

Coglink networks reveal computational mechanisms of uncertainty management and its perturbation in schizophrenia.

Control No. 4120

Mien Brabeeba Wang, Nancy Lynch, MIT CSAIL, USA Michael Halassa, Tufts University School of Medicine, USA

Introduction

- Genetic constitutes a significant risk in Schizophrenia (SZ) [1] and computational modeling has shown deficits in beliefupdating processes as a key aspect of the disorder [2].
- However, the intricate mechanisms bridging these genetic risk factors and belief updating deficits remain poorly understood.
- Our model, CogLinks, capable of linking mechanisms with normative behaviors, offers an avenue to study such connections.
- CogLinks model prefrontal cortex (PFC) and mediodorsal thalamus (MD) which not only involve in belief-updating processes but also show altered functional couplings in patients [3].
- We consider a probability reversal task in which patients show slow switching upon reversal and elevated win-switch rate (choosing an alternative action after receiving rewards) [4].

A mechanistic neural model Modulate cortical activities and plasticity via interneurons Learn contextual models of the environment

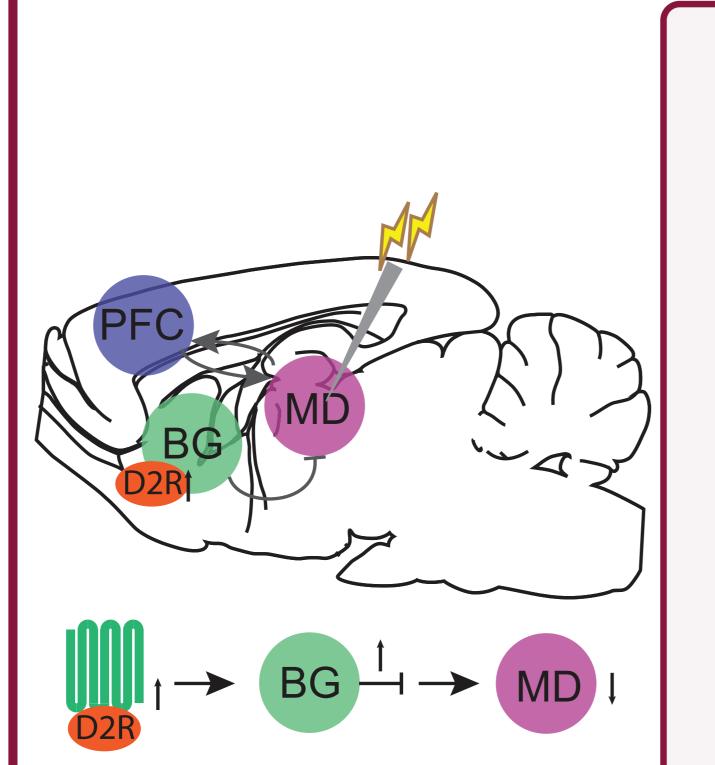
PFC-MD connections learn the contextual environmental model through Hebbian plasticity and infer the context in MD via recurrent dynamics. MD then contextually modulates cortical activity [5] and plasticity [6] through interneurons, in which VIP neurons amplify context-relevant cortical connectivity while PV neurons suppress context-irrelevant information.

