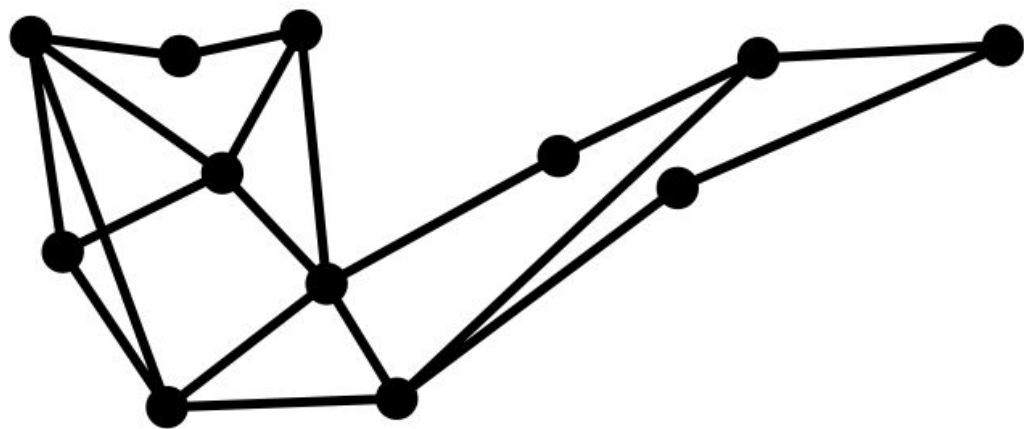


Topological Structure in the Study of Ablation and Desynchronization in Neural Networks

Mark Agrios
College of William & Mary

Quick Review of Graphs



Ceci n'est pas une graph

$$G = (V, E)$$

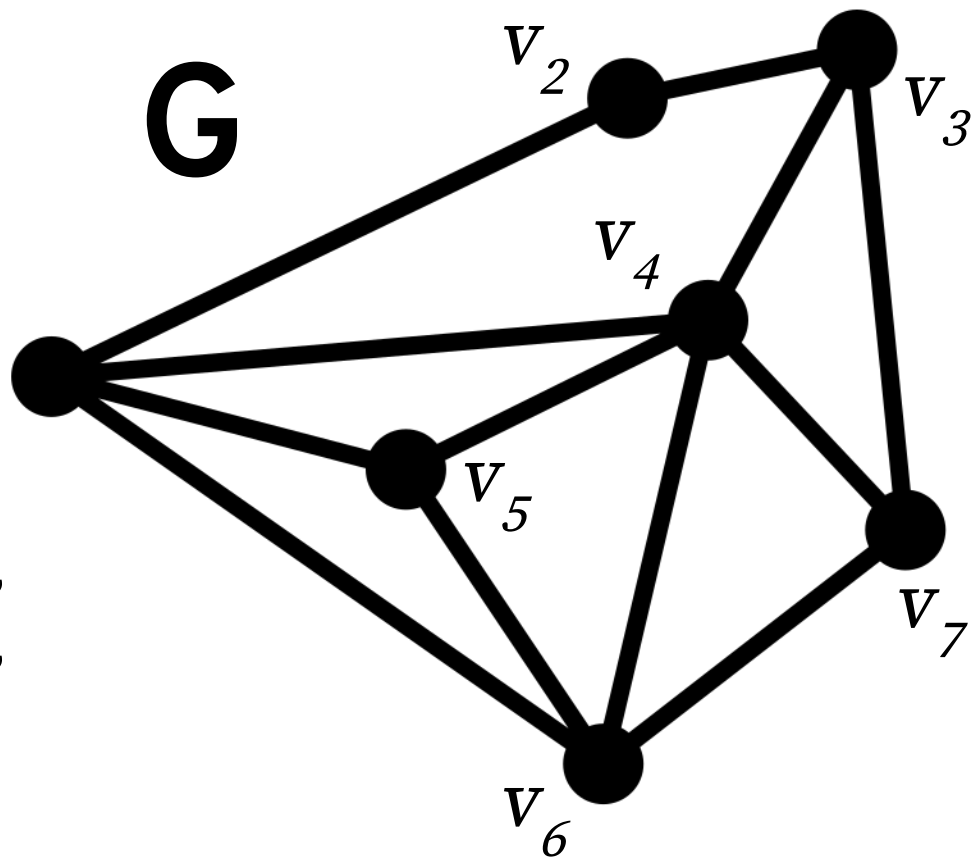
$$V = \{v_1, v_2, v_3, v_4, v_5, v_6, v_7\}$$

$$E = \{\{v_1, v_2\}, \{v_1, v_4\}, \{v_1, v_5\}, \{v_1, v_6\}, \\ \{v_2, v_3\}, \{v_3, v_4\}, \{v_3, v_7\}, \{v_4, v_5\}, \\ \{v_4, v_6\}, \{v_4, v_7\}, \{v_5, v_6\}, \{v_6, v_7\}\}$$

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Simplex: a generalization of the notion of a triangle or tetrahedron to arbitrary dimensions

An n -simplex is an n -dimensional polytope that is the convex hull of $n+1$ vertices.

Simplex: a generalization of the notion of a triangle or tetrahedron to arbitrary dimensions

An n -simplex is an n -dimensional polytope that is the convex hull of $n+1$ vertices.

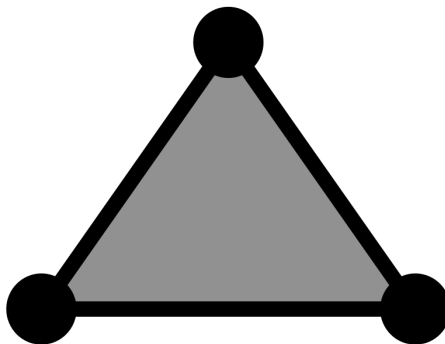
0-simplex



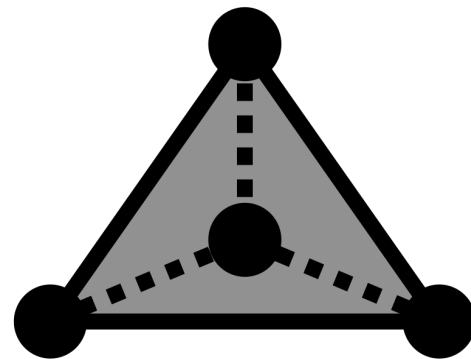
1-simplex



2-simplex



3-simplex



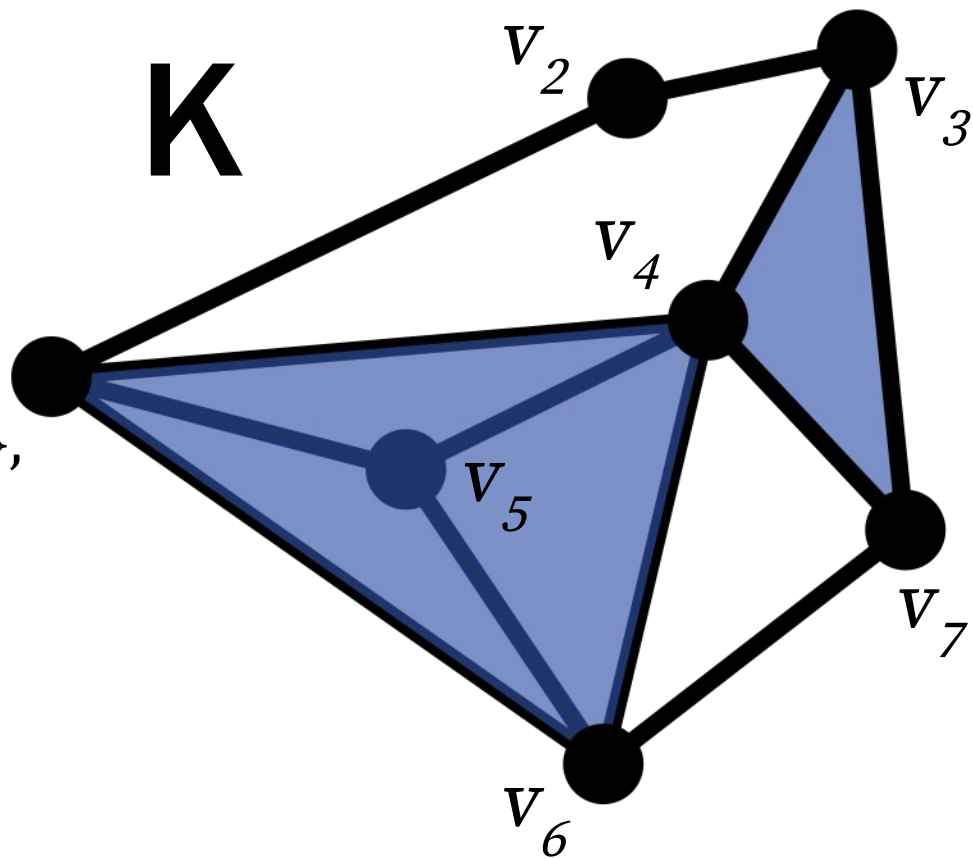
etc...

