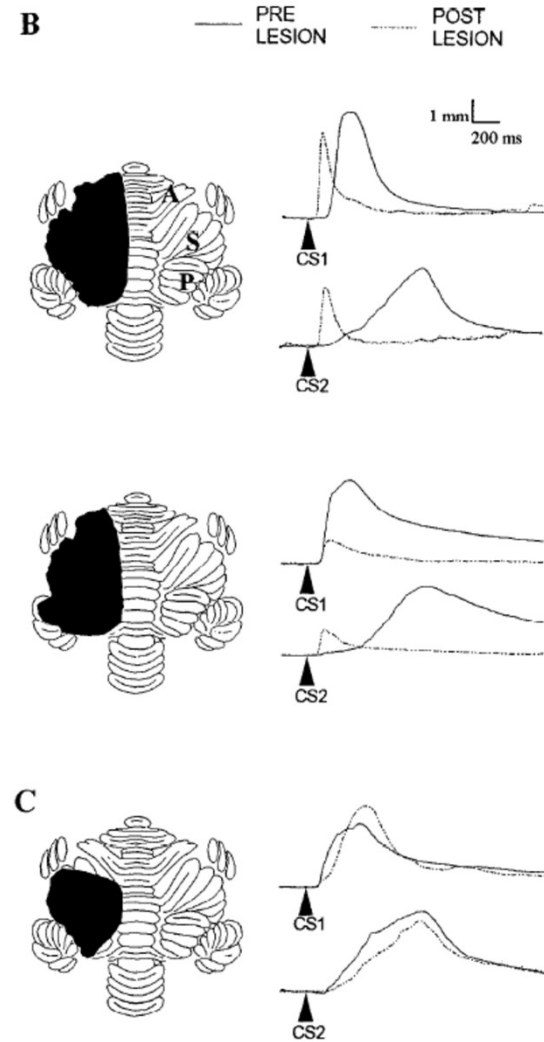
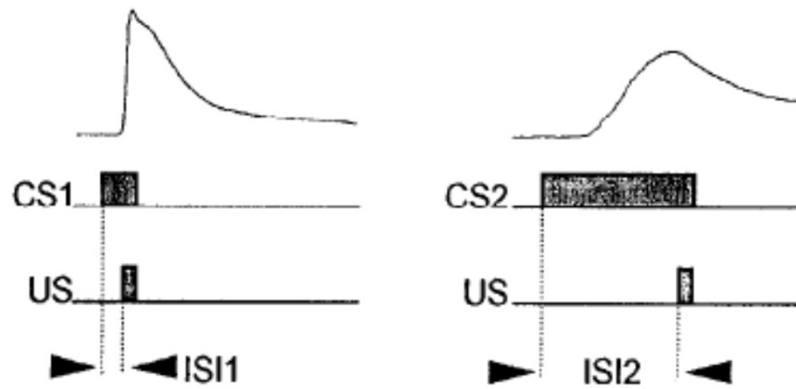
from Mauk et al.,1998

Learning is robust for CS-US intervals of 100 ms to 1 second



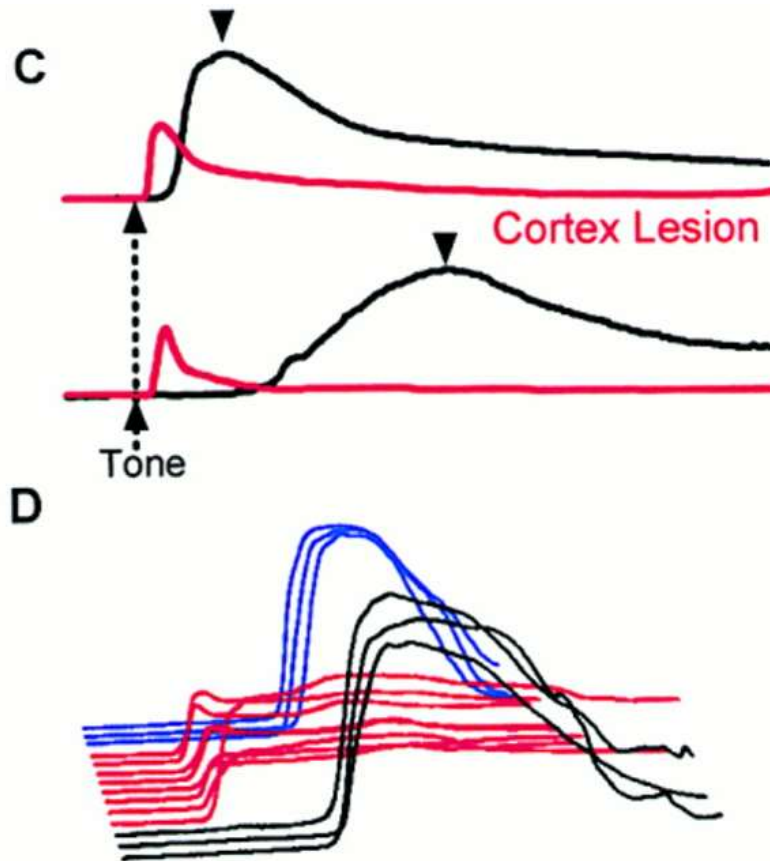
Ohyama and Mauk 2003

Lesions of cortex alter but do not block memories



Perrett et al., *J. Neurosci.* 13:1708, 1993

Lesions and pharmacological inactivation of cerebellar cortex cause improperly timed learned responses after eyeblink conditioning.

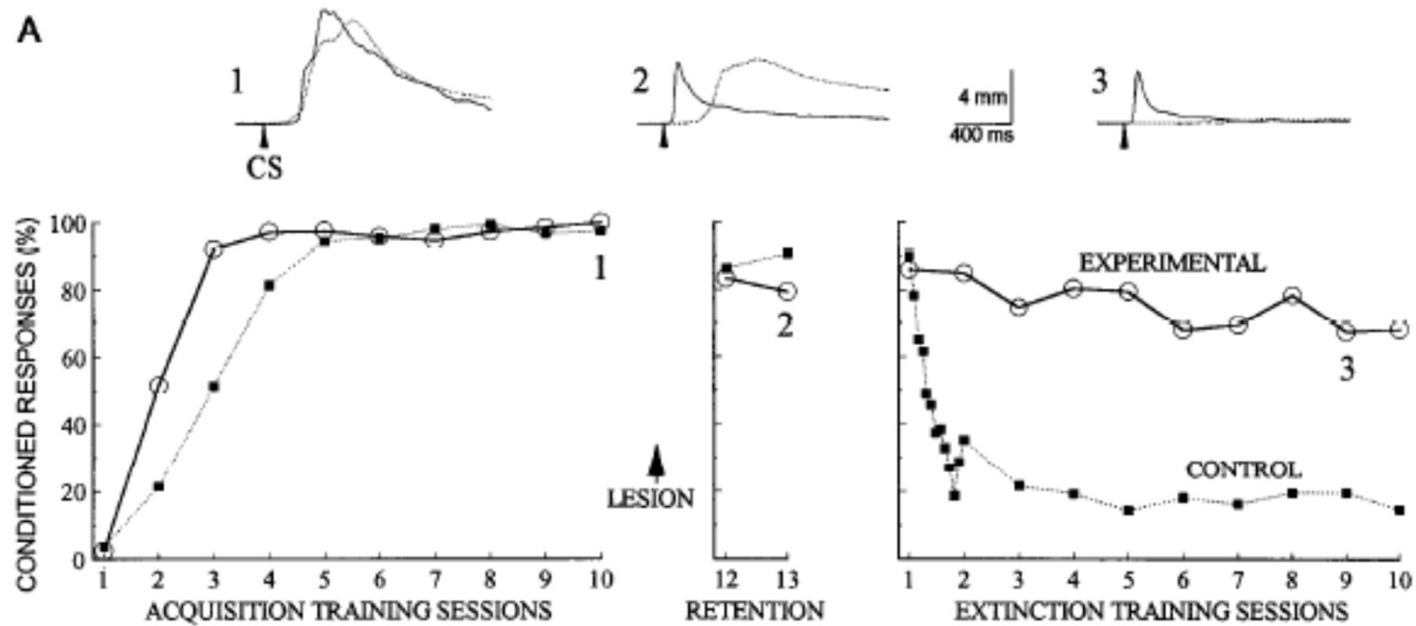


Lesions of cerebellar cortex (anterior lobe)

GABA_A receptor antagonist (picrotoxin) injected into interpositus nucleus

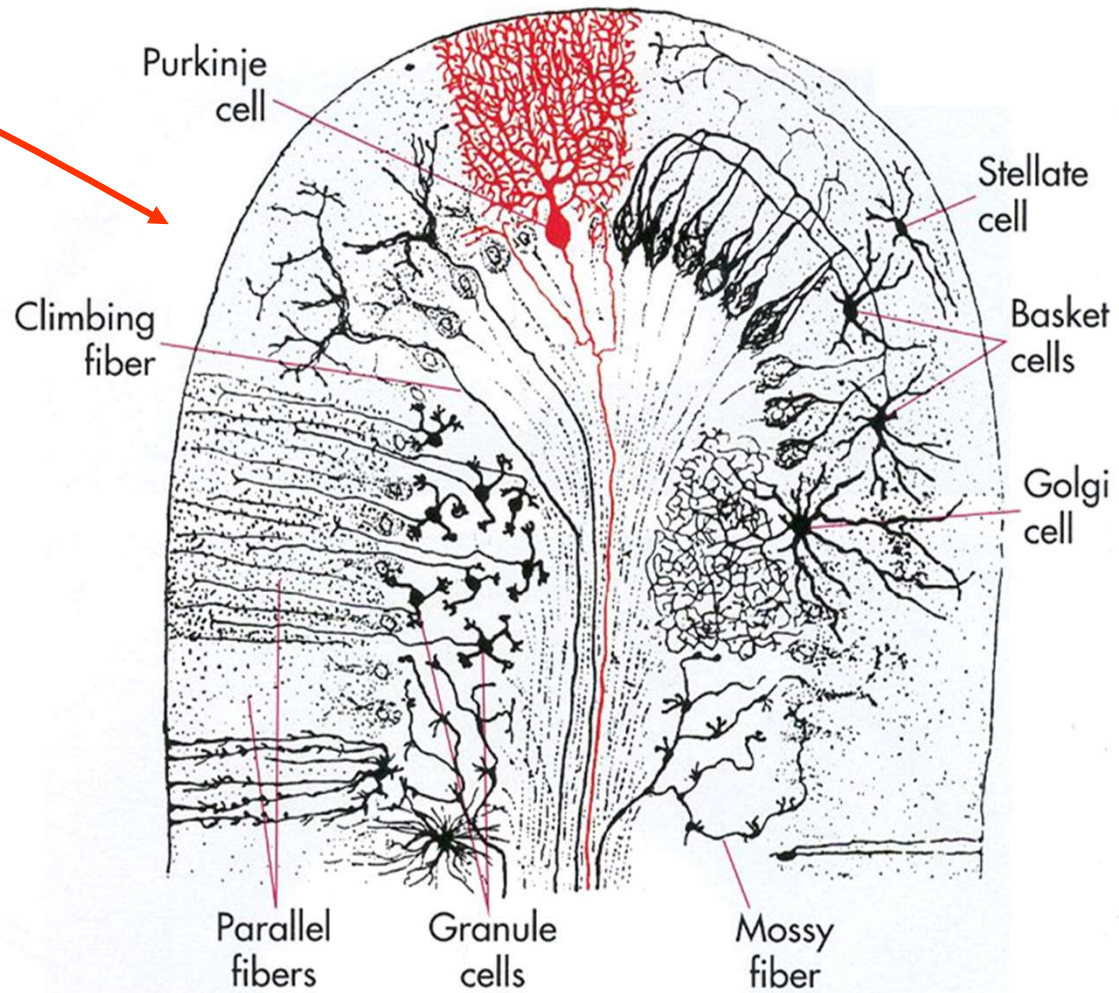
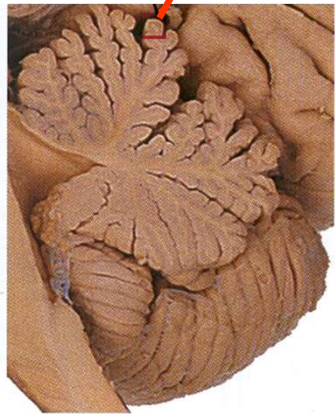
Mauk et al., 1998

