Synaptic Plasticity : Spike-timing dependent plasticity (STDP)

Oct 15th, 2018

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At which university am I working ?

At which university am I working ?

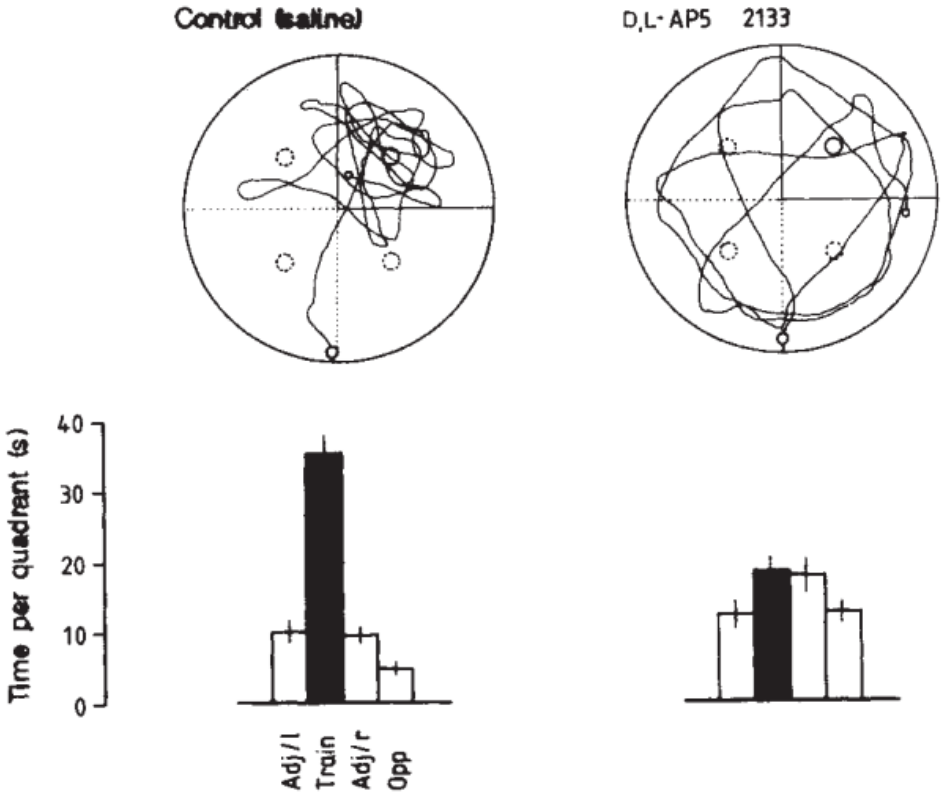
→ synaptic plasticity basis of learning

Why are we interested in synaptic plasticity ?

Morris water maze

[Morris *et al.*, 1986]

Relation between LTP and learning/memory



- NMDA receptor required to learn platform location [Morris *et al.*, 1986]
- NMDA receptor required to form spatial memories (place fields)

[McHugh *et al.* 1996]

Outline : STDP ... spike-timing dependent plasiticity

1. STDP : introduction and history
2. Phenomenology of STDP
3. Induction mechanisms
4. Biophysical models of STDP
5. STDP *in vivo*

Outline

1. STDP : introduction and history

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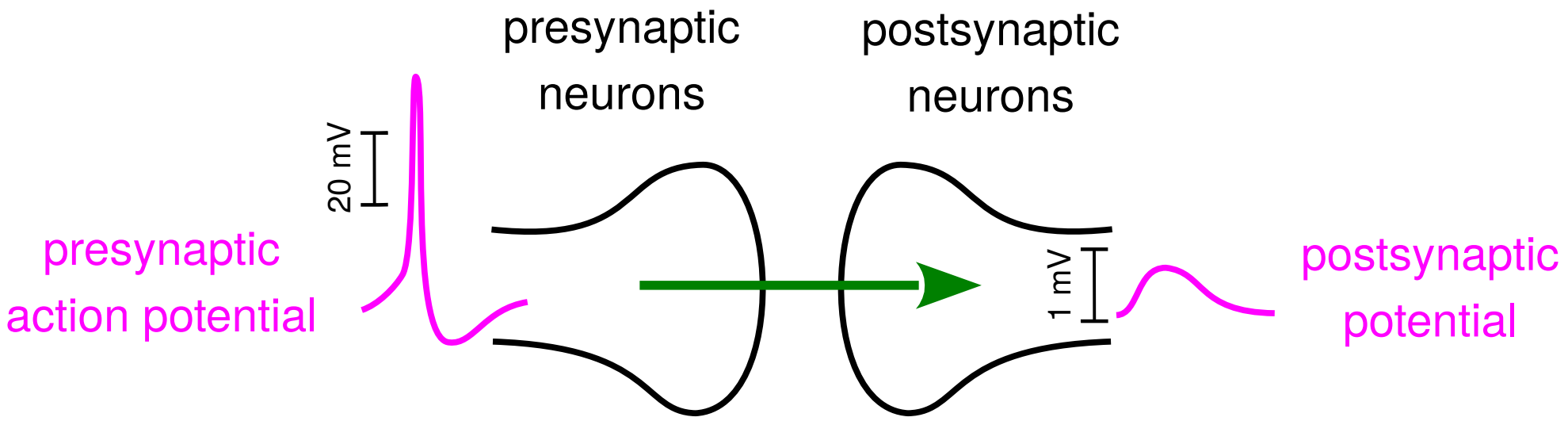
5. STDP *in vivo*

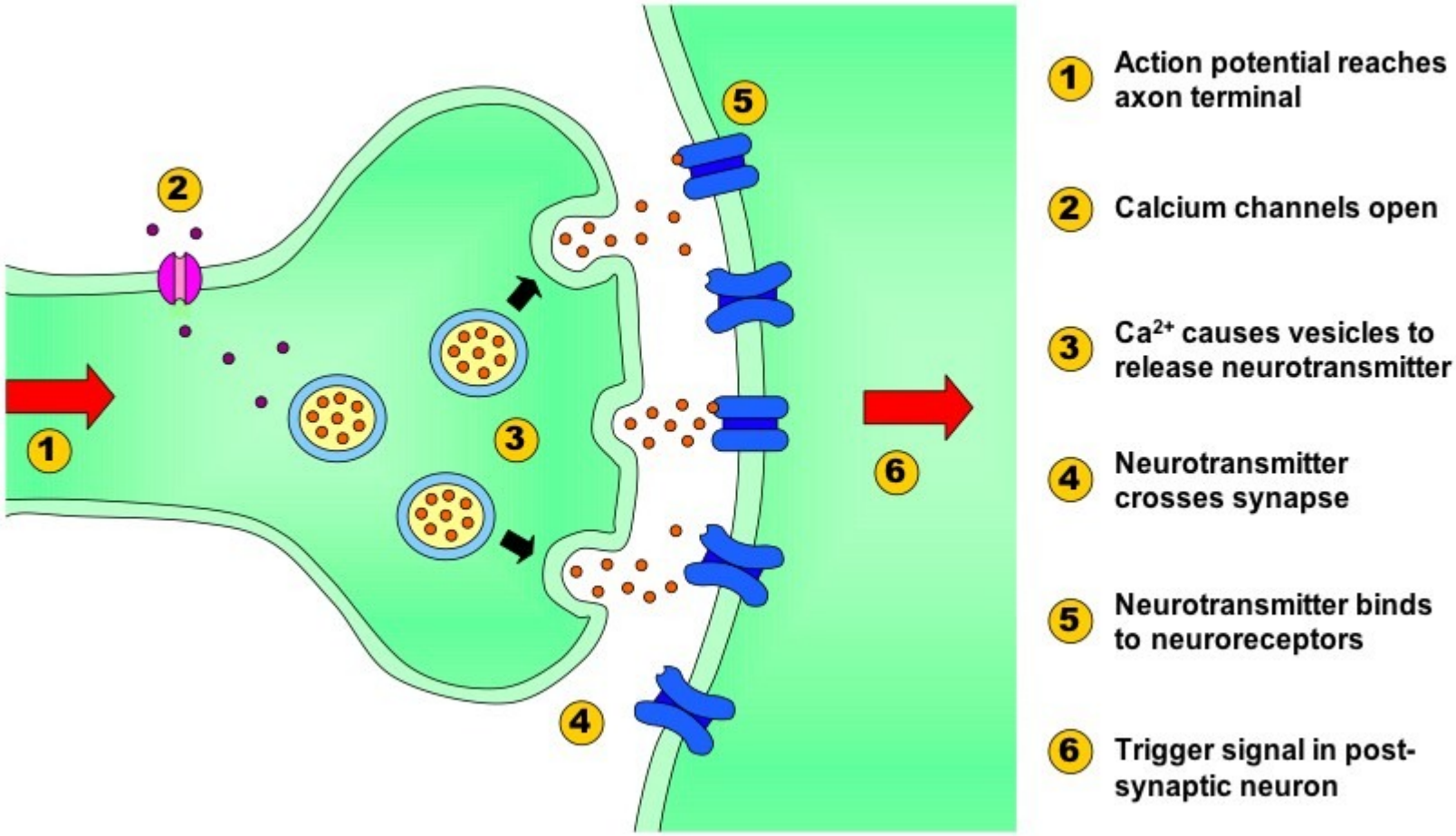
Chemical synapse : transmits electrical signals



electrical signal → chemical signal → electrical signal

- directional transmission
- conversion of signals allows for flexibility/plasticity

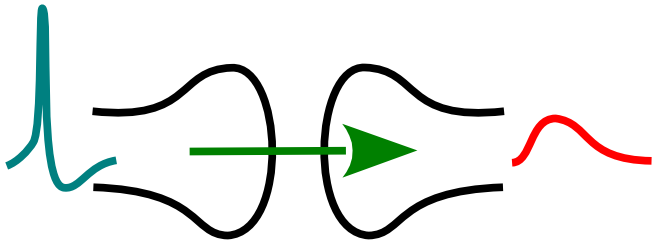
Chemical synapse : underlying biological machinery



http://outreach.mcb.harvard.edu/animations/synaptic.swf

Chemical synapse : excitatory or inhibitory

Excitatory synapse

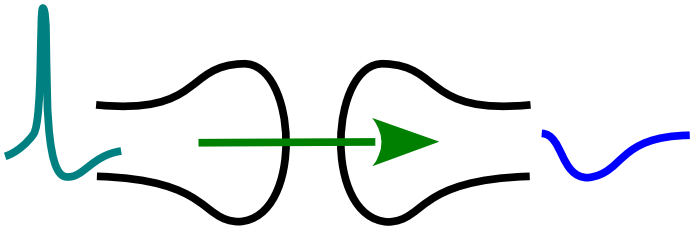


depolarization:

excitatory postsynaptic potential (EPSP)

neurotransmitter	receptor
glutamate	AMPA, NMDA
acetylcholine	nAChR, mACHR
catecholamines	G-protein-coupled receptors
serotonin	5-HT_3, ...
histamine	G-protein-coupled receptors

Inhibitory synapse



hyperpolarization:

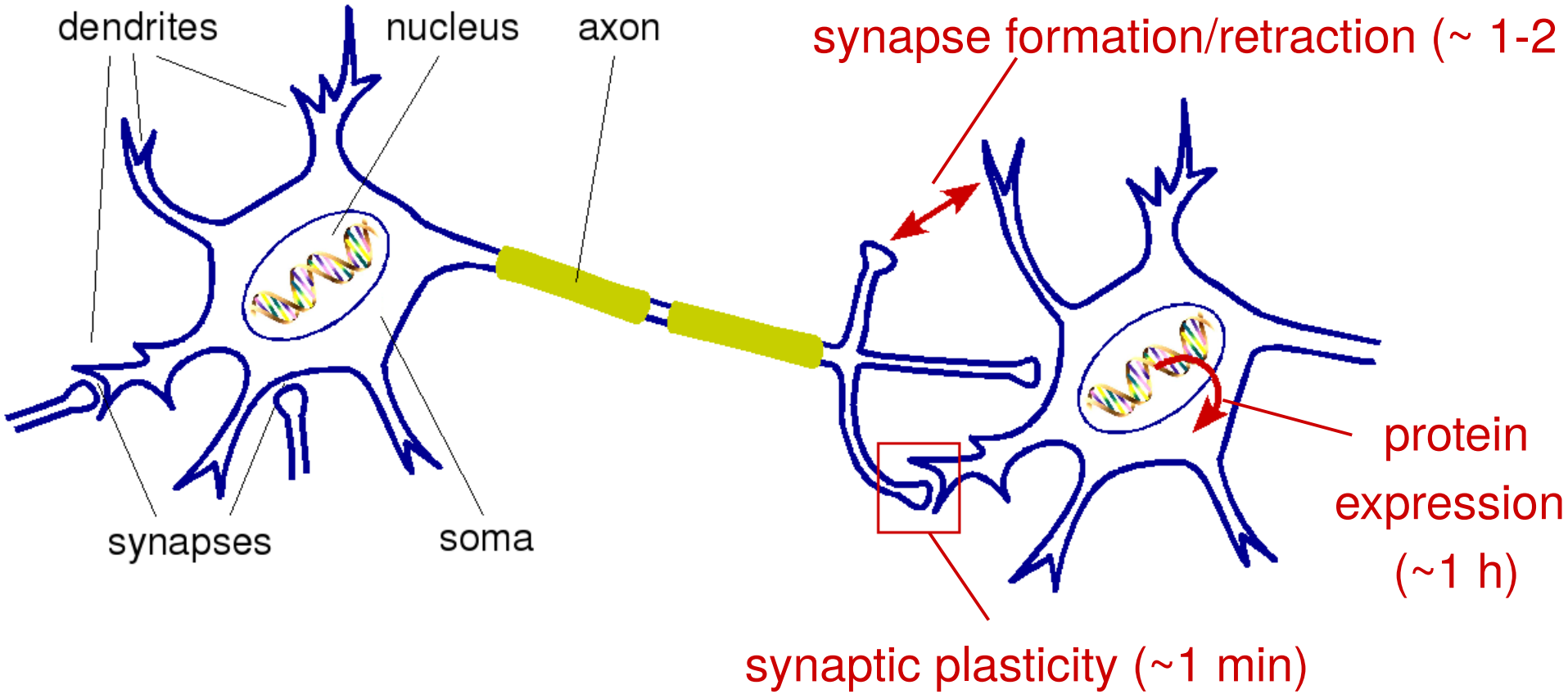
Inhibitory postsynaptic potential (IPSP)

neurotransmitter	receptor
GABA	GABA_A, GABA_B
glycine	GlyR

Different forms of plasticity

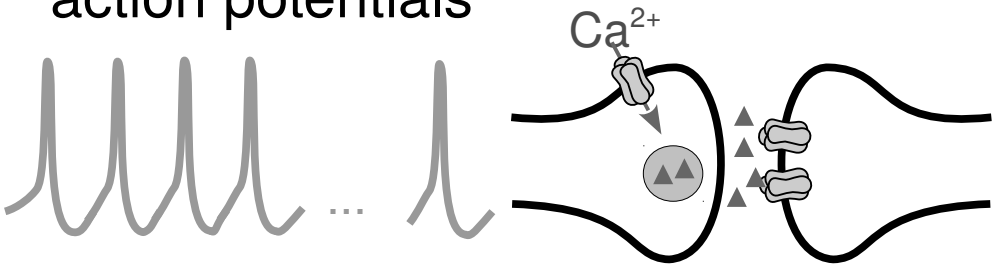
structure of neurons

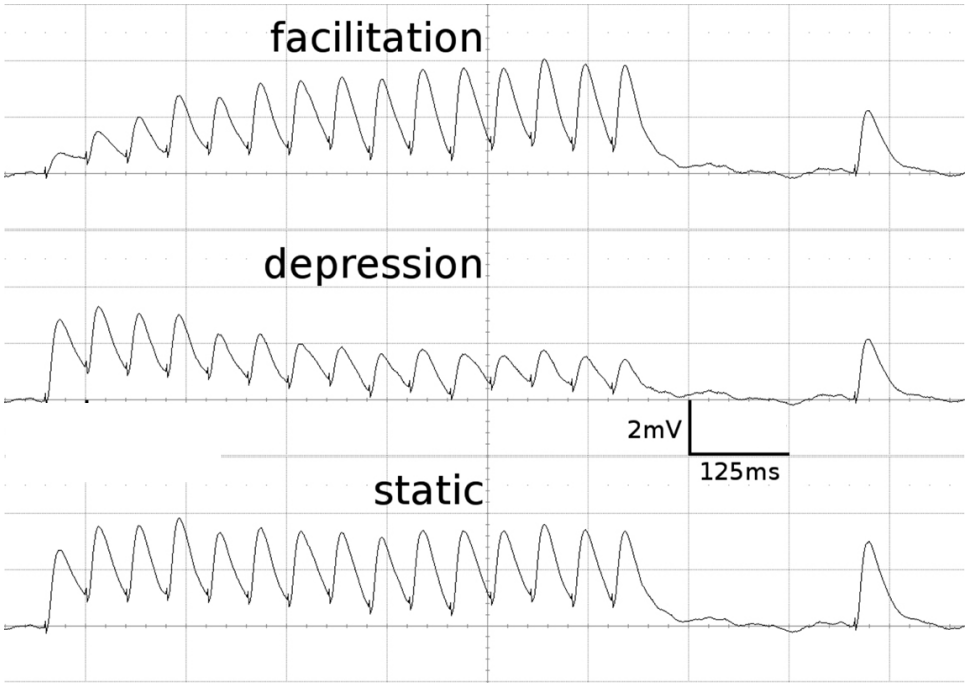
changes related to neural activity



Short-term synaptic plasticity

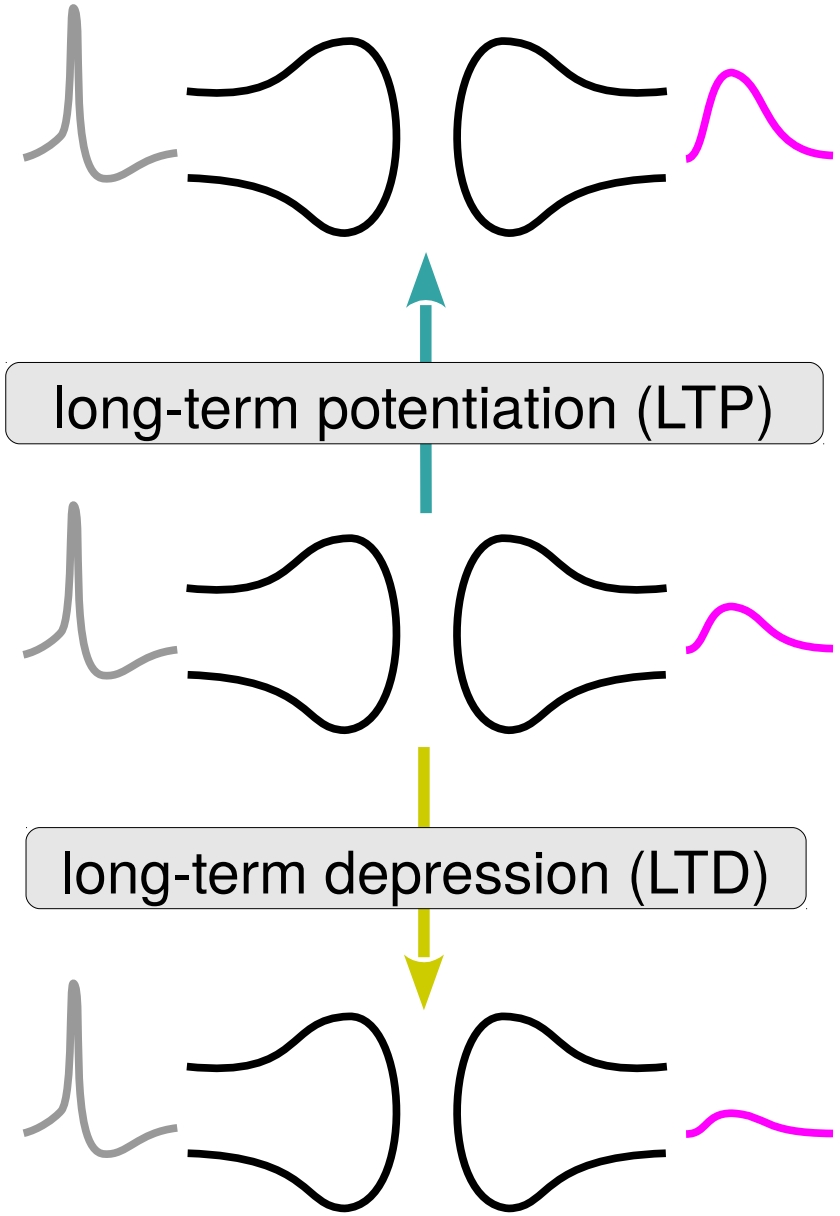
train of presynaptic action potentials





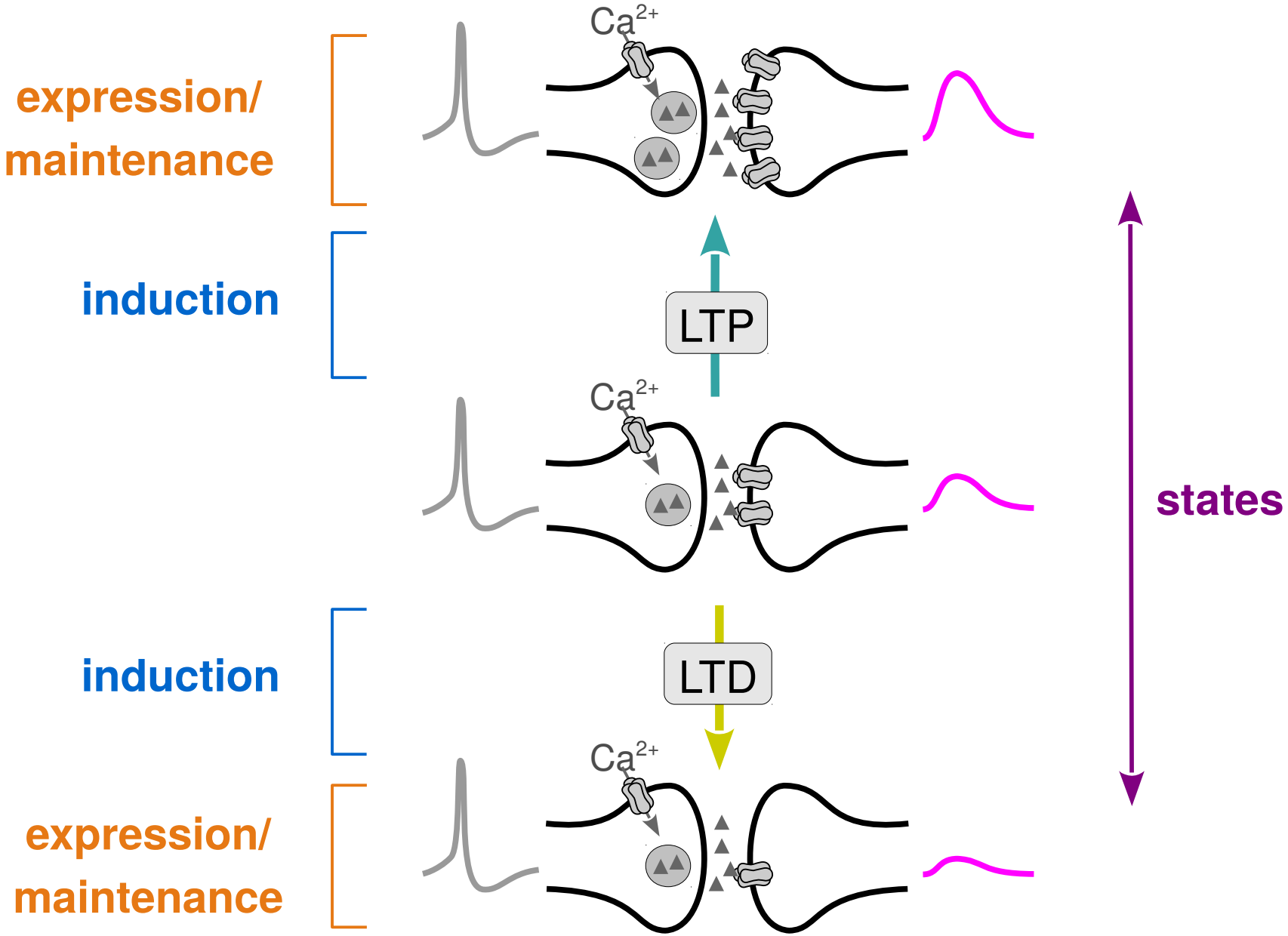
- transient change in transmission efficacy
- time scale of changes ~1 sec

