

# Offline Analysis Tools

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Working with TDT data in Python



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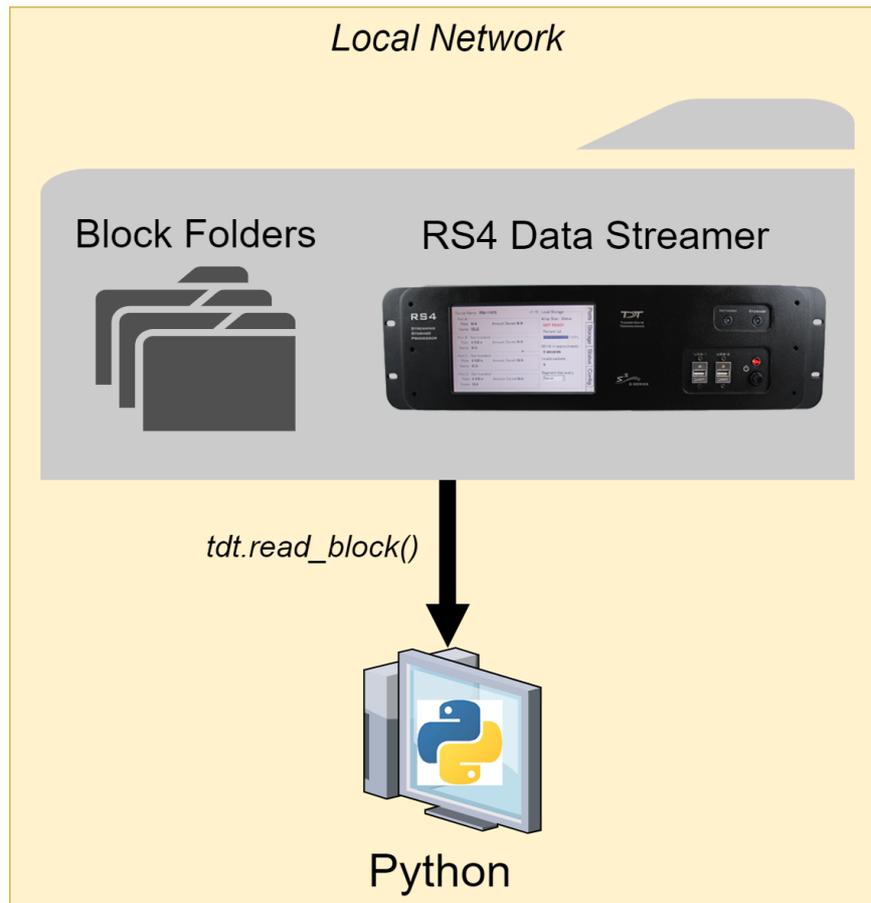
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# Overview of Python Offline Analysis Tools

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Read block files directly from disk or SEV files directly from RS4.



See [TDT Data Storage](#) for a description of the folder structure.

## read\_block

---

`read_block` is an all-in-one function for reading TDT data into Python. It needs only one input: the block path.

```
import tdt
data = tdt.read_block('C:\TDT\TDTEexampleData\Algernon-180308-130351')
```

`read_block` will return a structure containing all recorded data from that block, organized by type. See [TDT Data Types](#) for a description of the data types.

The returned structure also contains an **info** field with block start/stop times, duration, and information about the Subject, User, and Experiment that it came from (if the block was created in Synapse).

`read_block` uses input parameters to refine the imported data. To extract specific event types only, use the `evtype` parameter. For example, to import epochs and snippets only, use this:

```
data = tdt.read_block('C:\TDT\TDTEexampleData\Algernon-180308-130351',
                    evtype=['epochs', 'snips'])
```

Use the `store` parameter to extract a particular data store by name, in this example a streaming event called 'Wav1'. Combine this with the `channel` parameter to extract a single channel, or list of channels, in this case channels 2 and 4:

```
data = tdt.read_block('C:\TDT\TDTEexampleData\Algernon-180308-130351',
                    store='Wav1', channel=[2,4])
```

You can also filter by time, if you are only interested in portions of the recording, or if the entire recording won't fit into available memory (RAM) at one time. Use the `t1` and `t2` parameters to specify the start and stop time, in seconds, to retrieve from the block. This example reads only from time `t1=10s` to time `t2=20s` of the block into Python:

```
data = tdt.read_block('C:\TDT\TDTEexampleData\Algernon-180308-130351',
                    t1=10, t2=20)
```

`read_block` offers many more useful options that are described in its help documentation.

```
print(tdt.read_block.__doc__)
```

## epoc\_filter

---

`epoc_filter` applies advanced epoc filtering to extracted data. For example, if you only want to look at data around a certain epoc event, you will use `epoc_filter` to do this. See the [Raster/PSTH example](#) for a complete demonstration.

To only look at data around the epoc event timestamps, use a `t` parameter to set the filter time range. In this example, data from 20 ms before the *Levl* epoc to 50 ms after the onset is kept.

```
data = tdt.epoc_filter(data, 'Levl', t=[-0.02, 0.07])
```

To only look at data when an epoc was a certain value or values, use a `values` parameter. In this example, only data when the *Freq* epoc was equal to 9000 or 10000 is retained.

```
data = tdt.epoc_filter(data, 'Freq', values=[9000, 10000])
```

If you want to look for a particular behavioral response that occurs sometime during the allowed time range, use the `modifiers` filter. In this example, only data when the *Freq* epoc was 10000 **AND** the *Resp* epoc had a value of 1 sometime during the *Freq* epoc is retained.

```
data = tdt.epoc_filter(data, 'Freq', values=[10000])
data = tdt.epoc_filter(data, 'Resp', modifiers=[1])
```

As you can see, for complex filtering the output from one call to `epoc_filter` can become the input to the next call to `epoc_filter`. If your data sets are large, or if you are iterating through many combinations of epoc variables, it is preferred to extract only the epocs and do all of the epoc filtering first to find the valid time ranges that match the filter, and then use this as the `ranges` input to `read_block` to extract all events (including snips, streams) on only those valid time ranges.

```
# read just the epoc events
data = tdt.read_block(block_path, evtype=['epocs'])

# use the epocs to find time ranges we want
data = tdt.epoc_filter(data, 'Freq', values=[9000, 10000])
data = tdt.epoc_filter(data, 'Levl', values=[70, 80, 90])
data = tdt.epoc_filter(data, 'Resp', modifiers=[1])

# read just the value time ranges from the whole data set
data = tdt.read_block(block_path, ranges=data.time_ranges)
```

## read\_sev

---

`read_sev` reads SEV files into a Python structure. SEV files are created by the RS4 Data Streamer or by enabling the Discrete Files option when streaming continuous signals in Synapse. SEV files consist of a single channel of data per file, with a short header, so it is very fast to read them. `read_block` will automatically call `read_sev` if it finds SEV files in the block directory. Like `read_block`, `read_sev` needs only one input: the block path.

```
data = tdt.read_sev('C:\TDT\Synapse\Tanks\Exp1-160921-120606\Sub1-1')
```

`read_sev` will return a structure containing the streams that it found. Each stream field includes the data array and sampling rate.

# TDT Data Storage

---

Data collected or used by TDT software is stored in tanks - special directories on your hard drive. Each time you press 'Record' in Synapse, a new block is created within the tank. Within a block different stores can record different types of events at different rates. The blocks are special folders within the tank directories.

## TDT Data Types

---

1. **epochs** are values stored with onset and offset timestamps that can be used to create time-based filters on your data. They can be created by the [Epoch Data Storage gizmo](#), [Logic gizmos](#), [Stimulation gizmos](#), and many more.
  - a. If Runtime Notes were enabled in Synapse, they will appear in `data.epochs.Note`. The notes themselves will be in `data.epochs.Note.notes`.  
See the [Synapse Manual](#) for more information on Runtime Notes.
2. **streams** are continuous single channel or multichannel recordings, like those stored by the [Stream Data Storage gizmo](#), the [Fiber Photometry gizmo](#), and many others. The structure includes the data array and sampling rate.
3. **snips** are short snippets of data collected on a trigger. For example, action potentials recorded around threshold crossings in the [Spike Sorting gizmos](#), or fixed duration snippets recorded by the [Strobe Store gizmo](#). This structure includes the waveforms, channel numbers, sort codes, trigger timestamps, and sampling rate.
4. **scalars** are similar to epochs but can be single or multi-channel values and only store an onset timestamp when triggered. These can be created by the [Strobe Store gizmo](#).

The returned structure also contains an **info** field with block start/stop times, duration, and information about the Subject, User, and Experiment that it came from (if the block was created in Synapse).

## Merging Blocks

---

There is a tool called TankManager that installs with Synapse that allows you to concatenate blocks directly on disk. This can create a 'super block' by combining the snippets / epochs events from multiple blocks into a single block before you export the data to another format or import to OpenSorter. It shifts the timestamps of the second block (it adds ~10 second gap in between the recordings so there is no overlap).

TankManager is a command line executable. The format for merging blocks is:

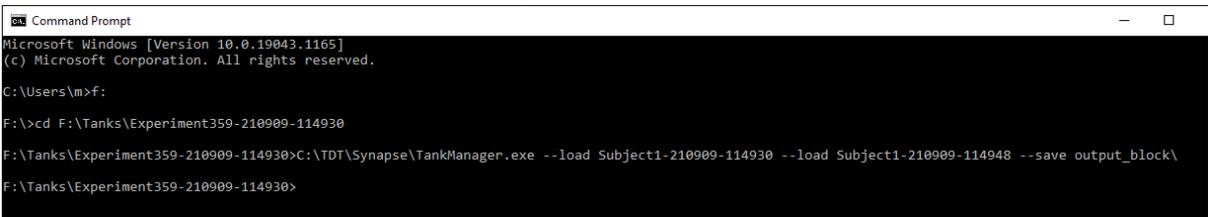
```
C:\TDT\Synapse\TankManager.exe --load block1 --load block2 --save outputblock
```

Replace the text with the two block names that you want to merge.

Here is an example that merges the two blocks called `Subject1-210909-114930` and `Subject1-210909-114948` in the tank `F:\Tanks\Experiment359-210909-114930` into a new block called `output_block` inside that same tank:

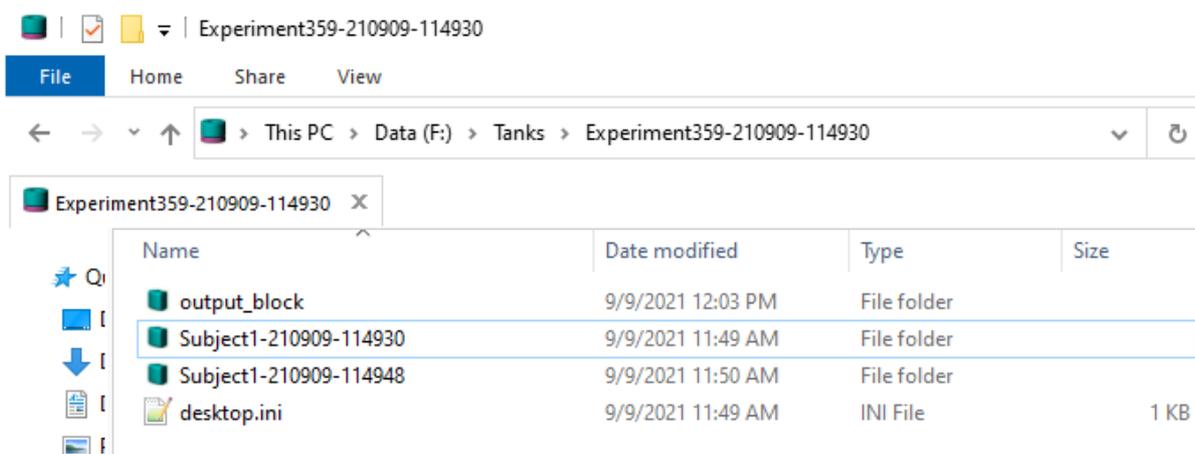
1. Press the Windows key (or Windows + R) and type `cmd` to enter the command line.
2. Change directory into your tank folder.
3. Run the TankManager command. In this example, it is

```
C:\TDT\Synapse\TankManager.exe --load "Subject1-210909-114930" --load "Subject1-210909-114948" --save "output_block"
```



```
Command Prompt
Microsoft Windows [Version 10.0.19043.1165]
(c) Microsoft Corporation. All rights reserved.
C:\Users\vm>f:
F:\>cd F:\Tanks\Experiment359-210909-114930
F:\Tanks\Experiment359-210909-114930>C:\TDT\Synapse\TankManager.exe --load Subject1-210909-114930 --load Subject1-210909-114948 --save output_block\
F:\Tanks\Experiment359-210909-114930>
```

This makes a new block called `output_block` inside the same tank folder, which you can then use like any other block.



## Splitting Blocks

You can extract time ranges of blocks and export them to smaller blocks using the same TankManager utility described above. This works best with snippet, epoc, and scalar data types.

Here's an example that will keep the data from time  $t=5s$  to  $t=10s$  from the block `C:\BLOCKPATH` and write it to a new block called `C:\BLOCKPATH_5_10`:

```
C:\TDT\Synapse\TankManager -l "C:\BLOCKPATH" -k "5-10" -s "C:\BLOCKPATH_5_10"
```

For example, you could use this to split up your blocks into hour long chunks like this:

```
C:\TDT\Synapse\TankManager -l "C:\BLOCKPATH" -k "0-3600" -s "C:\BLOCKPATH_0hr"
C:\TDT\Synapse\TankManager -l "C:\BLOCKPATH" -k "3600-7200" -s "C:\BLOCKPATH_1hr"
```

And so on. The data in these new blocks won't start until the first timestamp, so for later blocks there will be a large gap in the beginning. If you have an issue reading it with OpenSorter or another of our data utilities you might try following the steps in [Tech Note TN0909](#).

### Note

You might get a message during the block creation that says `Failed to set the system attribute on the block`. It is okay to ignore this.

# Importing TDT Data into Python for Offline Analysis

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TDT provides a set of cross-platform tools for reading TDT data files directly into Python offline. It is compatible with Python 3.

## Requirements

---

1. Python 3. We recommend installing [Anaconda](#).

## Installation

---

Just follow these three steps to get started:

1. Install the TDT Python package from [pypi](#). From the Anaconda Prompt, type:

```
pip install tdt
```

2. In Python, you can download the example data to the current directory with this:

```
import tdt
tdt.download_demo_data()
```

3. Importing data is easy

```
data = tdt.read_block('path/to/block')
```

# Example Notebooks

---

## Introduction to Python

---

### Using the tdt Package

This primer walks through installing the tdt Python package, shows how to convert existing MATLAB code to Python, and highlights some of the differences when working in Python.

### Installation

1. Make sure that you have **Anaconda** installed.
2. **Open an Anaconda Prompt** and type:

```
pip install tdt
```

3. **Done!**

### Converting Existing MATLAB Code to Python

The tdt Python library for reading TDT data is one-to-one compatible with the MATLAB library, however the function names and parameter names are different.