Extinction requires the cortex

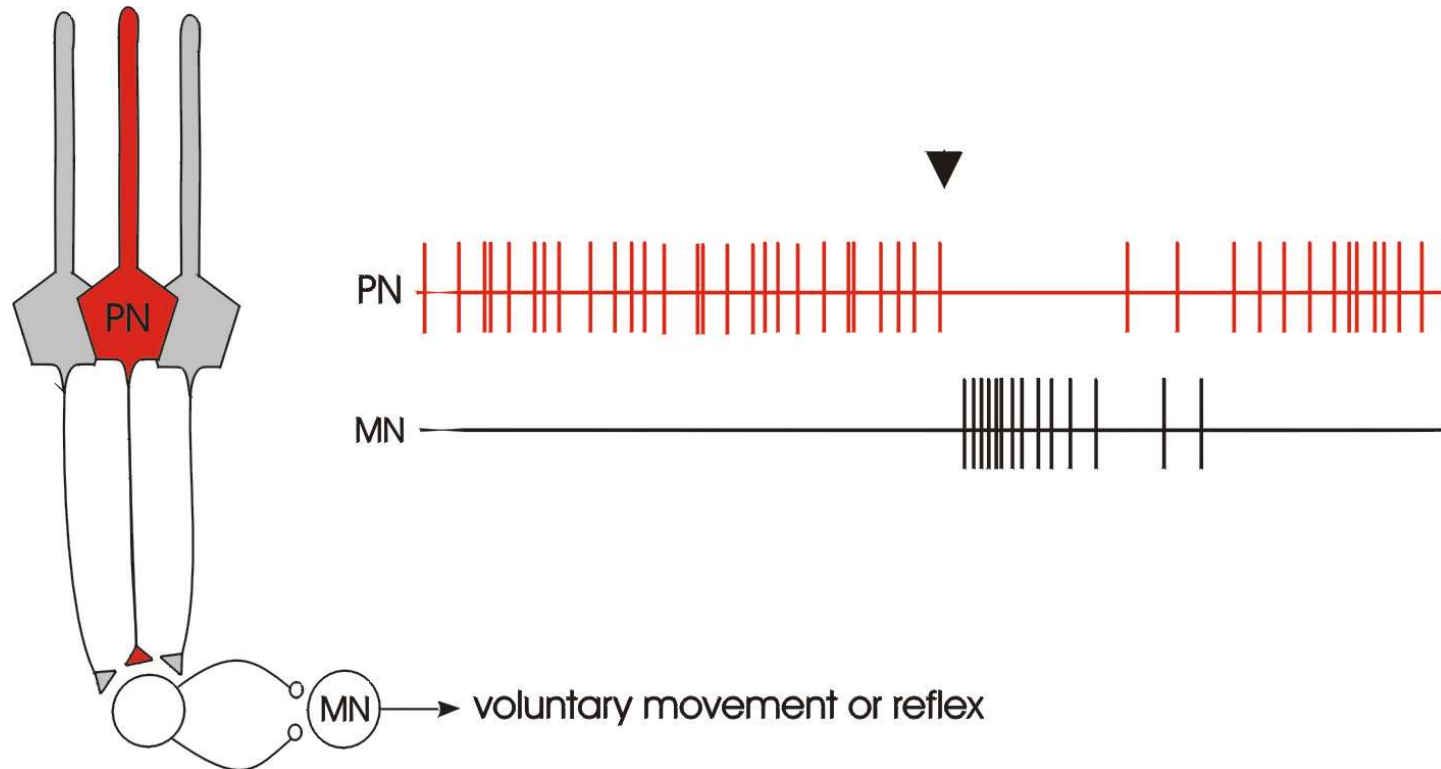


Perrett and Mauk, *J Neurosci.* 15:2074, 1995

Cellular anatomy of cerebellum



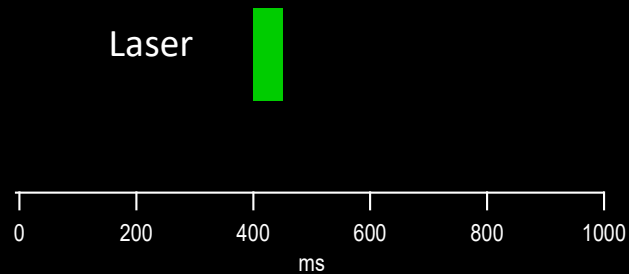
How does Purkinje neuron firing affect movement?



Purkinje neurons are inhibitory, thus when they slow or stop firing their targets are excited

Rapid, short latency arm movements triggered by brief PN inhibition

- Archearhodopsin (inhibitory opsin) expressed in PNs
- Optic fiber delivering 532nm laser light to forelimb region of cerebellar cortex

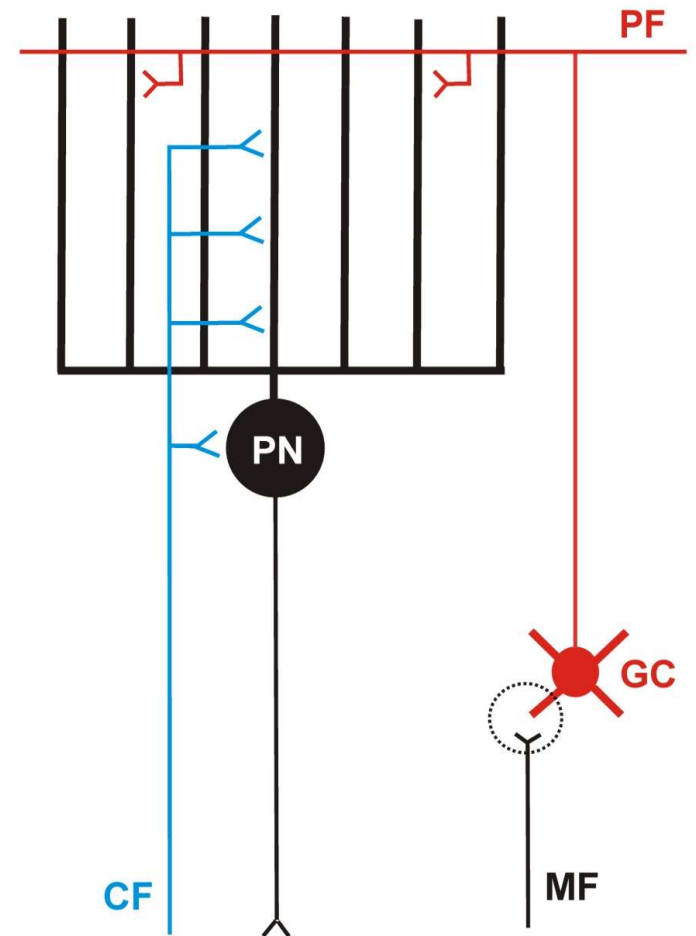


Circuit hypotheses for cerebellar associative learning

Two inputs to cerebellar cortex transmit distinct types of information

Mossy Fiber (MF) – **Parallel Fiber (PF)** system
the “sensorimotor context”

Climbing Fiber (CF) –
*the instructive signal, unexpected events
relevant to movement*



Some numbers: mossy fibers and climbing fibers

A mossy fiber excites
~30 granule cells.

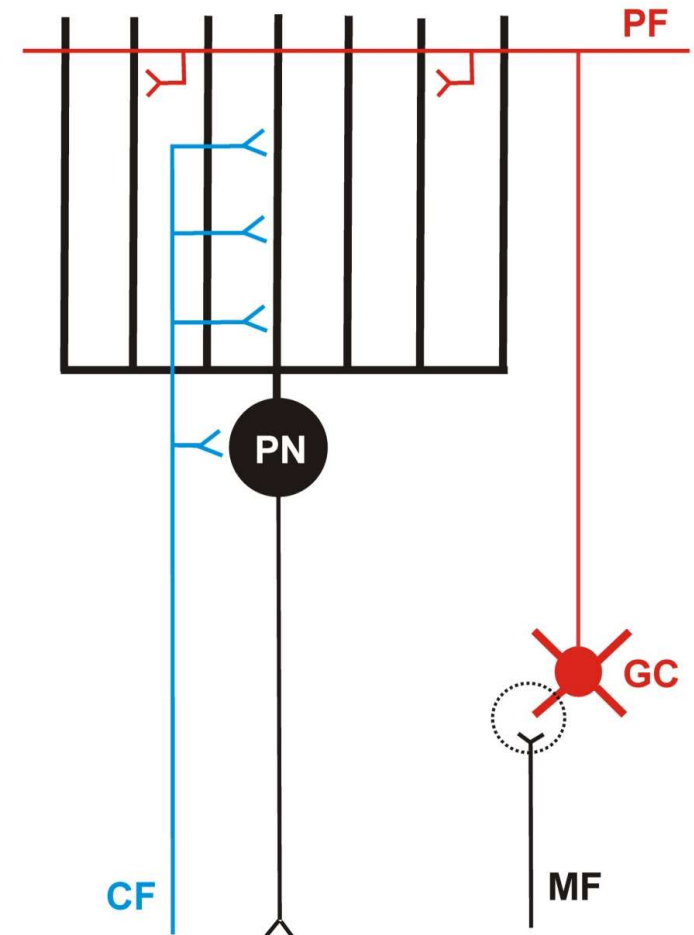
A granule cell is excited by
4-6 mossy fibers.

A parallel fiber excites ~300 PNs.

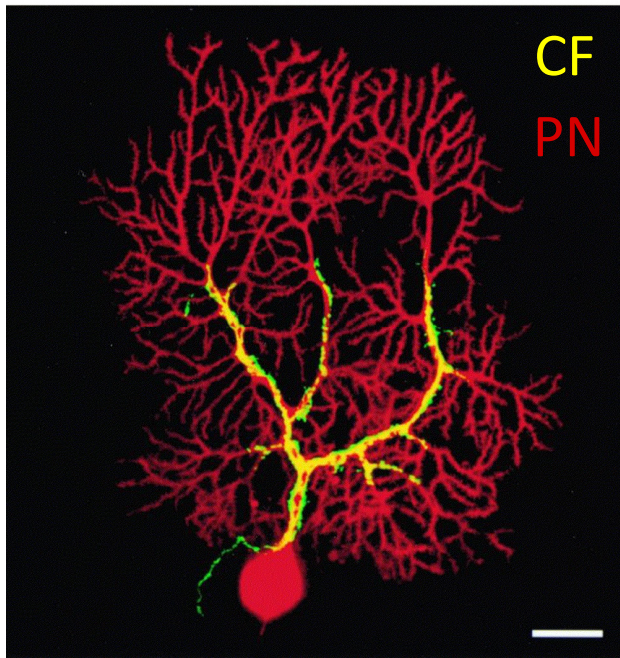
A PN is excited by ~100,000
parallel fibers.

A climbing fiber excites ~10 PNs.

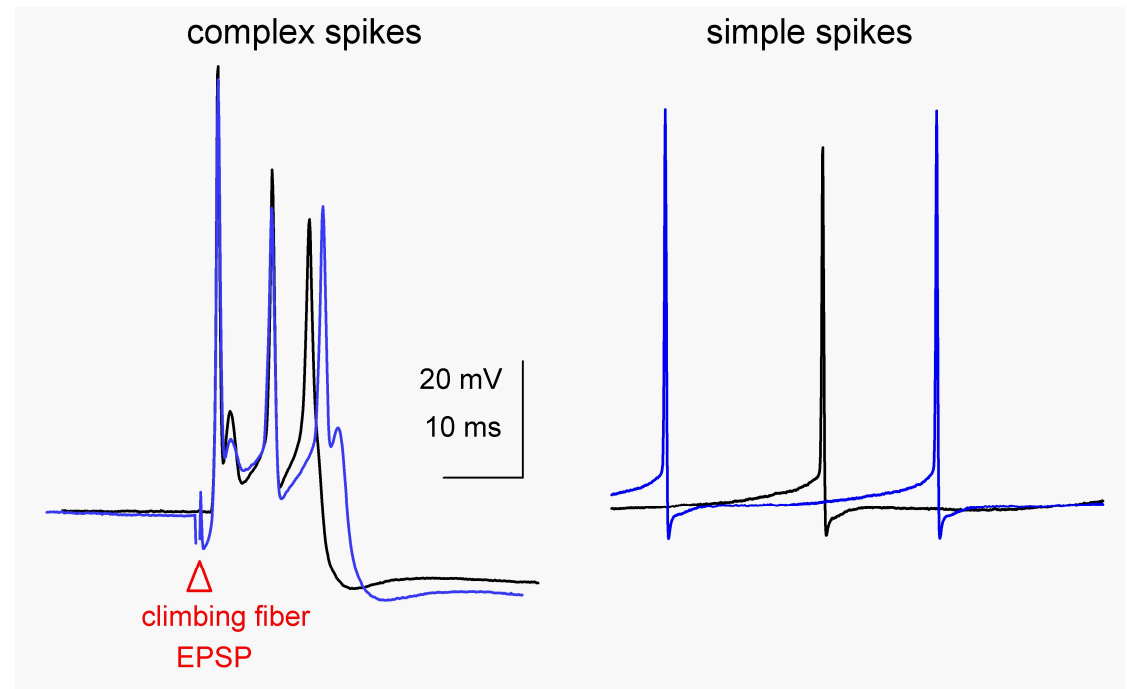
A PN is excited by 1 climbing fiber.



