Supplementary Material for H. E. Plesser (2024): Commentary on Vieth et al (2024)

This notebook provides code, simulations and analysis for the results reported in my *Comment* on the work by Vieth et al (2024). It is available at

https://github.com/heplesser/comment_vieth_code together with scripts used.

```
In [1]: import random
   import math
   import numpy as np
   import matplotlib.pyplot as plt
   import subprocess
   import pandas as pd
   import pickle
   from scipy.stats import norm, binom
   from pathlib import Path
```

The model

As pointed out in the *Comment*, we can limit ourselves to the dynamics of an isolated neuron to understand the discrepancy in simulation results between NEST and other simulators for the *Simple* network studied by Vieth and colleagues.

Ignoring recurrent input and focusing on a single neuron, the membrane potential in this model evolves according to

$$\frac{dv}{dt} = -\frac{v}{\tau} + \frac{I}{C} \tag{1}$$

$$\dot{I} = (\mathcal{U}[I_{\min}, I_{\max}) - I) \, \delta(t - jh)$$
 (2)

When the membrane potential reaches the threshold $V=\theta$, a spike is emitted and the membrane potential reset to 0mV. There is no refractory time. The input current changes in intervals of h=1ms, and this is the time step used for numerical integration.

Parameters used by Vieth et al are

```
In [2]: tau = 10 # ms

C = 1 # pF

I_min = 0 # pA

I_max = 1 # pA

theta = 6 # mV

h = 1 # ms
```

Time-discrete solution

For any ODE solution scheme linear in the membrane potential and input, we can write the update equation as

$$v_{k+1} = \beta v_k + \alpha I_k$$
.

This applies also to higher-order schemes such as RK4.

In particular, we have for exact (exponential) integration

$$lpha = rac{ au}{C} \Big(1 - e^{-h/ au} \Big) \,, \qquad eta = e^{-h/ au}$$

and for forward Euler integration

$$lpha = rac{h}{C}, \qquad eta = 1 - rac{h}{ au} \; .$$

Numerical values of integration coefficients

```
In [3]: alpha_exact, beta_exact = - tau / C * math.expm1(-h/tau), math.exp(-h/tau)
alpha_euler, beta_euler = h / C, 1 - h / tau

In [4]: print(f"Exact: alpha = {alpha_exact:6.4f}, beta = {beta_exact:6.4f}")
    print(f"Euler: alpha = {alpha_euler:6.4f}, beta = {beta_euler:6.4f}")

Exact: alpha = 0.9516, beta = 0.9048
Euler: alpha = 1.0000, beta = 0.9000
```

Actual model parameters for the Euler case

$$\tau_{\text{Euler}} = \frac{h}{1 - \beta_{\text{Exact}}} \tag{3}$$

$$C_{
m Euler} = rac{h}{lpha_{
m Exact}}$$
 (4)

```
In [5]: tau_euler = h / (1 - beta_exact)
C_euler = h / alpha_exact

In [6]: print(f"tau_euler = {tau_euler:5.2f} ms")
    print(f"C_euler = {C_euler:5.2f} pF")

    tau_euler = 10.51 ms
    C_euler = 1.05 pF
```

Mean and standard deviation of the free membrane potential

Mean and standard deviation of the noise input

```
In [7]: mu_I = (I_min + I_max) / 2
sigma_I = math.sqrt((I_max - I_min)**2/12)
print(f"mu = {mu_I:6.4f}pA, sigma = {sigma_I:6.4f}pA")
```

Free membrane potential under exact integration

Compute mean and standard deviation and survival probability S(v), i.e., the probability mass of the free membrane potential distribution above threshold θ .

```
In [8]: V_mean_exact = alpha_exact / (1 - beta_exact) * mu_I
         V_std_exact = alpha_exact / math.sqrt(1 - beta_exact**2) * sigma_I
         S_exact = norm.sf(theta, loc=V_mean_exact, scale=V_std_exact)
         V_mean_euler = alpha_euler / (1 - beta_euler) * mu_I
         V std euler = alpha euler / math.sqrt(1 - beta euler**2) * sigma I
         S_euler = norm.sf(theta, loc=V_mean_exact, scale=V_std_euler)
 In [9]: print(f"Exact <V> = {V_mean_exact:6.4f}mV")
         print(f"Exact <(DV)^2>^1/2 = \{V \text{ std exact:} 6.4f\}mV"\}
         print(f"Exact S(theta) = {S_exact:6.4f}")
         print()
         print(f"Euler <V> = {V mean euler:6.4f}mV")
         print(f"Euler <(DV)^2>^1/2 = {V_std_euler:6.4f}mV")
         print(f"Euler S(theta) = {S_euler:6.4f}")
        Exact <V>
                    = 5.0000 \, \text{mV}
        Exact <(DV)^2>^1/2 = 0.6452mV
        Exact S(theta) = 0.0606
       Euler <V> = 5.0000mV
        Euler <(DV)^2>^1/2 = 0.6623mV
        Euler S(theta) = 0.0655
In [10]: print(f"Excess survival prob for Euler: {(S euler/S exact-1)*100:.2f}%")
```

Excess survival prob for Euler: 8.15%

Numerical simulation

Simultaneous integration using both methods

```
if V_exact[ix] >= 6.0:
    V_exact[ix] = 0
    s_exact.append(t[ix])

V_euler[ix] = beta_euler * V_euler[ix-1] + alpha_euler * I
    if V_euler[ix] >= theta:
        V_euler[ix] = 0
        s_euler.append(t[ix])

return t, V_exact, V_euler, np.array(s_exact), np.array(s_euler)
```

