

The background of the slide is a complex, abstract network of blue, glowing, fiber-like structures that resemble a neural network or a biological structure. The lines are of varying thickness and are interconnected, creating a dense, web-like pattern against a dark blue background.

LEARNING IN SPIKING NEURAL NETWORKS
FOR BIOINSPIRED MOTION CONTROL

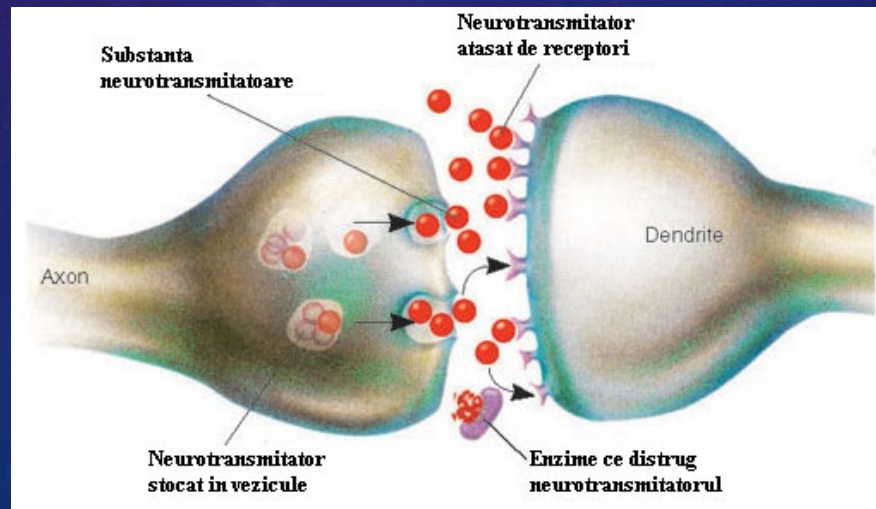
Mircea Hulea

Gheorghe Asachi Technical University of Iasi, Romania

SPIKE TIMING DEPENDENT PLASTICITY (STDP)

- Is related to learning in our brains
- Depends on the moments of neurons activation
- Is related to the degree of concurrence of PRE and POST neurons.
- Includes **long-term potentiation** and **long-term depression**.

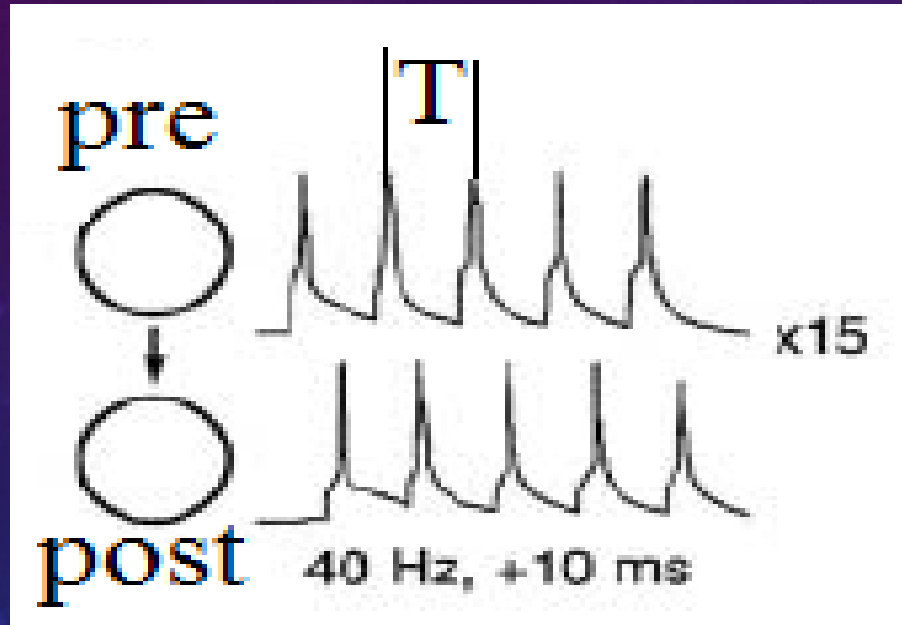
Presynaptic



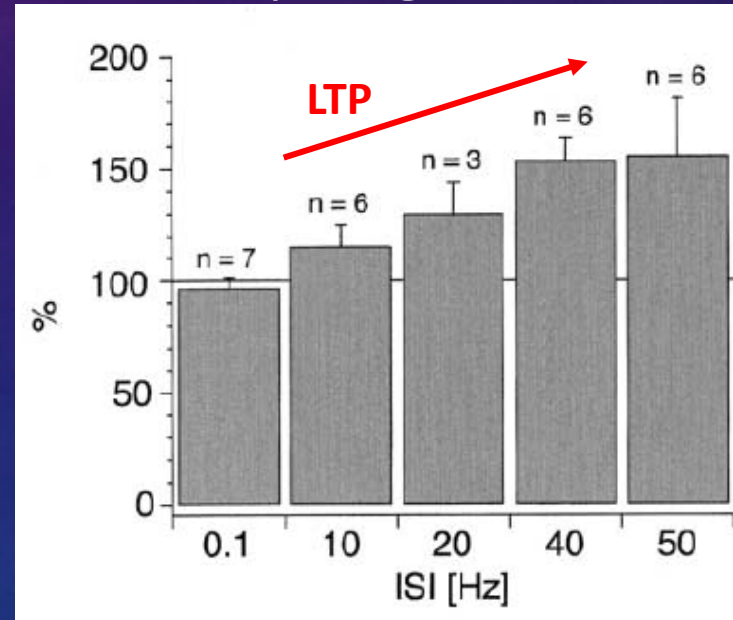
Postsynaptic

LTP AND LTD DEPENDS ON THE FREQUENCY OF ACTIVATION

Pair of stimuli *pre-post*



LTP increases with the spiking rate



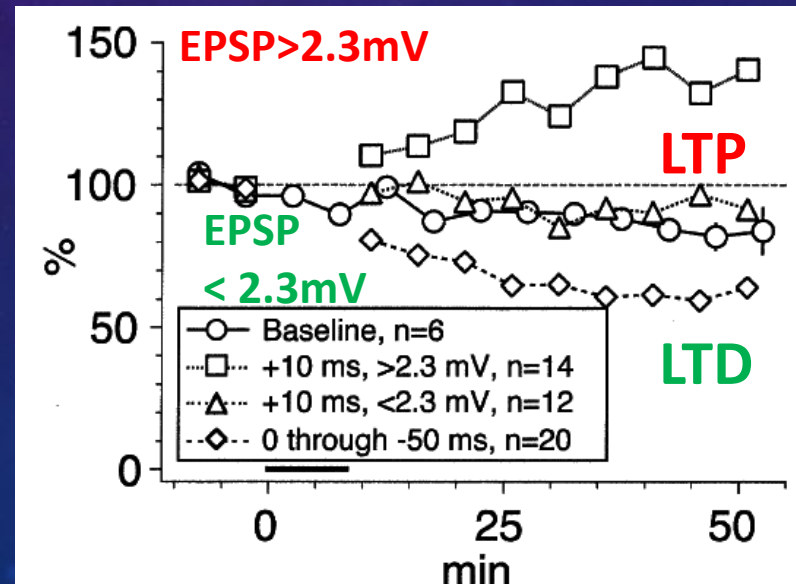
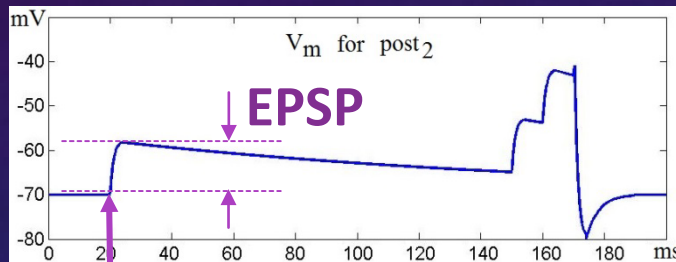
Sjostrom, et. al.

- For low stimulation frequency (0.1 Hz) LTP IS NOT observed for *pre-post*
- Value of LTP increases with the frequency of stimuli

POSTSYNAPTIC RESPONSE

2. LTP DEPENDS ON THE EPSP VALUE

- The synapses are potentiated by LTP if:
 - The value of EPSP > 2.3 mV even if the activation frequency is low
 - Pre-post succession
- Succession post-pre determines LTD



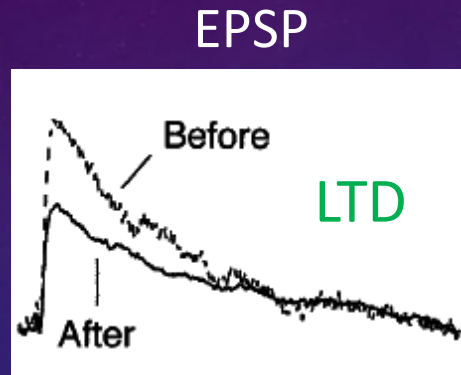
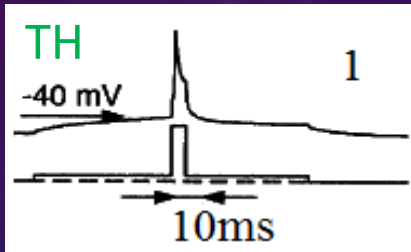
← If EPSP < 2.3 mV then no LTP
LTP is not produced at low values of EPSP

PRE activation

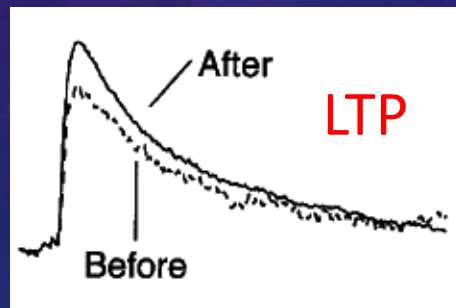
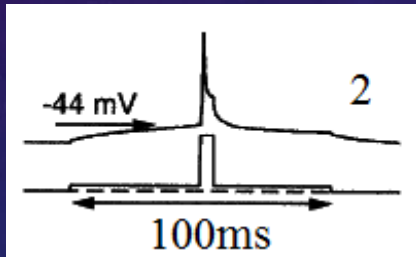
3. LTP-LTD DEPENDS ON INITIAL DEPOLARIZATION

Initial depolarization

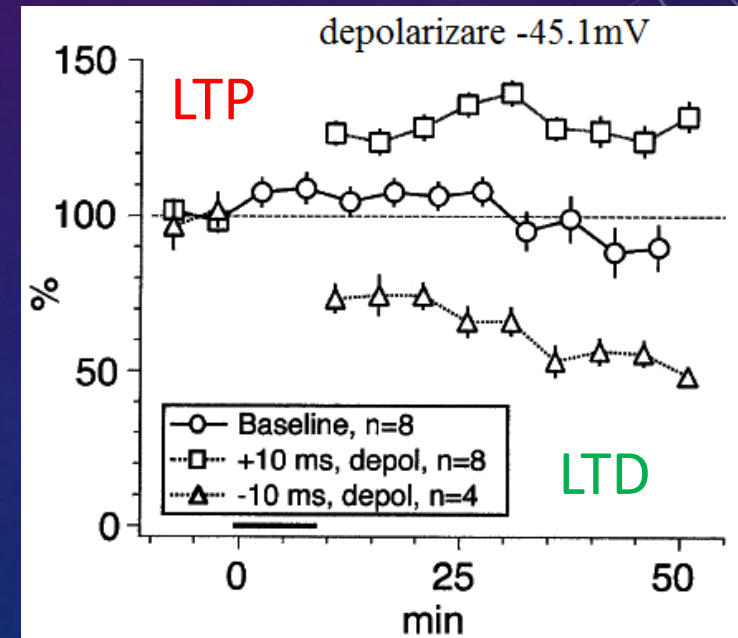
Just below the threshold (LTD)



Above the threshold (LTP)