CFs generate a unique, cell-wide signal

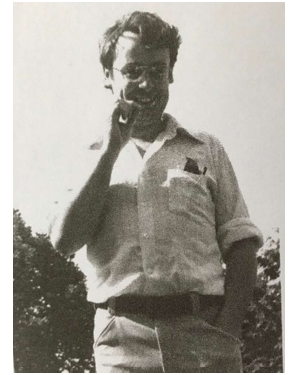


Kreitzer et al, 2000

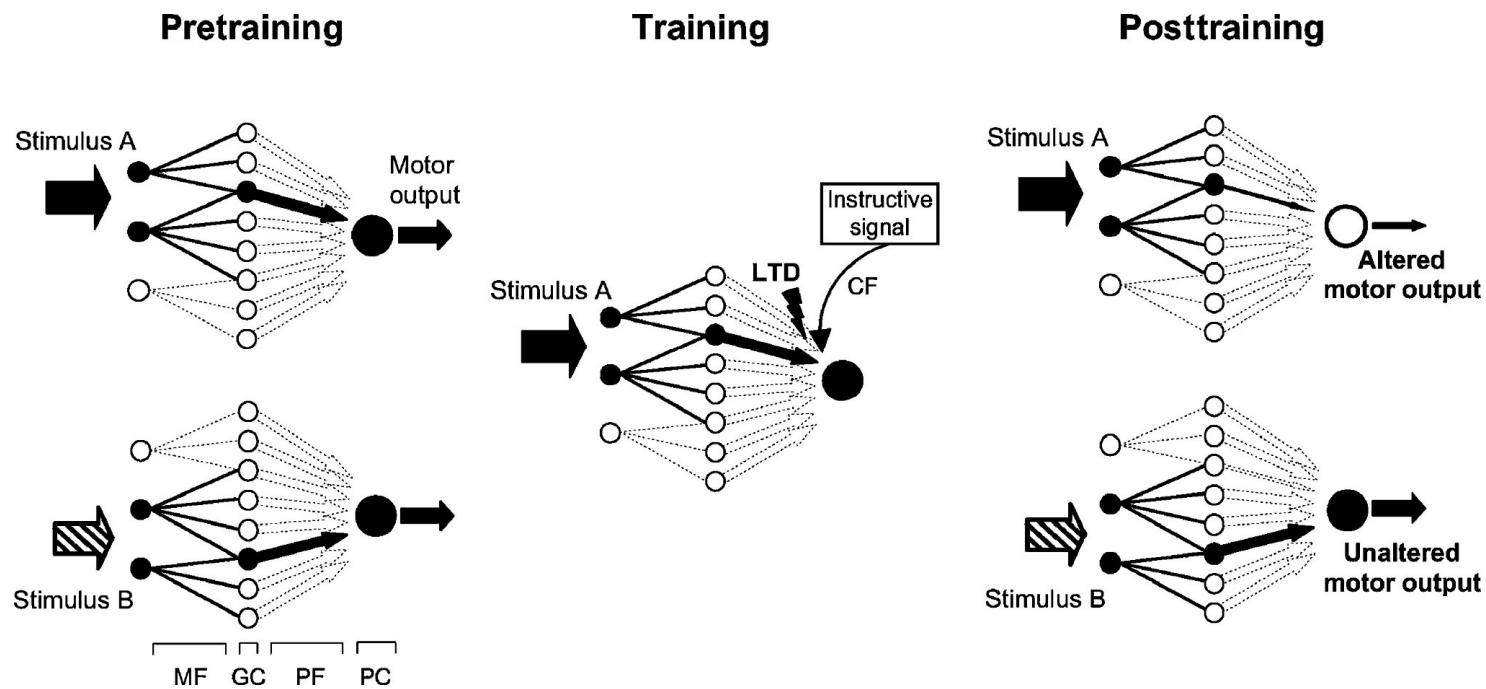


- Simple spikes are typical action potentials.
- Complex spikes occur in response to climbing fiber excitation.

The Marr/Ito/Albus model



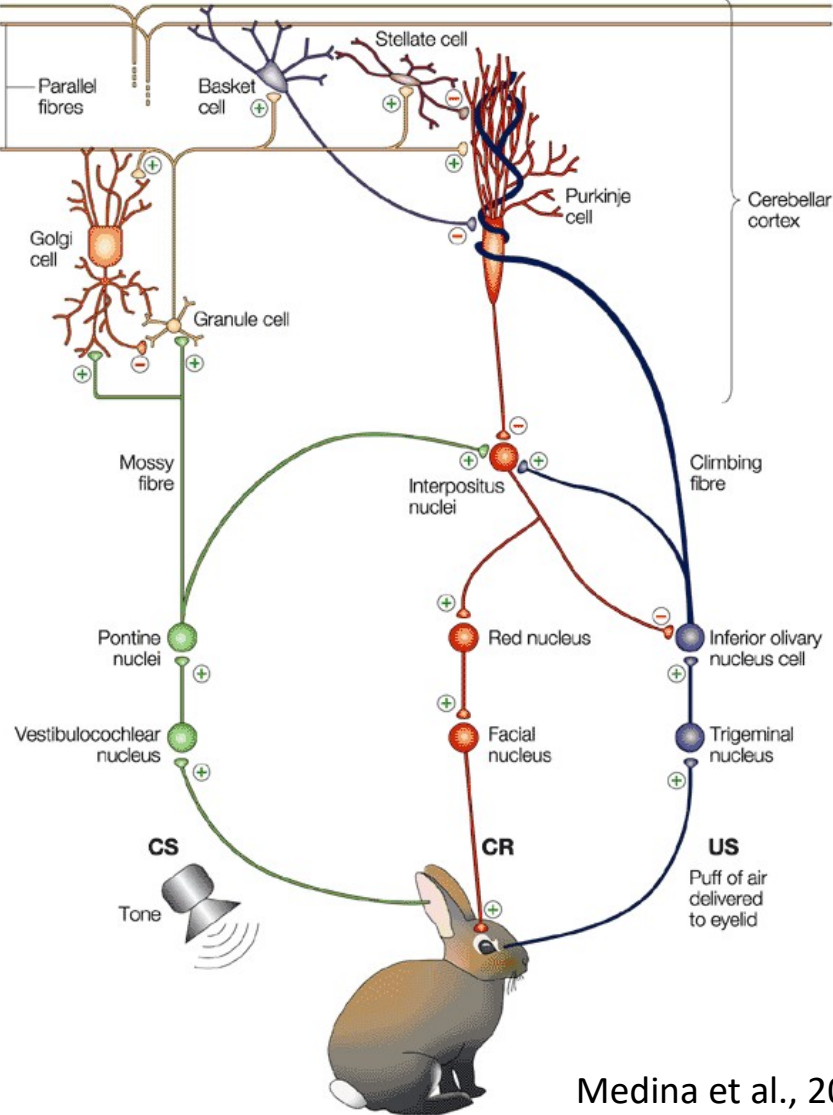
David Marr, 1970



from Boyden et al., 2004

for more on 'expansion recoding' see Kennedy et al., *Nat. Neurosci.*, 2014

Eyeblink conditioning circuitry



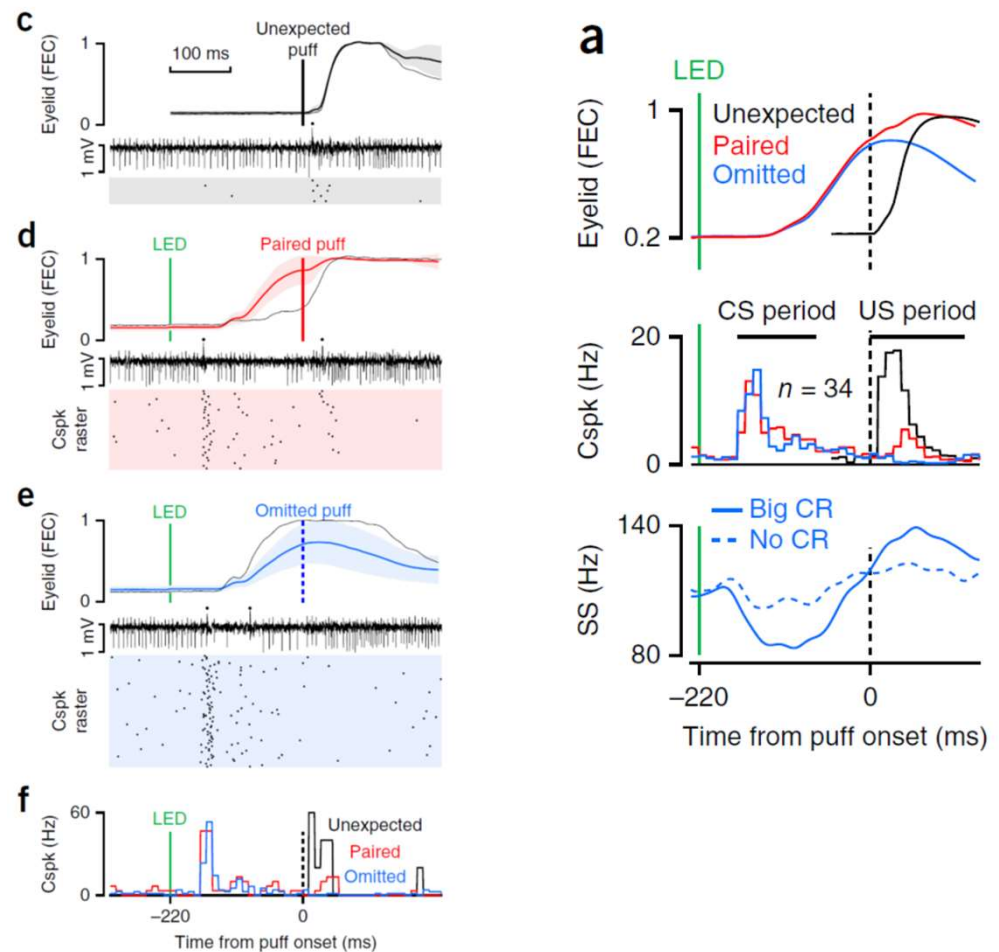
Medina et al., 2002

Evidence for the anatomical substrates of CS and US

- Lesions of the mossy fibers prevent learning (McCormick & Thompson, '84)
- Stimulation of the mossy fibers (pons) can substitute for the CS (Steinmetz et al, '89)
- Lesions of the olive (climbing fibers) prevent learning
- Stimulation of olive can substitute for the US (Mauk et al, '86)
- Inactivation of the climbing fibers extinguishes learning

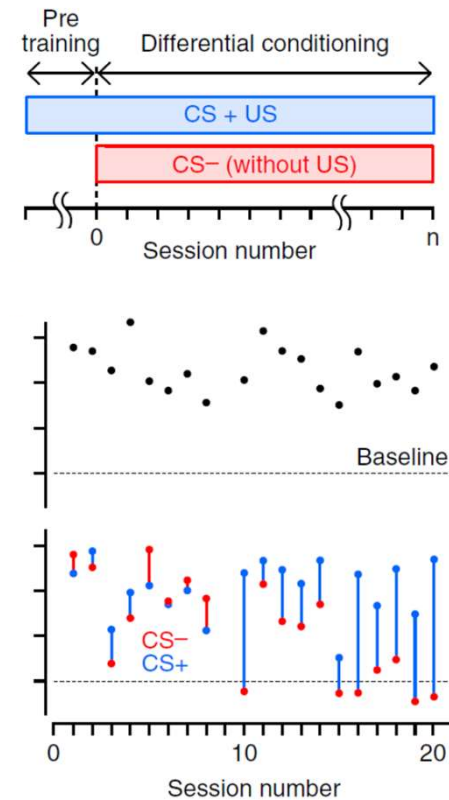
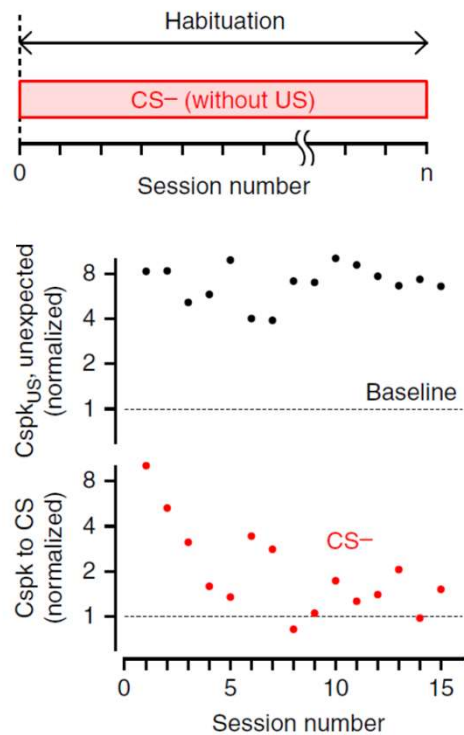
Complex spikes indicate errors or unexpected events

- Baseline rate of complex spikes $\sim 1 / s$
- Rate of complex spikes increases with errors in a novel task
- Complex spikes to unexpected events
- Rate of complex spikes decreases after learning corrects errors in performance

