

Selected resources for computational neuroscience

23 November 2020

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- Model name
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- Region(circuits)
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- Cell type
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- Find models of
- Realistic Networks
- Neurons
- Electrical synapses (gap junctions)
- Chemical synapses
- Ion channels
- Neuromuscular junctions

Amyloid beta (IA block) effects on a model CA1 pyramidal cell (Morse et al. 2010)

Download zip file Auto-launch

Help downloading and running models

Model Information Model File Citations Model Views Simulation Platform 3D Print

Accession:87284

The model simulations provide evidence oblique dendrites in CA1 pyramidal neurons are susceptible to hyper-excitability by amyloid beta block of I_A channel, I_A . See paper for details.

Reference:

1 . Morse TM, Carnevale NT, Mutalik PG, Migliore M, Shepherd GM (2010) Abnormal excitability of oblique dendrites implicated in early Alzheimer's computational study *Front. Neural Circuits* 4:16 [PubMed]

Model Information (Click on a link to find other models with that property)

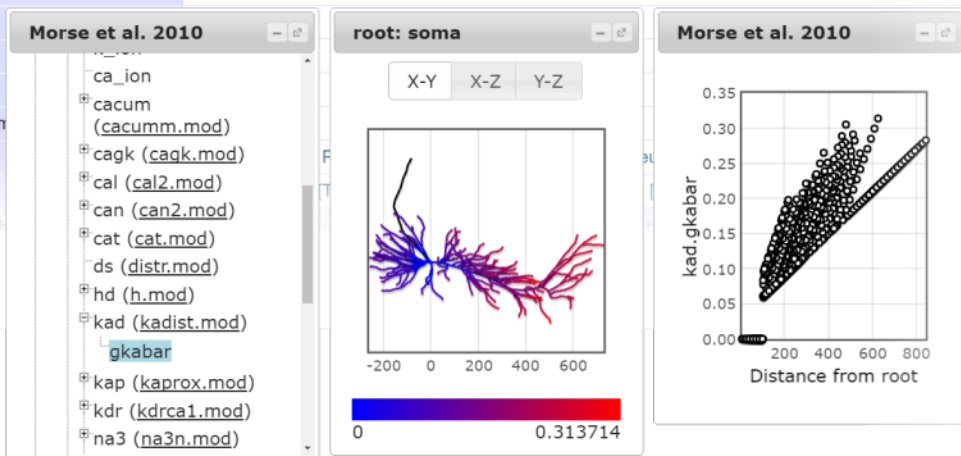
Model Type: Neuron or other electrically excitable cell;

Brain Region(s)/Organism:

Cell Type(s): Hippocampus CA1 pyramidal cell;

Channel(s): I Na,t; I L high threshold; I N; I T low threshold; I A; I K; I h;

Gap Junctions:



```
from neuron import h, rxd
import neuron.rxd.node as node
from matplotlib import pyplot
import time
```

```
h.load_file('stdrun.hoc')
```

```
soma = h.Section()
soma.L = 10
soma.diam = 10
soma.nseg = 11
dend = h.Section()
dend.connect(soma)
dend.L = 50
dend.diam = 2
dend.nseg = 51
```

```
def print_nodes():
    print ', '.join(str(v) for v in node._states)
```

```
print 'defining rxd'
region = rxd.Region(h.allsec(), nrn_region='i')
ca = rxd.Species(region, name='ca', d=1, charge=2, initial:
reaction = rxd.Rate(ca, -ca * (1 - ca) * (0.3 - ca))
```

```
print 'initializing'
h.initialize()
```

Morse TM, Carnevale NT, Mutalik PG, Migliore M, Shepherd GM (2010) Abnormal excitability of oblique dendrites implicated in early Alzheimer's: a computational study *Front. Neural Circuits* 4:16 [PubMed]

References and models cited by this paper

Acker CD, White JA (2007) Roles of $I(A)$ and morphology in action potential propagation in CA1 pyramidal cell dendrites. *J Comput Neurosci* 23(2):201-16 [Journal] [PubMed]

• Roles of $I(A)$ and morphology in AP prop. in CA1 pyramidal cell dendrites (Acker and White 2007) [Model]

Anderton BH, Callahan L, Coleman P, Davies P, Flood D, Jicha GA, Ohm T, Weaver C (1998) Dendritic changes in Alzheimer's disease and factors that may underlie these changes. *Prog Neurobiol* 55:595-609 [PubMed]

Andrasfalvy BK, Makara JK, Johnston D, Magee JC (2008) Altered synaptic and non-synaptic properties of CA1 pyramidal neurons in β -amyloid transgenic mice. *J Neurosci* 28(18):4550-4560 [Journal] [PubMed]

References and models that cite this paper

Culmone V, Migliore M (2012) Progressive effect of beta amyloid peptides accumulation on CA1 pyramidal neurons: a model study suggesting possible treatments *Front Comput Neurosci* 6:52 [Journal] [PubMed]

• CA1 pyramidal neurons: effects of Alzheimer (Culmone and Migliore 2012) [Model]

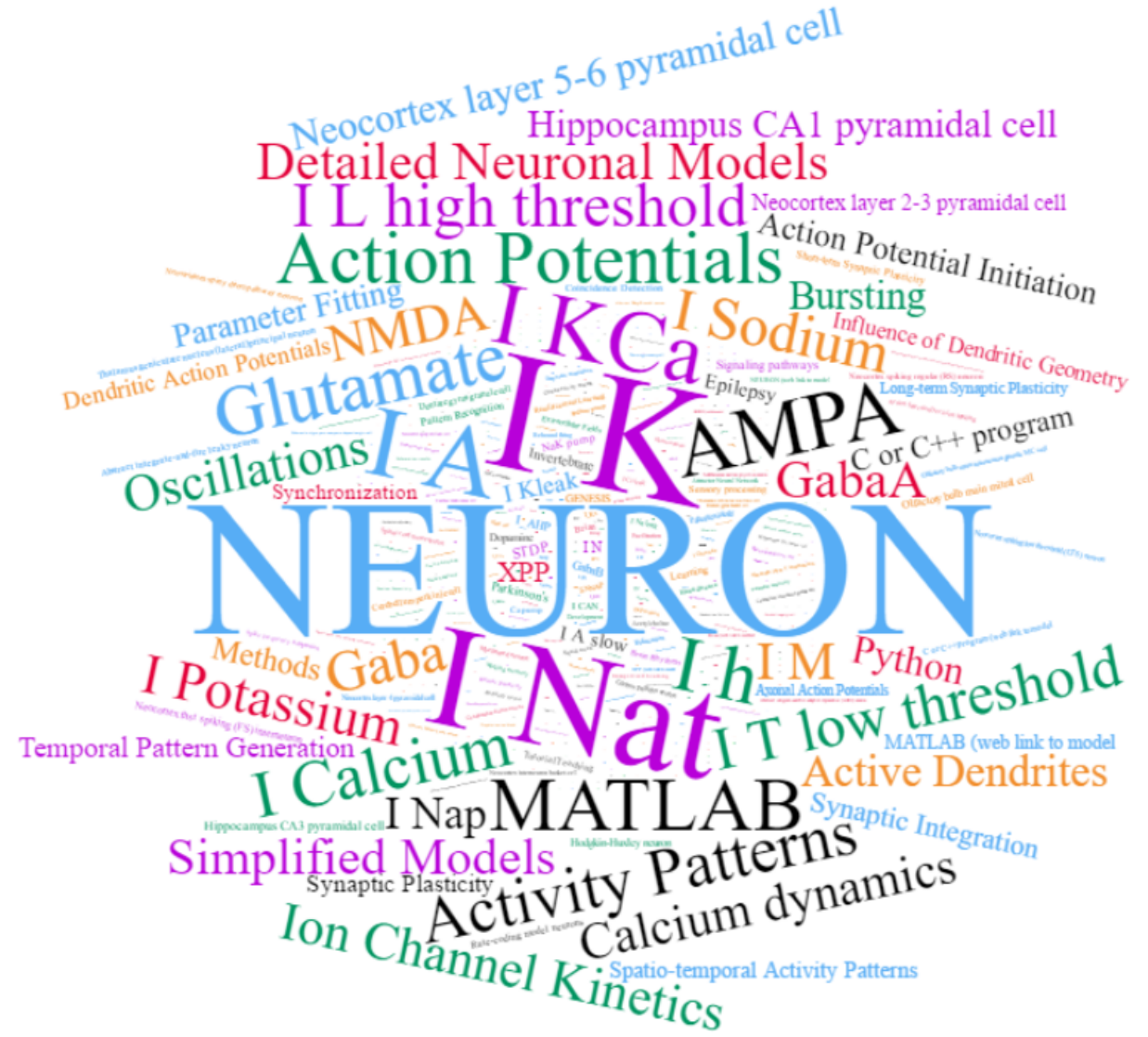
McDougal RA, Morse TM, Hines ML, Shepherd GM (2015) ModelView for ModelDB: online presentation of model structure *Neuroinformatics* 13(4):459-70 [Journal] [PubMed]