Long-term synaptic plasticity

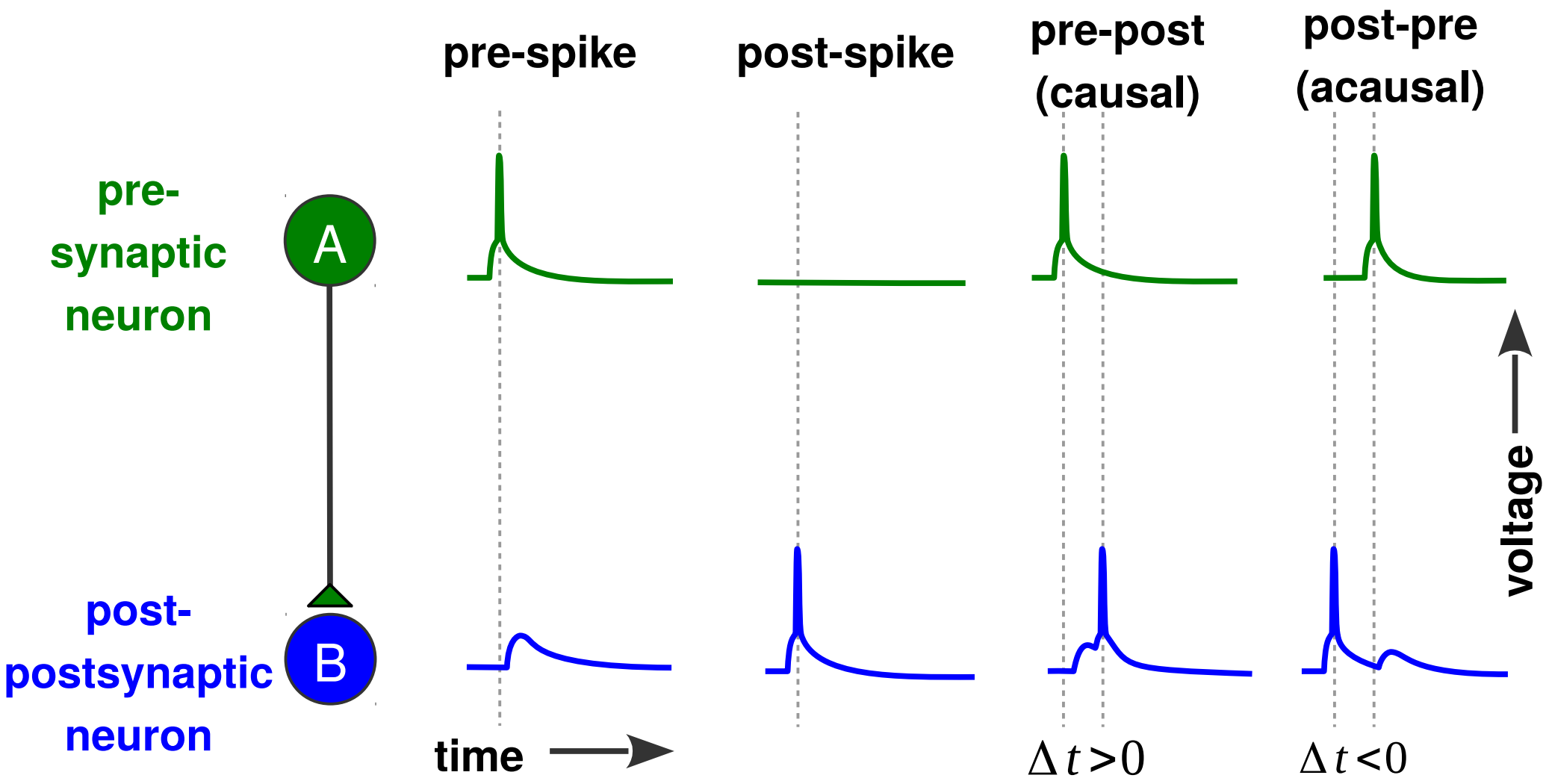


- long-lasting change (>60 min) in transmission efficacy
- time scale of induction ~ 1 min

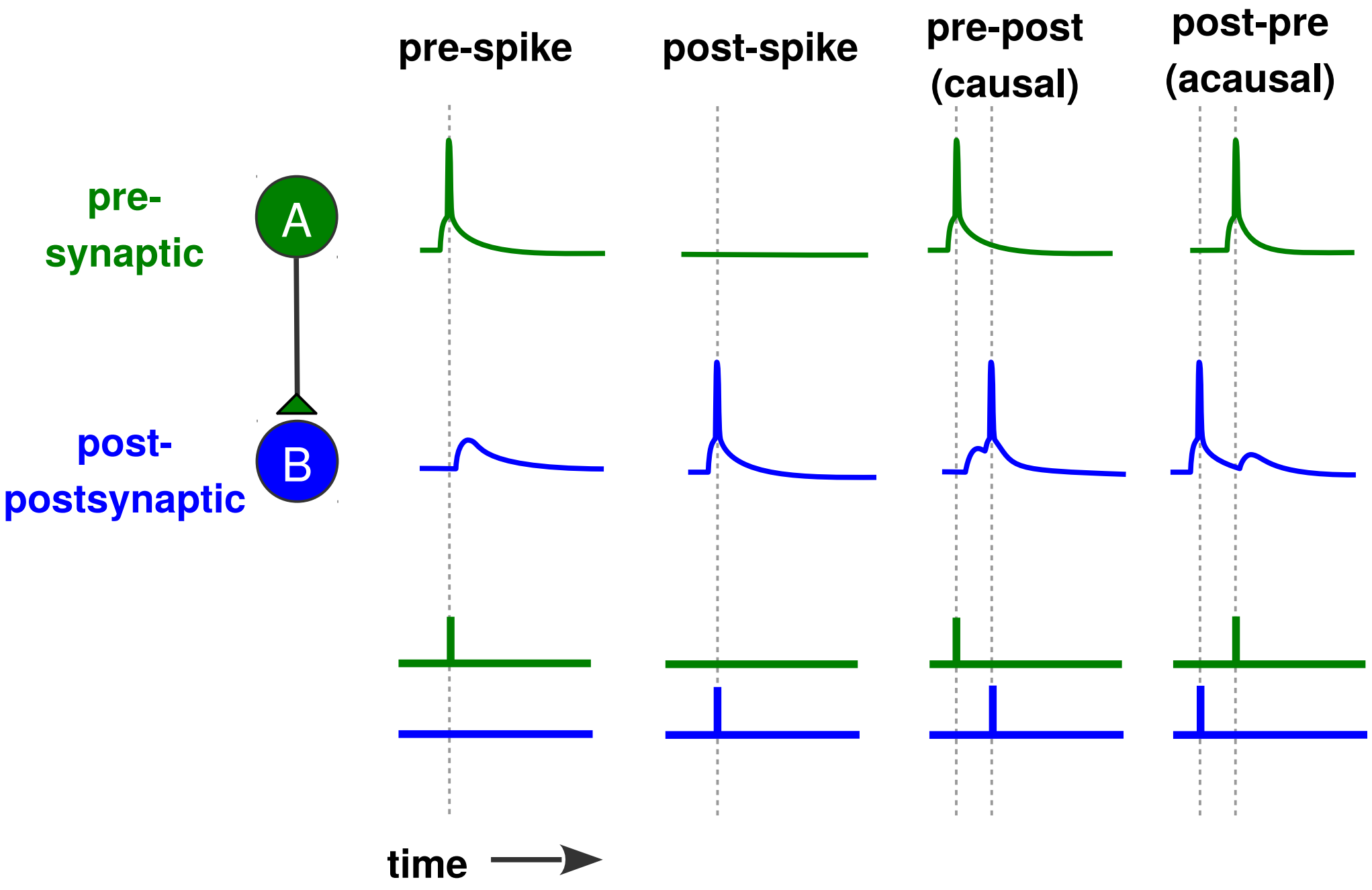
Synaptic plasticity: induction, maintenance & states



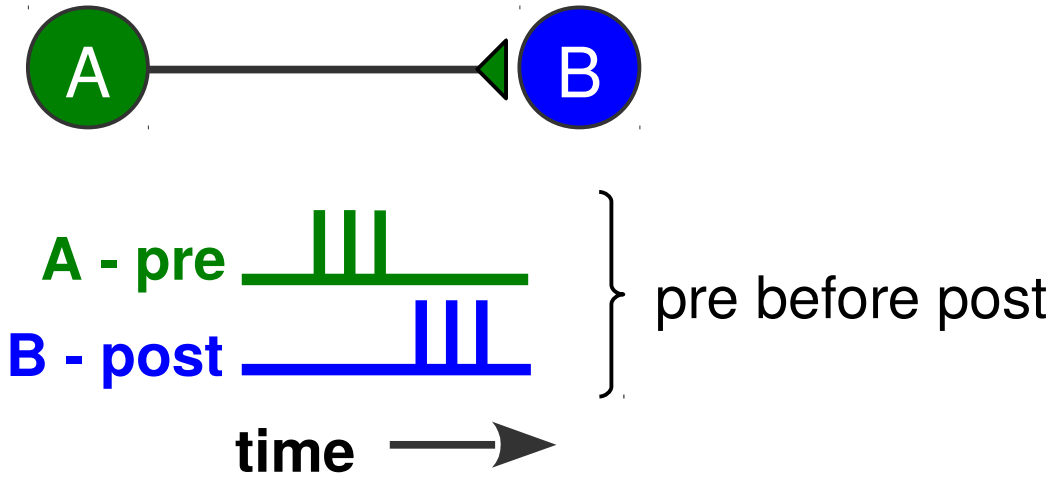
Spike timing : nomenclature



Spike timing : nomenclature



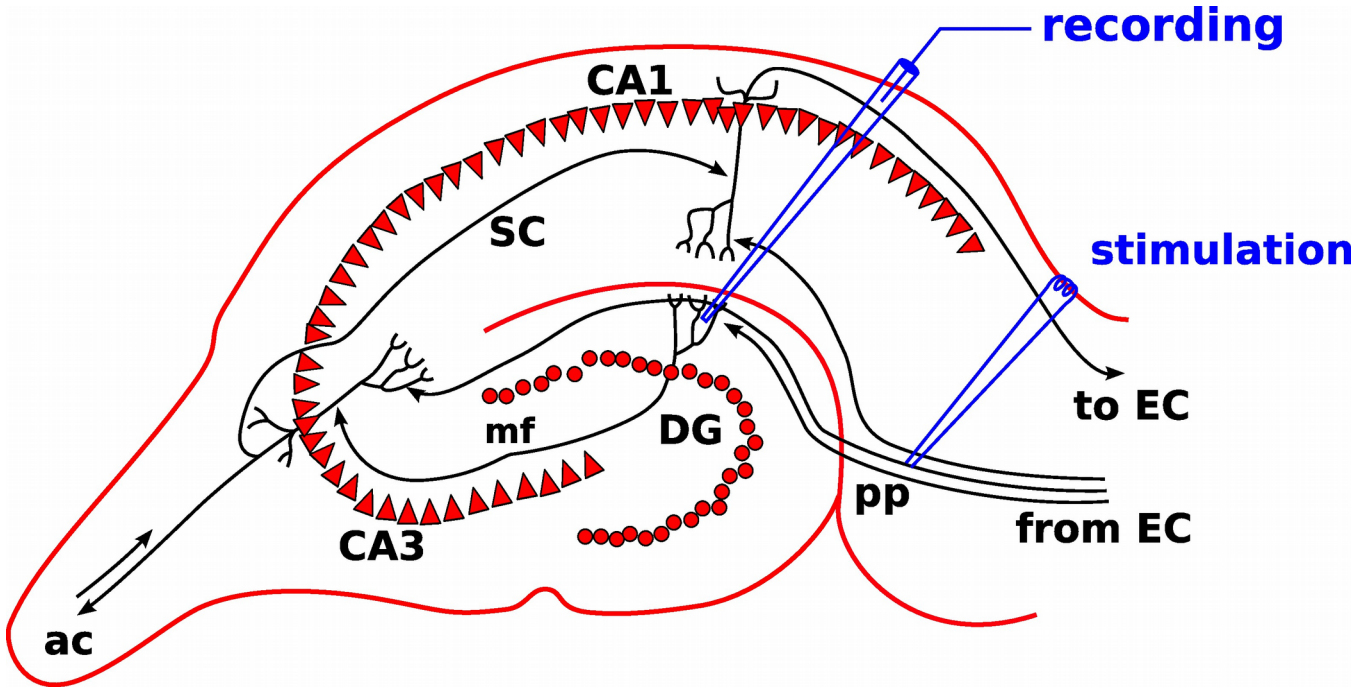
LTP induction: early conceptual work



"When an axon of cell A is near enough to excite a cell B and *repeatedly* and *persistently* takes part in firing it, some growth or metabolic changes take place in one or both cells such that A's efficiency, as one of the cells firing B, is *increased.*" [Hebb 1949; see also Konorski 1948]

Induction: first experimental work in hippocampus

hippocampus



EC	...	enthorhinal cortex
DG	...	dentate gyrus
CA3/1	...	cornu ammonis 3/1

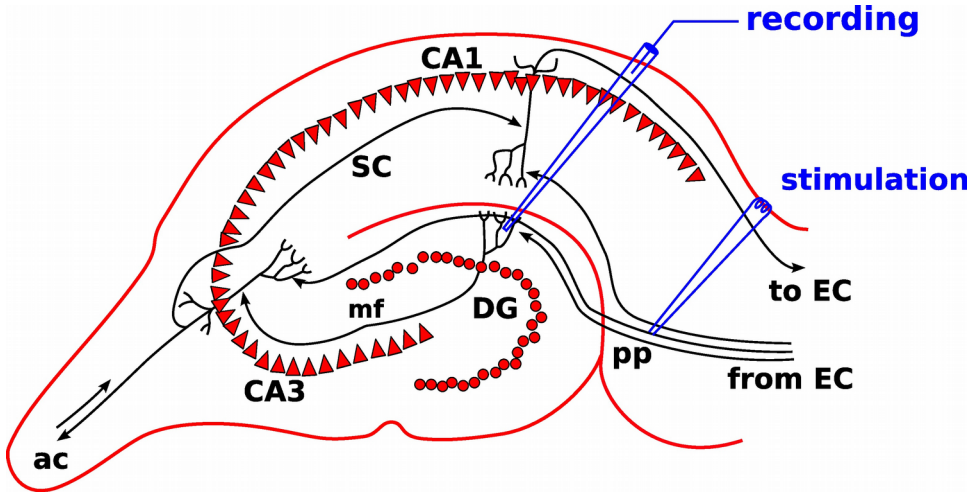
pp	...	perforant path
mf	...	mossy fibres
ac	...	associational commissural path
sc	...	Schaffer collateral

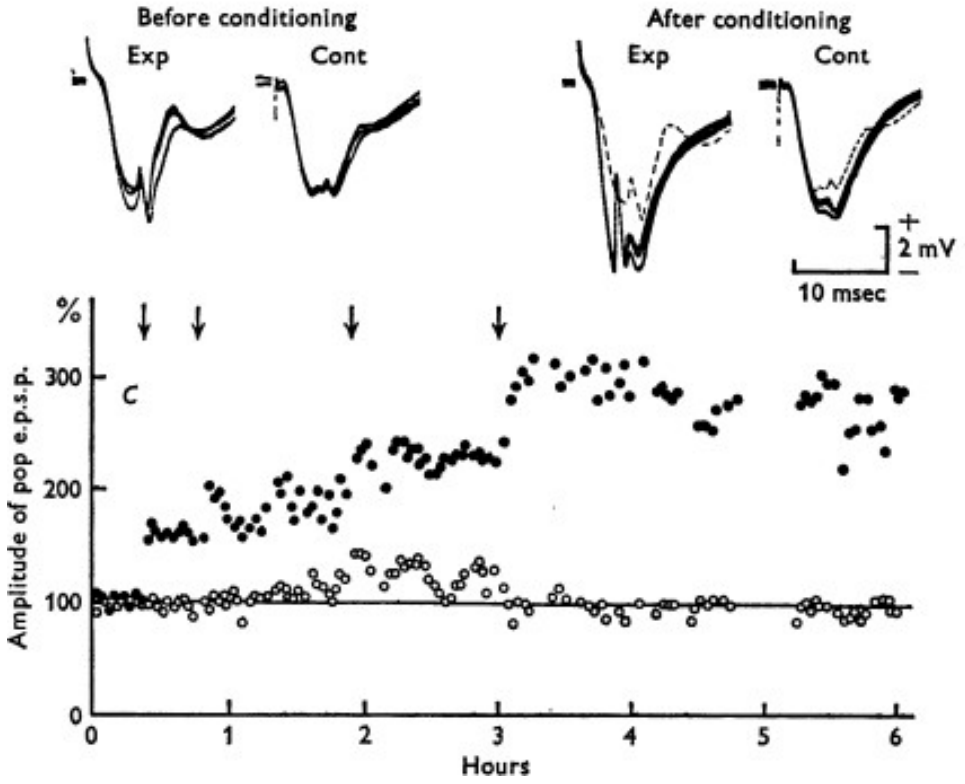
Induction: LTP through high frequency stimulation

hippocampus (*in vivo*)

pre 10-20 Hz for 10-15 sec

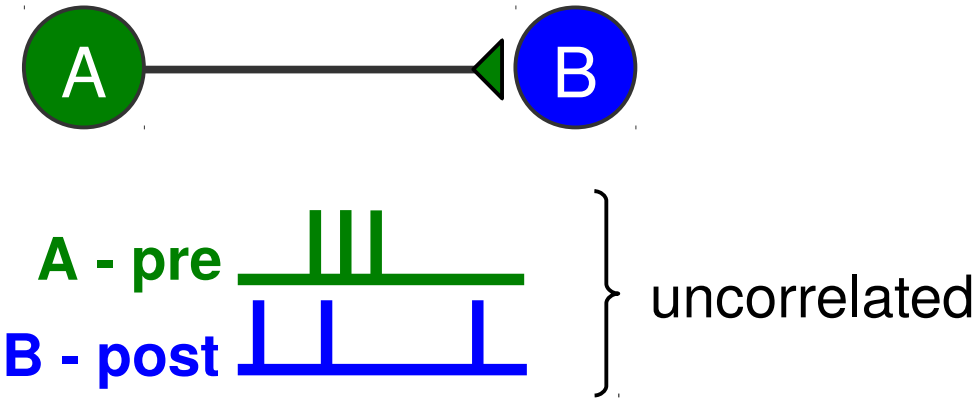
post or 100 Hz for 3-4 sec





[Bliss and Lømo 1973]

LTD induction: postulate of Stent

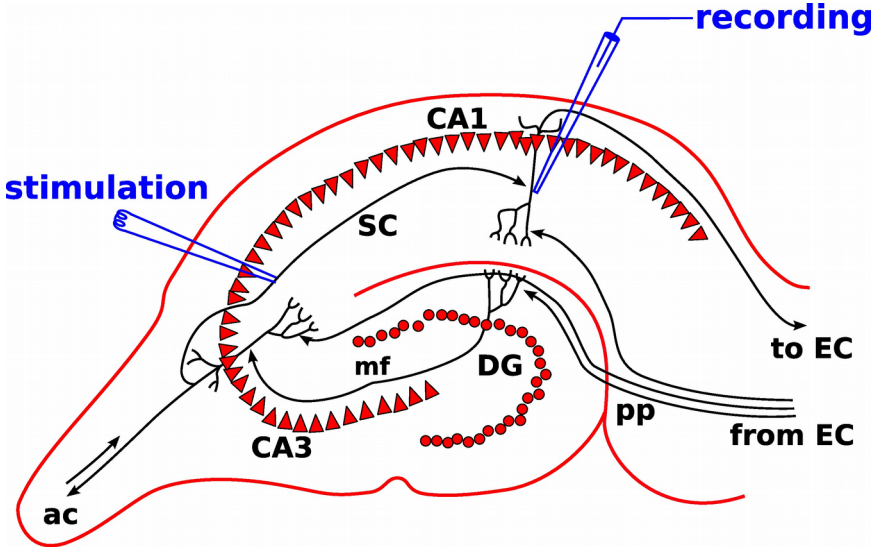


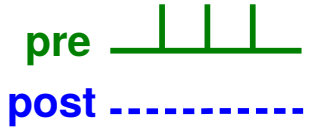
"When the presynaptic axon of cell A *repeatedly* and *persistently* fails to excite the postsynaptic cell B while cell B is firing under the influence of other presynaptic axons, metabolic change takes place in one or both cells such that A's efficiency, as one of the cells firing B, is *decreased*."

[G. Stent 1973;
see also Sejnowski 1977, von der Malsburg 1973, Bienenstock et al. 1982]

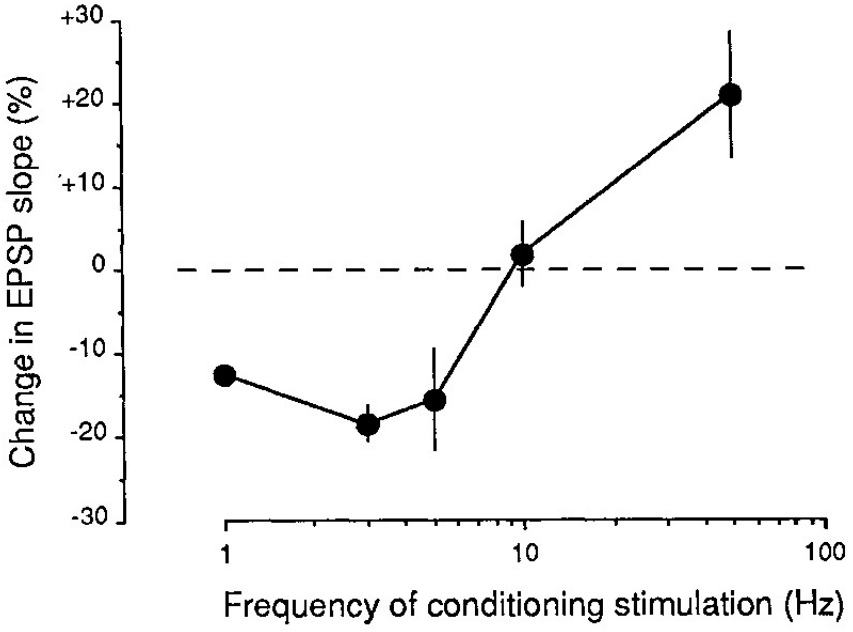
Plasticity induction: LTD obtained at low frequencies

hippocampus (slices)





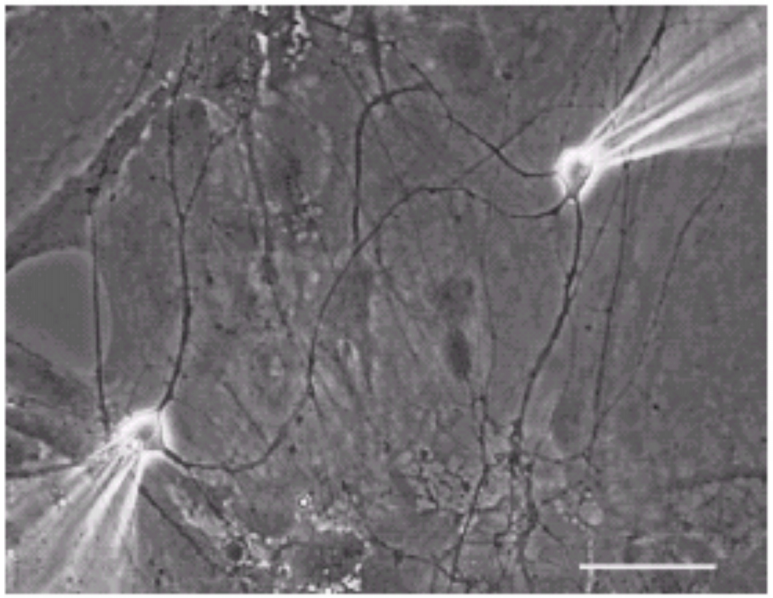
900 pulses at 1-50 Hz



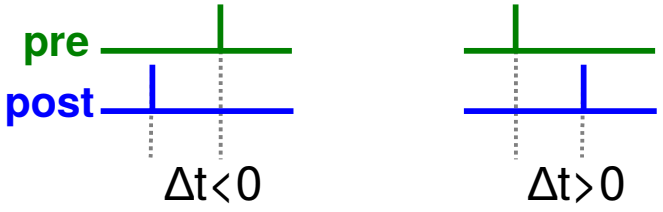
[Dudek and Bear 1992;
Dunwiddie and Lynch 1978]

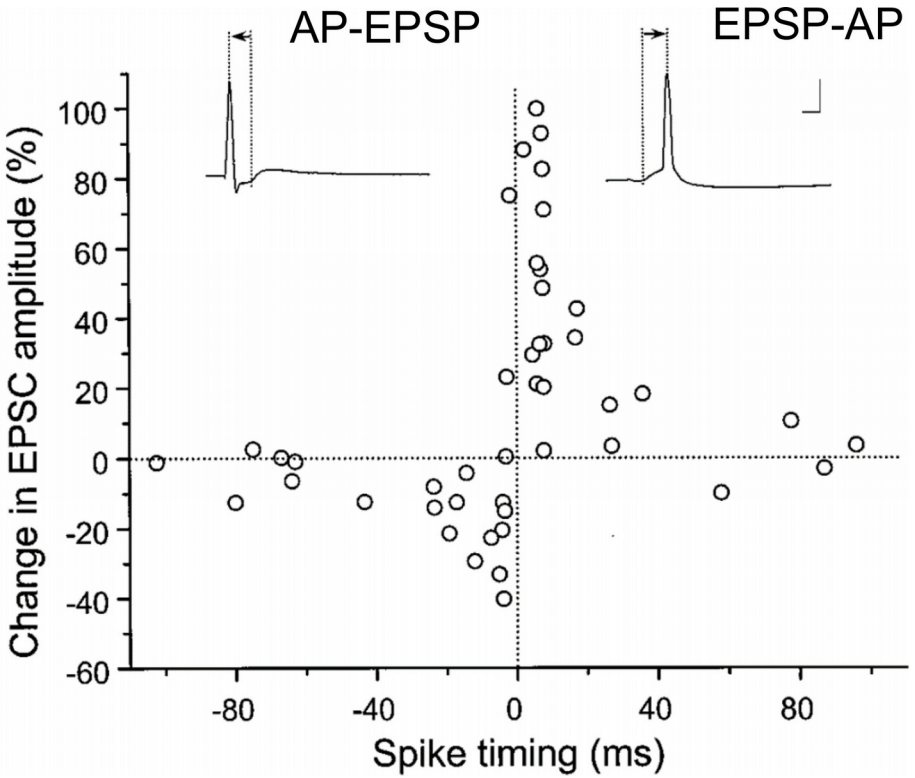
STDP : plasticity from single spike-pairs

hippocampal cultures



60 pairings @ 1 Hz





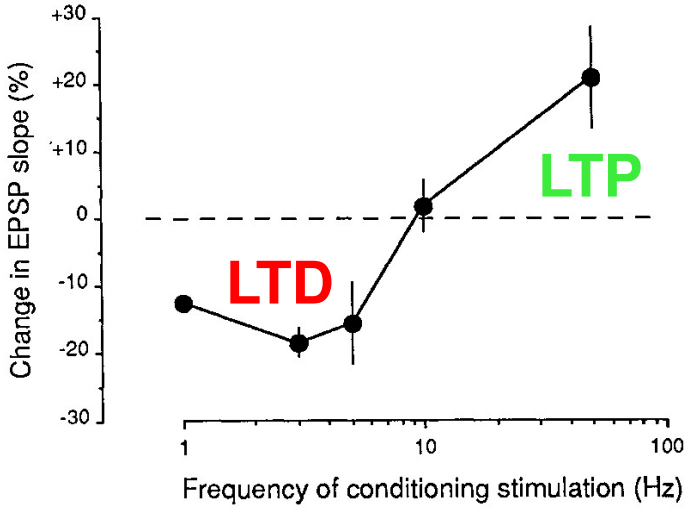
[Bi & Poo, J Neurosci 1998]