### Extracting Block Data

**TDTbin2mat** extracts block data into a MATLAB structure.

```
data = TDTbin2mat(BLOCK_PATH);
```

The Python equivalent is **read\_block**.

```
from tdt import read_block  
data = read_block(BLOCK_PATH)
```

**TDTbin2mat** and **read\_block** share parameters, but the parameters have different names.

### Parameter Name Translation Table

MATLAB	Python	Description
T1	t1	scalar, retrieve data starting at t1 (default = 0 for beginning of recording)
T2	t2	scalar, retrieve data ending at t2 (default = 0 for end of recording)
TYPE	evtype	array of strings, specifies what type of data stores to retrieve from the tank
SORTNAME	sortname	string, specify sort ID to use when extracting snippets (default = 'TankSort')
RANGES	ranges	array of valid time range column vectors
NODATA	nodata	bool, only return timestamps, channels, and sort codes for snippets (default = false)
STORE	store	string or list of strings, specify specific store(s) to extract
CHANNEL	channel	integer, choose a single channel to extract from stream or snippet events
BITWISE	bitwise	string, specify an epoc store or scalar store that contains a 32-bit integer. Onsets/offsets from individual bits will be extracted
HEADERS	headers	var, set to 1 to return only the headers for this block, if you need to make fast consecutive calls to read_block
COMBINE	combine	list, specify store(s) that were saved by the Strobed Data Storage gizmo in Synapse. It will intelligently combine data into snippets.

## Epoc Filtering

**TDTfilter** filters events around epoc events in MATLAB:

```
data = TDTbin2mat(BLOCK_PATH);
data = TDTfilter(data, 'Tick', 'TIME', [-0.3, 0.8], 'VALUES', [5, 10, 15]);
```

The Python equivalent is **epoc\_filter**:

```
from tdt import read_block, epoc_filter
data = read_block(BLOCK_PATH)
data = epoc_filter(data, 'Tick', t=[-0.3, 0.8], values=[5, 10, 15])
```

**TDTbin2mat** and **read\_block** share parameters, but the parameters have different names.

## Parameter Name Translation Table

MATLAB	Python	Description
VALUES	values	array of allowed epoc values
MODIFIERS	modifiers	array of allowed modifier values. For example, only allow time ranges when allowed modifier occurred sometime during that event, e.g. a correct animal response.
TIME	t	onset/offset pair, extracts events around epoc onsets only
TIMEREF	tref	boolean, set to True to use the epoc event onset as a time reference
KEEPDATA	keepdata	boolean, keep the original stream data array and add a field called 'filtered' that holds the data from each valid time range

## Extracting SEV Data

**SEV2mat** extracts SEV files from a given directory into a MATLAB structure. These files are created on the RS4 Data Streamer or by enabling the Discrete Files option in the Synapse Stream Data Storage gizmo. Each SEV file contains a header and the raw binary data from a single channel.

```
data = SEV2mat(BLOCK_PATH);
```

The Python equivalent is **read\_sev**.

```
from tdt import read_sev
data = read_sev(BLOCK_PATH)
```

**SEV2mat** and **read\_sev** share parameters, but the parameters have different names.

## Parameter Name Translation Table

MATLAB	Python	Description
T1	t1	scalar, retrieve data starting at t1 (default = 0 for beginning of recording)
T2	t2	scalar, retrieve data ending at t2 (default = 0 for end of recording)
CHANNEL	channel	integer, returns the SEV data from specified channel only (default = 0 for all channels)
RANGES	ranges	array of valid time range column vectors
JUSTNAMES	just_names	boolean, retrieve only the valid event names
EVENTNAME	event_name	string, specific event name to retrieve data from
VERBOSE	verbose	boolean, set to false to disable console output
FS	fs	float, sampling rate override. Useful for lower sampling rate recordings that aren't correctly written into the SEV header.

## Walkthrough

Let's look at some basic concepts for working with Python and the tdt library.

\*\*\*Python Tips!\*\*

Use `print` in Python in place of `disp` in MATLAB.

`%` is a special command used mainly in Python notebooks

`#` creates a single-line comment in Python

Use `'''` to make a multi-line comment

First we'll import the critical libraries.

```
# this is a single line comment

''' this is a comment
spanning multiple lines'''

# special call that tells notebook to show matplotlib figures inline
%matplotlib inline

import matplotlib.pyplot as plt # standard Python plotting library
import numpy as np # fundamental package for scientific computing, handles
arrays and maths

# import the tdt library
import tdt
```

\*\*\*Python Tip!\*\*

Use the `__doc__` function to get help on a function.

```
print(tdt.read_block.__doc__)
```

TDT tank data extraction.

`data = read_block(block_path)`, where `block_path` is a string, retrieves all data from specified block directory in struct format. This reads the binary tank data and requires no Windows-based software.

<code>data.epocs</code>	contains all epoc store data (onsets, offsets, values)
<code>data.snips</code>	contains all snippet store data (timestamps, channels, and raw data)
<code>data.streams</code>	contains all continuous data (sampling rate and raw data)
<code>data.scalars</code>	contains all scalar data (samples and timestamps)
<code>data.info</code>	contains additional information about the block

optional keyword arguments:

<code>t1</code>	scalar, retrieve data starting at <code>t1</code> (default = 0 for beginning of recording)
<code>t2</code>	scalar, retrieve data ending at <code>t2</code> (default = 0 for end of recording)
<code>sortname</code>	string, specify sort ID to use when extracting snippets (default = 'TankSort')
<code>evtype</code>	array of strings, specifies what type of data stores to retrieve from the tank. Can contain 'all' (default), 'epocs', 'snips', 'streams', or 'scalars'.

example:

```
data = read_block(block_path,
evtype=['epocs', 'snips'])
```

> returns only epocs and snips

<code>ranges</code>	array of valid time range column vectors.
---------------------	---

example:

```
tr = np.array([[1,3],[2,4]])
```

```
data = read_block(block_path, ranges=tr)
```

> returns only data on `t=[1,2)` and `[3,4)`

<code>nodata</code>	boolean, only return timestamps, channels, and sort codes for snippets, no waveform data (default =
---------------------	---

false).

Useful speed-up if not looking for waveforms

<code>store</code>	string, specify a single store to extract
--------------------	---

list of strings, specify multiple stores to extract

<code>channel</code>	integer, choose a single channel to extract from
----------------------	--

stream or snippet events. Default is 0, to extract all channels.

<code>bitwise</code>	string, specify an epoc store or scalar store that
----------------------	--

contains individual bits packed into a 32-bit integer. Onsets/offsets from individual bits will be extracted.

<code>headers</code>	var, set to 1 to return only the headers for this
----------------------	---

block, so that you can make repeated calls to read data without having to parse the TSQ file every time, for faster consecutive reads. Once created,

```

        pass in the headers using this parameter.
    example:
        heads = read_block(block_path, headers=1)
        data = read_block(block_path, headers=heads,
evtype=['snips'])
        data = read_block(block_path, headers=heads,
evtype=['streams'])
    combine    list, specify one or more data stores that were saved
                by the Strobed Data Storage gizmo in Synapse (or an
                Async_Stream_store macro in OpenEx). By default,
                the data is stored in small chunks while the strobe
                is high. This setting allows you to combine these
                small chunks back into the full waveforms that were
                recorded while the strobe was enabled.
    example:
        data = read_block(block_path, combine=['StS1'])
    export    string, choose a data exporting format.
        csv:        data export to comma-separated value files
                    streams: one file per store, one channel
per column
                    epochs: one column onsets, one column
offsets
        binary:    streaming data is exported as raw binary
files
                    one file per channel per store
                    interlaced: streaming data exported as raw binary
files
                    one file per store, data is interlaced
    scale    float, scale factor for exported streaming data. Default
= 1.
    dtype    string, data type for exported binary data files
        None: Uses the format the data was stored in (default)
        'i16': Converts all data to 16-bit integer format
        'f32': Converts all data to 32-bit integer format
    outdir   string, output directory for exported files. Defaults to
current
        block folder if not specified
    prefix   string, prefix for output file name. Defaults to None

```

```
print(tdt.epoc_filter.__doc__)
```

TDT tank data filter. Extract data around epoc events.

`data = epoc_filter(data, epoc)` where `data` is the output of `read_block`, `epoc` is the name of the epoc to filter on, and parameter value pairs define the filtering conditions.

If no parameters are specified, then the time range of the epoc event is used as a time filter.

Also creates `data.filter`, a string that describes the filter applied.

Optional keyword arguments:

<code>values</code>	specify array of allowed values ex: <code>tempdata = epoc_filter(data, 'Freq', values=[9000, 10000])</code>  > retrieves data when <code>Freq = 9000</code> or <code>Freq = 10000</code>
<code>modifiers</code>	specify array of allowed modifier values. For example, only allow time ranges when allowed modifier occurred sometime during that event, e.g. a correct animal response.  ex: <code>tempdata = epoc_filter(data, 'Resp', modifiers=[1])</code> > retrieves data when <code>Resp = 1</code> sometime during the allowed
<code>t</code>	time range. specify onset/offset pairs relative to epoc onsets. If the offset is not provided, the epoc offset is used. ex: <code>tempdata = epoc_filter(data, 'Freq', t=[-0.1, 0.5])</code> > retrieves data from 0.1 seconds before <code>Freq</code> onset to 0.4 seconds after <code>Freq</code> onset. Negative time ranges are discarded.
<code>tref</code>	use the epoc event onset as a time reference. All timestamps for epoc, snippet, and scalar events are then relative to epoc onsets.  ex: <code>tempdata = epoc_filter(data, 'Freq', tref=True)</code> > sets snippet timestamps relative to <code>Freq</code> onset
<code>keepdata</code>	keep the original stream data array and add a field called 'filtered' that holds the data from each valid time range.  Defaults to <code>True</code> .

**IMPORTANT!** Use a time filter (`t` argument) only after all value filters have been set.

```
print(tdt.read_sev.__doc__)
```

TDT sev file data extraction.

`data = read_sev(sev_dir)`, where `sev_dir` is a string, retrieves all sev data from specified directory in struct format. `sev_dir` can also be a single file. SEV files are generated by an RS4 Data Streamer, or by enabling the Discrete Files option in the Synapse Stream Data Storage gizmo, or by setting the Unique Channel Files option in `Stream_Store_MC` or `Stream_Store_MC2` macro to Yes in OpenEx.

If exporting is enabled, this function returns None.

`data` contains all continuous data (sampling rate and raw data)

optional keyword arguments:

<code>t1</code>	scalar, retrieve data starting at <code>t1</code> (default = 0 for beginning of recording)
<code>t2</code>	scalar, retrieve data ending at <code>t2</code> (default = 0 for end of recording)
<code>channel</code>	integer, returns the sev data from specified channel only (default = 0 for all channels)
<code>ranges</code>	array of valid time range column vectors
<code>just_names</code>	boolean, retrieve only the valid event names
<code>event_name</code>	string, specific event name to retrieve data from
<code>verbose</code>	boolean, set to false to disable console output
<code>fs</code>	float, sampling rate override. Useful for lower sampling rates that aren't correctly written into the SEV header.
<code>export</code>	string, choose a data exporting format. <code>csv</code> : data export to comma-separated value files streams: one file per store, one channel epocs: one column onsets, one column binary: streaming data is exported as raw binary files: one file per channel per store interlaced: streaming data exported as raw binary files: one file per store, data is interlaced
<code>per column</code>	
<code>offsets</code>	
<code>files</code>	
<code>files</code>	
<code>scale</code>	float, scale factor for exported streaming data. Default = 1.
<code>dtype</code>	string, data type for exported binary data files None: Uses the format the data was stored in (default) 'i16': Converts all data to 16-bit integer format 'f32': Converts all data to 32-bit integer format
<code>outdir</code>	string, output directory for exported files. Defaults to current block folder if not specified
<code>prefix</code>	string, prefix for output file name. Defaults to None

## Download demo data from the TDT website

```
tdt.download_demo_data()
```

```
demo data ready
```

This example uses our [example data sets](#). To import your own data, replace BLOCK\_PATH with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu > History. Find your block, then Right-Click > Copy path to clipboard.

```
BLOCK_PATH = 'data/Algernon-180308-130351'  
data = tdt.read_block(BLOCK_PATH)
```

```
read from t=0s to t=61.23s
```

```
print(data)
```

```
epocs    [struct]  
snips    [struct]  
streams  [struct]  
scalars  [struct]  
info     [struct]  
time_ranges:  array([[ 0.],  
                    [inf]])
```

**read\_block** returns a structured object. It is a Python dictionary but also allows you to use the dot syntax like in MATLAB, so you can access fields within the structure with either method. These two ways of looking at the block info field are equivalent:

```
data.info
```

```
tankpath: 'data'
blockname: 'Algernon-180308-130351'
start_date: datetime.datetime(2018, 3, 8, 13, 3, 53, 999999)
utc_start_time: '13:03:53'
stop_date: datetime.datetime(2018, 3, 8, 13, 4, 55, 233578)
utc_stop_time: '13:04:55'
duration: datetime.timedelta(seconds=61, microseconds=233579)
stream_channel: 0
snip_channel: 0
```

```
data['info']
```

```
tankpath: 'data'
blockname: 'Algernon-180308-130351'
start_date: datetime.datetime(2018, 3, 8, 13, 3, 53, 999999)
utc_start_time: '13:03:53'
stop_date: datetime.datetime(2018, 3, 8, 13, 4, 55, 233578)
utc_stop_time: '13:04:55'
duration: datetime.timedelta(seconds=61, microseconds=233579)
stream_channel: 0
snip_channel: 0
```

These three methods to access the 'Wav1' store sampling rate are equivalent:

```
data.streams.Wav1.fs # dot syntax
```

```
24414.0625
```

```
data['streams']['Wav1']['fs'] # dict keys only
```

```
24414.0625
```

```
data['streams'].Wav1['fs'] # mix of dot syntax and dict keys
```

```
24414.0625
```

\*\*\*Python Tip!\*\*

Spaces are important in python. Commands like `for`, `if`, `elif`, `while`, and others require indents to track their nests

```

for foo in foo_list:
    something
    if foo == check:
        conditional_something
    elif:
        still_in_for_loop
    else:
        still_in_for_loop

out_of_loop

```

Accessing a field with the string dictionary key method is useful when using a variable name, such as this example which loops through all the stream store names and prints their sampling rates.

```

print('Sampling rates in', data.info.blockname)
for store in data.streams.keys():
    print(store, '{:.4f} Hz'.format(data.streams[store].fs))

```

```

Sampling rates in Algernon-180308-130351
LFP1 3051.7578 Hz
pNe1 498.2462 Hz
Wav1 24414.0625 Hz