Simplicial Complex: A simplicial complex, K , is a set of “simplices” that are sets of finite order.

Simplicial Complex Condition:

1. If $a \in K$ then any subset (face) of a is also in K
2. The intersection of any two simplices $a, b \in K$ is either \emptyset or a face of both a and b

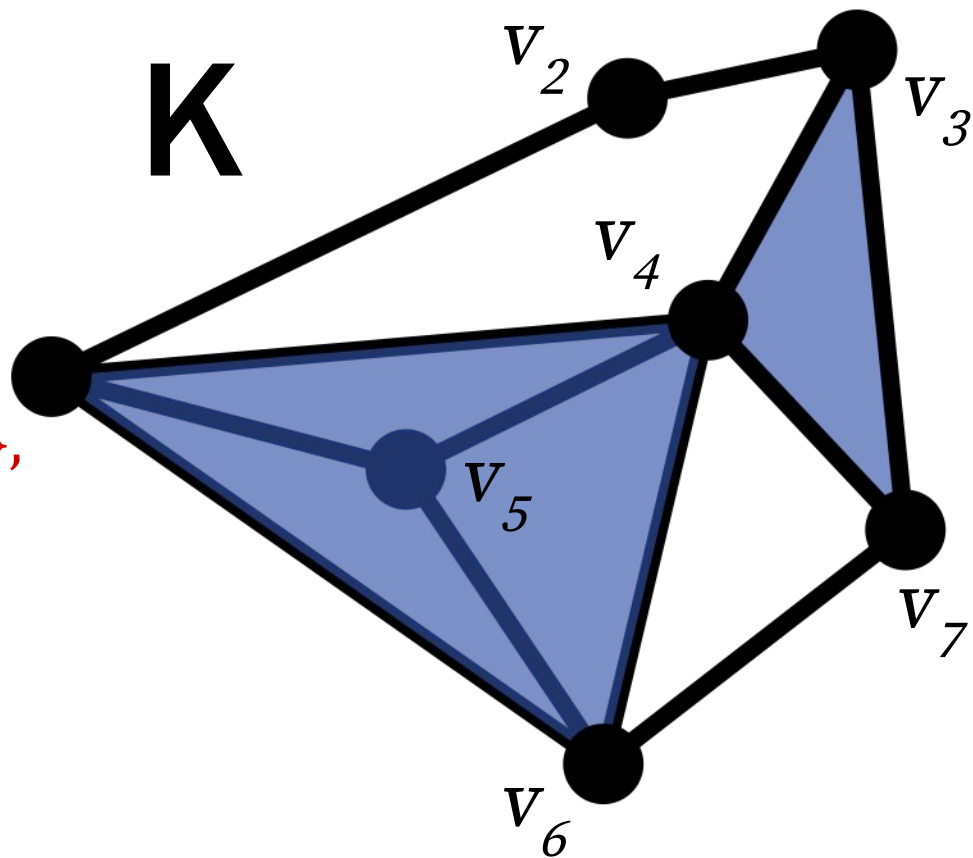
$K =$

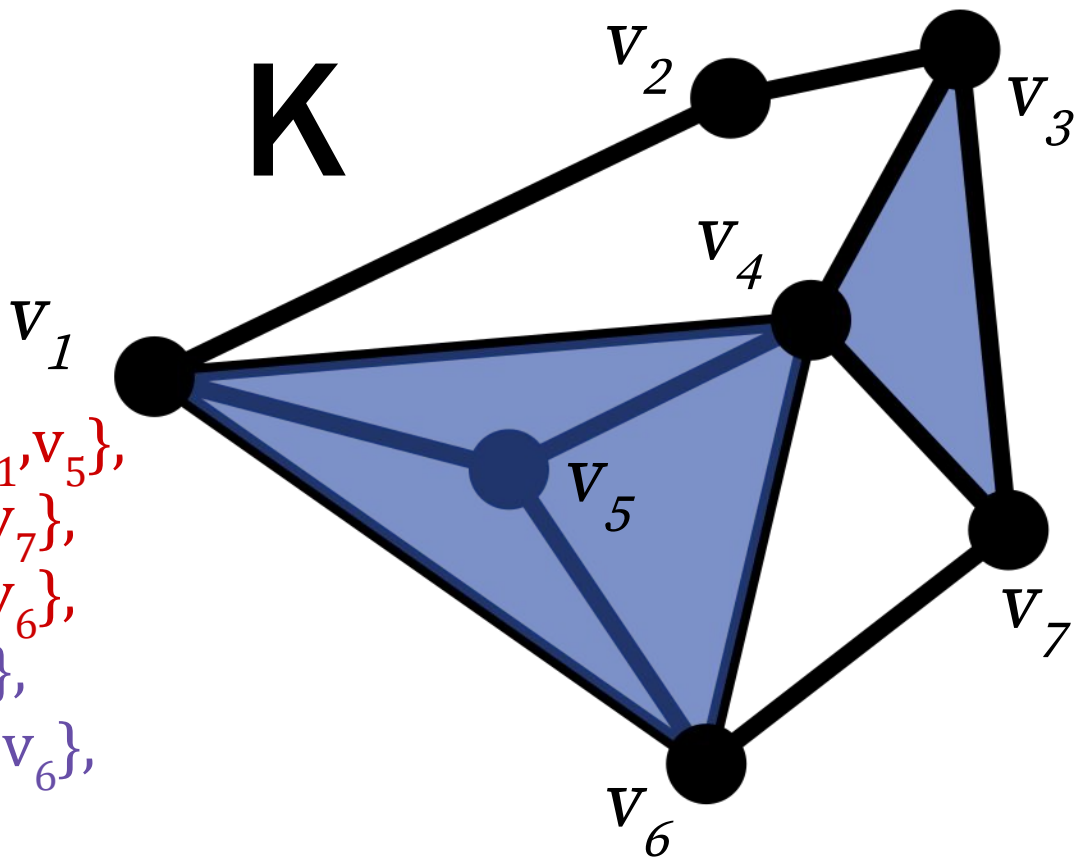
$$\begin{aligned} & \{\{v_1\}, \{v_2\}, \{v_3\}, \{v_4\}, \{v_5\}, \\ & \{v_6\}, \{v_7\}, \{v_1, v_2\}, \{v_1, v_4\}, \{v_1, v_5\}, \\ & \{v_1, v_6\}, \{v_2, v_3\}, \{v_3, v_4\}, \{v_3, v_7\}, \\ & \{v_4, v_5\}, \{v_4, v_6\}, \{v_4, v_7\}, \{v_5, v_6\}, \\ & \{v_6, v_7\}, \{v_1, v_4, v_6\}, \{v_1, v_5, v_6\}, \\ & \{v_1, v_4, v_5\}, \{v_1, v_4, v_6\}, \{v_4, v_5, v_6\}, \\ & \{v_3, v_4, v_7\} \} \end{aligned}$$