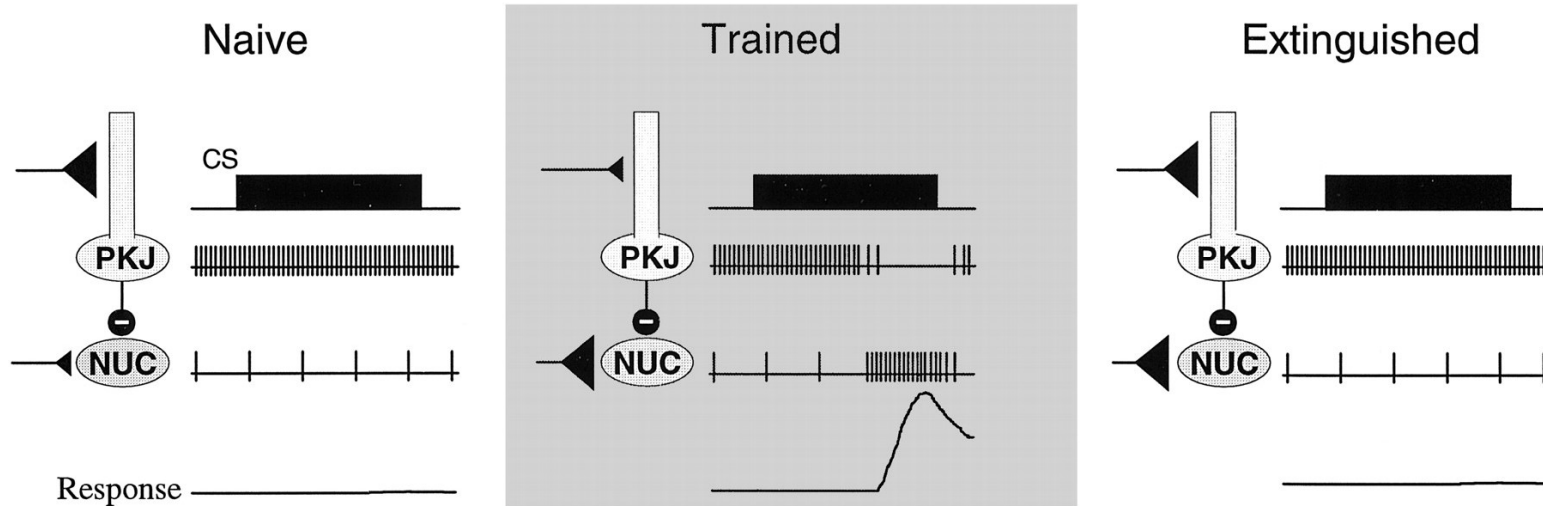


Ohmae & Medina, *Nat. Neurosci.*, 2015

Complex spikes to unexpected events habituate *unless they are predictive*



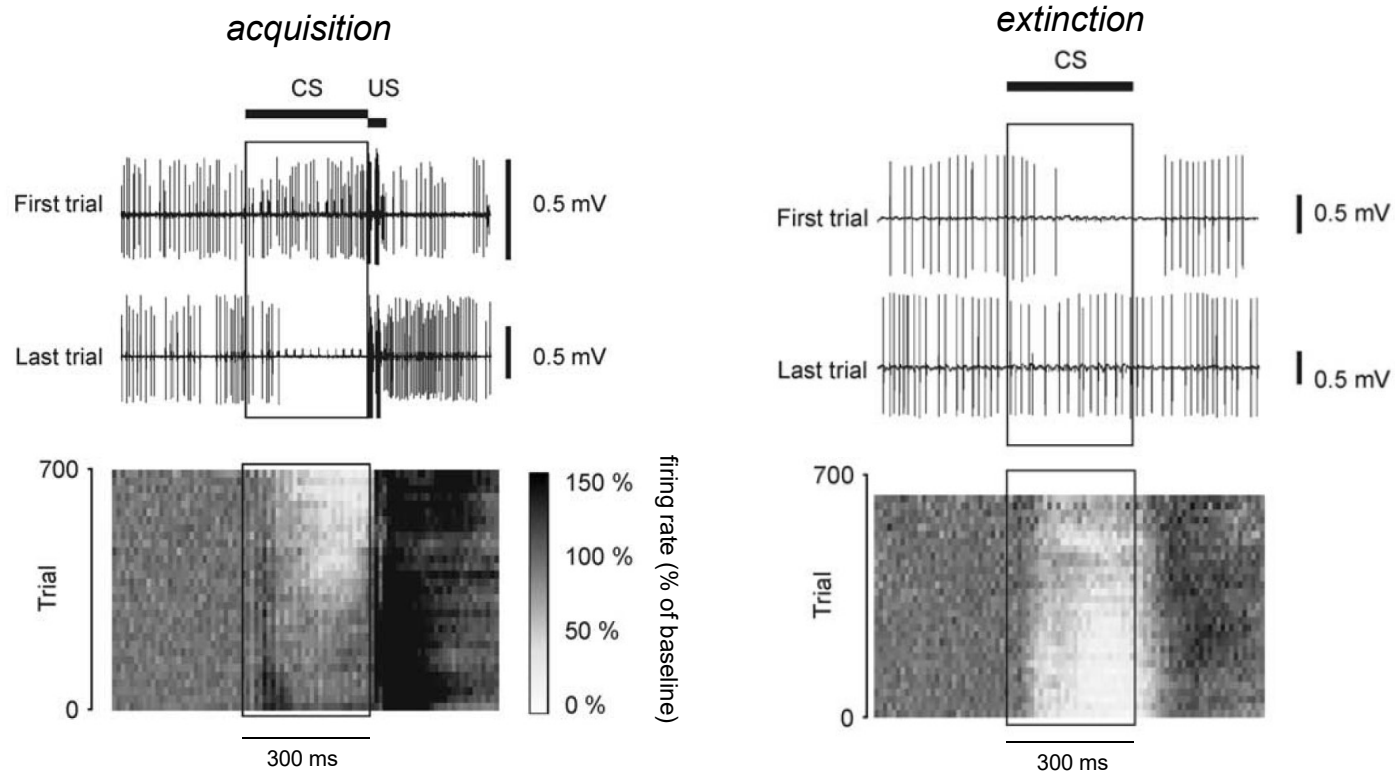
What does the CF 'teach' the Purkinje neuron?



Garcia, Steele, and Mauk, *J. Neurosci.* 19:10940, 1999

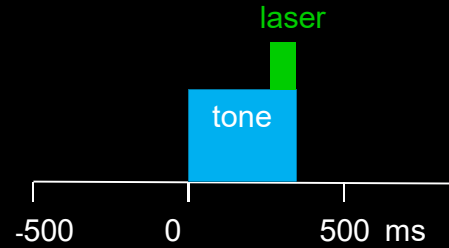
Acquisition, Extinction, and Reacquisition of a Cerebellar Cortical Memory Trace

Dan-Anders Jirenhed, Fredrik Bengtsson, and Germund Hesslow
Department of Experimental Medical Science, Lund University, 22184 Lund, Sweden

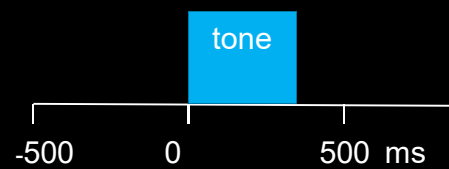


Pairing PC excitation with a tone leads to robust learned movements

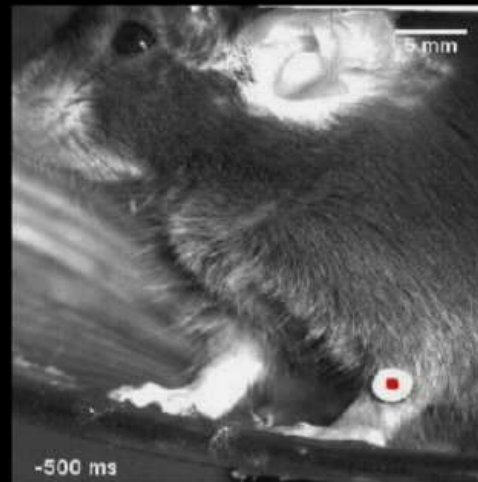
Training: 90 trials/day



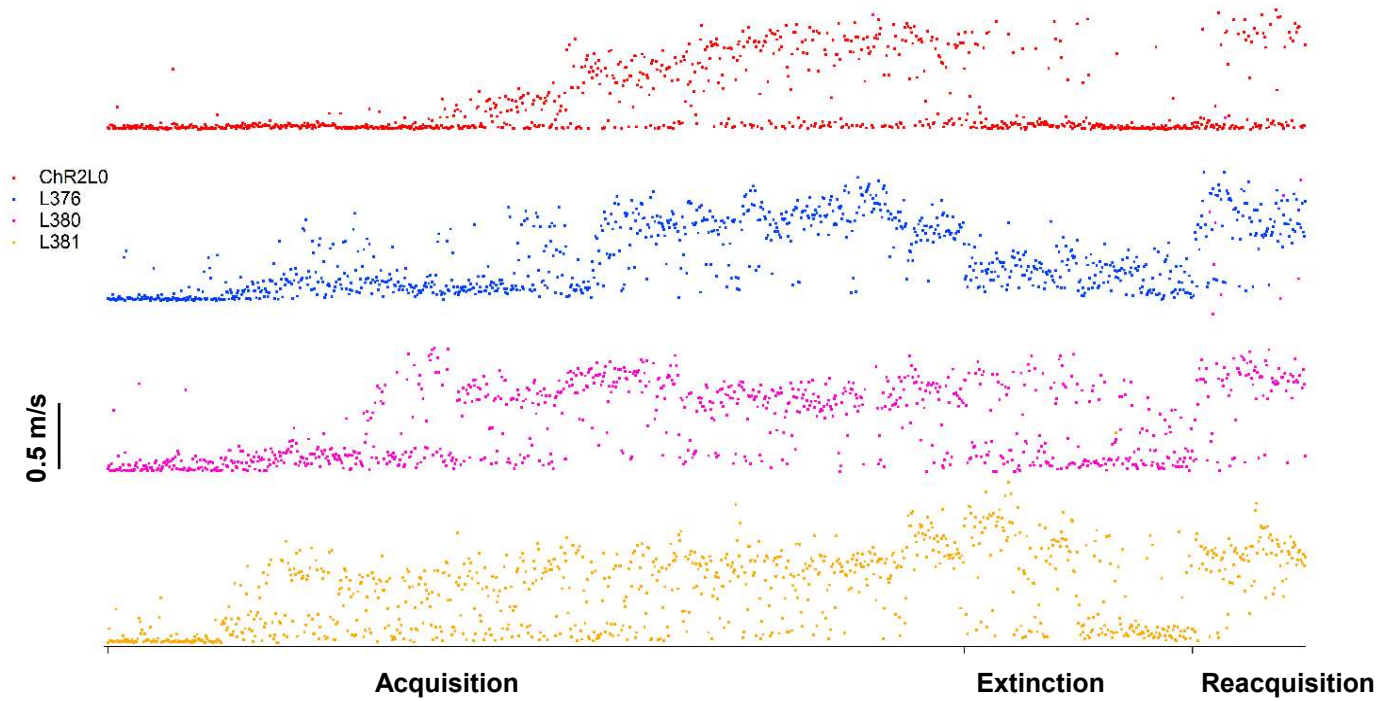
Testing:



