# nideconv Documentation

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# Getting started

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Nideconv is an easy-to use Python library that can perform automated deconvolution of (primarily) slowly fluctuating (proxies of) neural signals like pupil size and BOLD fMRI. It was developed at the Vrije Universiteit and the Spinoza Centre for Neuroimaging by Gilles de Hollander and Tomas Knapen.

# Installing nideconv

# 1.1 From PyPi

When the first Beta of nideconv will be released, it will be possible to install nideconv from pip:

pip install nideconv

## 1.2 From Github

Right now you can clone the main branch of nideconv using git

git clone https://github.com/VU-Cog-Sci/nideconv

Or download and unpack the zip file from Github under Clone and download -> Download ZIP

Then go to the directory in which the package was cloned

cd nideconv

and install the Python package

```
python setup.py install
```

Note: Click here to download the full example code

### What is Deconvolution?

Neuroscientists (amongst others) are often interested in time series that are derived from neural activity, such as fMRI BOLD and pupil dilation. However, for some classes of data (notably, pupil dilation and fMRI BOLD), neural activity gets temporally delayed and dispersed. This means that if the time series is related to some behavioral events that are close together in time, these event-related responses will contaminate each other.

```
from nideconv import simulate
import matplotlib.pyplot as plt
import seaborn as sns
sns.set_style('white')
sns.set_context('notebook')
```

## 2.1 Simulate data

Here we simulate fMRI data with a "cue - stimulus" design. There are four cues and stimulus pairs. The cue is always followed by a stimulus in 1, 2, 3, or 4 seconds. The cue leads to a small de-activation (0.5 % signal change), the stimulus a slight activation (1.0 % signal change)

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```
stim_pars]
```

## 2.2 Plot simulated data

```
data.plot()
sns.despine()
for onset in cue_onsets:
    l1 =plt.axvline(onset, c='r')
for onset in stim_onsets:
    l2 =plt.axvline(onset, c='g')
plt.legend([l1, l2], ['Cue', 'Stimulus'])
plt.gcf().set_size_inches(10, 4)
```



### 2.3 Underlying data-generating model

Because we simulated the data, we know that the event-related responses should exactly follow the *canonical Hemo*dynamic Response Function [1]\_are