```

## Explore Stream events

Let's look at the contents of the stream event structures.

```

print('all stream stores')
print(data.streams)

```

```

all stream stores
LFP1    [struct]
pNe1    [struct]
Wav1    [struct]

```

```

print(data.streams.Wav1)

```

```

name: 'Wav1'
code: 829841751
size: 2058
type: 33025
type_str: 'streams'
ucf: False
fs: 24414.0625
dform: 0
start_time: 0.0
data: array([[ 1.0028159e-03,  1.0012799e-03,  9.9590386e-04, ...,
              -1.5983999e-03, -1.5984639e-03, -1.5852799e-03],
             [ 5.5667193e-04,  5.6723197e-04,  5.6083198e-04, ...,
              -1.4531199e-03, -1.4584319e-03, -1.4480639e-03],
             [-4.6534397e-04, -4.5804796e-04, -4.6521597e-04, ...,
              -1.2184319e-03, -1.2098559e-03, -1.2177919e-03],
             ...,
             [ 2.9247998e-05,  2.3295999e-05,  3.2191998e-05, ...,
              -2.2208637e-03, -2.2241918e-03, -2.2300798e-03],
             [ 7.2191993e-04,  7.1571197e-04,  7.2358397e-04, ...,
              -2.1401597e-03, -2.1399679e-03, -2.1494399e-03],
             [ 2.3078399e-04,  2.3590398e-04,  2.4435198e-04, ...,
              -1.3180159e-03, -1.3103359e-03, -1.3012479e-03]], dtype=float32)

```

The actual data is store in numpy arrays. For a multi-channel stream store, each row is a channel. Python uses 0-based indexing, so we have to subtract 1 from our channel number when accessing the array

```
print('channel 1:', data.streams.Wav1.data[0,:])
```

```
channel 1: [ 0.00100282  0.00100128  0.0009959 ... -0.0015984 -0.00159846
           -0.00158528]
```

```
num_samples = len(data.streams.Wav1.data[0])
print('number of samples:', num_samples)
```

```
number of samples: 1490944
```

Create time vector for plotting by dividing the number of samples in the array by the sampling rate

```
Wav1_time = np.linspace(1, num_samples, num_samples) / data.streams.Wav1.fs
```

Plot the first 2 seconds from a single channel

```
t = int(2 * data.streams.Wav1.fs) # int rounds it to the nearest integer

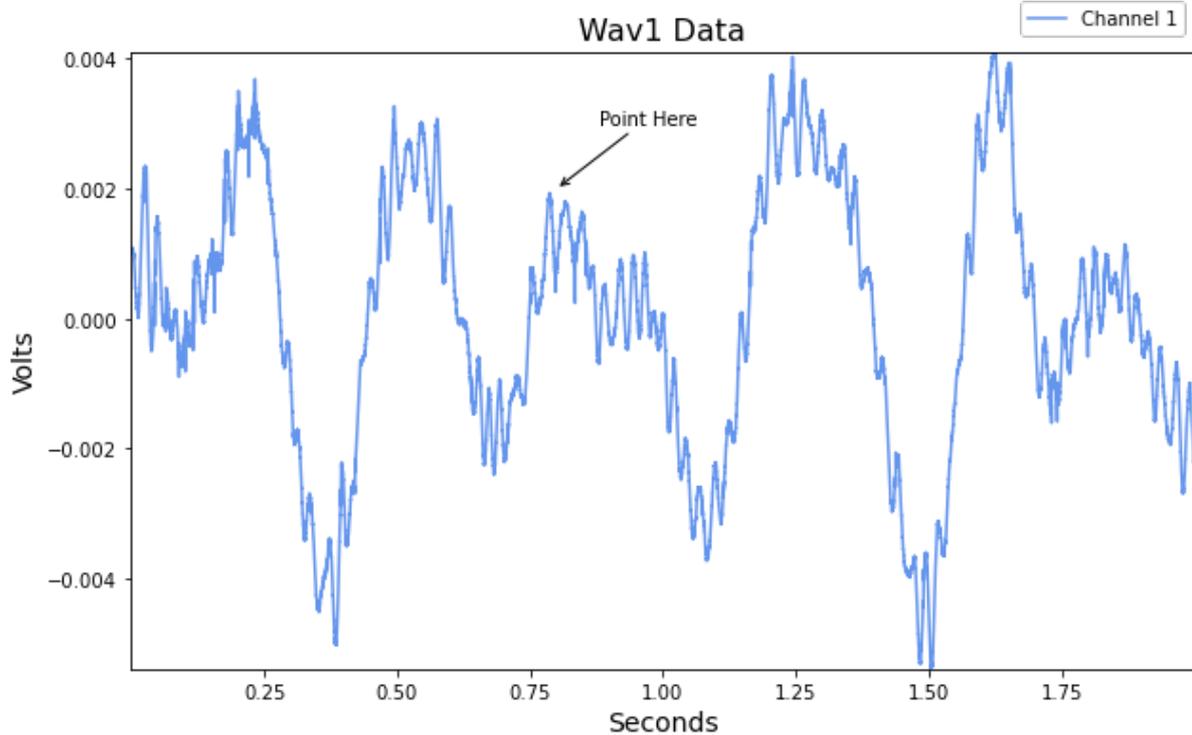
# declare the figure size
fig1 = plt.subplots(figsize=(10, 6))

channel = 1

# plot the line using slices
plt.plot(Wav1_time[0:t], data.streams.Wav1.data[channel-1,0:t],
color='cornflowerblue')

# Some matplotlib stuff
# add an annotation mark to the figure
plt.annotate('Point Here',
             xy=(0.8,0.002),
             xytext=(.88,.003),
             arrowprops=dict(arrowstyle='->', color='k')
            )

# create title, axis labels, and legend
plt.title('Wav1 Data', fontsize=16)
plt.xlabel('Seconds', fontsize=14)
plt.ylabel('Volts', fontsize=14)
plt.legend(('Channel {}'.format(channel)),
          loc='lower right',
          bbox_to_anchor=(1.0,1.01)
         )
plt.autoscale(tight=True)
plt.show()
```



### \*\*\*Python Tip!\*\*

Array slices in Python have some notable differences. 1. Zero-based indexing 2. Omit 'end' and just use a negative index to index starting from the end of the array 3. Omit the starting index if you want to include the first element

### MATLAB array slices

```
arr = 1:10;           % arr = [1 2 3 4 5 6 7 8 9 10]
b = arr(3:5);        % b = [3 4 5]
c = arr(1:end-2);    % c = [1 2 3 4 5 6 7 8]
d = arr(end-1:end); % d = [9 10]
```

### Python equivalent

```
arr = np.arange(1,11) # arr = [1 2 3 4 5 6 7 8 9 10]
b = arr[2:5]          # b = [3 4 5]
c = arr[:-2]          # c = [1 2 3 4 5 6 7 8]
d = arr[-2:]          # d = [9 10]
```

For reference, here are some matplotlib colors originally from [this stackoverflow answer](#)



## Explore Epcoc Events

Let's look at the contents of the epcoc event structures.

```
print('all epcoc events')
print(data.epocs)
```

```
all epcoc events
PC0_    [struct]
Pu1e    [struct]
```

```
print(data.epocs.Pu1e)
```

```

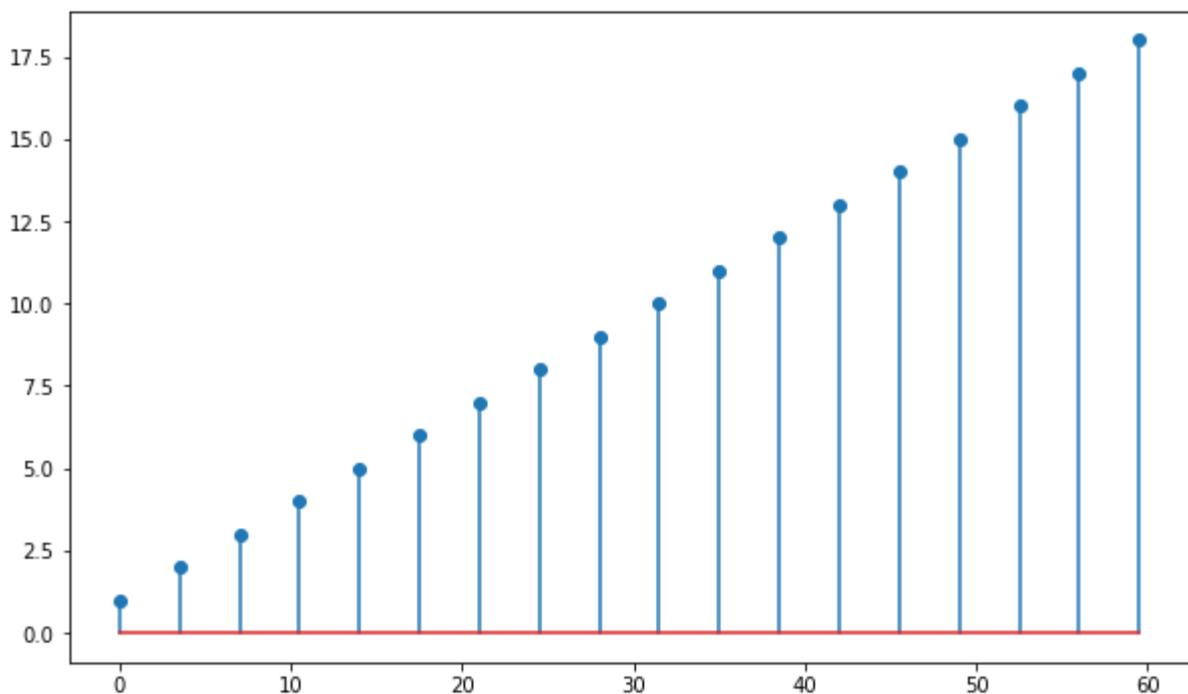
name: 'Pu1e'
onset: array([1.22880000e-04, 3.50011392e+00, 7.00010496e+00, 1.05000960e+01,
             1.40000870e+01, 1.75000781e+01, 2.10000691e+01, 2.45000602e+01,
             2.80000512e+01, 3.15000422e+01, 3.50000333e+01, 3.85000243e+01,
             4.20000154e+01, 4.55000064e+01, 4.89999974e+01, 5.24999885e+01,
             5.59999795e+01, 5.94999706e+01])
offset: array([ 0.30011392,  3.80010496,  7.300096  , 10.80008704,
              14.30007808,
              17.80006912, 21.30006016, 24.8000512  , 28.30004224, 31.80003328,
              35.30002432, 38.80001536, 42.3000064  , 45.79999744, 49.29998848,
              52.79997952, 56.29997056, 59.7999616  ])
type: 'onset'
type_str: 'epocs'
data: array([ 1.,  2.,  3.,  4.,  5.,  6.,  7.,  8.,  9., 10., 11., 12.,
            13.,
            14., 15., 16., 17., 18.])
dform: 4
size: 10

```

```

# a simple plot
fig1 = plt.subplots(figsize=(10, 6))
plt.stem(data.epocs.Pu1e.onset, data.epocs.Pu1e.data)
plt.show()

```



## Stream Plot Example

---

Import Continuous Data into Python

Plot a single channel of data with various filtering schemes

Good for first-pass visualization of streamed data

Combine streaming data and epochs in one plot

## Housekeeping

Import the tdt package and other python packages we care about.

```
# special call that tells notebook to show matplotlib figures inline
%matplotlib inline

import matplotlib.pyplot as plt # standard Python plotting library
import numpy as np # fundamental package for scientific computing, handles
arrays and math

# import the tdt library
import tdt
```

## Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCK_PATH = 'data/Algernon-180308-130351'
```

```
demo data ready
```

Now read channel 1 from all stream data into a Python structure called 'data'

```
data = tdt.read_block(BLOCK_PATH, evtype=['streams', 'epochs'], channel=1)
```

```
read from t=0s to t=61.23s
```

And that's it! Your data is now in Python. The rest of the code is a simple plotting example.

## Stream Store Plotting

Let's create time vectors for each stream store for plotting in time.

```
time_Wav1 = np.linspace(1, len(data.streams.Wav1.data),
len(data.streams.Wav1.data)) / data.streams.Wav1.fs
time_LFP1 = np.linspace(1, len(data.streams.LFP1.data),
len(data.streams.LFP1.data)) / data.streams.LFP1.fs
time_pNe1 = np.linspace(1, len(data.streams.pNe1.data),
len(data.streams.pNe1.data)) / data.streams.pNe1.fs
```

Plot five seconds of data from each store

```
fig, (ax1, ax2, ax3) = plt.subplots(nrows=3, ncols=1, figsize=(8, 8),
sharex=True)

ax1.plot(time_Wav1, data.streams.Wav1.data*1e6, color='cornflowerblue')
ax1.set_title('Basic Data Plotting: Ch 1\nRaw Waveform', fontsize=14)

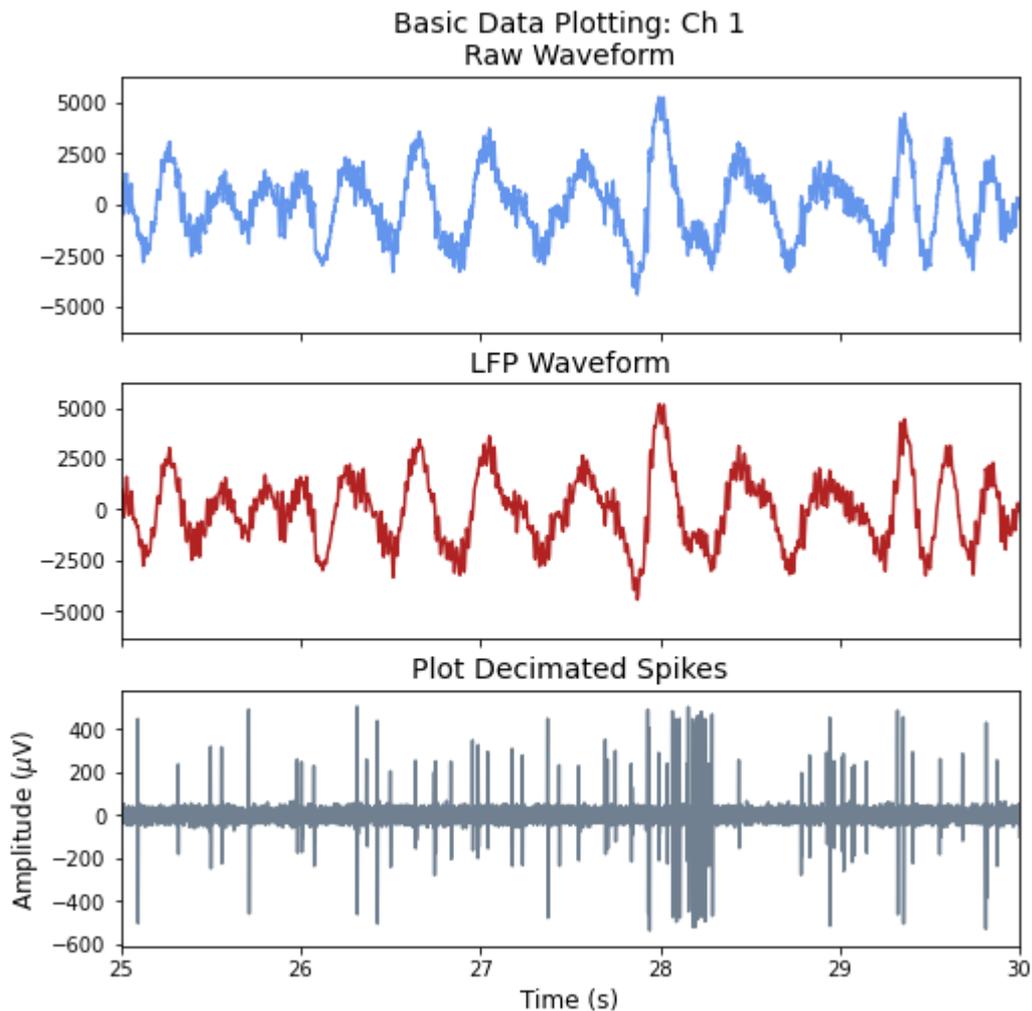
ax2.plot(time_LFP1, data.streams.LFP1.data*1e6, color='firebrick')
ax2.set_title('LFP Waveform', fontsize=14)

ax3.plot(time_pNe1, data.streams.pNe1.data, color='slategray')
ax3.set_title('Plot Decimated Spikes', fontsize=14)

ax3.set_xlabel('Time (s)', fontsize=12)
ax3.set_ylabel('Amplitude ( $\mu$ V)', fontsize=12)

ax1.set_xlim(25, 30)

plt.show()
```



## Epoc Events

Generate continuous time series for epoc data using epoc timestamps

