

[Re] Non-additive coupling enables propagation of synchronous spiking activity in purely random networks

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A reference implementation of

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Introduction

Dendritic non-linearities increase neurons' computation capacity, turning them into complex computing units [4]. However, network studies are usually based on point neuron models that do not incorporate dendrites and their non-linearities. In contrast, the study replicated here [2] uses a simple point-neuron model that contains an effective description of dendrites by a non-linear summation of its excitatory synaptic input. Due to the simplicity of the model, both large-scale parameter exploration of a medium-sized network, as well as an analytical investigation of its properties are feasible.

The original study was based on simulation and analysis code in C and Mathematica, but this code is not publicly available. Here, we replicate the study using the neural simulator Brian 2 [1, 6], a simulator based on the Python language that has become a common choice in computational neuroscience [3]. This simulator offers a good trade-off between flexibility and performance and is therefore a suitable choice for this study of a non-standard neuron model.

Methods

Neural network model

The simulations in the original paper were done in phase representation, but our implementation uses the more standard representation in terms of the neurons' membrane potential. The membrane potential ~~dynamic~~ V_i (of neuron i , measured relative to the resting potential) of neuron i , is described by ~~(a slightly rewritten version equivalent to equation (1) of [2]):~~

$$\tau_m \frac{dV_i(t)}{dt} = -V_i(t) + V_0 + \sigma \left(\sum_{j \in E_i} w_{\text{ex}} \epsilon_j(t - \tau) \right) - \sum_{j \in I_i} w_{\text{in}} \epsilon_j(t - \tau), \quad (1)$$

where $V(t)$ is the membrane voltage at time t , τ_m the membrane time constant, V_0 the displacement of the membrane potential due to a constant external input, w_{ex}

and w_{in} respectively the excitatory and inhibitory weights, τ the transmission delay, and E_i and I_i the indices of neurons that connect with respectively excitatory and inhibitory weights to neuron i . If the membrane potential crosses a threshold Θ , it emits the spike and its membrane potential is reset to its resting potential, i.e. 0 mV. The function ϵ_j is 1 whenever neuron j spikes, and 0 otherwise. [Equation 1 is an alternative, mathematically equivalent formulation of equation \(1\) of \[2\].](#)

The originality of this model lays in a filtering of the excitatory inputs by the function σ to model dendritic non-linearities. The study considers two functions σ (see Fig. 1 of the original paper and insets in Fig. 73). Linear coupling, defined by $\sigma(x) = x$, and non-linear coupling with:

$$\sigma(x) = \begin{cases} x, & \text{if } x \leq V_a \\ V_a + \frac{V_c - V_a}{V_b - V_a}(x - V_a), & \text{if } V_a < x \leq V_b \\ V_c, & \text{otherwise} \end{cases}$$

$$\sigma(x) = \begin{cases} x, & \text{if } x \leq V_a \\ V_a + \frac{V_c - V_a}{V_b - V_a}(x - V_a), & \text{if } V_a < x \leq V_b \\ V_c, & \text{otherwise.} \end{cases} \quad (2)$$

For this non-linear function, the biological interpretation is that below a first threshold V_a , excitatory inputs are propagated passively by the dendrite and sum linearly. Above this threshold, they trigger a dendritic spike, resulting in an amplified somatic voltage. If they are even higher and cross the threshold V_b , they stop increasing the somatic voltage and saturate. Note that this non-linearity only affects spikes received synchronously as has been observed experimentally [5]. With the introduction of this non-linearity, the modelled point neuron becomes an effective model of a neuron with one non-linear dendrite.

The network consists of N neurons, where a directed synaptic connection between a pair of neurons is established with probability p_0 . Each of these connections is taken to be excitatory with probability p_{ex} , and inhibitory otherwise. Note that this connection scheme implies that neurons cannot be separated into groups of excitatory and inhibitory neurons, a neuron typically projects with both excitatory and inhibitory connections to other neurons.

The parameter values used in the original study as well as in our replication are summarized in Table 1.

Parameter	Symbol	Value
Number of neurons	N	1000
Connection probability	p_0	0.3
Probability connection being excitatory	p_{ex}	0.5
Excitatory connection weight	w_{ex}	0.2 mV (varied in Fig. 2)
Inhibitory connection weight	w_{in}	0.2 mV (varied in Fig. 2)
Membrane time constant	τ_m	8 ms
Firing threshold	Θ	16 mV
External input	V₀	17.6 mV (effect on V)
Synaptic delay	τ	5 ms (but see Implementation section)
Threshold for supra-linear summation	V_a	2 mV
Threshold for saturation	V_b	4 mV
Saturation value	V_c	6 mV

Table 1: Parameter values used in the simulations

Implementation and numerical methods

The authors of the original study provided us with the principal C and Mathematica code used in their study. However, this code does not compile and run any more with recent Mathematica versions. Our implementation is not based on this code, but only on the descriptions in the original paper (with minor clarifications provided by the authors where necessary).

The authors of the original study used an event-based simulation scheme, exploiting the linearity of the sub-threshold dynamics. Due to this linearity, the membrane potential can be advanced for arbitrarily large time spans between spikes, and spike times do not have to be bound to any temporal grid. In contrast, the Brian simulator is built on a clock-driven paradigm where time is advanced in discrete steps. This is because the simulator supports a wide range of neural models of and event-driven methods can only be applied to a small number of simple neuron models, and also becomes computationally demanding in large networks with many spikes.