[Magee & Johnston 1997; Zhang et al. 1998; Markram et al. 1997; Sjöström et al. 2001; Feldman 200]

Frequency-dependent plasticity and STDP

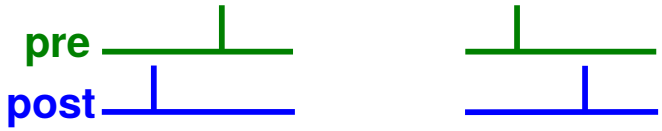
frequency-dependent plasticity

900 pulses at 1-100 Hz

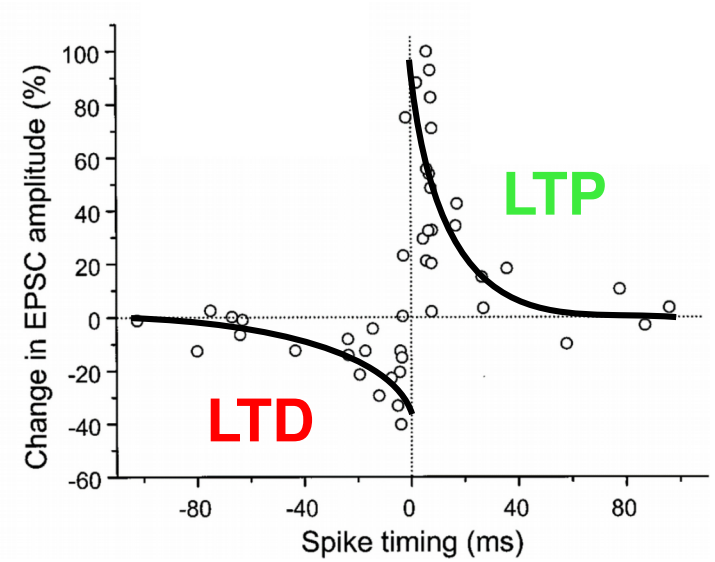


[Dudek and Bear 1992;
Dunwiddie and Lynch 1978]

spike timing-dependent plasticity



60 pairings @ 1 Hz



[Markram et al. 1997; Bi & Poo 1998;
Zhang et al. 1998]

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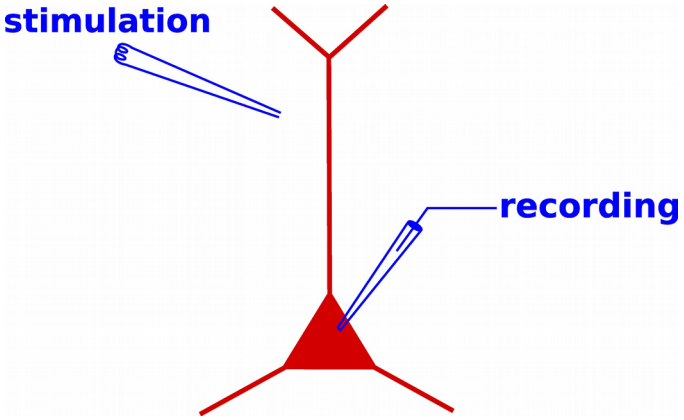
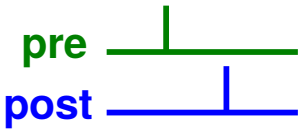
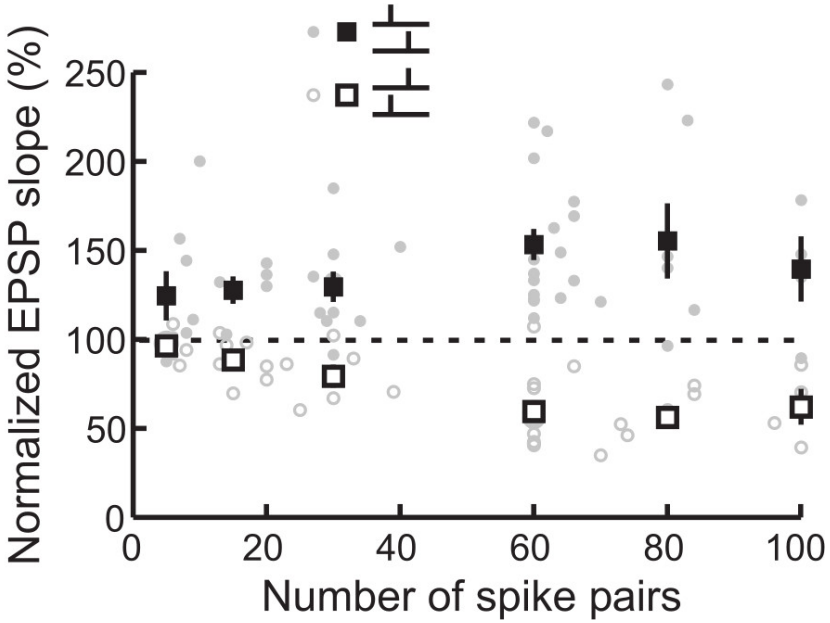
Number of pairing





XX pairings @ 0.2 Hz



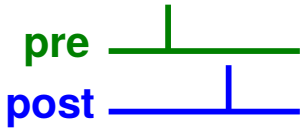
[Froemke et al. 2006]

- LTP induced with a few pairs
- LTD requires the presentation of ~20 stimulation pairs

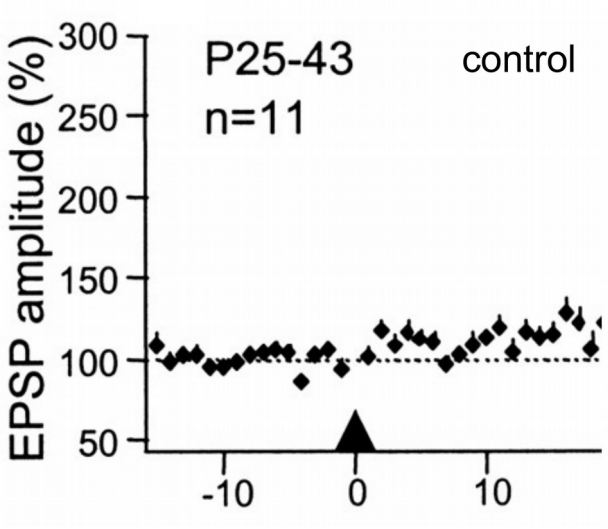
Role of synaptic inhibition

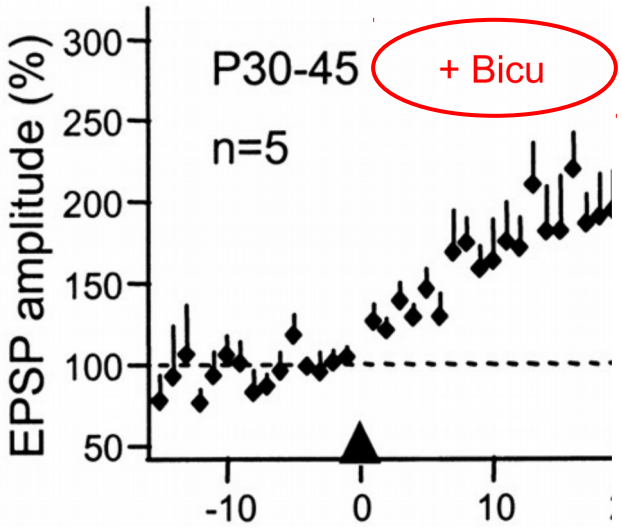
control
→ no plasticity

inhibition blocked
→ LTP



Δt = +10 ms

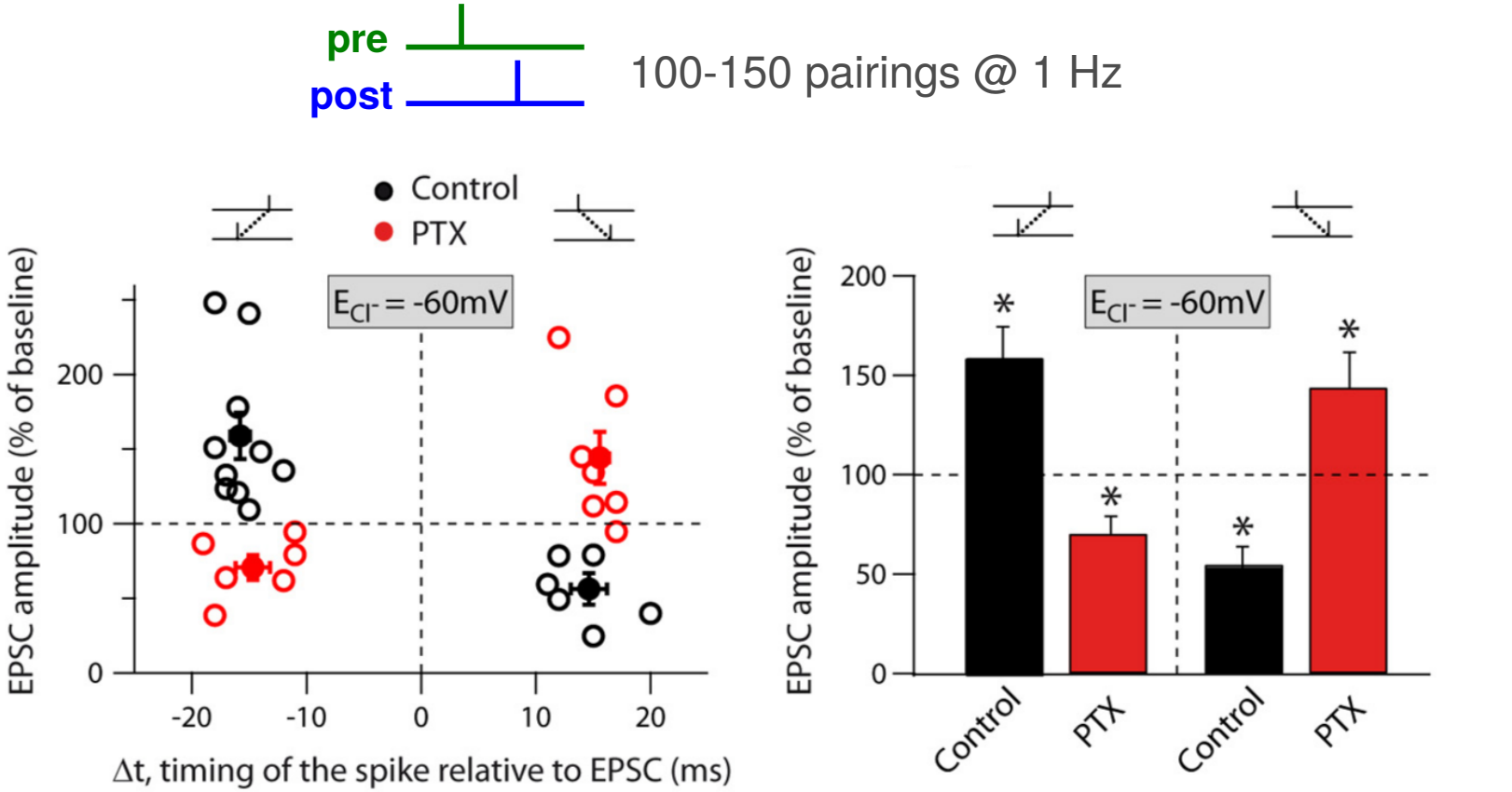




[Mederith et al. *J Neuroci* 2003]

Bicuculline is a competitive antagonist of $GABA_A$ receptors.

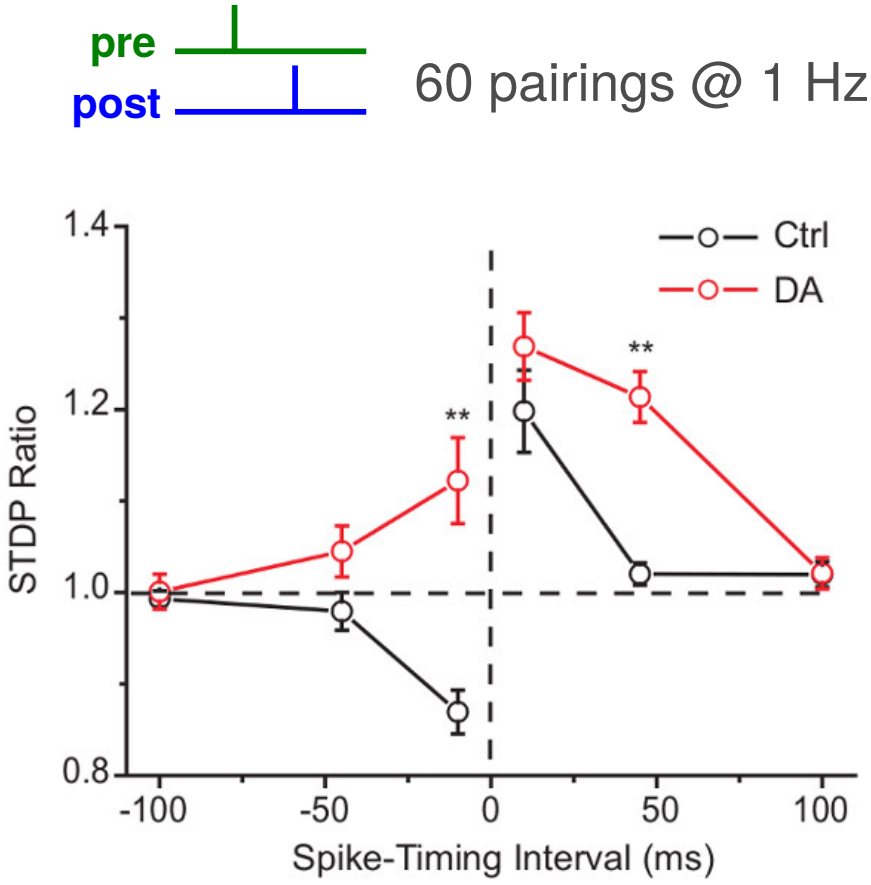
Role of synaptic inhibition



[Paille et al. *J Neurosci* 2013]

- inhibition inverts the STDP curve

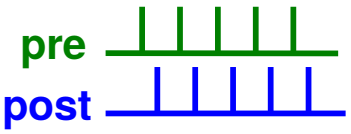
Role of neuromodulation - Dopamine



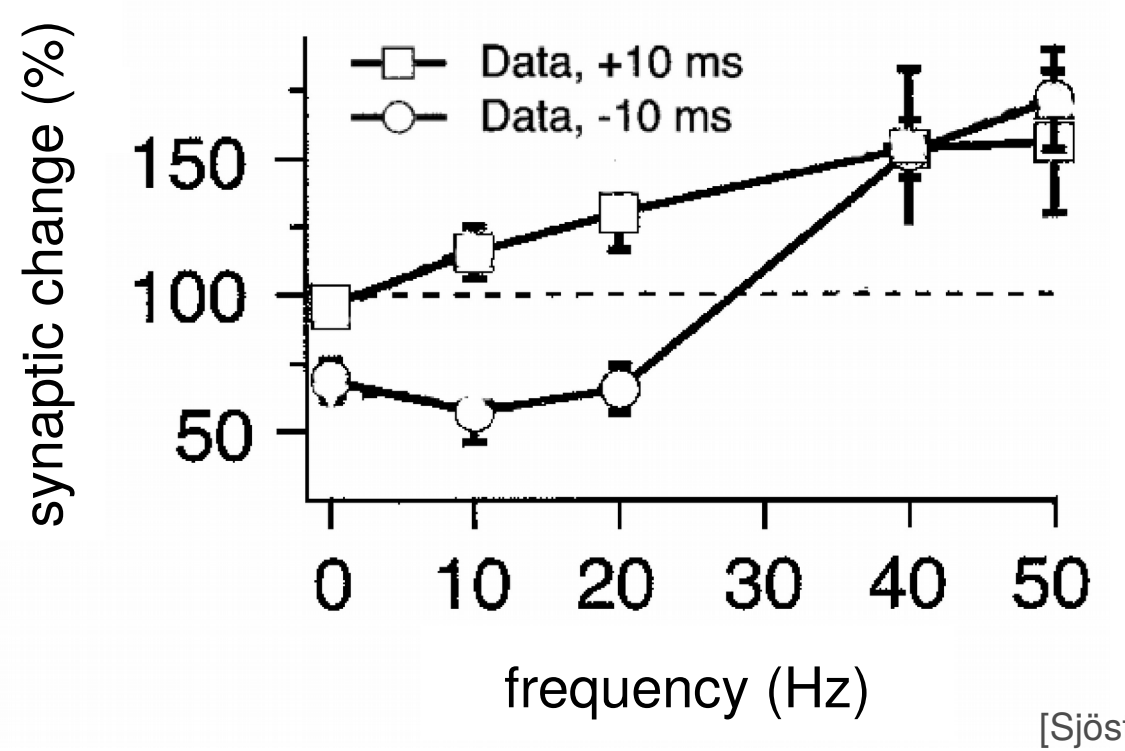
[Zhang et al. *PNAS* 2009]

- dopamine controls sign and magnitude of plasticity

STDP depends on frequency of spike-pairs



5 pairings, 15x @ 0.1 Hz

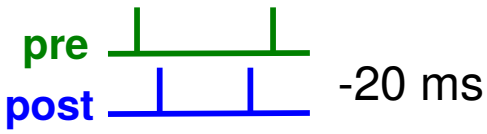


[Sjöström et al. *Neuron* 2001]

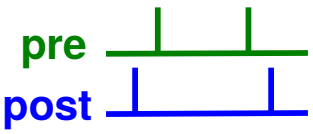
- post-pre pairings induce LTD at low and LTP at high frequencies

Non-linearity in STDP induction protocols

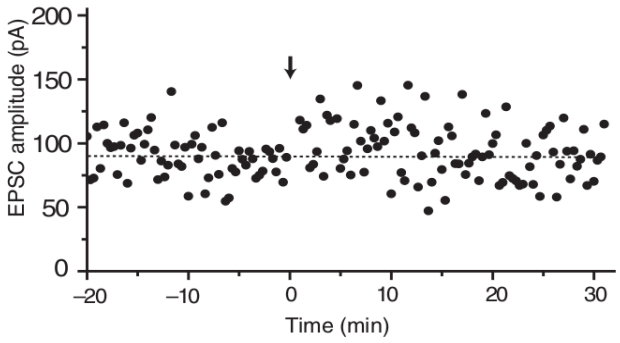
+5/-5 ms, 60 pairings @ 1 Hz

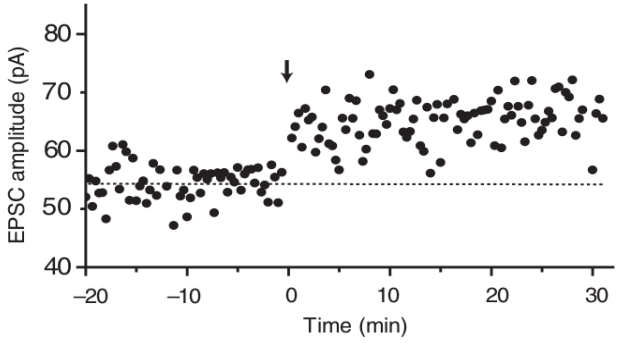


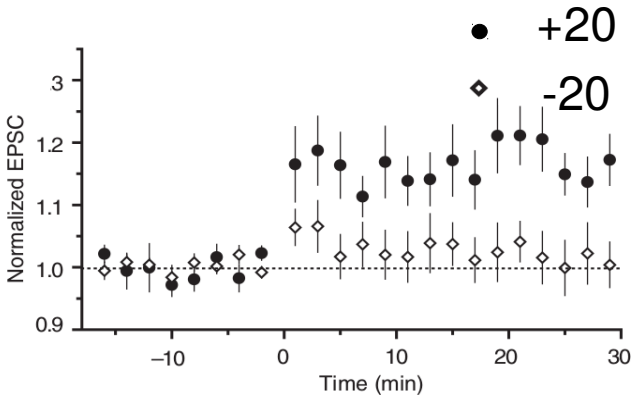
-20 ms



+20 ms



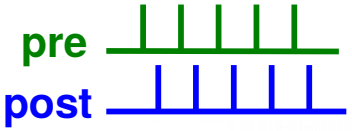




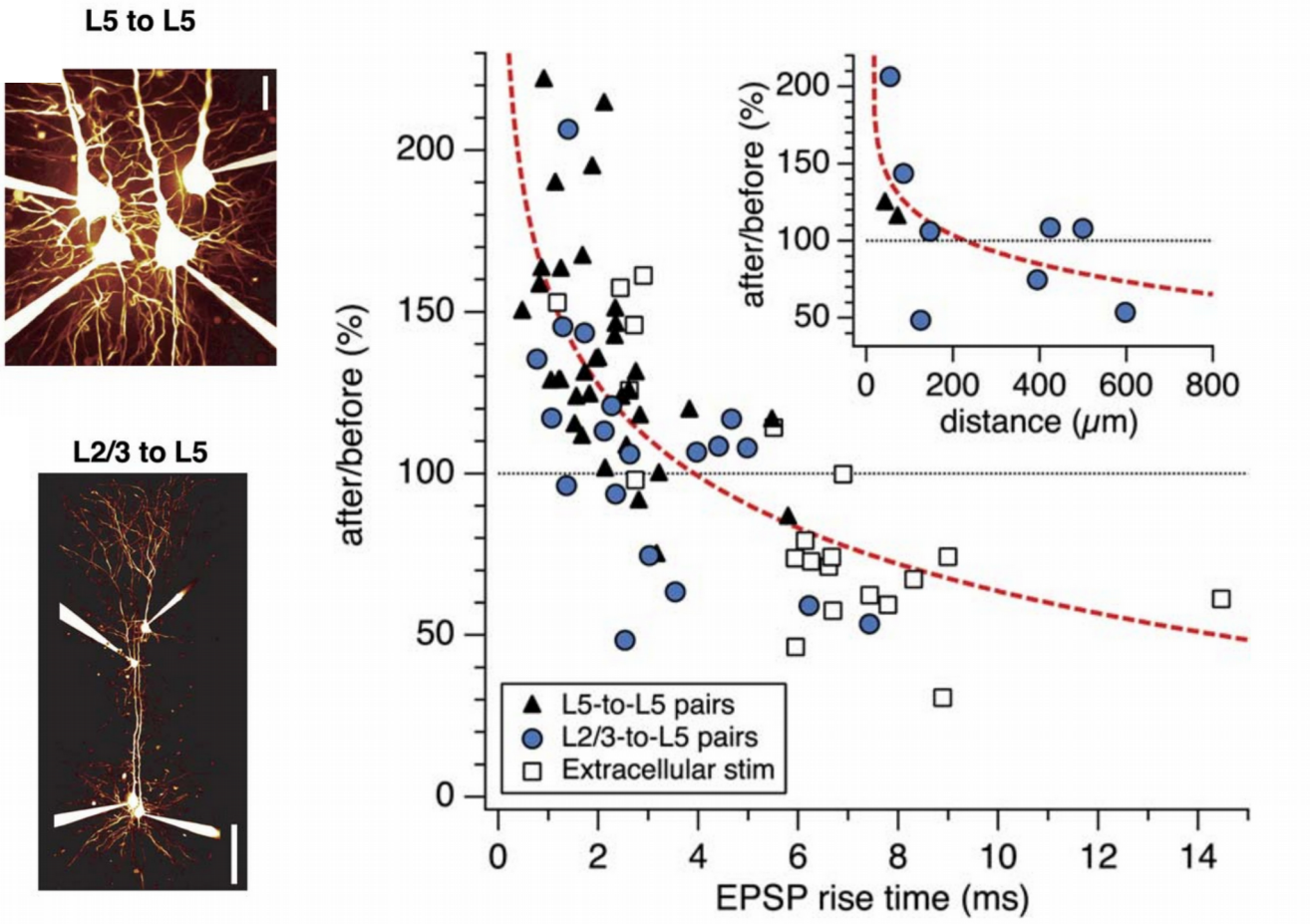
[Wang et al. *Nat Neurosci* 2005]

- Quadruplets : order of pre-post, post-pre pairs determines plasticity outcome

STDP depends on synaptic location



Δt=+10 ms, 5 pairings, 50 Hz, 15x @ 0.1 Hz

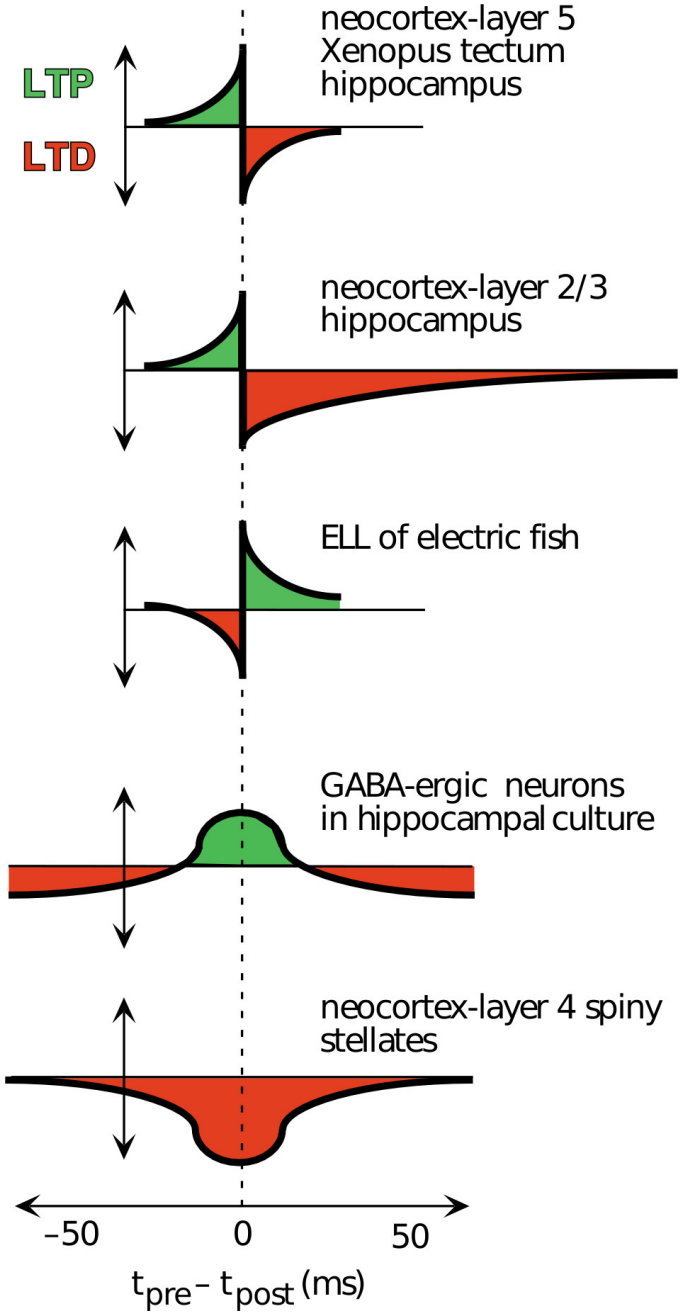


[Sjöström & Hausser, *Neuron* 2006]