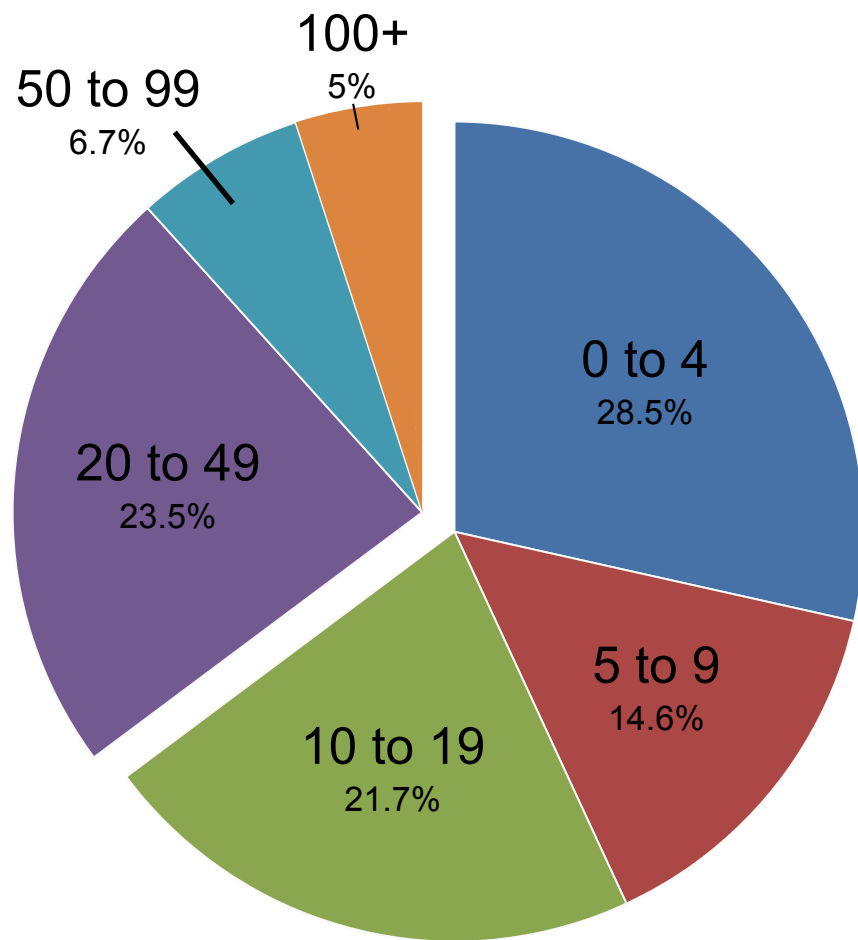
• ModelView: online structural analysis of computational models (McDougal et al. 2015) [Model]

modeldb.yale.edu

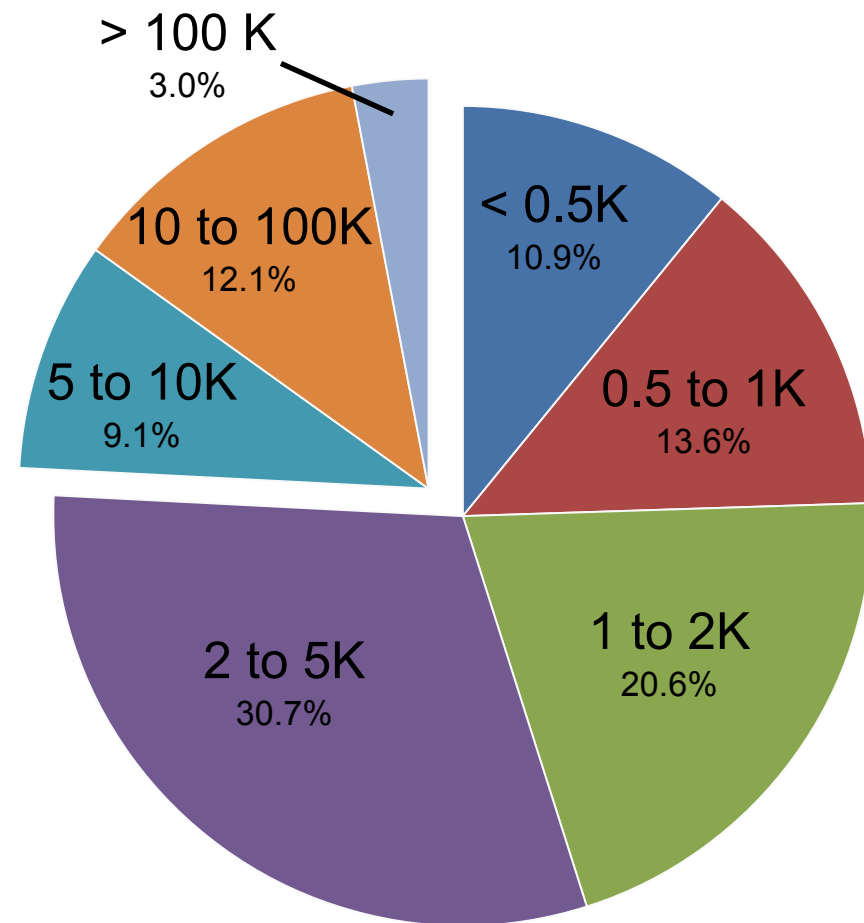
What is in ModelDB?

- Models for:
 - 181 cell types
 - 21+ species
 - 58 ion channels, pumps, etc
 - 169 topics (Alzheimer's, STDP, etc)
 - 25+ mammalian brain regions
- 1616 published models from 96 simulators/programming languages
- 737 NEURON models.
- 513 network models.





Files per Model



File Size

Only reuse
what you
understand

The easiest way to replicate someone else's results – a first step toward building on them – is to get their model code from a repository such as ModelDB.

But beware:



- They may be solving a different problem than you (with respect to species, temperature, age, etc).
- Their code may have bugs.

To reduce the risk of problems:



- Read the associated paper.
- Compare the model and results to other similar models.
- Examine the model with ModelView and/or psection.
- Test ion channels individually.
- Collaborate with an experimentalist.

Amyloid beta (IA block) effects on a model CA1 pyramidal cell (Morse et al. 2010)

Download zip file

Auto-launch

[Help downloading and running models](#)

Model Information

Model File

Citations

Model Views

Simulation Platform

3D Print

Download the displayed file

/

- CA1_abeta
 - translate
 - readme.html**
 - cacumm.mod
 - cagk.mod *
 - cal2.mod *
 - can2.mod *
 - cat.mod *
 - distr.mod *
 - h.mod
 - ipulse2.mod *
 - kadist.mod
 - kaprox.mod
 - kdrca1.mod
 - na3n.mod
 - naxn.mod ***
 - zcaquant.mod
 - aBeta.hoc

This is the readme file

Morse TM, et al. (2010) Abnormal Excitability in Alzheimer's: a model

The model code was contributed by Tom Morse. It was created (see paper for details) from earlier models (especially Migliore et al. 2005 and cal2.mod) and includes several modifications and interaction with other models to be installed.

To recreate figure 1, auto-launching the model

Under unix systems

In the expanded command window, run the simulation

Under Windows systems

Compile the model. A double click on mosinit.hoc will open the simulation window.

