Understanding impaired learning with enhanced plasticity based on work in preparation with: T.D. Barbara Nguyen-Vu, Grace Q. Zhao, Han-Mi Lee, Surya Ganguli, Carla J. Shatz, Jennifer L. Raymond

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Learning requires synaptic plasticity. Expect enhanced plasticity \rightarrow enhance learning.

[Tang et al. (1999), Malleret et al. (2001), Guan et al. (2009)]

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But often: \rightarrow impairment.

[Migaud et al. (1998), Uetani et al. (2000), Hayashi et al. (2004)] [Cox et al. (2003), Rutten et al. (2008), Koekkoek et al. (2005)] Learning requires synaptic plasticity. Expect enhanced plasticity \rightarrow enhance learning.

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Mice with enhanced cerebellar plasticity can show both impaired and enhanced learning.

Simple synapses cannot explain behaviour.

 \rightarrow Necessary & sufficient conditions on complex synapses to replicate this.

- Motor learning
 - Cerebellar learning of mice with enhanced plasticity
 - Complex synaptic models
- (Memory capacity of complex synapses)

Vestibulo-Occular Reflex



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Eye movements compensate for head movements to maintain fixation.

Requires control of VOR gain = $\frac{\text{eye velocity}}{\text{head velocity}}$

Needs to be adjusted as eye muscles age, etc.

VOR training



VOR Decrease Training





Impaired learning with enhanced plasticity

Enhanced plasticity impairs learning

Knockout of MHC-I $\mathsf{D}^\mathsf{b}\mathsf{K}^\mathsf{b}$ molecules in PF-Pk synapses

[McConnell et al. (2009)]

 \rightarrow lower threshold for LTD \rightarrow enhanced plasticity

Hypothesis: enhanced learning.



Depletion hypothesis

Learning rate \sim intrinsic plasticity rate \times # synapses available for LTD.



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Depletion hypothesis

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Impaired learning with enhanced plasticity

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Replenishment by reverse-training



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Questions:

• Can the depletion effect overcome enhanced intrinsic plasticity?



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Questions:

- Can the depletion effect overcome enhanced intrinsic plasticity?
- How can a little replenishment help, but too much hurt?



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VOR Increase Training



Complex synapses



Assumptions:

- Candidate plasticity events occur independently at each synapse,
- Each synapse responds with the same state-dependent rules,
- Keep track of distribution of synapses across states.

[Fusi et al. (2005), Fusi and Abbott (2007), Barrett and van Rossum (2008)]

Models of synaptic dynamics

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Models of synaptic dynamics



Simple synapses cannot explain the data



Complex synapses can explain the data



Enhanced plasticity can enhance or impair learning





Intrinsic plasticity dominates depletion ↓ enhanced plasticity enhances learning Depletion dominates intrinsic plasticity ↓ enhanced plasticity impairs learning

Reverse-training can impair or enhance learning





reverse-training depopulates boundary ↓ impaired learning reverse-training repopulates boundary ↓ enhanced learning

The success of the serial model relies on two features:

- Complexity needed to amplify the effect of depletion,
- Metaplasticity repeated potentiation makes subsequent depression harder.



 We find diverse behavioural patterns in these mutant mice: Enhanced plasticity → enhance/impair learning depending on prior experience.

Reverse-training \rightarrow enhance/impair learning depending on plasticity rates.

- We can explain these behavioural patterns using synaptic models.
- Key required synaptic properties are: Synaptic complexity: necessary to amplify depletion. Synaptic stubbornness: repeated potentiation makes subsequent depression harder.
- We used behaviour to constrain the dynamics of synaptic plasticity

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What about memory?

• Simple synapses have poor memory storage capacity. Synaptic complexity is needed for rescue.

[Amit and Fusi (1992), Amit and Fusi (1994)]

- Trade-off between learning and remembering: Too rigid → difficult to learn new memories. Too plastic → new memories quickly overwrite old.
- Exploring the *entire* space of complex synaptic models
 → upper bounds on their storage ability
 & the models that saturate them.

[Lahiri and Ganguli (submitted)]

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We have N synapses with M internal states each.

We study the decay of one memory over time due to corruption by subsequent memories.

We prove that, no matter what the structure, no synaptic model can have:

- initial fidelity (SNR) greater than \sqrt{N} .
- memory lifetime greater than $\sim \sqrt{N}M$.
- fidelity decay slower than $\sim \sqrt{N}M/t$.

At late times, fidelity is maximised by a model with a simple chain structure.

Surya Ganguli

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Jennifer Raymond Carl Barbara Nguyen-Vu Han Grace Zhao Aparna Suvrathan

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