

L. Colucci-D'Amato • V. Bonavita • U. di Porzio

The end of the central dogma of neurobiology: stem cells and neurogenesis in adult CNS

Is neuroplasticity in the central nervous system the missing link to our understanding of chronic musculoskeletal disorders?

René Pelletier^{1*}, Johanne Higgins^{1,2} and Daniel Bourbonnais^{1,2}

TOPICAL REVIEW

The olympic brain. Does corticospinal plasticity play a role in acquisition of skills required for high-performance sports?

Jens Bo Nielsen¹ and Leonardo G. Cohen²

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COLUCCI-D'AMATO L et al. 2006; PELLETIER R et al. 2015; NUDO RJ et al. 1996; NIELSON JB AND COHEN L 2007

Performance is a brain circuit forged through neuroimmune plasticity. #BSMPG2015

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Performance is a brain circuit forged through neuroimmune plasticity. #BSMPG2015

Neuroimmune Science

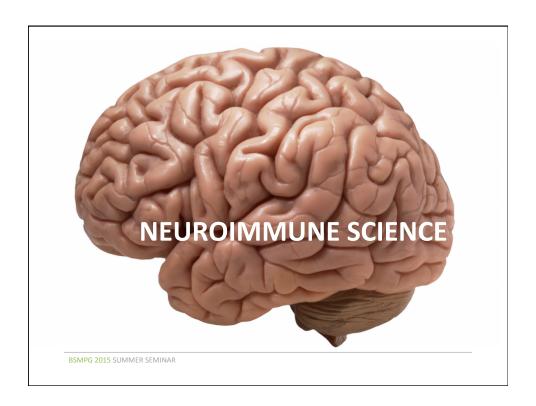
Neuroimmune Plasticity

#Priming

#Pruning

#Prefrontal

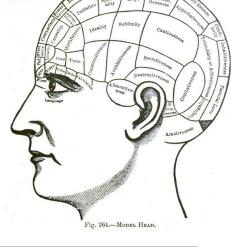
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A suggestion is that it works by being ... complex self-constructing distributed