All simulations presented here have been performed with a step size of 0.1 ms and the forward Euler integration algorithm (changing to an exact integration based on the analytical solution of the linear equations did not change the results).

Note that the event-based method used in the original study has important consequences for the specific implementation of non-linear synaptic summation: only spikes that arrive exactly synchronously can sum up non-linearly. This, together with the synaptic delay that is constant over all connections, implies that the non-linearity will only apply to spikes that originate from neurons that spiked perfectly synchronously one synaptic delay earlier; randomly occurring spikes have only an infinitesimally small chance to arrive at the receiving neuron synchronously. When using a clock-driven algorithm, as we do in this replication, this is no longer the case. All neurons that spike within the same time step will deliver their spikes to their target neurons at the same time step, potentially triggering non-linear summation. While the clock-driven method is a less accurate method in general, we would argue that in the context of non-linear summation it is actually closer to the biological phenomenon it models. While dendrites detect synchronous events with sub-millisecond precision [5], this precision is of course finite. With that said, the qualitative results of the original study can be replicated with a clock-driven method, as we will show in the Results section.

For linearly coupled networks, the neuron model is a standard leaky-integrate-and-fire neuron with “delta synapses”, i.e. incoming synaptic events instantaneously affect the membrane potential as soon as they are received. In Brian, this model is described as follows (omitting the code to connect the synapses and set their weights):

```
eq = "dV/dt = -gamma*V + V0 : volt"
G = br2.NeuronGroup(N, model=eq, threshold="V>Theta",
                    reset="V=0*mV", method='euler')
exc_syn = br2.Synapses(G, G, 'w : volt (constant, shared)',
                       on_pre='V += w', delay=5 * ms - dt)
inh_syn = br2.Synapses(G, G, 'w : volt (constant, shared)',
                       on_pre='V -= w', delay=5 * ms - dt)
```

Note that the delay has been set as one time step (dt) less than 5 ms. This is because in Brian, when a neuron receives synaptic input, this input can only trigger a threshold crossing of the receiving neuron in the following time step. To get chains of propagated synchronous spikes every 5 ms as in the original study, we therefore have to reduce the delay by one time step.

~~For non-linearly coupled networks, we have to use a slightly more complicated approach where we sum up all the received excitatory input in a~~ We used a the predefined clip function to implemented non-linear coupling into a network. The function clips a value to a given range, following the syntax

```
clip(value, lower_bound, upper_bound)
```

We employed this functions on the received excitatory inputs stored in a placed holder variable `ve`, ~~apply the non-linear function σ to this input, and~~. The variable corresponds to x in eq. 2. We then reset the variable `ve` back to 0 to prepare for the next time step. In Brian syntax (the definition of the inhibitory synapses is identical to the linearly coupled network):

```
eq = """dV/dt = -gamma*V + V0 : volt
        ve : volt
        """
G = br2.NeuronGroup(N, model=eq, threshold="V>Theta",
                    reset="V=0*mV", method='euler')
exc_syn = br2.Synapses(G, G, 'w : volt (constant, shared)',
                       on_pre='ve += w', delay=5 * ms - dt)
G.run_regularly(''
                V += clip(ve, 0*mV, 2*mV) + clip(2*(ve-2*mV), 0*mV, 4*mV)
                ve = 0*mV
                '', when='after_synapses')
```

The `run_regularly` operation ~~that is run~~ applies the non-linearity to the summed input at every time step ~~to implement the non-linear σ function refers to the predefined clip function which clips the value to a given range, following the syntax `clip(value, lower_bound, upper_bound)`~~.

In addition to random initial conditions, i.e. random initial values of the membrane potential, the simulation includes 1–50 random spikes “initially in transit”. The original article does not go into further details about their implementation, so we decided to use the following: at the beginning of the simulation, a uniformly distributed number between 1 and 50 is drawn to determine the total number of initial spikes. These spikes are “in transit”, meaning that they are modelled as arising from events before the start of the simulation. Since the synaptic delay in the network is 5 ms, we draw the arrival times for these spikes between 0 ms and 5 ms. Consistent with the probability for excitatory connections in the network ($p_{ex} = 0.5$), we take half of these spikes as arising from excitatory and half from inhibitory sources, modelling their effect as described above.

To facilitate simulation, we use the *joblib* library to automatically store results and to distribute computation over processor cores. Several simulation and analysis functions are annotated with a `@mem.cache` decorator, meaning that their input arguments and outputs will be automatically stored to disk and calling the function will return the stored values if available. This feature does not only ease development (e.g. plotting functions can be changed without re-running the underlying simulations), but also allows to interrupt long-running parameter explorations without losing the results obtained so far. The same result could be achieved by manually storing results to disk, but *joblib*’s caching mechanism helps to avoid common errors (e.g. to re-use an old result obtained with an outdated version of the code) while keeping the code readable. To reduce the total simulation time, the grid exploration in Fig. 2, and the analysis of group size evolution in Fig. 7–3 have been distributed over processor cores using *joblib*’s `Parallel` construct. For the simulations presented in Fig. 2, multiple repetitions for the same combination of synaptic weights are distributed over processor cores, for the simulations presented in Fig. 7–3, simulations are evaluated in parallel over group sizes. In both cases, a number of $C - 1$ parallel processes is used, where C is the number of processor cores in the user’s system.

We wanted reproducible simulations despite the use of random connectivity and initial conditions. Setting a single global seed was not sufficient because we used parallel simulations executed in non-deterministic order. Instead, we used a set of pre-determined seeds, one for each figure, and used those “meta-seeds” to generate a set of random seeds for each individual simulation.