KEY REFERENCES

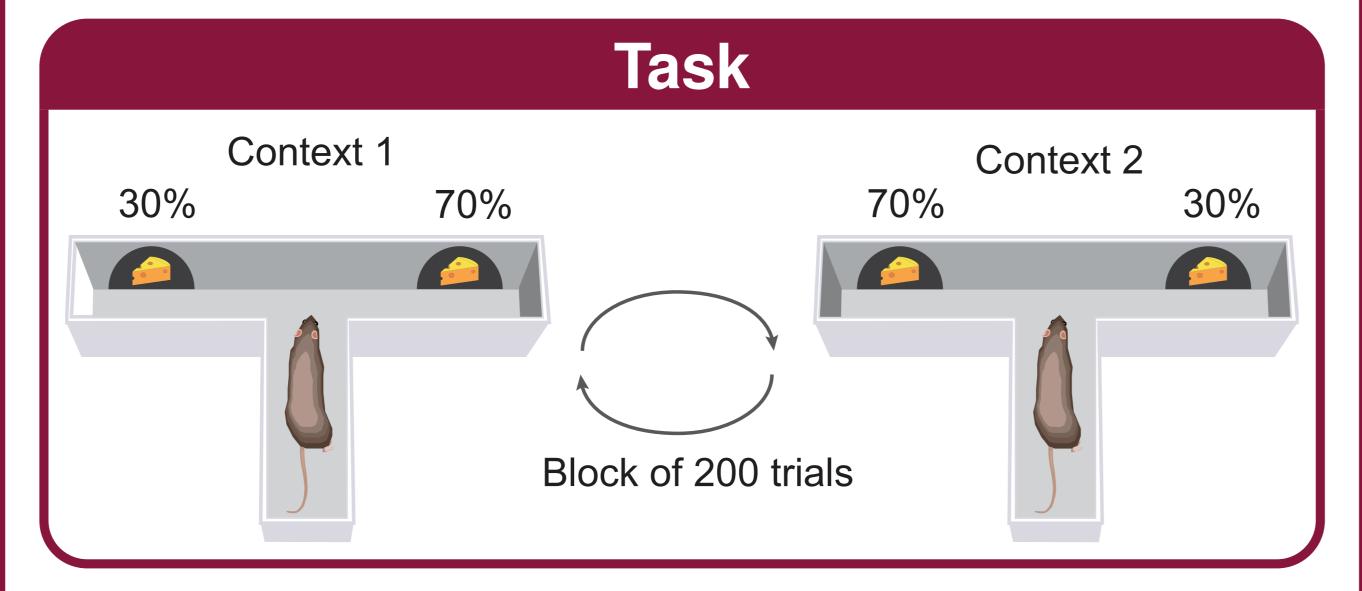
- [1] Gejman, P. V. et al.. Psychiatr Clin North Am. 2010
- [2] Baker, S. C.. Brain. 2019.
- [3] Paz, R. and Amaya, J. M.. J Psychiatry Neurosci. 2008. [4] Waltz, J. A.. Neuroscience. 2017.
- [8] Zhou, T. et al.. bioRxiv. 2024.

[6] Canto-Bustos, M. et al.. J Neurosci. 2022. [7] Abi-Dargham, A. et al.. PNAS. 2000.

Schizophrenia model and rescue model



We consider a model with hyperactivation of striatal D2 receptors (D2Rs) because most SZ patients show elevated striatal D2Rs expression [7]. the abundance of D2Rs increases the inhibition from BG to thalamus, we model SZ by reducing the excitability of MD to mimic strong BG inhibition. To rescue the model, we inject a small current in MD.