Perform simulations in Python

The code below simulates 10000 individual neurons five times over and reports firing rates reported. Firing rates are computed for the [100, 300] ms interval to obtain the firing rate for the stationary state.

```
In [12]: random.seed(12345)

t_sim = 300
t_min = 100
n_sims = 10000

rates_exact, rates_euler = [], []
for _ in range(5):
    num_exact, num_euler = [], []
    for _ in range(n_sims):
        t, V_exact, V_euler, s_exact, s_euler = sim(t_sim)
            num_exact.append(len(s_exact[s_exact > t_min]))
        num_euler.append(len(s_euler[s_euler > t_min]))

rates_exact.append(sum(num_exact) / n_sims / ( t_sim - t_min ) * 1000)
rates_euler.append(sum(num_euler) / n_sims / ( t_sim - t_min ) * 1000)
```

Resulting firing rates

```
In [13]: print(f'Rate Exact: {np.mean(rates_exact):6.2f} ± {np.std(rates_exact):4.2f}
print(f'Rate Euler: {np.mean(rates_euler):6.2f} ± {np.std(rates_euler):4.2f}

Rate Exact: 9.96 ± 0.04 sp/s
Rate Euler: 10.93 ± 0.04 sp/s
```

Markov chain model

Given v_k , the smallest possible v_{k+1} is obtained for $I_k = I_{\min}$ during the time step and the largest for $I_k = I_{\max}$. In between the maximum and minimum possible values, any value of v_{k+1} is attained with equal probability.

We can thus write the transition probability for the membrane potential as

$$\hat{p}(v'|v) = \begin{cases}
0 & \text{if } v' < \alpha I_{\min} + \beta v \Leftrightarrow v > (v' - \alpha I_{\min})/\beta \\
\frac{1}{\tau(I_{\max} - I_{\min})} & \text{else} \\
0 & \text{if } v' > \alpha I_{\max} + \beta v \Leftrightarrow v < (v' - \alpha I_{\max})/\beta \\
= \frac{1}{\tau(I_{\max} - I_{\min})} \left[\Theta\left(v - \frac{v' - \alpha I_{\min}}{\beta}\right) - \Theta\left(v - \frac{v' - \alpha I_{\max}}{\beta}\right)\right] . (6)$$

To include the effect of the spiking threshold θ , we define

$$p(v'|v) = egin{cases} \hat{p}(0|v) + \int_{ heta}^{\infty} \hat{p}(v'|v) dv' & v' = 0 \ 0 & v' > heta \ \hat{p}(v'|v) & ext{else.} \end{cases}$$

If $q_k(v)$ is the membrane potential distribution at time step k, then the distribution at step k+1 is given by

$$q_{k+1}(v') = \int_{-\infty}^{\infty} p(v'|v) q_k(v) dv \;.$$

Discrete Markov chain

We discretize the transition operator p(v'|v) and the membrane potential distribution $q_k(v)$ evenly along the voltage axis between 0 and θ to solve for the stationary distribution numerically.

```
In [14]: def markov(kind, dv=0.01):
             Input "kind" is either "exact" or "euler".
             Returns the stationary distribution, the transition matrix and the volta
                     : discretization of matrix, in mV
             delta_v : membrane potential lift due to recurrent input in single time
             if kind == 'exact':
                 alpha = alpha exact
                 beta = beta_exact
             elif kind == 'euler':
                 alpha = alpha_euler
                 beta = beta_euler
             else:
                 raise ValueError(f'Unknow kind "{kind}"')
             I_min, I_max = 0, 1
             # Create a voltage axis that will contain all non-vanishing parts
             # of the free membrane potential distribution.
             vv = np.arange(0, max(V_mean_exact, theta) + 6 * V_std_exact, dv)
```

```
# Find location of threshold
ix theta = sum(vv < theta)</pre>
# Create Markov matrix for free membrane potential distribution (normali
# This matrix extends beyond the threshold in order to compute the super
# elements that then are integrated to represent re-injection after firi
# Each element of M represents the probability that if the membrane pote
# [v from, v from+dv), then it will be in [v to, v to+dv) in the next st
# possible voltage step is from v_from+dv -> v_to, while the largest is
# and we calculate the corresponding input currents below.
# M is first filled with relative probabilities and normalized columnwis
M = np.zeros((len(vv), len(vv)))
v from min, v to min = np.meshqrid(vv, vv)
v_from_max = v_from_min + dv
v_{to} = v_{to} + dv
I_from_max_to_min = np.clip((v_to_min - beta * v_from_max) / alpha, I_mi
I_from_min_to_max = np.clip((v_to_max - beta * v_from_min) / alpha, I_mi
M = abs(I_from_min_to_max - I_from_max_to_min)
# Sum superthreshold values representing spiking and add
# to row representing transition to 0 \le v < dv
p_spike = M[ix_theta:, :].sum(axis=0)
M[0, :] += p_spike
# Restrict matrix and voltage axis to subthreshold parts
M = M[:ix theta, :ix theta]
vv = vv[:ix\_theta]
# Normalize matrix columnwise
M /= M.sum(axis=0)
# Compute eigenvalues, largest will come first and should be 1
lam, ev = np.linalq.eiq(M)
assert np.isclose(lam[0], 1), "Largest eigenvector == 1 expected"
# Stationary distribution is normalized first eigenvector
phi = np.abs(ev[:, 0]) / np.sum(np.abs(ev[:, 0]))
return phi, M, vv
```