For the grid search presented in Fig. 2, we varied inhibitory and excitatory synaptic weights between 0.16 mV and 0.4 mV ~~(given~~. Given that each neuron receives

on average 150 excitatory and 150 inhibitory inputs, this corresponds to a “total weight” between 24 mV and 60 mV, as used on the axes of Fig. 2), ~~using~~. We used a resolution of 100 steps for each variable, comparable to the 96 steps used in the original publication. Each network was simulated 10 times with different initial conditions (synaptic connections and initial membrane potential values), whereas the original publication used 20 simulations per weight combination ~~and additionally added “1–50 random spikes initially in transit”~~.

To numerically derive the evolution of synchronous pulse sizes presented in Fig. 7 3 (for the semi-analytical approach, see the next section), we performed simulations of 56 ms each. During each simulation, we stimulated the network at ~~50ms~~ 50 ms with a synchronous pulse of activity, consisting of between 1 and ~~176~~ 181 neurons (varied in steps of 5 – the original study used group sizes ~~between 1 and 181 in the same range~~ varied in steps of 6). We then observed the number of active neurons at 55 ms, i.e. at one τ after the stimulation. For each stimulation group size, we ran the simulations 50 times with different initial conditions, consistent with the original study. To achieve a better quantitative match with the original study, we applied a correction for the stimulation group size g'_0 and the observed group size g'_1 in the next iteration (for the uncorrected results, see Fig. 7). We refer to the corrected values as \hat{g}_0 and \hat{g}_1 , respectively. These corrections are necessary because of the background activity and our clock-based simulation method and are motivated as follows. The activity of the full network in the absence of stimulation is about 55 kHz (see Fig. 1). Given that our simulation uses a time step of 0.1 ms, this means that at every time step about 5.5 neurons on average are synchronously active. When we stimulate a network with a pulse of g'_0 synchronous neurons, the actual number of synchronously active neurons is therefore higher than this stimulation. Similarly, we have to correct the size of the observed group in the next iteration. In contrast to the simulation strategy of the original publication that could make use of an “infinite precision” to detect synchronous spikes triggered by the stimulation, our simulation will also detect all the synchronous spikes resulting from the background activity. In the corrected values shown in Fig. 3 we therefore used the actually observed synchronous spikes $\hat{g}'_0 \geq g'_0$ at the time of the stimulation, and the number of added spikes due to the stimulation $\hat{g}'_1 = g'_{1stim} - g'_{1no-stim}$, where g'_{1stim} is the number of synchronous spikes we observe in a network with stimulation, and $g'_{1no-stim}$ the number of observed spikes in a network without stimulation. Both networks are simulated with the same random seed and therefore have the same network connectivity and the same initial conditions; the only difference lies in the presence or absence of external synchronous stimulation. Note that the correction for \hat{g}'_1 slightly “overcorrects”, as it discards all spikes that are resulting from background activity, even those that also would have been triggered by the stimulation.

Semi-analytical and analytical methods

We did not attempt to replicate the analytical solution for the evolution of synchronous pulses based on diffusion approximation (blue dots in Fig. 4 of [2]), ~~since the original publication did not give a detailed description of this approach. However, we replicate the semi-analytical work that was described in more detail.~~

To calculate the semi-analytical solutions for Fig. 7 3, based on a numerical estimate of the distribution $P(V)$, we proceeded as follows. First, we ran 2×50 (once for the linear, once for the non-linear network) simulations of 250 ms ~~each of a network with the default parameters but without any~~. Each of these simulations was randomly connected and had random initial conditions, but received no additional synchronous stimulation. ~~We~~ The original study additionally simulated each of the 100 networks 10 times, with the same synaptic connectivity but differing in the initial conditions. We then binned all membrane potential values over all time steps and neurons into 100 bins in the voltage interval $[-\Theta/8, \Theta]$, to obtain an estimate of $P(V)$, the distribution

of the membrane potential. ~~From this distribution~~ Note that the original study did not further specify how $P(V)$ was estimated from the simulation results.

We then calculate the expected synchronous group size after one iteration as in the original publication. To make this calculation clearer, we now describe its derivation in detail. From the distribution $P(V)$ we can estimate the probability F that an additional input ϵ drives the neuron over its threshold (or equivalently, that the neuron is less than ϵ away from its threshold):

$$F(\epsilon) = P(V > \Theta - \epsilon) = \int_{\Theta - \epsilon}^{\Theta} P(V) dV \quad (3)$$

Let $\epsilon(n_{\text{ex}}, n_{\text{in}})$ be the total input to a neuron that receives n_{ex} excitatory and n_{in} inhibitory spikes, i.e. $\epsilon(n_{\text{ex}}, n_{\text{in}}) = \sigma(n_{\text{ex}}w_{\text{ex}}) + n_{\text{in}}w_{\text{in}}$. Let us assume that in the ~~last~~ previous iteration g_i neurons spiked and that these neurons are refractory now, i.e. that $N - g_i$ neurons are available to spike in this iteration. To calculate the probability for a single (non-refractory) neuron j to spike in response to these g_i neurons, we will first calculate the probability that out of the g_i neurons that spiked, exactly n_{ex} neurons are connected with excitatory connections and n_{in} neurons are connected with inhibitory connections to neuron j . There are $\binom{g_i}{n_{\text{ex}} + n_{\text{in}}}$ possibilities to chose $n_{\text{ex}} + n_{\text{in}}$ out of g_i neurons, and $\binom{n_{\text{ex}} + n_{\text{in}}}{n_{\text{ex}}}$ possibilities to divide these neurons into groups of size n_{ex} and n_{in} . This yields the total number of combinations $c(g_i, n_{\text{ex}}, n_{\text{in}})$:

$$c(g_i, n_{\text{ex}}, n_{\text{in}}) = \binom{g_i}{n_{\text{ex}} + n_{\text{in}}} \binom{n_{\text{ex}} + n_{\text{in}}}{n_{\text{ex}}} = \frac{g_i!}{n_{\text{ex}}!n_{\text{in}}!(g_i - n_{\text{ex}} - n_{\text{in}})!} \quad (4)$$