Under MAC OS X:

Other models using cagk.mod:

- [A model of unitary responses from A/C and PP synapses in CA3 pyramidal cells \(Baker et al. 2010\)](#)
- [CA1 pyramidal neuron: effects of R213Q and R312W Kv7.2 mutations \(Miceli et al. 2013\)](#)
- [CA3 pyramidal neuron \(Safiulina et al. 2010\)](#)
- [CA3 pyramidal neuron: firing properties \(Hemond et al. 2008\)](#)
- [Neuronal dendrite calcium wave model \(Neymotin et al, 2015\)](#)

Other models using naxn.mod:

- [CA1 pyramidal neuron: effects of R213Q and R312W Kv7.2 mutations \(Miceli et al. 2013\)](#)
- [CA1 pyramidal neuron: functional significance of axonal Kv7 channels \(Shah et al. 2008\)](#)
- [CA1 pyramidal neuron: rebound spiking \(Ascoli et al.2010\)](#)
- [CA1 pyramidal neuron: schizophrenic behavior \(Migliore et al. 2011\)](#)
- [CA1 pyramidal neuron: signal propagation in oblique dendrites \(Migliore et al 2005\)](#)
- [CA1 pyramidal neurons: binding properties and the magical number 7 \(Migliore et al. 2008\)](#)
- [CA1 pyramidal neurons: effect of external electric field from power lines \(Cavarretta et al. 2014\)](#)
- [CA1 pyramidal neurons: effects of Alzheimer \(Culmone and Migliore 2012\)](#)
- [CA1 pyramidal neurons: effects of Kv7 \(M-\) channels on synaptic integration \(Shah et al. 2011\)](#)
- [CA1 pyramidal neurons: effects of a Kv7.2 mutation \(Miceli et al. 2009\)](#)
- [Ca1 pyramidal neuron: reduction model \(Marasco et al. 2012\)](#)
- [Effect of the initial synaptic state on the probability to induce LTP and LTD \(Migliore et al. 2015\)](#)
- [Effects of electric fields on cognitive functions \(Migliore et al 2016\)](#)
- [Neuronal morphology goes digital ... \(Parekh & Ascoli 2013\)](#)
- [Spine head calcium in a CA1 pyramidal cell model \(Graham et al. 2014\)](#)

Amyloid beta (IA block) effects on a model CA1 pyramidal cell (Morse et al. 2010)

[Download zip file](#)[Auto-launch](#)[Help downloading and running models](#)**Model Information**[Model File](#)[Citations](#)[Model Views](#)[Simulation Platform](#)[3D Print](#)**Accession:**87284

The model simulations provide evidence oblique dendrites in CA1 pyramidal neurons are susceptible to hyper-excitability by amyloid beta block of the transient K⁺ channel, IA. See paper for details.

Reference:

1 . Morse TM, Carnevale NT, Mutalik PG, Migliore M, Shepherd GM (2010) Abnormal excitability of oblique dendrites implicated in early Alzheimer's: a computational study *Front. Neural Circuits* 4:16 [PubMed]

Model Information (Click on a link to find other models with that property)

Model Type: [Neuron or other electrically excitable cell;](#)

Brain Region(s)/Organism:

Cell Type(s): [Hippocampus CA1 pyramidal cell;](#)

Channel(s): [I Na,t; I L high threshold; I N; I T low threshold; I A; I K; I h;](#)

Gap Junctions:

Receptor(s):

Gene(s):

Transmitter(s):

Simulation Environment: [NEURON;](#)

Model Concept(s): [Dendritic Action Potentials; Active Dendrites; Detailed Neuronal Models; Pathophysiology; Aging/Alzheimer's;](#)

Implementer(s): [Carnevale, Ted \[Ted.Carnevale at Yale.edu\]; Morse, Tom \[Tom.Morse at Yale.edu\];](#)

Search NeuronDB for information about: [Hippocampus CA1 pyramidal cell; I Na,t; I L high threshold; I N; I T low threshold; I A; I K; I h;](#)

Morse et al. 2010



- 194 sections; 974 segments
- + 1 cell with morphology
- 0 artificial cells
- 0 NetCon objects
- 0 LinearMechanism objects
- + Temperature: 35°C
- + Density Mechanisms
- + 1 point processes (0 can receive events) of 1 base classes
- + 7 files shared with other ModelDB models
- + References

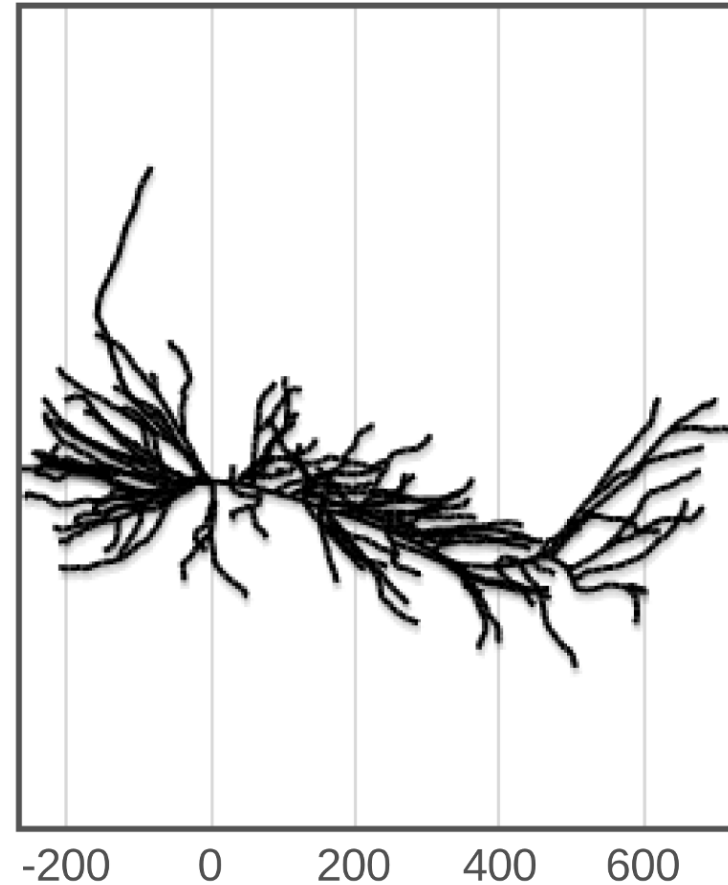
root: soma



X-Y

X-Z

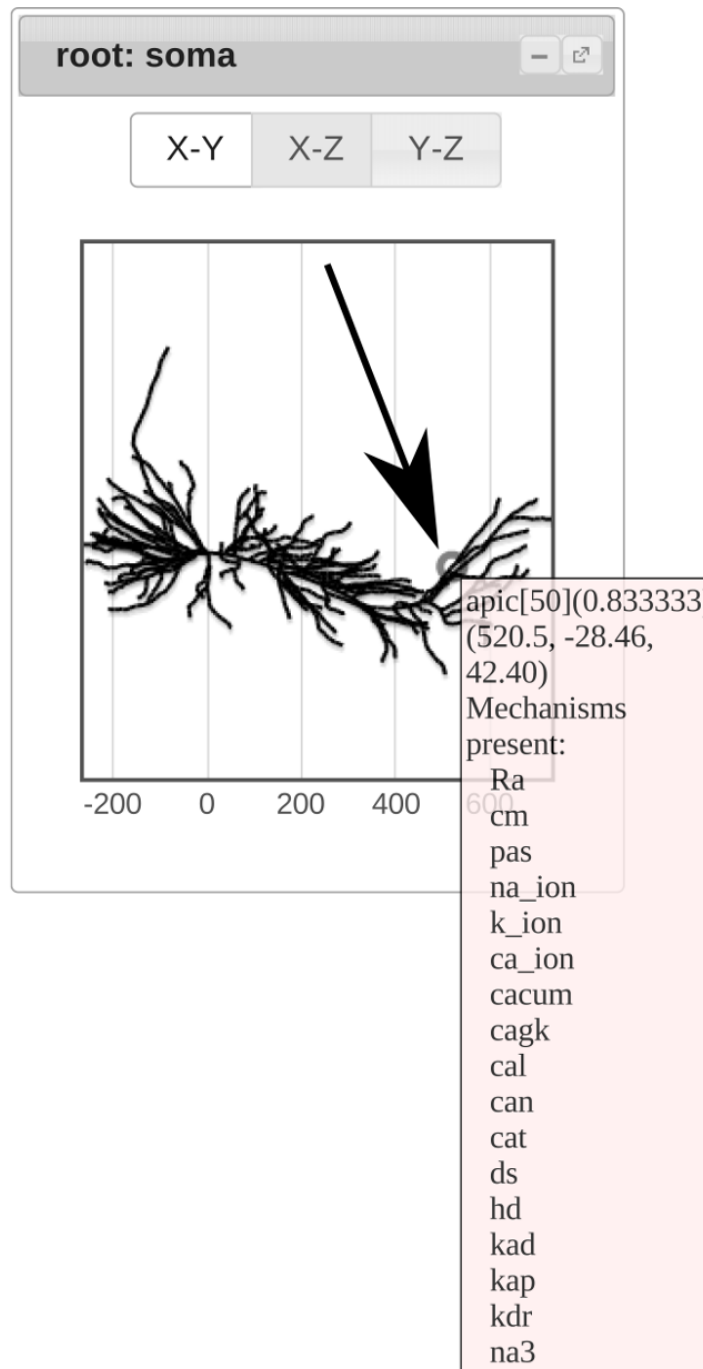
Y-Z



a

Morse et al. 2010

- 0 distinct values of nseg
- 18 inserted mechanisms
 - Ra
 - cm
 - + pas
 - + na_ion
 - + k_ion
 - ca_ion
 - + cacum (cacumm.mod)
 - + cagk (cagk.mod)
 - + cal (cal2.mod)
 - + can (can2.mod)
 - + cat (cat.mod)
 - ds (distr.mod)
 - + hd (h.mod)
 - + kad (kadist.mod)
 - + kap (kaprox.mod)
 - + kdr (kdrca1.mod)
 - + na3 (na3n.mod)
 - + nax (naxn.mod)