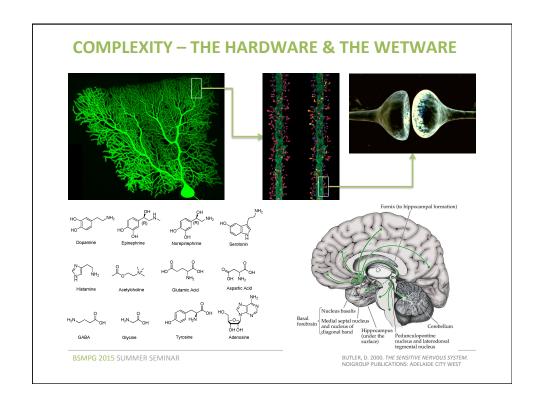
representational

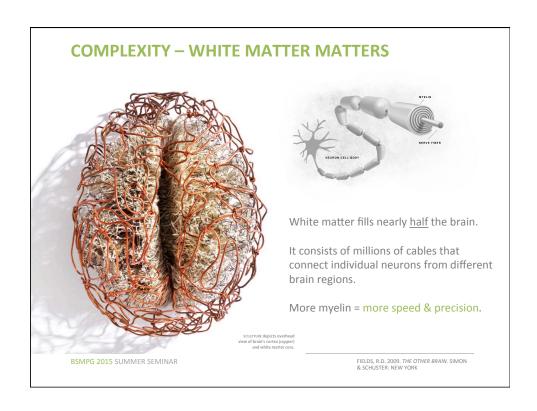
plastic



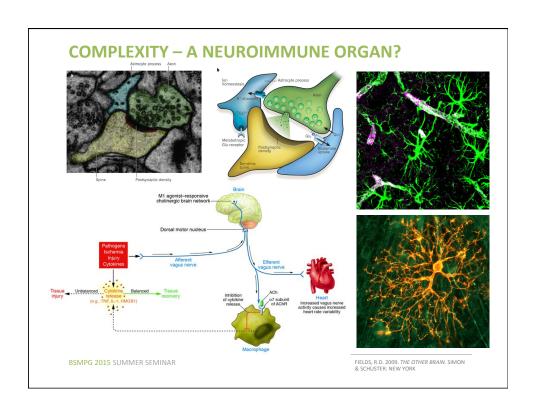
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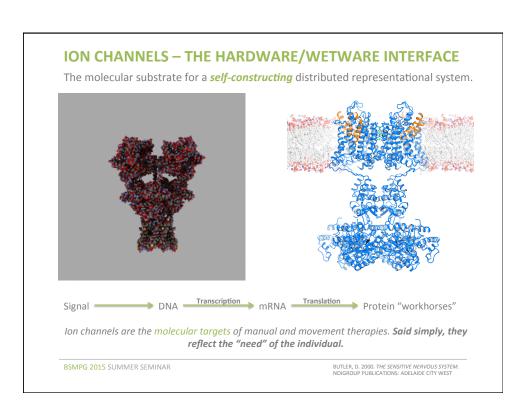
BUTLER, D. 2000. THE SENSITIVE NERVOUS SYSTEM. NOIGROUP PUBLICATIONS: ADELAIDE CITY WEST

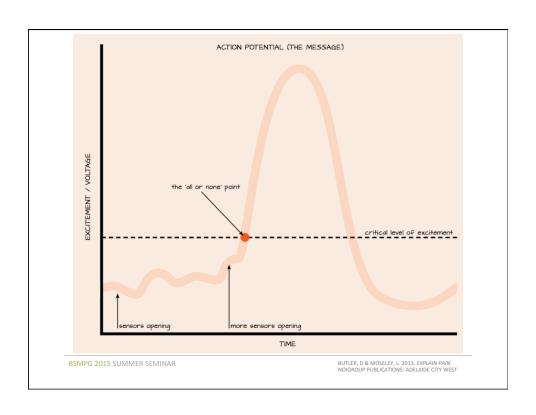


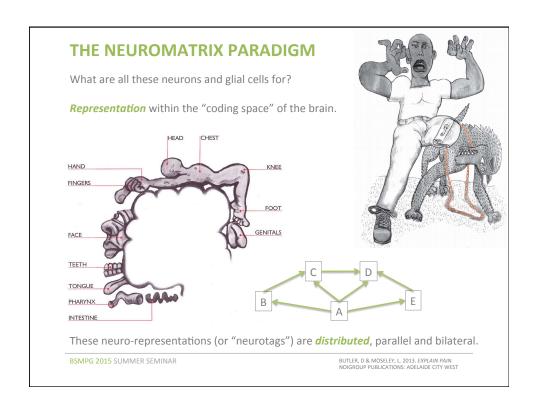


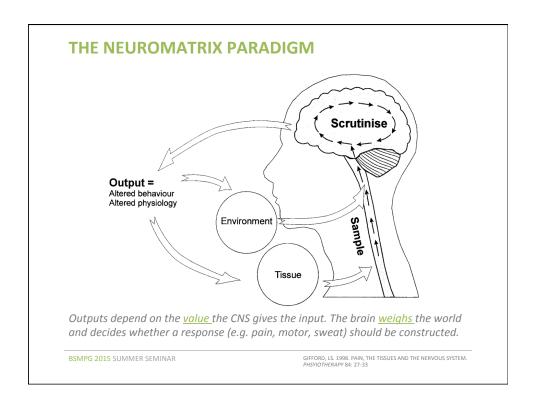
COMPLEXITY — A NEUROIMMUNE ORGAN? **THE PROPERTY OF THE BRAIN. SIMON & SCHUSTER NEW YORK**