Normal behavior for far depolarization



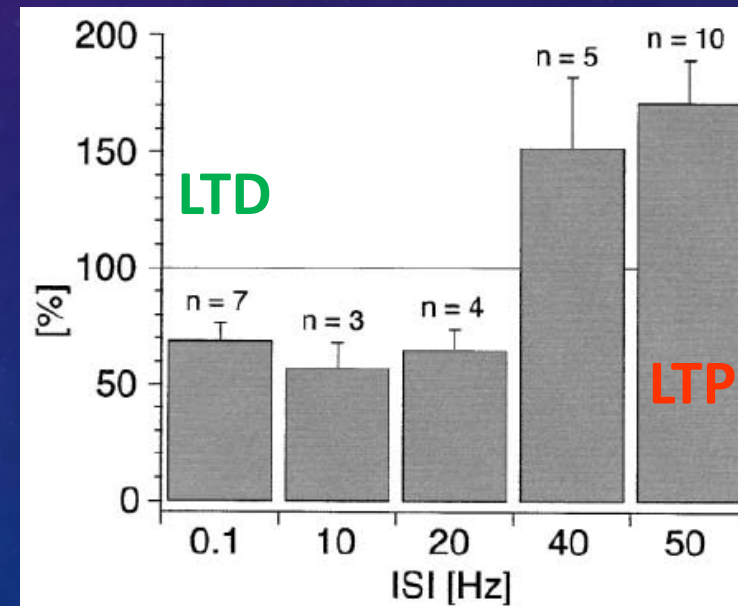
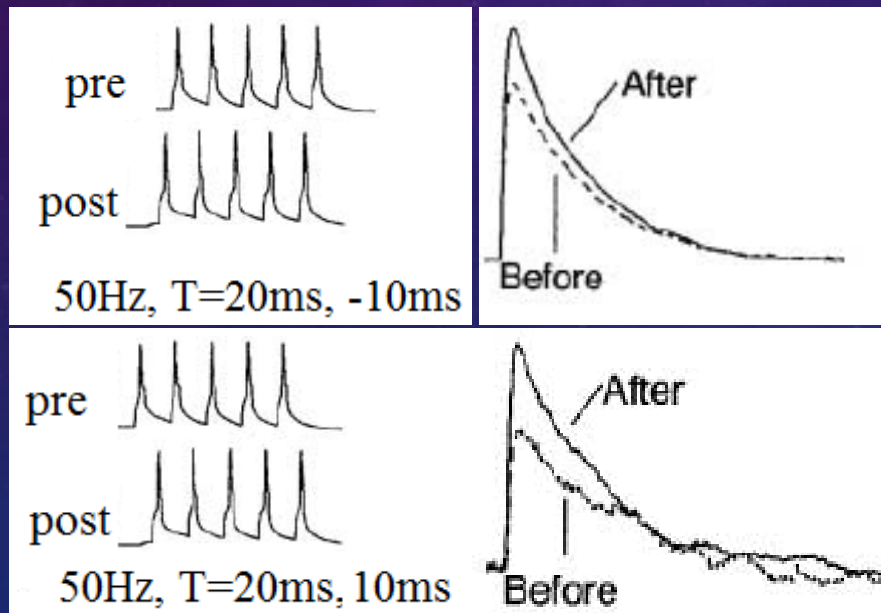
Sjostrom, et. al.

The initial depolarization should be far from the activation threshold to produce **LTP**. Otherwise **LTD** is produced.

LTP and LTD depends also on the residual depolarization (data not shown)

LTD (DEPRESSION) PRODUCES FOR LOW FREQUENCIES

- For frequency **below 20 Hz** the **LTD** is produced and it does not depend on the **frequency of activation**
- For frequency **above 40 Hz** only **LTP** is produced
- Note that **LTP** occurs when the frequency of *post-pre* is above 40 Hz

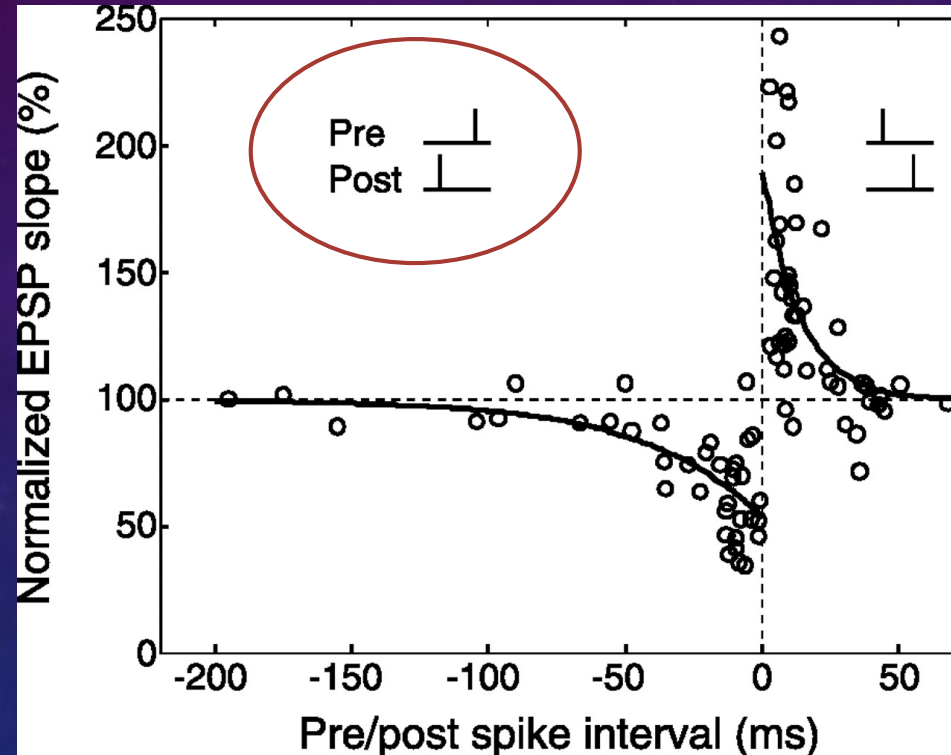


The synaptic plasticity **depends on the initial value** of the weight only for **LTP** and not for **LTD**.

THE REAL **STDP** WINDOWS

For high frequencies
Hebbian learning occurs
because of LTP

Post $\xrightarrow{\text{LTD}}$ pre



Pre $\xrightarrow{\text{LTP}}$ post

- The LTP window is shorter but **LTP** is more powerful than **LTD** for short time intervals
- **LTP** dominates the **LTD**

CONSTRUIREA MODELULUI PENTRU T-STDP

- For the synaptic transmission we consider the following **variables**:
 - r_1, r_2 – detectors of presynaptic events
 - Possible biological meaning:
 - **The quantity of NMDA** that stimulates the postsynaptic membrane
 - The number of NMDA receptors that are activated
 - Time constants for the variation of presynaptic detectors: τ_+, τ_x
 - o_1, o_2 – detectors for postsynaptic events
 - Biological plausibility
 - **The flux of Ca^{2+} ions** through the NMDA channels.
 - **The electric field** which is determined by the retro-propagation of the action potential.
 - Time constants for the variation of the postsynaptic detectors: τ_-, τ_y

VARIATION OF $R_{1,2}$ AND $O_{1,2}$ DURING NEURON ACTIVATION

Variation of detectors for neurons activation:

pre: $\text{if } t = t^{\text{pre}}, \text{ then } r_i \rightarrow r_i + 1$

post: $\text{if } t = t^{\text{post}} \text{ then } o_i \rightarrow o_i + 1$

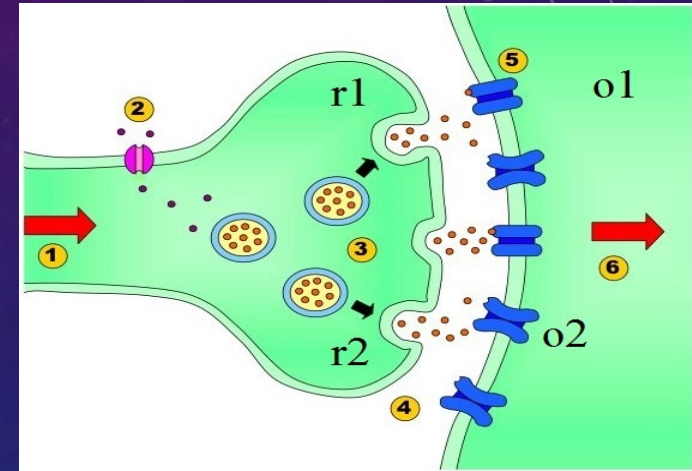
pre

post

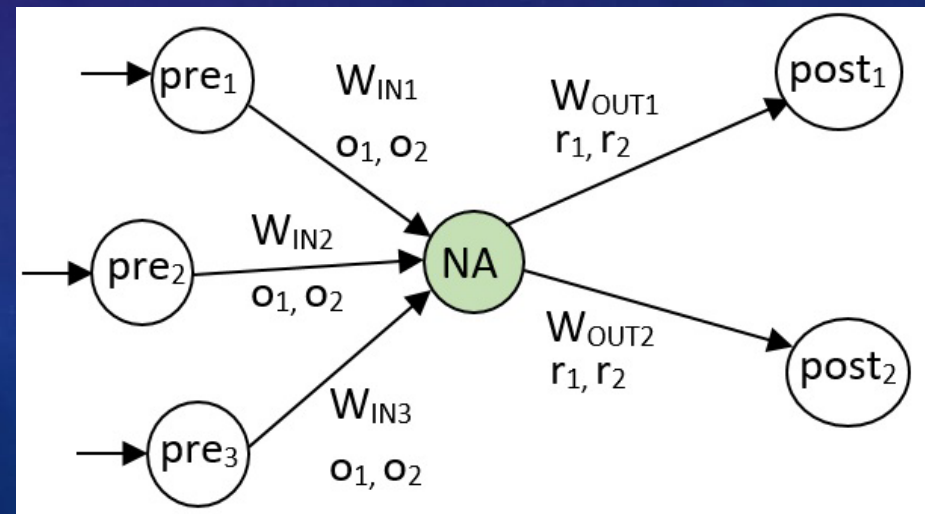
Inactivation

$$\frac{dr_i(t)}{dt} = -\frac{r_i(t)}{\tau_{+,x}}$$

$$\frac{do_i(t)}{dt} = -\frac{o_i(t)}{\tau_{-,y}}$$



When **NA** is activated r si o are activated for the adjacent synapses



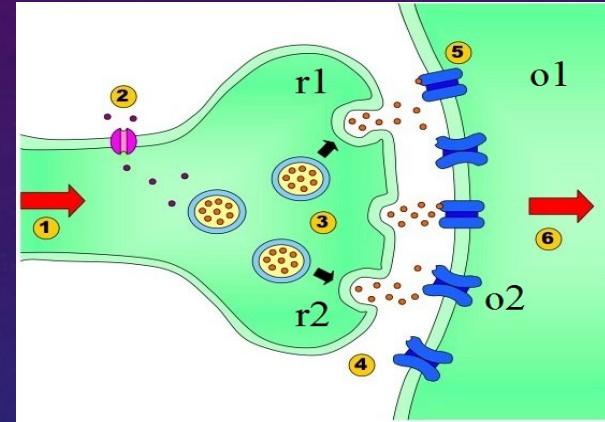
POTENTIATION AND DEPRESSION OF THE SYNAPSES

Variation of activation detectors:

$$\begin{array}{l} \text{pre:} \\ \text{post:} \end{array} \quad \frac{\text{if } t = t^{\text{pre}}, \text{ then } r_i \rightarrow r_i + 1}{\text{if } t = t^{\text{post}} \text{ then } o_i \rightarrow o_i + 1}$$

Inactivation

$$\begin{array}{cc} \text{pre} & \text{post} \\ \frac{dr_i(t)}{dt} = -\frac{r_i(t)}{\tau_{+,x}} & \frac{do_i(t)}{dt} = -\frac{o_i(t)}{\tau_{-,y}} \end{array}$$



- LTD: w **decreases** for **post** that are activated previously

$$w(t) \rightarrow w(t) - o_1(t)[A_2^- + A_3^- r_2(t - \epsilon)] \text{ if } t = t^{\text{pre}}$$

- Depends on o_1 – (principal **post**) si r_2 – prior to activation of **pre**

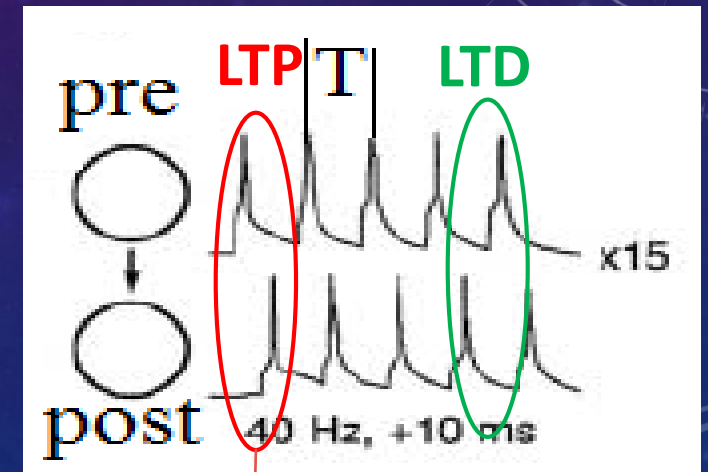
- LTP: w **increases** for **pre** activated previously

$$w(t) \rightarrow w(t) + r_1(t)[A_2^+ + A_3^+ o_2(t - \epsilon)] \text{ if } t = t^{\text{post}}$$

- Depends on r_1 – (principal **pre**) si o_2 – prior to **post** activation

QUESTION ?

- The effects of STP and LTP seems antagonistic
- This rises the question:
- STP and LTP compensates each other on long term?

