

# Programming neuromorphic computers: PyNN and beyond

Andrew Davison  
Paris-Saclay Institute of Neuroscience  
CNRS - Université Paris-Saclay



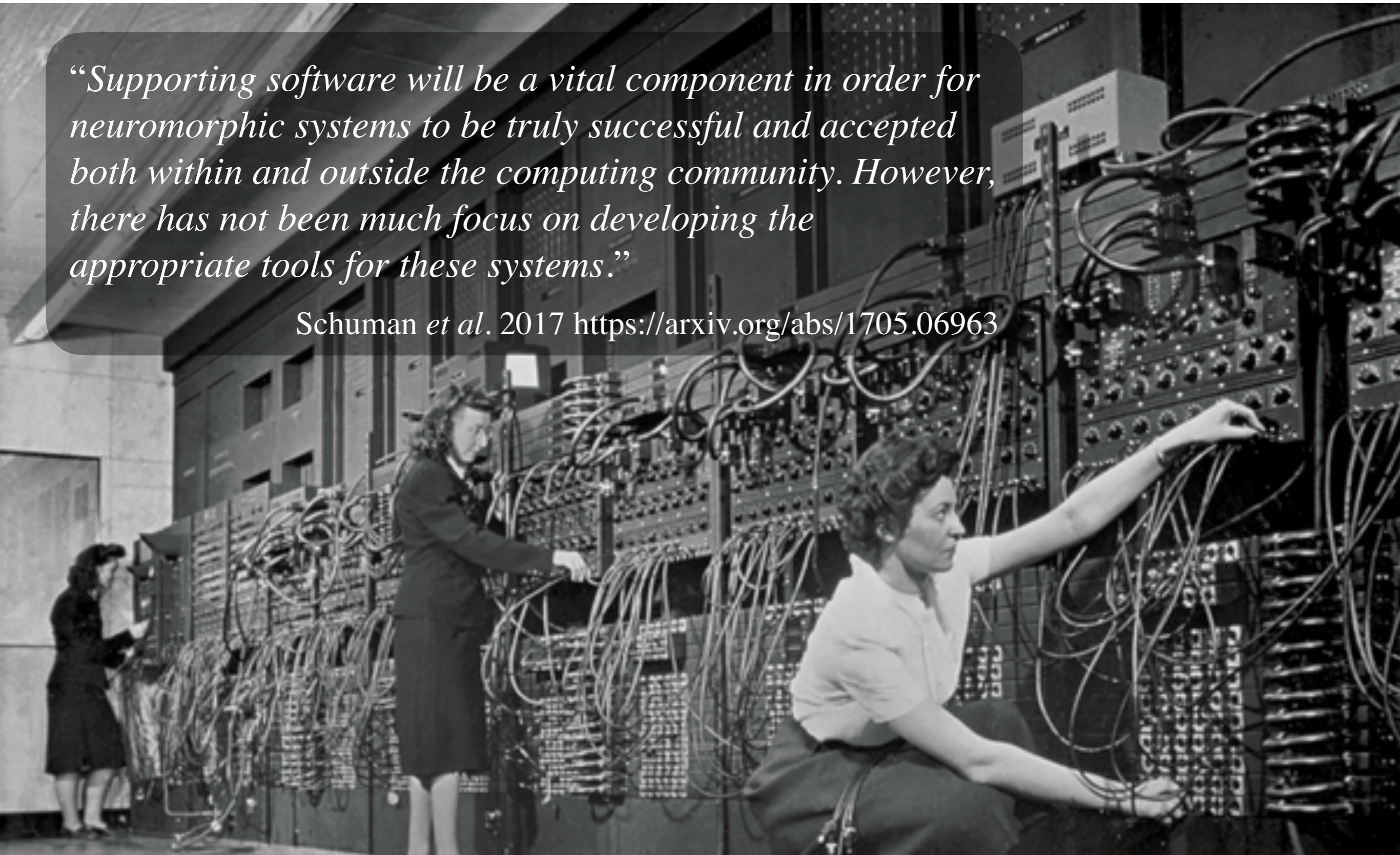
NICE 2021

19th March 2021

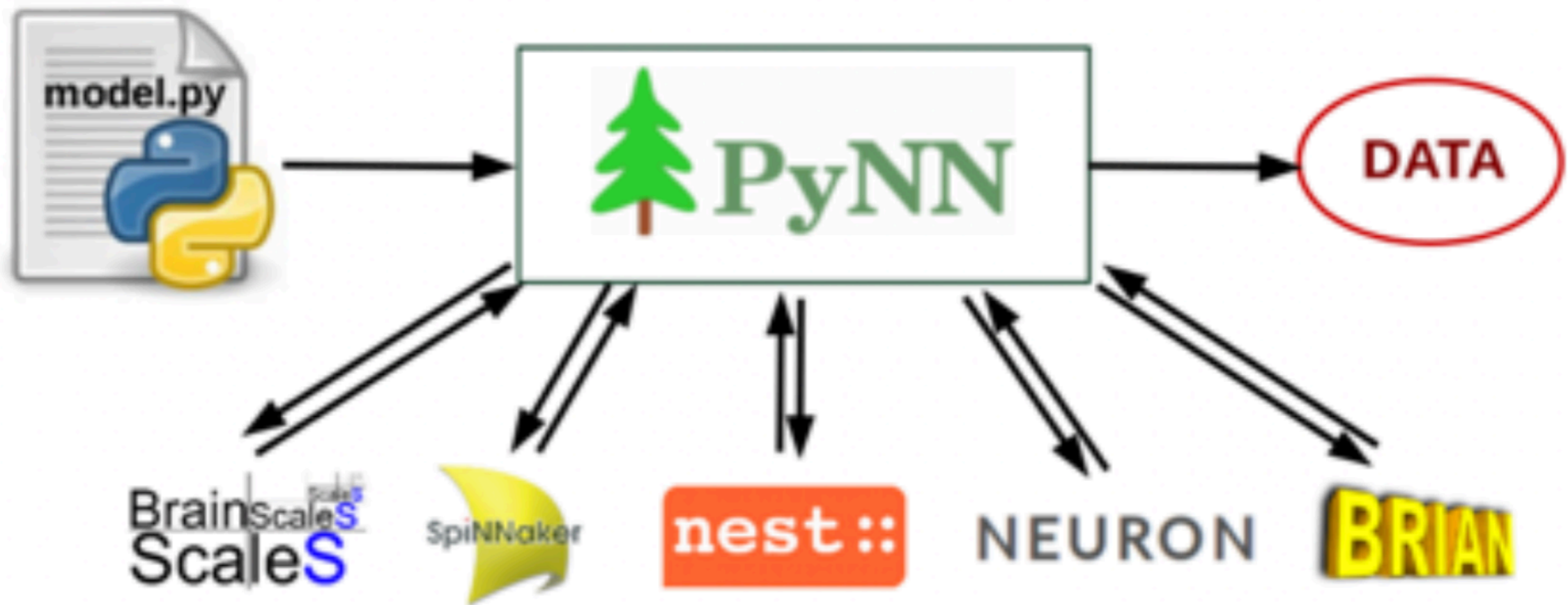
# Programming neuromorphic computers

*“Supporting software will be a vital component in order for neuromorphic systems to be truly successful and accepted both within and outside the computing community. However, there has not been much focus on developing the appropriate tools for these systems.”*

Schuman *et al.* 2017 <https://arxiv.org/abs/1705.06963>



# Introduction to PyNN



```
sim.setup(timestep=0.1)

cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
all = pE + pI
input = sim.Population(100, sim.SpikeSourcePoisson(
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))

weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
DDPC = sim.DistanceDependentProbabilityConnector
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,
    delays="0.5+0.01d")
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)

exc = sim.Projection(pE, all, connector,
    synapse_type="depressing", receptor_type="excitatory")
inh = sim.Projection(pI, all,
    connector, receptor_type="inhibitory")
```

```
import pyNN.nest as sim

sim.setup(timestep=0.1)

cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
all = pE + pI
input = sim.Population(100, sim.SpikeSourcePoisson(
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))

weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
DDPC = sim.DistanceDependentProbabilityConnector
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,
    delays="0.5+0.01d")
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)

exc = sim.Projection(pE, all, connector,
    synapse_type="depressing", receptor_type="excitatory")
inh = sim.Projection(pI, all,
    connector, receptor_type="inhibitory")
```

```
import pyNN.neuron as sim
```

```
sim.setup(timestep=0.1)
```

```
cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
```

```
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
```

```
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
```

```
all = pE + pI
```

```
input = sim.Population(100, sim.SpikeSourcePoisson(  
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
```

```
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))
```

```
weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
```

```
DDPC = sim.DistanceDependentProbabilityConnector
```

```
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,  
    delays="0.5+0.01d")
```

```
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)
```

```
exc = sim.Projection(pE, all, connector,  
    synapse_type="depressing", receptor_type="excitatory")