$K =$

$\{\{v_1\}, \{v_2\}, \{v_3\}, \{v_4\}, \{v_5\},$
 $\{v_6\}, \{v_7\}, \{v_1, v_2\}, \{v_1, v_4\}, \{v_1, v_5\},$
 $\{v_1, v_6\}, \{v_2, v_3\}, \{v_3, v_4\}, \{v_3, v_7\},$
 $\{v_4, v_5\}, \{v_4, v_6\}, \{v_4, v_7\}, \{v_5, v_6\},$
 $\{v_6, v_7\}, \{v_1, v_4, v_6\}, \{v_1, v_5, v_6\},$
 $\{v_1, v_4, v_5\}, \{v_1, v_4, v_6\}, \{v_4, v_5, v_6\},$
 $\{v_3, v_4, v_7\}\}$

v_1



$$K =$$
$$\begin{aligned} & \{\{v_1\}, \{v_2\}, \{v_3\}, \{v_4\}, \{v_5\}, \\ & \{v_6\}, \{v_7\}, \{v_1, v_2\}, \{v_1, v_4\}, \{v_1, v_5\}, \\ & \{v_1, v_6\}, \{v_2, v_3\}, \{v_3, v_4\}, \{v_3, v_7\}, \\ & \{v_4, v_5\}, \{v_4, v_6\}, \{v_4, v_7\}, \{v_5, v_6\}, \\ & \{v_6, v_7\}, \{v_1, v_4, v_6\}, \{v_1, v_5, v_6\}, \\ & \{v_1, v_4, v_5\}, \{v_1, v_4, v_6\}, \{v_4, v_5, v_6\}, \\ & \{v_3, v_4, v_7\}\} \end{aligned}$$


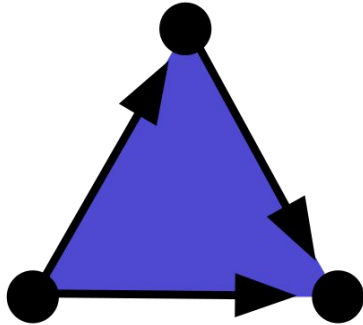
$$\beta_0 = 1, \quad \beta_1 = 2, \quad \beta_2 = 1$$

Directed cliques and why we care.

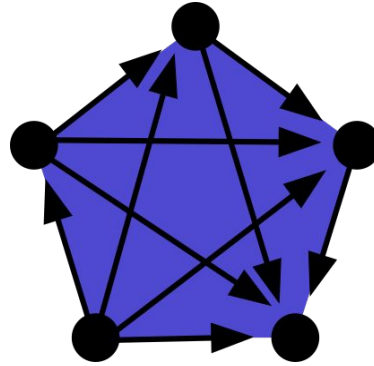
Directed clique: An all-to-all connected subgraph of the network with a clear “source” and “sink”

Directed cliques and why we care.

Directed clique: An all-to-all connected subgraph of the network with a clear “source” and “sink”



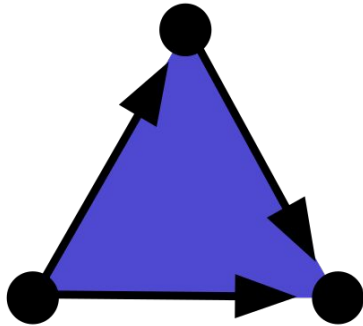
Directed
3-clique



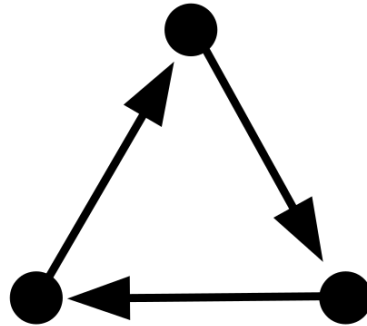
Directed
5-clique

Directed cliques and why we care.

Directed clique: An all-to-all connected subgraph of the network with a clear “source” and “sink”



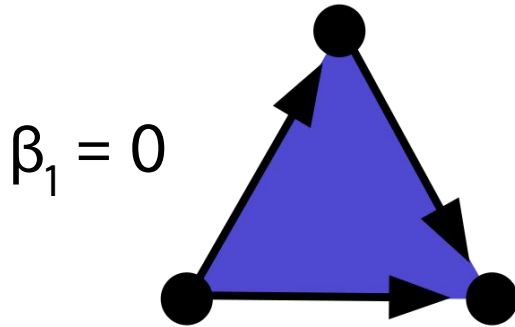
Directed
3-clique



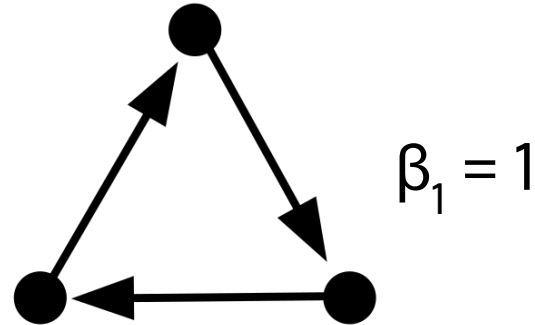
Directed
3-cycle

Directed cliques and why we care.

Directed clique: An all-to-all connected subgraph of the network with a clear “source” and “sink”



Directed
3-clique



Directed
3-cycle

Directed cliques give us a condition to create simplicial complexes which emphasize the direction that information flows in neural networks.

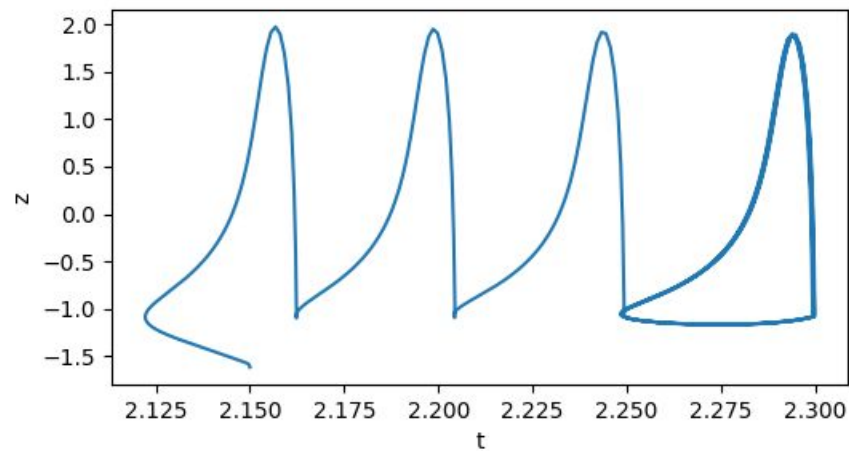
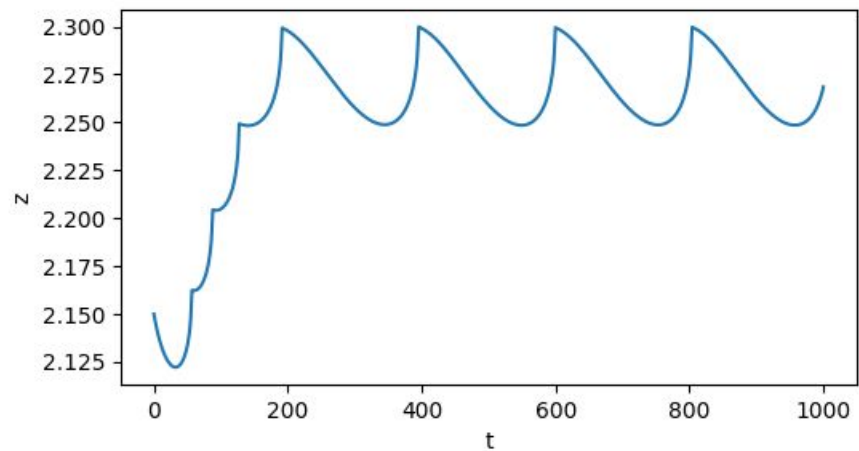
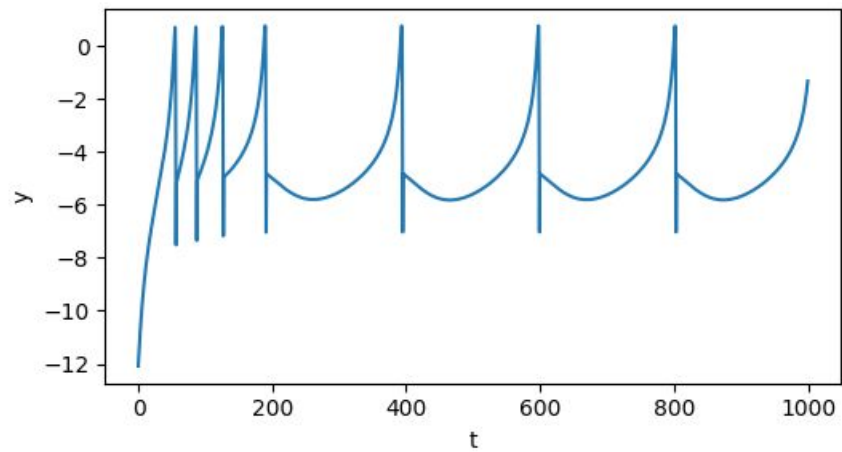
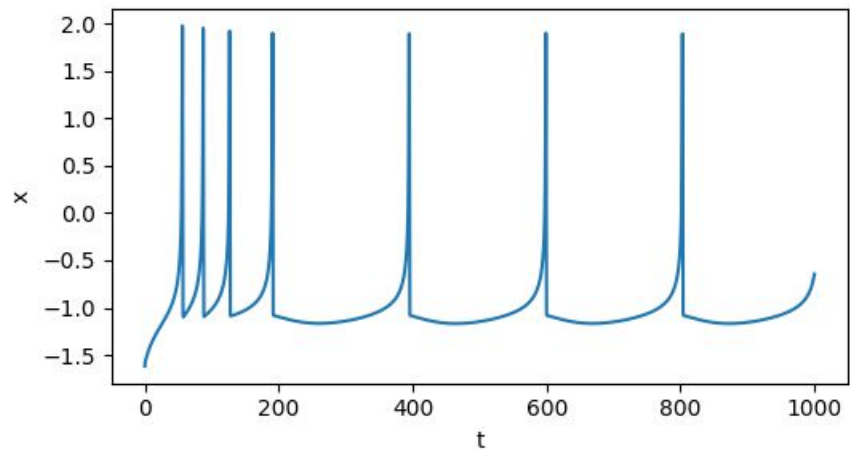
Hindmarsh-Rose Neuron Model