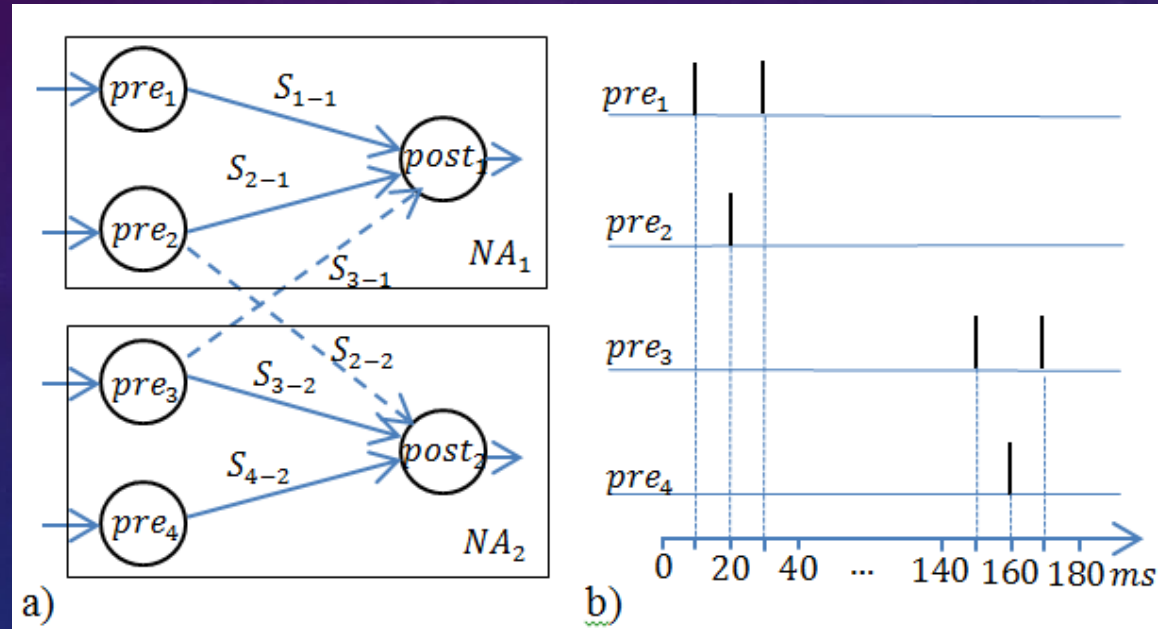


NEURAL NETWORK STRUCTURE

- Allow evaluation of the synaptic weights variation
- Main principle for the weights adjustment
 - Causality
 - potentiate the synapses that participates to the neuron action potential (activated before *post*)
 - Depress the synapses that did not participate to postsynaptic neuron activation (activates after *post*)
 - Triplets and quadruplets interaction
 - *Pre-post-pre* and *pre-post-post-pre* produces depression or leave the synaptic weights unchanged
 - *Post-pre-post* and *post-pre-pre-post* produces potentiation

TESTING THE NEURON MODEL

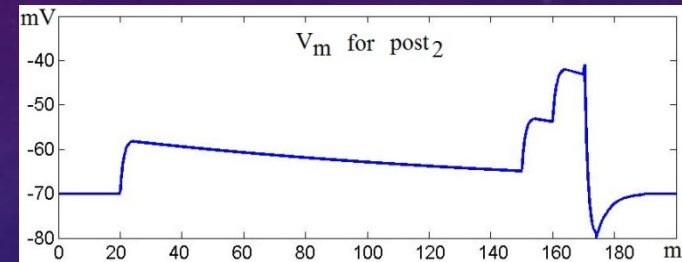
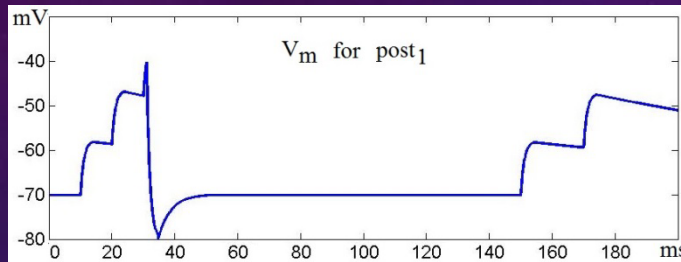
- Simple neural network structure
- Input pattern of stimuli



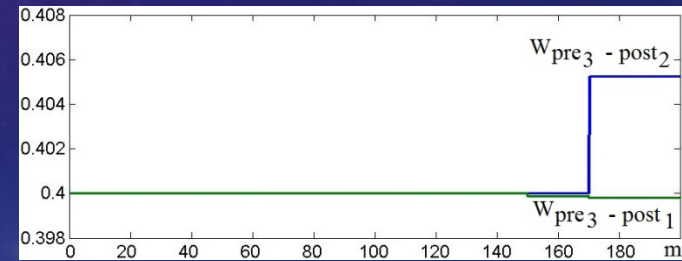
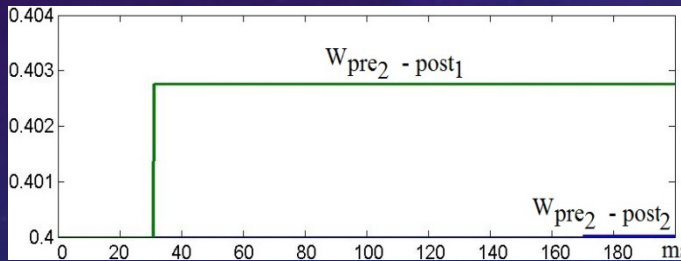
- The Neural Network is split in two areas

NETWORK RESPONSE FOR THE COMBINATION OF STIMULI

- Membrane potential variation



- Synaptic weights adjustment

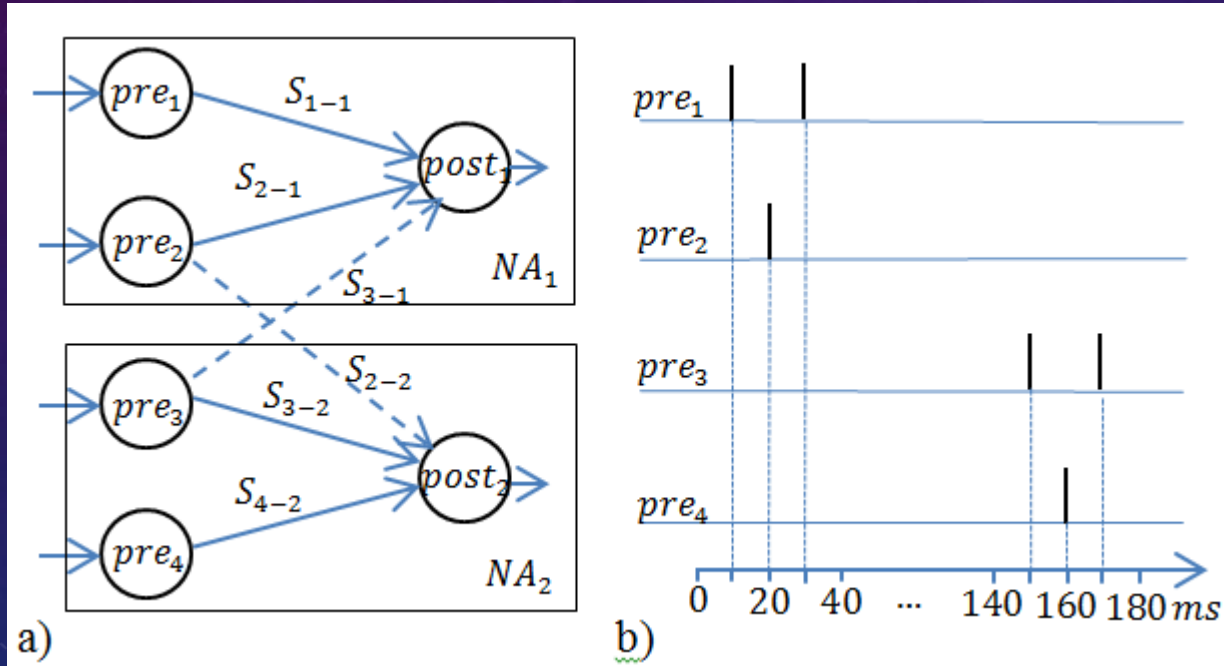


	Weights (initial value is 0.4)					
Patt. rep.	S1-1	S2-1	S2-2	S3-1	S3-2	S4-2
1	0.4047	0.4028	0.4000	0.3998	0.4053	0.4029
5	0.4246	0.4146	0.4000	0.3989	0.4265	0.4146
10	0.4273	0.4338	0.4001	0.3980	0.4529	0.4292