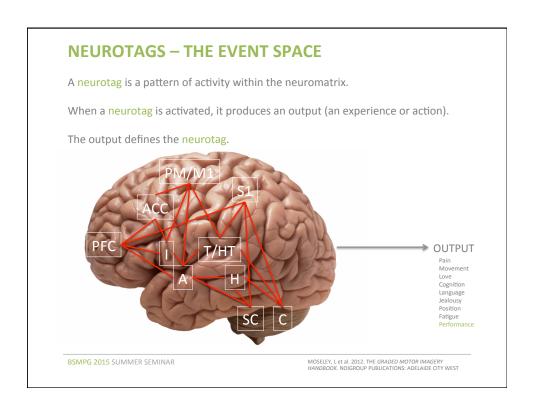


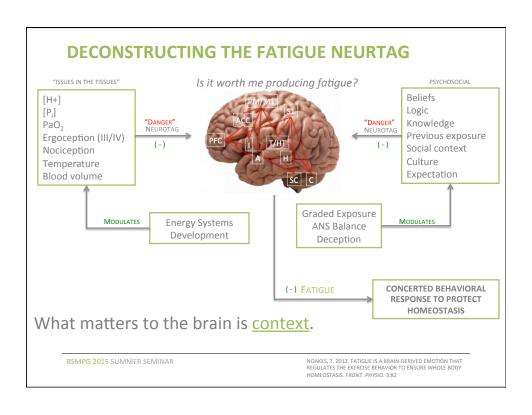














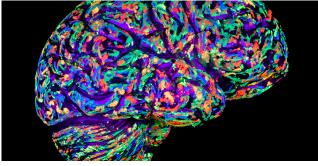


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PLASTICITY IN THE CNS

Neuroplasticity means a change in function of a neuron or group of neurons.

These changes are <u>use-dependent</u>.

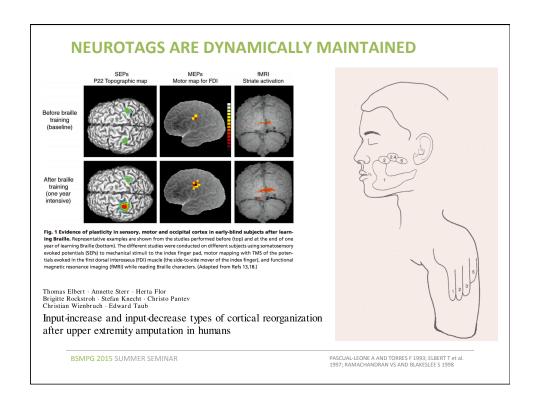


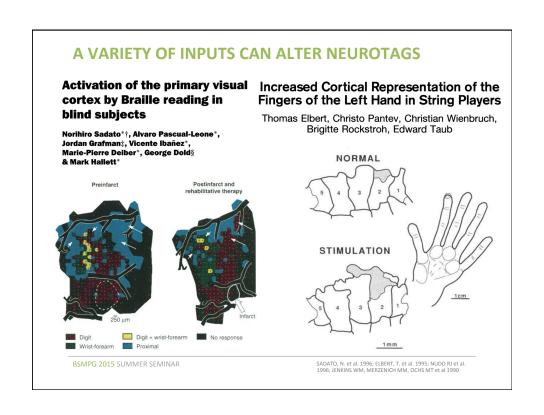
[plas-tik]

 $\underline{\text{All}}$ learning is represented by structural and functional changes to $\underline{\text{all}}$ levels of the neuroimmune system.

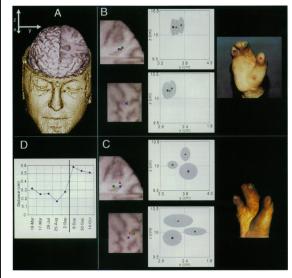
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NEUROTAG REORGANIZATION HAPPENS QUICKLY



Cerebral Cortex September 2007;17:2134-2142 doi:10.1093/cercor/bhl120

Temporal Dynamics of Plastic Changes in Human Primary Somatosensory Cortex after Finger Webbing

The primary somatosensory cortex (SI) exhibits a detailed topographic organization of the hand and fingers, which has been found to undergo plastic changes following modifications of the sensory input. Although the spatial properties of these changes have been extensively investigated, little is known about their temporal dynamics. In this study, we adapted the paradigm of finger webbing, in which 4 fingers are temporarily webbed together, hence modifying their sensory feedback. We used magnetoence-phalography, to measure changes in the hand representation in SI, before, during, and after finger webbing for about 5 h. Our results showed a decrease in the Euclidean distance (ED) between cortical sources activated by electrical stimuli to the index and small finger 30 min after webbing, followed by an increase lasting for about 2 h after webbing, which was followed by a return toward baseline values. These results provide a unique frame in which the different representational changes occur, merging previous findings that were only apparently controversial, in which either increases or decreases in ED were reported after sensory manipulation for relatively long or short duration, respectively. Moreover, these observations further confirm that the mechanisms that underlie cortical reorganization are extremely rapid in their expression and, for the first time, show how brain reorganization occurs over time.