[Froemke et al. *Nature,* 2005; Letzkus et al. *J Neurosci* 2006]

- Proximal synapse : LTP
- Distal synapse : LTD

STDP windows depends on brain structure, synapse type



[Abbott & Nelson *Nat Neurosci* 2000]

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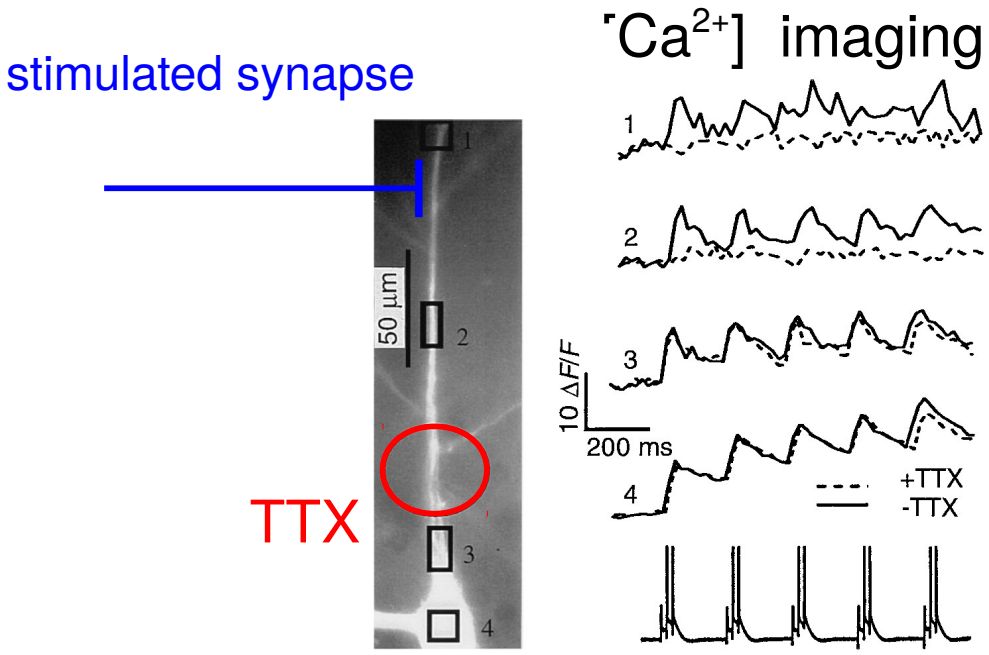
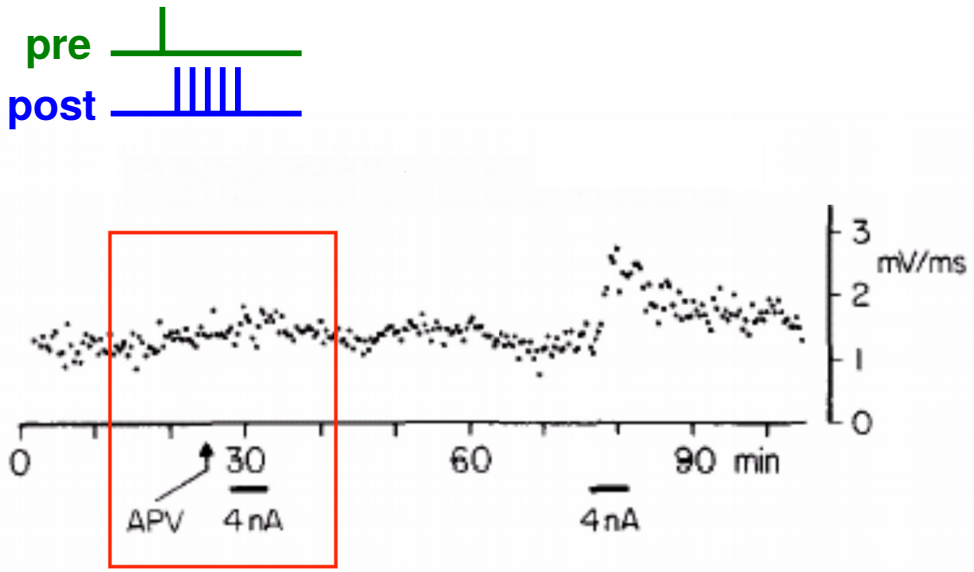
Backpropagating action potential required for STDP



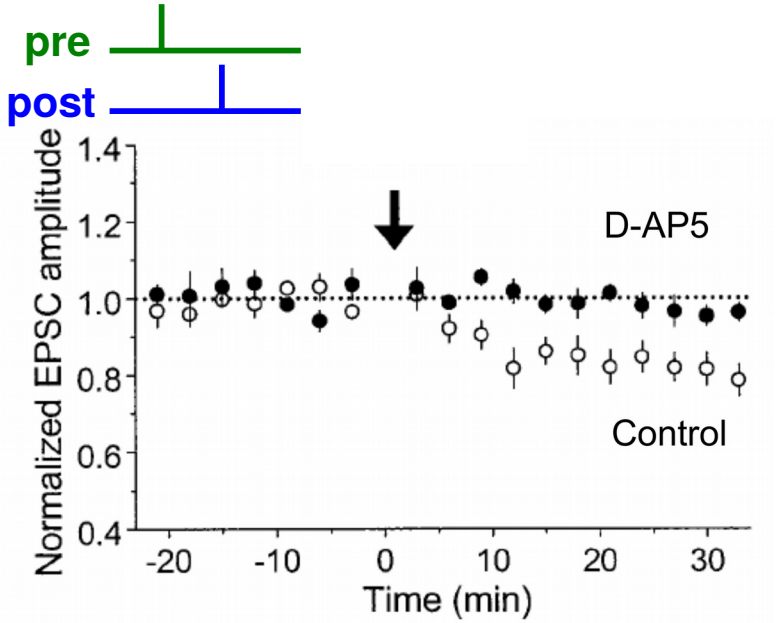
[Magee & Johnston *Science* 1997]

- Backpropagating action potential provides postsynaptic depolarization required for STDP

STDP requires NMDA receptor activation



[Gustafsson et al. *J Neurosci* 1987]

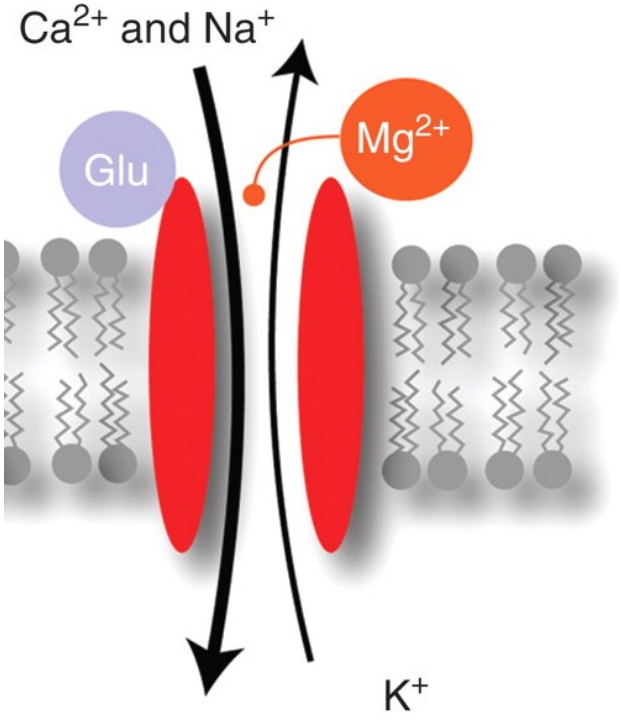


[Bi & Poo *J Neurosci* 1998]

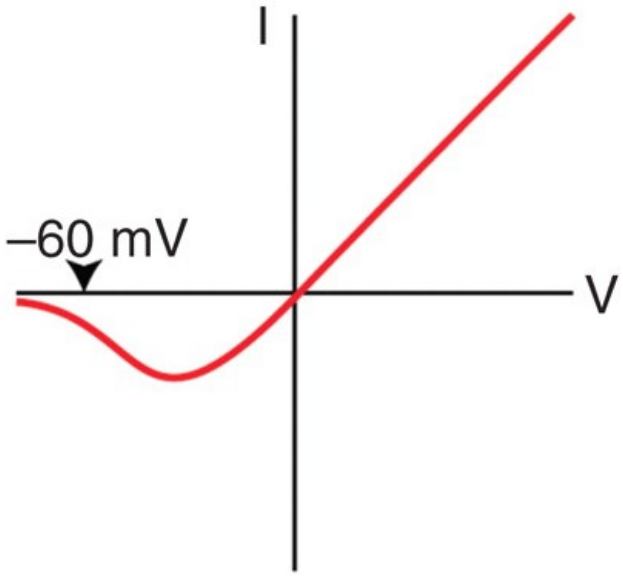
- NMDAR antagonist blocks STDP induction

  (D-AP5 or APV is a selective NMDA receptor antagonist)

Postsynaptic NMDA receptor



current-voltage relationship



- coincidence detector :

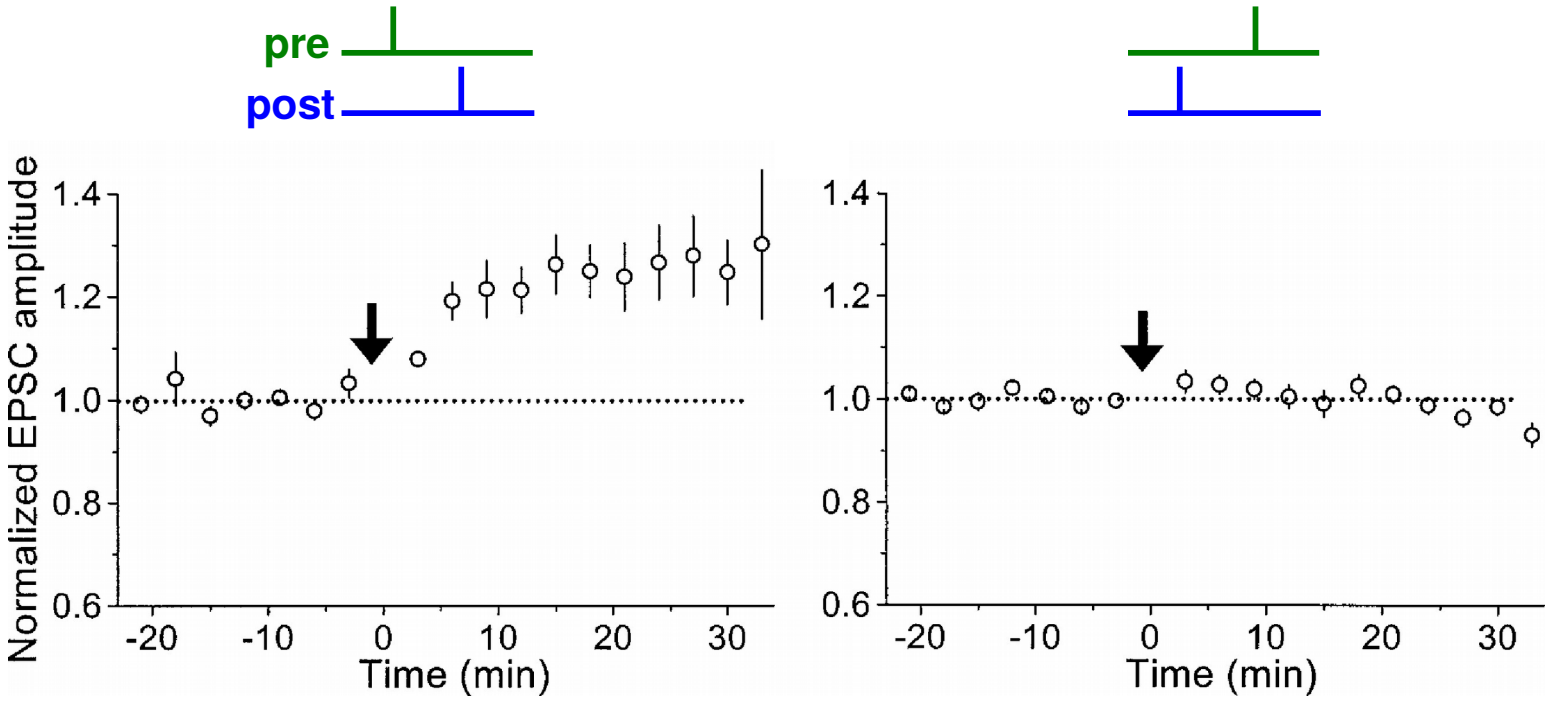
  presynaptic action potential → glutamate (Glu)

  postsynaptic depolarization → Mg^{2+} block is expelled

- calcium permeable

Voltage-dependent Ca channels required for LTD

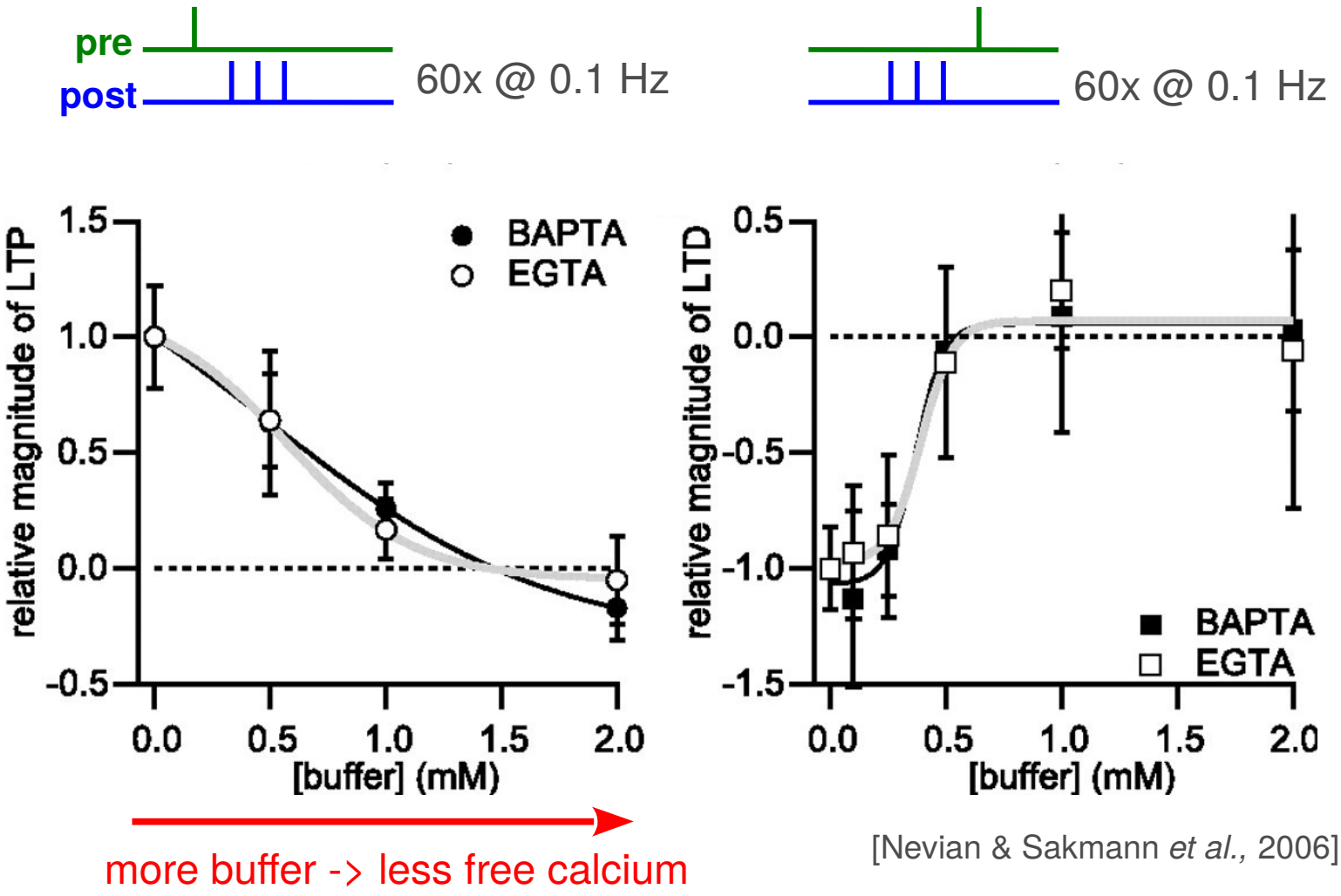
+ nimodipine (L-type calcium channel blocker)



[Bi & Poo *J Neurosci* 1998]

- LTD but not LTP involves the activation of L-type calcium channels

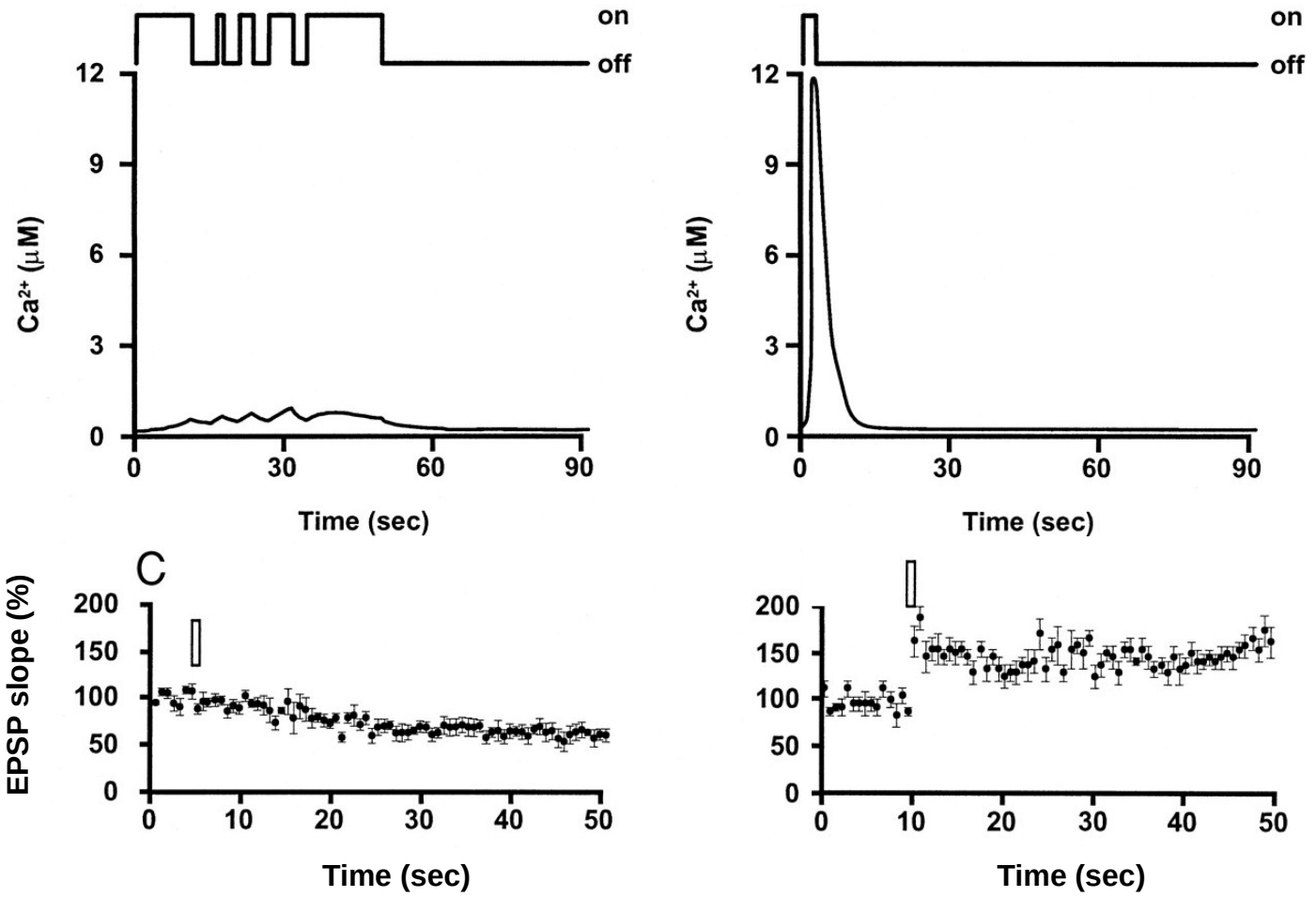
Postsynaptic calcium *required* for plasticity



[Nevian & Sakmann *et al.*, 2006]

- LTP/LTD equally sensitive to fast and slow [Ca^{2+}] buffers

Postsynaptic calcium *sufficient* for plasticity

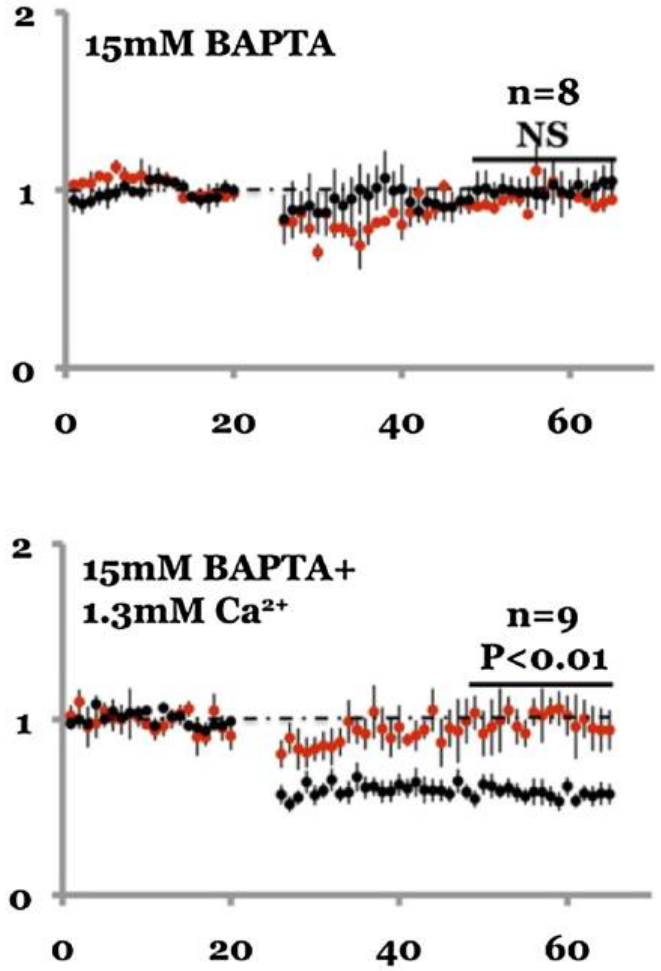


[Malenka *et al. Science* 1988; Yang *et al.*, *J Neurophysiol* 1999]

- LTP induced by brief, large amplitude [Ca^{2+}] increases
- prolonged, modest rise in [Ca^{2+}] elicits LTD

NMDA receptor activation but no calcium required for LTD

[Nabavi *et al.*, 2013, PNAS]