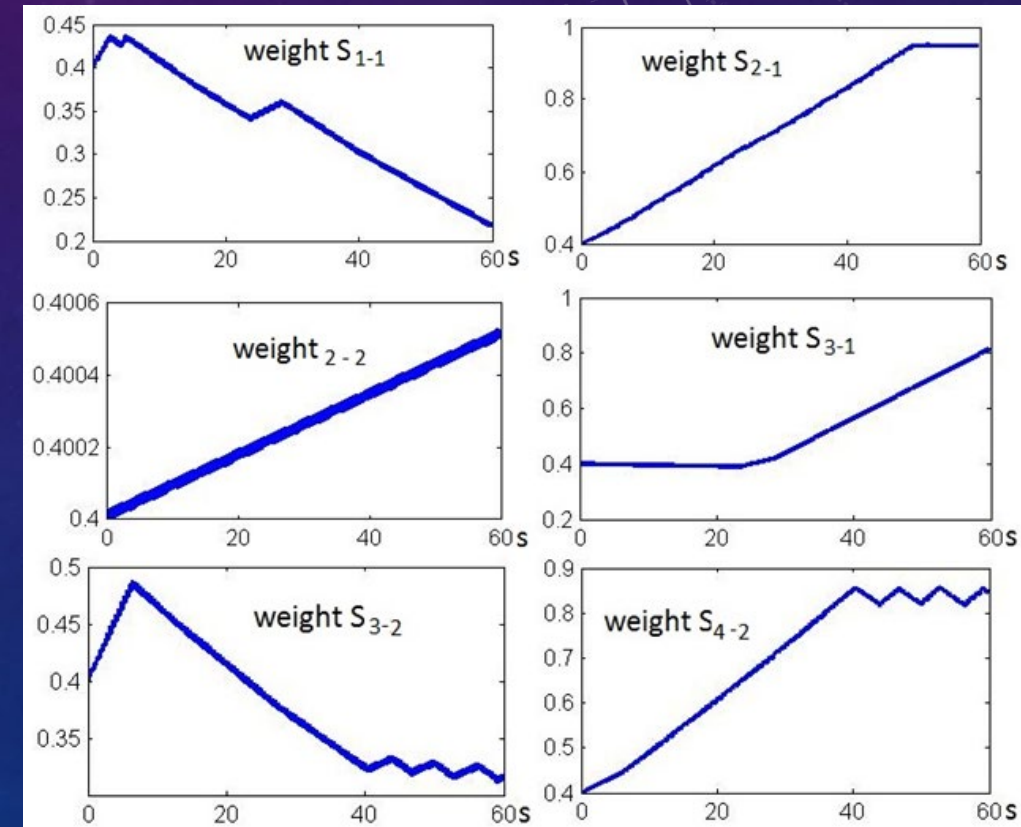
WEIGHTS EVOLUTION

- Weights variation during the first 60 seconds

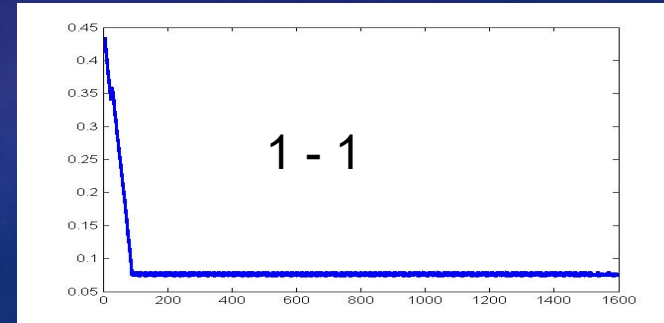
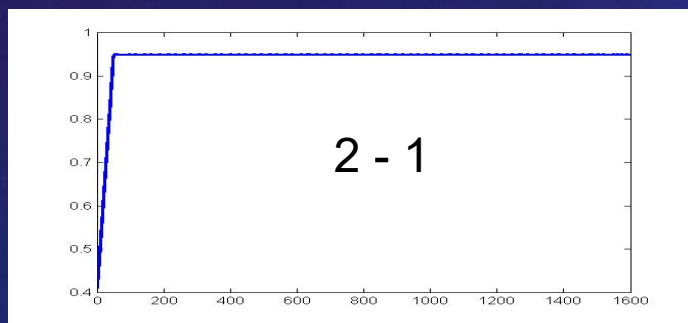
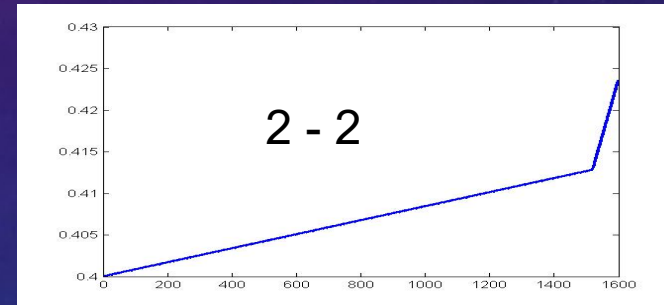
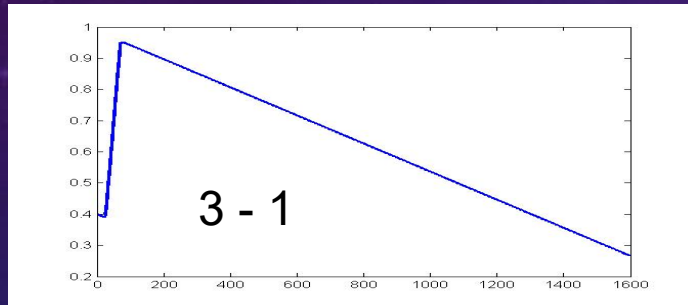
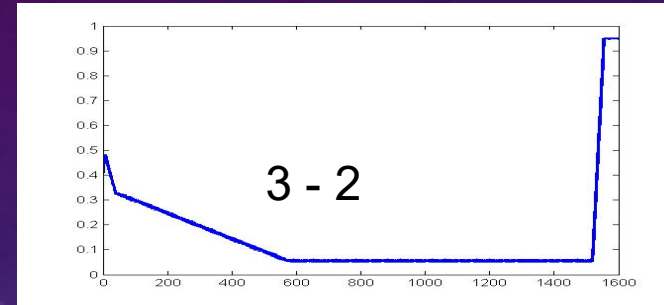
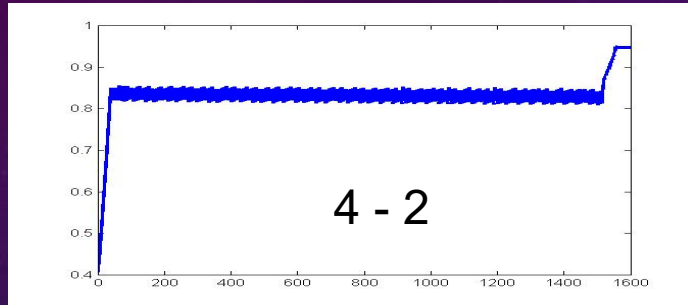
Spiking Neural Network



Weights variation



SYNAPTIC EVOLUTION



REMARKS

- Usually the weights tend to stabilize to minimum or to maximum value of the variation interval.
- In some conditions the weights tend to values that are different then the weight variation intervals limits
- Around these values the weights oscillate in small intervals
- In these cases the STP and STD effects compensate each other.
- **Future work:** These weights tend to input specific values ?

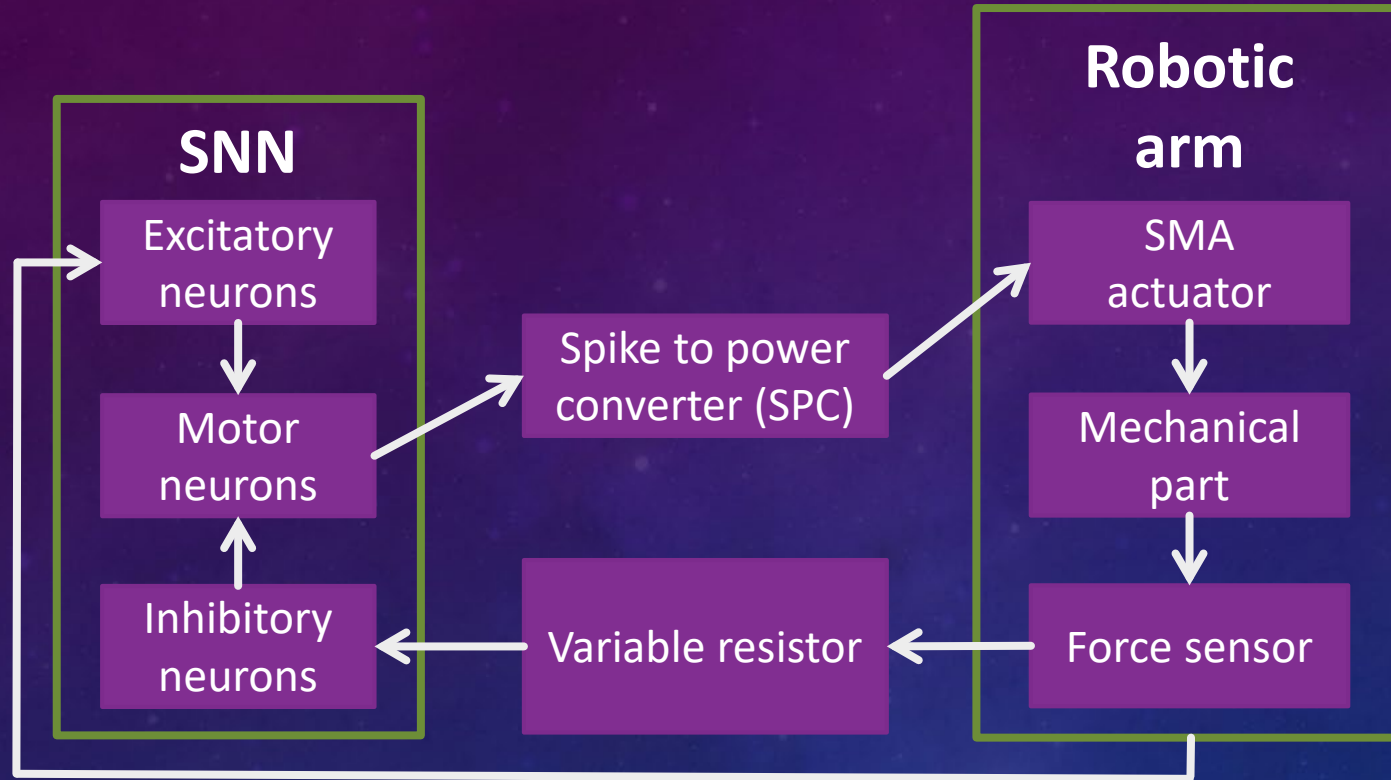
MUSCLE CONTROL

BIOLOGICAL BACKGROUND

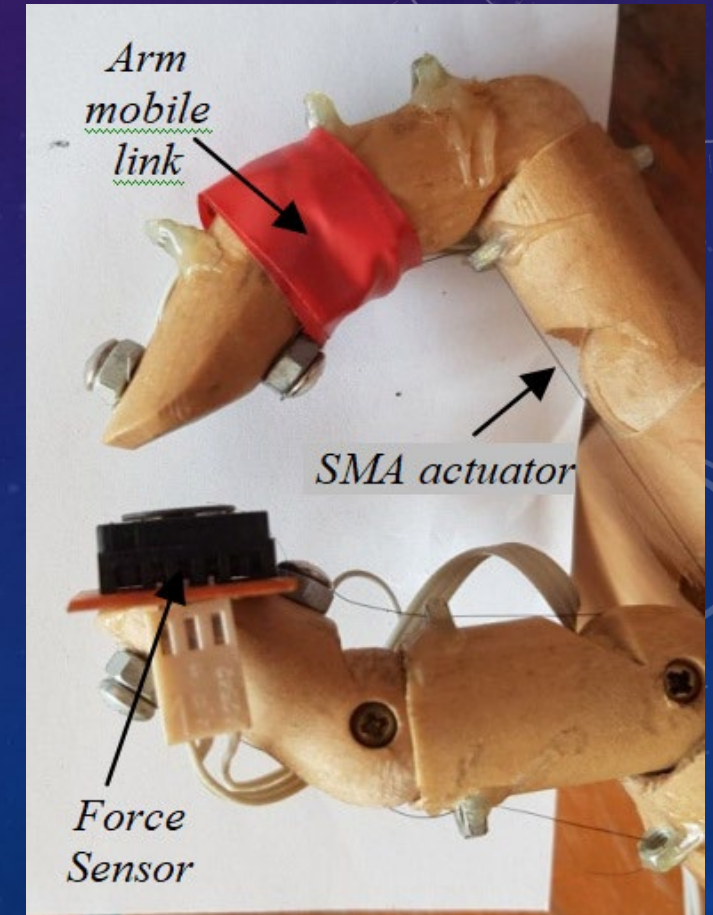
- Muscle control is one of the most important functions of the cerebral cortex
 - Provides the organisms with the ability to mechanically interact with the external environment
- Muscle control is bidirectional (**in biology**)
 - Muscles contraction is determined by the spiking frequency of the motor neurons
 - Neural network receives input related to elongation and contraction force from the spindles
- Limbs movements
 - Multiple muscles are controlled by the central pattern generators (CPG)



GENERAL CONCEPT



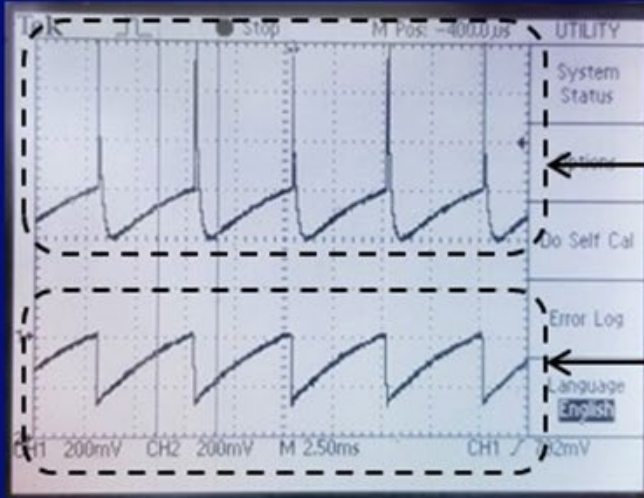
- Two opposing fingers that are actuated by SMA actuators



- **Robotic fingers:**
 - Flexed by SMA actuators
 - SMA contracts because of heating
 - Force sensor stops the finger motion

ELECTRONIC NEURON OPERATION

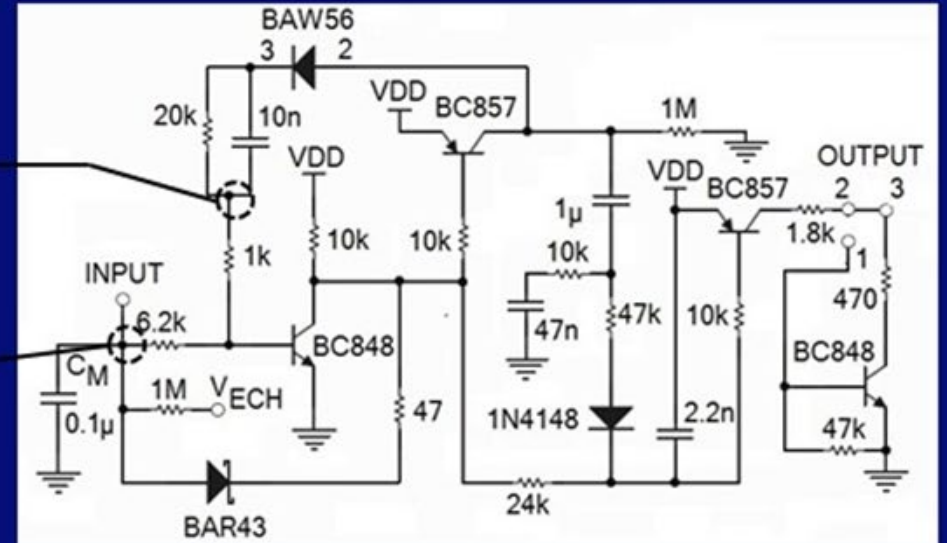
Spikes are the neuron activations



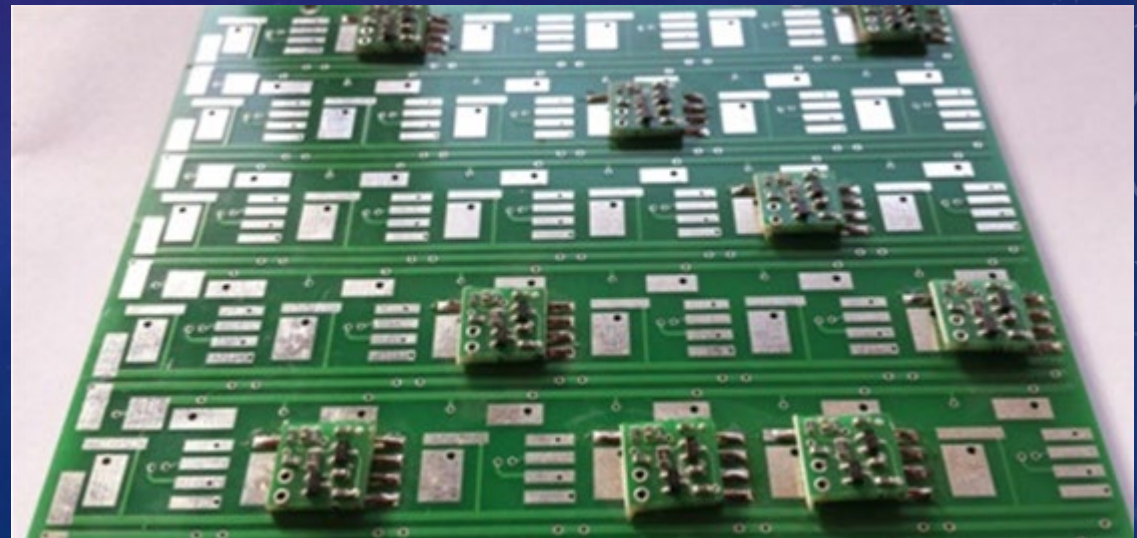
Bioinspired
signal

Monitored
signal
(neuron input)