```

```
inh = sim.Projection(pI, all,  
    connector, receptor_type="inhibitory")
```

```
import pyNN.brian as sim
```

```
sim.setup(timestep=0.1)
```

```
cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
```

```
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
```

```
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
```

```
all = pE + pI
```

```
input = sim.Population(100, sim.SpikeSourcePoisson(  
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
```

```
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))
```

```
weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
```

```
DDPC = sim.DistanceDependentProbabilityConnector
```

```
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,  
    delays="0.5+0.01d")
```

```
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)
```

```
exc = sim.Projection(pE, all, connector,  
    synapse_type="depressing", receptor_type="excitatory")
```

```
inh = sim.Projection(pI, all,  
    connector, receptor_type="inhibitory")
```

```
import pyNN.spiNNaker as sim

sim.setup(timestep=0.1)

cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
all = pE + pI
input = sim.Population(100, sim.SpikeSourcePoisson(
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))

weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
DDPC = sim.DistanceDependentProbabilityConnector
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,
    delays="0.5+0.01d")
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)

exc = sim.Projection(pE, all, connector,
    synapse_type="depressing", receptor_type="excitatory")
inh = sim.Projection(pI, all,
    connector, receptor_type="inhibitory")
```



```
import pyNN.brainscales as sim
```

```
sim.setup(timestep=0.1)
```

```
cell_parameters = {"tau_m": 12.0, "cm": 0.8, "v_thresh": -50.0}
```

```
pE = sim.Population(2e4, sim.IF_cond_exp(**cell_parameters))
```

```
pI = sim.Population(5e3, sim.IF_cond_exp(**cell_parameters))
```

```
all = pE + pI
```

```
input = sim.Population(100, sim.SpikeSourcePoisson(  
    rate=random.RandomDistribution("normal", (10.0, 2.0))))
```

```
all.inject(sim.NoisyCurrentSource(mean=0.1, stdev=0.01))
```

```
weight_distr = random.RandomDistribution("uniform", (0.0, 0.1))
```

```
DDPC = sim.DistanceDependentProbabilityConnector
```

```
connector = DDPC("exp(-d**2/400.0)", weights=weight_distr,  
    delays="0.5+0.01d")
```

```
depressing = sim.TsodyksMarkramMechanism(U=0.5, tau_rec=800.0)
```

```
exc = sim.Projection(pE, all, connector,  
    synapse_type="depressing", receptor_type="excitatory")
```

```
inh = sim.Projection(pI, all,  
    connector, receptor_type="inhibitory")
```

# The PyNN API

- neuron and synapse models
- populations
- connectivity
- recording & data handling