```
# StimSync epoc event
STIM_SYNC = 'PC0_'
pc0_on = data.epocs[STIM_SYNC].onset
pc0_off = data.epocs[STIM_SYNC].offset
pc0_x = np.reshape(np.kron([pc0_on, pc0_off], np.array([[1], [1]])).T, [1,
-1])[0]
```

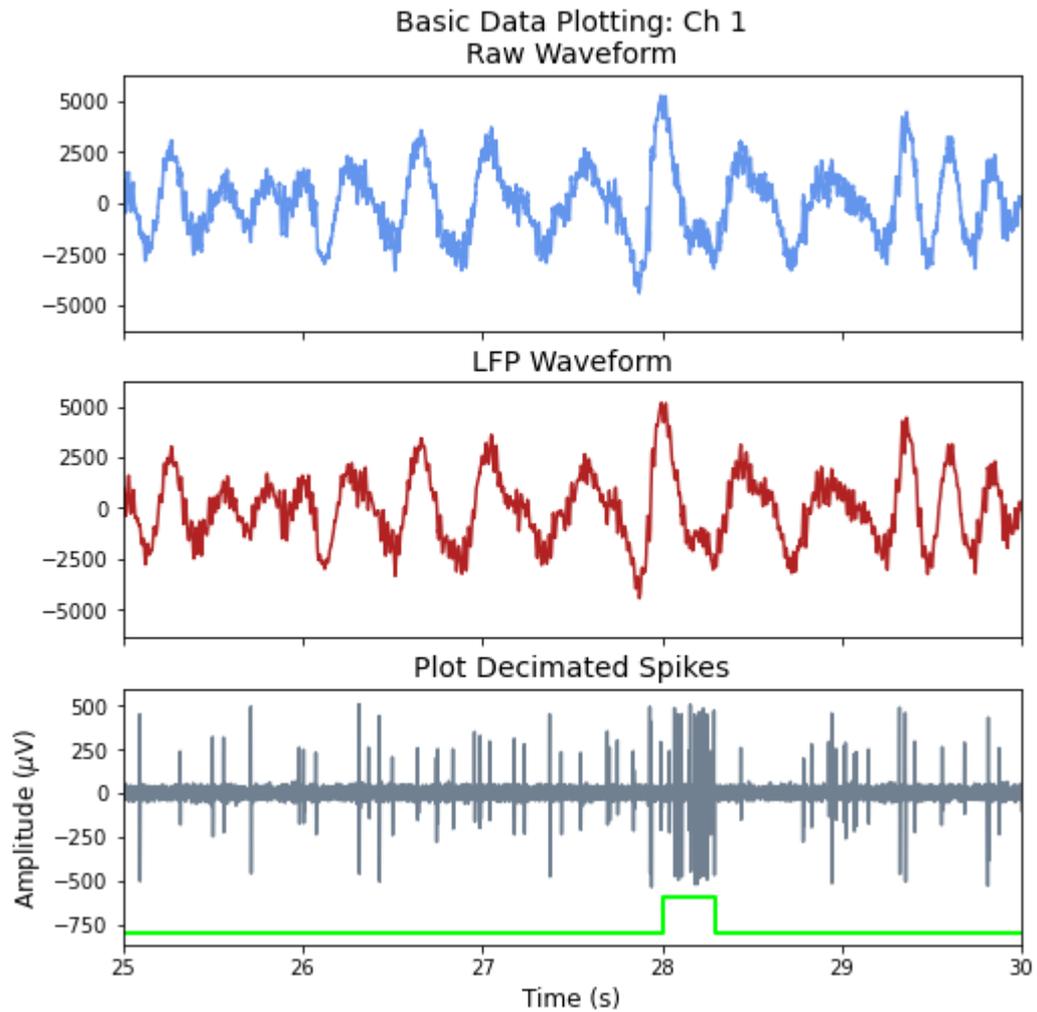
Make a time series waveform of epoc values and plot them.

```

sz = len(pc0_on)
d = data.epocs[STIM_SYNC].data
pc0_y = np.reshape(np.vstack([np.zeros(sz), d, d, np.zeros(sz)]).T, [1,-1])[0]

ax3.plot(pc0_x, 200*(pc0_y) - 800, color='lime', linewidth=2)
fig

```



## Averaging Example

---

Import stream and epoc data into Python using **read\_block**

Plot the average waveform around the epoc event using **epoc\_filter**

Good for Evoked Potential detection

## Housekeeping

Import the tdt package and other python packages we care about.

```
# special call that tells notebook to keep matplotlib figures open
%config InlineBackend.close_figures = False

# special call that tells notebook to show matplotlib figures inline
%matplotlib inline

import matplotlib.pyplot as plt # standard Python plotting library
import numpy as np # fundamental package for scientific computing, handles
arrays and math

# import the tdt library
import tdt
```

## Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCK_PATH = 'data/Algernon-180308-130351'
```

```
demo data ready
```

Set up the variables for the data you want to extract. We will extract channel 3 from the LFP1 stream data store, created by the Neural Stream Processor gizmo, and use our PulseGen epoc event ('PC0/') as our stimulus onset.

```
REF_EPOC      = 'PC0/'
STREAM_STORE  = 'LFP1'
ARTIFACT     = np.inf          # optionally set an artifact rejection level
CHANNEL      = 3
TRANGE       = [-0.3, 0.8]    # window size [start time relative to epoc onset,
                               window duration]
```

Now read the specified data from our block into a Python structure.

```
data = tdt.read_block(BLOCK_PATH, evtype=['epocs', 'scalars', 'streams'],
                     channel=CHANNEL)
```

```
read from t=0s to t=61.23s
```

## Use epoc\_filter to extract data around our epoc event

Using the 't' parameter extracts data only from the time range around our epoc event. For stream events, the chunks of data are stored in a list.

```
data = tdt.epoc_filter(data, 'PC0/', t=TRANGE)
```

Optionally remove artifacts.

```
art1 = np.array([np.any(x>ARTIFACT) for x in
                 data.streams[STREAM_STORE].filtered], dtype=bool)
art2 = np.array([np.any(x<-ARTIFACT) for x in
                 data.streams[STREAM_STORE].filtered], dtype=bool)
good = np.logical_not(art1) & np.logical_not(art2)
data.streams[STREAM_STORE].filtered = [data.streams[STREAM_STORE].filtered[i]
for i in range(len(good)) if good[i]]
num_artifacts = np.sum(np.logical_not(good))
if num_artifacts == len(art1):
    raise Exception('all waveforms rejected as artifacts')
```

Applying a time filter to a uniformly sampled signal means that the length of each segment could vary by one sample. Let's find the minimum length so we can trim the excess off before calculating the mean.

```
min_length = np.min([len(x) for x in data.streams[STREAM_STORE].filtered])
data.streams[STREAM_STORE].filtered = [x[:min_length] for x in
data.streams[STREAM_STORE].filtered]
```

Find the average signal.

```
all_signals = np.vstack(data.streams[STREAM_STORE].filtered)
mean_signal = np.mean(all_signals, axis=0)
std_signal = np.std(all_signals, axis=0)
```

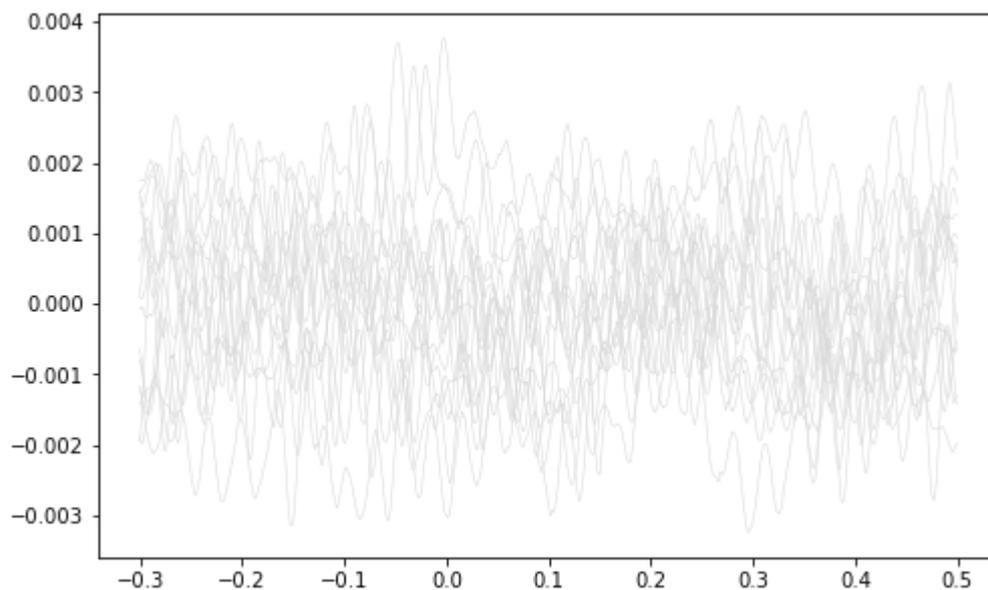
## Ready to plot

Create the time vector.

```
ts = TRANGE[0] + np.arange(0, min_length) / data.streams[STREAM_STORE].fs
```

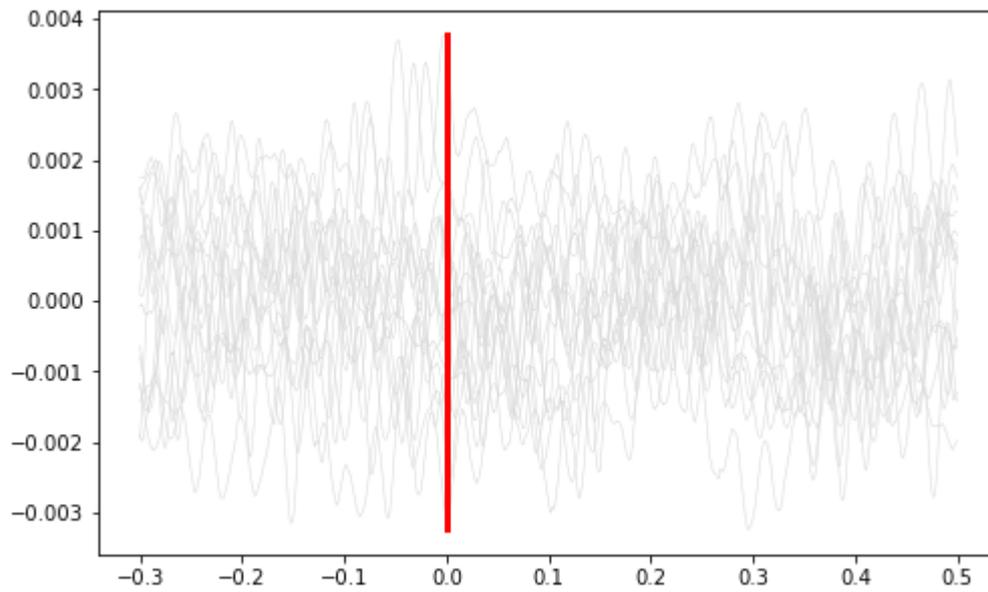
Plot all the signals as gray.

```
fig, ax1 = plt.subplots(1, 1, figsize=(8,5))
ax1.plot(ts, all_signals.T, color=(.85,.85,.85), linewidth=0.5)
plt.show()
```



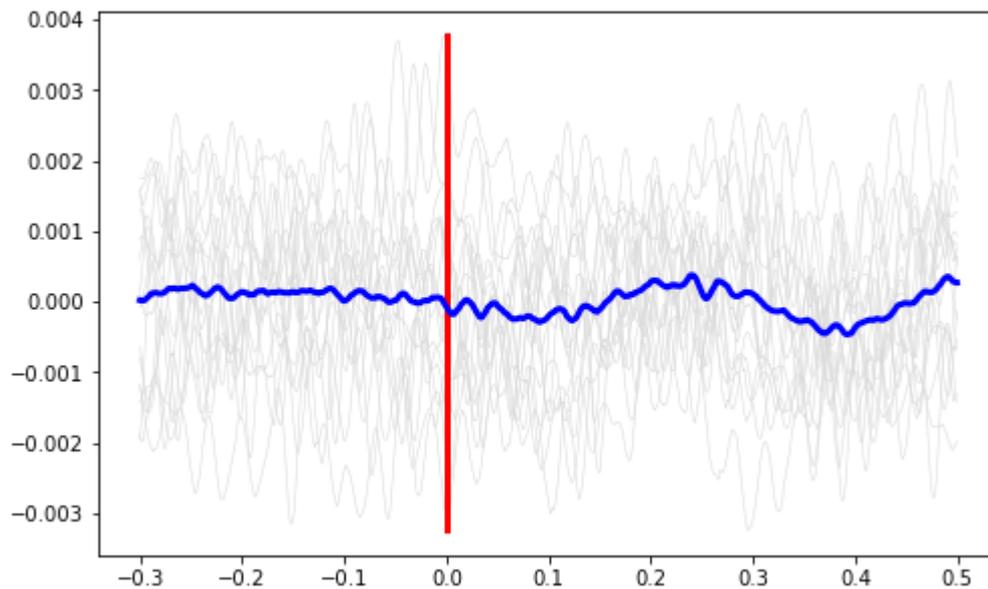
Plot vertical line at time=0.

```
ax1.plot([0, 0], [np.min(all_signals), np.max(all_signals)], color='r',  
linewidth=3)  
plt.show()
```