The probability that these selected n_{ex} neurons are actually connected via excitatory connections to neuron j , that the n_{in} neurons are actually connected via inhibitory connections to neuron j , and that the remaining $g_i - n_{\text{ex}} - n_{\text{in}}$ neurons are *not* connected to neuron j is:

$$p(g_i, n_{\text{ex}}, n_{\text{in}}) = (p_0 p_{\text{ex}})^{n_{\text{ex}}} (p_0 p_{\text{in}})^{n_{\text{in}}} (1 - p_0)^{g_i - n_{\text{ex}} - n_{\text{in}}} \quad (5)$$

Finally, we need to sum these probabilities up over all possible values for n_{ex} and n_{in} , multiplying it with the probability that the generated input elicits a spike. This yields the total probability $P^s(g_i)$ that a neuron j spikes in response to the activity of g_i neurons in the previous iteration:

$$P^s(g_i) = \sum_{n_{\text{ex}}=1}^{g_i} \sum_{n_{\text{in}}=0}^{g_i - n_{\text{ex}}} F(\epsilon(n_{\text{ex}}, n_{\text{in}})) c(g_i, n_{\text{ex}}, n_{\text{in}}) p(g_i, n_{\text{ex}}, n_{\text{in}}) \quad (6)$$

In Fig. 73, we plot the expected value of spiking neurons during the next iteration, ~~i.e. given a certain initial synchronous pulse g'_i :~~

$$E(g_{i+1} | g_i = g'_i) = (N - g'_i) P^s(g'_i) \quad (7)$$

Results

We first replicate a simulation showing the propagation of synchronous spiking activity in networks using linear (Fig. 1A) and non-linear (Fig. 1B) coupling. This figure corresponds to Fig. 2 of the original article. Given that the simulations are based on random synaptic connections and random initial conditions, we do not expect exact replication of spike times, but the qualitative features of the simulation are faithfully replicated: synchronous activity quickly dies out and disappears in the background activity for linearly coupled networks (Fig. 1A) but is persistently propagated in non-linearly coupled networks (Fig. 1B).

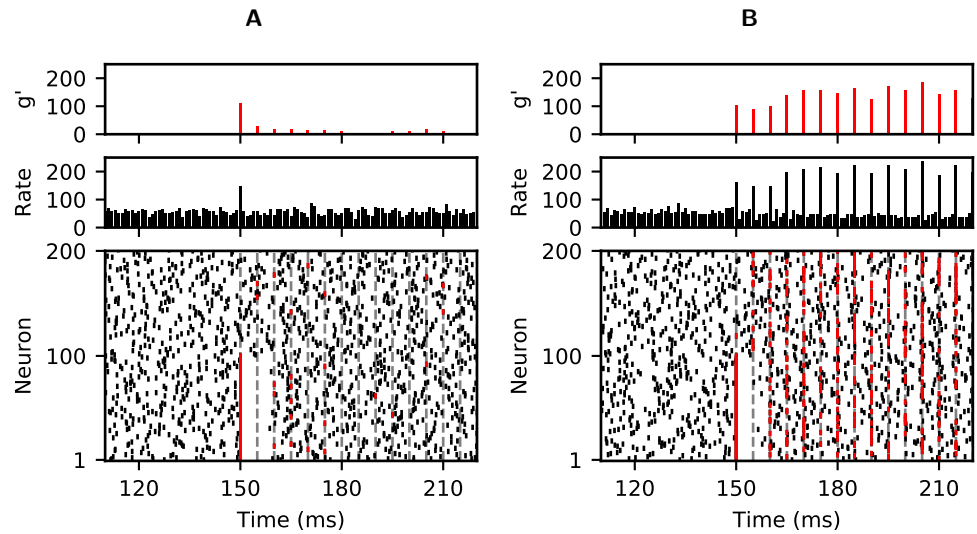


Figure 1: Propagation of synchronous spiking in linearly (A) and non-linearly (B) coupled networks. We initiate a chain of synchronous pulses by applying external supra-threshold inputs to the first 100 neurons at time $t = 150$ ms (red coloured spikes, grey vertical dotted lines indicate times where spikes occur as part of the chain). Top row: total size g' of synchronized groups within the chain. Middle row: network rate over all neurons (rate in kHz, bin size 1 ms). Bottom row: spiking activity of the first 200 neurons in a network of a 1000 neurons versus time. Replication of Fig. 2 in [2] (<https://doi.org/10.1371/journal.pcbi.1002384.g002>).