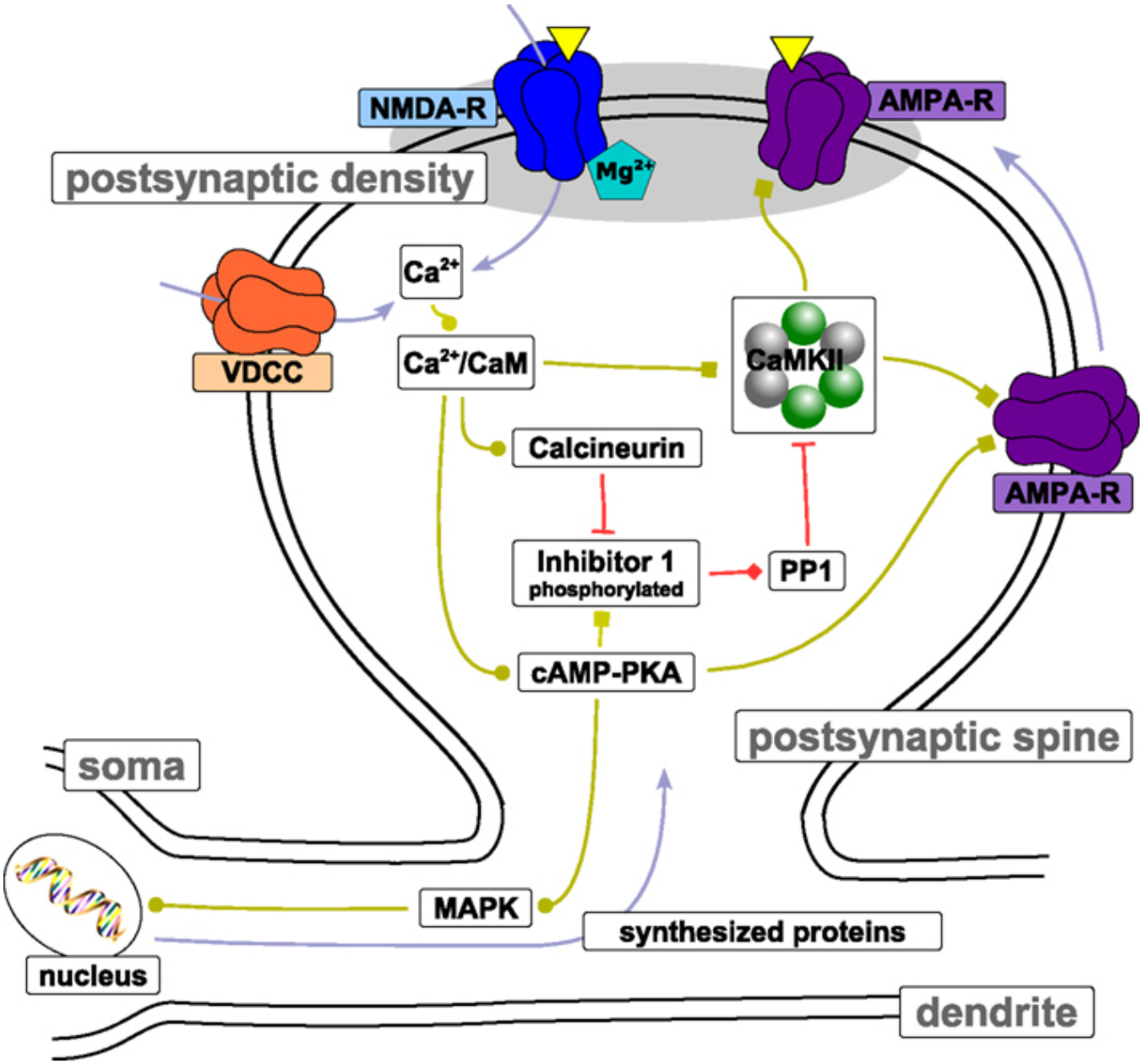


complete block of calcium

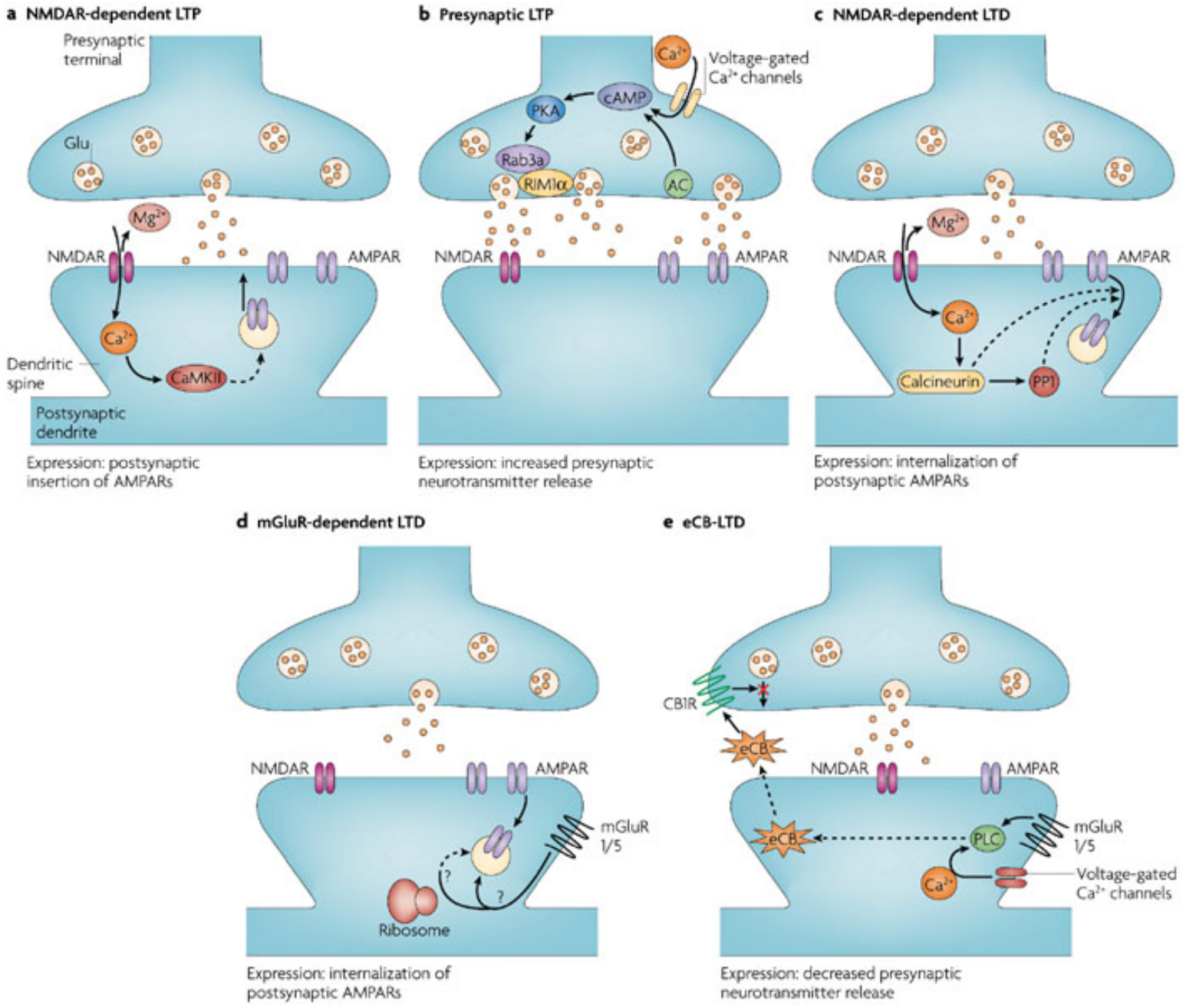
maintaining resting calcium levels

- ligand binding to NMDA receptors is sufficient for LTD
- basal levels of [Ca^{2+}] are permissively required

Signal pathways downstream of Calcium



Diversity of induction and expression pathways



[Kauer, Malenka. *Nat Rev Neurosci* 2010]

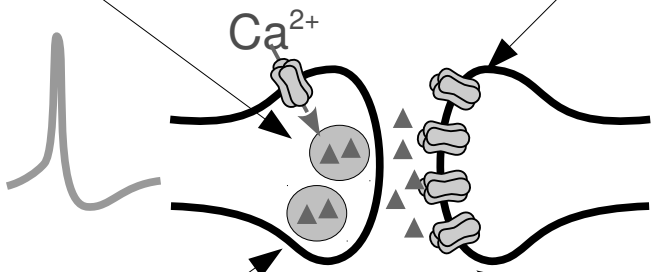
Expression of long-term changes

presynaptic

postsynaptic

neurotransmitter vesicle number

number of AMPA receptors



probability of vesicle release

conductance of AMPA receptors

Outline

1. STDP : introduction and history

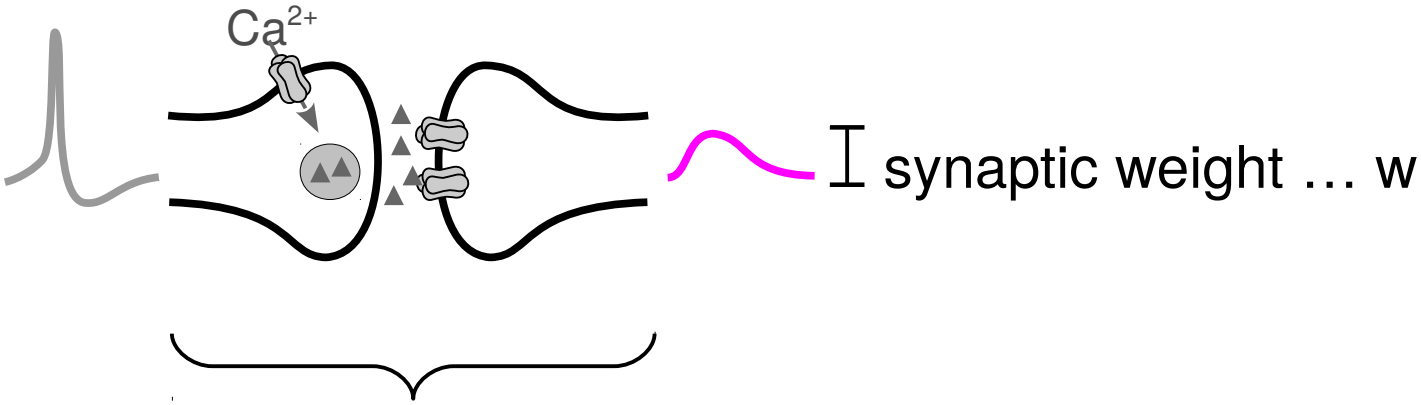
2. Phenomenology of STDP

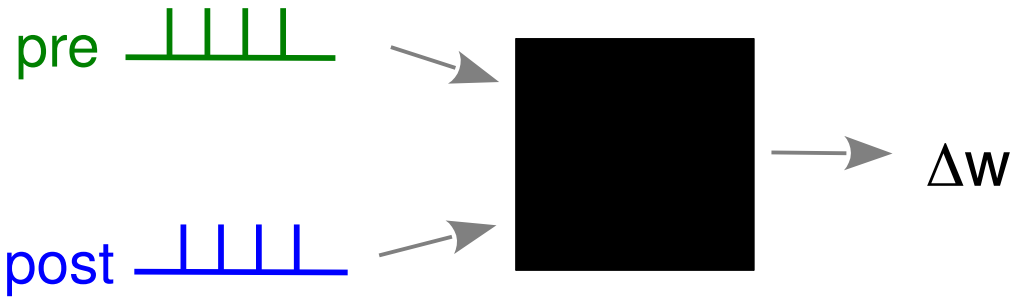
3. Induction mechanisms

4. Biophysical models of STDP

5. STDP *in vivo*

Modeling : translation from spikes to plasticity results





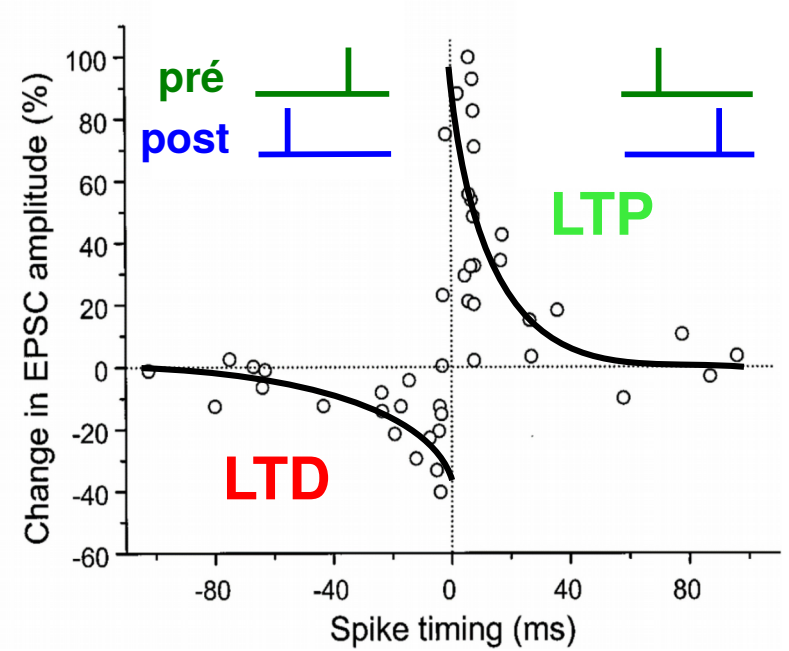
“Standard” STDP model

- spike-timing based rules : $\Delta w_{ij} = f\left(\{t_{ik}\},\{t_{jk}\}\right)$

- “standard” STDP :

$$f\left(\{t_{ik}\},\{t_{jk}\}\right) = \sum_{k,k'} F\left(t_{ik} - t_{jk'}\right)$$

$$F(\Delta t) = \begin{cases} A_+ \exp\left(-\Delta t/\tau_+\right) & \Delta t > 0 \\ A_- \exp\left(-\Delta t/\tau_-\right) & \Delta t < 0 \end{cases}$$



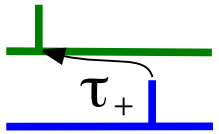
- Variations of the rule :
  * additive/multiplicative
  * All-to-all spike pairings / nearest neighbors

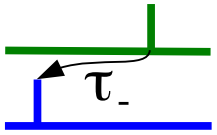
- **Problems :** does not depenend on firing rate
  does not resolve the nonlinearities of plasticity

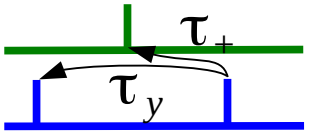
More recent plasticity models

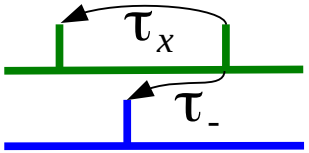
Triplet-based model

[Pfister & Gerstner, 2006; Clopath et al., 2010]

LTP : A_2^+ 

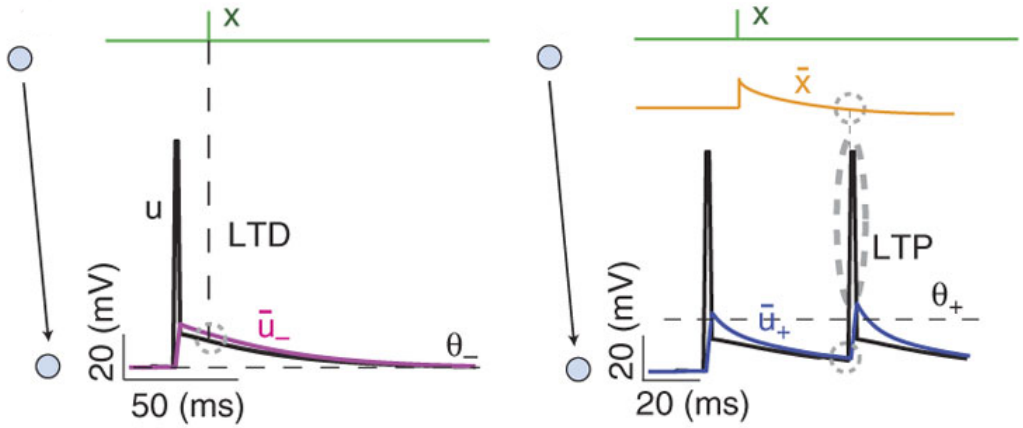
LTD : A_2^- 

LTP : A_3^+ 

LTD : A_3^- 

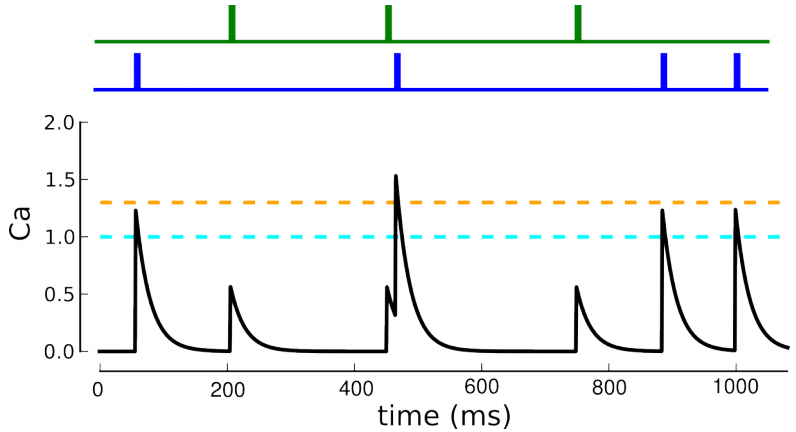
Model based on postsynaptic potential

[Clopath *et al.*, 2010]



Calcium-based model

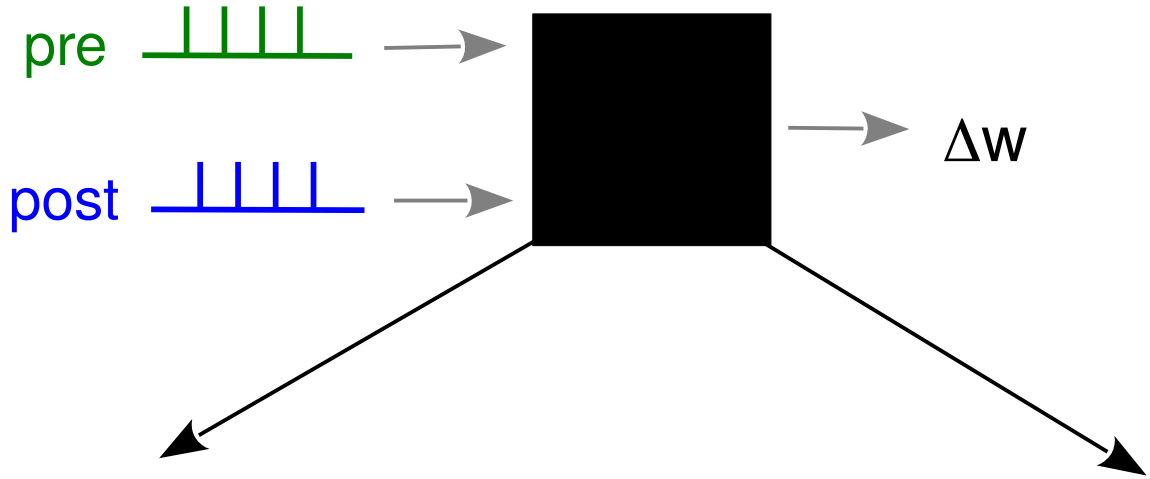
[Shouval *et al.* 2002,Graupner &Brunel 2012]



potentiation

depression

Modeling approaches : phenomenological vs. biophysical



phenomenological models of LTP/LTD

- use pre- and postsynaptic spike times or rate to calculate change in synaptic strength
- conversion can involve arbitrarily complex mathematical models

biophysical models of LTP/LTD

- resolve *parts* of the underlying biological machinery involved in the induction of plasticity
- degree of biological detail varies largely

Modeling studies : phenomenological vs. biophysical

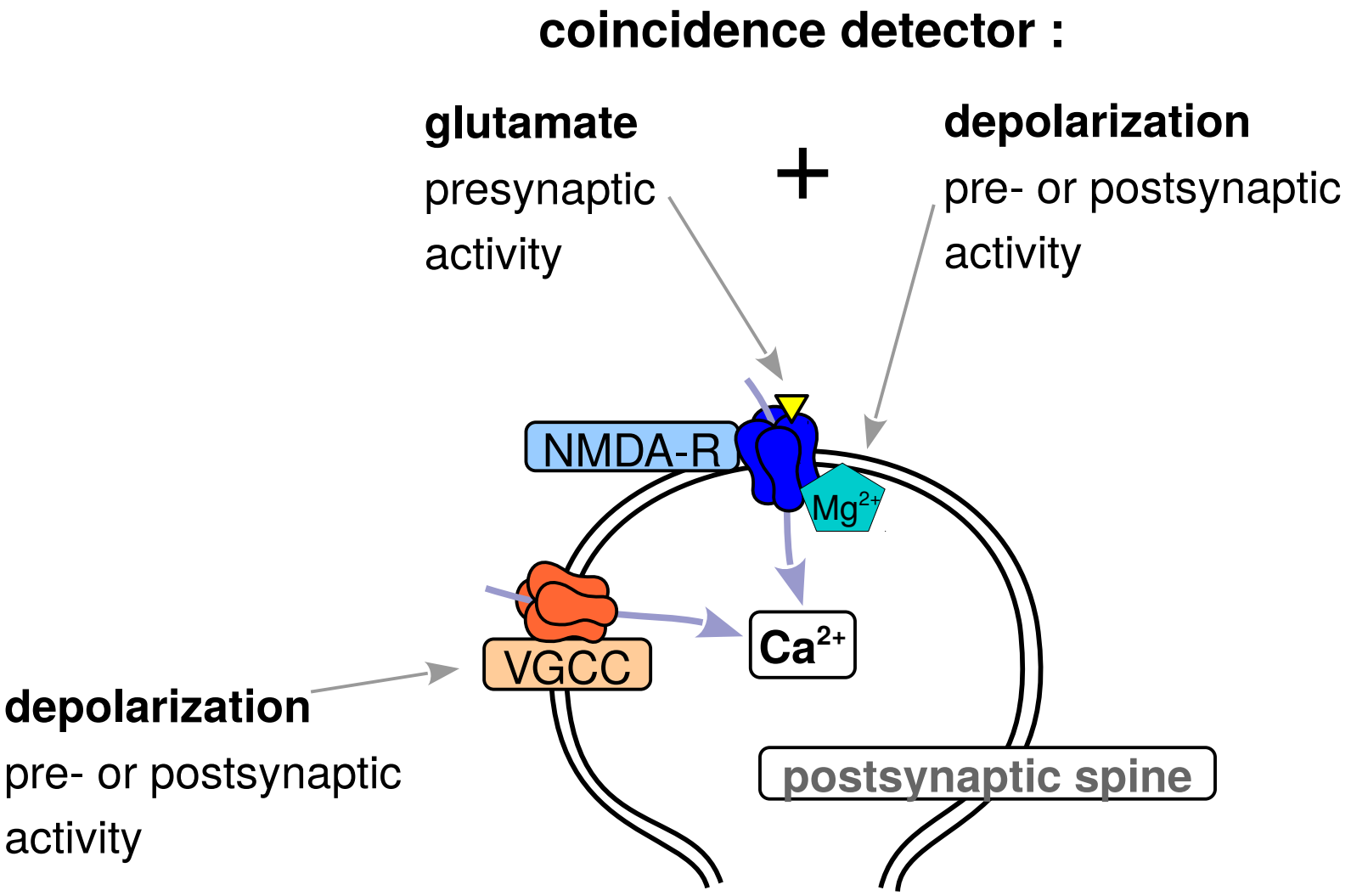
phenomenological models of LTP/LTD

- rate-based plasticity models
  [Hebb, 1949; Bienenstock *et al.*, 1982; Oja, 1982]
- spike-timing based models
  [Gerstner *et al.*, 1996; van Rossum *et al*. 2000; Song, 2000; Pfister & Gerstner, 2006]

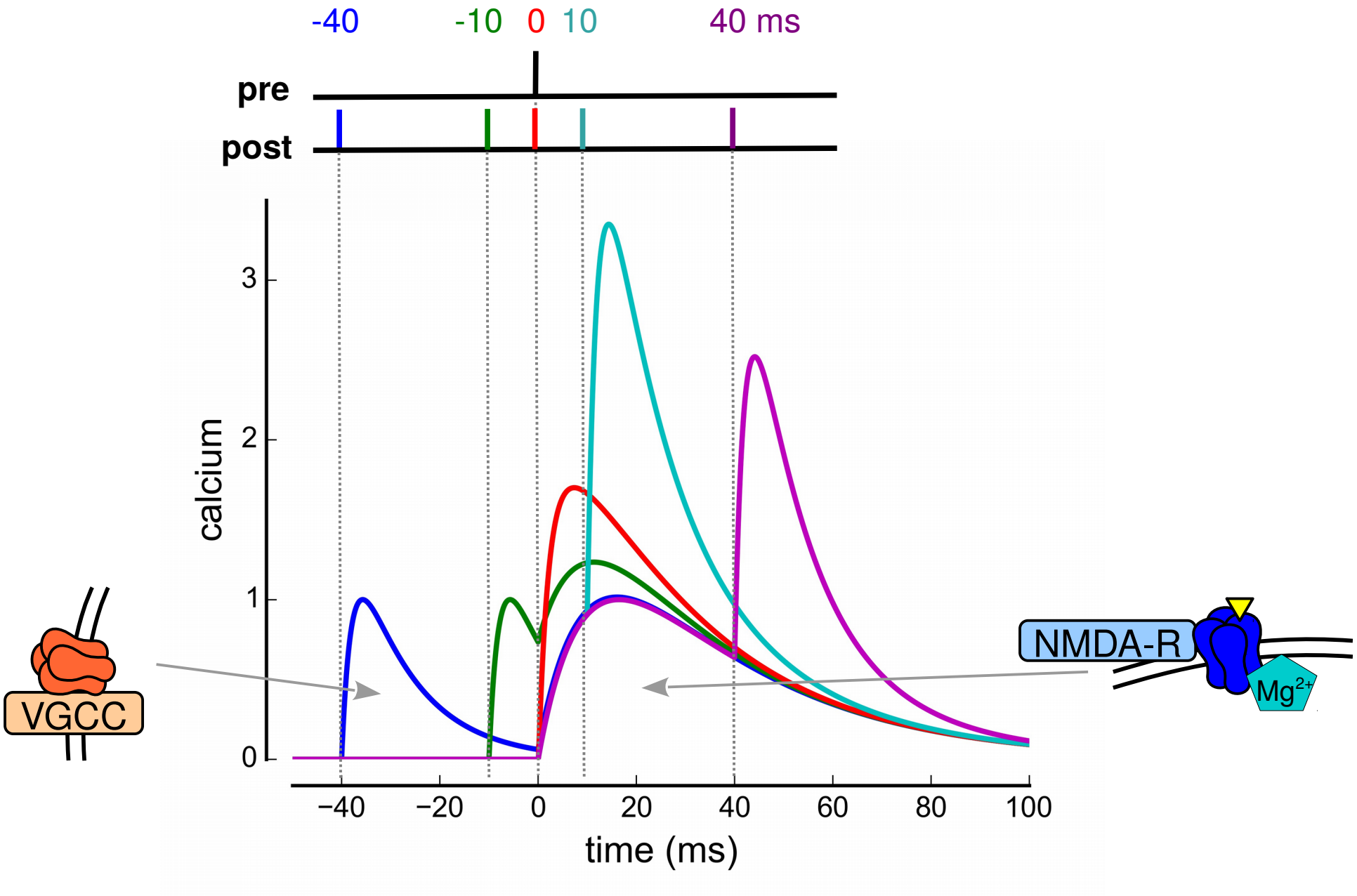
biophysical models of LTP/LTD

- Ca^{2+} – dynamics based models
  [Karmarkar *et al*., 2002; Shouval *et al.*, 2002; Rubin *et al*., 2005; Graupner & Brunel 2012]
- CaMKII kinase-phosphatase system
  [Crick 1984; Lisman, 1985; Okamoto & Ichikawa, 2000; Zhabotinsky, 2000; Graupner & Brunel, 2007; Urakubo *et al.*, 2008]
- extensive protein networks
  [Bhalla & Iyengar, 1999; Hayer & Bhalla, 2005]
- local clustering of receptors
  [Shouval, 2005]