Evaluate for both rules

```
In [15]: phi_exact, M_exact, vv_exact = markov('exact')
    phi_euler, M_euler, vv_euler = markov('euler')
```

Illustrate evolution of membrane potential

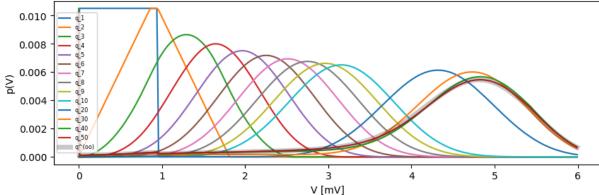
Starting from a $q_0(V) = \delta(V)$, the membrane potential evolves quickly towards a Gaussian-like shape while still too far from the threshold to feel its effects and then converges rather quickly to the stationary distribution.

```
In [16]: plt.figure(figsize=(10, 3))

q = np.zeros_like(vv_exact)
q[0] = 1
for i in range(51):
    q = M_exact @ q
    if i < 10 or (i+1) % 10 == 0:
        plt.plot(vv_exact, q, label=f'q_{i+1}');

plt.plot(vv_exact, phi_exact, 'k-', lw=5, alpha=0.2, label='q^(oo)');
plt.legend(fontsize='xx-small');
plt.xlabel('V [mV]');
plt.ylabel('p(V)');
plt.ylabel('p(V)');
plt.title('Evolution of membrane potential distribution (exact integration)'</pre>
```





Compute firing rates from Markov analysis in absence of recurrent input

The stationary-state firing rate is determined by the stationary membrane potential distribution and the integral term in the first clause of Eq. 15 in the paper. This integral term will have non-zero contributions only for v sufficiently close to the threshold that v can reach θ in a single step of duration h. To extract this part of p(v'=0|v) from the matrix \mathbf{M} , we zero out the first part of the first row of the matrix, i.e., the contribtions of $\hat{q}(0|v)$. Multiply by 1000 to convert from per ms to per s.

Rate Exact: 9.93 sp/s Rate Euler: 10.92 sp/s

Simulations with NEST and Brian2

The following simulations have been performed with

- NEST 3.7
- NESTML 7.0.2
- Brian2 2.6.0
- Code by Vieth et al (2024) from https://github.com/saeedark/Fast-SNN-PymoNNto-rch/tree/main at c9c59f0c6cc2cb3f52875ea1615e4fc9acdd7904.

Vieth et al (2024) used a developer version of NESTML downstream of NESTML 5.3.0. The precise version used (Git hash) is not available from their paper. I use NESTML 7.0.2 here, which requires minor changes to the input specification in the NESTML code from

```
input:
    spikes mV <- spike
    I_e pA <- continuous

to

input:
    spikes <- spike
    I_e pA <- continuous</pre>
```

I also extended Python scripts

Benchmarks/Simples/{brian_LIF,nest_native_LIF,globparams}.py to support disabling connectivity entirely, to disable STDP, to change the Brian integration rule and to record spikes and optionally weights before and after simulation.

```
In [19]: def run_sims(code, n_runs, n_neurons, *args):
             Simulate network by running given code.
             code
                       - name of Python script to run; must write output to tmpdata
                       - number of simulations to run
             n neurons - network size

    these options can be passed

             args
                         no_plot - do not plot at end of simulation
                         no conn - do not create any connections
                         no stdp - disable STDP by setting stdp speed to 0
                         rec_weights - record all weights before and after simulation
                         psc delta — use delta synapses instead of exponential in E
                         exponential_euler - if given, use this method instead of eul
             Returns list of lists of spike times and neuron ids and optionally, init
             if code.startswith('nest_'):
```

```
fb = f"nest native LIF {n neurons} Conn {'no conn' not in args} STDF
             else:
                 EE = 'exponential_euler'
                 fb = f"brian_LIF_Delta_{'psc_delta' in args}_{EE if EE in args else
             base = Path('Benchmarks/Simple/')
             cmd = ['python', base / code, str(n_neurons), 'no_plot']
             cmd.extend(args)
             spikes t = []
             spikes qid = []
             w ini = []
             w_post = []
             w src = []
             w_tgt = []
             wrec = []
             for in range(n runs):
                 subprocess.check call(cmd)
                 s = pd.read_csv(next(Path('tmpdata').glob(f'{fb}_spikes.dat')), comm
                 spikes t.append(s.times)
                 spikes gid.append(s.senders)
                 if 'rec_weights' in args:
                     w = pd.read_csv(next(Path('tmpdata').glob(f'{fb}_weights.dat')),
                     w ini.append(w.w ini)
                     w post.append(w.w post)
                     if code.startswith('nest '):
                         w_src.append(w.source)
                         w tqt.append(w.target)
                         wr = pd.read_csv(next(Path('tmpdata').glob(f'{fb}_wr.dat')),
                         wrec.append(wr)
             return {'spikes_t': spikes_t, 'spikes_id': spikes_gid, 'w_ini': w_ini,
                      'w src': w src, 'w tgt': w tgt, 'wrec': wrec}
In [20]: def run_simset(n_runs, n_size, *args):
             Run set of simulations for NEST, Brian with Euler and Exponential Euler,
             Same parameters as for run_sims(), except that args shall not contain ex
             Returns dictionary of results and writes results to data in pickled form
             nest = run_sims('nest_native_LIF.py', n_runs, n_size, *args)
             brian_exp_euler = run_sims('brian_LIF.py', n_runs, n_size, *args)
             brian_exp_expeuler = run_sims('brian_LIF.py', n_runs, n_size, 'exponenti
             brian_delta_euler = run_sims('brian_LIF.py', n_runs, n_size, 'psc_delta'
             brian_delta_expeuler = run_sims('brian_LIF.py', n_runs, n_size, 'psc_del
             res = {'nest': nest,
                     'brian_exp_euler': brian_exp_euler, 'brian_exp_expeuler': brian_e
                     'brian_delta_euler': brian_delta_euler, 'brian_delta_expeuler': b
             fn = f"sim_{n_size}_Conn_{'no_conn' not in args}_STDP_{'no_stdp' not in
             pickle.dump(res, open(Path('data') / fn, 'wb'))
```