To confirm the equivalence of our network's behaviour with the original study over a wide range of synaptic connection strength we ran a grid search exploration for linearly (Fig. 2A) and non-linearly (Fig. 2B) coupled networks. Network behaviour is colour-coded as being part of one out of four classes: stable activity, but no persistent propagation (green); stable activity with persistent propagation (blue); unstable activity after synchronous stimulation (yellow); unstable activity before synchronous stimulation (red). For quantitative definitions, see Table 2. We run each combination of connection strengths 10 times, and combine the results to yield a color in RGB notation (values between 0 and 1 for the red, green, and blue component), according to:

$$R = U_1 + U_2; \quad G = E + U_2; \quad B = S, \quad (8)$$

where U_1 , and U_2 stand for the fraction of trials displaying unstable network activity before respectively after onset of propagation, and S and E for the fraction of trials displaying stable background activity with respectively without persistent propagation. We obtained this scheme directly from the authors of the original study, it was only coarsely described in the original publication. Combining individual results in this way does not necessarily lead to easily interpretable colours, but using a different scheme, e.g. colouring according to the class that occurred the most often, leads to visually similar results (~~not shown~~ Fig. 4).

Our replication does not reproduce all the quantitative details of the original plots (in particular at the top of the plots), but. The parameter range where simulations are only unstable after they receive the synchronous stimulation (yellow) is smaller than in the original article. For linear networks, we do observe almost no simulations classified in this category, while the original article has a thin line of such classifications between the stable regime without propagation (green) and the regime where simulations are unstable even without stimulation (red). Similarly, for non-linearly coupled networks the parameter exploration in the original article shows for each inhibitory connection weight a range of excitatory connection weights of about 5 mV where the network is unstable only after the synchronous stimulation (yellow). In contrast, our simulations

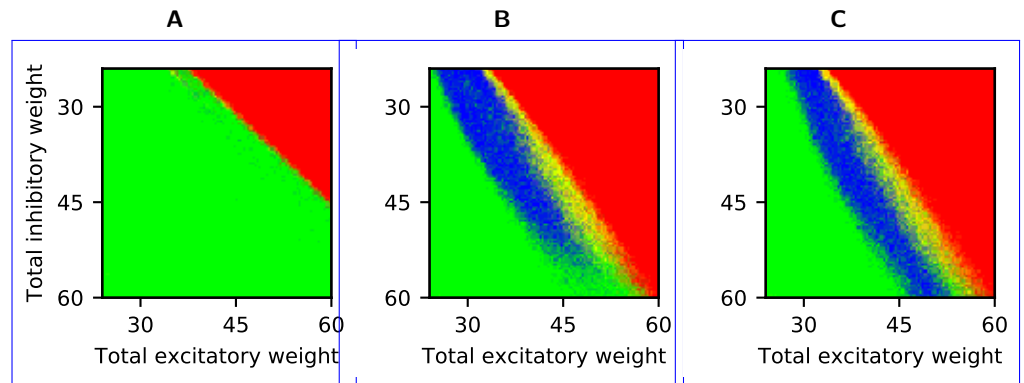


Figure 2: Qualitative network behaviour as a function of excitatory and inhibitory connection strength for linearly (A) and non-linearly (B, C) coupled networks. For each combination, we initiated synchronous activity with 100 (A, B) or 75 (C) neurons and assessed the stability of the temporal evolution. Blue colouring indicates stable propagation of synchrony, red indicates unstable background activity before onset of propagation, yellow indicates unstable background activity after onset of propagation, and green colouring indicates the absence of propagation (see Methods for details).

Replication of Fig. 3 in [2] (<https://doi.org/10.1371/journal.pcbi.1002384.g003>)

Class		Definition
U_1 (red)	Unstable before stimulation	$\exists t < t_{\text{stim}} : a_b(t) \geq 100$
U_2 (yellow)	Unstable after stimulation	$\exists t > t_{\text{stim}} : a_b(t) \geq 100$
E (green)	Stable activity, no propagation	$\forall t : a_b(t) < 100 \wedge g_p < 10$
S (blue)	Stable activity, persistent propagation	$\forall t : a_b(t) < 100 \wedge g_p \geq 10$

Table 2: Classification of network behaviour for Fig. 2. Here, a_b refers to the background activity, the number of neurons active during a 1 ms time bin, ignoring the time steps at $t_p^k = t_{\text{stim}} + k \cdot \tau_s$, i.e. time steps that follow the synchronous stimulation by multiples of the synaptic delay. The variable g_p refers to the number of successful propagations of the synchronous activity, i.e. the number of time steps t_p^k where the number of active neurons exceeds the maximal background activity a_b .

show a much smaller such zone when the inhibitory connection strength is small (Fig. 2B). It seems that these differences mostly stem from the fact that the transition to the regime where simulations are unstable without stimulation (red) happens for lower excitatory and higher inhibitory connection strength in our simulations than in the original article, i.e. that the red zone is shifted towards the bottom left. This is particularly evident for strong excitatory and inhibitory connections in non-linearly coupled networks (bottom right in Fig. 2B and C) which are unstable in our simulations, but stable or only unstable after stimulation in the original publication. Simulating the parameter combination corresponding to the point in the bottom right corner of the plot at different time resolutions (Fig. 6), we find that this difference does indeed seem to be triggered by limited time resolution in the model. The forced synchronisation of all spikes to a grid of 0.1 ms, together with the homogeneity of the synaptic delays, leads to strong oscillations in our simulation (Fig. 6A), which is consequently classified as unstable. When we use a simulation time step of 0.01 ms, these oscillations disappear and the network activity therefore gets classified as “stable without persistent propagation”. Finally, non-linearly coupled networks show a discontinuity for weak inhibitory connection strengths in the original publication. For increasing excitatory connection strengths, network behaviour goes from stable (blue) to unstable after stimulation (yellow) to unstable before stimulation (red), but then get classified again as stable (blue) in a small range of parameters before finally transitioning to unstable activity (yellow and red). We do not observe this in our simulations, but simulations with a finer time resolution (Fig. 5C) indicate that this difference could be caused by the limited time resolution of our simulations as well.