b

Morse et al. 2010

- Density Mechanisms
 - 18 mechanisms in use
 - Ra
 - cm
 - pas
 - na_ion
 - k_ion
 - ca_ion
 - cacum (cacumm.mod)
 - READs: ica
 - WRITEs: cai, Nonspecific Current
 - Present in 193 sections
 - cagk (cagk.mod)
 - READs: cai, ek
 - WRITEs: ik
 - Present in 193 sections
 - Possibly temperature dependent
 - cal (cal2.mod)

Morse et al. 2010

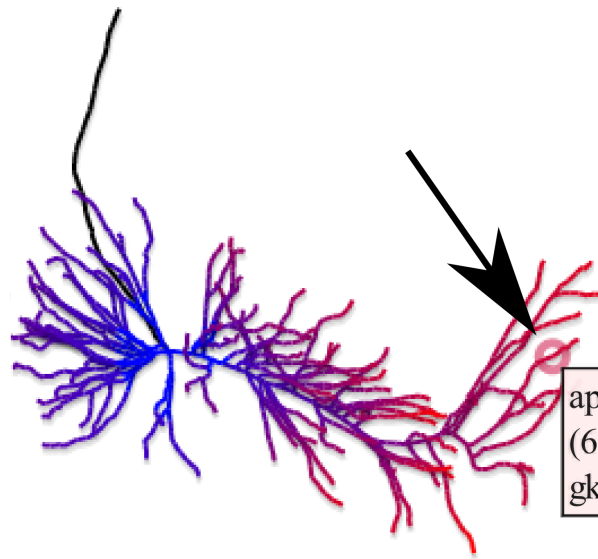
- can (can2.mod)
 - gcanbar
- cat (cat.mod)
 - gcatbar
- ds (distr.mod)
- hd (h.mod)
 - ghdbar
 - vhalfl
- kad (kadist.mod)
 - gkabar
- kap (kaprox.mod)
 - gkabar
- kdr (kdrca1.mod)
 - gkdrbar
- na3 (na3n.mod)
 - sh
 - gbar
 - ar
- nax (naxn.mod)

root: soma

X-Y

X-Z

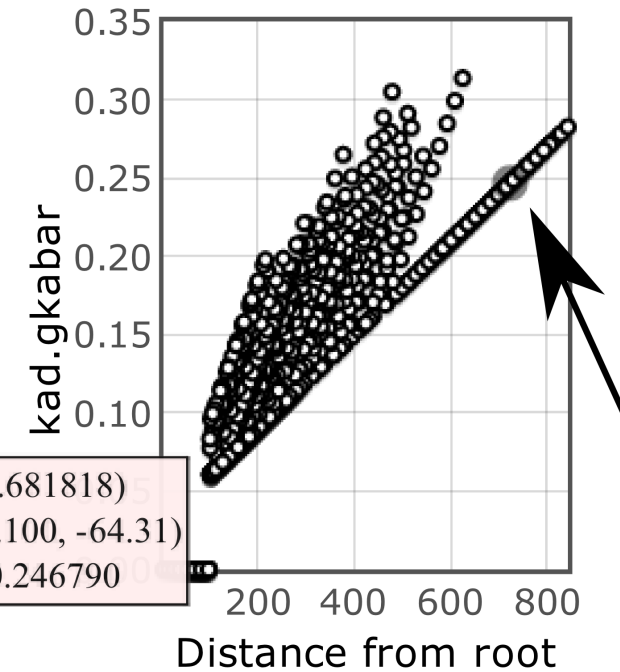
Y-Z



0

0.313714

Morse et al. 2010



Morse et al. 2010

- 7 files shared with other ModelDB models
 - cagk.mod
 - A model of unitary responses from A/C and PP synapses in CA3 pyramidal cells (Baker et al. 2010)
 - CA1 pyramidal neuron: effects of R213Q and R312W Kv7.2 mutations (Miceli et al. 2013)
 - CA3 pyramidal neuron (Safiulina et al. 2010)
 - CA3 pyramidal neuron: firing properties (Hemond et al. 2008)
 - distr.mod
 - cal2.mod
 - can2.mod
 - cat.mod
 - ipulse2.mod
 - naxn.mod

Morse et al. 2010

- ModelDB Entry
 - Run Protocol
 - Compiling
 - cd CA1_abeta
 - nrnivmodl
 - Launching NEURON
 - nrngui -python
 - Running
 - from neuron import h
 - h.load_file("mosinit.hoc")
 - h.fig1and2()

The screenshot shows the ICGenealogy web interface. At the top, there are tabs: 'Model Information', 'Model File' (selected), 'Citations', 'Model Views', 'Simulation Platform', and '3D Print'. Below the tabs, there is a button 'Download the displayed file' and a button 'ICGenealogy' which is circled in red. On the left, a file tree shows a directory structure with files like 'CA1_abeta', 'translate', 'readme.html', 'cacumm.mod', 'cagk.mod' (highlighted), 'cal2.mod', 'can2.mod', 'cat.mod', 'distr.mod', 'h.mod', 'ipulse2.mod', 'kadist.mod', 'kaprox.mod', 'kdrca1.mod', 'na3n.mod', 'naxn.mod', 'zcaquant.mod', 'aBeta.hoc', 'add_ca.hoc', 'bAP_peak_vecs.hoc', 'c91662.ses', 'c91662_Link.txt', 'cond_report.hoc', 'control_boxes.hoc', and 'distribute_currents.hoc'. The main panel on the right displays the contents of 'cagk.mod', including a title, units, neuron definition, and parameters.

Model Information | **Model File** | Citations | Model Views | Simulation Platform | 3D Print

Download the displayed file | **ICGenealogy**

[] /
 [] CA1_abeta
 [] translate
 [] readme.html
 [] cacumm.mod
 [] **cagk.mod ***
 [] cal2.mod *
 [] can2.mod *
 [] cat.mod *
 [] distr.mod *
 [] h.mod
 [] ipulse2.mod *
 [] kadist.mod
 [] kaprox.mod
 [] kdrca1.mod
 [] na3n.mod
 [] naxn.mod *
 [] zcaquant.mod
 [] aBeta.hoc
 [] add_ca.hoc
 [] bAP_peak_vecs.hoc
 [] c91662.ses
 [] c91662_Link.txt
 [] cond_report.hoc
 [] control_boxes.hoc
 [] distribute_currents.hoc

TITLE Cagk
 : Calcium activated K channel.
 : Modified from Moczydlowski and Latorre (1983) J. Gen. Physiol. 82

 UNITS {
 (molar) = (1/liter)
 }

 UNITS {
 (mV) = (millivolt)
 (mA) = (milliamp)
 (mM) = (millimolar)
 }

 NEURON {
 SUFFIX cagk
 USEION ca READ cai
 USEION k READ ek WRITE ik
 RANGE gbar,gkca,ik
 GLOBAL oinf, tau
 }

 UNITS {
 FARADAY = (faraday) (kilocoulombs)
 R = 8.313424 (joule/degC)
 }

 PARAMETER {
 celsius (degC)
 v (mV)
 gbar=.01 (rho/cm2) : Maximum Permeability
 cai (mM)
 ek (mV)

 d1 = .84
 d2 = 1.
 k1 = .48e-3 (mM)
 k2 = .13e-6 (mM)
 obar = .28 (/ms)
 bbar = .48 (/ms)
 st=1 (1)
 }

General data

- **ICG id:** 2464
- **ModelDB id:** [87284](#)
- **Reference:** Morse TM, Carnevale NT, Mutalik PG, Migliore M, Shepherd GM (2010): [Abnormal Excitability of Oblique Dendrites Implicated in Early Alzheimer's: A Computational Study](#).