Next cell takes long to run, data are loaded form pickle in following cell

```
In [21]: if False:
    run_simset(5, 10000) # full size, complete network
if False:
    run_simset(5, 10000, 'no_stdp') # full size, complete network, no pl
if False:
    run_simset(5, 10000, 'no_conn') # full size, no connections
```

Load simulation results from above

Firing rates from simulations

Compute rates over $[t_{\min}, t_{\max}]$ with t_{\max}, t_{\min} to avoid initial transient and differences between simulators on what gets recorded from last time step.

```
In [23]: def compute_rates(data, t_min=100, t_max=295, n_size=10000):
    mn, std = {}, {}
    for name, res in data.items():
        rates = [len(d[(t_min < d) & (d <= t_max)]) / ((t_max-t_min)/1000) /
        mn[name] = np.mean(rates)
        std[name] = np.std(rates)

    return {'mean': mn, 'std': std}

def print_rates(res):
    for name in res['mean'].keys():
        print(f"Firing rate {name:20}: {res['mean'][name]:5.2f} ± {res['std']</pre>
```

Full model

```
In [24]: rates_full = compute_rates(full)
    print_rates(rates_full)

Firing rate nest : 11.56 ± 0.07 sp/s
    Firing rate brian_exp_euler : 12.86 ± 0.05 sp/s
    Firing rate brian_exp_expeuler : 13.79 ± 0.06 sp/s
    Firing rate brian_delta_euler : 12.62 ± 0.04 sp/s
    Firing rate brian_delta_expeuler: 11.66 ± 0.07 sp/s
```

Model without STDP

```
In [25]: rates_no_stdp = compute_rates(no_stdp)
print_rates(rates_no_stdp)
```

```
Firing rate nest : 11.01 \pm 0.05 sp/s

Firing rate brian_exp_euler : 12.10 \pm 0.04 sp/s

Firing rate brian_exp_expeuler : 11.73 \pm 0.06 sp/s

Firing rate brian_delta_euler : 12.02 \pm 0.06 sp/s

Firing rate brian_delta_expeuler: 10.86 \pm 0.04 sp/s
```

Model without connections

```
In [26]: rates_no_conn = compute_rates(no_conn)
print_rates(rates_no_conn)

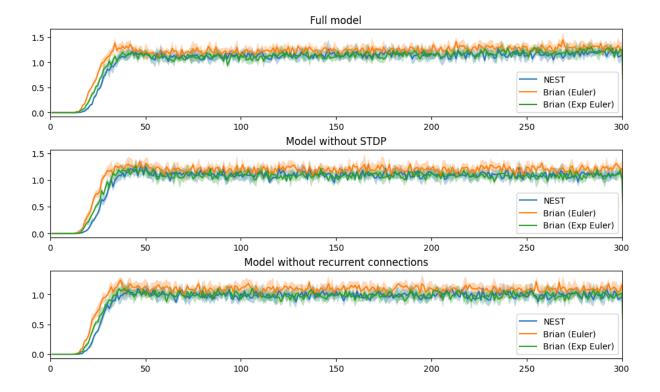
Firing rate nest : 9.92 ± 0.07 sp/s
Firing rate brian_exp_euler : 10.90 ± 0.05 sp/s
Firing rate brian_exp_expeuler : 9.89 ± 0.04 sp/s
Firing rate brian_delta_euler : 10.93 ± 0.04 sp/s
Firing rate brian_delta_expeuler: 9.86 ± 0.05 sp/s
```

Plot results

```
In [27]: def plot_activity(spks, bins, n_size, ax, lbl, between=True):
    cnts = [np.histogram(s, bins)[0] for s in spks]
    ca = np.vstack(cnts) / n_size*100
    mc = ca.mean(axis=0)
    sc = ca.std(axis=0)
    ax.plot(bins[1:], mc, label=f'{lbl}');
    if between:
        ax.fill_between(bins[1:], mc-sc, mc+sc, alpha=0.3);
    ax.set_xlim(0, 300);

In [28]: def plot_set(data, bins, n_size, ax, cases, legend=True, between=True):
    for key, label in cases.items():
        plot_activity(data[key]['spikes_t'], bins, n_size, ax, label, between if legend:
        ax.legend();
```