Despite these differences, the qualitative network behaviour looks very similar on a coarse scale. Importantly, it strongly supports the original paper’s main conclusion that non-linearly coupled networks have a large parameter range of stable propagation (i.e., blue-coloured points), while linearly coupled networks do not. The ~~minor~~ differences in our replication ~~could be due to either~~ are most likely due to our use of a clock-based simulation method ~~, or to the slightly different initial conditions (no spikes “with a limited time resolution (see Fig. 5 and Fig. 6). In principle, another reason could be implementation details of the initial “spikes in transit”).~~ **Class Definition** U_1 (red) Unstable before stimulation $\exists t < t_{\text{stim}} : a_b(t) \geq 100$ U_2 (yellow) Unstable after stimulation $\exists t > t_{\text{stim}} : a_b(t) \geq 100$ E (green) Stable activity, no propagation $\forall t : a_b(t) < 100 \wedge g_p < 10$ S (blue) Stable activity, persistent propagation $\forall t : a_b(t) < 100 \wedge g_p \geq 10$ of network behaviour for Fig. 2. Here, a_b refers to the background activity, the number of neurons active during a 1 ms time bin, ignoring the time steps at $t_p^k = t_{\text{stim}} + k \cdot \tau$, i.e. time steps that follow the synchronous stimulation by multiples of the synaptic delay. The variable g_p refers to the number of successful propagations of the synchronous activity, i.e. the number of time steps t_p^k where the number of active neurons exceeds the maximal background activity a_b . ~~that were not fully specified in the original article, but this seems unlikely since completely removing them did not change the results in a noticeable way (not shown).~~

Evolution of synchronous pulses in linearly (A) and non-linearly (B) coupled networks. To numerically estimate the probability distribution, we show occurrences of pulse-sizes g_1' in response to a pulse of size g_0' by grey lines; green squares represent the corresponding mean response group sizes. Replication of Fig. 4 in [2] (–)

Finally, we looked at the evolution of synchronous group sizes for one combination of synaptic weights ($w_{\text{ex}} = w_{\text{in}} = 0.2$ mV) in detail (Fig. 73). The results confirm our previous conclusion that we can faithfully replicate the network behaviour. ~~They match both qualitatively and quantitatively, which shows identical qualitative behaviour;~~ the group size consistently declines with each iteration in linearly coupled networks (Fig. 73A), whereas it increases for intermediate group sizes and declines otherwise in non-linearly coupled networks (Fig. 73B). ~~While the semi-analytic solution (red dotted line) shows a good match to the numerical results (grey lines, means~~

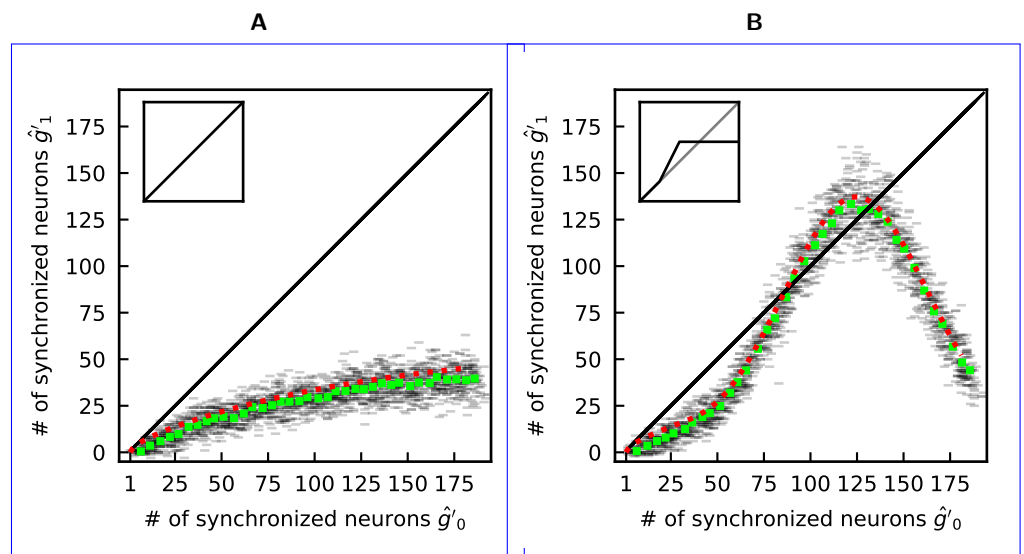


Figure 3: Qualitative network behaviour as a function of excitatory and inhibitory connection strength for linearly (A) and non-linearly (B) coupled networks. For each combination of excitatory and inhibitory connection strength, we initiated synchronous activity with 100 neurons and assessed the stability of the temporal evolution. Blue colouring indicates stable propagation pulse sizes g'_1 in response to a pulse of synchrony, red indicates unstable background activity before onset of propagation, yellow indicates unstable background activity after onset of propagation, and size g'_0 by grey lines; green colouring indicates squares represent the absence of propagation (see Methods for details) corresponding mean response group sizes.