Metadata classes

- **Animal Model:** rat
- **Brain Area:** hippocampus, CA1
- **Classes:** KCa
- **Ion Type:** K
- **Neuron Region:** unspecified
- **Neuron Type:** pyramidal cell
- **Runtime Q:** Q4 (slow)
- **Subtype:** not specified

Metadata generic

- **Age:** 7-14 weeks old.
- **Comments:** Calcium activated k channel, modified from moczydlowski and latorre (1983). From hemond et al. (2008), model no. 101629, with no changes (identical mod file). Animal model taken from chen (2005) which is used to constrain model. Channel kinetics from previous study on hippocampal pyramidal neuron (hemond et al. 2008)
- **Runtime:** 76.722

ICGenealogy:
ion channel metadata

When viewing most mod files describing an ion channel, an ICGenealogy button appears. Clicking this button loads the corresponding page of the ICGenealogy derived, information about the underlying data, etc) and response curves.

ICGenealogy

Browse ICG

Database

About

Contact

Sign In

Sign Up

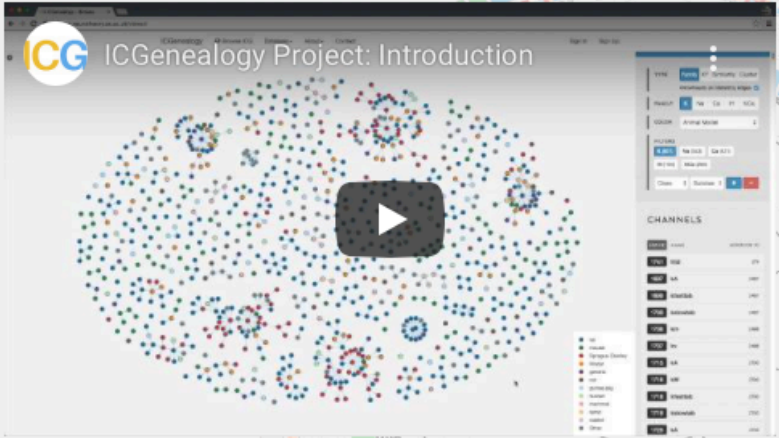
IonChannelGenealogy

Our database provides a comprehensive and quantitative assay of ion channel models currently available in the neuroscientific modeling community, all browsable in interactive visualizations.

Currently, the database contains 4815 models with **3706 quantitatively evaluated ion channels** for the [NEURON](#) simulator.

Learn more »

ICG ICGenealogy Project: Introduction



The ICG Project is an initiative of the [CNCB @ University of Oxford](#) in collaboration with the [LCN @ EPFL](#).

Channel Browser

A graphical user interface to all channels currently available in our database. We offer several interactive

Contribute

Together we can improve ICG! Upload your own channel models or submit tickets to correct existing ones should

API

All our data is accessible via an API. This enables you to run automated evaluations against current traces, or

ModelDB for meta- literature review

- Every model can be considered a review of the literature.
- ModelDB reveals what has been modeled in each cell type.
- Comparing models shows what mechanisms are considered critical by the community.

Hippocampus CA1 Pyramidal Cells

IA

- 47 models: 2796, 7386, 9769, 19696, 20212, 32992, 44050, 55035, ...

IK,Ca

- 11 models: 20212, 87284, 115356, 119266, 123927, 125152, ...

IM

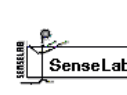
- 16 models: 2937, 20212, 66268, 112546, 115356, 118986, 119266, ...

26 currents, 6 transmitters, 10 receptors

Sharing your models



Advanced search



ModelDB

SimToolDB

ModelDB Help

User account

Login

Register

Find models by

Model name

First author

Each author

Region(circuits)

Find models for

Cell type

Current

Receptor

Gene

Transmitters

Topic

Simulators

Methods

Find models of

Realistic Networks

Neurons

Electrical synapses
(gap junctions)

Chemical synapses

Ion channels

Neuromuscular junctions

Axons

Other resources

ModelDB related
resources


Models in mercurial
repository

Submit Model

ModelDB provides an accessible location for storing and efficiently retrieving computational neuroscience models. ModelDB is tightly coupled with [NeuronDB](#). Models can be coded in any language for any environment. Model code can be viewed before downloading and browsers can be set to auto-launch the models. For further information, see [model sharing in general](#) and [ModelDB in particular](#).

Browse or search through over 1000 models using the navigation on the left bar or in the menu button on a mobile device. To search papers instead of models, go [here](#); this may be used to identify models whose paper cites or is cited by a given paper.

Tweets by @SenseLabProject

 **SenseLab** @SenseLabProject
New in #ModelDB: A Layer V CCS type pyramidal cell, inhibitory synapse current conduction (Kubota Y et al., 2015)
modeldb.yale.edu/183424
19 Apr

 **SenseLab** @SenseLabProject
Embed View on Twitter



@SenseLabProject



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Questions, comments, problems? Email the [ModelDB Administrator](#)

[How to cite ModelDB](#) [ModelDB Credits](#)

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Registered with NIF

[ModelDB Help](#)**User account**[Login](#)[Register](#)**Find models by**[Model name](#)[First author](#)[Each author](#)[Region\(circuits\)](#)**Find models for**[Cell type](#)[Current](#)[Receptor](#)[Gene](#)[Transmitters](#)[Topic](#)[Simulators](#)[Methods](#)**Find models of**[Realistic Networks](#)[Neurons](#)[Electrical synapses \(gap junctions\)](#)[Chemical synapses](#)[Ion channels](#)[Neuromuscular junctions](#)[Axons](#)**Other resources**[ModelDB related resources](#)[Computational neuroscience](#)

Submit New Model

Required information:

Your full name:**Your email address:****Zip file of model code:** No file chosen**Read-Write access code (15 character max):***Used as a password to only access this model***PubMed ID(s) or citation(s) associated with the model:***Only required for publicly shared models.**Citation(s) can be in any bibliographic format.*

You may with just the above information, but **to make your model more discoverable, please fill out as much of the next section as you can.** *Note:*

Your model will remain private until you request the ModelDB administrator make it public.

Let us find ModelDB keywords for you!

Click the button to automatically find, approve, and populate model entry keywords based on your paper abstract.

Additional information: *More information will help your model more discoverable*

Required

Your full name

Your email

Zip file of model

Read-Write

Used as a paper

PubMed ID

Only required

Citation(s) can

You may

Your model will

Automatic keyword identifier

Please paste your paper abstract here.

The integrative properties of cortical pyramidal dendrites are essential to the neural basis of cognitive function, but the impact of amyloid beta protein (abeta) on these properties in early Alzheimer's is poorly understood. In animal models, electrophysiological studies of proximal dendrites have shown that abeta induces hyperexcitability by blocking A-type K+ currents (I(A)), disrupting signal integration. The present study uses a computational approach to analyze the hyperexcitability induced in distal dendrites beyond the experimental recording sites. The results show that back-propagating action potentials in the dendrites induce hyperexcitability and excessive calcium concentrations not only in the main apical trunk of pyramidal cell dendrites, but also in their oblique dendrites. Evidence is provided that these thin branches are particularly sensitive to local reductions in I(A). The results suggest the hypothesis that the oblique branches may be most vulnerable to disruptions of I(A) by early exposure to abeta, and point the way to further experimental analysis of these actions as factors in the neural basis of the early decline of cognitive function in Alzheimer's.

Cancel

Submit

Let us find ModelDB keywords for you!

Click the button to automatically find, approve, and populate model entry keywords based on your paper abstract.

Additional information: More information will help your model more discoverable

Model Name

out as much of the next section as you can. Note:

Required

Your full name

Your email

Zip file of model

Read-Write

Used as a paper

PubMed ID

Only required

Citation(s) can

You may

Submit

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Let us find ModelDB keywords for you!

Click the button to automatically find, approve, and populate model entry keywords based on your paper abstract.

Additional information: More information will help your model more discoverable

Model Name

Automatic keyword identifier: results

Deselect keywords that do not describe the model, then press the button to accept the rest.