Hindmarsh-Rose Neuron Model

$$\dot{x} = y - ax^3 + bx^2 + I - z$$

$$\dot{y} = c - dx^2 - y$$

$$\dot{z} = r(s(x - x_{rest}) - z)$$



Hindmarsh-Rose Neuron Model

$$\dot{x}_{post} = y_{post} - ax_{post}^3 + bx_{post}^2 + I - z_{post} - gA_{i,j}(x_{post} - V_0) \sum \Gamma(x_{pre})$$

$$\dot{y}_{post} = c - dx_{post}^2 - y_{post}$$

$$\dot{z}_{post} = r(s(x_{post} - x_{rest}) - z_{post})$$

Hindmarsh-Rose Neuron Model

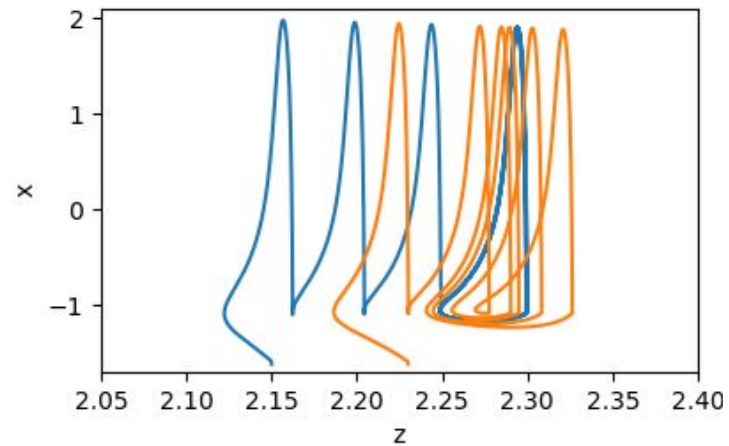
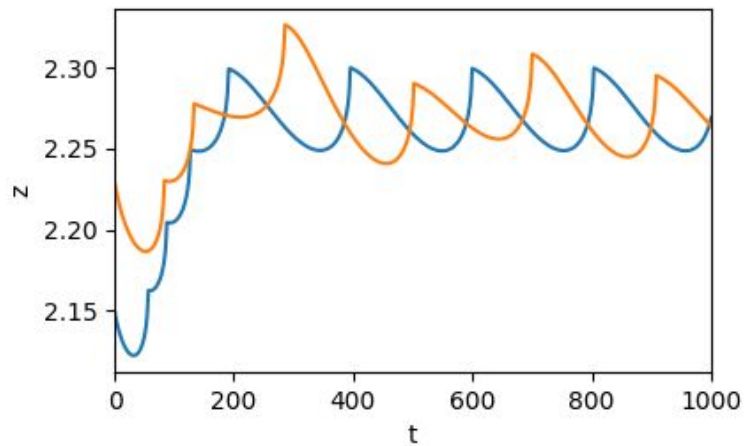
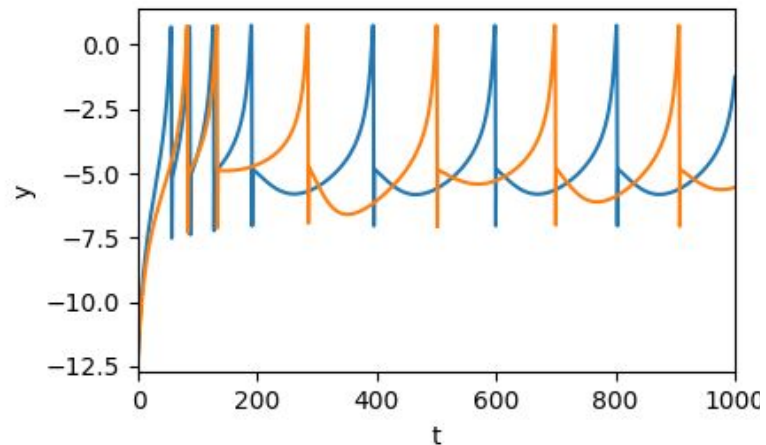
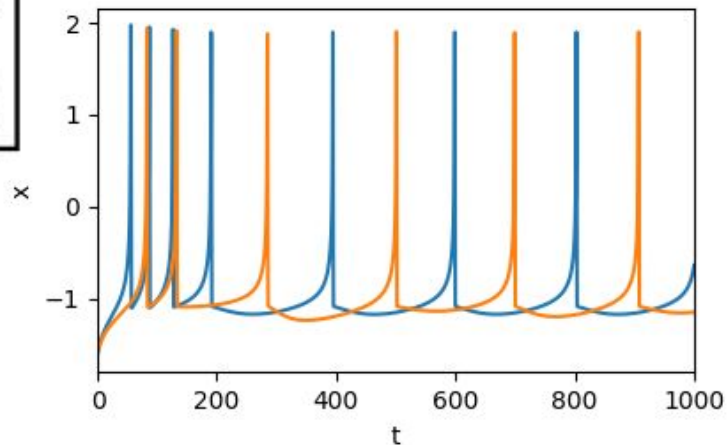
$$\dot{x}_{post} = y_{post} - ax_{post}^3 + bx_{post}^2 + I - z_{post} - \underline{gA_{i,j}(x_{post} - V_0) \sum \Gamma(x_{pre})}$$