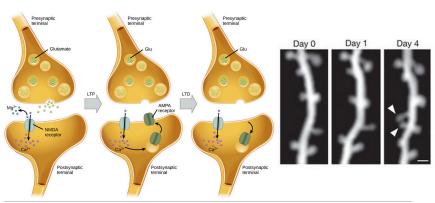
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MOLINGER A et al. 1993; STAVRINOU M et al. 2006

GREY MATTER PLASTICITY

"Neurons that fire together, wire together."

Long-term potentiation requires cooperativity, associativity, and specificity.



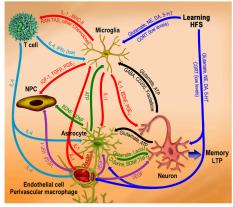
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Immune modulation of learning, memory, neural plasticity and neurogenesis

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ABSTRACT

A B S T R A C T

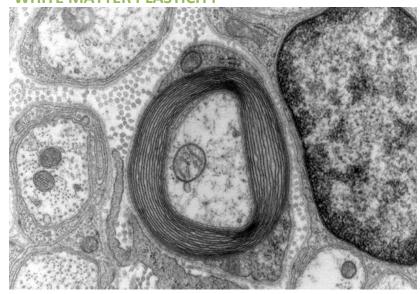
Over the past two decades it became evident that the immune system plays a central role in modulating learning, memory and neural plasticity. Under normal quiescent conditions, immune mechanisms are activated by environmental/psychological stimuli and positively regulate the remodeling of neural circuits, promoting memory consolidation, hippocampal (long-term potentiation (LTP) and neuropenesis. These beneficial effects of the immune system are mediated by complex interactions among brain cells. These heneficial effects of the immune system are mediated by complex interactions among brain cells researched to the control of the proposition of the cells and macrophages heurons, and neural precursor cells. These interactions involve the responsiveness of non-neuronal cells to classical neurotransmitters (e.g., glutamate and monoamine) and bornomes (e.g., glutocorticoids), as well as the secretion and responsiveness of neurons and glis to low levels of inflammatory cytokines, such as pincategoral positions under which the immune system is strongly activated by infection or injury, as well as by severe or chronic stressful conditions, glia and other brain immune cells change their morphology and functioning and secrete high levels of pro-inflammatory cytokines and prostaglandins. The production of these inflammatory mediators disrupts the delicate balance needed for the neurophysiological actions of immune processes and produces direct detrimental effects on memory, neural plasticity and neurogenesis. These effects are mediated by inflammation-induced neuronal hyper-excitability and adrenocordical stimulation, followed by reduced production of neurotrophins and other plasticity-related molecules, facilitating many forms of neuropathology associated with normal aging as well as neurodegenerative and neuropsychiators diseases.

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WHITE MATTER PLASTICITY



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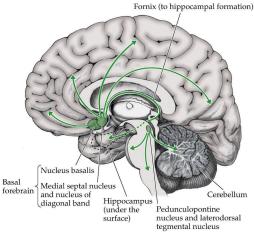
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FIELDS, R.D. 2009. THE OTHER BRAIN. SIMON & SCHUSTER: NEW YORK

PRINCIPLES OF NEUROIMMUNE PLASTICITY



Fornix (to hippocampal formation) Principles of Experience-Dependent
Neural Plasticity: Implications for
Rehabilitation After Brain Damage

SUPPLEMENT

Use it or lose it.

Use it and improve it.

Specificity.

Repetition.

Intensity.

Time.

Salience.

Age.

Transference.

Interference.

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KLEIM JA AND JONES TA. 2008. PRINCIPLES OF EXPERIENCE-DEPENDENT NEURAL PLASTICITY: IMPLICATIONS FOR REHABILITATION AFTER BRAIN DAMAGE. *J SPEECH LANG HEAR RES.* 51(1): S225-39