Spike activity with delta synapses in NEST and Brian



Spike activity with delta synapses in NEST and exponential synapses in Brian

```
In [30]: cases_exp = {'nest': 'NEST', 'brian_exp_euler': 'Brian (Euler)', 'brian_exp_n_size = 10000
    bins = np.arange(0, 305, 1)

plt.figure(figsize=(10, 6), layout='compressed');
    ax = plt.subplot(3, 1, 1)
    plot_set(full, bins, n_size, ax, cases_exp)
    ax.set_title('Full model')

ax = plt.subplot(3, 1, 2)
    plot_set(no_stdp, bins, n_size, ax, cases_exp)
    ax.set_title('Model without STDP')

ax = plt.subplot(3, 1, 3)
    plot_set(no_conn, bins, n_size, ax, cases_exp)
    ax.set_title('Model without recurrent connections');
```

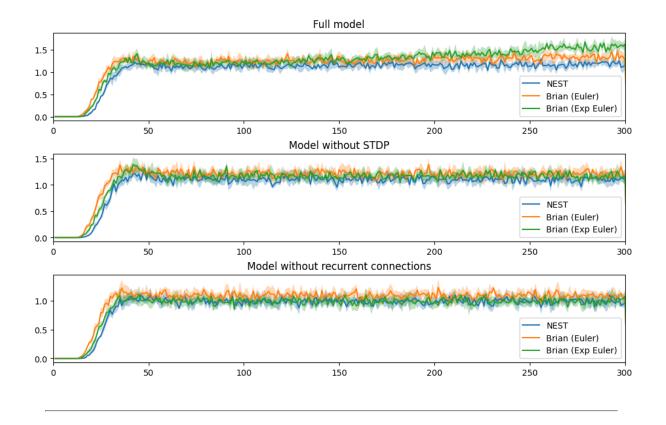


Figure for paper

Perform single simulation over 1000s to obtain membrane potential distribution (Python code from above)

```
In [31]: __, V_exact, V_euler, sp_exact, sp_euler = sim(1000000)

In [32]: hc_eu, be_eu = np.histogram(V_euler, bins=600, density=True);
hc_ex, be_ex = np.histogram(V_exact, bins=600, density=True);
```

Plot functions slightly modified from above for better control over line properties

```
In [33]: def plot_pV(ax):
    ax.stairs(hc_ex, be_ex, label='Exact simul', lw=2, ec=(1, 0.4, 0.), zord
    ax.stairs(hc_eu, be_eu, label='Euler simul', lw=2, ec=(0.18, 0.55, 1.),
    ax.step(vv_exact, phi_exact*100, where='post', label='Exact Markov', lw=
    ax.step(vv_euler, phi_euler*100, where='post', label='Euler Markov', lw=

In [34]: def pplot_activity(spks, bins, n_size, ax, lbl, params):
    cnts = [np.histogram(s, bins)[0] for s in spks]
    ca = np.vstack(cnts) / n_size*100
    mc = ca.mean(axis=0)
    ax.stairs(mc, bins, **params, label=f'{lbl}');
    ax.set_xlim(0, 300);

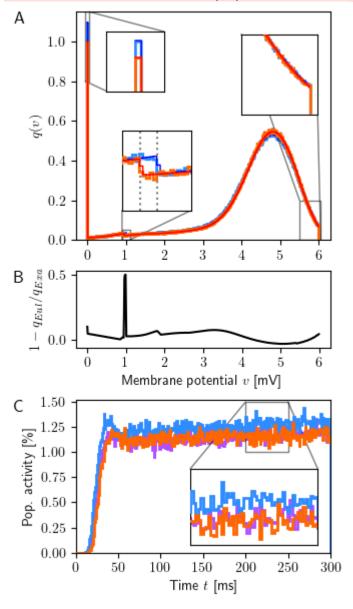
In [35]: def pplot_set(data, bins, n_size, ax, cases, legend=True):
    for key, (label, params) in cases.items():
```

```
pplot_activity(data[key]['spikes_t'], bins, n_size, ax, label, param
             if legend:
                 ax.legend();
In [36]: pcases = {
             'nest': ('NEST', {'lw': 2, 'ec': (1, 0.4, 0.), 'zorder': 100}),
             'brian_delta_euler': ('Brian (Euler)', {'lw': 2, 'ec': (0.18, 0.55, 1.),
              'brian_delta_expeuler': ('Brian (Exp Euler)', {'lw': 2, 'ec': (0.7, 0.3,
In [37]: plt.rc('text', usetex=True);
         fig, axd = plt.subplot_mosaic([['top'],
                                         ['middle'],
                                        ['bottom']],
                                        height ratios=[3, 1, 2],
                                        figsize=(85/25.4, 150/25.4), layout="constrain
         ax = axd['top'];
         plot pV(ax);
         ax.set ylabel('$q(v)$');
         ax.set_xticks(range(7));
         ix1 = ax.inset_axes([0.65, 0.55, 0.3, 0.35], xlim=[5.5, 6.05], ylim=[0, 0.2]
         plot_pV(ix1)
         _, cl1 = ax.indicate_inset_zoom(ix1, edgecolor="black");
         for cl in cl1:
             cl.set_visible(not cl.get_visible())
         ax.text(-0.2, 1, 'A', transform=ax.transAxes, va='top', ha='right', fontsize
         ix2 = ax.inset_axes([0.9, 0.15, 1.8, 0.4], xlim=[0.9, 1.1], ylim=[0, 0.05],
                            transform=ax.transData);
         plot pV(ix2)
         _, cl2 = ax.indicate_inset_zoom(ix2, edgecolor="black");
         ix2.vlines([alpha exact, alpha euler], ymin=0, ymax=0.05, colors='gray', ls=
         ix3 = ax.inset_axes([0.5, 0.75, 1.5, 0.3], xlim=[-0.05, 0.05], ylim=[0.8, 1.5]
                            transform=ax.transData);
         plot pV(ix3)
         _, cl3 = ax.indicate_inset_zoom(ix3, edgecolor="black");
         ax4 = axd['middle']
         ax4.plot(vv_exact, phi_euler/phi_exact-1, c='k');
         ax4.set_xlabel('Membrane potential $v$ [mV]');
         ax4.set ylabel('$1-q {{Eul}}/q {{Exa}}$');
         ax4.set_xticks(range(7));
         ax4.text(-0.2, 1, 'B', transform=ax4.transAxes, va='top', ha='right', fontsi
         ax5 = axd['bottom']
         bins = np.arange(0, 305, 1);
         pplot set(full, bins, 10000, ax5, pcases, legend=False);
         ax5.set_xlabel('Time $t$ [ms]');
         ax5.set_ylabel(r'Pop. activity [\%]');
         ax5.set_yticks(np.arange(0, 1.51, 0.25));
         ax5.set_xlim(0, 300);
         ax5.text(-0.2, 1, 'C', transform=ax5.transAxes, va='top', ha='right', fontsi
```

```
ix5= ax5.inset_axes([0.45, 0.05, 0.5, 0.5], xticks=[], yticks=[]);
pplot_set(full, bins, 10000, ix5, pcases, legend=False);
ix5.set_xlim(200, 250);
ix5.set_ylim(1, 1.5);
_, cl5 = ax5.indicate_inset_zoom(ix5, edgecolor="black");

plt.savefig('figures/markov_analysis.eps', bbox_inches='tight');
```