# Neuron and synapse models

- “standard” models

available on at least 2  
simulators/hardware platforms

- native models

use the PyNN API with  
simulator-specific models: useful  
in migrating to “full PyNN”

- NineML, NeuroML,  
NESTML models

code generation for custom,  
simulator-independent models

# “Native” models

can wrap any model provided by/buildable with a given simulator to use with PyNN:

```
from pyNN.nest import native_cell_type, native_synapse_type

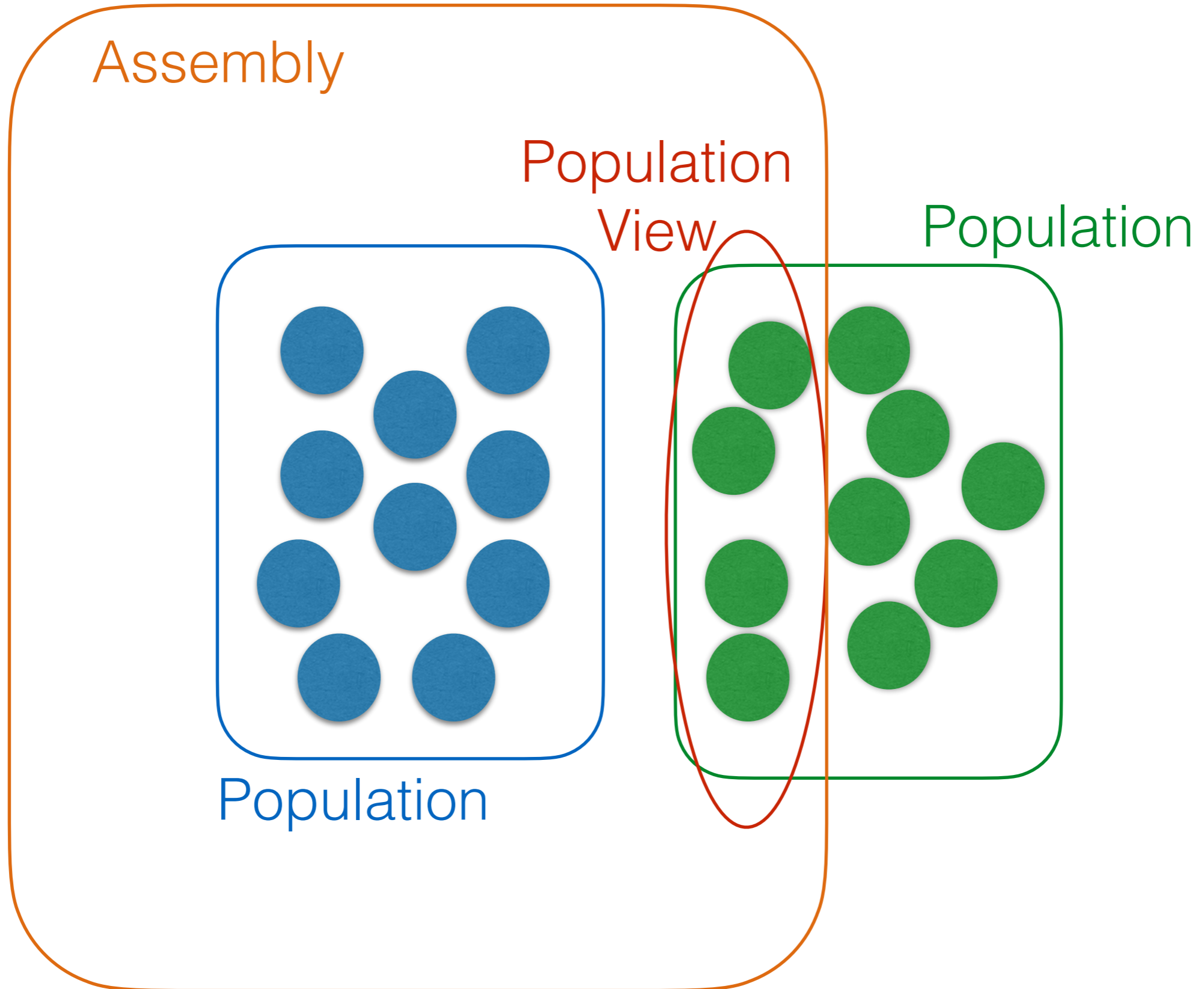
ht_neuron = native_cell_type("ht_neuron")
poisson = native_cell_type("poisson_generator")

cell_type = ht_neuron(Tau_m=20.0)
input_type = poisson(rate=200.0)

stdp = native_synapse_type("stdp_synapse")

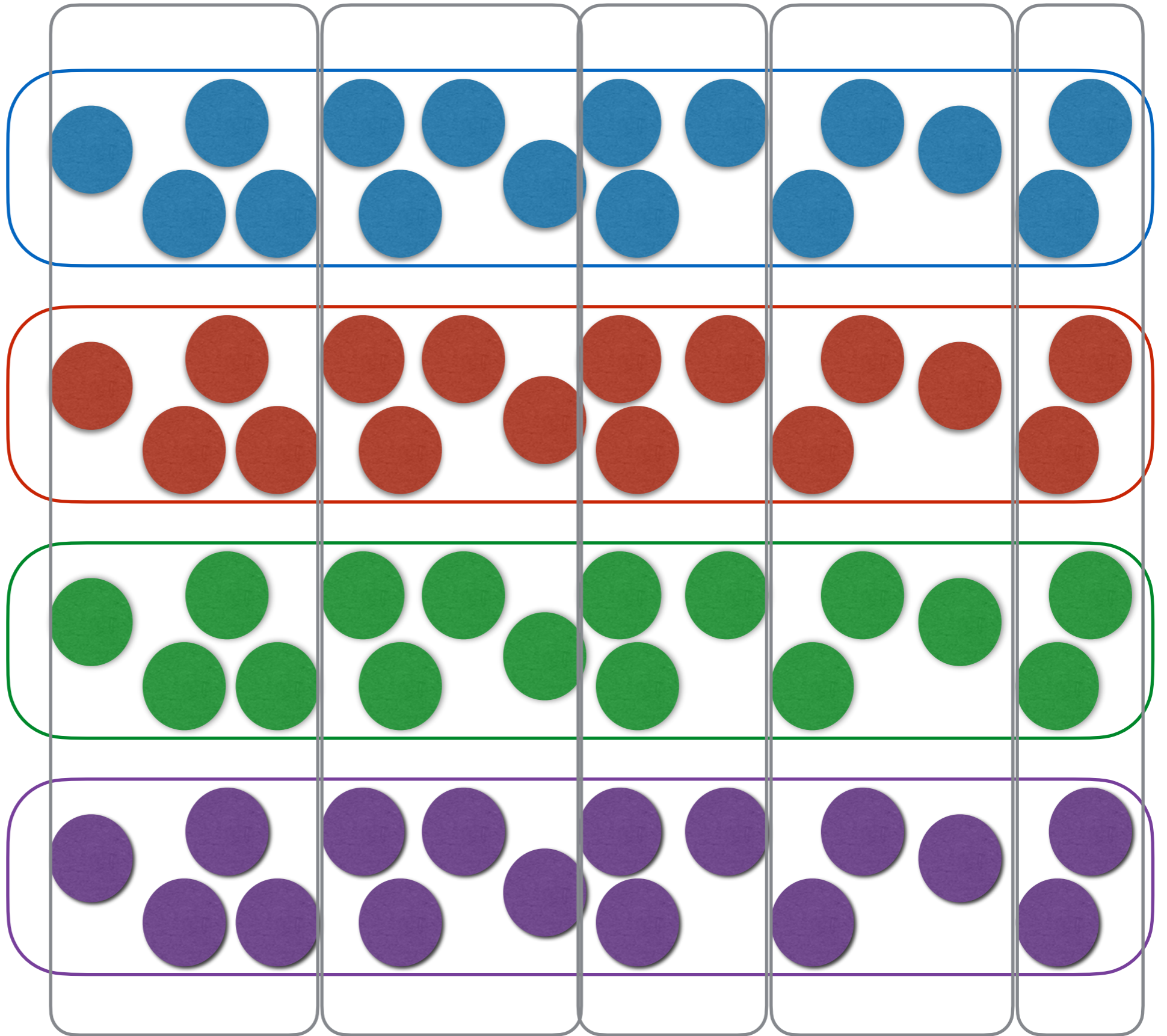
synapse_type = stdp(Wmax=50.0, lambda=0.015)
```