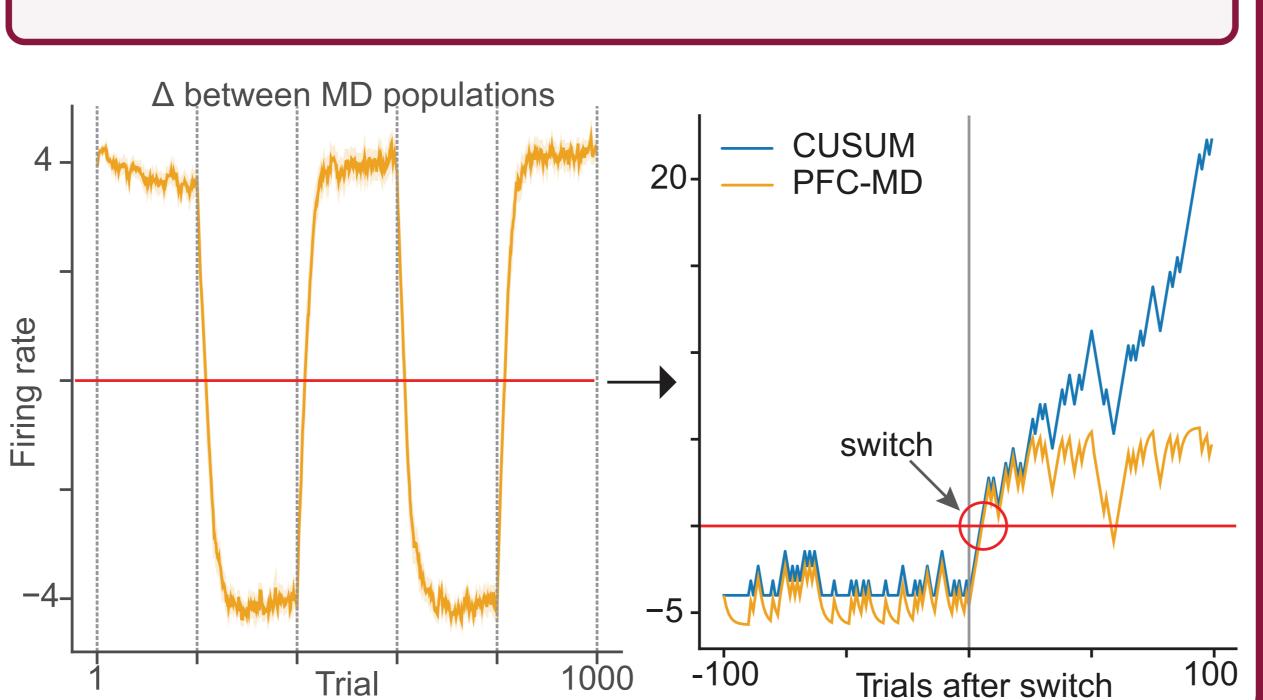


Approximation to normative model

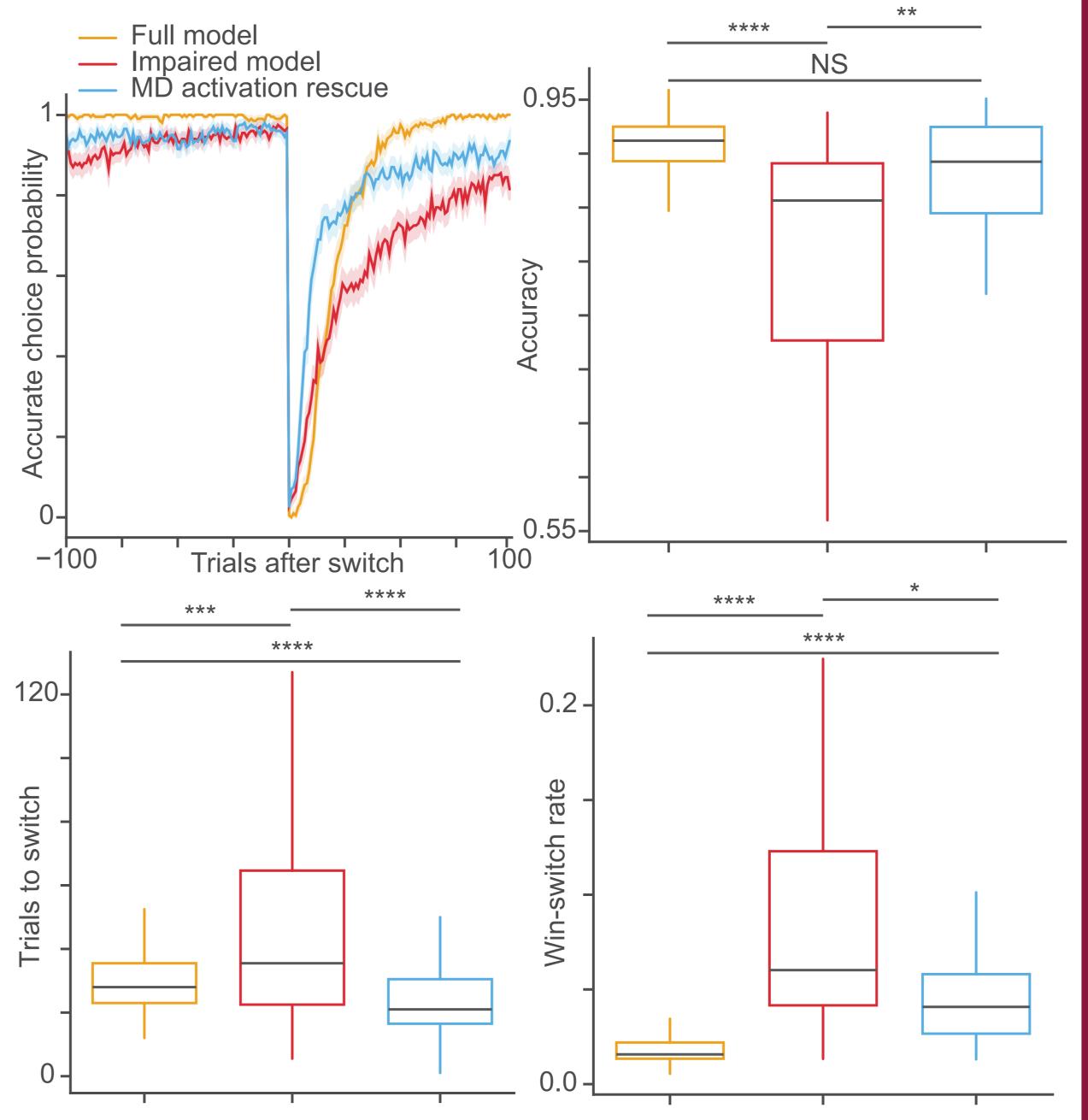
- It is usually difficult to understand a mechanistic model on a computational level due to its complexity.
- To overcome this, we mathematically approximate our mechanistic model to a novel normative model and analyze its functions and performance.