$$\dot{y}_{post} = c - dx_{post}^2 - y_{post}$$

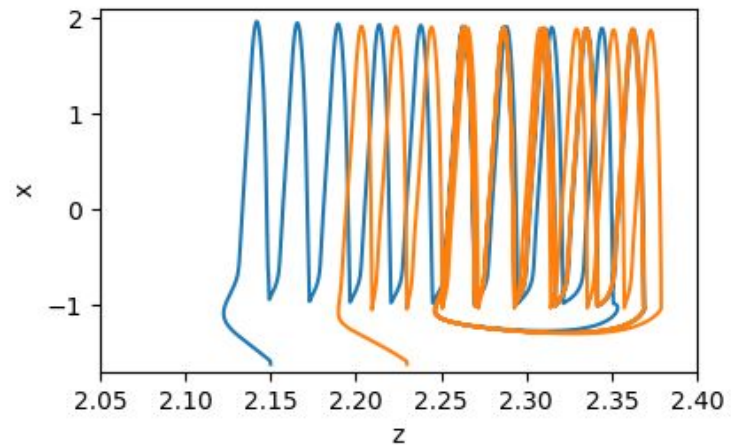
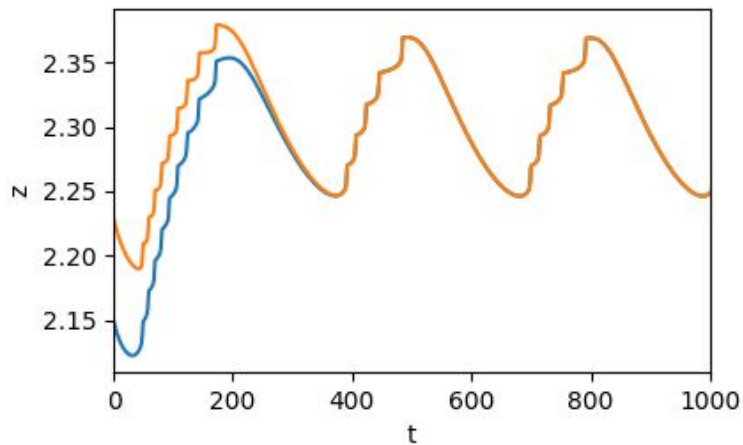
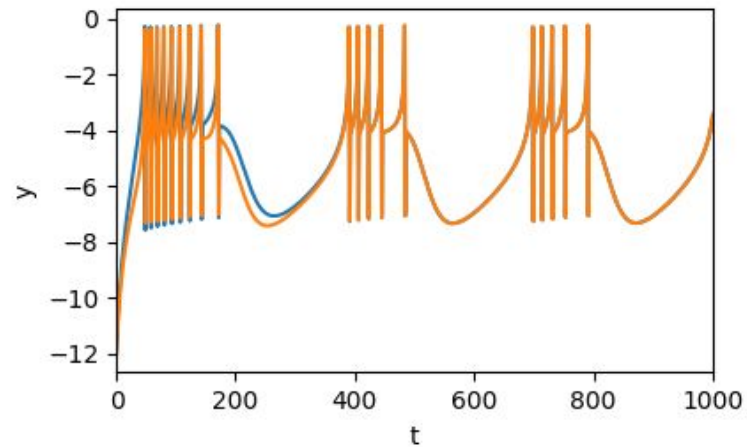
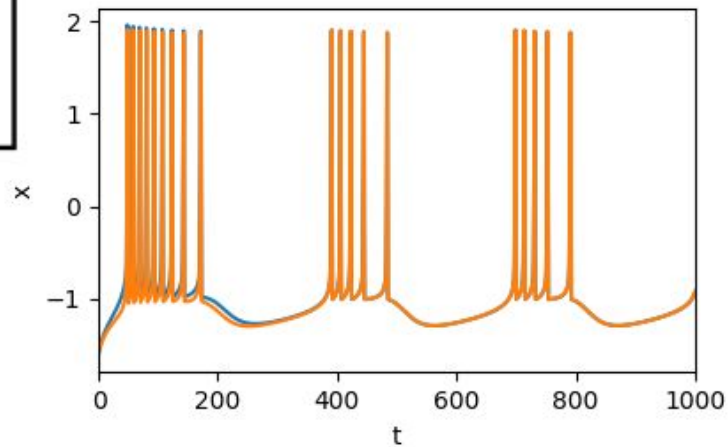
$$\dot{z}_{post} = r(s(x_{post} - x_{rest}) - z_{post})$$

$$\Gamma(x) = \frac{1}{1 + e^{-\lambda(x - \Theta)}}$$

$$A = \begin{bmatrix} 0 & 0 \\ 0 & 0 \end{bmatrix}$$



$$A = \begin{bmatrix} 0 & 1 \\ 1 & 0 \end{bmatrix}$$



Measuring synchrony of a network

For a time t , we can measure the synchrony of a network by looking at the variable z of each neuron because this one dictates when the neuron will spike or burst.

Measuring synchrony of a network

First, transform each neuron's z variable to a number between 0 and π using the following formula, where z_{min} and z_{max} are the extreme value z reaches during the simulation.

$$f(z) = \frac{z - z_{min}}{z_{max} - z_{min}} \cdot \pi$$

Measuring synchrony of a network

Now with each $\theta_i = f(z_i)$ for each neuron i in the network, $\theta_i \in [0, \pi]$ and so we can map each to a vector on the complex unit circle by $v_i = e^{i\theta_i}$

Measuring synchrony of a network

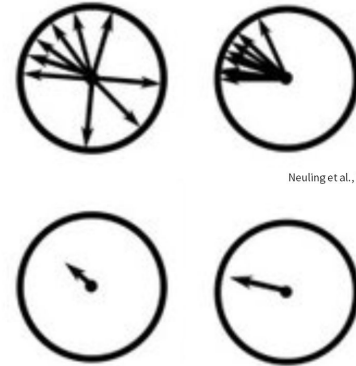
Then when all the vectors are averaged, the resulting vector will have a magnitude close to 1 if the starting vectors all point in the same direction and 0 if they point in different directions.

Measuring synchrony of a network

Then when all the vectors are averaged, the resulting vector will have a magnitude close to 1 if the starting vectors all point in the same direction and 0 if they point in different directions.

We take the resulting vector's magnitude as ϕ which gives us a numerical value of synchrony between 0 and 1 at each timestep in the simulation.