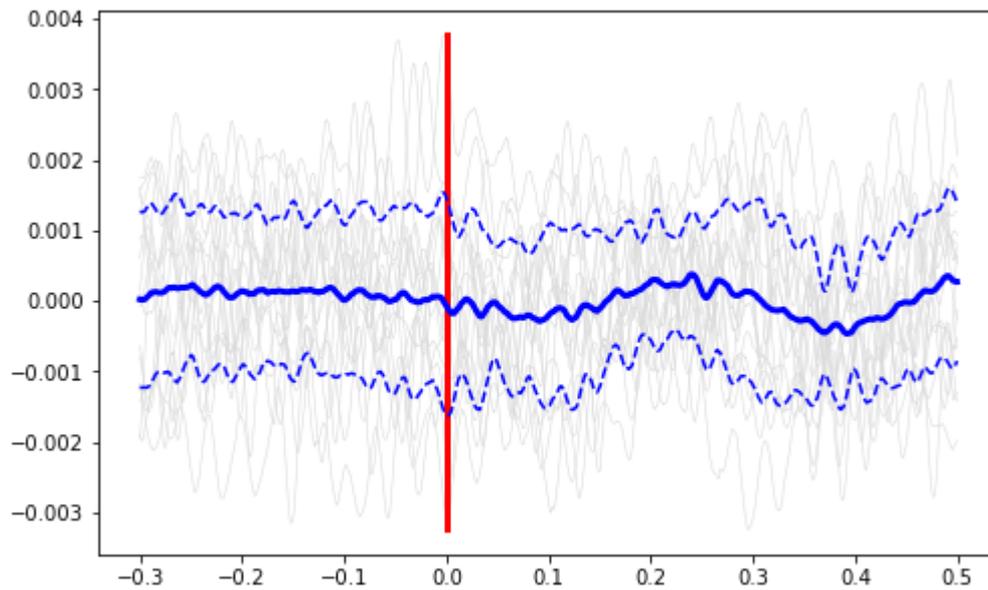
Plot the average signal.

```
ax1.plot(ts, mean_signal, color='b', linewidth=3)  
plt.show()
```



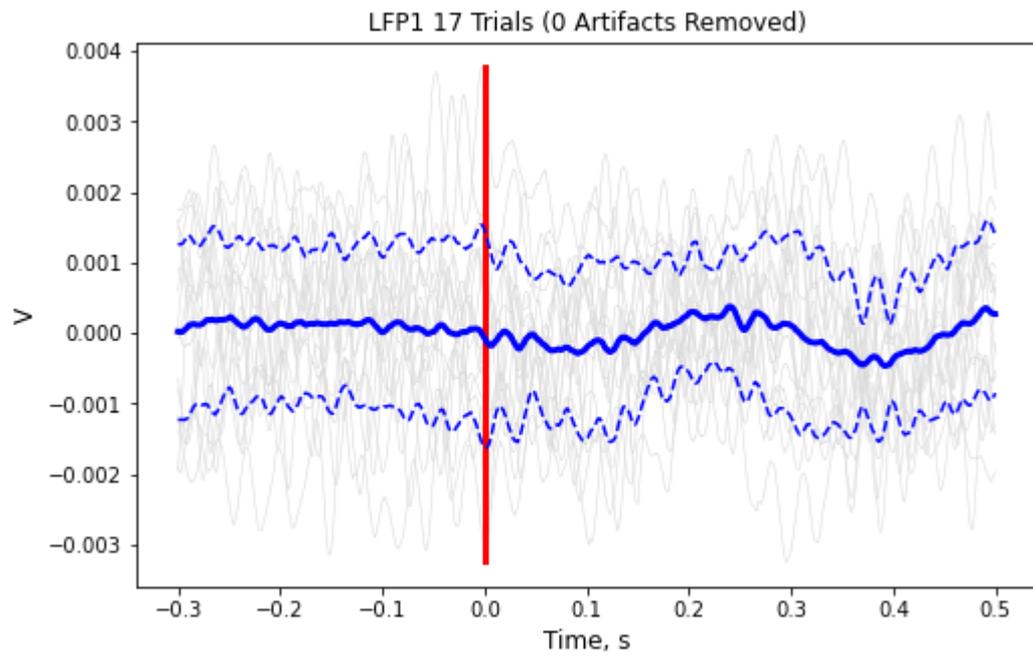
Plot the standard deviation bands.

```
ax1.plot(ts, mean_signal + std_signal, 'b--', ts, mean_signal - std_signal,
        'b--')
plt.show()
```



Finish up the plot

```
ax1.axis('tight')
ax1.set_xlabel('Time, s', fontsize=12)
ax1.set_ylabel('V', fontsize=12)
ax1.set_title('{0} {1} Trials ({2} Artifacts Removed)'.format(
    STREAM_STORE,
    len(data.streams[STREAM_STORE].filtered),
    num_artifacts))
plt.show()
```



## Raster Peristimulus Time Histogram (PSTH) Example

---

Import snippet and epoc data into Python using **read\_block**

Generate peristimulus raster and histogram plots over all trials using **epoc\_filter**

Good for stim-response experiments, such as optogenetic or electrical stimulation

### Housekeeping

Import the tdt package and other python packages we care about.

```
%config InlineBackend.close_figures = False
%matplotlib inline

import numpy as np # fundamental package for scientific computing, handles
arrays and math
import matplotlib.pyplot as plt # standard Python plotting library
from matplotlib.ticker import MaxNLocator # so we can force integer tick
labels later

# import the tdt library
import tdt
```

### Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCK_PATH = 'data/Algernon-180308-130351'
```

```
demo data ready
```

Set up the variables for the data you want to extract. We will extract channel 1 from the eNe1 snippet data store, created by the PCA Sorting gizmo, and use our PulseGen epoc event `PC0/` as our stimulus onset.

```

REF_EPOC = 'PC0/'
SNIP_STORE = 'eNe1'
SORTID = 'TankSort'
CHANNEL = 3
SORTCODE = 0          # set to 0 to use all sorts
TRANGE = [-0.3, 0.8]

```

Now read the specified data from our block into a Python structure. The `nodata` flag means that we are only interested in the snippet timestamps, not the actual snippet waveforms in this example.

```

data = tdt.read_block(BLOCK_PATH, evtype=['epochs', 'snips', 'scalars'],
                    sortname=SORTID, channel=CHANNEL, nodata=1)

```

```

read from t=0s to t=61.23s

```

## Use `epoc_filter` to extract data around our epoc event

Using the `t` parameter extracts data only from the time range around our epoc event.

```

raster_data = tdt.epoc_filter(data, REF_EPOC, t=TRANGE)

```

Adding the `tref` flag makes all of the timestamps relative to the epoc event, which is ideal for generating histograms.

```

hist_data = tdt.epoc_filter(data, REF_EPOC, t=TRANGE, tref=1)

```

And that's it! Your data is now in Python. The rest of the code is a simple plotting example. First, we'll find matching timestamps for our selected sort code (unit).

```

ts = raster_data.snips[SNIP_STORE].ts
if SORTCODE != 0:
    i = np.where(raster_data.snips[SNIP_STORE].sortcode == SORTCODE)[0]
    ts = ts[i]
if len(ts) == 0:
    raise Exception('no matching timestamps found')

num_trials = raster_data.time_ranges.shape[1]

```

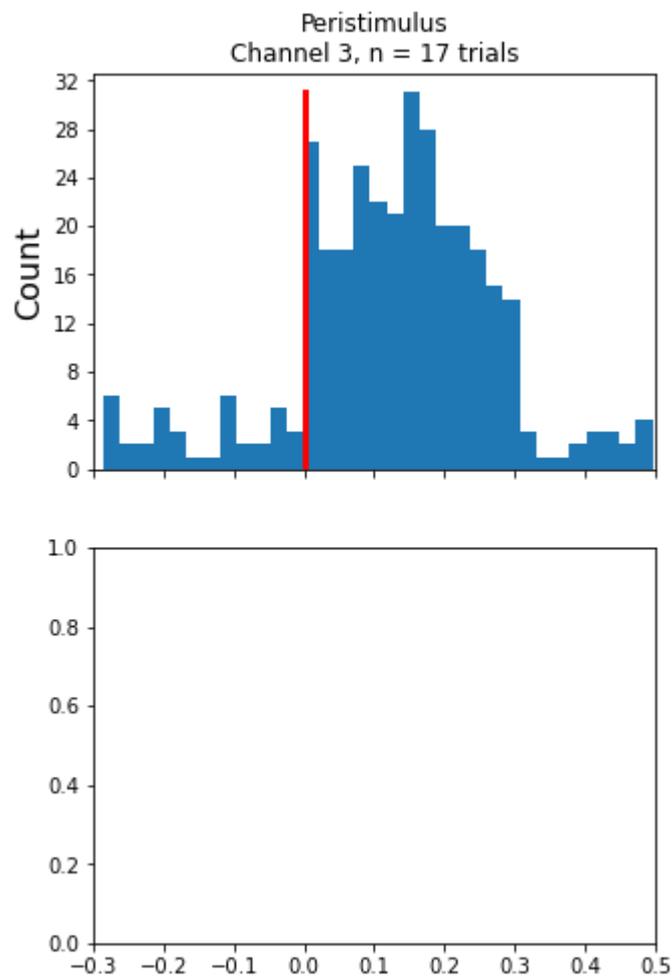
## Make the histogram plot

```
fig, (ax1, ax2) = plt.subplots(2, 1, sharex=True, figsize=(5, 8))

hist_ts = hist_data.snips[SNIP_STORE].ts
nbins = np.int64(np.floor(len(hist_ts)/10.))
hist_n = ax1.hist(hist_ts, nbins)[0]

ax1.axis('tight')
ax1.set_xlim(left=TRANGE[0], right=TRANGE[0]+TRANGE[1])
ax1.set_ylabel('Count', fontsize=16)
ax1.set_title('Peristimulus\nChannel {0}, n = {1} trials'.format(CHANNEL,
num_trials))
ax1.yaxis.set_major_locator(MaxNLocator(integer=True))

# Draw a vertical line at t=0.
ax1.plot([0, 0], [0, np.max(hist_n)], 'r-', linewidth=3)
plt.show()
```



## Creating the Raster Plot

```

# For the raster plot, make an array of lists containing timestamps for each
trial.
all_ts = [[] for x in range(num_trials)]
all_y = [[] for x in range(num_trials)]
for trial in range(num_trials):
    trial_on = raster_data.time_ranges[0, trial]
    trial_off = raster_data.time_ranges[1, trial]
    ind1 = ts >= trial_on
    ind2 = ts < trial_off
    trial_ts = ts[ind1 & ind2]
    all_ts[trial] = trial_ts - trial_on + TRANGE[0]
    all_y[trial] = (trial+1) * np.ones(len(trial_ts))

all_x = np.concatenate(all_ts)
all_y = np.concatenate(all_y)

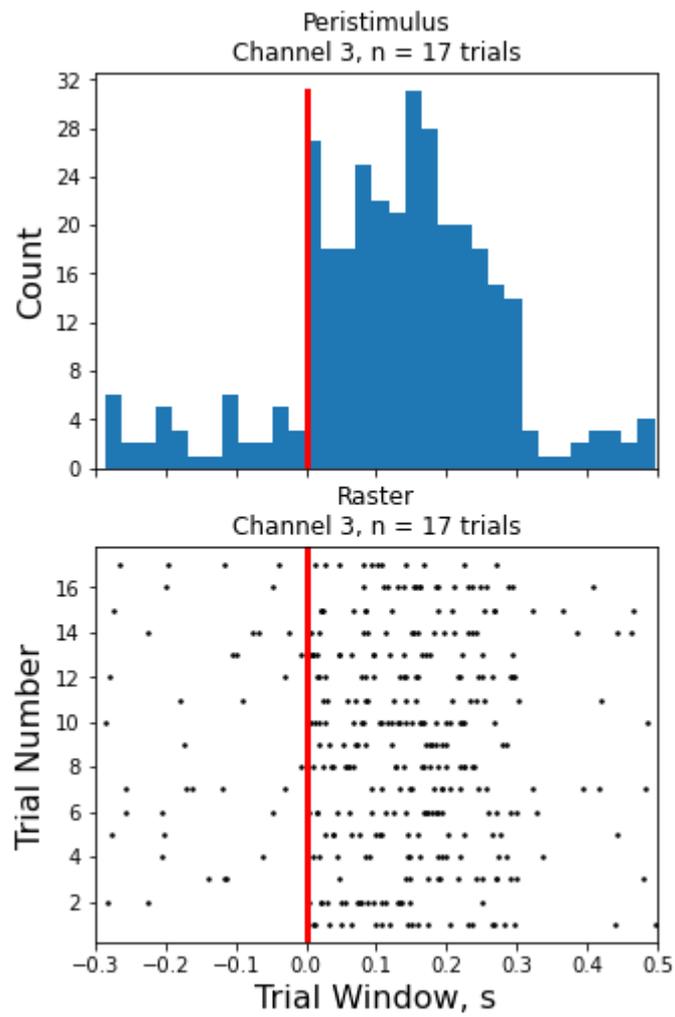
# Make the raster plot.
ax2.plot(all_x, all_y, 'k.', markersize=3)
ax2.axis('tight')
ax2.set_xlim(left=TRANGE[0], right=TRANGE[0]+TRANGE[1])
ax2.set_xlabel('Trial Window, s', fontsize=16)
ax2.set_ylabel('Trial Number', fontsize=16)
ax2.set_title('Raster\nChannel {0}, n = {1} trials'.format(CHANNEL,
num_trials))

# Draw a vertical line at t=0.
ax2.plot([0, 0], [0, trial+2], 'r-', linewidth=3)

ax2.yaxis.set_major_locator(MaxNLocator(integer=True))

plt.show()

```



## Note Filter Example

---

Import streaming EEG data into Python using **read\_block**

Filter around behavioral events that were timestamped by the user using the Run-time Notes feature in Synapse, using **epoc\_filter**

Plot each occurrence in a subplot organized by Note type

Good for sleep scoring and behavioral discrimination

### Housekeeping

Import the tdt package and other python packages we care about

```
# magic for Jupyter
%matplotlib inline

import matplotlib.pyplot as plt # standard Python plotting library
import numpy as np

import tdt
```

### Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCK_PATH = 'data/Subject1-180426-120951'
```

```
demo data ready
```

## Set up the variables for the data you want to extract.

We will extract channel 1 from the EEG1 stream data store.

```
STORE = 'EEG1'  
CHANNEL = 1  
ONSET = [-3] # relative onset, in seconds, from the note timestamp
```