# Populations



# Assembly

Population



# Populations

```
structure = RandomStructure(boundary=Sphere(radius=200.0))

cells = Population(100, thalamocortical_type,
                  structure=structure,
                  initial_values={'v': -70.0},
                  label="Thalamocortical neurons")

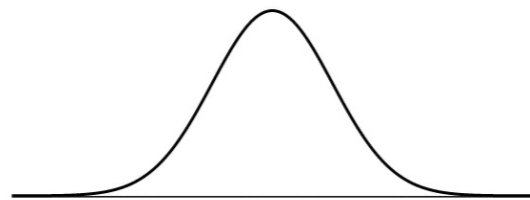
view = cells[:80]           # the first eighty neurons
view = cells[::2]          # every second neuron
view = cells[45, 91, 7]    # a specific set of neurons
view = cells.sample(50)    # 50 neurons at random

layer4 = spiny_stellates + l4_interneurons # an Assembly
```

# Parameterization

```
parameter_space = {  
    'tau_m': RandomDistribution('uniform', (10.0, 15.0)),  
    'cm':    0.85,  
    'v_rest': lambda i: np.cos(i*pi*10/n),  
    'v_reset': np.linspace(-75.0, -65.0, num=n)}  
  
cell_type = IF_cond_alpha(**parameter_space)
```