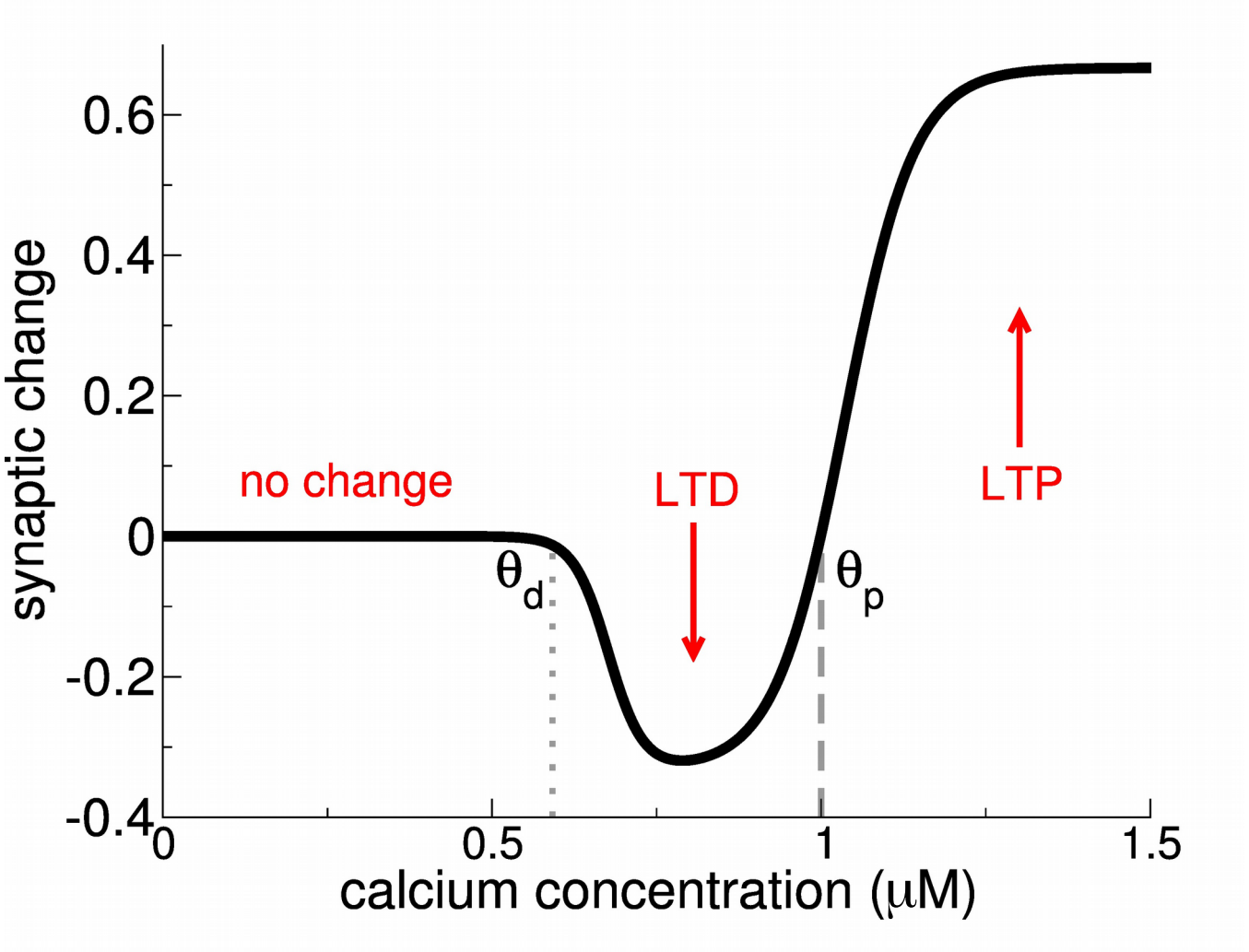
Calcium influx



Calcium transients from spike-pair stimulation

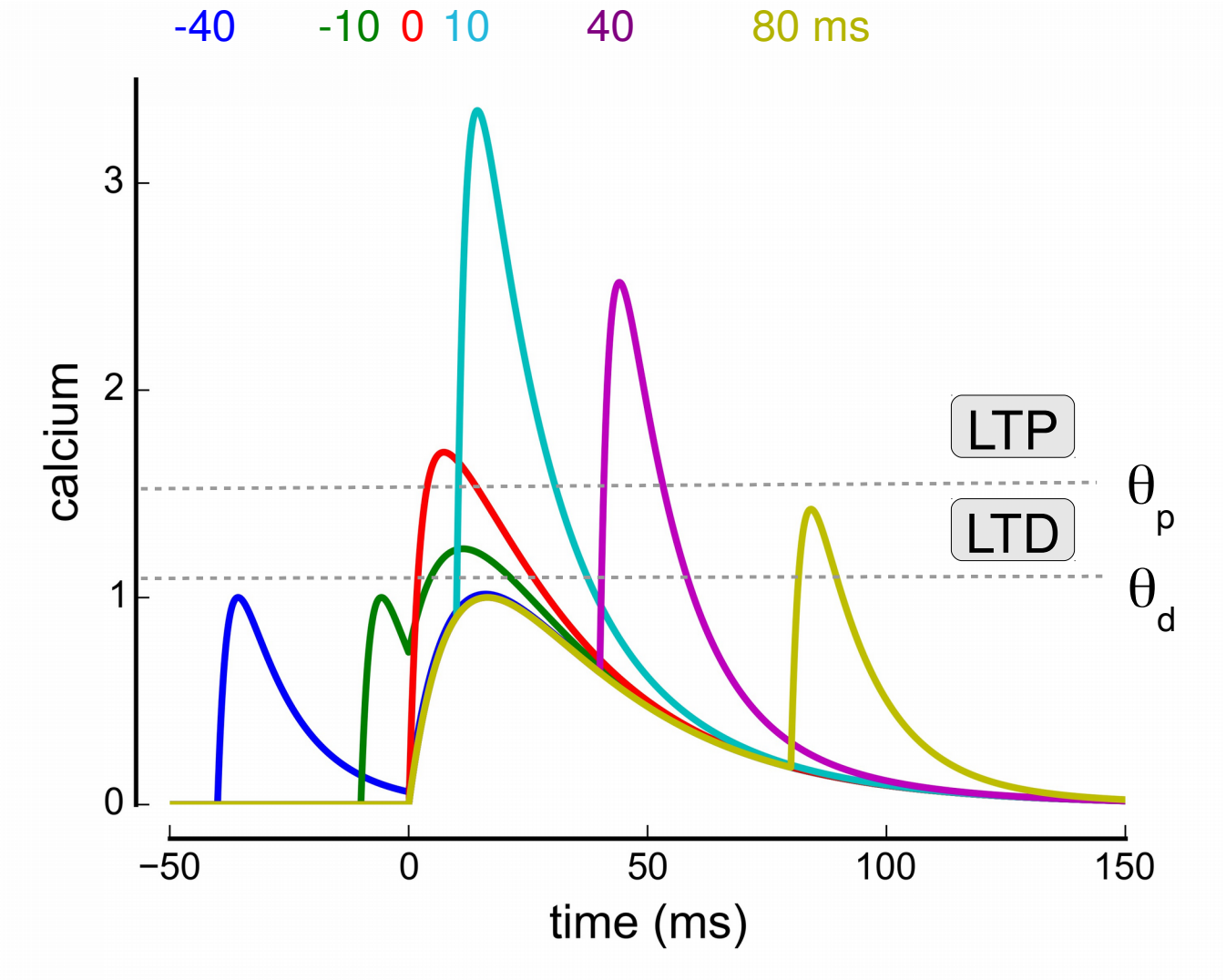


Calcium control hypothesis

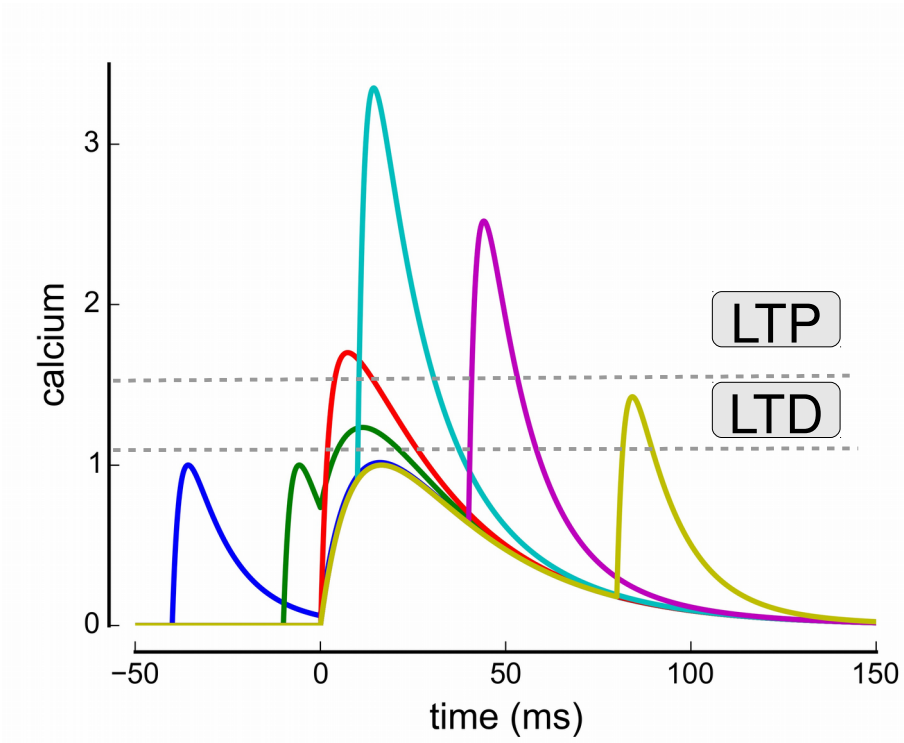


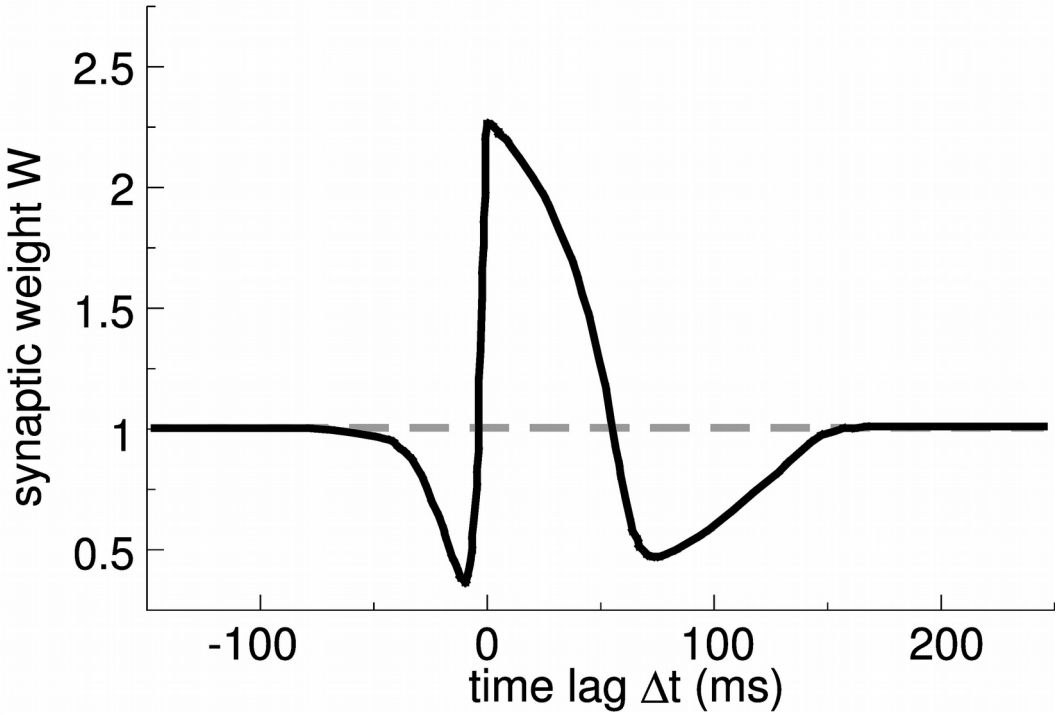
[Shouval *et al.*, *PNAS* 2002]

Calcium transients from spike-pair stimulation



STDP curve from calcium control hypothesis

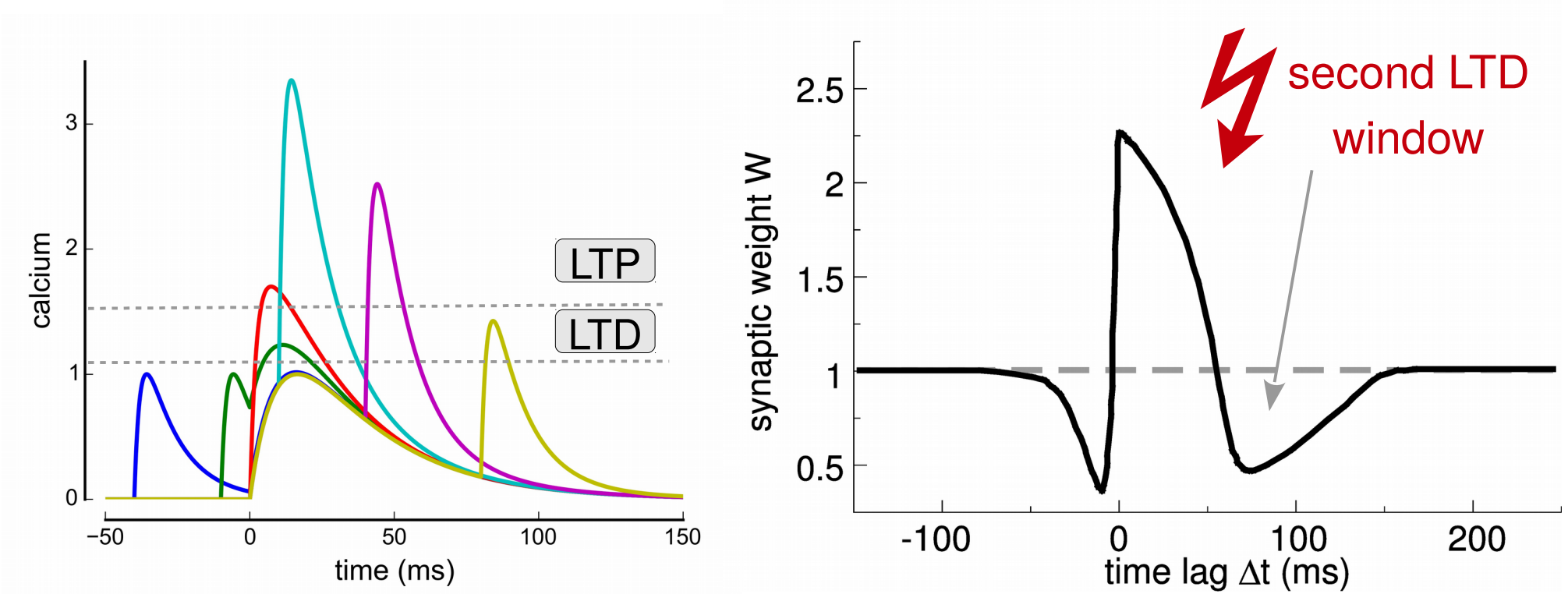




[Shouval *et al.*, *PNAS* 2002]

STDP curve from calcium control hypothesis

[Shouval *et al.*, 2002]

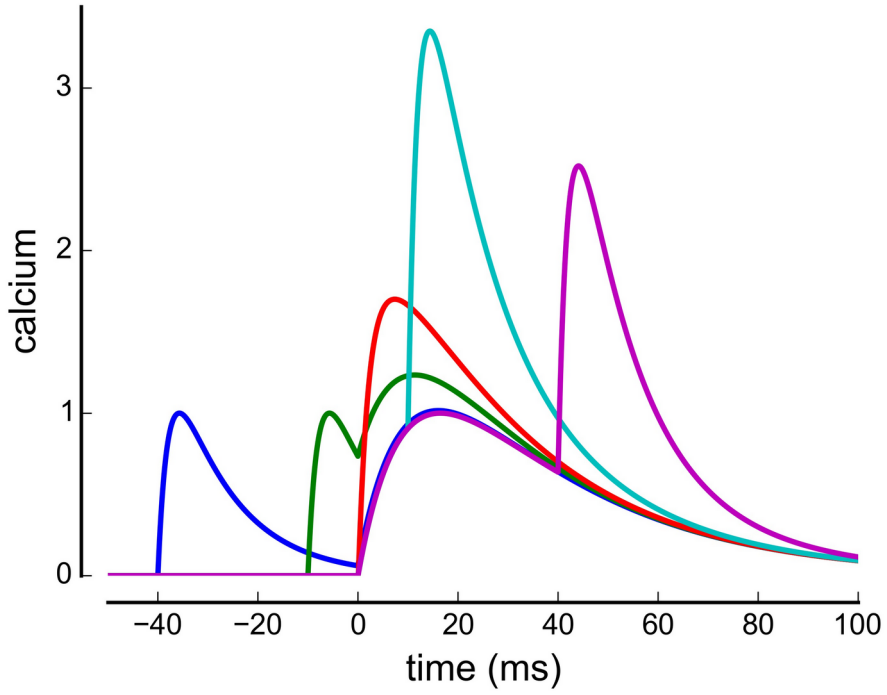


→ [Karmarkar *et al.*, 2002; Badoual *et al.*, 2006; Rubin *et al.*, 2005; Urakubo *et al.*, 2008; Graupner & Brunel 2012 ...]

Question : role of calicum in shaping STDP

I. Can the dynamics of the postsynaptic calcium account for synaptic plasticity induced by spike-pairs ?

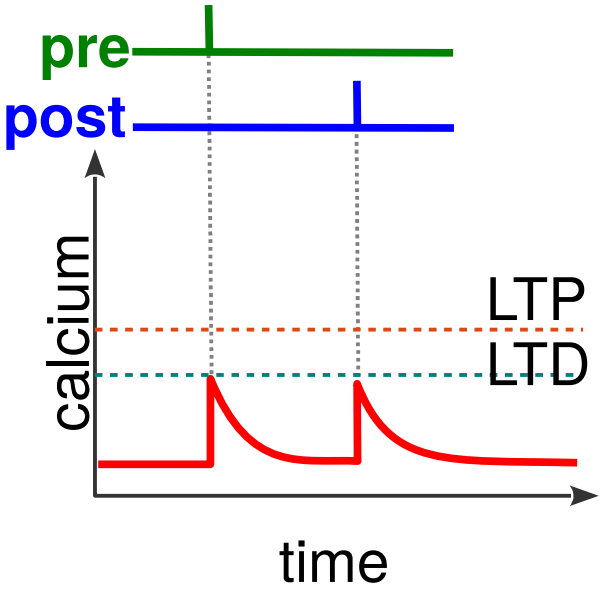
II. To which extent can the STDP phenomenology be explained by calcium ?

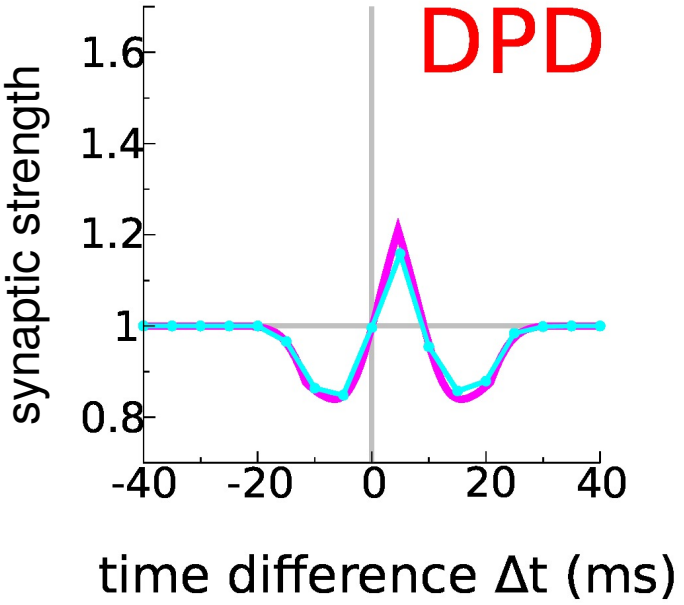


Calcium amplitudes determine shape of STDP curve

$C_{pre} < \theta_d$

$C_{post} < \theta_d$





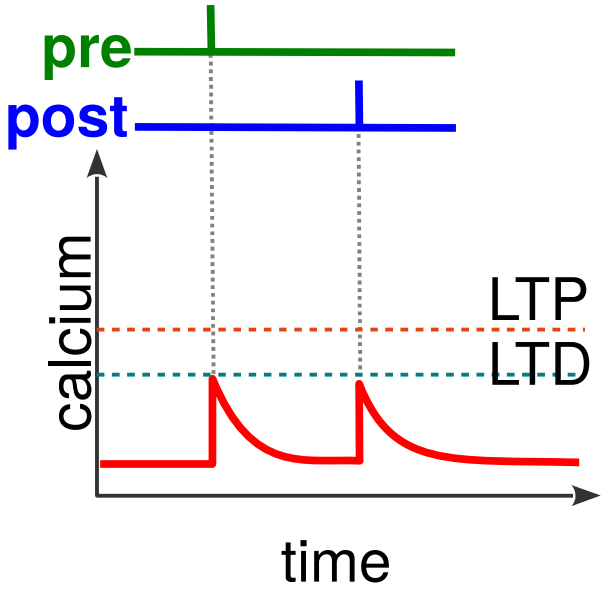
Calcium amplitudes determine shape of STDP curve