ChR2-induced learning
before training after training

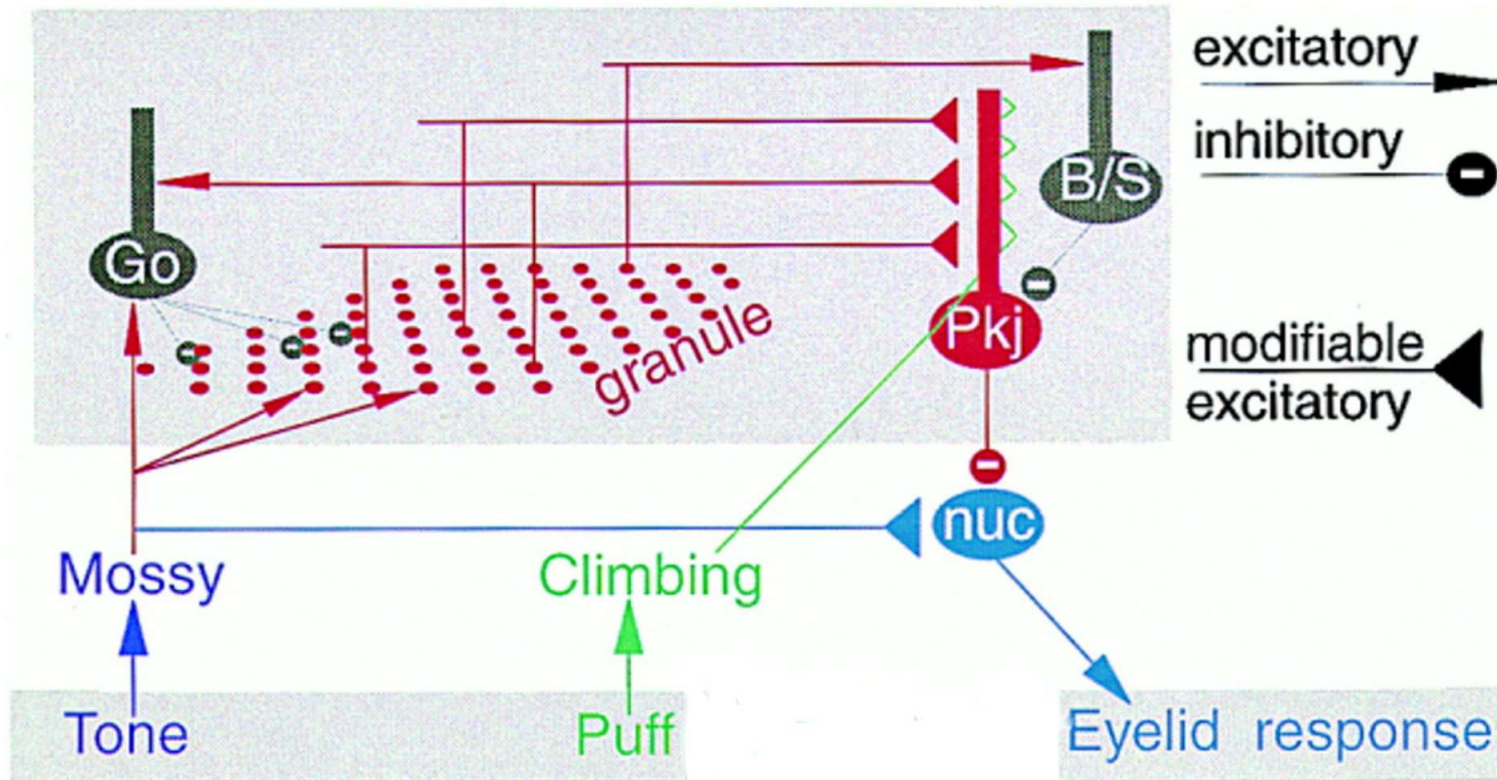


Chr2 training, individual mice



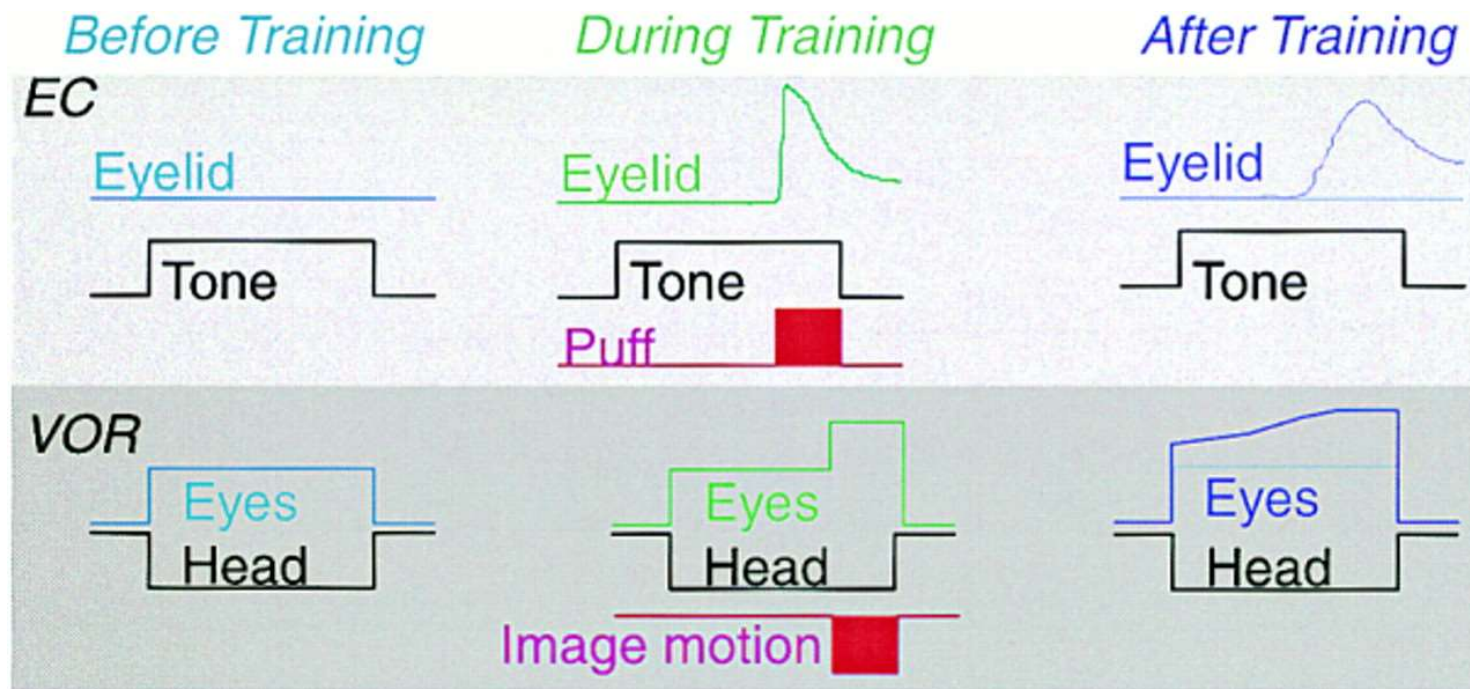
A. Reeves, unpublished

Which pathways carry the information critical for learning?



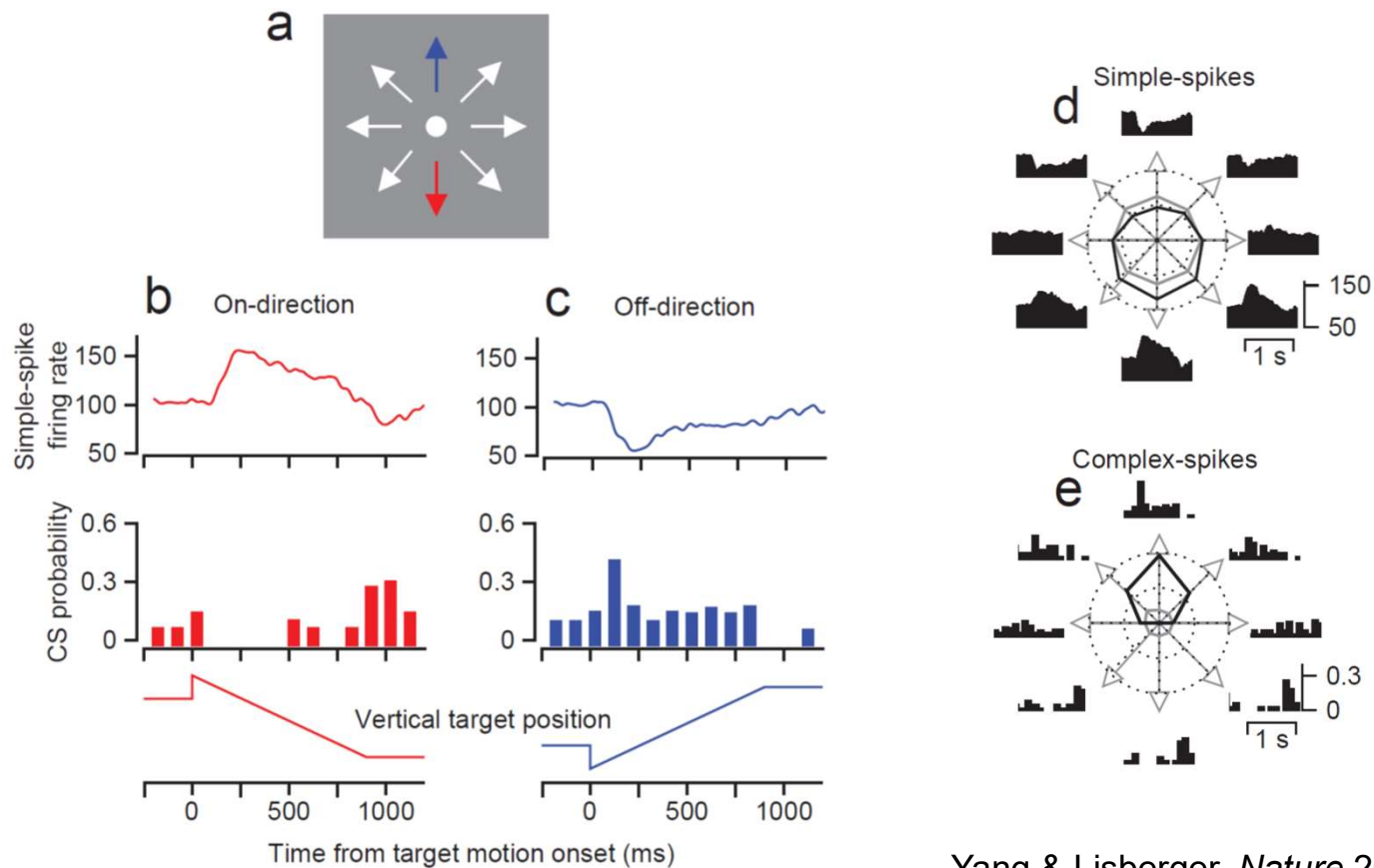
Mauk, 1997

Similarities between classical eyeblink conditioning (*EC*) and plasticity of the vestibulo-ocular reflex (*VOR*)



Mauk, 1997

PNs in flocculus are directionally tuned to smooth pursuit eye movements



Yang & Lisberger, *Nature* 2014

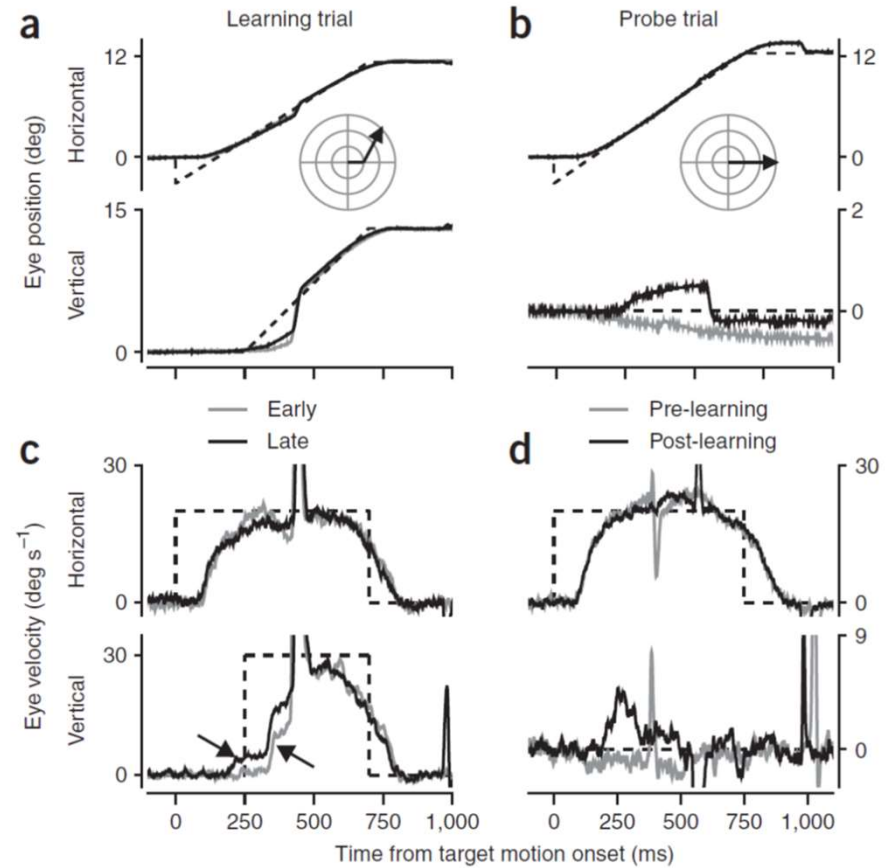
Smooth pursuit learning task



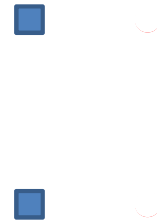
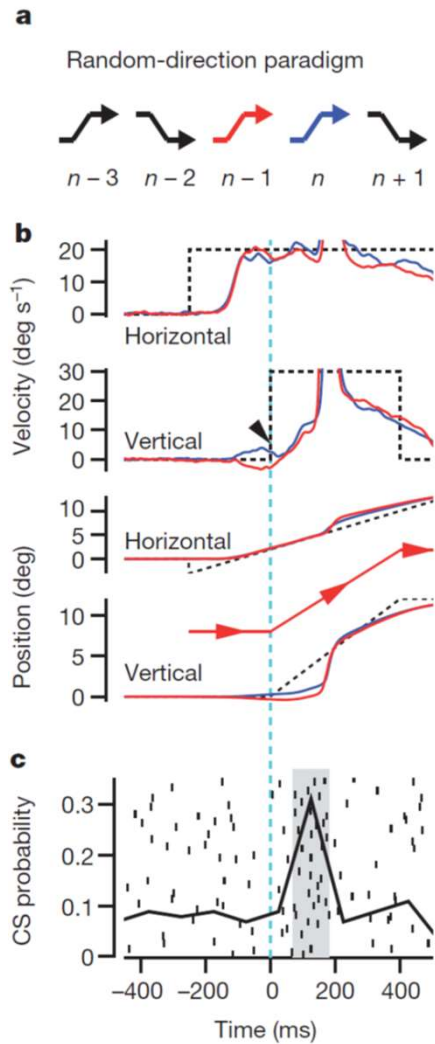
Medina & Lisberger, *Nat. Neurosci.* 2008

Smooth pursuit learning task

- task shows single trial learning
- complex spikes predict learning on a trial by trial basis