42

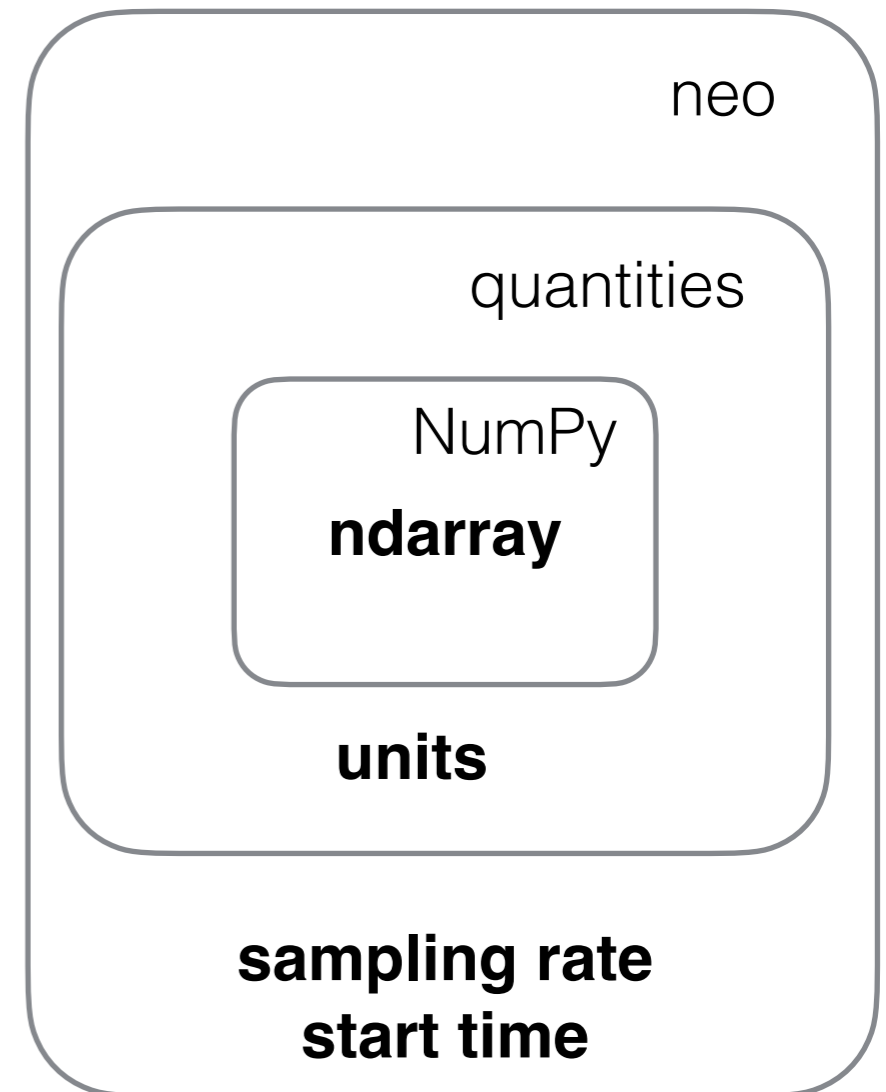
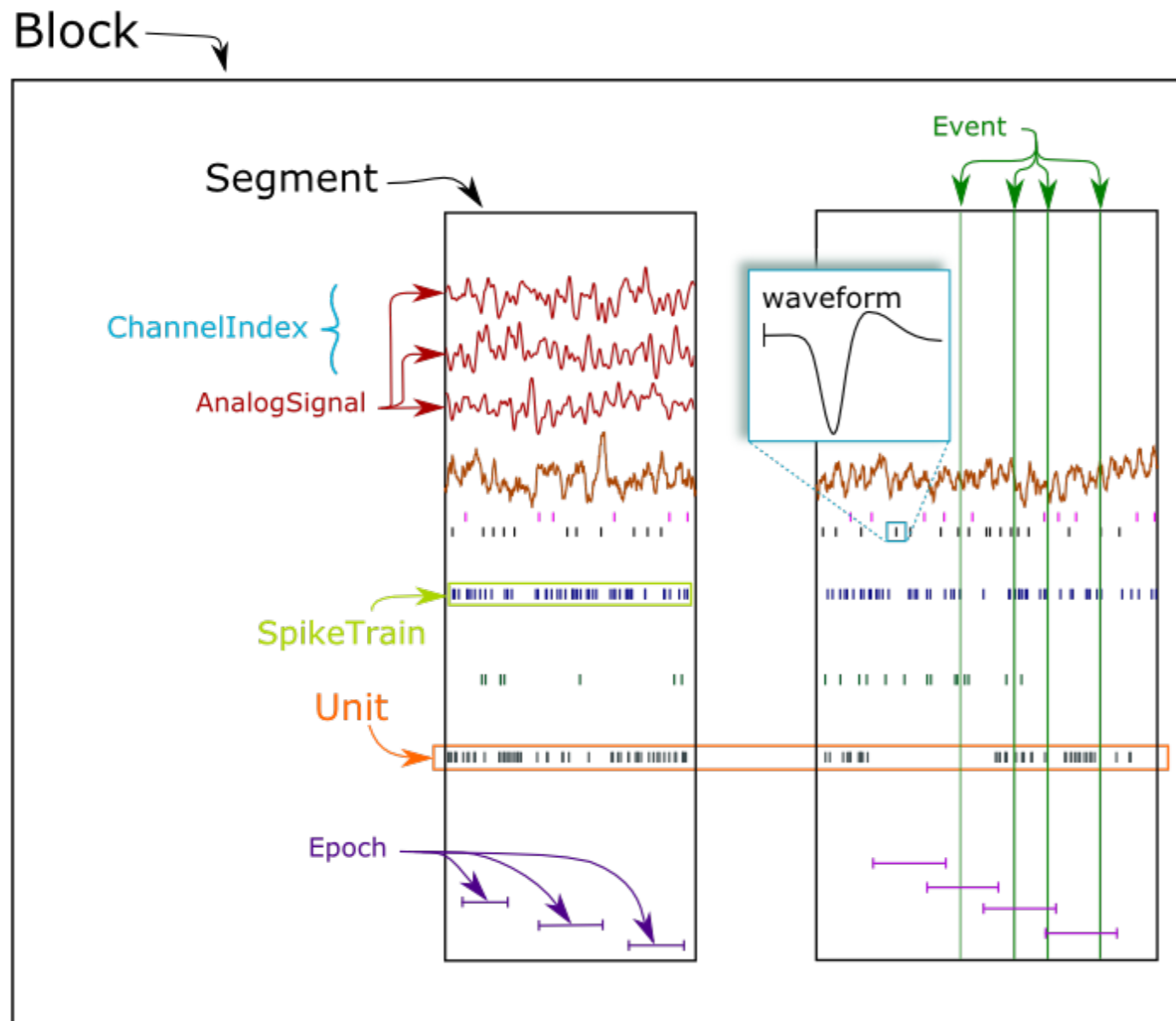