Now read the specified data from our block into a Python structure

```
data = tdt.read_block(BLOCK_PATH, channel=CHANNEL)
```

```
Found Synapse note file: data/Subject1-180426-120951\Notes.txt  
read from t=0s to t=31.81s
```

All user notes are stored in a special epoc event called 'Note'

```
# find all the unique note values  
notes, counts = np.unique(data.epocs.Note.notes, return_counts=True)  
  
# find the highest number of occurrences (to inform our plot)  
maxOccur = np.max(counts)
```

## Loop through the notes for plotting

```

# some useful variables
num_notes = len(notes)
fs = data.streams[STORE].fs

fig = plt.figure(figsize=(14, 8))
for ind, note in enumerate(notes,1):

    print('Reading note:', note)

    # look at only the data around this note type
    filtered = tdt.epoc_filter(data, 'Note', values=note, t=ONSET)

    # for each note occurrence, plot the data from
    # the note onset to the next note onset
    n = len(filtered.streams[STORE].filtered)
    for j in range(n):
        plotInd = j * num_notes + ind
        ax = fig.add_subplot(maxOccur, num_notes, plotInd)

        # x-axis is the valid time ranges, in seconds
        len_wav = len(filtered.streams[STORE].filtered[j])
        ts = filtered.time_ranges[0,j] + np.linspace(1, len_wav, len_wav) / fs

        # plot the snippet, in microvolts
        y = np.transpose(1e6 * filtered.streams[STORE].filtered[j])
        trace1 = ax.plot(ts, y, lw=1, color='cornflowerblue')

        # if we specified an ONSET, draw the vertical line at the note onset
        if ONSET != 0:
            trace2 = ax.axvline(x = (ts[0]-ONSET),
                               color='slategray',
                               linewidth=3)

        # plot labels
        if j == 0:
            ax.set_title(note, fontsize=14)
        elif j == (n-1):
            if ind == 1:
                ax.set_ylabel('\u03BCV', fontsize=12)
                ax.set_xlabel('Time (s)', fontsize=12)

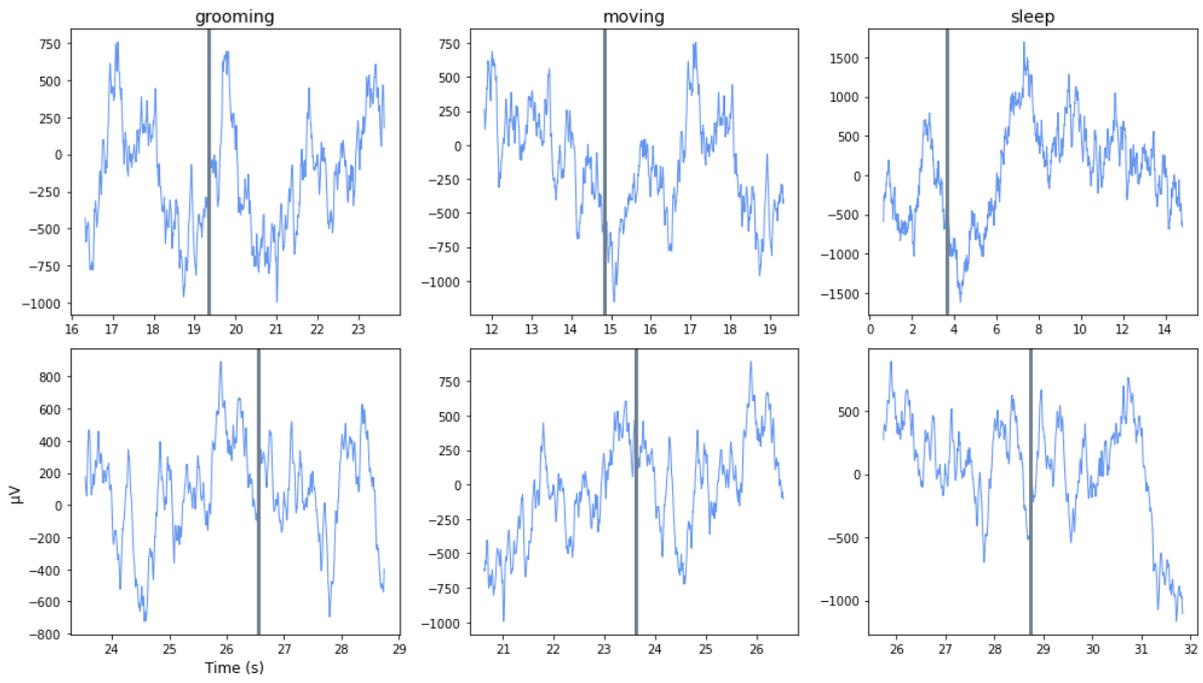
fig.tight_layout()

```

```

Reading note: grooming
Reading note: moving
Reading note: sleep

```



## Fiber Photometry Epoch Averaging Example

---

This example goes through fiber photometry analysis using techniques such as data smoothing, bleach detrending, and z-score analysis.

The epoch averaging was done using `epoc_filter`.

### Author Contributions:

TDT, David Root, and the Morales Lab contributed to the writing and/or conceptualization of the code.

The signal processing pipeline was inspired by the workflow developed by David Barker et al. (2017) for the Morales Lab.

The data used in the example were provided by David Root.

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### About the authors:

The Root lab and Morales lab investigate the neurobiology of reward, aversion, addiction, and depression.

TDT edits all user submissions in coordination with the contributing author(s) prior to publishing.

## Housekeeping

Import the tdt package and other python packages we care about.

```
# magic for Jupyter
%matplotlib inline

#import the read_block function from the tdt package
#also import other python packages we care about
import numpy as np
from sklearn.metrics import auc
import matplotlib.pyplot as plt # standard Python plotting library
import scipy.stats as stats

import tdt
```

## Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCKPATH = 'data/FiPho-180416'
```

```
demo data ready
```

```
# Jupyter has a bug that requires import of matplotlib outside of cell with
# matplotlib inline magic to properly apply rcParams
import matplotlib
matplotlib.rcParams['font.size'] = 16 #set font size for all plots
```

## Setup the variables for the data you want to extract

We will extract two different stream stores surrounding the 'PtAB' epoch event. We are interested in a specific event code for the shock onset.

```
REF_EPOC = 'PtAB' #event store name. This holds behavioral codes that are
# read through ports A & B on the front of the RZ
SHOCK_CODE = [64959] #shock onset event code we are interested in

# make some variables up here to so if they change in new recordings you won't
# have to change everything downstream
ISOS = '_4054' # 405nm channel. Formally STREAM_STORE1 in maltab example
GCaMP = '_4654' # 465nm channel. Formally STREAM_STORE2 in maltab example
TRANGE = [-10, 20] # window size [start time relative to epoc onset, window
duration]
BASELINE_PER = [-10, -6] # baseline period within our window
ARTIFACT = float("inf") # optionally set an artifact rejection level

#call read block - new variable 'data' is the full data structure
data = tdt.read_block(BLOCKPATH)
```

```
read from t=0s to t=583.86s
```

## Use epoc\_filter to extract data around our epoc event

Using the 't' parameter extracts data only from the time range around our epoc event.

Use the 'values' parameter to specify allowed values of the REF\_EPOC to extract.

For stream events, the chunks of data are stored in cell arrays structured as

```
data.streams[GCaMP].filtered
```

```
data = tdt.epoc_filter(data, REF_EPOC, t=TRANGE, values=SHOCK_CODE)
```

```
# Optionally remove artifacts. If any waveform is above ARTIFACT level, or
# below -ARTIFACT level, remove it from the data set.
total1 = np.size(data.streams[GCaMP].filtered)
total2 = np.size(data.streams[ISOS].filtered)

# List comprehension checking if any single array in 2D filtered array is >
Artifact or < -Artifact
data.streams[GCaMP].filtered = [x for x in data.streams[GCaMP].filtered
                                if not np.any(x > ARTIFACT) or np.any(x < -
ARTIFACT)]
data.streams[ISOS].filtered = [x for x in data.streams[ISOS].filtered
                                if not np.any(x > ARTIFACT) or np.any(x < -
ARTIFACT)]

# Get the total number of rejected arrays
bad1 = total1 - np.size(data.streams[GCaMP].filtered)
bad2 = total2 - np.size(data.streams[ISOS].filtered)
total_artifacts = bad1 + bad2
```

Applying a time filter to a uniformly sampled signal means that the length of each segment could vary by one sample. Let's find the minimum length so we can trim the excess off before calculating the mean.

```

# More examples of list comprehensions
min1 = np.min([np.size(x) for x in data.streams[GCaMP].filtered])
min2 = np.min([np.size(x) for x in data.streams[ISOS].filtered])
data.streams[GCaMP].filtered = [x[1:min1] for x in
data.streams[GCaMP].filtered]
data.streams[ISOS].filtered = [x[1:min2] for x in data.streams[ISOS].filtered]

# Downsample and average 10x via a moving window mean
N = 10 # Average every 10 samples into 1 value
F405 = []
F465 = []
for lst in data.streams[ISOS].filtered:
    small_lst = []
    for i in range(0, min2, N):
        small_lst.append(np.mean(lst[i:i+N-1])) # This is the moving window
mean
    F405.append(small_lst)

for lst in data.streams[GCaMP].filtered:
    small_lst = []
    for i in range(0, min1, N):
        small_lst.append(np.mean(lst[i:i+N-1]))
    F465.append(small_lst)

#Create a mean signal, standard error of signal, and DC offset
meanF405 = np.mean(F405, axis=0)
stdF405 = np.std(F405, axis=0)/np.sqrt(len(data.streams[ISOS].filtered))
dcF405 = np.mean(meanF405)
meanF465 = np.mean(F465, axis=0)
stdF465 = np.std(F465, axis=0)/np.sqrt(len(data.streams[GCaMP].filtered))
dcF465 = np.mean(meanF465)

```

## Plot epoc averaged response

```

# Create the time vector for each stream store
ts1 = TRANGE[0] + np.linspace(1, len(meanF465), len(meanF465))/
data.streams[GCaMP].fs*N
ts2 = TRANGE[0] + np.linspace(1, len(meanF405), len(meanF405))/
data.streams[ISOS].fs*N

# Subtract DC offset to get signals on top of one another
meanF405 = meanF405 - dcF405
meanF465 = meanF465 - dcF465

# Start making a figure with 4 subplots
# First plot is the 405 and 465 averaged signals
fig = plt.figure(figsize=(9, 14))
ax0 = fig.add_subplot(411) # work with axes and not current plot (plt.)

# Plotting the traces
p1, = ax0.plot(ts1, meanF465, linewidth=2, color='green', label='GCaMP')
p2, = ax0.plot(ts2, meanF405, linewidth=2, color='blueviolet', label='ISOS')

# Plotting standard error bands
p3 = ax0.fill_between(ts1, meanF465+stdF465, meanF465-stdF465,
                      facecolor='green', alpha=0.2)
p4 = ax0.fill_between(ts2, meanF405+stdF405, meanF405-stdF405,
                      facecolor='blueviolet', alpha=0.2)

# Plotting a line at t = 0
p5 = ax0.axvline(x=0, linewidth=3, color='slategray', label='Shock Onset')

# Finish up the plot
ax0.set_xlabel('Seconds')
ax0.set_ylabel('mV')
ax0.set_title('Foot Shock Response, %i Trials (%i Artifacts Removed)'
              % (len(data.streams[GCaMP].filtered), total_artifacts))
ax0.legend(handles=[p1, p2, p5], loc='upper right')
ax0.set_ylim(min(np.min(meanF465-stdF465), np.min(meanF405-stdF405)),
             max(np.max(meanF465+stdF465), np.max(meanF405+stdF405)))
ax0.set_xlim(TRANGE[0], TRANGE[1]+TRANGE[0]);

plt.close() # Jupyter cells will output any figure calls made, so if you
don't want to see it just yet, close existing axis
# https://stackoverflow.com/questions/18717877/prevent-plot-from-
showing-in-jupyter-notebook
# Note that this is not good code practice - Jupyter lends it
self to these types of bad workarounds

```

## Fitting 405 channel onto 465 channel to detrend signal bleaching

Scale and fit data. Algorithm sourced from Tom Davidson's Github: [https://github.com/tjd2002/tjd-shared-code/blob/master/matlab/photometry/FP\\_normalize.m](https://github.com/tjd2002/tjd-shared-code/blob/master/matlab/photometry/FP_normalize.m)

```

Y_fit_all = []
Y_dF_all = []
for x, y in zip(F405, F465):
    x = np.array(x)
    y = np.array(y)
    bls = np.polyfit(x, y, 1)
    fit_line = np.multiply(bls[0], x) + bls[1]
    Y_fit_all.append(fit_line)
    Y_dF_all.append(y-fit_line)

# Getting the z-score and standard error
zall = []
for dF in Y_dF_all:
    ind = np.where((np.array(ts2)<BASELINE_PER[1]) &
(np.array(ts2)>BASELINE_PER[0]))
    zb = np.mean(dF[ind])
    zsd = np.std(dF[ind])
    zall.append((dF - zb)/zsd)

zerror = np.std(zall, axis=0)/np.sqrt(np.size(zall, axis=0))

```

## Heat Map based on z score of 405 fit subtracted 465

```

ax1 = fig.add_subplot(412)
cs = ax1.imshow(zall, cmap=plt.cm.Greys, interpolation='none', aspect="auto",
                extent=[TRANGE[0], TRANGE[1]+TRANGE[0], 0,
len(data.streams[GCaMP].filtered)])
cbar = fig.colorbar(cs, pad=0.01, fraction=0.02)

ax1.set_title('Individual z-Score Traces')
ax1.set_ylabel('Trials')
ax1.set_xlabel('Seconds from Shock Onset')

plt.close() # Suppress figure output again

```

## Plot the z-score trace for the 465 with std error bands

```

ax2 = fig.add_subplot(413)
p6 = ax2.plot(ts2, np.mean(zall, axis=0), linewidth=2, color='green',
label='GCaMP')
p7 = ax2.fill_between(ts1, np.mean(zall, axis=0)+zerror
                    ,np.mean(zall, axis=0)-zerror, facecolor='green',
alpha=0.2)
p8 = ax2.axvline(x=0, linewidth=3, color='slategray', label='Shock Onset')
ax2.set_ylabel('z-Score')
ax2.set_xlabel('Seconds')
ax2.set_xlim(TRANGE[0], TRANGE[1]+TRANGE[0])
ax2.set_title('Foot Shock Response')

plt.close()

```

## Quantify changes as an area under the curve for cue (-5 sec) vs shock (0 sec)

```

AUC = [] # cue, shock
ind1 = np.where((np.array(ts2)<-3) & (np.array(ts2)>-5))
AUC1= auc(ts2[ind1], np.mean(zall, axis=0)[ind1])
ind2 = np.where((np.array(ts2)>0) & (np.array(ts2)<2))
AUC2= auc(ts2[ind2], np.mean(zall, axis=0)[ind2])
AUC.append(AUC1)
AUC.append(AUC2)