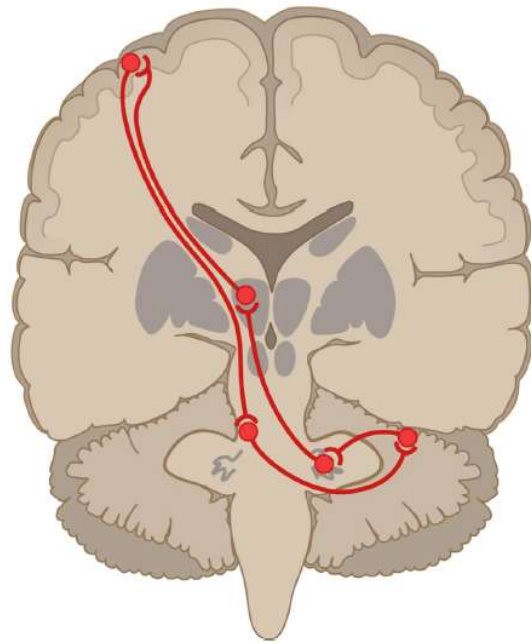


Complex spike signals predict single trial learning

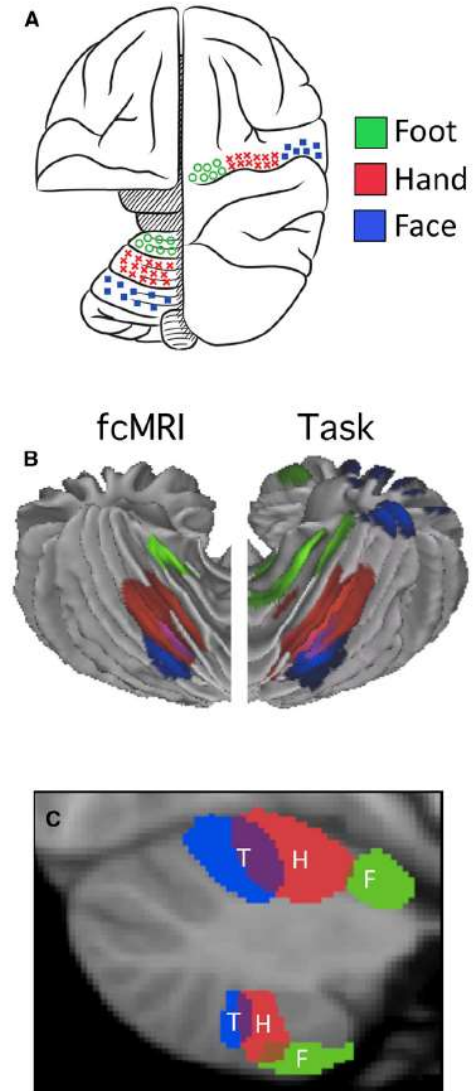


Yang & Lisberger, *Nature* 2014

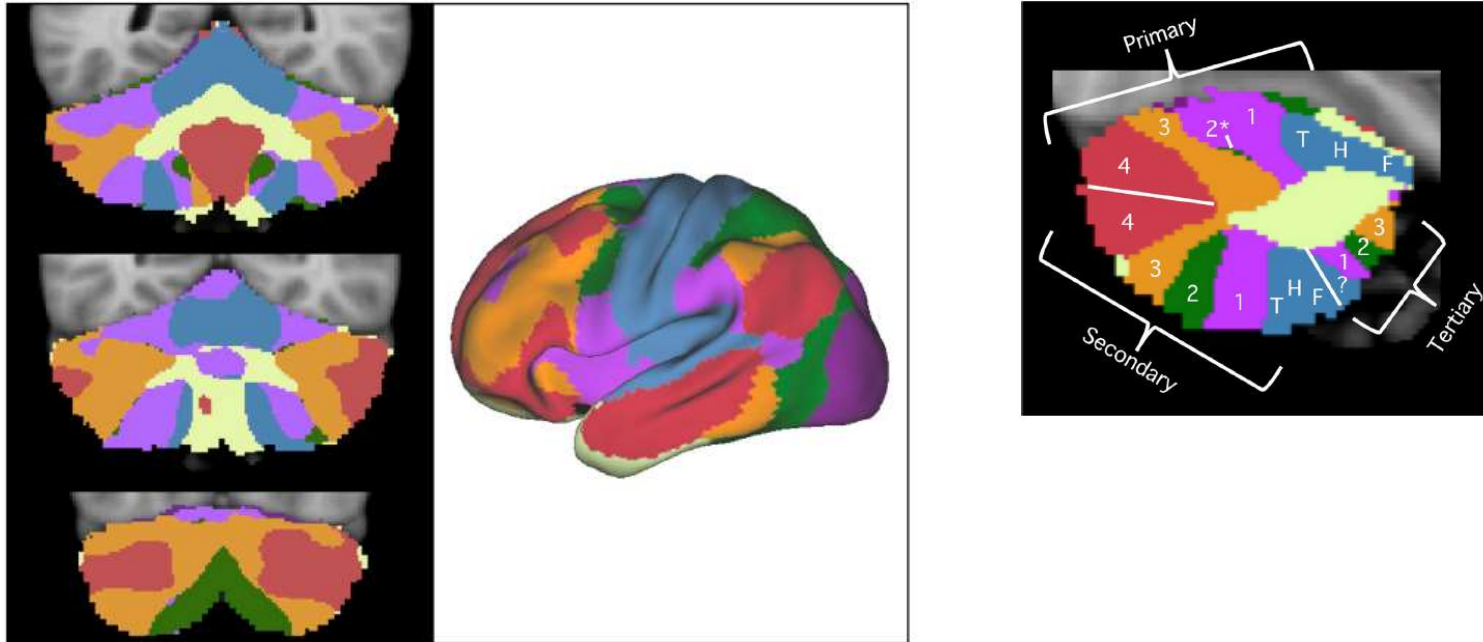
Reciprocal disynaptic connections between motor areas of cerebellum and neocortex



Buckner, *Neuron* 80:807-815, 2013



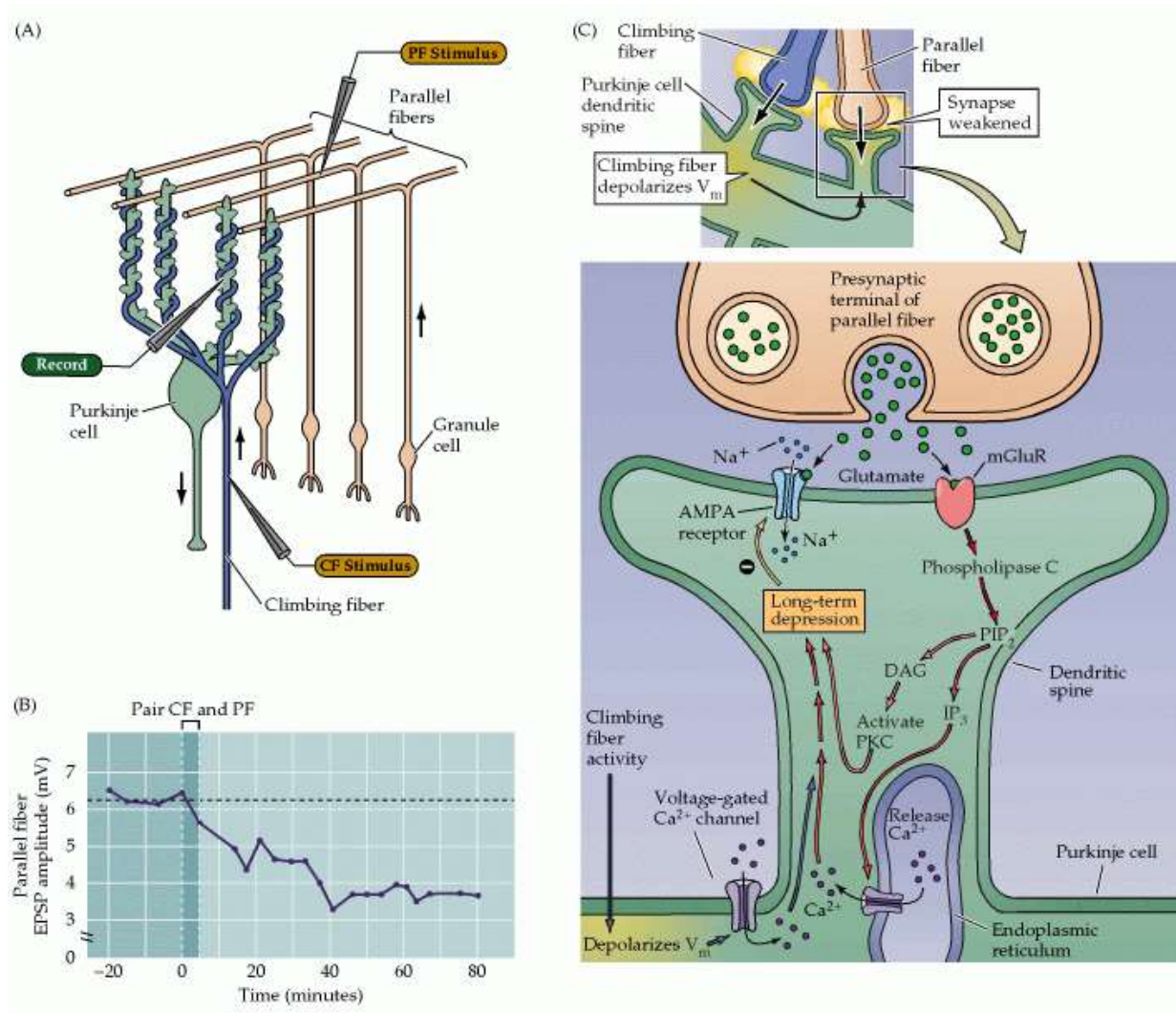
Reciprocal connections between cerebellum and all of neocortex



Buckner, *Neuron* 80:807-815, 2013; see also work by Strick and colleagues, and Schmahmann on cerebellar cognitive syndrome & “*dysmetria of thought*”

Cellular mechanisms of cerebellar LTD

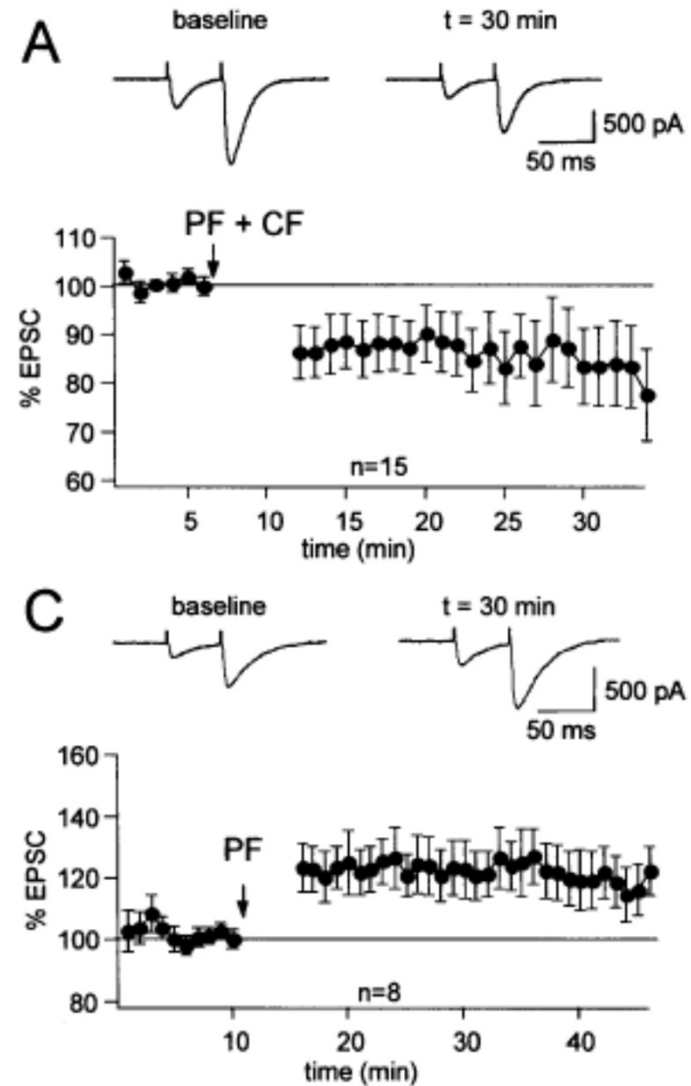
Long term depression (LTD) of PF synapses