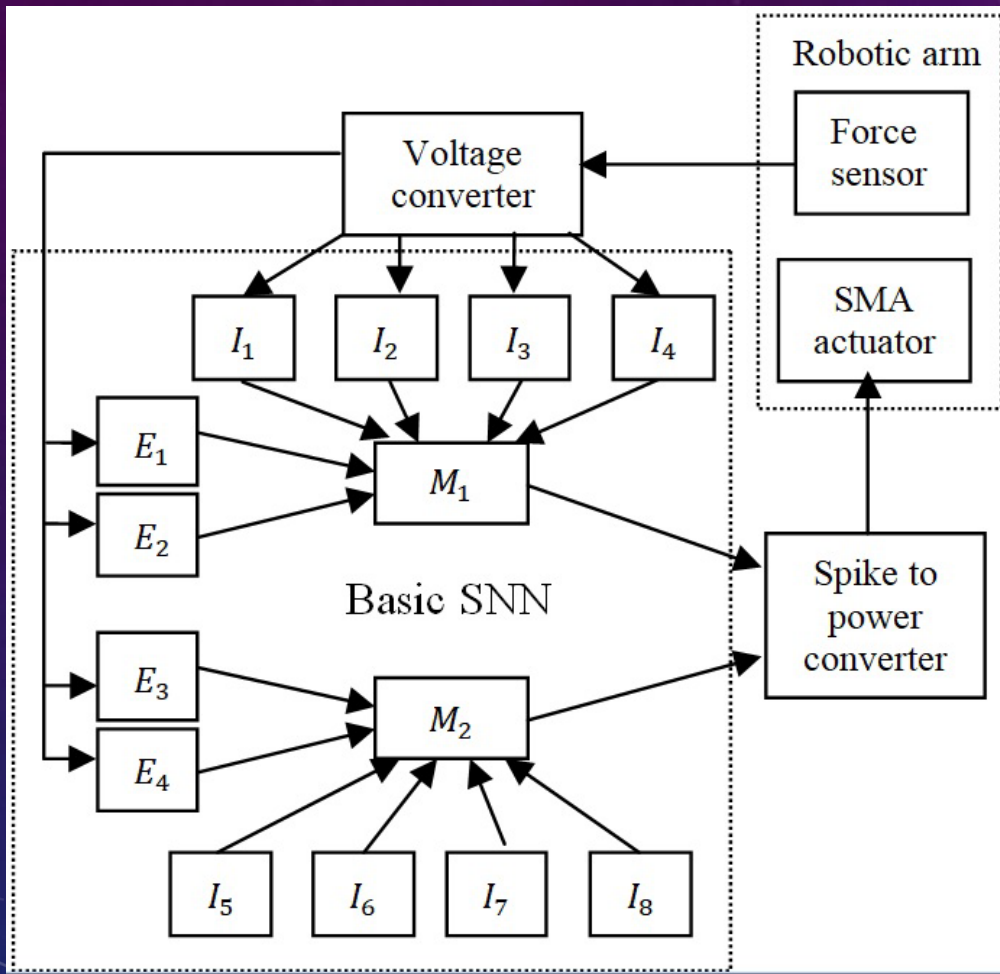
Electronic neuron schematic



- PCB implementation of unconnected neurons



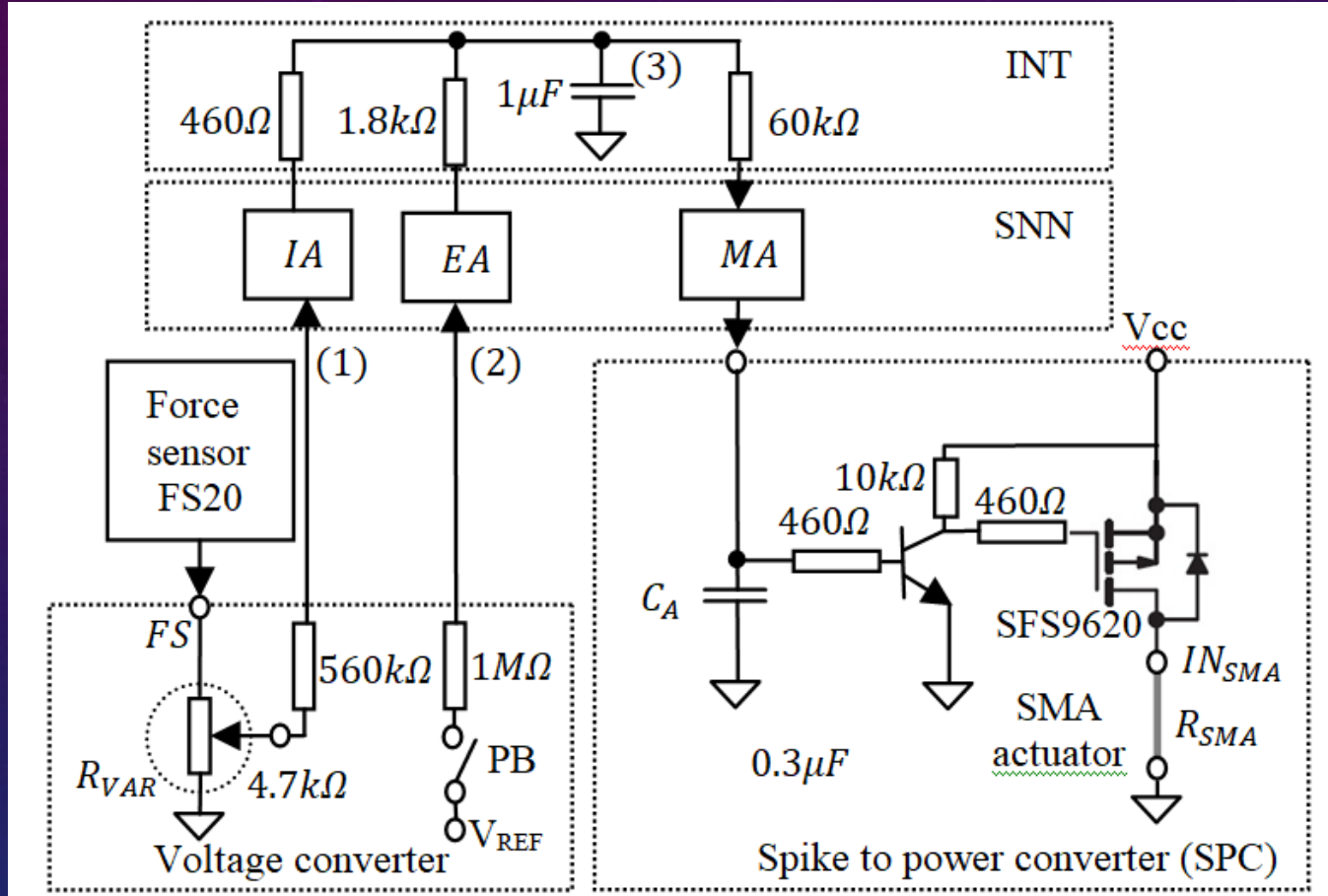
NEURAL NETWORK



Basic SNN includes:

- two motor neurons (M)
- 4 excitatory neurons (E)
- 8 inhibitory neurons (I)
- SMA actuator is driven by the SPC
- Integrated excitatory output of M
- Inhibitory neurons stimulated by FS
- SNN controls the contraction force

STRUCTURE OF THE BIOINSPIRED SYSTEM



ANALOGUE ELECTRONICS

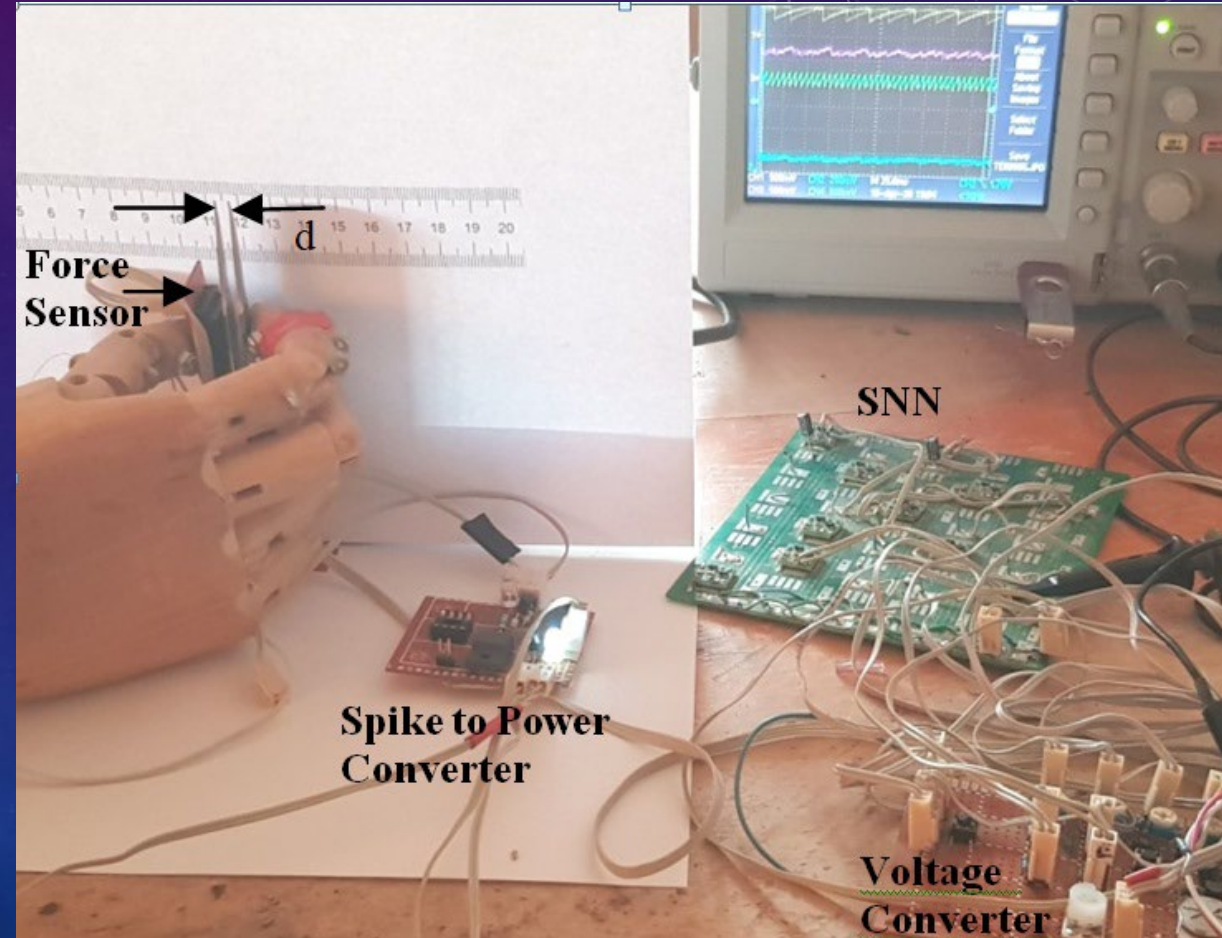
- Voltage converter
- Spike to power converter
- Integrator (INT)

- SMA converts current into force
- Force sensor (FS) converts force into voltage