- ☒ Neuron or other electrically excitable cell
- ☒ Dendritic Action Potentials
- ☒ I Potassium
- ☒ Action Potentials
- ☒ Calcium dynamics
- ☒ I A
- ☒ Active Dendrites
- ☒ Aging/Alzheimer`s

Accept selected keywords

Other Neuron	
Model Neurotransmitters	<div></div>
Other Neurotransmitter	
Model Receptors	<div></div>
Other Receptor	
Model Currents	<div><div>x I Potassium</div><div>x I A</div><div></div></div>
Other Current	
Gap Junctions	<div></div>
Gene	<div></div>
Other Gene	
Model Type	<div><div>x Neuron or other electrically excitable cell</div><div></div></div>
Other Model Type	
Model Concept	<div><div>x Dendritic Action Potentials</div><div>x Action Potentials</div><div>x Calcium dynamics</div><div>x Active Dendrites</div><div>x Aging/Alzheimer`s</div><div></div></div>
Other Concept	
Simulator software	<div></div>
Other Simulator	
Region Organism	<div></div>
Implemented by	<div></div>

ModelDB redesign

- Updated look.
- More integrated ModelView data, with support for more simulator types.
- More emphasis on analysis.
- Mobile device friendly.
- Simpler API.
- (Soon to be) public GitHub; submit issues and pull requests.
- Current status: see <http://52.90.37.175>

 ModelDB Login Browse ▾ Analysis ▾ More ▾

Amyloid beta (IA block) effects on a model CA1 pyramidal cell (Morse et al. 2010)

Overview Files Mechanisms Parameters Citations

The model simulations provide evidence oblique dendrites in CA1 pyramidal neurons are susceptible to hyper-excitability by amyloid beta block of the transient K⁺ channel, IA. See paper for details.

Model Type: [Neuron or other electrically excitable cell](#)

Cell Type(s): [Hippocampus CA1 pyramidal GLU cell](#)

Currents: [I_{Na,t}](#) ; [I_L high threshold](#) ; [I_N](#) ; [I_T low threshold](#) ; [I_A](#) ; [I_K](#) ; [I_h](#) ; [I_{K,Ca}](#)


Model Concept(s): [Dendritic Action Potentials](#) ; [Active Dendrites](#) ; [Detailed Neuronal Models](#) ; [Pathophysiology](#) ; [Aging/Alzheimer`s](#)

Simulation Environment: [NEURON](#)

Implementer(s): [Carnevale, Ted \[Ted.Carnevale at Yale.edu\]](#) ; [Morse, Tom \[Tom.Morse at Yale.edu\]](#)

References:

[Morse TM, Carnevale NT, Mutalik PG, Migliore M, Shepherd GM. \(2010\). Abnormal Excitability of Oblique](#)



[Show Diameter](#)

[Simulation Platform !\[\]\(cc550a6475ae0a4b1b30a2c3ecc0f2f6_img.jpg\)](#)
[View on GitHub !\[\]\(c69b0103e9852262815d3edabc991a19_img.jpg\)](#)



NeuroMorpho.Org



Version 8.0 - Released: 6/29/2020 - Content: 131960 neurons

Total number of downloads: 13894177

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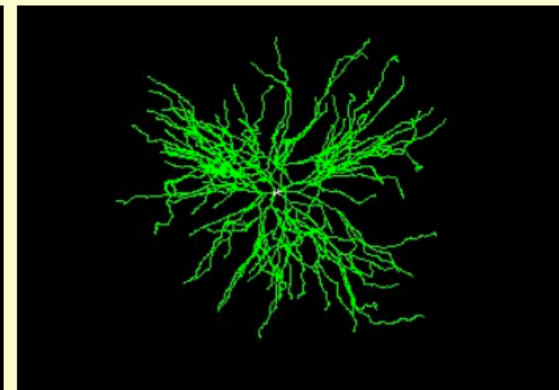
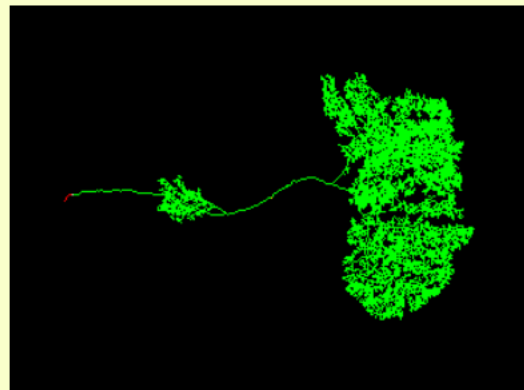
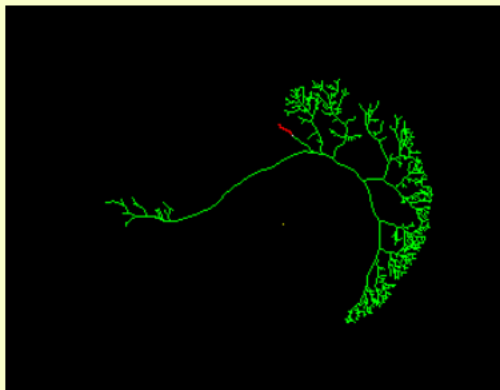
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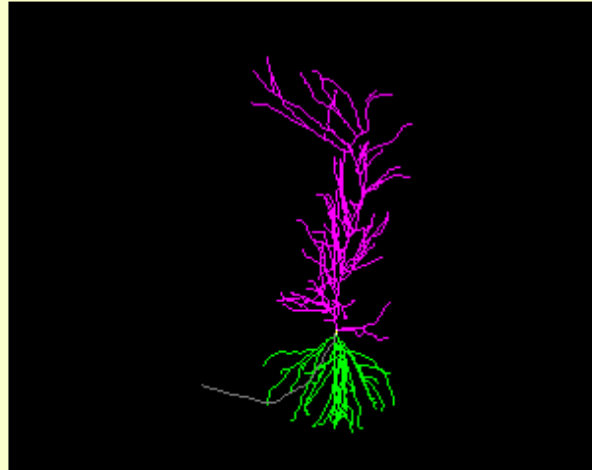
0

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Reconstructions from 384 brain regions >3.57million hours of manual reconstructions



131,960 reconstructions · 641 cell types · 384 brain regions

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Standardized:
Always SWC

Original format:
Could be anything

Details about selected neuron

NeuroMorpho.Org ID : NMO_00227

Neuron Name : c91662

Archive Name : Amaral

Species Name : rat

Strain : Sprague-Dawley

Structural Domains : Dendrites, Soma, Axon

Physical Integrity : Dendrites Complete, Axon Incomplete

Morphological Attributes : Diameter, 3D, Angles

Min Age : 33.0 days

Metadata

NeuroMorpho.Org ID : NMO_00082
Neuron Name : n401
Archive Name : Turner
Species Name : rat
Strain : Fischer 344
Structural Domains : Dendrites, Soma, No Axon
Physical Integrity : Dendrites Complete
Morphological Attributes : Diameter, 3D, Angles
Min Age : 2.0 months
Max Age : 8.0 months
Gender : Male/Female
Min Weight : 200 grams
Max Weight : 350 grams
Development : young
Primary Brain Region : hippocampus
Secondary Brain Region : CA1
Tertiary Brain Region : Not reported
Primary Cell Class : principal cell
Secondary Cell Class : pyramidal
Tertiary Cell Class : Not reported
Original Format : CVAPP.swc
Experiment Protocol : in vivo
Experimental Condition : Control
Staining Method : biocytin
Slicing Direction : coronal
Slice Thickness : 80.00 μm
Tissue Shrinkage : Reported 25% in xy, 75% in z
Corrected 133% in xy, 400% in z
Objective Type : oil
Magnification : 100x
Reconstruction Method : Neurolucida
Date of Deposition : 2005-12-31
Date of Upload : 2006-08-01

Soma Surface : 903.25 μm^2
Number of Stems : 7
Number of Bifurcations : 113
Number of Branches : 233
Overall Width : 363.7 μm
Overall Height : 717.18 μm
Overall Depth : 364.21 μm
Average Diameter : 1.16 μm
Total Length : 22216.3 μm
Total Surface : 84796.1 μm^2
Total Volume : 30674.3 μm^3
Max Euclidean Distance : 668.56 μm
Max Path Distance : 1893.37 μm
Max Branch Order : 25
Average Contraction : 0.7
Total Fragmentation : 5460
Partition Asymmetry : 0.56
Average Rall's Ratio : 1.78
Average Bifurcation Angle Local : 89.59°
Average Bifurcation Angle Remote : 75.23°
Fractal Dimension : 1.07

THE JOURNAL OF COMPARATIVE NEUROLOGY 391:335-352 (1998)

Dendritic Properties of Hippocampal CA1 Pyramidal Neurons in the Rat: Intracellular Staining In Vivo and In Vitro

G.K. PYAPALI,^{1,2} A. SIK,³ M. PENTTONEN,³ G. BUZSAKI,² AND D.A. TURNER^{1,2,4*}

¹Department of Neurosurgery, Duke University, Durham, North Carolina 27710

²Durham Veterans Affairs Medical Center, Durham, North Carolina 27710

³Center for Molecular and Behavioral Neuroscience, Rutgers,

The State University of New Jersey, Newark, New Jersey 07102

⁴Department of Neurobiology, Duke University, Durham, North Carolina 27710

Not
everything
was made for
you

Not every morphology was reconstructed with the intent of being in a simulation.