# Run a two-sample T-test
t_stat,p_val = stats.ttest_ind(np.mean(zall, axis=0)[ind1],
                             np.mean(zall, axis=0)[ind2], equal_var=False)

```

## Make a bar plot

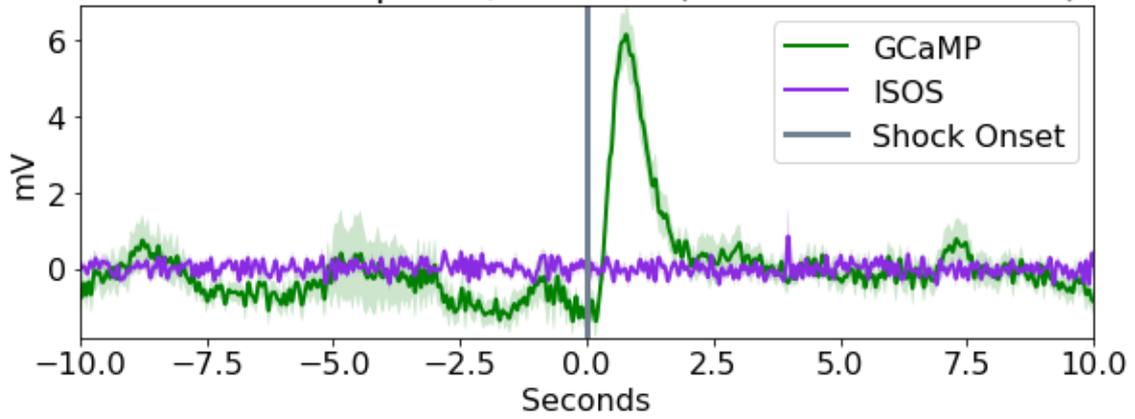
```
ax3 = fig.add_subplot(414)
p9 = ax3.bar(np.arange(len(AUC)), AUC, color=[.8, .8, .8], align='center',
alpha=0.5)

# statistical annotation
x1, x2 = 0, 1 # columns indices for labels
y, h, col = max(AUC) + 2, 2, 'k'
ax3.plot([x1, x1, x2, x2], [y, y+h, y+h, y], lw=1.5, c=col)
p10 = ax3.text((x1+x2)*.5, y+h, "*", ha='center', va='bottom', color=col)

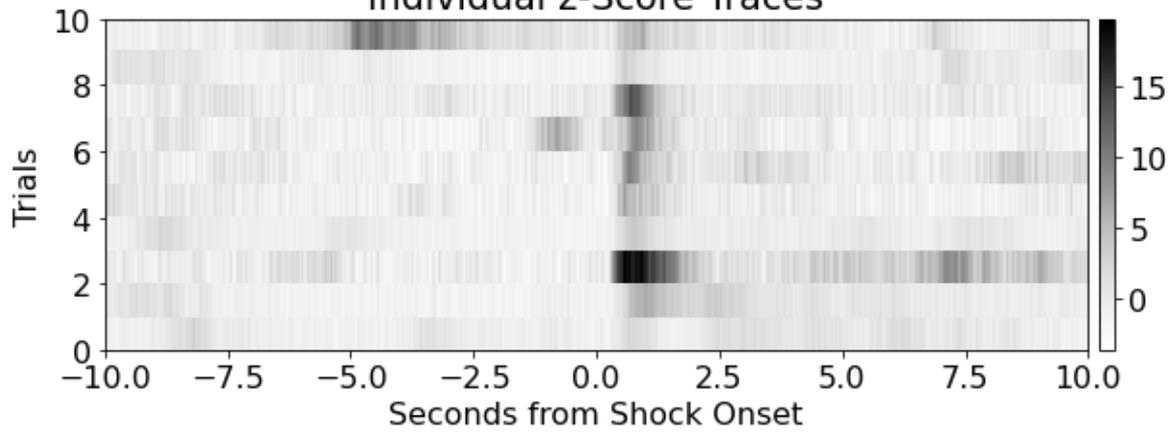
# Finish up the plot
ax3.set_ylim(0, y+2*h)
ax3.set_ylabel('AUC')
ax3.set_title('Cue vs Shock Response Changes')
ax3.set_xticks(np.arange(-1, len(AUC)+1))
ax3.set_xticklabels(['', 'Cue', 'Shock', ''])

fig.tight_layout()
fig
```

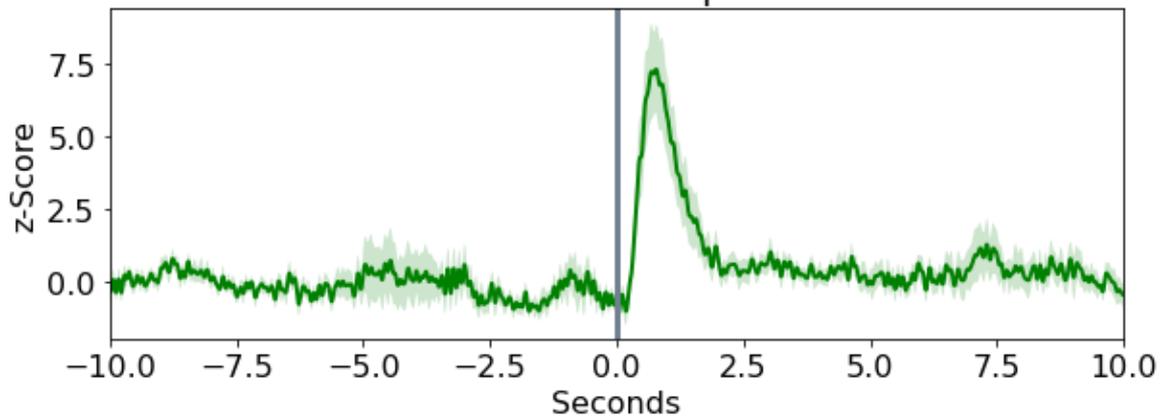
Foot Shock Response, 10 Trials (0 Artifacts Removed)



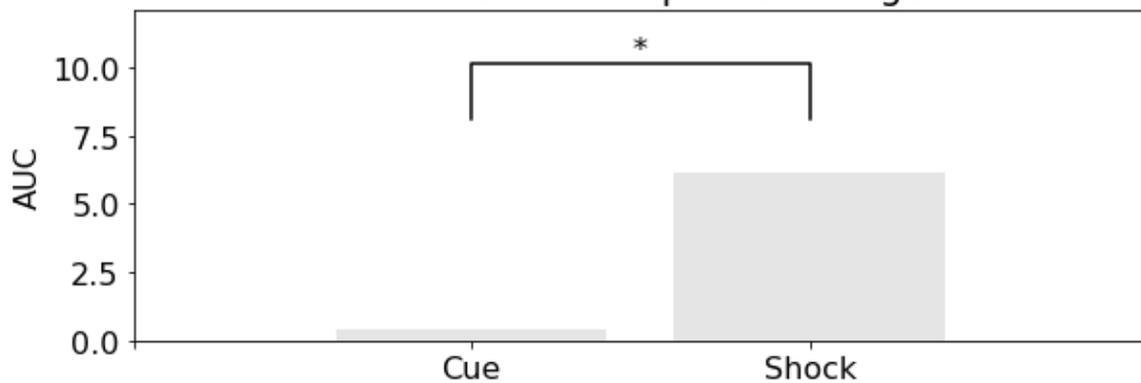
Individual z-Score Traces



Foot Shock Response



Cue vs Shock Response Changes



## Licking Bout Epoc Filtering

---

This example looks at fiber photometry data in the VTA where subjects are provided sucrose water after a fasting period.

Lick events are captured as TTL pulses.

Objective is to combine many consecutive licking events into a single event based on time difference and lick count thresholds.

New lick bout events can then be used for clear peri-event filtering.

## Housekeeping

Import the tdt package and other python packages we care about.

```
# Jupyter magic
%matplotlib inline

import numpy as np
import matplotlib.pyplot as plt # standard Python plotting library

# import the tdt library
import tdt
```

## Importing the Data

This example uses our [example data sets](#). To import your own data, replace `BLOCK_PATH` with the full path to your own data block.

In Synapse, you can find the block path in the database. Go to Menu → History. Find your block, then Right-Click → Copy path to clipboard.

```
tdt.download_demo_data()
BLOCKPATH = 'data/VTA4-190125-100559'
data = tdt.read_block(BLOCKPATH)
```

```
demo data ready
Found Synapse note file: data/VTA4-190125-100559\Notes.txt
read from t=0s to t=785.44s
```

## Basic plotting and artifact removal

```
#Jupyter has a bug that requires import of matplotlib outside of cell with
matplotlib inline magic to properly apply rcParams
```

```
import matplotlib
matplotlib.rcParams['font.size'] = 18 # set font size for all figures
```

```
# Make some variables up here to so if they change in new recordings you
won't have to change everything downstream
```

```
GCAMP = '_480G' # GCaMP channel
ISOS = '_405G' # Isosbestic channel
LICK = 'Ler_'
```

```
# Make a time array based on the number of samples and sample freq of
# the demodulated streams
```

```
time = np.linspace(1, len(data.streams[GCAMP].data),
len(data.streams[GCAMP].data))/data.streams[GCAMP].fs
```

```
# Plot both unprocessed demodulated stream
```

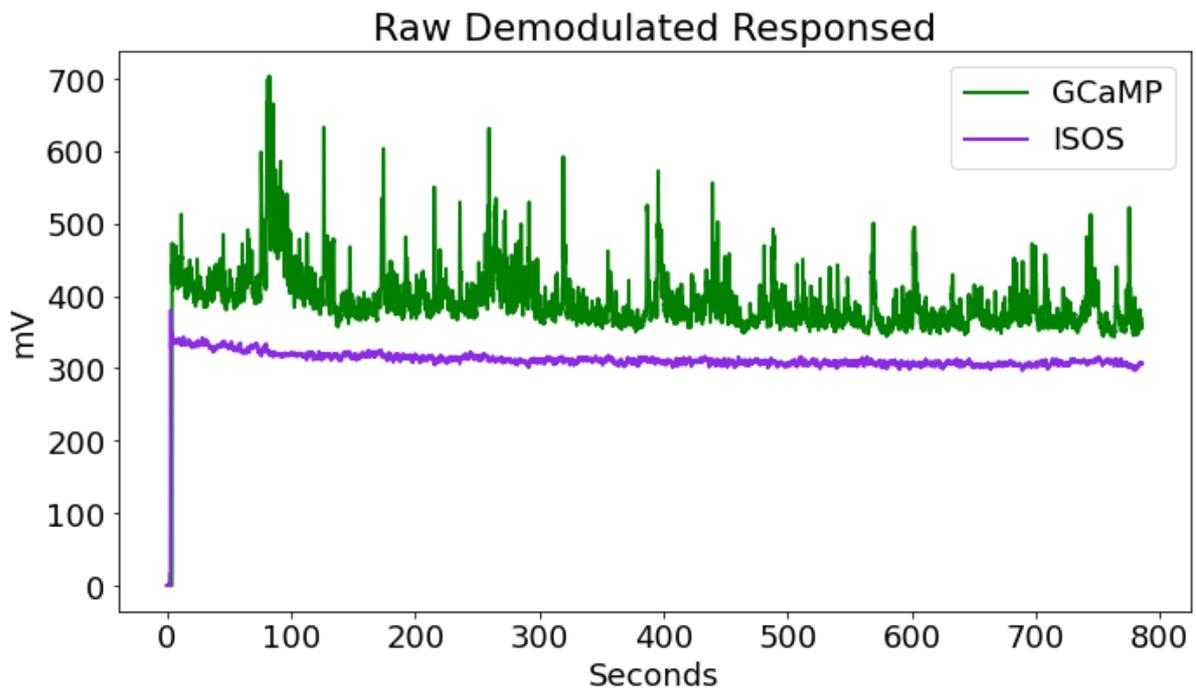
```
fig1 = plt.figure(figsize=(10,6))
ax0 = fig1.add_subplot(111)
```

```
# Plotting the traces
```

```
p1, = ax0.plot(time, data.streams[GCAMP].data, linewidth=2, color='green',
label='GCaMP')
p2, = ax0.plot(time, data.streams[ISOS].data, linewidth=2, color='blueviolet',
label='ISOS')
```

```
ax0.set_ylabel('mV')
ax0.set_xlabel('Seconds')
ax0.set_title('Raw Demodulated Responses')
ax0.legend(handles=[p1,p2], loc='upper right')
fig1.tight_layout()
```

```
# Jupyter for some reason (sometimes) shows the figure without be called,
# Likely when plt.figure() is called
# otherwise you would call fig in a line by itself like:
# fig
```



## Artifact Removal

```

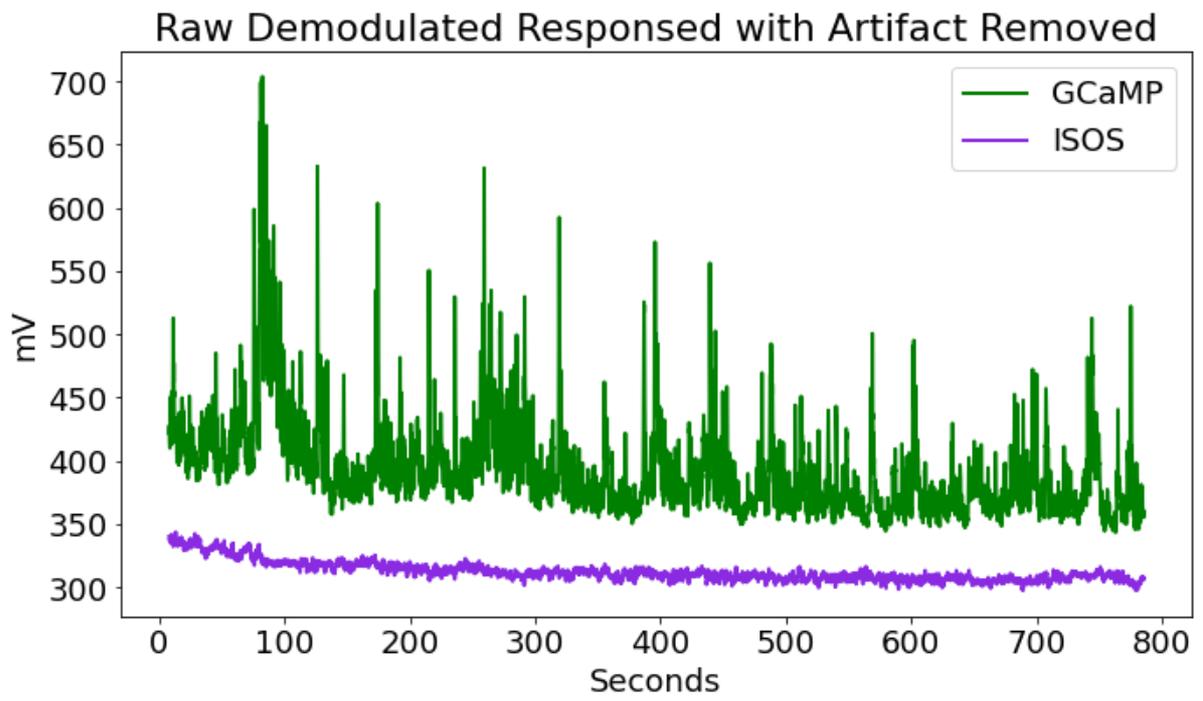
# There is often a large artifact on the onset of LEDs turning on
# Remove data below a set time t
t = 8
inds = np.where(time>t)
ind = inds[0][0]
time = time[ind:] # go from ind to final index
data.streams[GCAMP].data = data.streams[GCAMP].data[ind:]
data.streams[ISOS].data = data.streams[ISOS].data[ind:]