The PostScript backend does not support transparency; partially transparent artists will be rendered opaque.

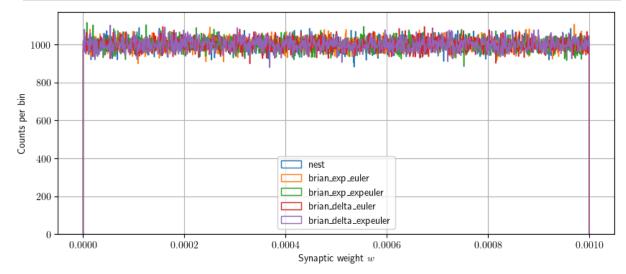


STDP analysis

Based on 1000 neuron network to avoid overly large weight matrices, single run only

```
In [38]: if False:
    run_simset(1, 1000, 'rec_weights') # smaller network, record weights
In [39]: small = pickle.load(open('data/sim_1000_Conn_True_STDP_True.pkl', 'rb'))
```

Initial membrane potential distribution



Final weight distribution

Expected weight distribution

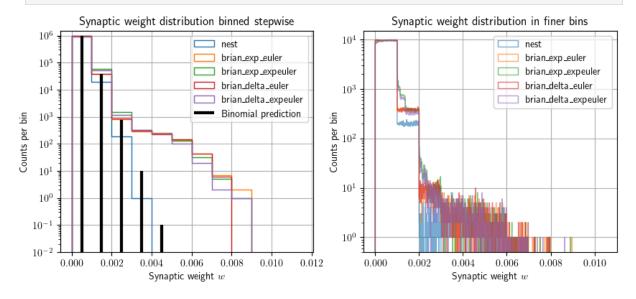
- A synapse weight is increased by 0.001 if the pre- and post-synaptic neuron fire in subsequent time steps
- Since firing is rare (ISI approx 100 ms), we can treat pre- and post-synaptic spikes as independent
- The probability that a specific synapse is increased in a given time step is

$$p_{ ext{STDP}} = \left(rac{h}{r^{(\infty)}}
ight)^2$$

- ullet In 300 ms simulation time there are $n_{
 m STDP}=299$ time steps with opportunity for plasticity
- Thus, the expected distribution of weights after a 300 ms simulation is given by the following binomial distribution

```
In [41]: stdp_steps = 299
p_spike = rates_full['mean']['nest'] / h / 1000 # 1000 for ms->s
p_stdp = p_spike**2
expected_weight_dist = binom(stdp_steps, p_stdp)
```

```
In [42]: wbins = np.arange(0, 0.0105, 0.001)
         plt.figure(figsize=(10, 4));
         plt.subplot(1, 2, 1)
         for name, data in small.items():
             data['w_post'][0].hist(bins=wbins, histtype='step', label=name, log=True
         x_{stdp} = np.arange(0, 11.4, 1)
         plt.gca().vlines(0.0005 + x_stdp*0.001, 0, expected_weight_dist.pmf(x_stdp)*
                           colors='k', linestyles='-', lw=3, label='Binomial prediction
         plt.ylim(1e-2, 1.5e6)
         plt.legend();
         plt.xlabel('Synaptic weight $w$');
         plt.ylabel('Counts per bin');
         plt.title('Synaptic weight distribution binned stepwise');
         plt.subplot(1, 2, 2)
         for name, data in small.items():
             data['w post'][0].hist(bins=np.arange(0, 0.0105, 0.00001), histtype='ste
         plt.legend();
         plt.ylim(5e-1, 1.5e4)
         plt.xlabel('Synaptic weight $w$');
         plt.ylabel('Counts per bin');
         plt.title('Synaptic weight distribution in finer bins');
```



- Columns in the left graph represent from left to right synapses that have undergone no, one, two, etc weight increases.
- Initially, we only had a single flat column for weights in [0,0.001). Since weights always increase by 0.001, all subsequent steps must also be flat.
- In the left graph, each column is 0.001 wide, thus by construction flat. In the right graph, we have 100 bins per step, more finely resolving the weight distribution.