f0



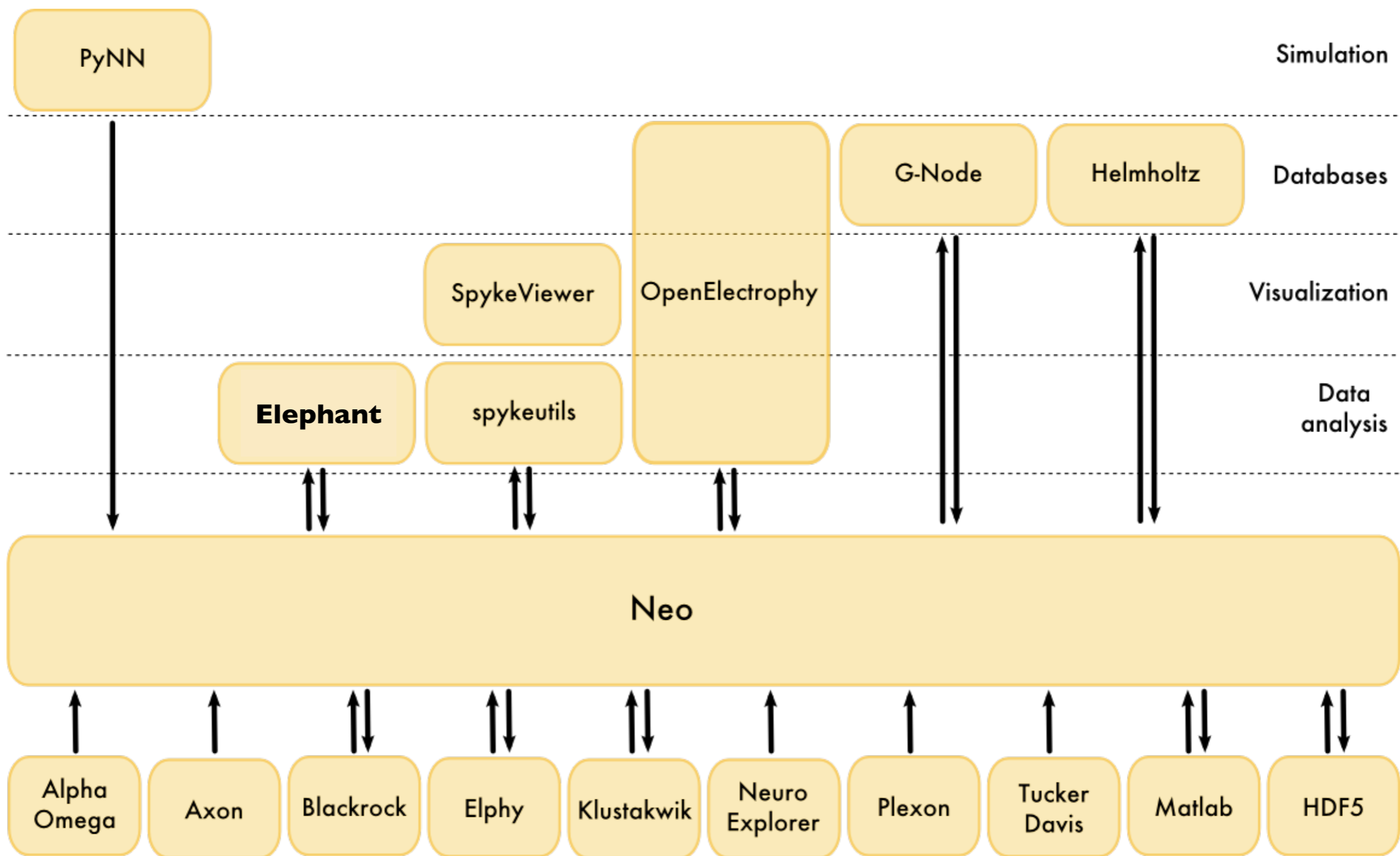


# Data handling



<http://neuralensemble.org/neo>





# PyNN usage and uptake



☆ Star 192    🍴 Fork 104

## Dependency graph

Dependencies    Dependents    Dependabot Beta

### Repositories that depend on pynn

56 Repositories    5 Packages    ⓘ

jakobj / UP-Tasks	☆ 0	🍴 5
ac13711 / Spike_Sorting_GUI	☆ 0	🍴 0
SpiNNakerManchester / SpiNNCer	☆ 1	🍴 0
CSNG-MFF / mozaik	☆ 12	🍴 16
NeuroML / NeuroMLlite	☆ 7	🍴 6
Katterrina / mff_nprg045	☆ 0	🍴 0
NeuromorphicProcessorProject / snn_toolbox	☆ 187	🍴 74
SpiNNakerManchester / sPyNNaker	☆ 75	🍴 32
ruthvik92 / SpykeFlow	☆ 3	🍴 4
MetaCell / scidash	☆ 1	🍴 2

PyNN: a common interface for neuronal network simulators

AP Davison, D Brüderle, JM Eppler, J Kremkow, E Muller, D Pecevski, ...  
Frontiers in neuroinformatics 2, 11

592

2009

# Implementing the PyNN API

- Each backend is a separate Python package
- Implementation choices:
  - entirely independent implementation from scratch (e.g. in C++ with Python wrapper)
  - implement minimal hooks for the “common” implementation
  - anywhere in between

New!

## PyCARL: A PyNN Interface for Hardware-Software Co-Simulation of Spiking Neural Network

Adarsha Balaji<sup>1</sup>, Prathyusha Adiraju<sup>2</sup>, Hiram J. Kashyap<sup>3</sup>, Anup Das<sup>1,2</sup>,  
Jeffrey L. Krichmar<sup>3</sup>, Nikil D. Dutt<sup>3</sup>, and Francky Catthoor<sup>2,4</sup>

<sup>1</sup>Electrical and Computer Engineering, Drexel University, Philadelphia, USA

<sup>2</sup>Neuromorphic Computing, Stichting Imec Nederlands, Eindhoven, Netherlands

<sup>3</sup>Cognitive Science and Computer Science, University of California, Irvine, USA

<sup>4</sup>ESAT Department, KU Leuven and IMEC, Leuven, Belgium

<https://arxiv.org/abs/2003.09696>

# Hooks for the common implementation

```
class Population(common.Population):  
  
    def _create_cells(self):  
        ...  
  
    def _get_parameters(self, *names):  
        ...  
  
    def _set_parameters(self, parameter_space):  
        ...
```