Replication of Fig. 3a and b in [2] (<https://doi.org/10.1371/journal.pcbi.1002384.g004>)

as green dots), it shows some quantitative differences to the semi-analytic solution presented in Fig. 4 of the original article. In particular, our Quantitatively, the semi-analytical solution seems to mostly slightly underestimate the size of the group results also match well. Based on the plotted values in the original publication (Fig. 4 in [2]), the curve based on the semi-analytical solution crosses the identity line at about 85 (G_1) and 135 (G_2) for non-linearly coupled networks, peaking in between for $g'_0 \approx 125$ and $g'_1 \approx 139$. In our simulations, it crosses the line at about 86 and 134, and peaks at $g'_0 \approx 126$ and $g'_1 \approx 137$. As in the original article, our numerical results show slightly lower values for the group size in the next iteration while the, with the biggest difference for values around the peak of the function for non-linearly coupled networks (Fig. 3B). However, we see a quantitative difference for low initial group sizes. In these simulations, the original publication shows a close match between the numerical results and the semi-analytical solution presented in the original article seems to slightly overestimate it. In our simulation, the semi-analytical solution overestimates the numerical one. Be reminded that to obtain this good quantitative match, we had to apply a correction to the numerical results, as explained in the implementation section (for the uncorrected values, see Fig. 7).

Despite the small quantitative differences, our work strengthens the original result by demonstrating a qualitative difference between linearly coupled and non-linearly coupled networks.

Discussion

We managed to qualitatively reproduce the main results of Raoul-Martin Memmesheimer and Marc Timme [2] despite some slight quantitative differences. These difference could be The grid search over parameters showed a higher prevalence of networks with unstable activity before the onset of stimulation compared to the original study. In the analysis of the evolution of synchronous pulses of different sizes, our results need a correction for the use of a clock-based simulation method to show a good quantitative match with the original results. However, a marked difference remains for small initial group sizes. The differences we see are most likely due to our use of a clock-based instead of event-driven simulations, as it forces all activity to a temporal grid of finite precision. We tried However, the main qualitative finding—non-linearly coupled networks have a large parameter range supporting persistent propagation of synchronous activity whereas linearly coupled networks do not—remains if we use finer (0.05 ms) and/or coarser (0.2 ms, 0.5 ms) temporal resolutions and obtained qualitatively the same results (data not shown). Another potential source for the differences is that our initial conditions do not include spikes “in transit” at the start of the simulation, which might further contribute to the quantitative differences we see.

The reusable code we provide with this work paves the way to future studies of networks of non-linearly coupled integrate-and-fire neurons. It gives researchers the means to verify that the original study’s conclusion hold under variations of its model assumptions, such as the details of the connectivity scheme or the synaptic model. This article has already provided a first such verification, demonstrating that the original studies main findings do not critically depend on the event-based method and the entailing “perfect time resolution”. Finally, we hope that the provided code can also serve as a basis for future studies of functional paradigms going beyond the mere propagation of synchronous activity.

Acknowledgements

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Supplementary Figures

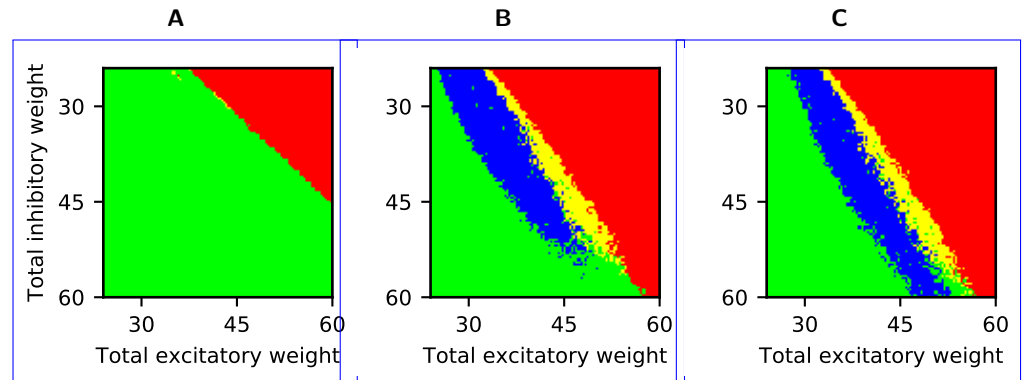


Figure 4: Most-often observed qualitative network behaviour as a function of excitatory and inhibitory connection strength All conventions as in Fig. 2, but points are coloured according the observed qualitative behaviour that occurred the most often among the 10 independent simulations for given connection strengths.