- NEST (blue) shows too few neurons with weight increases. This is primarily due to the fact that a weight change in NEST first becomes measurable with the next spike of the postsynaptic neuron.
- Brian simulations show too many synapses with three and more weight increases.
 The reason for this is unclear.
- Brian using exponential Euler integration shows different weight distributions than with plain Euler (left graph) and resolving weights more finely, one sees that instead of flat stairs as expected we see transients in case of exponential Euler. This is most clearly visible in the [0.001, 0.002) interval. The reason for this is that the dspiked/dt = -spiked / (1*ms) : 1 equation in Brian only resets spiked instantaneously from one time step to the next when integrating with forward Euler.

Replay STDP development

- Given spike trains from actual simulations, we can replay the STDP process in
 Python by finding all situation in which a spike in one neuron is followed by a spike in
 another neuron in the next time step.
- To account for NEST's approach to effectuate the weight change caused by a prepost spike pair only when the next spike passes through the synapse, we must only include a plasticity event if it is followed by a later presynaptic spike when replaying NEST simulations.

```
In [43]: def stdp_replay(data, wbins, nest=False):
             If nest is True, include pre-post pair only if followed by another pre s
             Returns histogram over weights and full matrix.
             n_rep = len(data['spikes_t'])
             hists = np.zeros((n_rep, len(wbins)-1), dtype=int)
             W all = []
             for rep in range(n rep):
                 s_t, s_id = data['spikes_t'][rep].values, data['spikes_id'][rep].val
                 if nest:
                     t_max = np.zeros(1000, dtype=int)
                     for t, gid in zip(s_t, s_id):
                         if t == 300:
                             continue
                         t_max[gid-1] = t # exploit that spikes for a given neuron
                 W = np.zeros((1000, 1000))
                 id_pre = []
                 for t_post in range(1, 300):
                     t_pre = t_post - 1
                     id_post = s_id[np.argwhere(s_t == t_post)]
```

Replay results

- We obtain the full replay matrix only for NEST to later explore it in detail
- For Brian, we focus on weight distributions and delta synapses
- We expect that the replay produces *exactly* the same weights and thus weight distributions as the simulation
- Note that because the simulations scripts provided by Vieth et al obtain random seeds from the system clock, detailed numbers below can vary if simulations are rerun.

	Theory	NEST Sim	NEST Replay	Brian Euler Sim	Brian Euler Replay	Brian ExpEul Sim	Brian ExpEul Replay
n_stdp							
0	960847	979691	977684	960256	961308	946946	966858
1	38378	20121	22108	38156	38094	51173	32678
2	763	187	207	858	593	1199	462
3	10	1	1	302	5	320	2
4	0	0	0	225	0	236	0
5	0	0	0	152	0	103	0
6	0	0	0	45	0	20	0
7	0	0	0	6	0	2	0
8	0	0	0	0	0	1	0
9	0	0	0	0	0	0	0

- The NEST simulation and replay results show far more synapses with unchanged weight that expected (approx 978.000 vs 960.000). This is due to the fact that only pre-post spike pairings followed by another presynaptic spike with the simulation time are observable.
- In the NEST simulation, approximately 2000 more synapses (out of 1 million) remain unchanged compared to the replay. This turned out to be a bug in the code generated by NESTML for the synapse model (see https://github.com/nest/nestml/issues/1057, case 1).
- The Brian simulation results show long tails (four or more STDP changes), while the Brian replay results (based on the spikes simulated by Brian) show no more than three STDP changes. The latter is in accordance with the prediction from the binomial distribution. This may indicate an error in the Brian implementation of the STDP rule.
- Brian with exponential Euler shows results consistent with theory on replay, while simulation results show far too few synapses with unchanged weights (946.946 vs expected 960.832). This is due to the fact that equation

dspiked/dt = -spiked / (1*ms) : 1
in the Brian implementation of the plasticity rule when integrated with exponential
Euler will not lead to an instantaneous, complete reset of spiked.

- Why does NEST Sim show about 2000 more unchanged weights than NEST replay?
- We consider NEST replay here the ground truth.
- The weight matrix W_replay obtained from stdp_replay() contains all changes due to STDP, but not the intial values.
- We create a corresponding matrix from simulated data using the source and target indices stored with the weights (-1 to convert from 1-based neuron IDs in NEST to 0-based Python indices; rows: targets, columns: sources):

```
In [46]: W_sim = np.zeros((1000, 1000))
    W_sim[small['nest']['w_tgt'][0]-1, small['nest']['w_src'][0]-1] = small['nest']
In [47]: deltaW = W_replay[0] - W_sim
    too_many = np.argwhere(deltaW < -1e-6)
    too_few = np.argwhere(deltaW > 1e-6)
    print(f"Synapses with too many weight increases: {len(too_many):4d}")
    print(f"Synapses with too few weight increases: {len(too_few):4d}")

Synapses with too many weight increases: 0
Synapses with too few weight increases: 2027
```