PROTOTYPE OF BIOINSPIRED SYSTEM

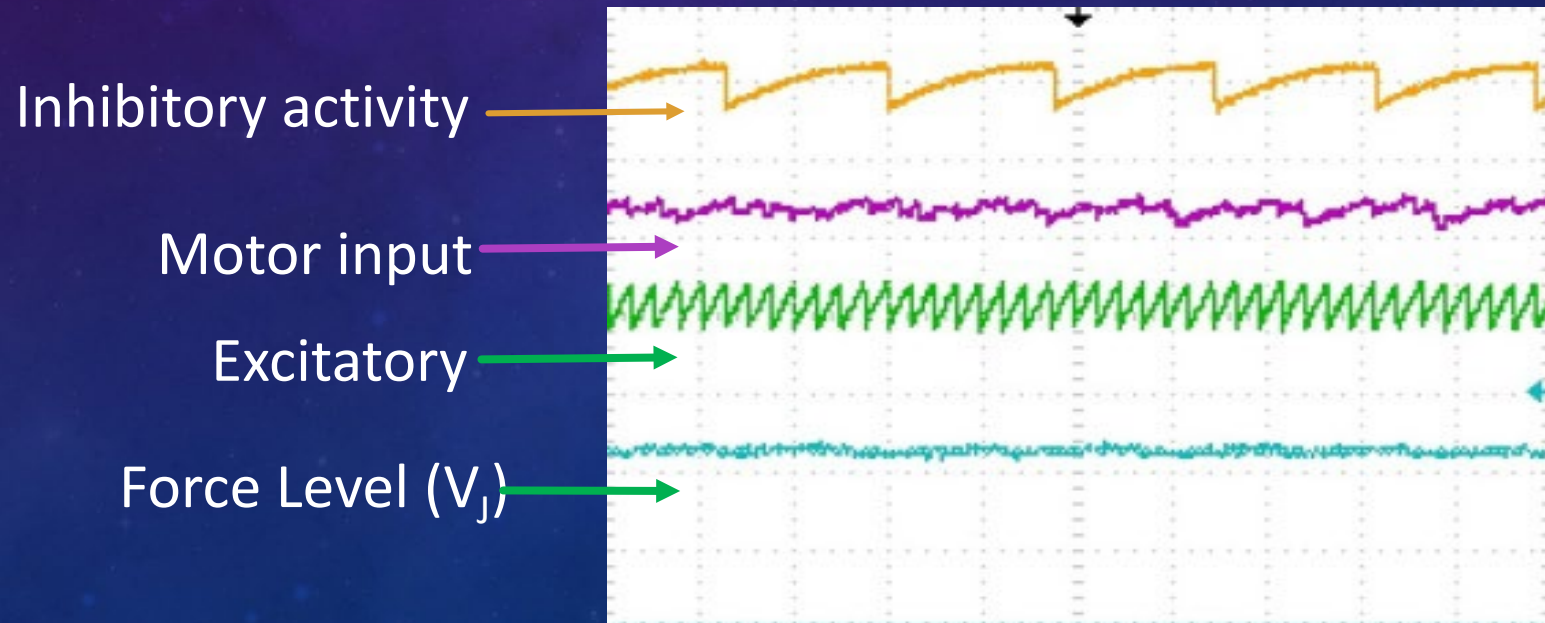
Experimental setup:

- Robotic hand holding a tweeters
- Distance between heads (d)
- Spiking neural network
- Auxiliary electronics
 - Spike to power converter
 - Voltage converter

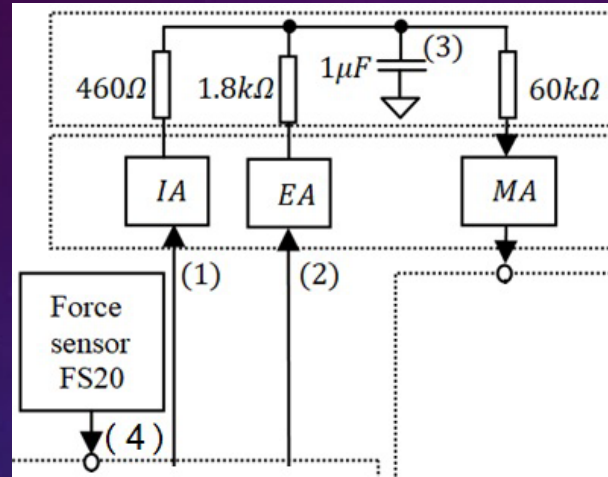
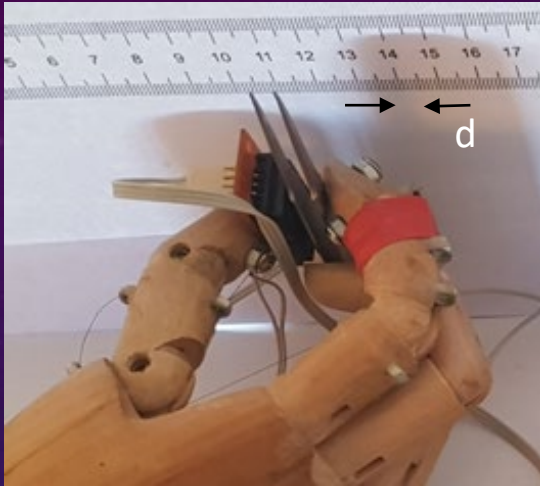


RESULTS

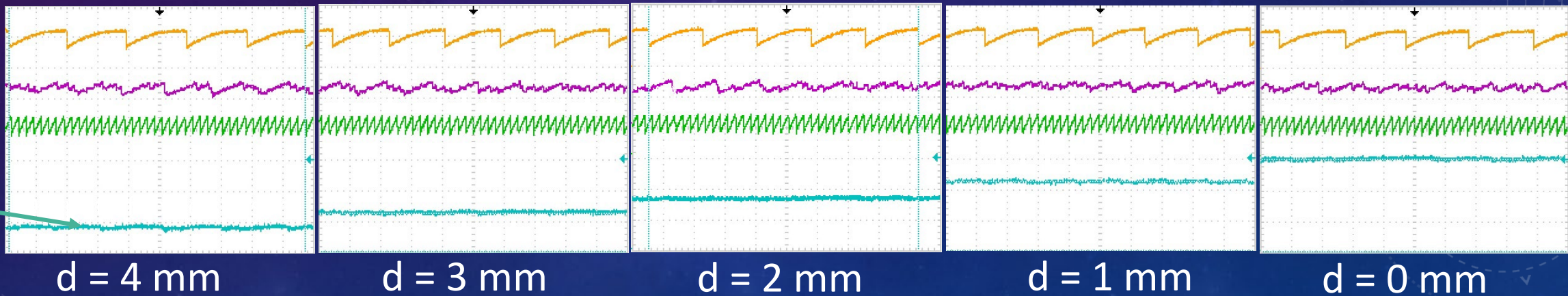
- The following tests were performed:
 - Force sensor response
 - Possibility to adjust force strength by adjusting system parameters
 - Regulatory performance of the neural network



RESULTS



- Observations:
 - Higher motor frequency generates more force
 - Frequency of the inhibitory neurons oscillates



Force increases

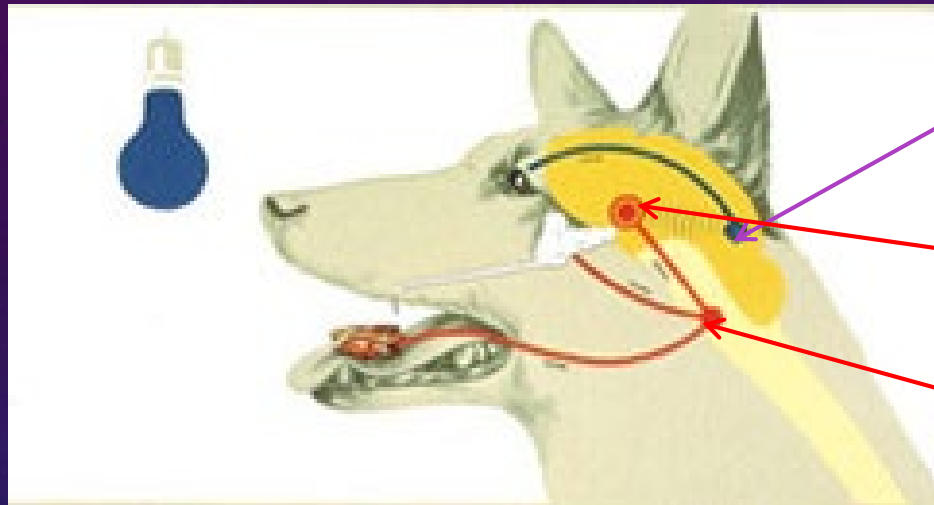


THUS ... THE SNN IS SMALL

- The SNN includes a **few excitatory neurons** that determine SMA **actuator contraction**
- And a few **inhibitory neurons** that are driven by a **force sensor**
- With a few neurons **SNN** is able to control the force applied on an object by the two opposing fingers
- SNN is a good regulator for the contraction force of SMA actuators



HEBBIAN LEARNING

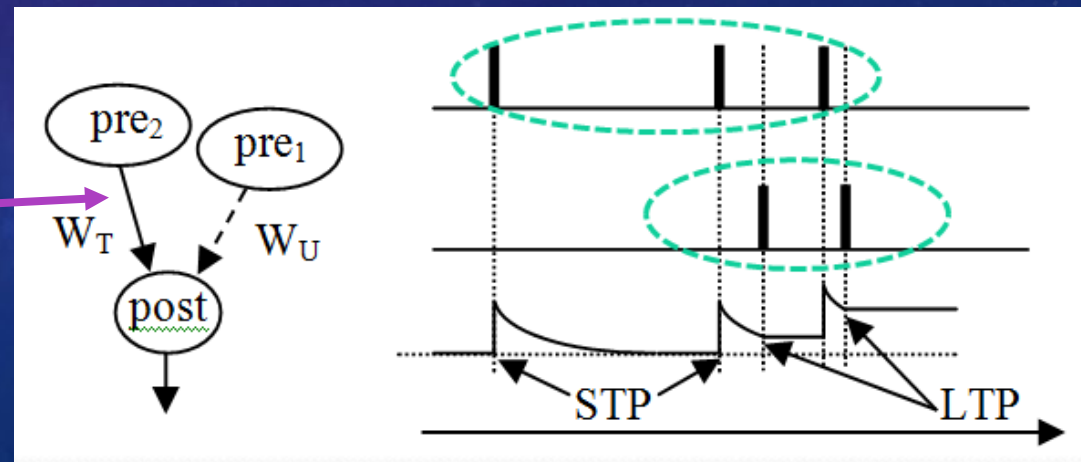


Occipital lobe *Image processing*

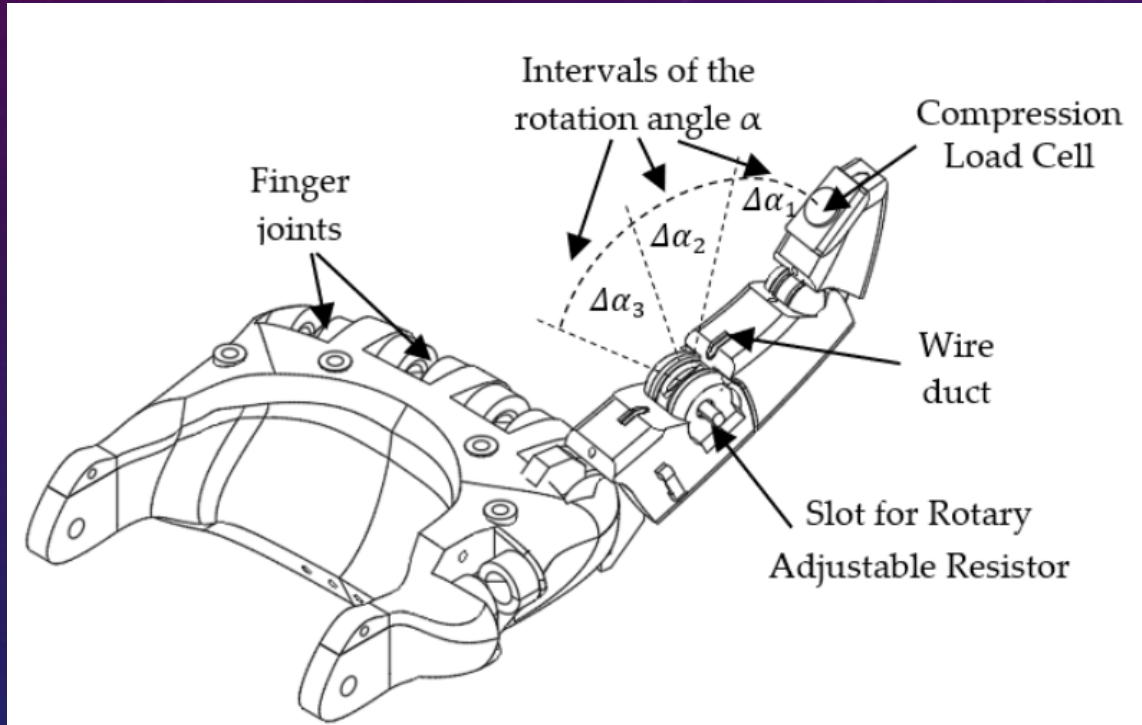
Salivary nucleus: *Activation of glands*