```
from nideconv.utils import double_gamma_with_d
import numpy as np
plt.figure(figsize=(12, 4))
t = np.linspace(0, 20, 100)
ax1 = plt.subplot(121)
plt.title('Ground truth cue-related response')
```

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```
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```



### 2.4 Naive approach: epoched averaging

A simple approach that is more appropriate for fast electrphysiological signals like EEG and MEG would be to select little chunks of the time series, corresponding to the onset of our events-of-interest and the first 20 seconds ("epoching").

nideconv

## 2.5 References

Total running time of the script: (0 minutes 1.155 seconds)

# CHAPTER $\mathbf{3}$

Usage examples

Note: Click *here* to download the full example code

# 3.1 Cortical depth estimation from MGDM segmentation

This example shows how to obtain a cortical laminar depth representation from blabla.

This is like super cool.

import response\_fytter

Total running time of the script: (0 minutes 0.000 seconds)

# Deconvolution of a single time series

# 4.1 ResponseFitter

ResponseFitter takes an input signal and performs deconvolution on it. To do this, it requires event times, and possible covariates. ResponseFytter can, for each event\_type, use different basis function sets, see Event.

### **Methods**

add_confounds(name, confound)	Add a timeseries or set of timeseries to the general
	design matrix as a confound
<pre>add_event(event_name[, onset_times,])</pre>	create design matrix for a given event_type.
<pre>get_epochs(onsets, interval[,])</pre>	Return a matrix corresponding to specific onsets,
	within a given interval.
get_rsq()	calculate the rsq of a given fit.
<pre>predict_from_design_matrix([X])</pre>	predict a signal given a design matrix.
regress([type, cv, alphas, store_residuals])	Regress a created design matrix on the input_data.
ridge_regress([cv, alphas, store_residuals])	run CV ridge regression instead of ols fit.

add_intercept	
get_residuals	
get_time_to_peak	
get_timecourses	
plot_timecourses	

### add\_confounds (name, confound)

Add a timeseries or set of timeseries to the general design matrix as a confound

### **Parameters**

**confound** [array] Confound of (n\_timepoints) or (n\_timepoints, n\_confounds)

add\_event (event\_name, onset\_times=None, basis\_set='fir', interval=[0, 10], n\_regressors=None, durations=None, covariates=None, \*\*kwargs)

create design matrix for a given event\_type.

### Parameters

event\_name [string] Name of the event\_type, used as key to lookup this event\_type's characteristics

**\*\*kwargs** [dict] keyward arguments to be internalized by the generated and internalized Event object. Needs to consist of the necessary arguments to create an Event object, see Event constructor method.

### get\_epochs (onsets, interval, remove\_incomplete\_epochs=True)

Return a matrix corresponding to specific onsets, within a given interval. Matrix size is (n\_onsets, n\_timepoints\_within\_interval).

Note that any events that are in the ResponseFitter-object will be regressed out before calculating the epochs.

#### get\_rsq()

calculate the rsq of a given fit. calls predict\_from\_design\_matrix to predict the signal that has been fit

### predict\_from\_design\_matrix(X=None)

predict a signal given a design matrix. Requires regression to have been run.

#### **Parameters**

X [np.array, (timepoints, n\_regressors)] the design matrix for which to predict data.

**regress** (*type='ols'*, *cv=20*, *alphas=None*, *store\_residuals=False*) Regress a created design matrix on the input\_data.

Creates internal variables betas, residuals, rank and s. The beta values are then injected into the event\_type objects the ResponseFitter contains.

### Parameters

**type** [string, optional] the type of fit to be done. Options are 'ols' for np.linalg.lstsq, 'ridge' for CV ridge regression.

### **ridge\_regress** (*cv*=20, *alphas=None*, *store\_residuals=False*)

run CV ridge regression instead of ols fit. Uses sklearn's RidgeCV class

### Parameters

cv [int] number of cross-validation folds

alphas [np.array] the alpha/lambda values to try out in the CV ridge regression

# Deconvolution of a group of timeseries (level 2 analysis)

In most neuroscience studies, you want to analyze a group of subjects, each of which has one or multiple timeseries (often called 'runs' in fMRI).

To analyze those, you can use the GroupResponseFitter

## 5.1 GroupResponseFitter

class	nideconv.GroupResponseFitter(timeseries	, behavior,	input_sample_rate,	oversam-
	ple_desig	$m_matrix=20$ ,	confounds=None,	concate-
	nate_runs	=True, *args,	**kwargs)	
C	an fit a group of individual subjects and/or runs using a	high-level int	erface.	

### Methods

add_event	
fit	
get_conditionwise_timecourses	
get_subjectwise_timecourses	
get_timecourses	
plot_groupwise_timecourses	

# Voxelwise deconvolution on Nifti Images

# 6.1 NiftiResponseFytter

### Methods

add_confounds(name, confound)	Add a timeseries or set of timeseries to the general	
	design matrix as a confound	
<pre>add_event(event_name[, onset_times,])</pre>	create design matrix for a given event_type.	
<pre>get_epochs(onsets, interval[,])</pre>	Return a matrix corresponding to specific onsets,	
	within a given interval.	
get_rsq()	calculate the rsq of a given fit.	
regress([type, cv, alphas, store_residuals])	Regress a created design matrix on the input_data.	

add_intercept	
get_residuals	
get_time_to_peak	
get_timecourses	
plot_timecourses	
predict_from_design_matrix	
ridge_regress	

### add\_confounds (name, confound)

Add a timeseries or set of timeseries to the general design matrix as a confound

### Parameters

confound [array] Confound of (n\_timepoints) or (n\_timepoints, n\_confounds)

add\_event (event\_name, onset\_times=None, basis\_set='fir', interval=[0, 10], n\_regressors=None, durations=None, covariates=None, \*\*kwargs) create design matrix for a given event\_type.

#### **Parameters**

- event\_name [string] Name of the event\_type, used as key to lookup this event\_type's characteristics
- **\*\*kwargs** [dict] keyward arguments to be internalized by the generated and internalized Event object. Needs to consist of the necessary arguments to create an Event object, see Event constructor method.
- get\_epochs (onsets, interval, remove\_incomplete\_epochs=True)

Return a matrix corresponding to specific onsets, within a given interval. Matrix size is (n\_onsets, n\_timepoints\_within\_interval).

Note that any events that are in the ResponseFitter-object will be regressed out before calculating the epochs.

### get\_rsq()

calculate the rsq of a given fit. calls predict\_from\_design\_matrix to predict the signal that has been fit

### predict\_from\_design\_matrix(X=None)

predict a signal given a design matrix. Requires regression to have been run.

#### **Parameters**

X [np.array, (timepoints, n\_regressors)] the design matrix for which to predict data.

**regress** (*type='ols'*, *cv=20*, *alphas=None*, *store\_residuals=False*) Regress a created design matrix on the input\_data.

Creates internal variables betas, residuals, rank and s. The beta values are then injected into the event\_type objects the ResponseFitter contains.

#### **Parameters**

**type** [string, optional] the type of fit to be done. Options are 'ols' for np.linalg.lstsq, 'ridge' for CV ridge regression.

### ridge\_regress(\*args, \*\*kwargs)

run CV ridge regression instead of ols fit. Uses sklearn's RidgeCV class

#### **Parameters**

cv [int] number of cross-validation folds

alphas [np.array] the alpha/lambda values to try out in the CV ridge regression

### Simulate data

### 7.1 Simulate fMRI data

### 

Simulates an fMRI experiment and returns a pandas DataFrame with the resulting time series in an analysisready format.

By default a single run of a single subject is simulated, but a larger number of subjects, runs, and ROIs can also be simulated.

### Parameters

**conditions** [list of dictionaries or *None*] Can be used to customize different conditions. Every conditions is represented as a dictionary in this list and has the following form:

```
[{'name':'Condition A',
  'mu_group':1,
  'std_group':0.1},
  {'name':'Condition B',
  'mu_group':1,
  'std_group':0.1}]
```

*mu\_group* indicates the mean amplitude of the response to this condition across subjects. *std\_group* indicates the standard deviation of this amplitude across subjects.

Potentially, customized onsets can also be used as follows:

```
{'name':'Condition A',
    'mu_group':1,
    'std_group':0.1
    'onsets':[10, 20, 30]}
```

**TR** [float] Indicates the time between volume acquistisions in seconds (Inverse of the sample rate).

n\_subjects [int] Number of subjects.

n\_runs [int] Number of runs per subject.

**n\_trials** [int] Number of trials *per condition per run*. Only used when no custom onsets are provided (see *conditions*).

run\_duration [float] Duration of a single run in seconds.

noise\_level [float] Standard deviation of Gaussian noise added to time series.

**n\_rois** [int] Number of regions-of-interest. Determines the number of columns of *data*.

#### Returns

**data** [DataFrame] Contains simulated time series with subj\_idx, run and time (s) as index. Columns correspond to different ROIs

**onsets** [DataFrame] Contains used event onsets with subj\_idx, run and trial type as index.

parameters [DataFrame] Contains parameters (amplitude) of the different event type.

### **Other Parameters**

- **oversample** [int] Determines how many times the kernel is oversampled before convolution. Should usually not be changed.
- **kernel** [str] Sets which kernel to use for response function. Currently only '*double\_hrf*' can be used.

#### **Examples**

By default, simulate\_fmri\_experiment simulates a 5 minute run with 40 trials for one subject

```
>>> data, onsets, params = simulate_fmri_experiment()
>>> print(data.head())
                    area 1
subj_idx run t
     1 0.0 -1.280023
1
             1.0 0.908086
             2.0 0.850847
             3.0 -1.010475
             4.0 -0.299650
>>> print(data.onsets)
                              onset
subj_idx run trial_type
        1
                         94.317361
1
           A
            А
                         106.547084
                         198.175115
             А
                          34.941112
             А
                          31.323272
             А
>>> print(params)
                        amplitude
subj_idx trial_type
                           1.0
1
         Α
         В
                           2.0
```

With n\_subjects we can increase the number of subjects

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