Theorem 1. After PFC-MD synapses learn the contextual generative model $P(a_t, r_t|c)$, our PFC-MD circuit approximates to a multiple change points generalization of CUSUM algorithm, an algorithm that is known to detect single environmental changes optimally [10].

Our PFC-MD circuit approximates a normative model that detects sequential environmental changes optimally.

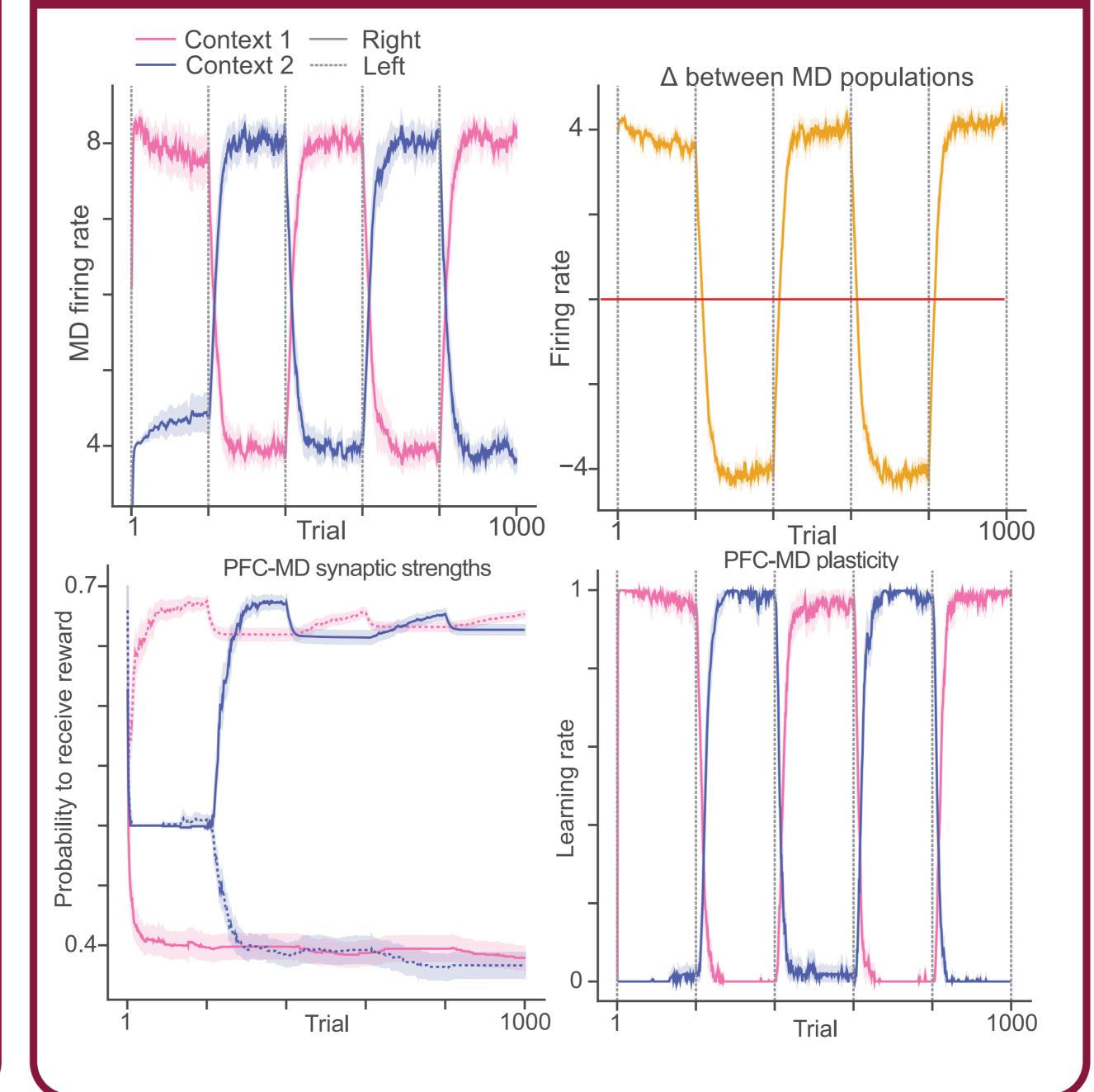


Behaviors

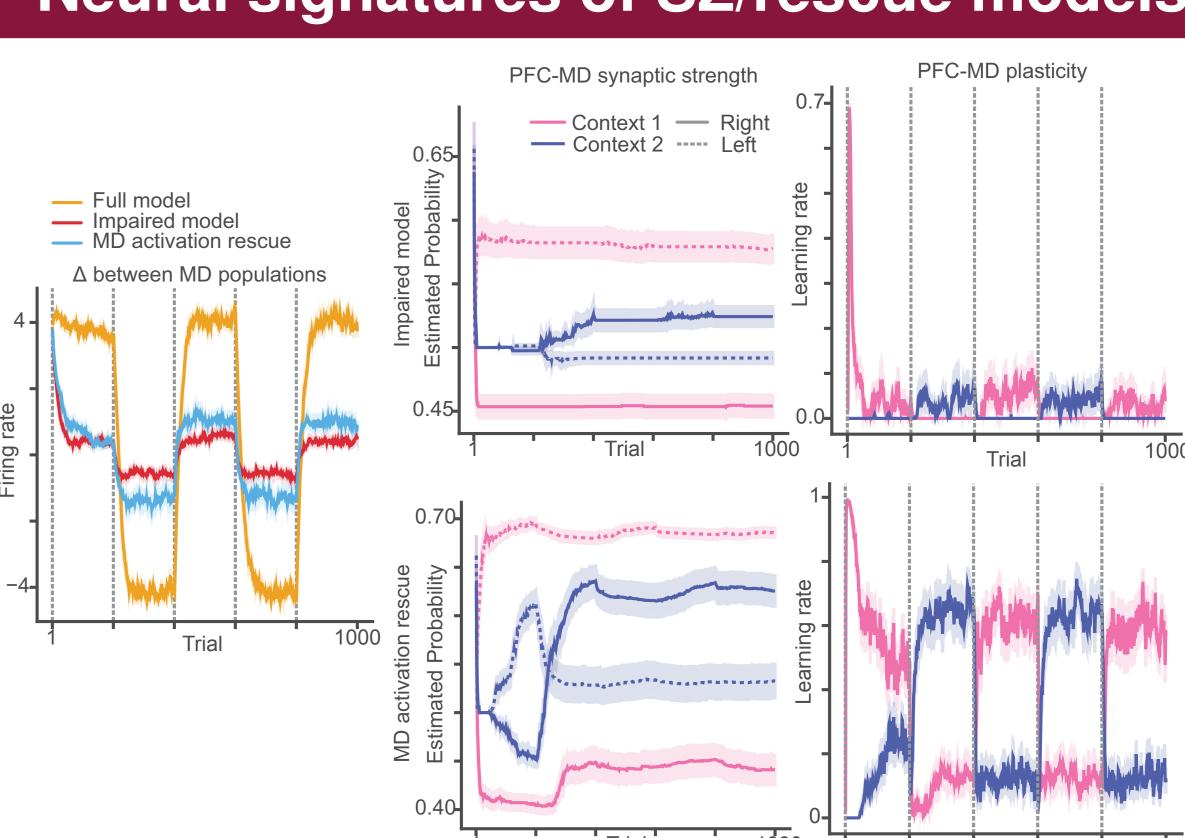


- Our SZ model has much lower accuracy with high variance compared to the normal model.
- It also shows slow switching upon reversal and elevated winswitch rates, consistent with the behavioral findings in patients.
- Our rescue model rescues the behaviors, consistent with the MD activation experiment on SZ-related mouse model [8].

MD activity tunes to contexts and contextually modulates PFC-MD plasticity



Neural signatures of SZ/rescue models



We first investigate the MD activity and the normative model:

- The normative model of SZ model has a much lower threshold.
- This shows that SZ model has a strong prior on environmental volatility and therefore much higher win-switch rates.

Theorem 2. The threshold of the normative theory of the SZ model is $T_{SZ} = \frac{2\beta_{d2}}{1-\beta_{d2}} |\langle I_1^{pfc/md} - I_2^{pfc/md} \rangle| \approx 0.64 \ll 4 = T_{normal}$ much smaller than the threshold of the normal model. Furthermore, the corresponding normative model is a leaky integrator, further strengthening the prior on environmental volatility.

Both MD activity and the corresponding normative model show SZ model has a strong prior on environmental volatility, potentially contributing to paranoia.

We then examine PFC-MD connectivity and plasticity:

- Indeed, compared to the normal model, SZ model struggles to learn the correct contextual model of the environments.
- The improper learning happens because of the abnormally low learning rate, potentially due to low MD excitability.
- By injecting the current into MD, the rescue model reinstates the proper learning of the environmental model.

The PFC-MD connections in SZ model are unable to learn proper environmental model, potentially contributing to delusion. By injecting current in MD, the rescue model recovers the proper learning of contextual model.

Conclusion

- Our work links neural mechanisms, normative models and cognitive (dys)functions together in a single framework.
- Our CogLink network approximates a normative model that detects environmental changes optimally.
- Our SZ model shows idiosyncratic behaviors observed in patients, including slow switching and elevated win-switch rates.
- Both MD activity and the normative model shows SZ model has a strong prior on environmental volatility (potentially paranoia).
- PFC-MD connectivity in SZ model cannot learn the proper environmental model, potentially contributing to delusion.
- By injecting the current in MD, the rescue model not only rescues the behaviors but also reinstates proper PFC-MD learning in environmental models.