AMPA receptors are removed at PF synapses

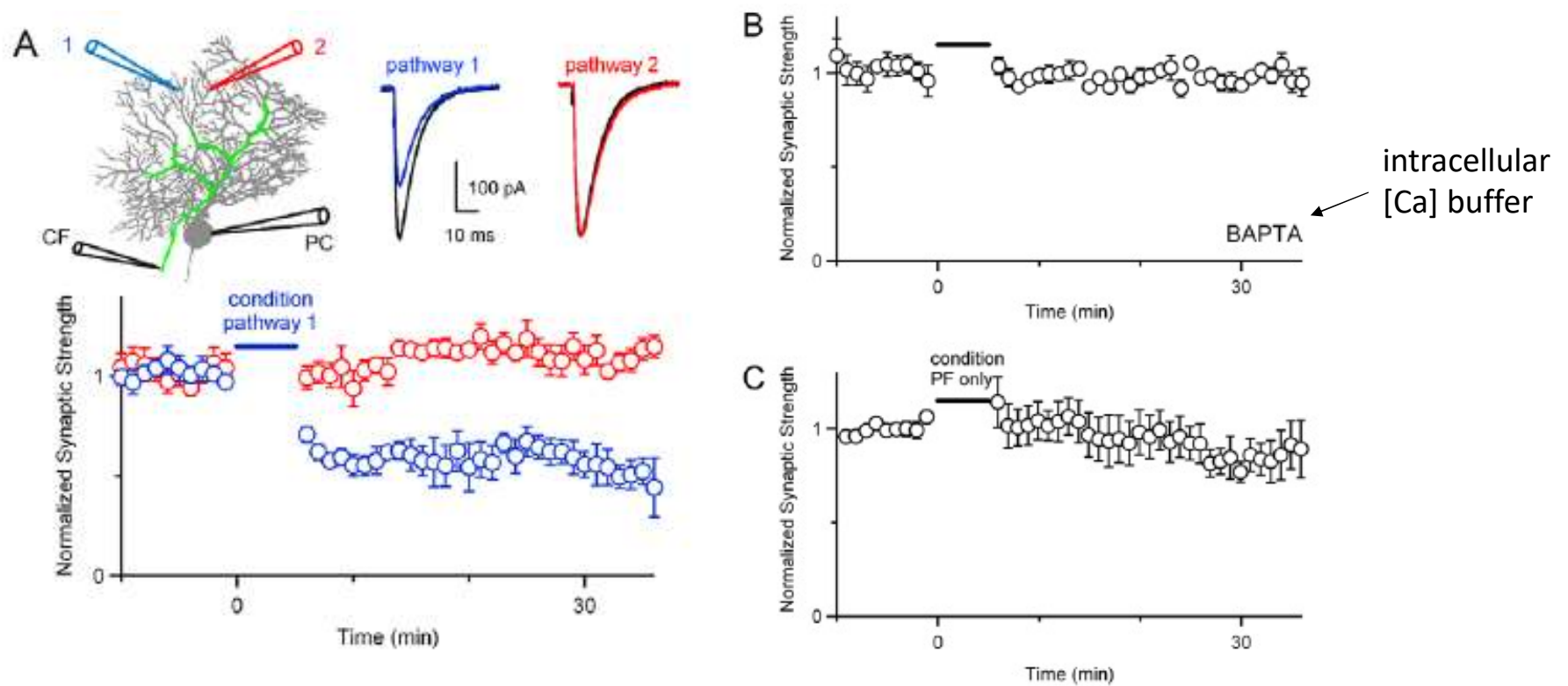
Fig.24-13, Purves

The direction of plasticity is determined by whether CF is stimulated



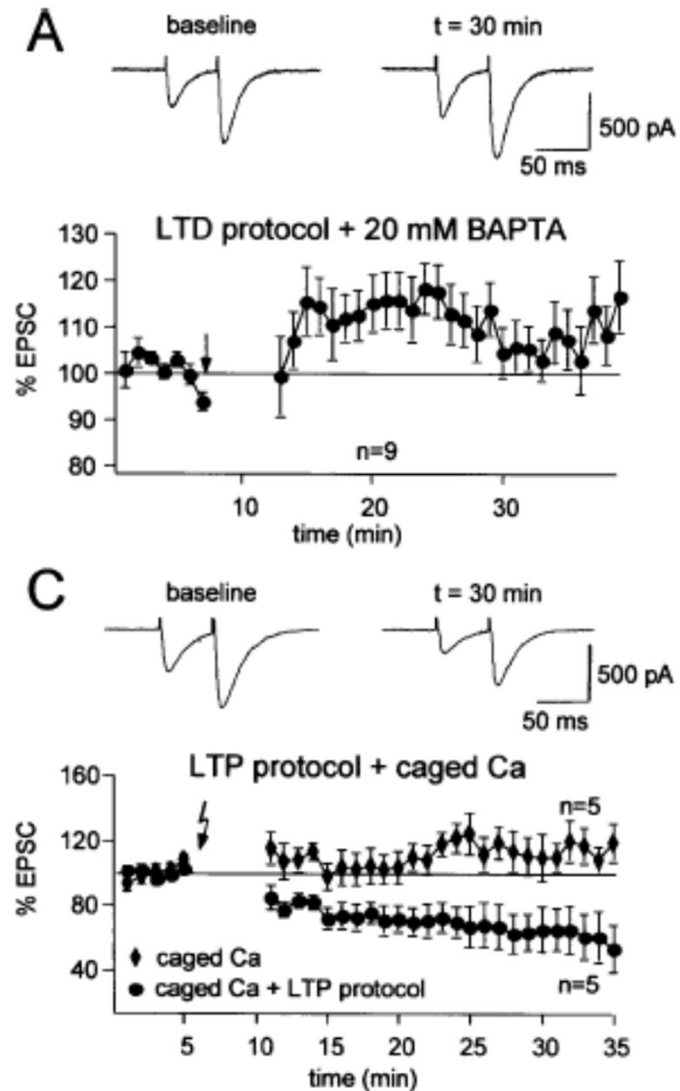
Coesmans et al., *Neuron* 44:691, 2004

LTD is synapse specific & requires an rise in $[Ca^{2+}]_i$



Safo and Regehr, *Neuron* 48:647, 2005

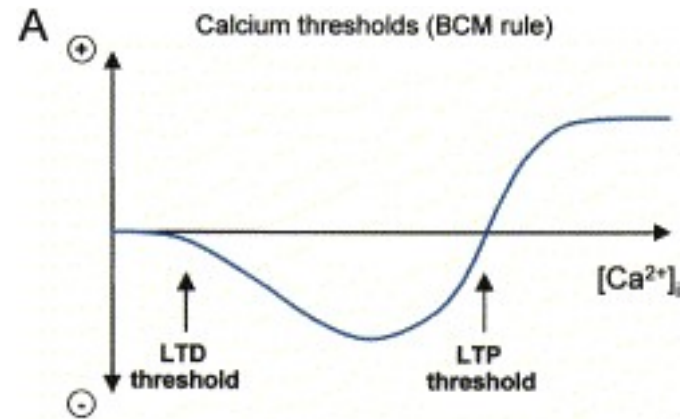
The direction of plasticity is determined by the amount of calcium



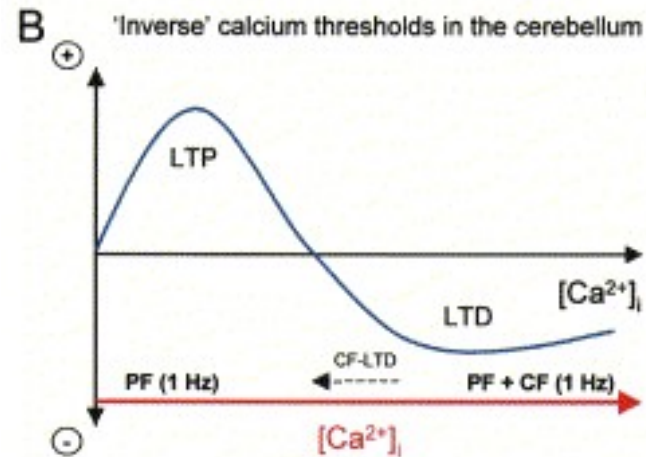
Coesmans et al., *Neuron* 44:691, 2004

An inverse $[Ca^{2+}]_i$ dependence in cerebellum?

Schaffer-collateral synapse

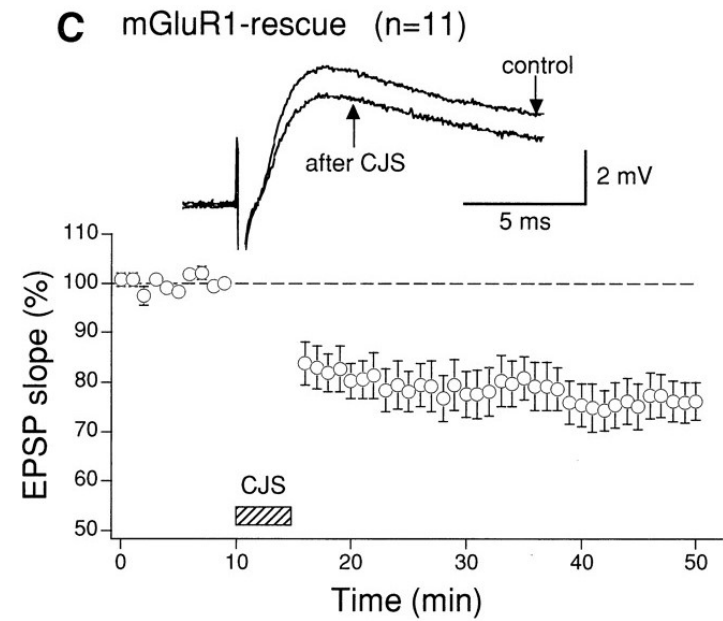
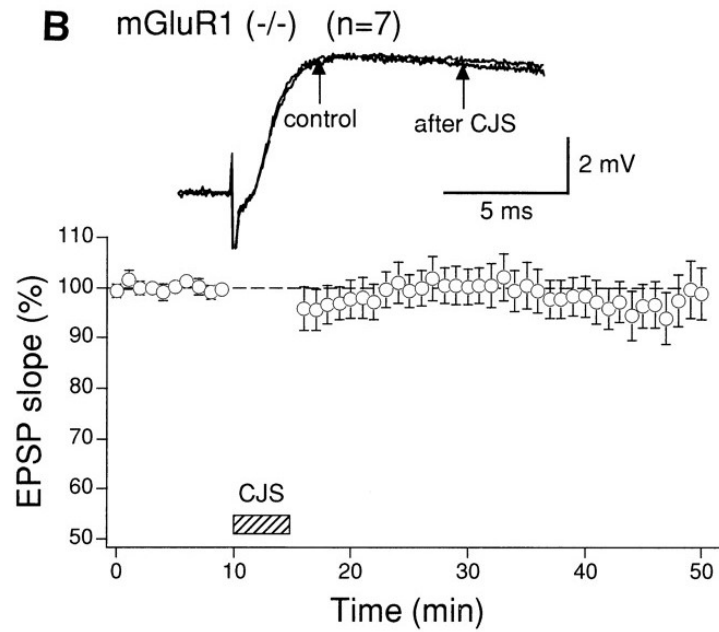


parallel fiber synapse



Coesmans et al., *Neuron* 44:691, 2004

mGluR1 function is required for LTD



Ichise et al., *Science* 288:1832, 2000

Coincidence detection mechanisms

- 1) $PF \rightarrow mGluR1\alpha \rightarrow PLC\beta \rightarrow DAG$
 $CF \rightarrow VGCC \rightarrow [Ca^{2+}] \uparrow$
 PKC α

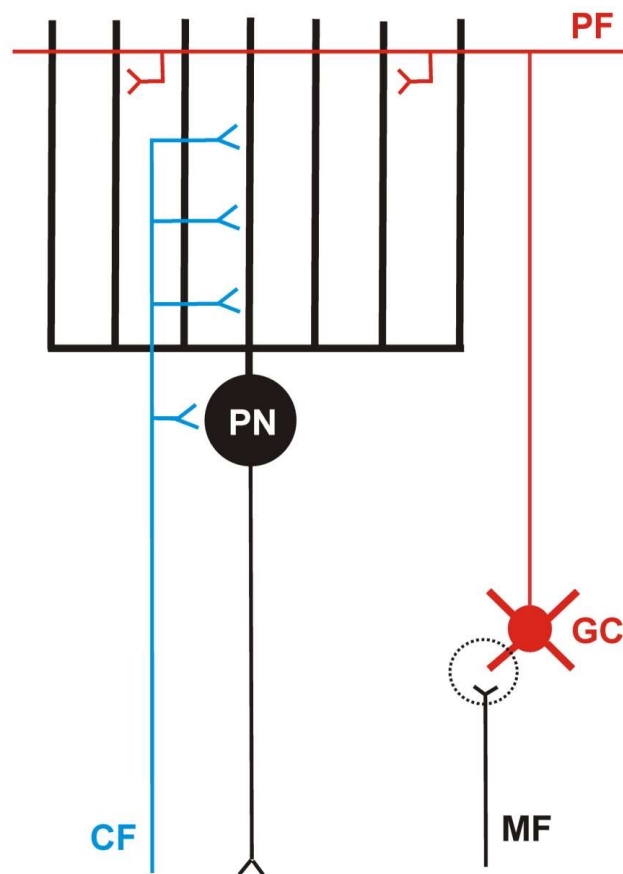
Linden & colleagues

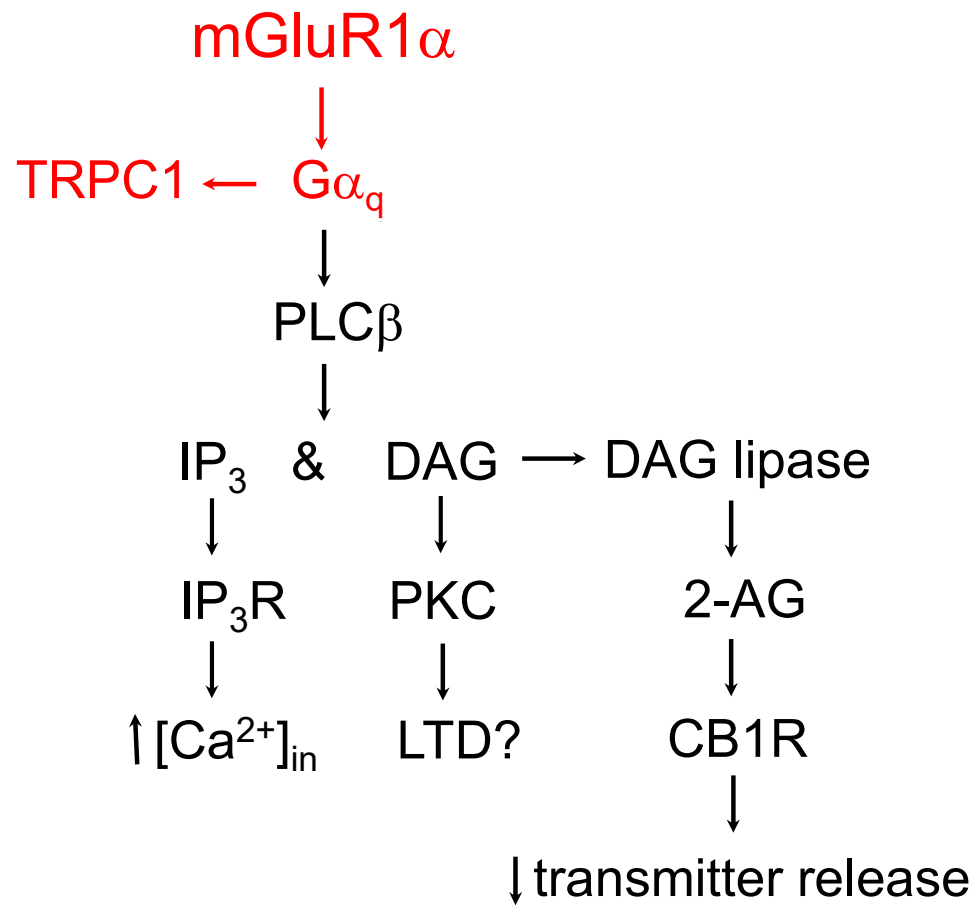
- 2) $PF \rightarrow mGluR1\alpha \rightarrow PLC\beta \rightarrow IP_3$
 $CF \rightarrow VGCC \rightarrow [Ca^{2+}] \uparrow$
 IP $_3$ R