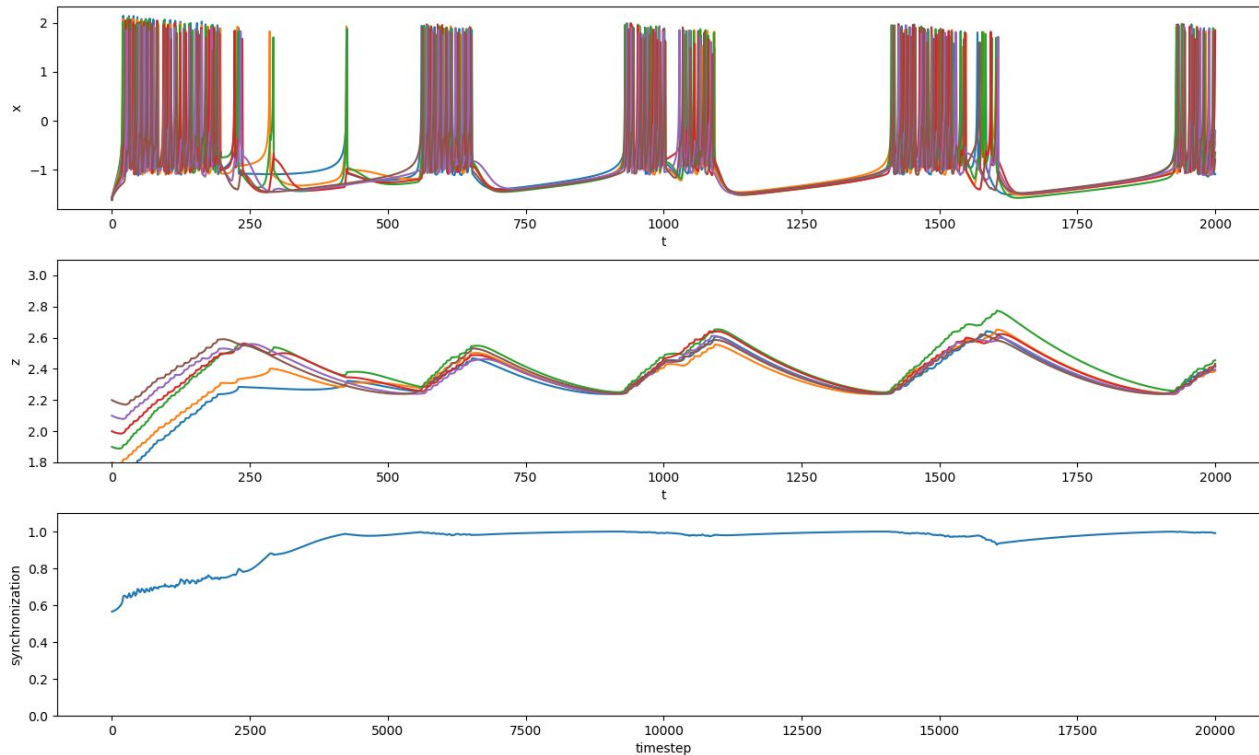
unsynchronized *synchronized*



$\phi = 0.21$

$\phi = 0.93$

Measuring synchrony of a network



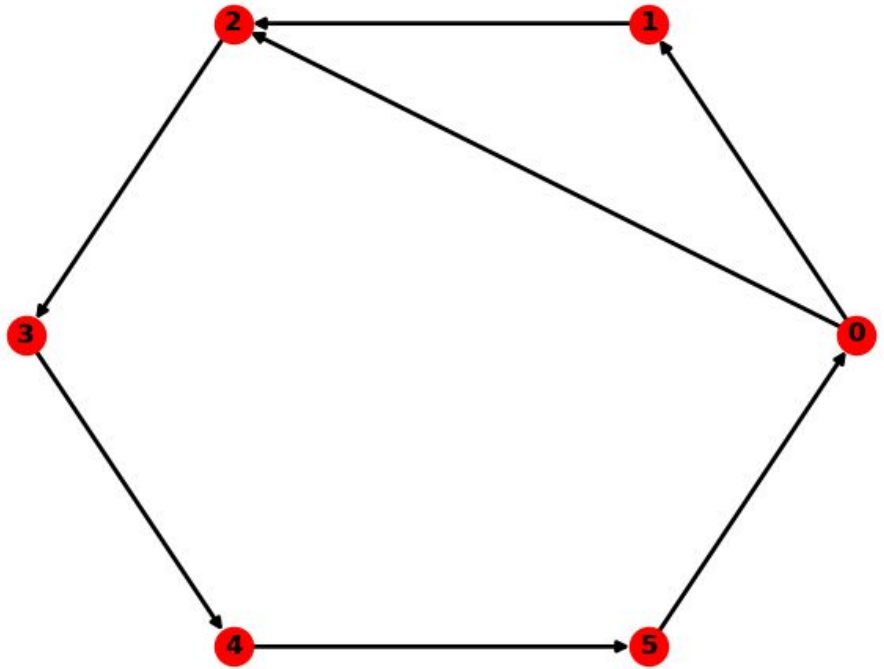
Measuring synchrony of a network extra part

To get another measure of synchrony for a network, we can also take the total variation of our time series of synchronization values. For $t = 0, 1, \dots, T$

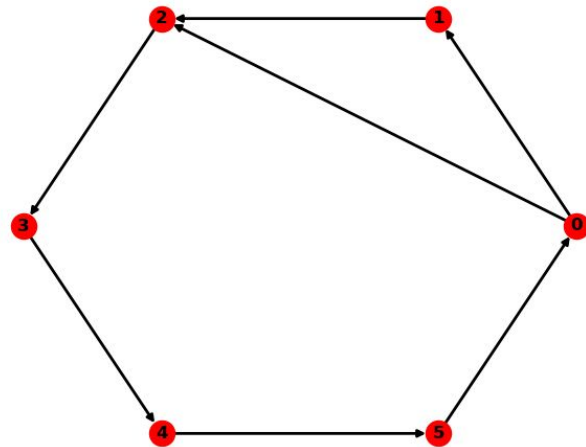
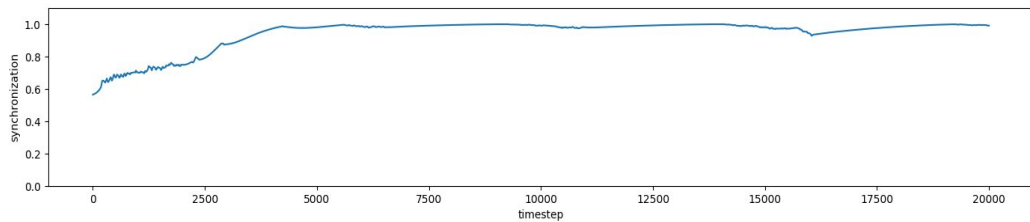
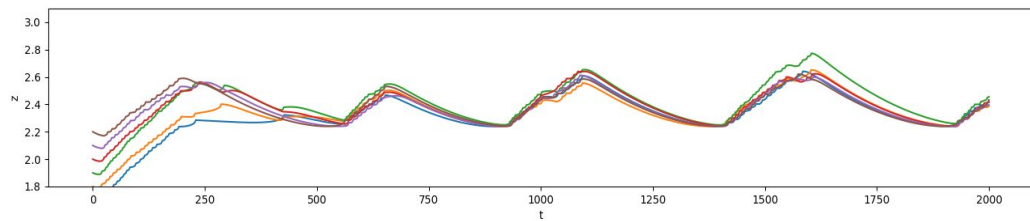
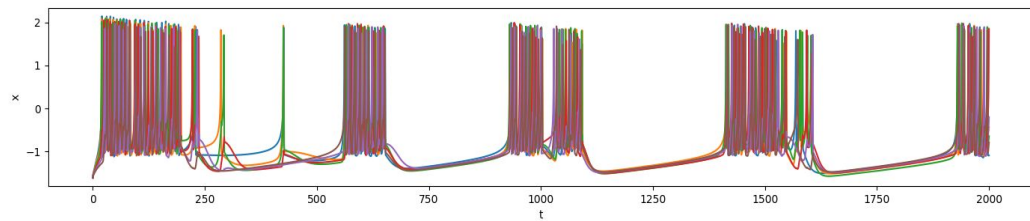
$$\Phi = \sum_{i=0}^T |\phi_{i+1} - \phi_i|$$

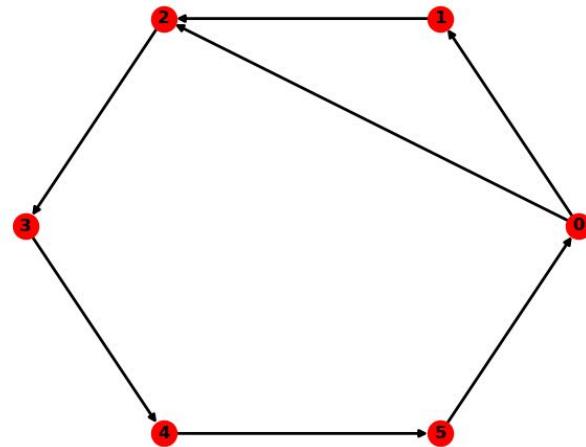
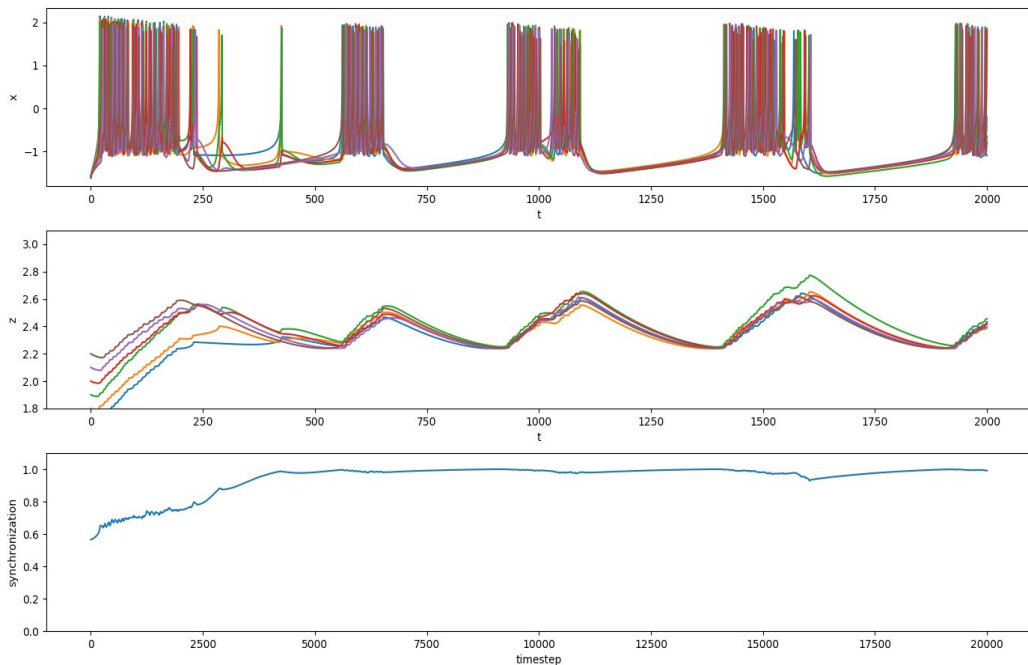
Small networks