- In the simulations, we had no extra weight increase, but some 2000 increases are missing
- Thus, the NEST simulations missed some 2000 pre-post spike pairings
- There is no obvious pattern in the misses:

- We take the first case as an example, i.e., [0, 18].
- Since these are target-source indicies, they correspond to target neuron 1, source neuron 19.
- We now use weight recorder data to look more closely at pertaining patterns

```
In [49]: wr = small['nest']['wrec'][0].set_index(['targets', 'senders']).drop(['Unnam
In [50]: tx, sx = too_few[0, :] + np.array([1, 1], dtype=int)
In [51]: tx, sx
Out[51]: (1, 19)
In [52]: wr.loc[(tx, sx)]
```

Out[52]:			ports	times	weights
	targets	senders			
	1	19	2250	52.0	0.000208
		19	2250	178.0	0.000208

- Three spikes passed the 19 -> 1 synapse, the weight retained its initial value
- We now look at the spike times of the postsynaptic neuron, neuron 1

19 2250 229.0 0.000208

- After the sender (19) spiked at 52ms, the target (1) spiked at 53 ms, and this should have caused a weight increase by 0.001.
- In NEST, this weight increase should have been visible in the weight recorder entry for the spike from 19 to 1 at 178 ms.
- We note that the postsynaptic neuron spiked two more times (82 and 160 ms) before the next spike through the 19->1 synpse at 178 ms.

Good vs bad cases

- The following analysis depends on the 52-53 ms spike timing. For different simulation data, this may need to be adjusted. Choose the pair of spike times that should have led to stdp.
- We now look in the full spike data for all pairs of neurons in which one fired at 52 ms and the other at 53 ms and group them as follows:
 - Good: synapse weight has increased
 - Bad: synapse weight has not increased
 - Latent: synapse weight increase is not visible since no more spike through synapse after 53 ms

```
In [55]: def group_events(spks, wr, pre_time):
    pre_spikes = spks.loc[spks.times==pre_time]
    post_spikes = spks.loc[spks.times==pre_time+1]
    pairs = [(post, pre) for pre in pre_spikes.index for post in post_spikes
    res = {'good': [], 'bad': [], 'latent': []}
```

```
for ix in pairs:
                  elem = wr.loc[ix] # all spikes between pre and post neuron
                  tix = np.argwhere(elem.times.values == pre_time) # index for spike
                  if tix < len(elem) - 1:</pre>
                      if elem.weights.values[tix+1] > elem.weights.values[tix]:
                          res['good'].append(ix)
                      else:
                          res['bad'].append(ix)
                  else:
                      res['latent'].append(ix)
              return res
In [56]: gr = group_events(spks, wr, 52)
         Some good examples
In [57]: gr['good'][0]
Out[57]: (660, 463)
In [58]: wr.loc[gr['good'][0]]
Out[58]:
                                         weights
                           ports times
          targets senders
                     463 57784 52.0 0.000689
            660
                     463 57784 162.0 0.001689
In [59]: spks.loc[660].T
Out[59]: senders 660
                        660
                              660
                                     660
                                           660
            times 53.0 125.0 191.0 244.0 275.0
           • For the 463 -> 660 pair, there is only one postsynaptic spike (at 125) before the
             next presynaptic (at 162)
In [60]: gr['good'][15]
Out[60]: (287, 515)
```

In [61]: wr.loc[gr['good'][15]]

```
Out[61]:
                          ports times weights
         targets senders
            287
                                  20.0 0.000313
                     515
                          64285
                                  52.0 0.000313
                     515 64285
                     515
                         64285
                                 122.0 0.001313
                     515 64285
                                158.0 0.001313
                     515 64285
                                 211.0 0.001313
                     515 64285
                                 244.0 0.001313
In [62]: spks.loc[287].T
Out[62]: senders
                       287 287
                                        287
                  287
                                  287
```

times 29.0 53.0 81.0 126.0 204.0

 Also for the 515 -> 287, only a single spike (at 81) between pairing and readout (at 122)

Some bad cases

```
gr['bad'][0]
In [63]:
Out[63]: (231, 463)
In [64]: wr.loc[gr['bad'][0]]
Out[64]:
                          ports times
                                        weights
          targets senders
             231
                     463
                          57851
                                  52.0
                                       0.000804
                     463 57851
                                 162.0 0.000804
In [65]: spks.loc[231].T
Out[65]: senders
                   231
                         231
                               231
            times 53.0 102.0 132.0
```

 For the 463 -> 231 pair, the postsynaptic neuron fires twice (102, 132) before the "readout" spike at 162

```
In [66]: gr['bad'][15]
```

```
Out[66]: (797, 392)

In [67]: wr.loc[gr['bad'][15]]

Out[67]: ports times weights

targets senders

797 392 48920 52.0 0.000742

392 48920 271.0 0.000742
```

```
In [68]: spks.loc[797].T

Out[68]: senders 797 797 797

times 53.0 93.0 182.0 281.0
```

- For the 392 -> 797 pair, the postsynaptic neuron fires twice (93, 182) before the "readout" spike at 271
- Thus, the occurence of more than one post-synaptic spike between the pre-post pairing and the next pre-synaptic spike (aka readout spike) makes NEST miss an stdp event.
- See https://github.com/nest/nestml/issues/1057

Note for debugging

• The (target id, port) pair can be used to uniquely identify a synapse in calls to send() to activate debugging output only for specific synapses.

```
In []:
```