Augustine, Finch, Wang

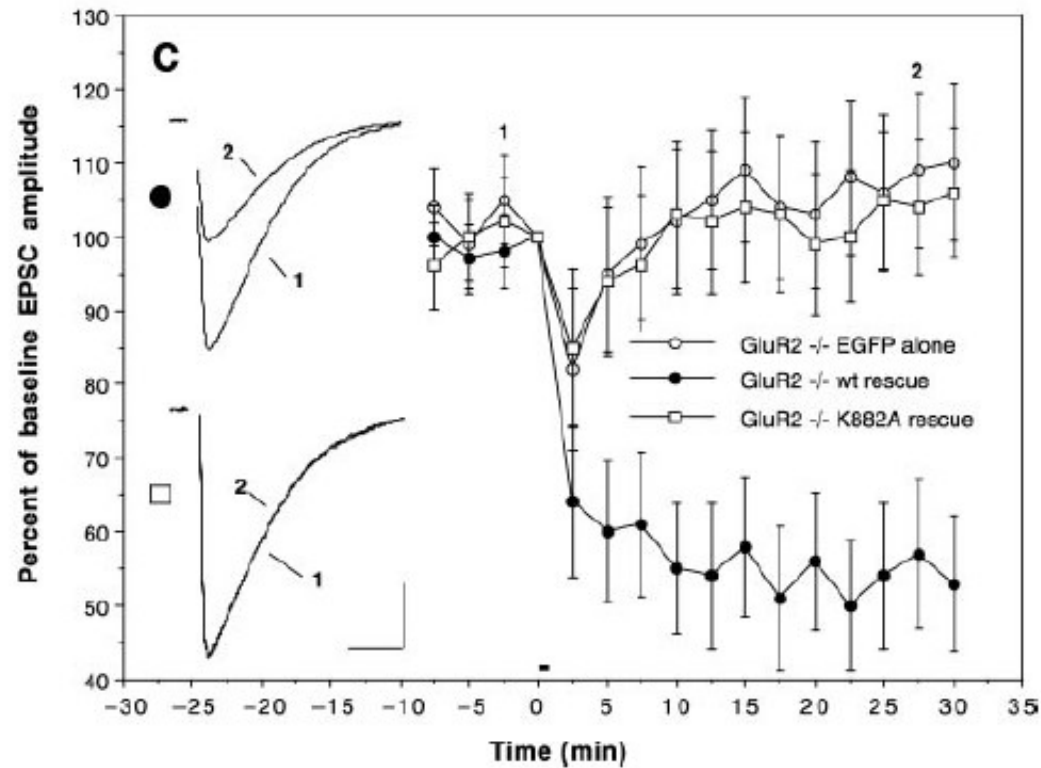
- 3) $PF \rightarrow NO \rightarrow sGC \rightarrow cGMP$
 $CF \rightarrow VGCC \rightarrow [Ca^{2+}] \uparrow$
 PKG?

Lev Ram, Hartell, Crepel



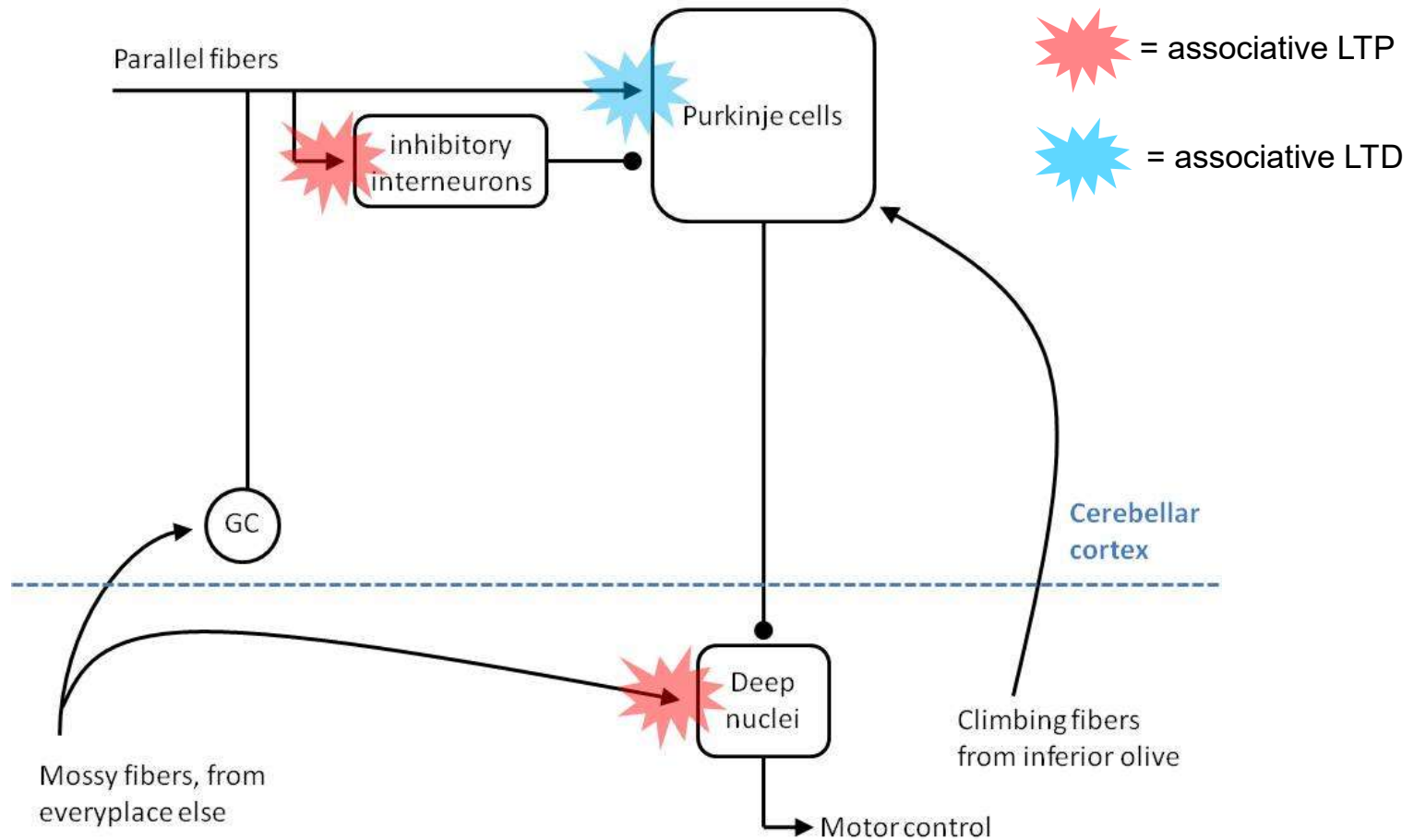


Endocytosis of GluR2-containing AMPARs is the basis for LTD



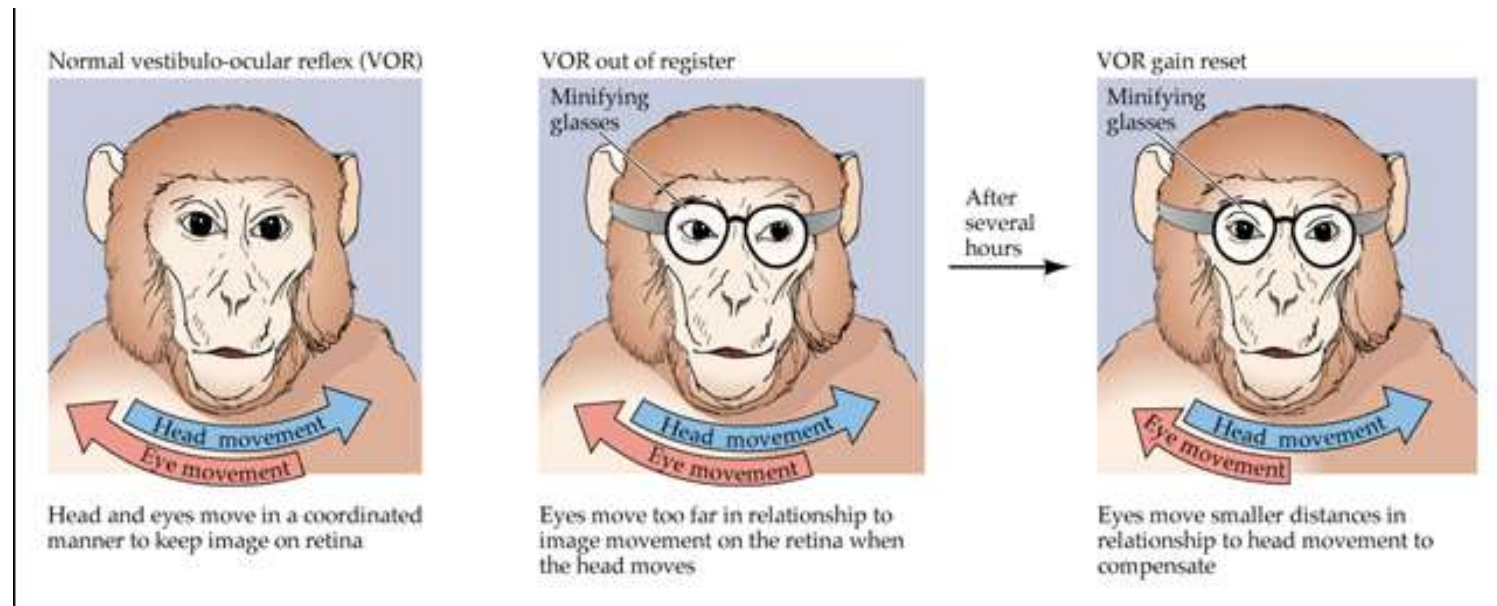
Chung et al., *Science* 300:1751, 2003

Summary: sites of plasticity



Backup, extra slides

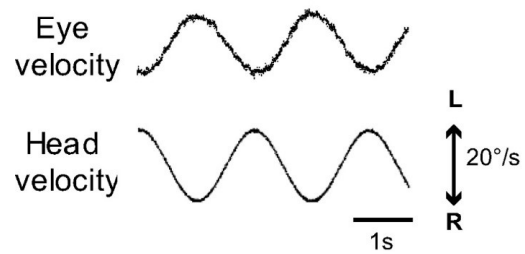
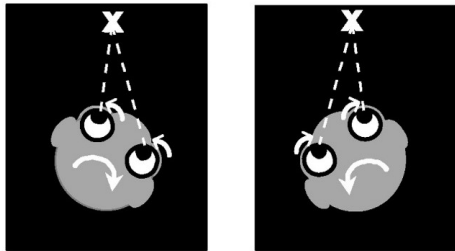
VOR plasticity can be induced by minimizing or magnifying spectacles.



From Purves et al., 1997

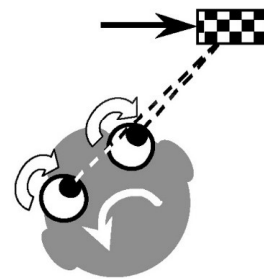
VOR learning

Pretraining
(dark)

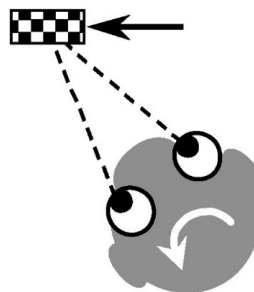


Training

gain-up stimulus

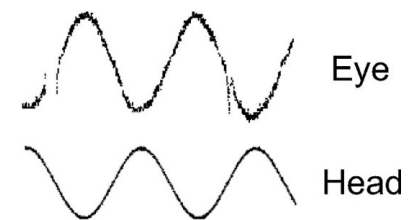


gain-down stimulus

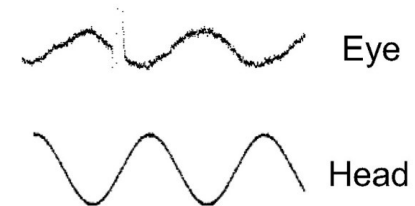


Posttraining
(dark)

increase in VOR gain



decrease in VOR gain



Boyden et al., 2004