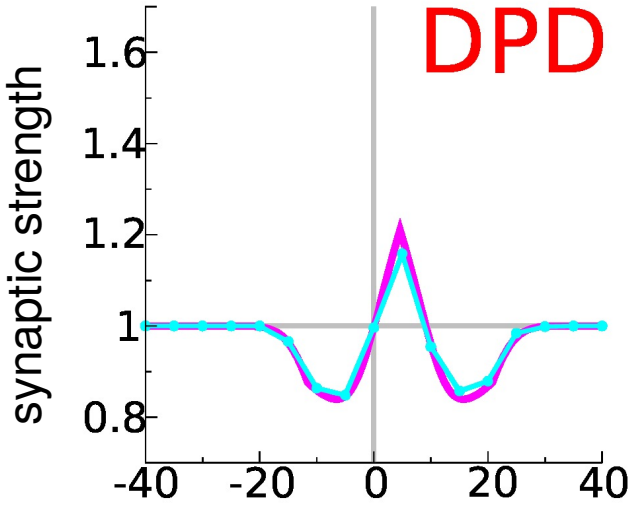
simulation I

Calcium amplitudes determine shape of STDP curve

$C_{pre} < \theta_d$

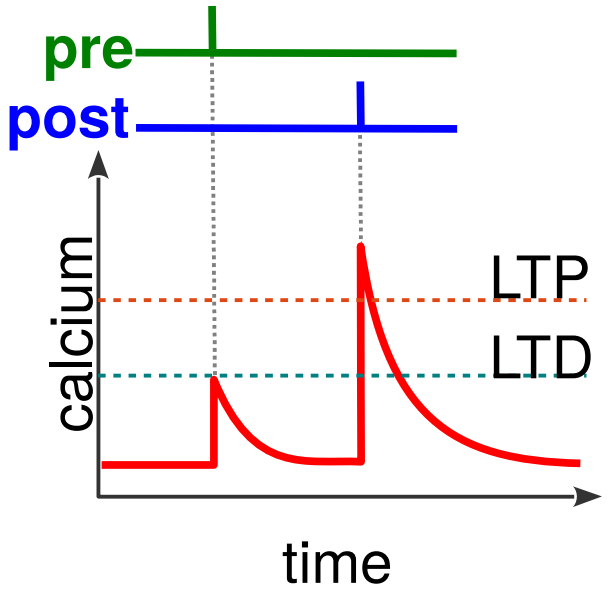
$C_{post} < \theta_d$

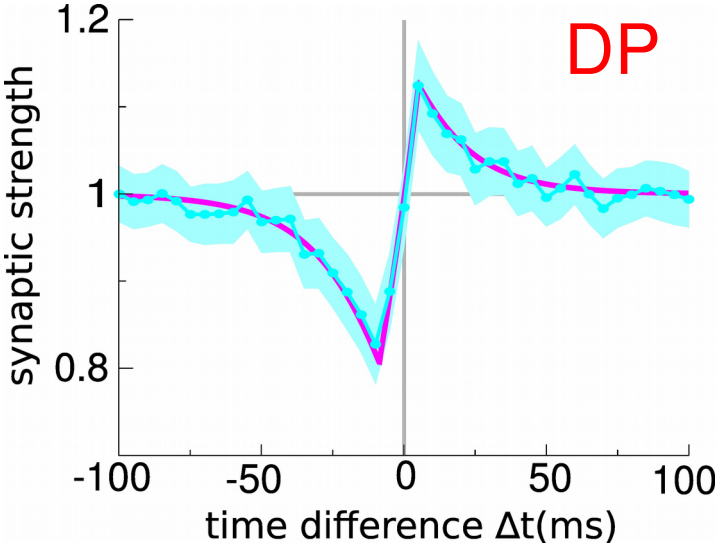




$C_{pre} < \theta_p$

$C_{post} > \theta_p$

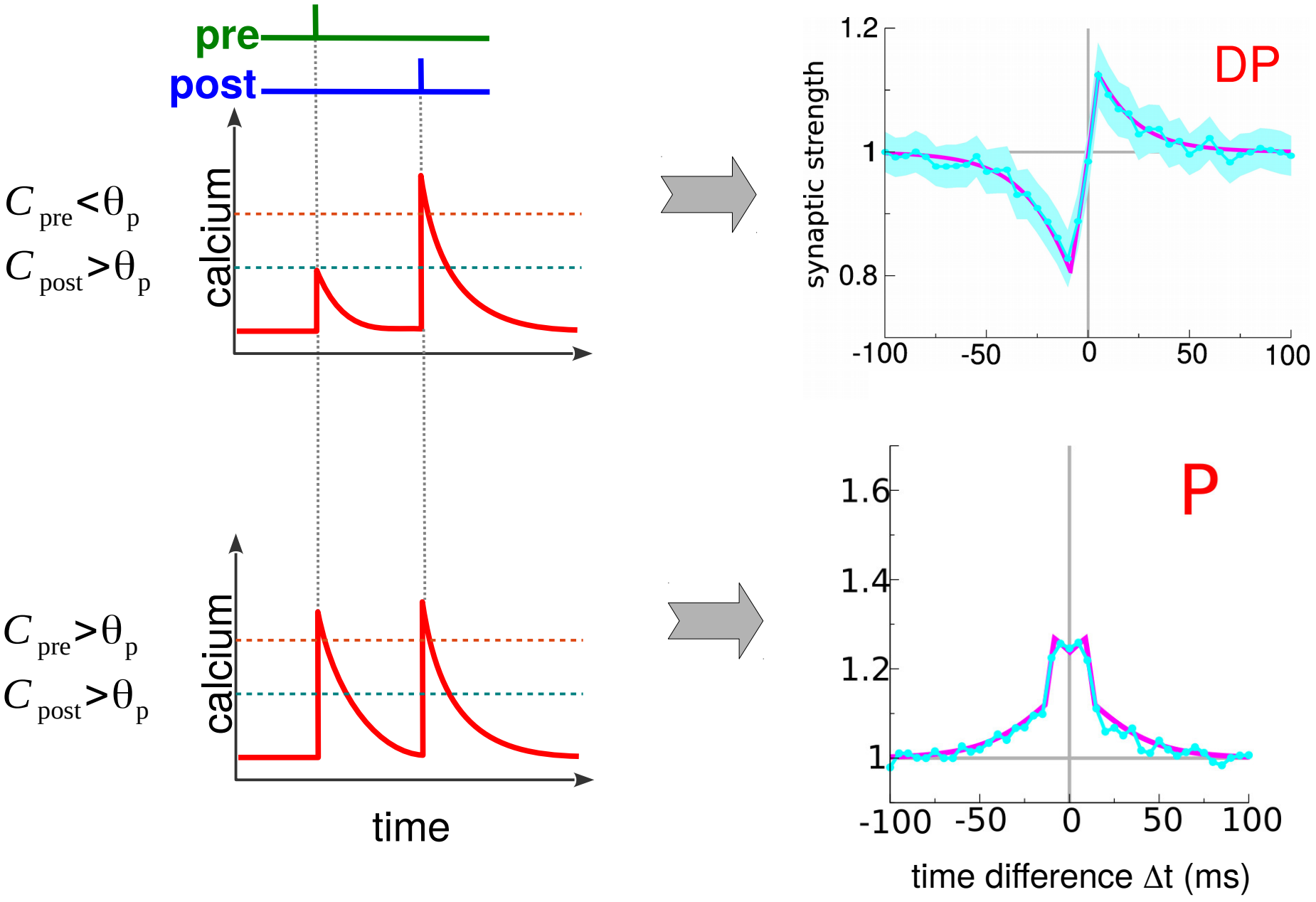




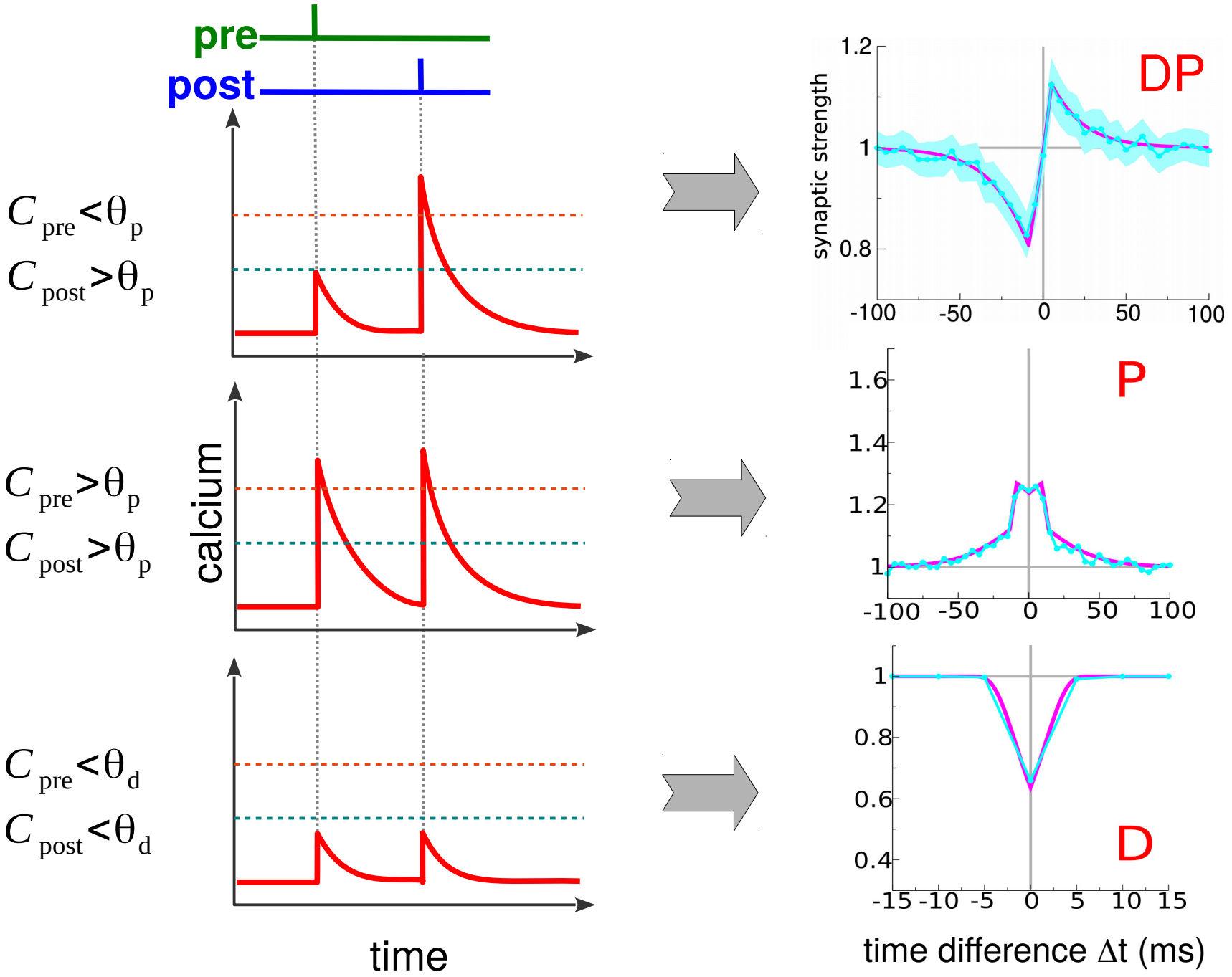
Calcium amplitudes determine shape of STDP curve

simulation II

Calcium amplitudes determine shape of STDP curve

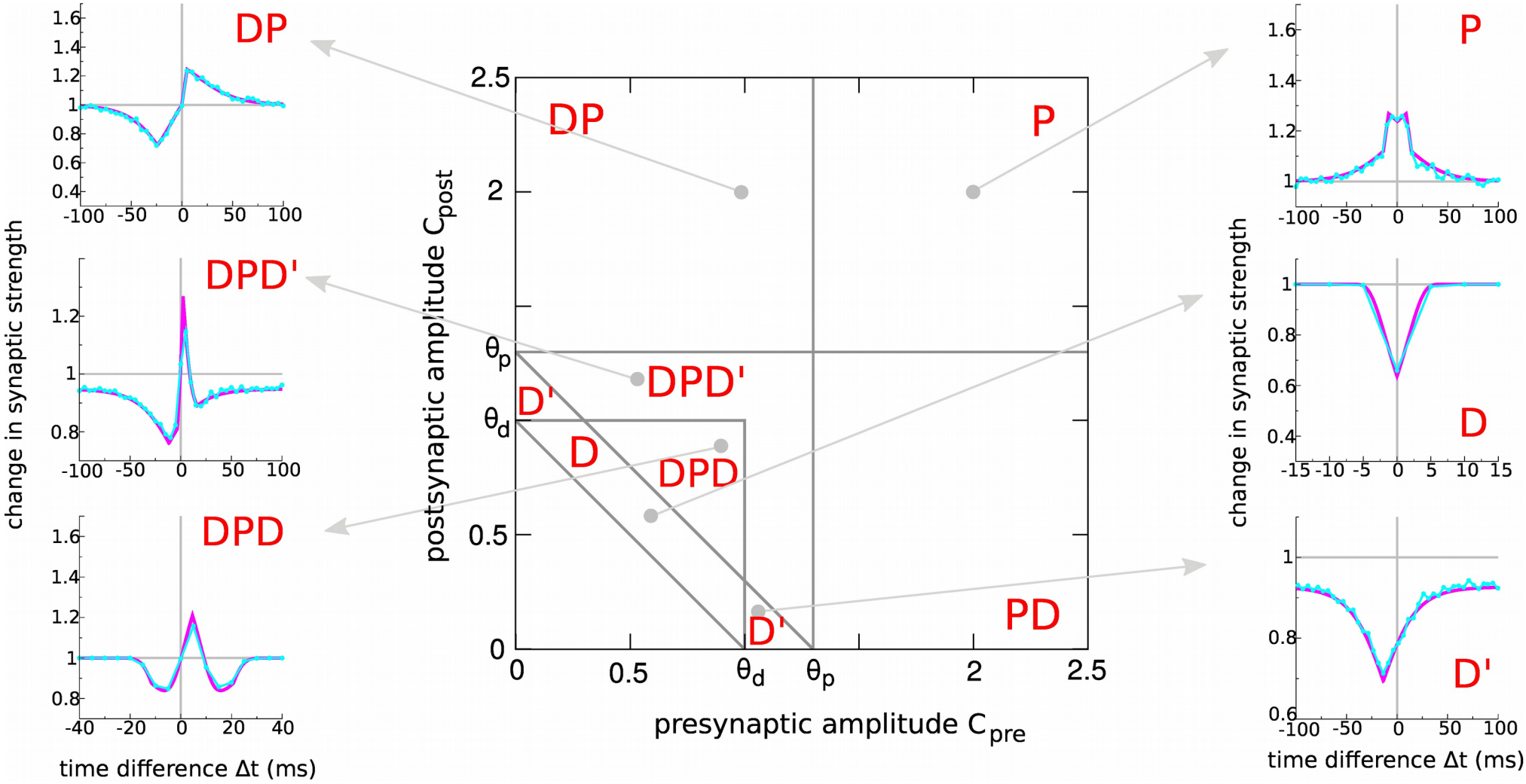


Calcium amplitudes determine shape of STDP curve



Diversity of STDP curves : spike-pair stimulation

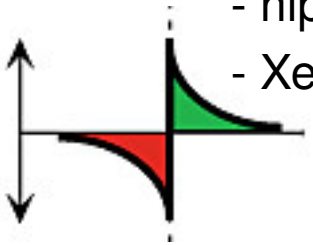
D ... depression , P ... potentiation



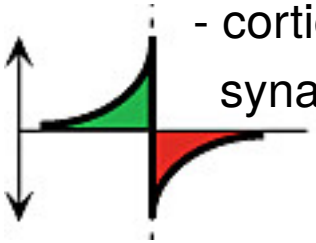
[Graupner & Brunel, *PNAS* 2012]

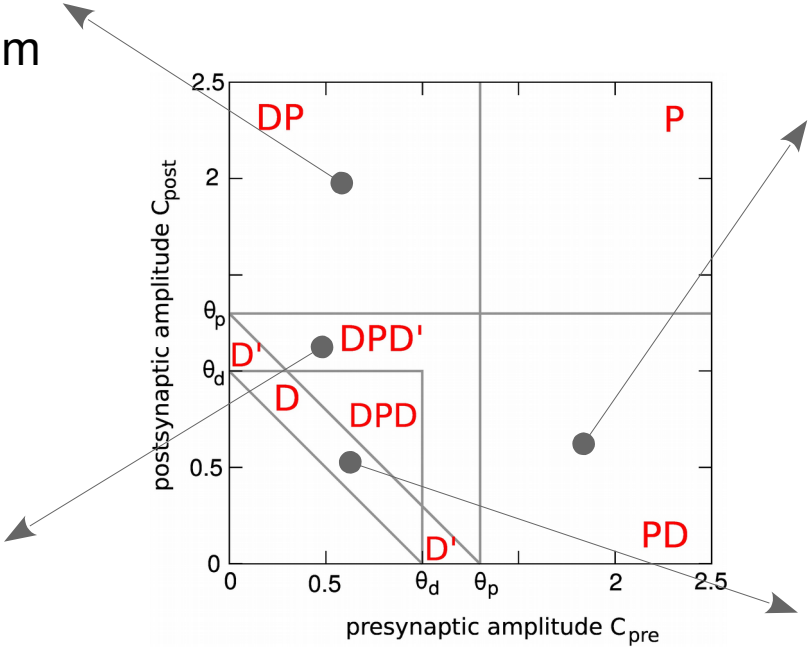
Diversity of STDP curves : experimental results

- neocortex-layer V
- hippocampus
- Xenopus tectum



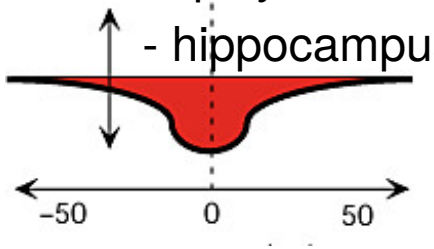
- ELL of electric fish
- corticostriatal synapse





- GABA-ergic neurons in hippocampal cultures

- neocortex-layer IV spiny stellates
- hippocampus



Outline

1. STDP : introduction and history

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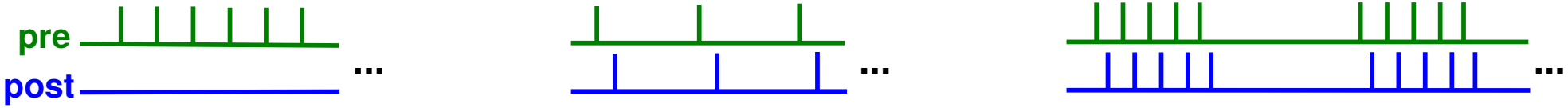
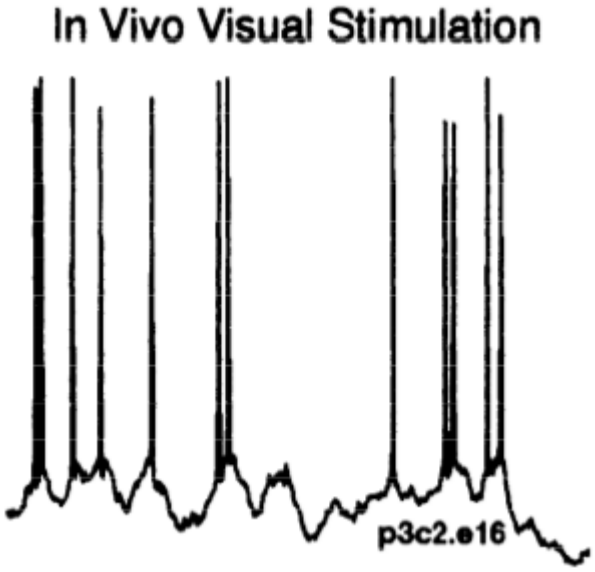
5. STDP *in vivo* ?

Firing patterns : Realistic firing is highly irregular

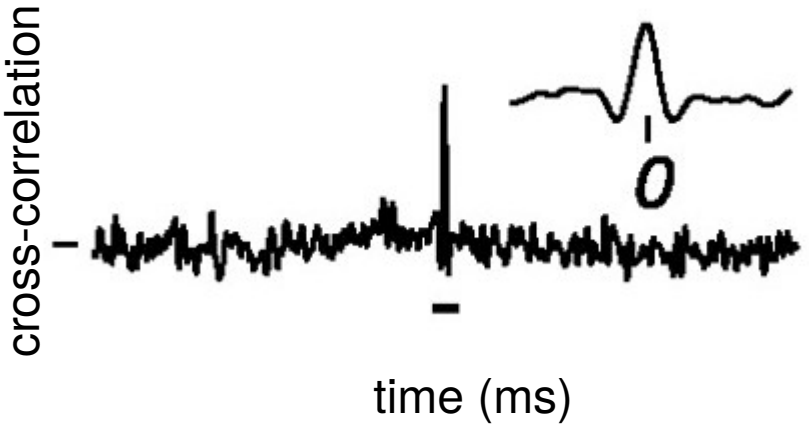
- stimulation protocols used to induce plasticity



- *in vivo* firing patterns

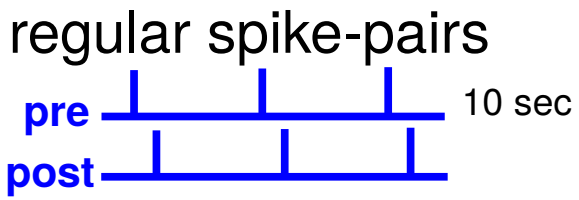


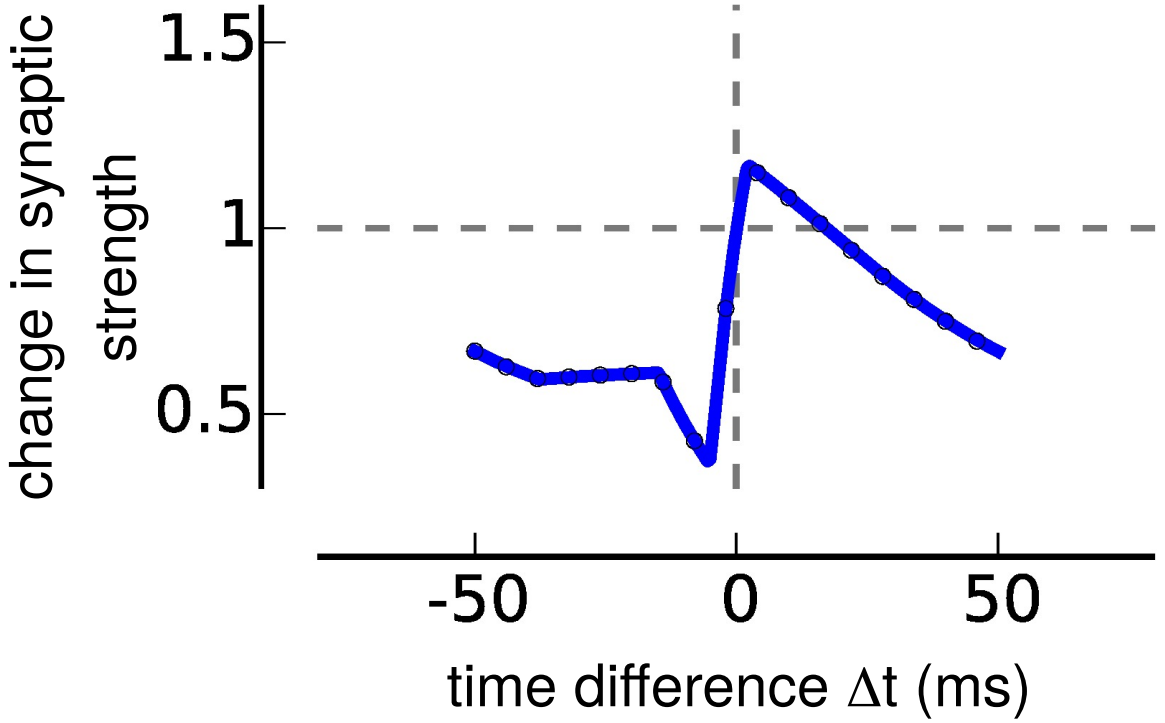
[Holt *et al.,* 1996]



[Kohn and Smith, 2005]

Regular vs. irregular spike-pairs

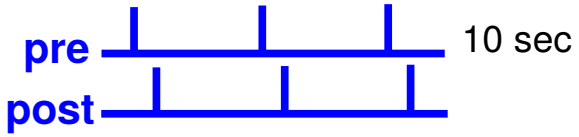




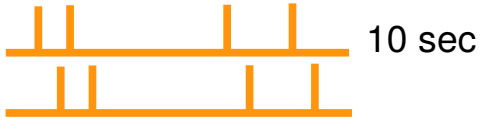
$\nu_{pre} = \nu_{post} = 10\ \mathrm{Hz}$

Regular vs. irregular spike-pairs

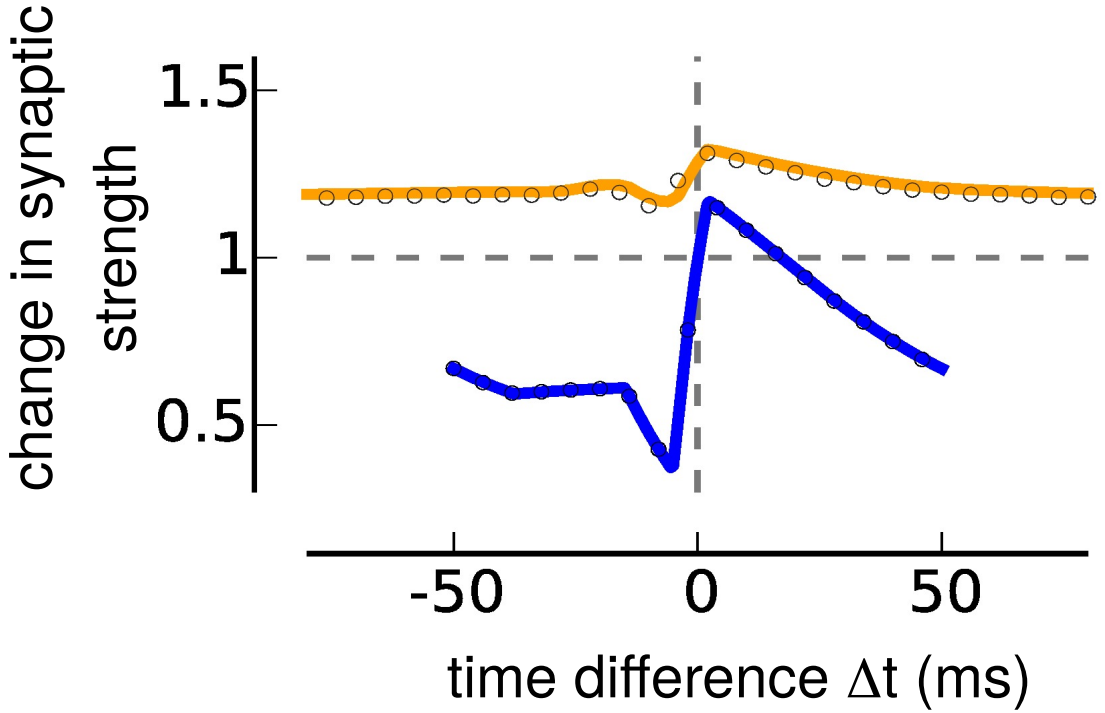
regular spike-pairs



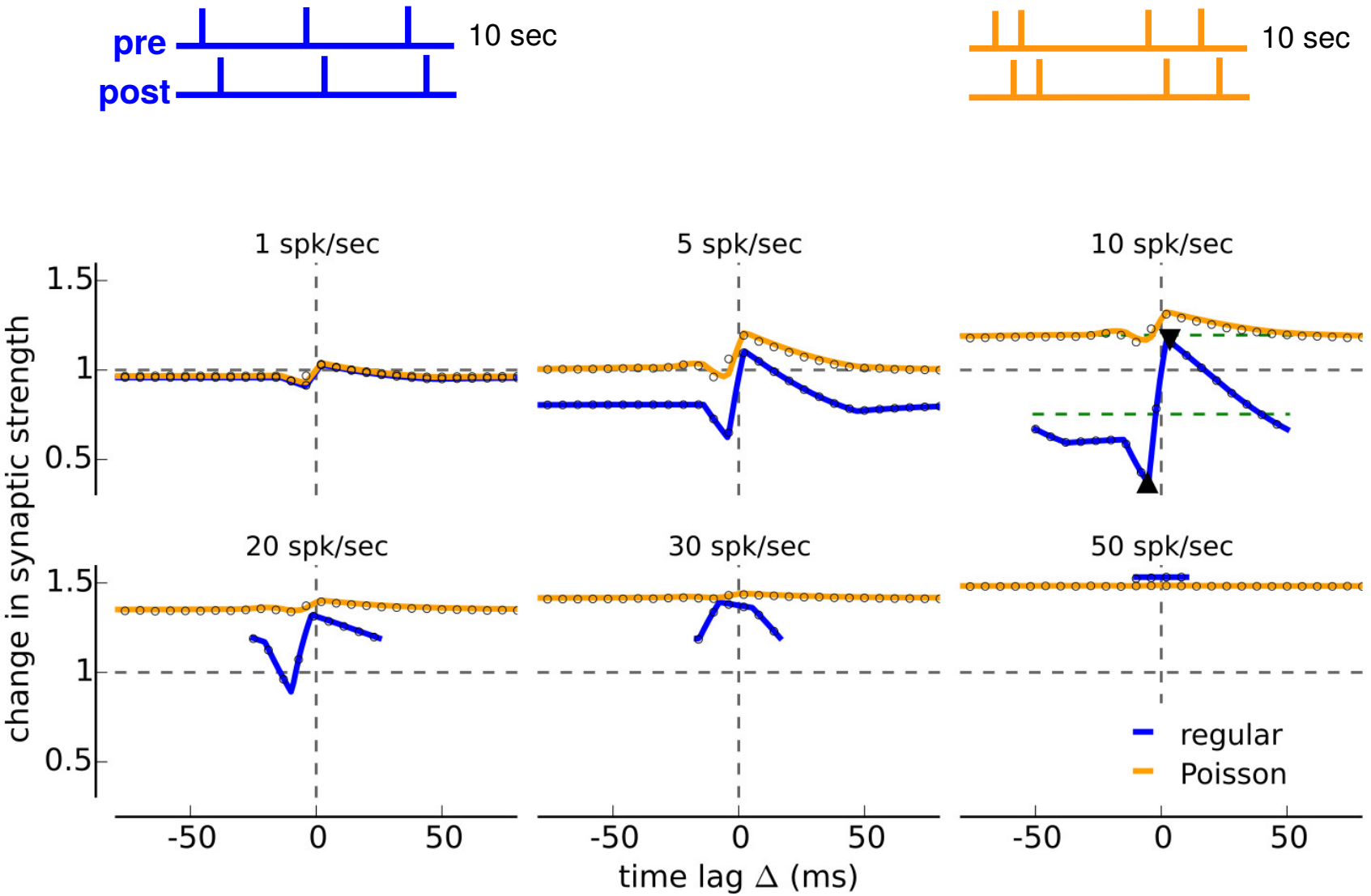
Poisson distributed spike-pairs



$\nu_{pre} = \nu_{post} = 10\,\text{spk/sec}$



Irregular spike-pairs flatten STDP curve

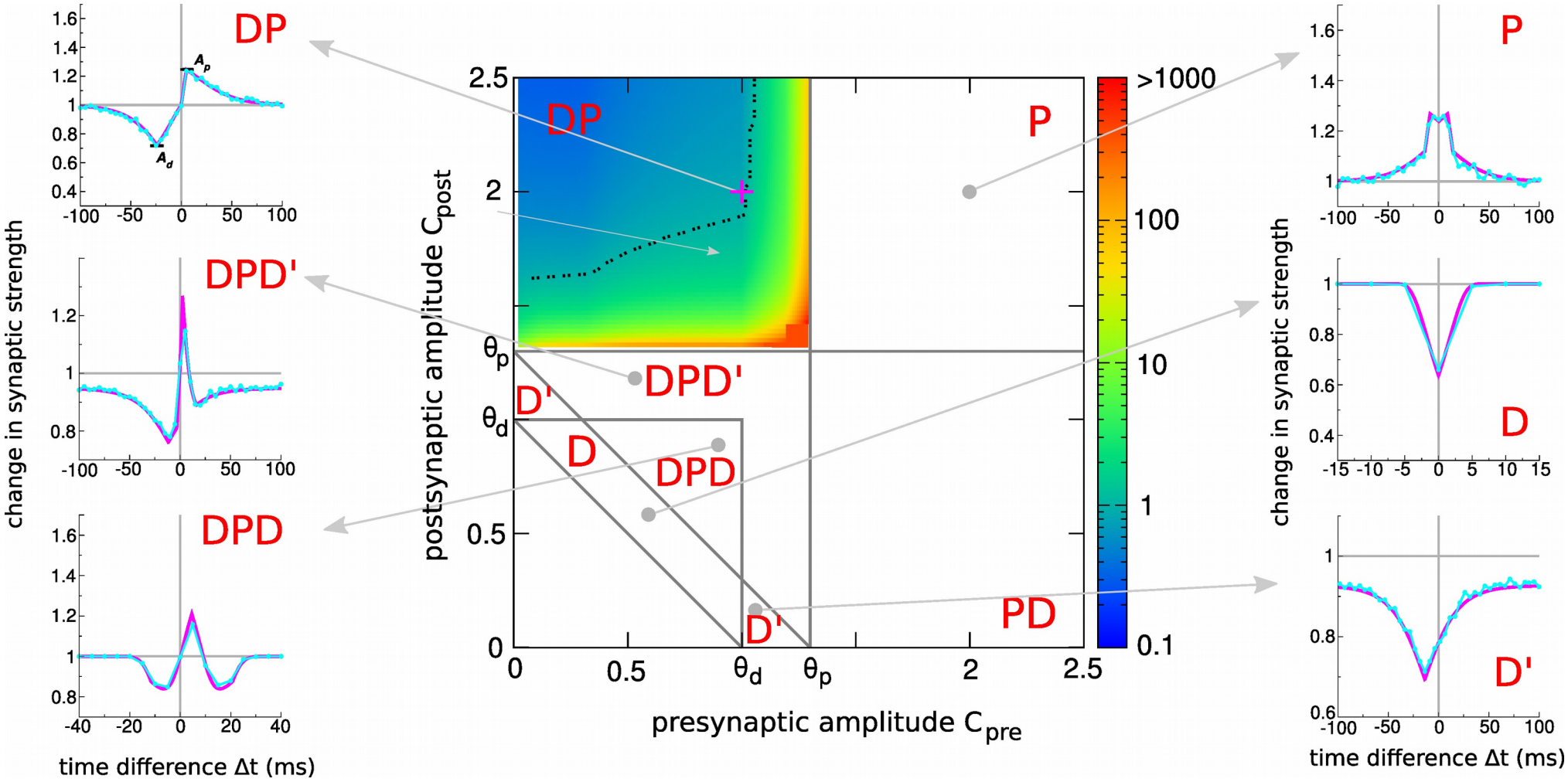


[Graupner, Wallisch & Ostojic. *J Neurosci* in press]