# Hooks for the common implementation

```
class Projection(common.Projection):  
  
    def _convergent_connect(self, presynaptic_indices,  
                           postsynaptic_index,  
                           **connection_parameters):  
  
        ...
```

# Hooks for the common implementation

```
class Recorder(recording.Recorder):  
  
    def _record(self, variable, new_ids):  
        ...  
  
    def _get_spiketimes(self, id):  
        ...  
  
    def _get_all_signals(self, variable, ids, clear=False):  
        ...  
  
    def _local_count(self, variable, filter_ids=None):  
        ...
```



# Hooks for the common implementation

```
class SpikeSourcePoisson(cells.SpikeSourcePoisson):  
  
    translations = build_translations(  
        ('start', 'START'),  
        ('rate', 'INTERVAL', "1000.0/rate", "1000.0/INTERVAL"),  
        ('duration', 'DURATION'),  
    )
```

# Hooks for the common implementation

```
class State(common.control.BaseState):
```

```
    def run_until(self, tstop):
```

```
        ...
```

```
    def clear(self):
```

```
        ...
```

```
    def reset(self):
```

```
        ...
```

# Development model

- open source
- open community
- community governance\*

**CONTRIBUTIONS  
WELCOME!**

Pierre Yger	Eilif Muller
Daniel Brüderle	Jochen Eppler
Jens Kremkow	Dejan Pecevski
Mike Hull	Michael Schmuker
Mikael Djurfeldt	Bernhard Kaplan
Subhasis Ray	Yury Zaytsev
Jan Antolik	Alexandre Gravier
Thomas Close	Oliver Breitwieser
Jannis Schücker	Maximilian Schmidt
Christian Rössert	Shailesh Appukuttan
Elodie Legouée	Joffrey Gonin
Ankur Sinha	Håkon Mørk

<https://github.com/NeuralEnsemble/PyNN/>

\*Any contributor who has had at least three pull requests accepted may be nominated as a maintainer.

# Ongoing and future work

- separation of API specification from reference implementation
- revised and extended documentation
- API support for cleaner separation of model and experiment descriptions
- multicompartmental models

# Separation of API specification from reference implementation

- “PyNN-like”
- API currently defined by the reference implementation
- Plan: use Python `abc` module, Python 3 type annotations to separate API from implementation
- API simplification? core & extensions



# Multicompartmental models

- dendrites offer rich possibilities for computation

Review Article | Published: 11 May 2020

## **Illuminating dendritic function with computational models**

Panayiota Poirazi  & Athanasia Papoutsis

*Nature Reviews Neuroscience* 21, 303–321(2020) | [Cite this article](#)

- an increasing number of software simulators and neuromorphic systems now support multicompartment models
- time to extend PyNN from point neurons to neurons with detailed morphologies, ion channels...

<http://neuralensemble.org/docs/PyNN/2.0/>

# Design goals

- maintain the same main level of abstraction: populations of neurons and the sets of connections between populations (projections);
- backwards compatibility (point neuron models created with PyNN 1.0 (not yet released) or later should work with no, or minimal, changes);
- integrate with other open-source simulation tools and standards (e.g. NeuroML) wherever possible, rather than reinventing the wheel;
- support neuromorphic hardware systems.

# Example: ball-and-stick model

```
from neuroml import Segment, Point3DWithDiam as P
from pyNN.morphology import NeuroMLMorphology, uniform
from pyNN.parameters import IonicSpecies
import pyNN.neuron as sim

sim.setup(timestep=0.025)

soma = Segment(proximal=P(x=0, y=0, z=0, diameter=18.8),
               distal=P(x=18.8, y=0, z=0, diameter=18.8),
               name="soma", id=0)
dend = Segment(proximal=P(x=0, y=0, z=0, diameter=2),
               distal=P(x=-500, y=0, z=0, diameter=2),
               name="dendrite",
               parent=soma, id=1)

cell_class = sim.MultiCompartmentNeuron
cell_class.label = "ExampleMultiCompartmentNeuron"
cell_class.ion_channels = {"pas": sim.PassiveLeak, "na": sim.NaChannel,
                          "kdr": sim.KdrChannel}

cell_type = cell_class(morphology=NeuroMLMorphology(segments=(soma, dend)),
                      cm=1.0, Ra=500.0,
                      pas={"conductance_density": uniform("all", 0.0003), "e_rev": -54.3},
                      na={"conductance_density": uniform("soma", 0.120), "e_rev": 50.0},
                      kdr={"conductance_density": uniform("soma", 0.036), "e_rev": -77.0})

cells = sim.Population(2, cell_type, initial_values={'v': [-60.0, -70.0]})
```

morphology defined  
using libNeuroML

standard library  
of ion channels



# Example: morphology from SWC

```
from pyNN.morphology import load_morphology, uniform, random_section, dendrites,
apical_dendrites, by_distance
from pyNN.parameters import IonicSpecies
import pyNN.neuron as sim

sim.setup(timestep=0.025)

pyr_morph = load_morphology("oi15rpy4-1.CNG.swc")

cell_class = sim.MultiCompartmentNeuron
cell_class.label = "ExampleMultiCompartmentNeuron"
cell_class.ion_channels = {"pas": sim.PassiveLeak, "na": sim.NaChannel,
                           "kdr": sim.KdrChannel}
cell_class.post_synaptic_entities = {"AMPA": sim.CondExpPostSynapticResponse,
                                     "GABA_A": sim.CondExpPostSynapticResponse}

cell_type = cell_class(morphology=pyr_morph),
                    cm=1.0, Ra=500.0,
                    pas={"conductance_density": uniform("all", 0.0003), "e_rev": -54.3},
                    na={"conductance_density": uniform("soma", 0.120), "e_rev": 50.0},
                    kdr={"conductance_density": uniform("soma", 0.036), "e_rev": -77.0}
                    AMPA={"density": uniform('all', 0.05), # number per  $\mu\text{m}$ 
                          "e_rev": 0.0, "tau_syn": 2.0},
                    GABA_A={"density": by_distance(dendrites(), lambda d: 0.05 * (d < 50.0)),
                           "e_rev": -70.0, "tau_syn": 5.0})
```

morphology read from  
SWC

standard library  
of ion channels  
and synaptic  
receptors

# Recording and injecting current

## named segments

```
step_current = sim.DCSource(amplitude=0.1, start=50.0, stop=150.0)
step_current.inject_into(cells[0:1], location="soma")

cells.record('spikes')
cells.record(['na.m', 'na.h', 'kdr.n'], locations=['soma'])
cells.record('v', locations=['soma', 'dendrite'])
```