We start by running simulations on very small networks because ablating single synapses has a very small chance to have any detectable effect on large networks.



$$\beta_0 = 1 \quad \beta_1 = 1$$

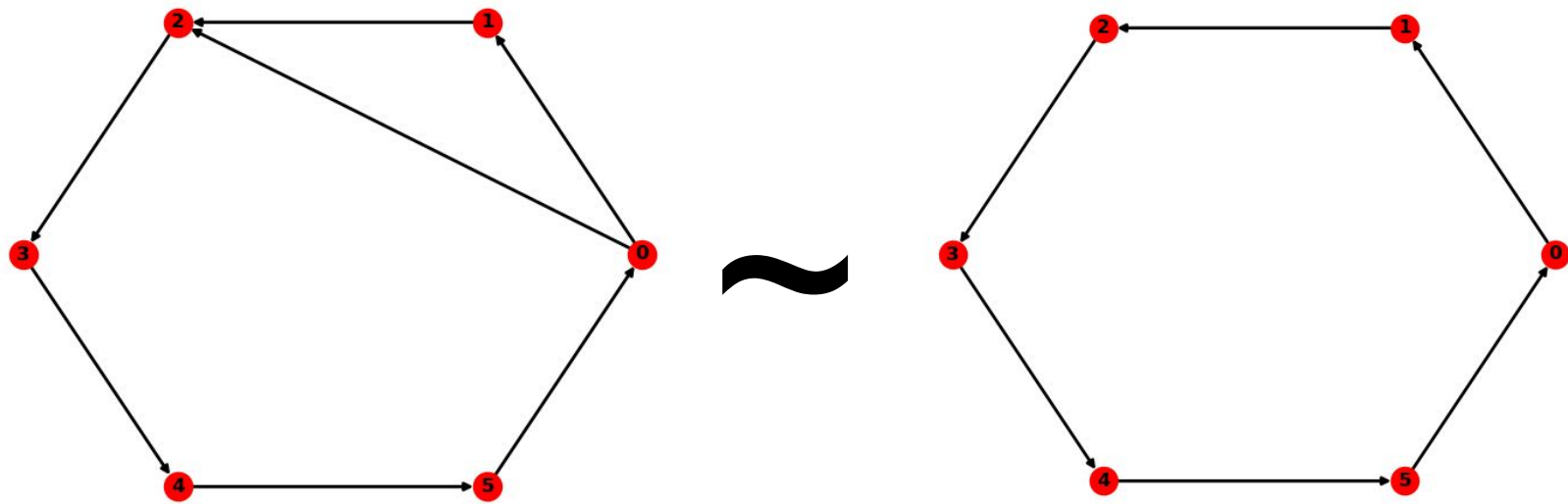


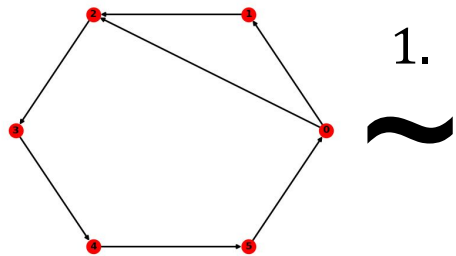


Let's ablate a synapse
that preserves the
homology of this
simplicial complex

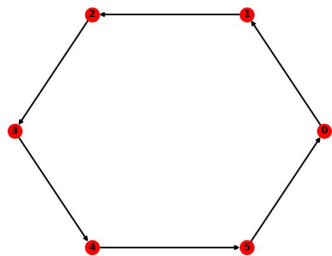
$$\beta_0 = 1$$

$$\beta_1 = 1$$

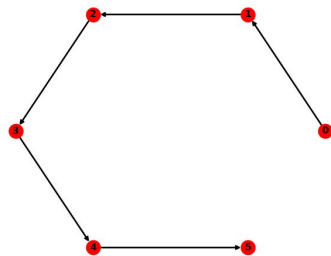




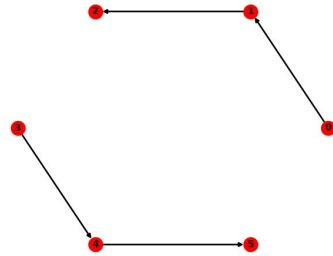
1.



2.



3.



$$\beta_0 = 1$$

$$\beta_1 = 1$$

$$\beta_0 = 1$$

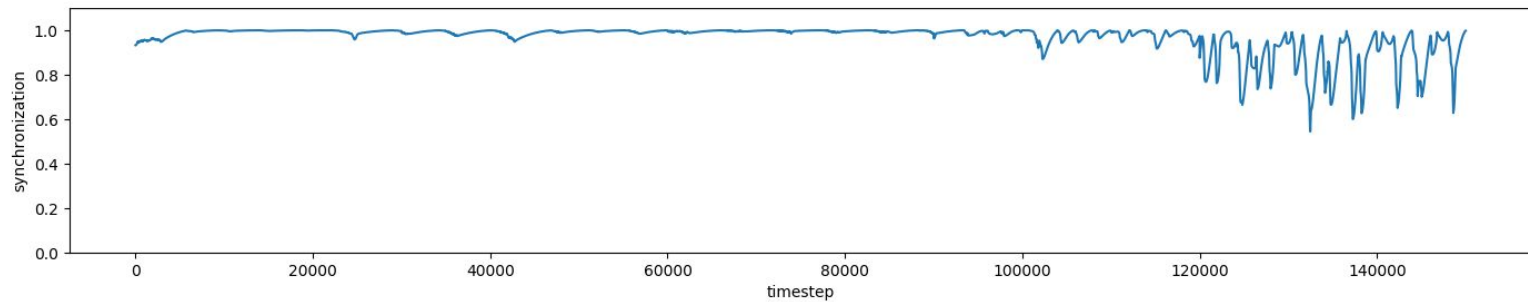
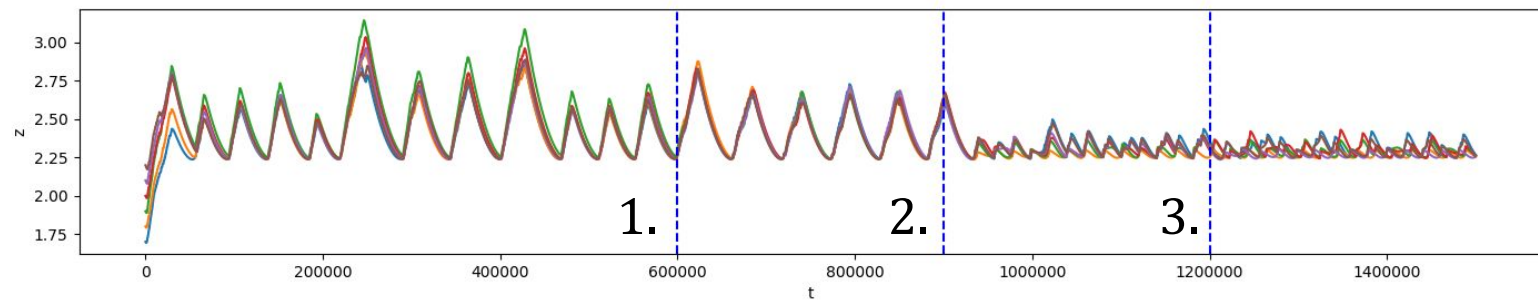
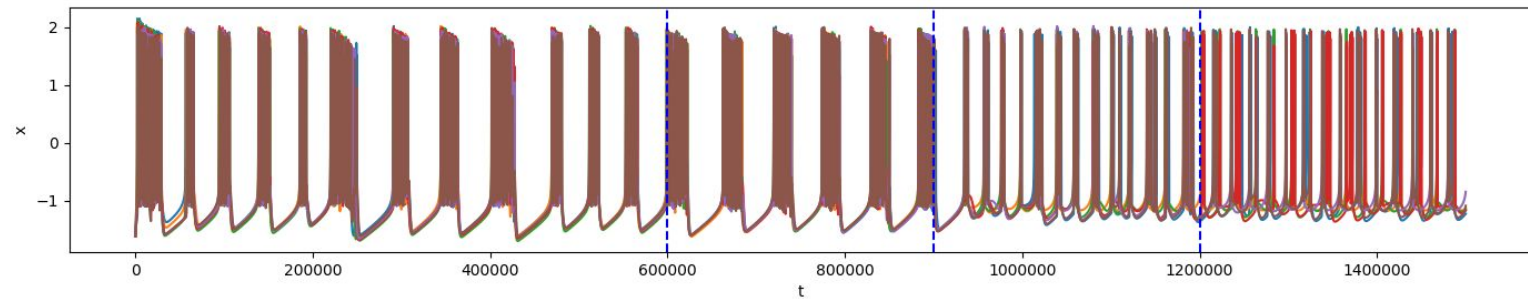
$$\beta_1 = 1$$