Conclusions

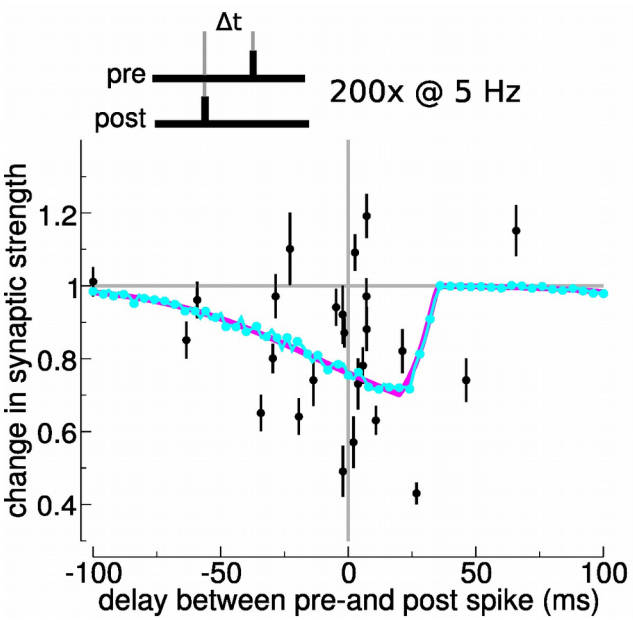
- STDP : temporally asymmetric form of synaptic plasticity induced by tight temporal correlations between the spikes of pre- and postsynaptic neurons
- induction: coincident pre- and postsynaptic activity lead to calcium influx through NMDA receptors, triggering intracellular signaling cascades
- biophysical model resolve various aspects of the synaptic machinery involved in plasticity induction, most commonly the postsynaptic calcium dynamics
- the role of STDP for learning in the living animal remains elusive

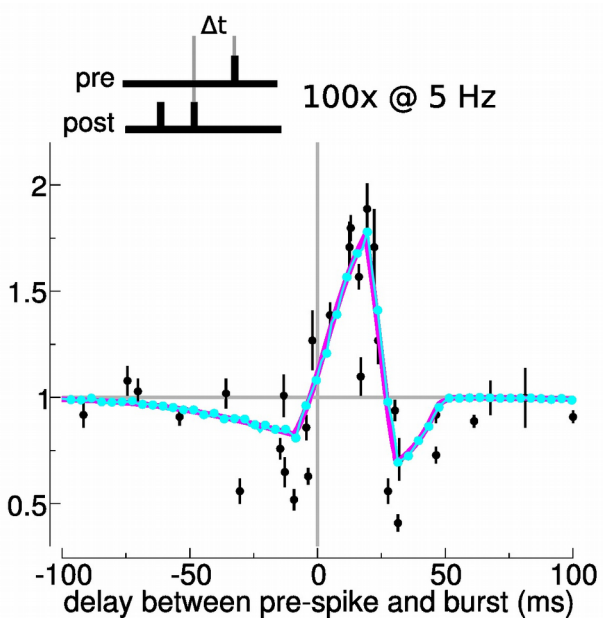
Diversity of STDP curves : spike-pair stimulation

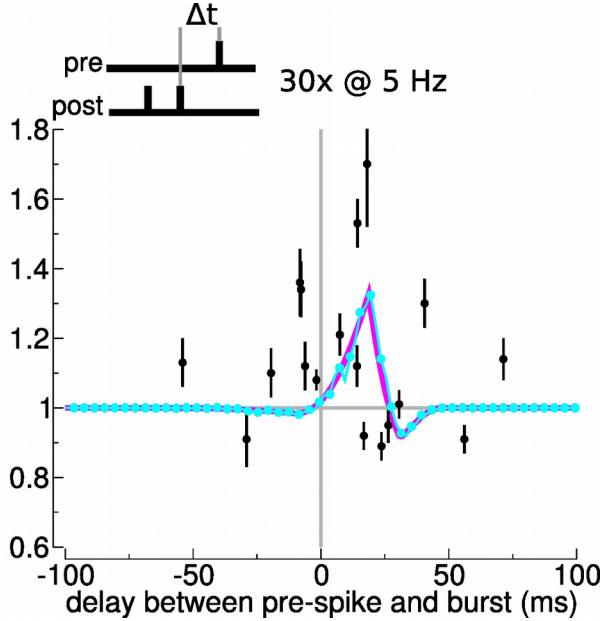


[Graupner & Brunel, 2012]

Malleability of hippocampal STDP explained by Ca^{2+}

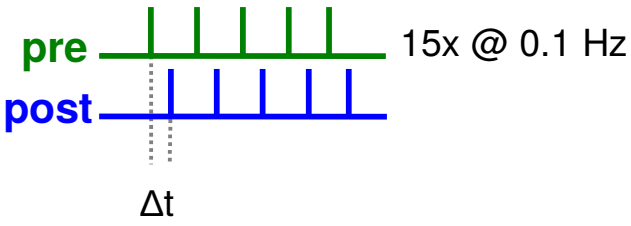


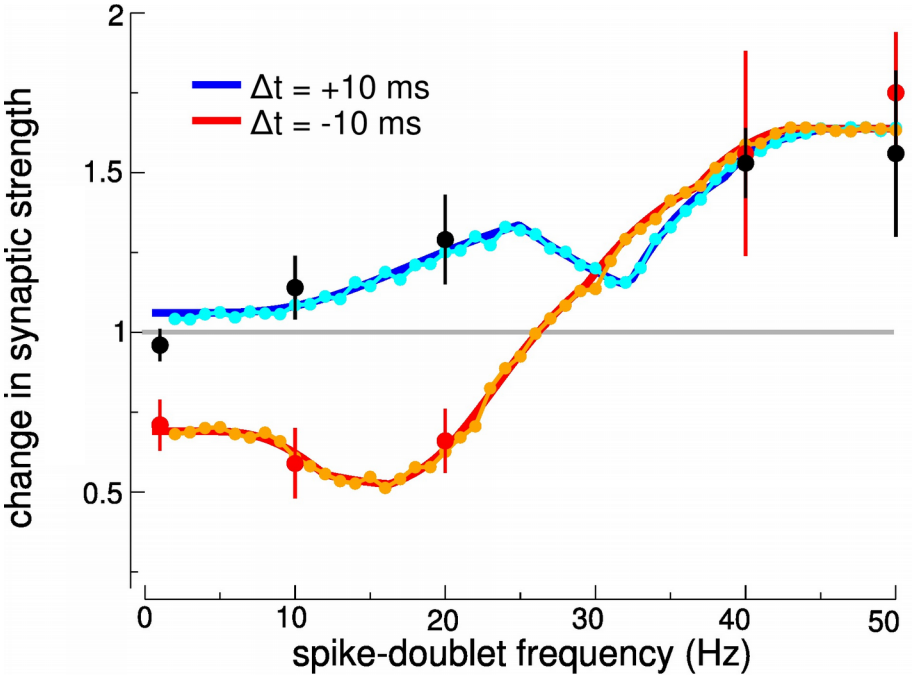


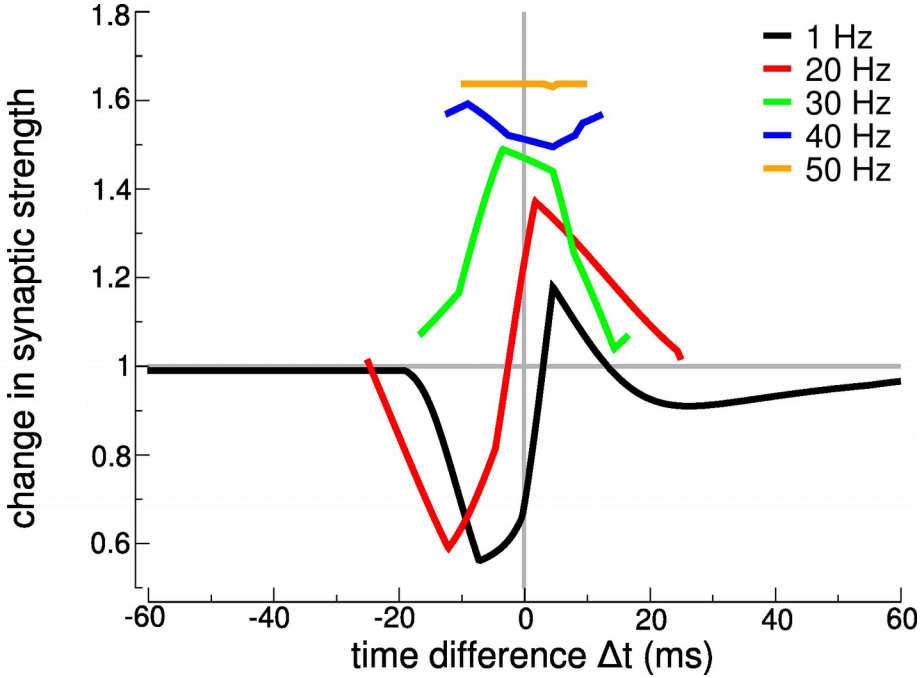


[Wittenberg & Wang, 2006]

Firing rate dependence in cortical slices

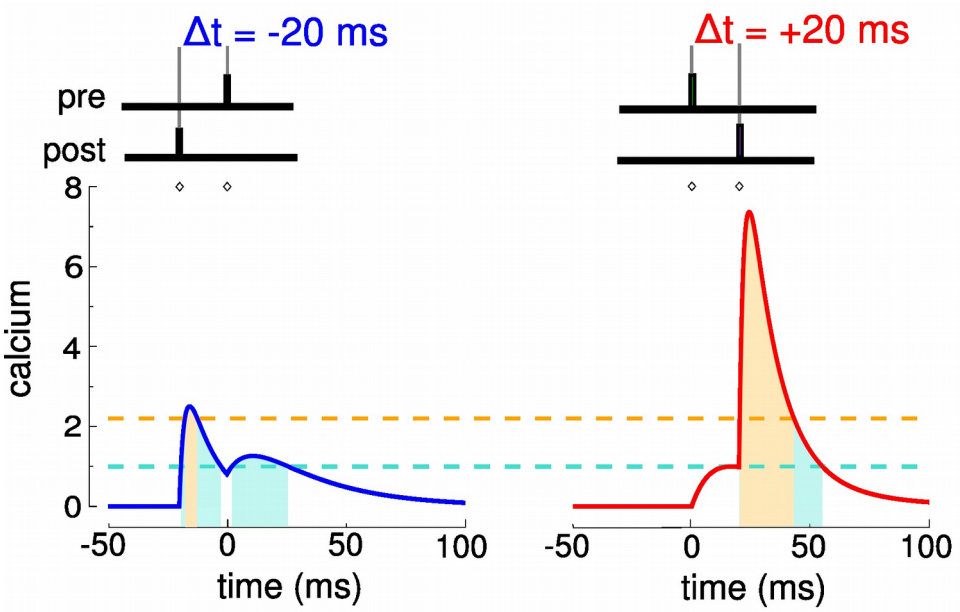


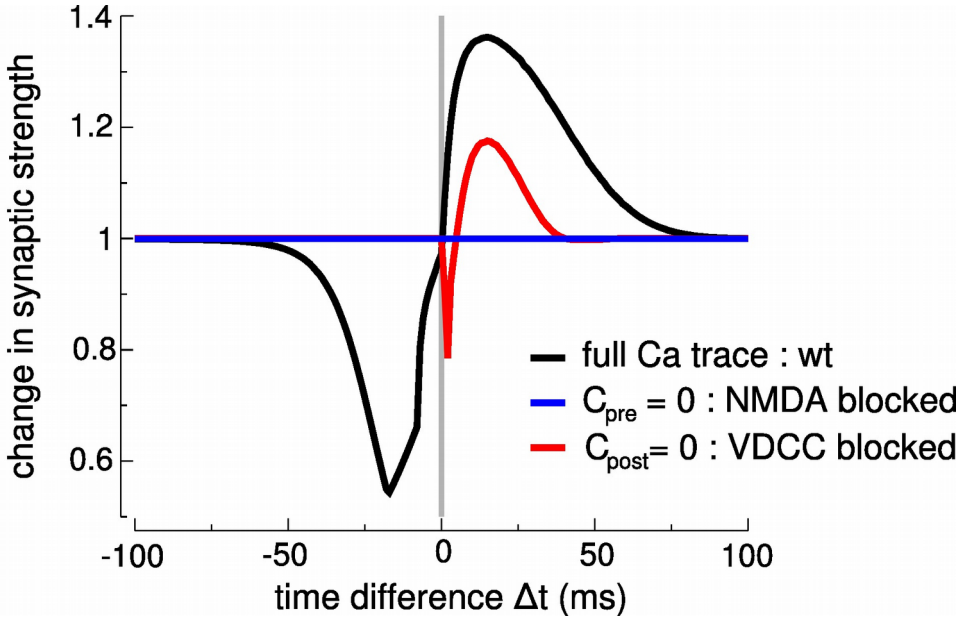




[Sjöström *et al.*, 2001]

Pharmacological manipulations explained by Ca^{2+}

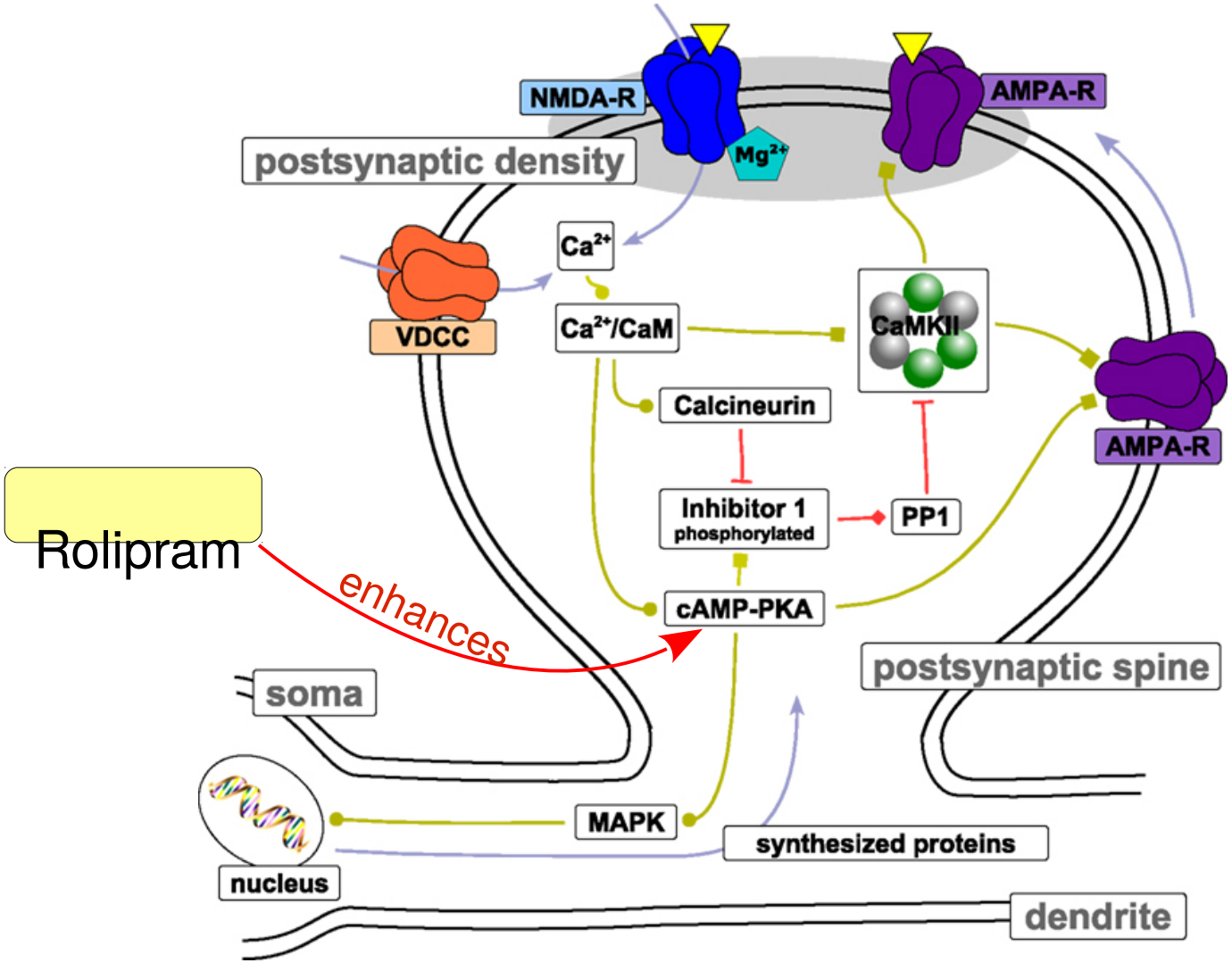




[Bi & Poo, 1998; Nevian & Sakmann, 2006]

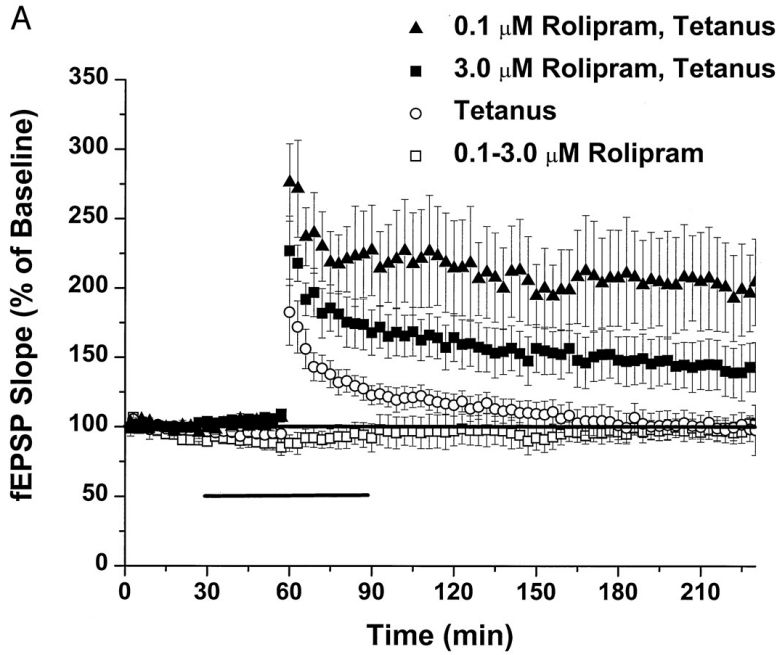
- nonlinear, finite rise time calcium transients necessary to reproduced pharmacological block experiments

Study the effect of nootropic drugs (memory enhancer)



Rolipram … selective phosphodiesterase-4 inhibitor

Study the effect of nootropic drugs

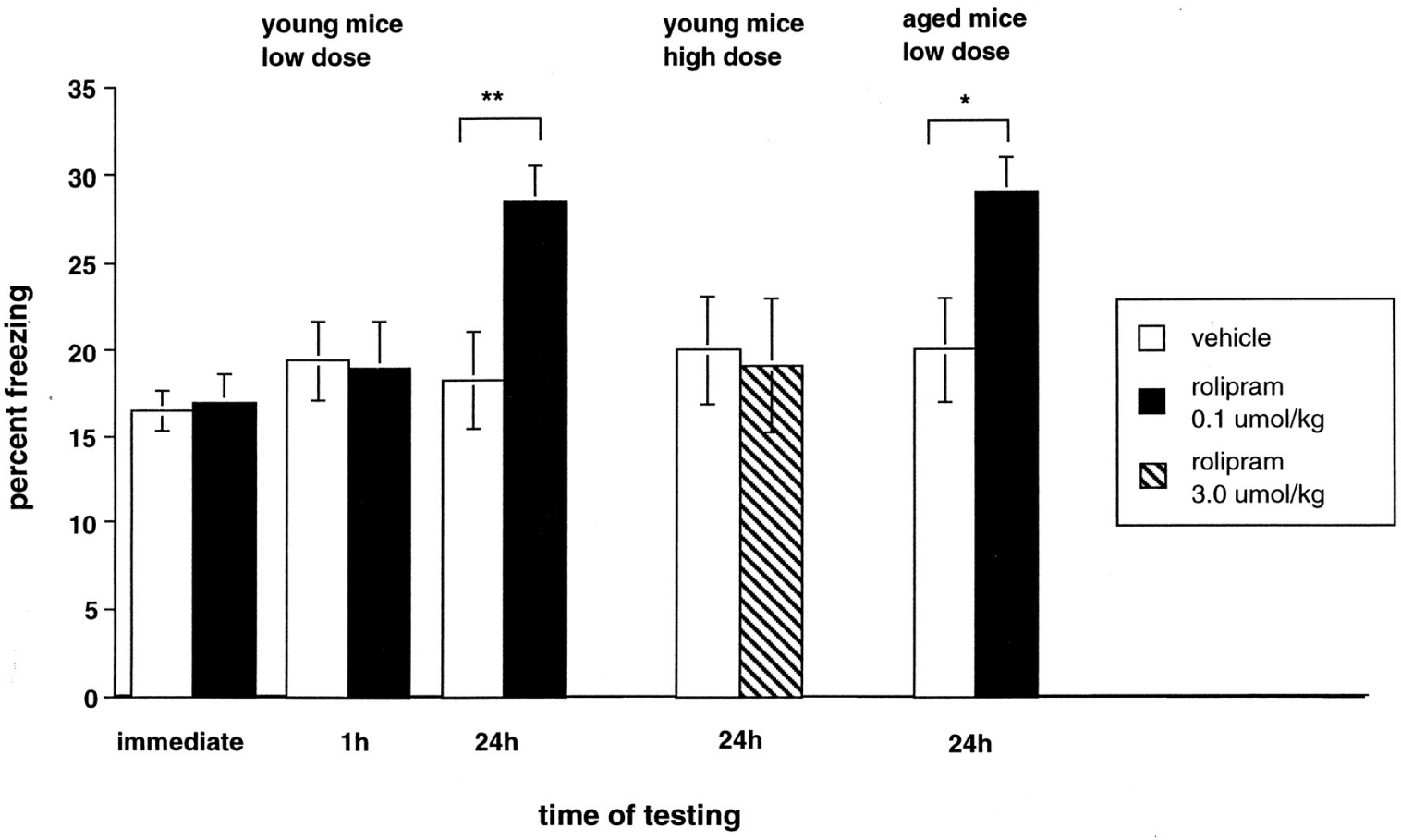


- boosting of cAMP *during* stimulation increases LTP

[Barath *et al.*, 1998]

Study the outcome of nootropic drugs

- Rolipram enhances memory



[Barath *et al.*, 1998]