## selecting neurite locations

```
step_current = sim.DCSource(amplitude=5.0, start=50.0, stop=150.0)
step_current.inject_into(cells[1:2], location=random_section(apical_dendrites()))

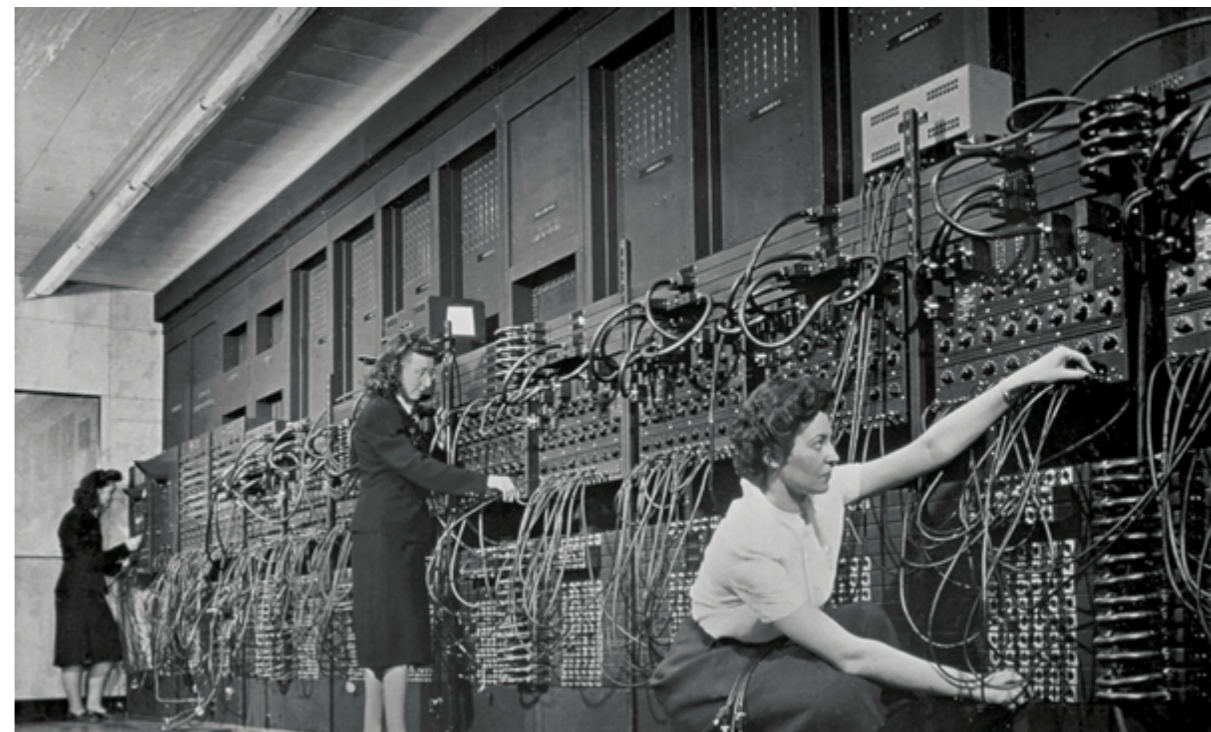
cells.record('spikes')
cells.record(['na.m', 'na.h', 'kdr.n'], locations={'soma': 'soma'})
cells.record('v', locations={'soma': 'soma', 'dendrite': random_section(apical_dendrites())})
```

# Networks

```
i2p = sim.Projection(  
    inputs,  
    pyramidal_cells,  
    connector=sim.AllToAllConnector(  
        location_selector=random_section(apical_dendrites()),  
        synapse_type=sim.StaticSynapse(weight=0.5, delay=0.5),  
        receptor_type="AMPA"  
    )  
)
```

# Beyond PyNN

- if PyNN is assembly / VHDL / C, we need:
  - a standard library
  - higher-level languages
  - exploration of algorithms
  - very preliminary work in my group on this. Open to collaboration



# More information

## *Documentation*

<http://neuralensemble.org/PyNN/>

## *Licence*

CeCILL (GPL-equivalent)

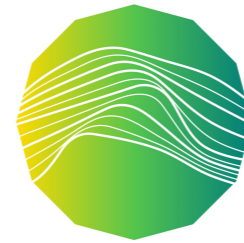
## *Mailing list*

<https://groups.google.com/forum/#!forum/neuralensemble>





Human Brain Project



EBRAINS

[andrew.davison@cnr.fr](mailto:andrew.davison@cnr.fr)

[@apdavison](https://twitter.com/apdavison)

Pierre Yger	Eilif Muller
Daniel Brüderle	Jochen Eppler
Jens Kremkow	Dejan Pecevski
Mike Hull	Michael Schmuker
Mikael Djurfeldt	Bernhard Kaplan
Subhasis Ray	Yury Zaytsev
Jan Antolik	Alexandre Gravier
Thomas Close	Oliver Breitwieser
Jannis Schücker	Maximilian Schmidt
Christian Rössert	Shailesh Appukuttan
Elodie Legouée	Joffrey Gonin
Ankur Sinha	Håkon Mørk



Co-funded by  
the European Union