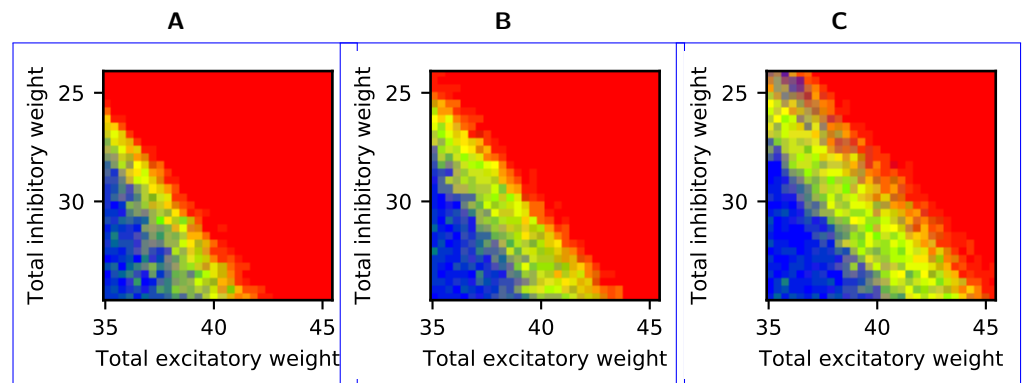


Figure 5: Zoom on qualitative network behaviour with different simulation time steps. Color code as in Fig. 2. Results are shown for a smaller range of connection strengths than in Fig. 2, and only for non-linearly coupled networks with an synchronous stimulation of 100 neurons. The total inhibitory connection strength was varied between 24 mV and 34.55 mV, the total excitatory connection strength was varied between 34.91 mV and 45.45 mV, i.e. the parameter range corresponds to a square region at the top of Fig. 2B. Simulations where performed with simulation time steps of 0.1 ms (A; also the value used for all other simulations shown), 0.05 ms (B), and 0.01 ms (C).

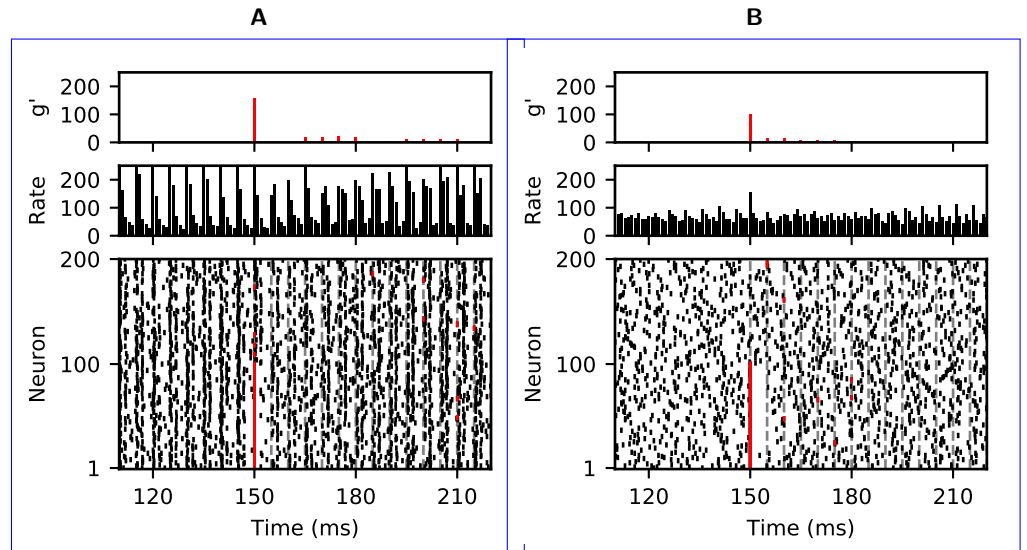


Figure 6: Propagation of synchronous spiking in strongly connected networks when simulated with a time step of 0.1 ms (A), or 0.01 ms (B). All conventions as in Fig. 1, but for strong recurrent connections ($w_{ex} = w_{in} = 0.4$ mV). Both simulated networks use non-linearly coupling, the network shown in panel (B) is simulated with a 10-times finer temporal resolution compared to the other results shown.

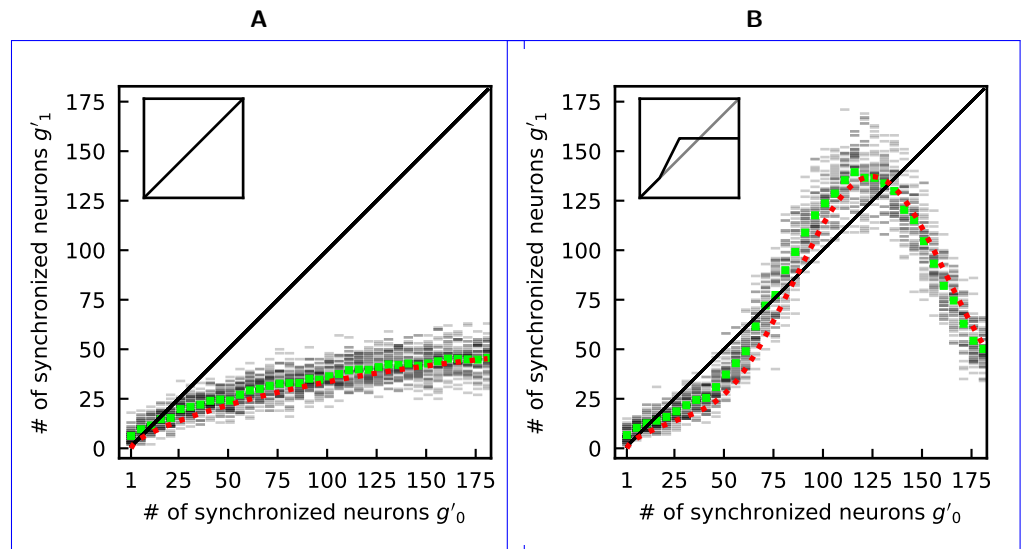


Figure 7: Evolution of synchronous pulses without correction in linearly (A) and non-linearly (B) coupled networks. All conventions as in Fig. 3. The numerical results shown here (gray lines and green squares) are not corrected for the bias introduced by the clock-driven integration method (see text for details).