Potential factors affecting the quality of the data:

- histology
 - staining, amputation, shrinkage
- physics
 - diameter
- spines

Before using a morphology found online, always read the associated paper(s) to make sure you understand any limitations of the reconstruction.

For example, why did they make this? Were they studying a disease (e.g. Alzheimer's) that alters morphology?

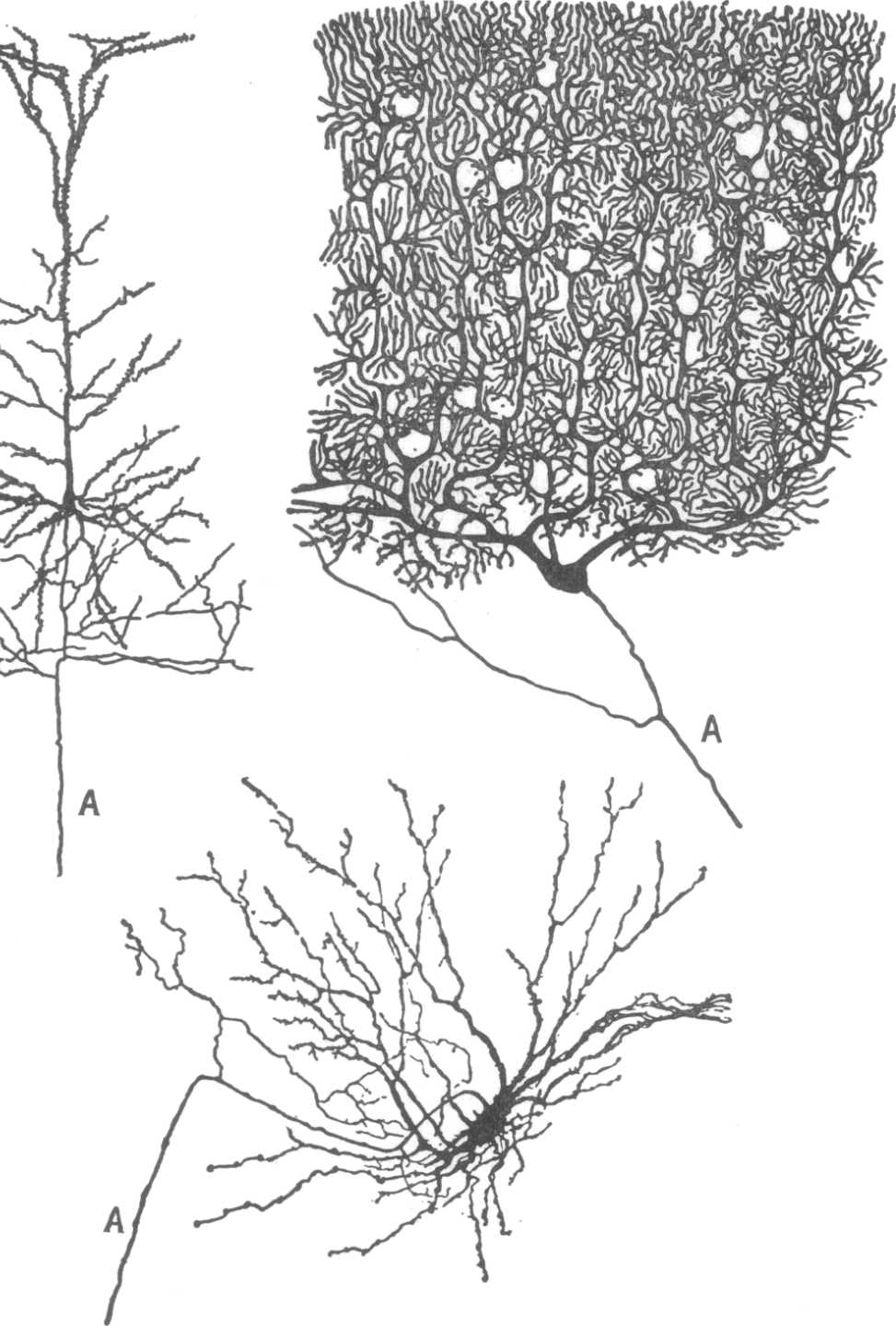
Qualitative tests

Look for orphan sections and bottlenecks.

Insert pas, set Ra and g_pas = pas.g low. Inject large depolarizing current at soma. Examine a PlotShape of v.

Look for z-axis drift and backlash.
Rotate the cell on a PlotShape and look for abrupt jumps.

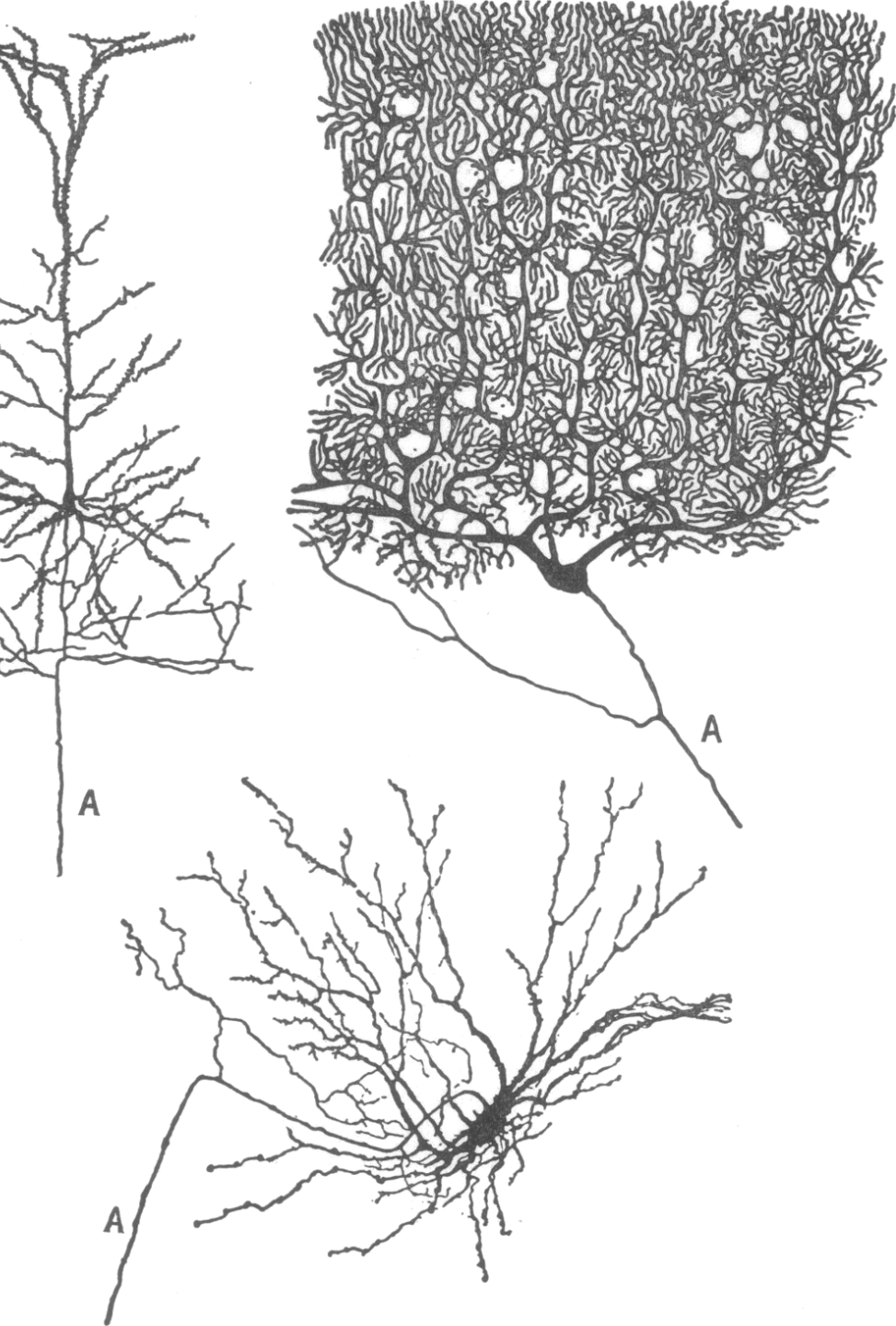
Are diameters constant or varying? Are they reasonable?