Parietal lobe: *Taste detection*

Trained



MORE COMPLEX CONTROL OF FINGER'S MOTION



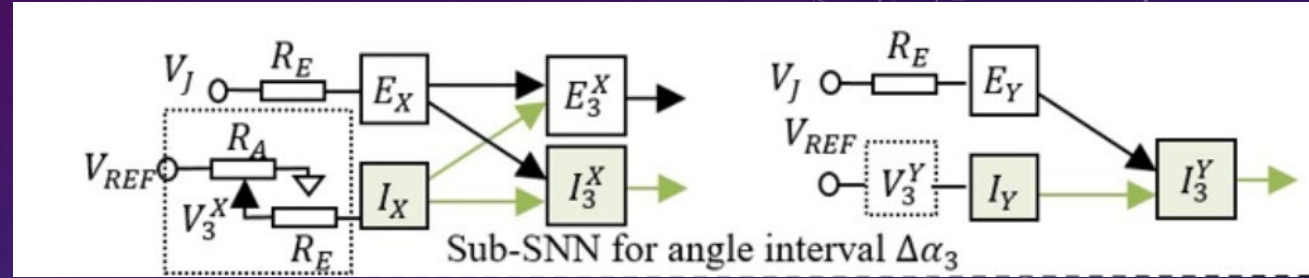
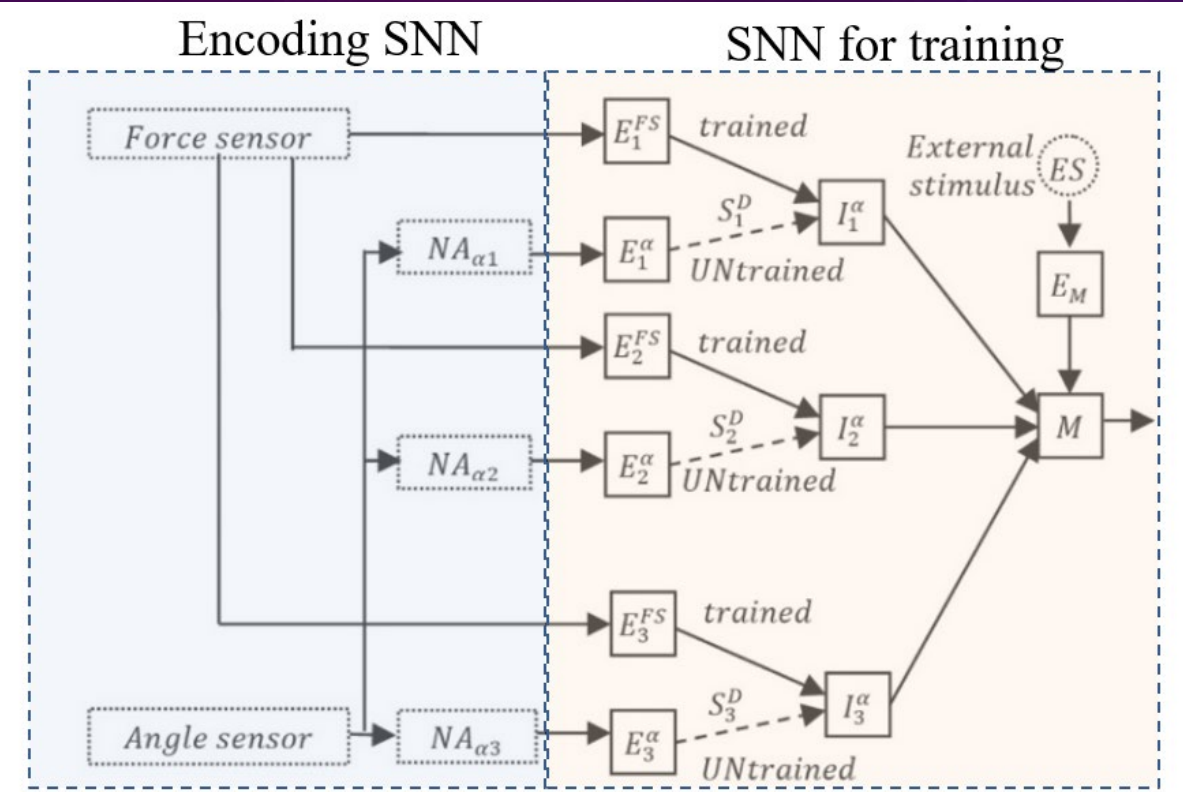
The finger can be stopped in target angles of rotation

The finger motion is stopped where the finger tip is blocked

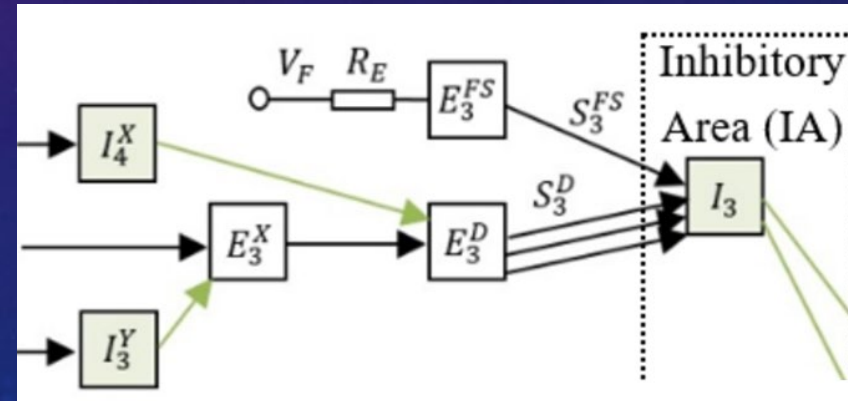
- Anthropomorphic finger which is actuated by SMA and have two force sensors on the tip and rotary sensor in the junction.

SNN STRUCTURE – HIGHER COMPLEXITY

Encoding SNN for a single α



Inhibitory motor neuron for one α



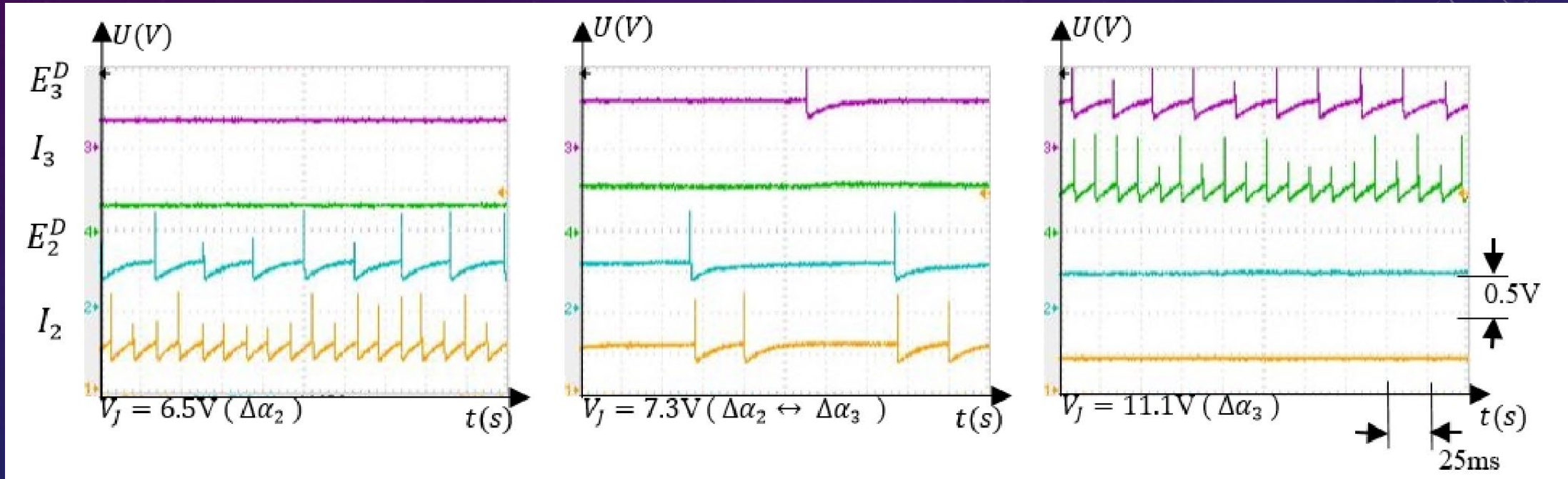
- The **Spiking Neural Network** includes an:
 - Encoding SNN module for the angle of rotation.
 - A decoding SNN that can be trained.

RESULTS – ENCODING LAYER

First interval

Between the angle intervals

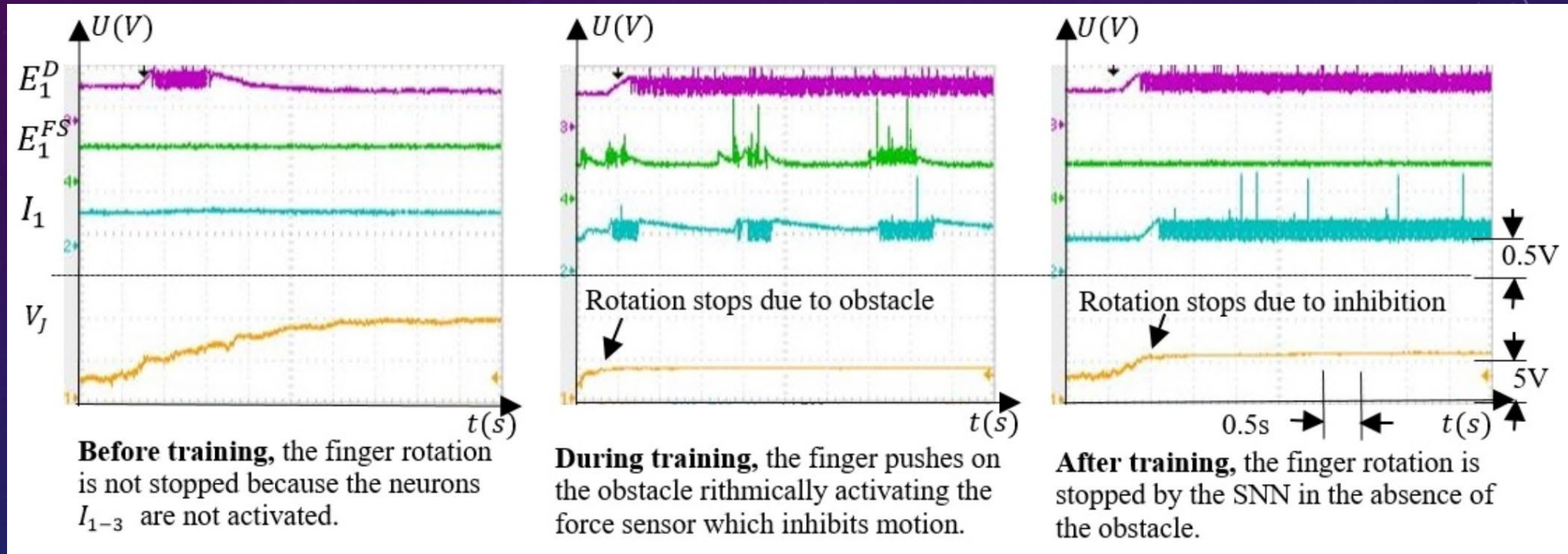
Second interval



The activity of the inhibitory neurons when the finger crosses between two angle intervals.