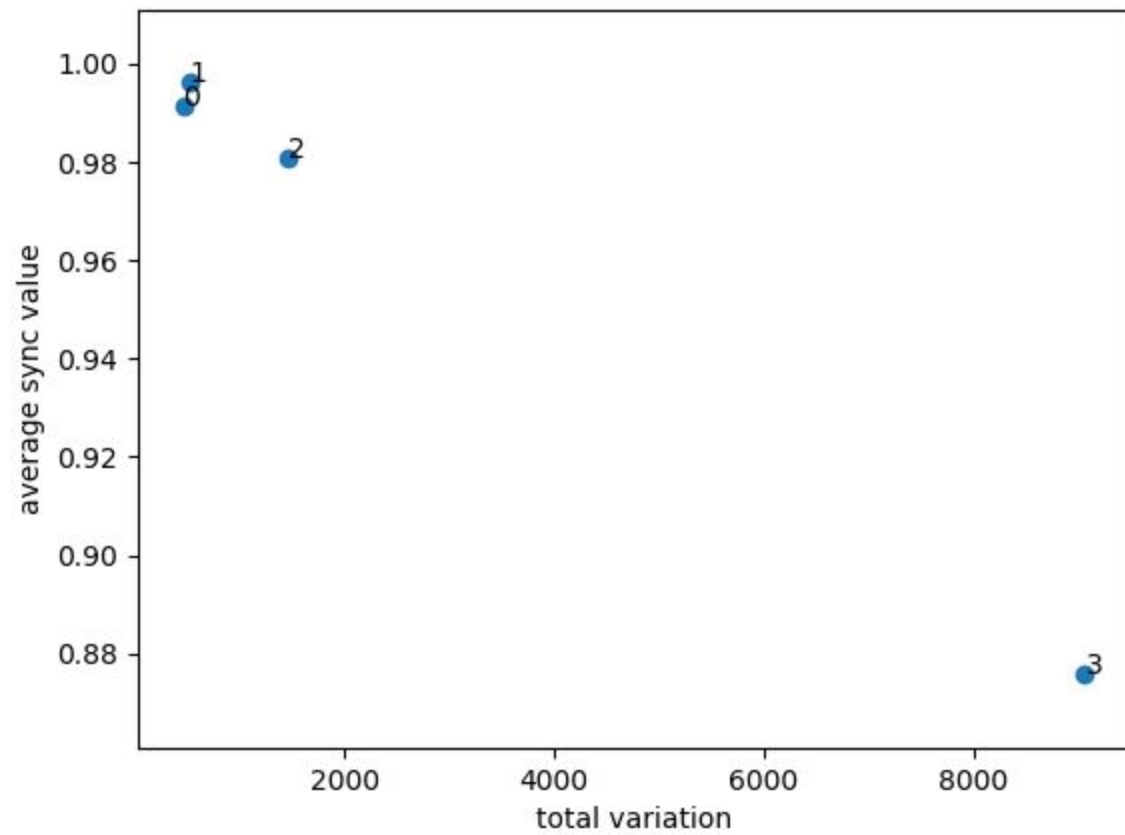
$$\beta_0 = 1$$

$$\beta_1 = 0$$

$$\beta_0 = 2$$

$$\beta_1 = 0$$

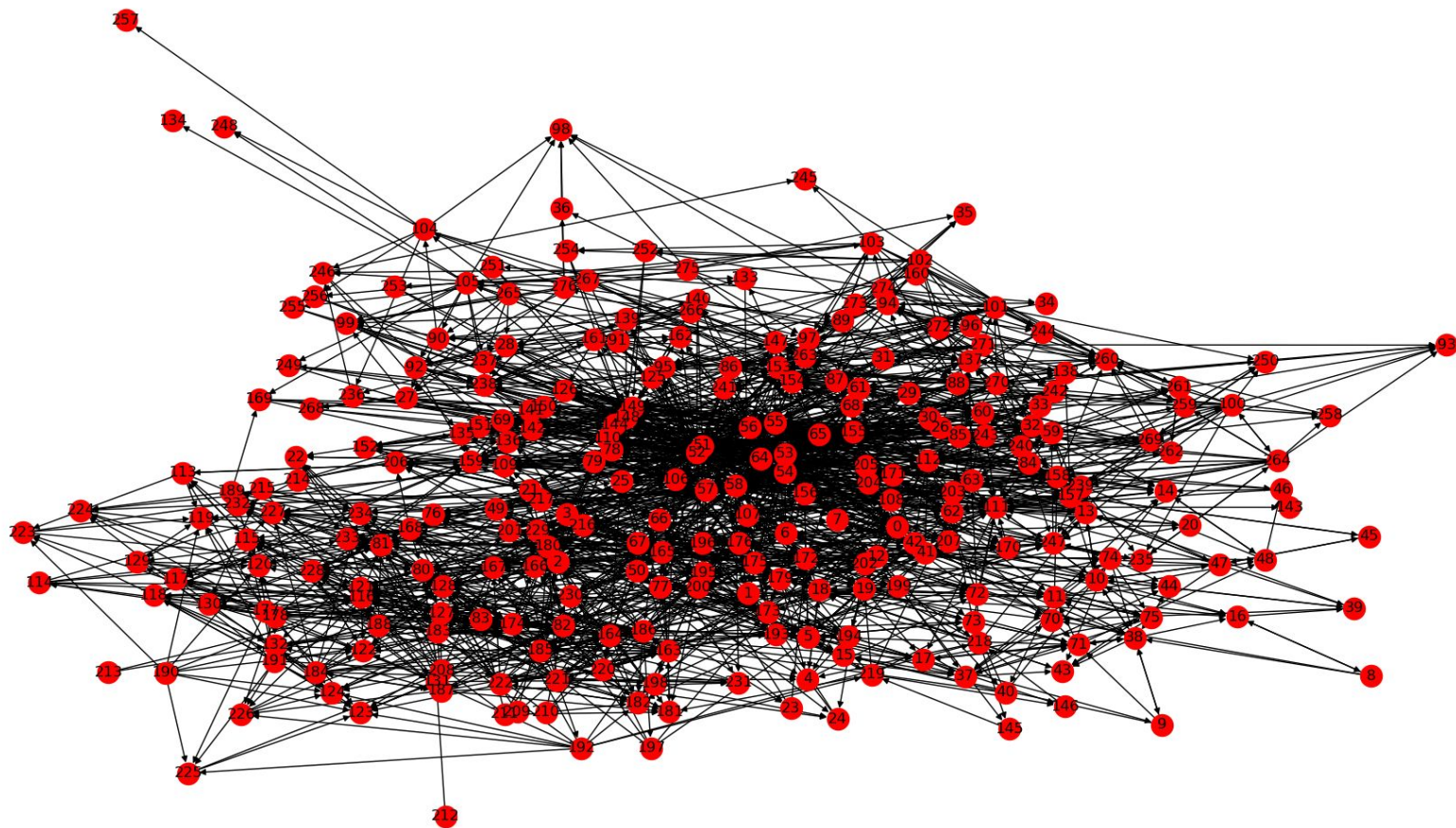




Who does this help?

Computationally, running a neural network simulation is a very intense process.

If we could conclude that “homology matters” for any sized network, then reducing a network to its essential connections while preserving the homology would increase computability while exhibiting the same dynamic behavior.



Thank you!