Loading Morphologies

```
from neuron import h
h.load_file('import3d.hoc')

cell = h.Import3d_SWC_read()
cell.input('filename.swc')
i3d = h.Import3d_GUI(cell, False)
i3d.instantiate(None) # or i3d.instantiate(self)
```



Plotting Morphologies

```
import plotly
```


```
ps = h.PlotShape(False)
```

```
ps.scale(-80, 40)
```

```
ps.variable('v')
```


```
ps.plot(plotly).show()
```

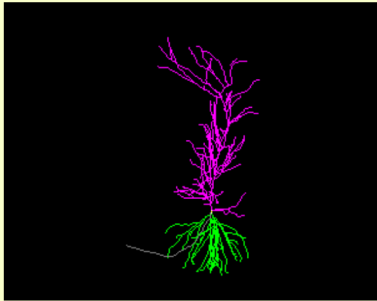
Example




NeuroMorpho.Org

Version 8.0 - Released: 6/29/2020 - Content: 131960 neurons

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[3D Neuron Viewer - Java, legacy](#)
[3D Neuron Viewer - WebGL, novel](#)
[Animation](#)

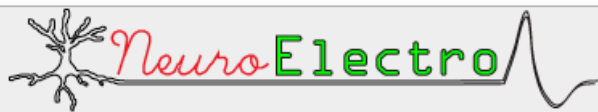
Details about selected neuron

NeuroMorpho.Org ID	: NMO_00227
Neuron Name	: c91662
Archive Name	: Amaral
Species Name	: rat
Strain	: Sprague-Dawley
Structural Domains	: Dendrites, Soma, Axon
Physical Integrity	: Dendrites Complete, Axon Incomplete
Morphological Attributes	: Diameter, 3D, Angles
Min Age	: 33.0 days

```
# Original file c91662.swc edited using StdSwc version 1.31 on 11/10/13.
# Irregularities and fixes documented in c91662.swc.std. See StdSwc1.31.doc for more information.
#
# Neurolucida to SWC conversion from L-Measure. Sridevi Polavaram: spolavar@gmu.edu
# Original fileName:C:\Users\praveen\Desktop\Uzma>ErrorArchives\ToBeProcessed\Amaral\asc\c91662.asc
#The original file has a single soma contour that is averaged into 3 soma points
# NEUROMANTIC V1.6.3 (10/18/2013 6:55:13 PM): Saved to c91662-T1.swc
1 1 0.0 0.0 0.0 8.8677 -1
2 1 1.13 8.71 1.2 8.8677 1
3 1 -1.13 -8.71 -1.2 8.8677 1
4 4 -1.86 11.06 -0.47 1.85 1
5 4 -1.94 19.75 -0.65 1.6 4
6 4 -2.52 31.1 -1.23 1.35 5
7 4 -2.94 39.91 -2.02 1.35 6
8 4 -2.55 49.45 -1.47 1.1 7
9 4 -2.61 56.49 -0.77 1.1 8
10 4 -1.17 70.15 -1.59 1.1 9
11 4 2.04 83.45 -1.43 1.1 10
12 4 1.89 91.65 -1.68 1.1 11
13 4 4.35 106.58 -1.57 1.1 12
14 4 5.09 115.06 -1.02 1.1 13
15 4 7.16 126.11 -1.93 1.1 14
16 4 7.13 129.63 -1.58 1.1 15
17 4 9.19 135.02 -2.02 1.1 16
18 4 11.82 145.77 -1.23 1.1 17
19 4 13.47 151.73 -1.73 0.9 18
20 4 14.65 157.05 -0.86 0.9 19
21 4 15.6 164.42 0.15 0.9 20
22 4 17.22 166.37 -0.76 0.9 21
23 4 17.27 175.11 -1.42 0.8 22
24 4 17.43 180.1 -0.87 0.8 23
25 4 18.41 192.2 -0.94 0.8 24
26 4 20.93 207.62 -0.75 0.8 25
27 4 22.64 214.07 -1.18 0.8 26
28 4 26.23 231.47 2.1 0.8 27
29 4 28.89 246.23 3.3 0.8 28
30 4 31.83 252.62 2.17 0.8 29
31 4 33.06 266.68 2.37 0.8 30
32 4 36.17 276.41 2.67 0.8 31
33 4 38.23 281.8 2.23 0.8 32
34 4 43.26 297.81 3.18 0.8 33
35 4 49.51 314.69 4.04 0.8 34
36 4 51.98 319.51 3.66 0.8 35
37 4 55.37 329.56 5.11 0.8 36
38 4 59.09 339.26 5.05 0.8 37
39 4 63.87 351.94 0.9 0.8 38
40 4 65.01 361.5 0.62 0.8 39
41 4 64.84 372.28 0.1 0.6 40
42 4 63.56 393.06 -1.8 0.6 41
43 4 63.28 401.93 -3.07 0.6 42
44 4 62.98 405.13 -3.87 0.6 43
45 4 61.56 411.24 -2.61 0.6 44
46 4 57.99 423.02 -3.47 0.6 45
```

<http://tinyurl.com/neuromorpho-c91662>

<http://tinyurl.com/neuromorpho-c91662-swc>



NeuroElectro Publications

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NeuroElectro: *organizing information on cellular neurophysiology.*

The goal of the NeuroElectro Project is to extract information about the electrophysiological properties (e.g. [resting membrane potentials](#) and [membrane time constants](#)) of diverse neuron types from the existing literature and place it into a centralized database.

Published literature

Novel subcellular distribution pattern of A-type K⁺ channels on neuronal surface.
Unique clustering of A-type potassium channels on different cell types of the main olfactory bulb.
Kollo M, Holderith N, Antal M, Nusser Z.
Theoretical and functional studies predicted a highly non-uniform distribution of voltage-gated ion channels on the neuronal surface. This was confirmed by recent immunolocalization experiments for Na⁺, Ca²⁺, hyperpolarization activated mixed cation and K⁺ channels. These experiments also indicated that some K⁺ channels were clustered in synaptic or non-synaptic membrane specializations. Here we analysed the subcellular distribution of Kv4.2 and Kv4.3 subunits in the rat main olfactory bulb at high resolution to address whether clustering characterizes their distribution, and whether they are concentrated in synaptic or non-synaptic junctions. The cell surface distribution of the Kv4.2 and Kv4.3 subunits is highly non-uniform. Strong Kv4.2 subunit-immunopositive clusters were detected in intercellular junctions made by mitral, external puffed and granule cells (GECs). We also found Kv4.3 subunit-immunopositive clusters in periglomerular (PGC), deep short-axon and GECs. In the juxttaglomerular region some calretinin-immunopositive glial cells envelop neighboring PGC somata in a cap-like manner. Kv4.3 subunit clusters are present in the cap membrane that directly contacts the PGC, but not the one that faces the neuropil. In membrane specializations established by members of the same cell type, K⁺ channels are enriched in both membranes, whereas specializations between different cell types contain a high density of channels asymmetrically. None of the K⁺ channel-rich junctions showed any of the ultrastructural features of known chemical synapses. Our study provides evidence for highly non-uniform subcellular distributions of A-type K⁺ channels and predicts their involvements in novel

Physiology database

Olfactory Bulb Mitral Cell

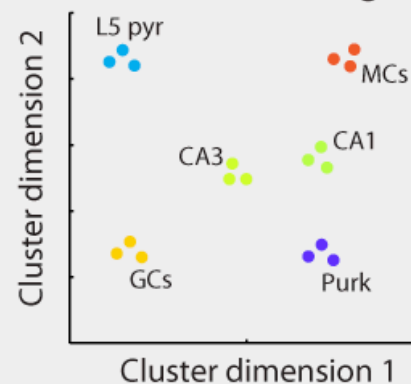
Input resistance	200 MΩ
V _{rest}	-65 mV
Spike width	1 ms
...	

CA1 Pyramidal Cell

Input resistance	400 MΩ
V _{rest}	-70 mV
Spike width	.5 ms
...	

Extracted from Literature

Neuron clustering



Our goal is to facilitate the discovery of [neuron-to-neuron relationships](#) and better understand the role of functional diversity across neuron types.


New here? Watch a short video to see what we're about.



NeuroElectro.org description



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Learn about the Hodgkin Huxley model



Simulate electrophysiologically detailed cell models



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