LEARNING TO STOP THE FINGER

Rhythmic actuation – The finger tries to push on the obstacle

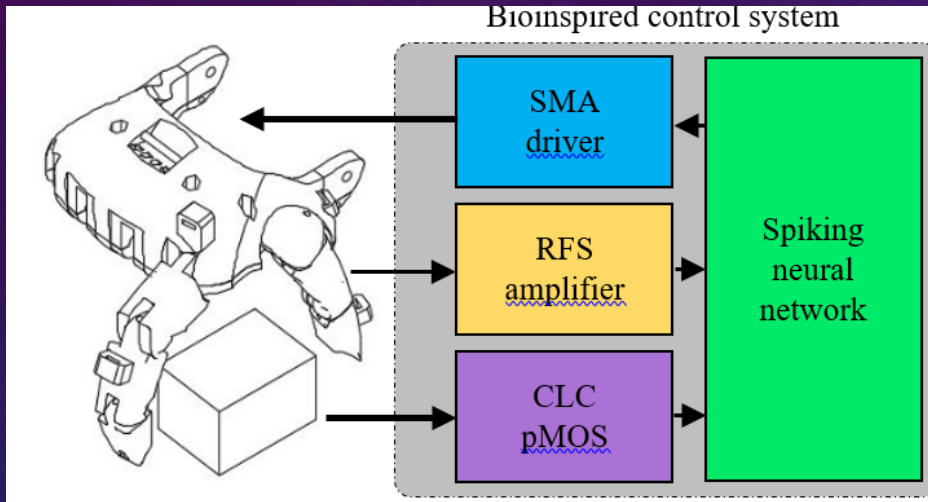


Initial state: With no obstacle the finger does not stop

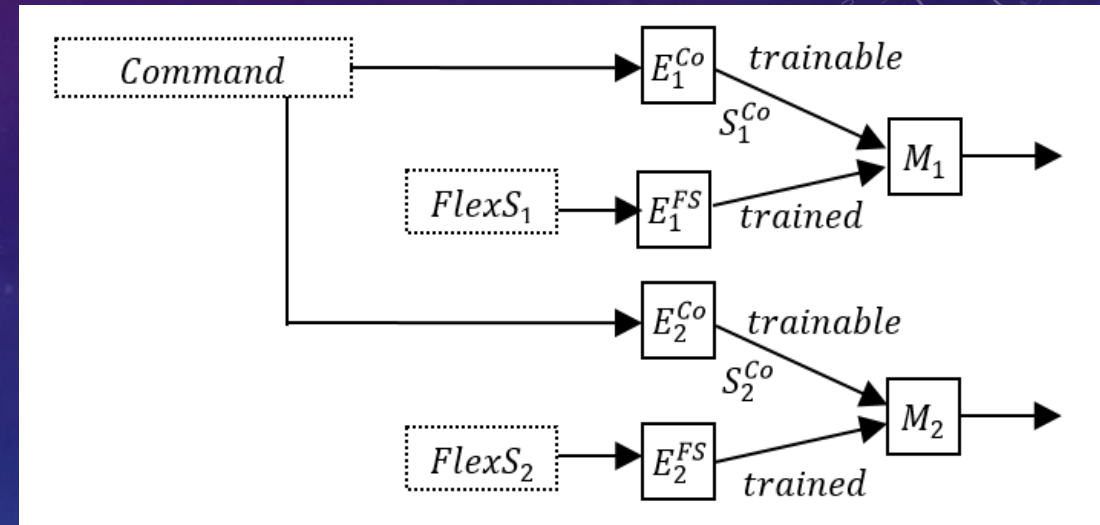
After learning: The finger stops without the obstacle

LEARNING WHICH MOTION TO INITIATE

Robotic hand with flex and force sensors

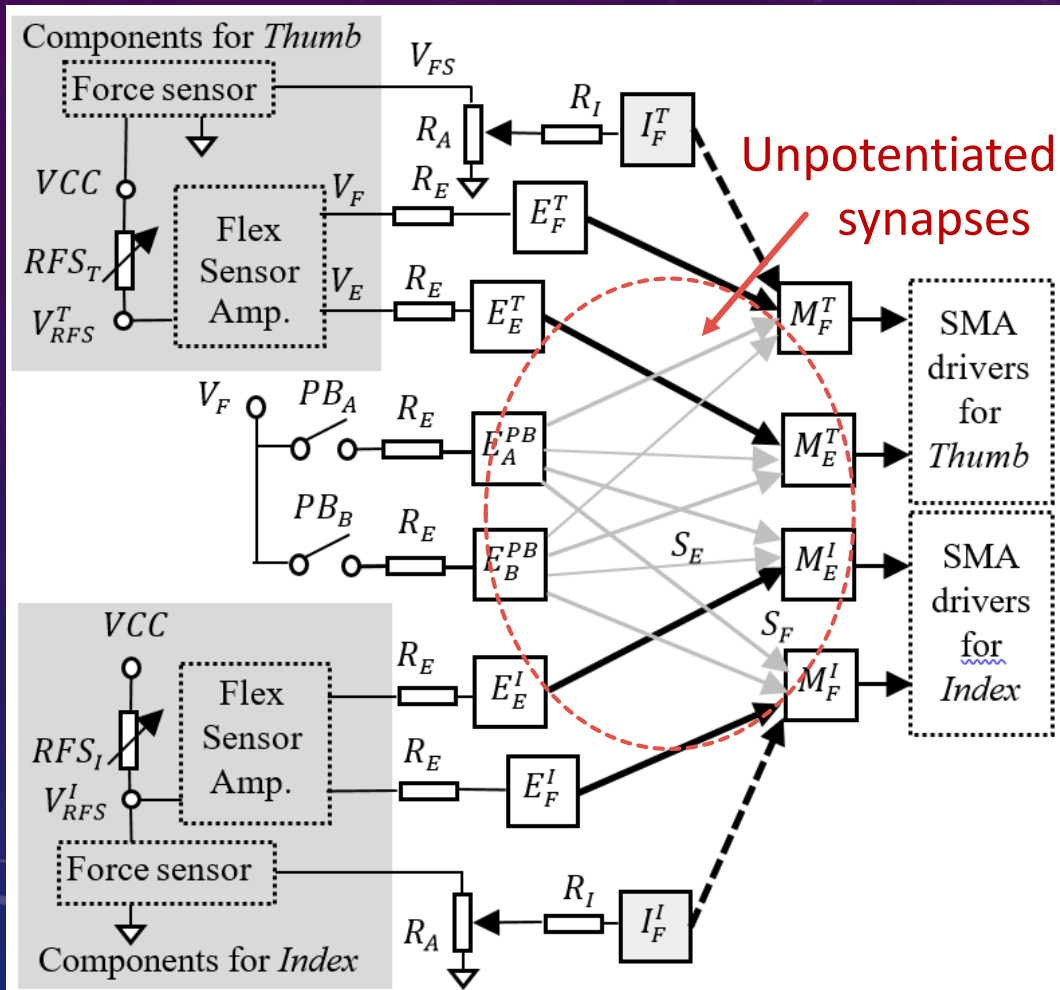


The general structure of the SNN



- Flex sensors detect which finger is moved and in which direction
- Force sensors detect if the finger touches an object

SNN STRUCTURE



- The SNN includes excitatory, inhibitory and **motor neurons**
- The neurons with **potentiated synapses** are connected to the sensors that detect motion.
- The **un-potentiated synapses** are connected command
- Concurrent activation of potentiated and un-potentiated synapses determine learning

Synapses representation:

Inhibitory potentiated \dashrightarrow

Excitatory potentiated \longrightarrow

Excitatory un-potentiated \rightarrow

Neurons:

M - motor

E - excitatory

I - inhibitory

Indices:

T - thumb

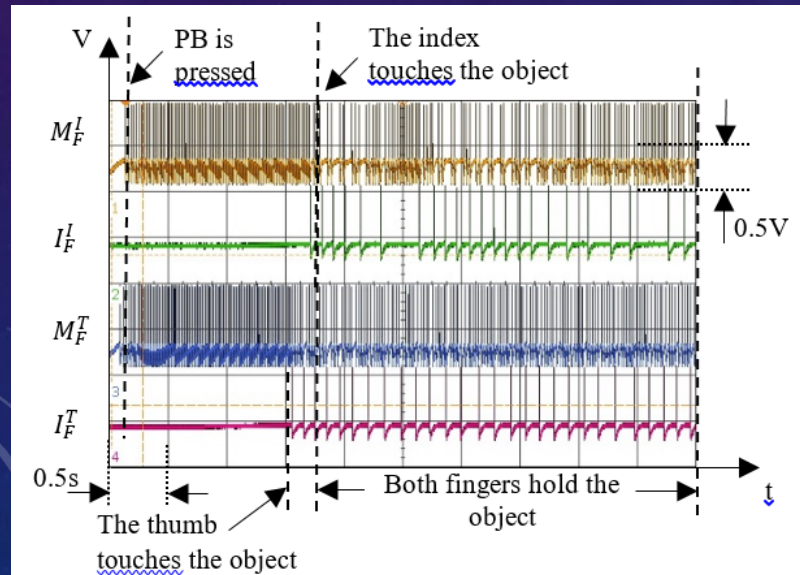
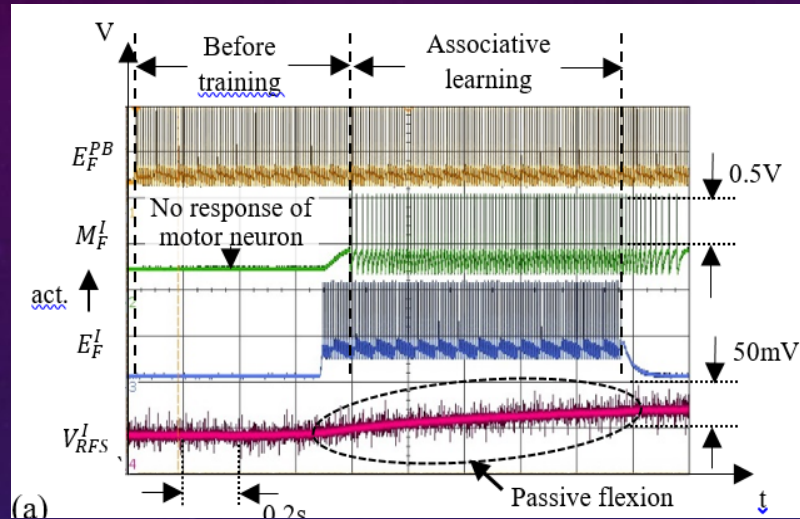
I - index

F - flexion

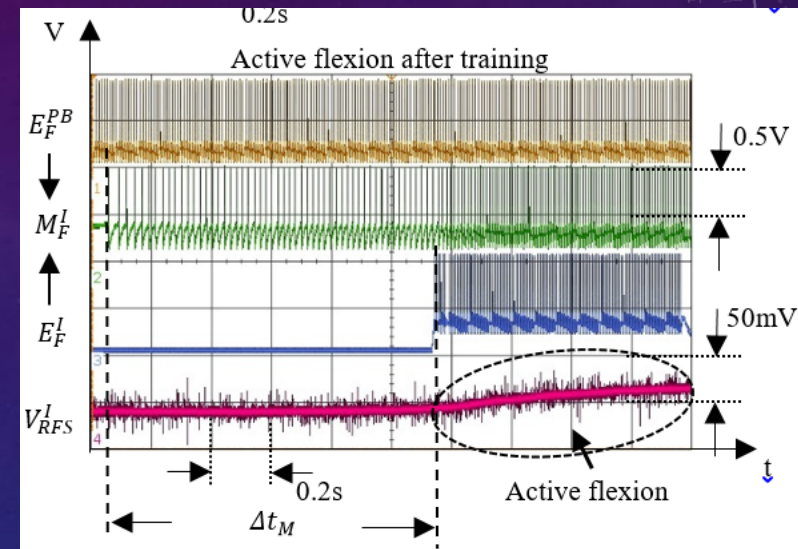
E - extension

RESULTS

During training



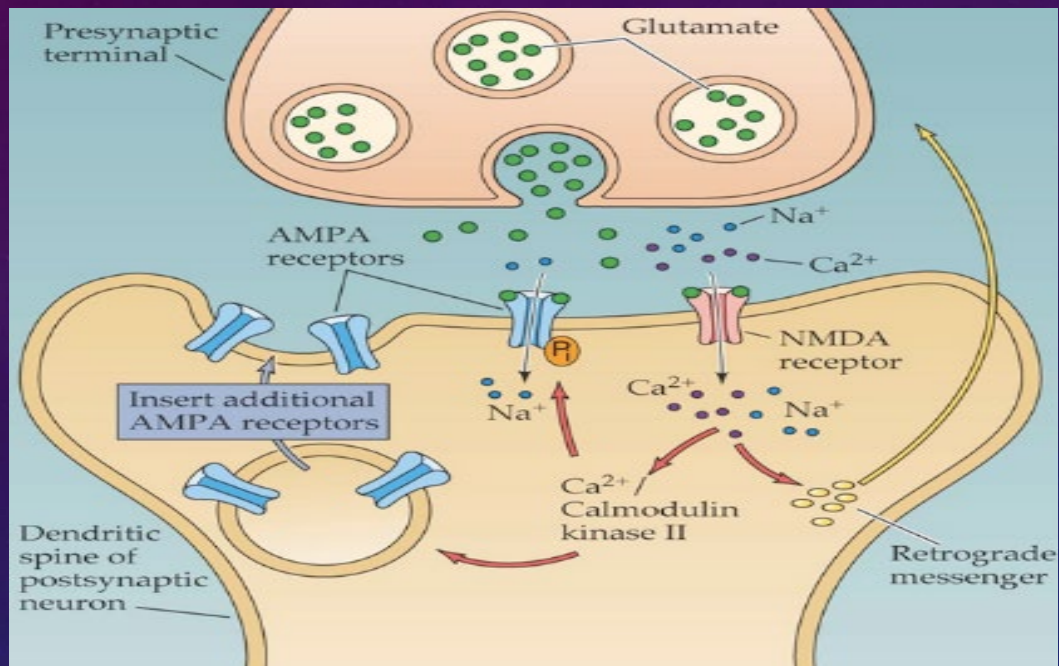
After training



- During training the finger is moved by hand
- After training the same motion is controlled by the SNN
- The finger is stopped when it touches an object

REMARKS

- The SNN can be trained by **physical guidance**
 - Good to teach the children handwriting (for biological networks)
 - Special skills such as walking on a rope
- SNN learns to:
 - Start motion of the fingers – that were executed passively
 - Stop motion in the target intervals
- The synapses are potentiated by the mechanisms of Hebbian learning
- SNN is simple with just a few neurons



THANKS FOR YOUR ATTENTION