# Plot again at new time range
fig2 = plt.figure(figsize=(10, 6))
ax1 = fig2.add_subplot(111)

# Plotting the traces
p1, = ax1.plot(time,data.streams[GCAMP].data, linewidth=2, color='green',
label='GCaMP')
p2, = ax1.plot(time,data.streams[ISOS].data, linewidth=2, color='blueviolet',
label='ISOS')

ax1.set_ylabel('mV')
ax1.set_xlabel('Seconds')
ax1.set_title('Raw Demodulated Responed with Artifact Removed')
ax1.legend(handles=[p1,p2],loc='upper right')
fig2.tight_layout()
# fig

```



## Downsample Data Doing Local Averaging

```

# Average around every Nth point and downsample Nx
N = 10 # Average every 10 samples into 1 value
F405 = []
F465 = []

for i in range(0, len(data.streams[GCAMP].data), N):
    F465.append(np.mean(data.streams[GCAMP].data[i:i+N-1])) # This is the
moving window mean
data.streams[GCAMP].data = F465

for i in range(0, len(data.streams[ISOS].data), N):
    F405.append(np.mean(data.streams[ISOS].data[i:i+N-1]))
data.streams[ISOS].data = F405

#decimate time array to match length of demodulated stream
time = time[:N] # go from beginning to end of array in steps on N
time = time[:len(data.streams[GCAMP].data)]

# Detrending and dFF
# Full trace dFF according to Lerner et al. 2015
# https://dx.doi.org/10.1016/j.cell.2015.07.014
# dFF using 405 fit as baseline

x = np.array(data.streams[ISOS].data)
y = np.array(data.streams[GCAMP].data)
bls = np.polyfit(x, y, 1)
Y_fit_all = np.multiply(bls[0], x) + bls[1]
Y_dF_all = y - Y_fit_all

dFF = np.multiply(100, np.divide(Y_dF_all, Y_fit_all))
std_dFF = np.std(dFF)

```

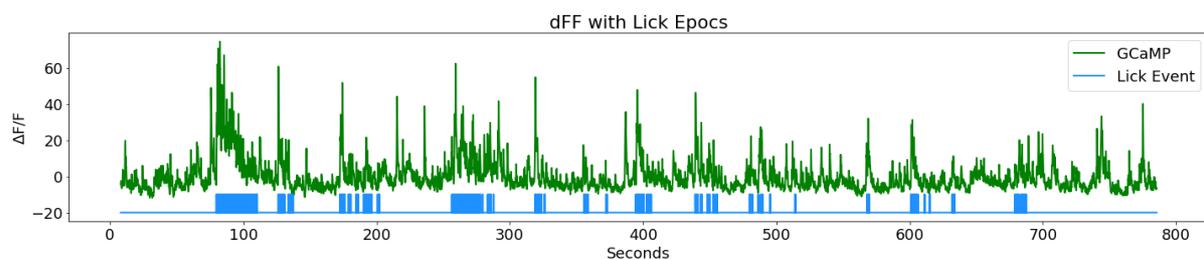
## Turn Licking Events into Lick Bouts

```
# First make a continuous time series of Licking TTL events (epocs) and plot
LICK_on = data.epocs[LICK].onset
LICK_off = data.epocs[LICK].offset
# Add the first and last time stamps to make tails on the TTL stream
LICK_x = np.append(np.append(time[0], np.reshape(np.kron([LICK_on, LICK_off],
    np.array([[1], [1]])).T, [1,-1])[0]), time[-1])
sz = len(LICK_on)
d = data.epocs[LICK].data
# Add zeros to beginning and end of 0,1 value array to match len of LICK_x
LICK_y = np.append(np.append(0, np.reshape(np.vstack([np.zeros(sz),
    d, d, np.zeros(sz)]).T, [1, -1])[0]), 0)

y_scale = 10 #adjust according to data needs
y_shift = -20 #scale and shift are just for aesthetics
```

```
# First subplot in a series: dFF with lick epocs
fig3 = plt.figure(figsize=(20,12))
ax2 = fig3.add_subplot(311)

p1, = ax2.plot(time, dFF, linewidth=2, color='green', label='GCaMP')
p2, = ax2.plot(LICK_x, y_scale*LICK_y+y_shift, linewidth=2,
color='dodgerblue', label='Lick Event')
ax2.set_ylabel(r'$\Delta F/F$')
ax2.set_xlabel('Seconds')
ax2.set_title('dFF with Lick Epocs')
ax2.legend(handles=[p1,p2], loc='upper right')
fig3.tight_layout()
```



## Lick Bout Logic

Now combine lick epochs that happen in close succession to make a single on/off event (a lick BOUT). Top view logic: if difference between consecutive lick onsets is below a certain time threshold and there was more than one lick in a row, then consider it as one bout, otherwise it is its own bout. Also, make sure a minimum number of licks was reached to call it a bout.

```

LICK_EVENT = 'LICK_EVENT'

LICK_DICT = {
    "name":LICK_EVENT,
    "onset":[],
    "offset":[],
    "type_str":data.epocs[LICK].type_str,
    "data":[]
}
# pass StructType our new dictionary to make keys and values
data.epocs.LICK_EVENT = tdt.StructType(LICK_DICT)

lick_on_diff = np.diff(data.epocs[LICK].onset)
BOUT_TIME_THRESHOLD = 10
lick_diff_ind = np.where(lick_on_diff >= BOUT_TIME_THRESHOLD)[0]
#for some reason np.where returns a 2D array, hence the [0]

# Make an onset/ offset array based on threshold indicies
diff_ind = 0
for ind in lick_diff_ind:
    # BOUT onset is thresholded onset index of lick epoc event
    data.epocs[LICK_EVENT].onset.append(data.epocs[LICK].onset[diff_ind])
    # BOUT offset is thresholded offset of lick event before next onset
    data.epocs[LICK_EVENT].offset.append(data.epocs[LICK].offset[ind])
    # set the values for data, arbitrary 1
    data.epocs[LICK_EVENT].data.append(1)
    diff_ind = ind + 1

# special case for last event to handle lick event offset indexing
data.epocs[LICK_EVENT].onset.append(data.epocs[LICK].onset[lick_diff_ind[-1]
+1])
data.epocs[LICK_EVENT].offset.append(data.epocs[LICK].offset[-1])
data.epocs[LICK_EVENT].data.append(1)

# Now determine if it was a 'real' lick bout by thresholding by some
# user-set number of licks in a row
MIN_LICK_THRESH = 4 #four licks or more make a bout
licks_array = []

# Find number of licks in licks_array between onset and offset of
# our new lick BOUT LICK_EVENT
for on, off in
zip(data.epocs[LICK_EVENT].onset,data.epocs[LICK_EVENT].offset):
    licks_array.append(
        len(np.where((data.epocs[LICK].onset >= on) & (data.epocs[LICK].onset
<= off))[0]))

# Remove onsets, offsets, and data of thrown out events
licks_array = np.array(licks_array)
inds = np.where(licks_array<MIN_LICK_THRESH)[0]

```

```

for index in sorted(inds, reverse=True):
    del data.epocs[LICK_EVENT].onset[index]
    del data.epocs[LICK_EVENT].offset[index]
    del data.epocs[LICK_EVENT].data[index]

# Make a continuous time series for lick BOUTS for plotting
LICK_EVENT_on = data.epocs[LICK_EVENT].onset
LICK_EVENT_off = data.epocs[LICK_EVENT].offset
LICK_EVENT_x = np.append(time[0], np.append(
    np.reshape(np.kron([LICK_EVENT_on, LICK_EVENT_off], np.array([[1],
[1]])).T, [1,-1])[0], time[-1]))
sz = len(LICK_EVENT_on)
d = data.epocs[LICK_EVENT].data
LICK_EVENT_y = np.append(np.append(
    0, np.reshape(np.vstack([np.zeros(sz), d, d, np.zeros(sz)]).T, [1,-1])
[0]), 0)

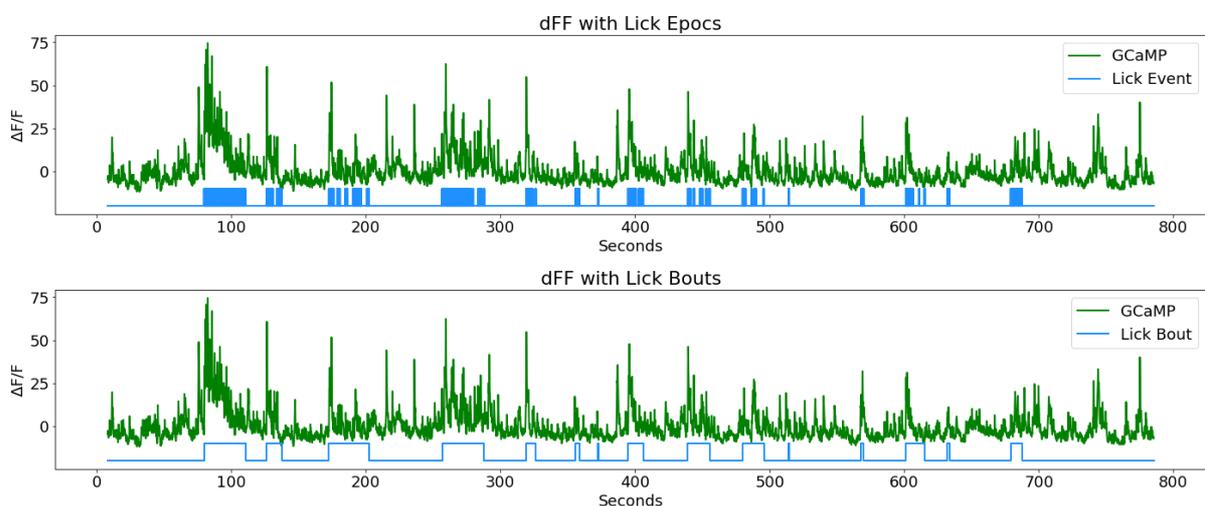
```

## Plot dFF with newly defined lick bouts

```

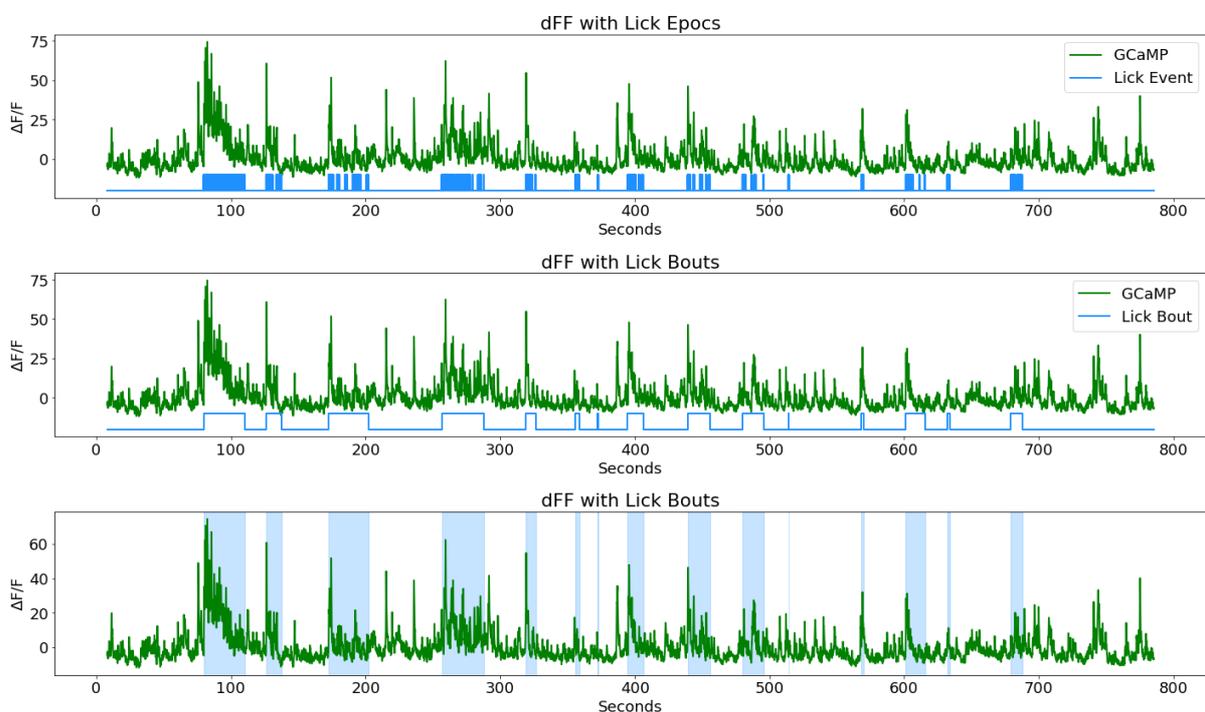
ax3 = fig3.add_subplot(312)
p1, = ax3.plot(time, dFF, linewidth=2, color='green', label='GCaMP')
p2, = ax3.plot(LICK_EVENT_x, y_scale*LICK_EVENT_y+y_shift, linewidth=2,
color='dodgerblue', label='Lick Bout')
ax3.set_ylabel(r'$\Delta F/F$')
ax3.set_xlabel('Seconds')
ax3.set_title('dFF with Lick Bouts')
ax3.legend(handles=[p1, p2], loc='upper right')
fig3.tight_layout()
fig3

```



## Make nice area fills instead of epocs for aesthetics

```
ax4 = fig3.add_subplot(313)
p1, = ax4.plot(time, dFF, linewidth=2, color='green', label='GCaMP')
for on, off in zip(data.epocs[LICK_EVENT].onset,
data.epocs[LICK_EVENT].offset):
    ax4.axvspan(on, off, alpha=0.25, color='dodgerblue')
ax4.set_ylabel(r'\Delta F/F')
ax4.set_xlabel('Seconds')
ax4.set_title('dFF with Lick Bouts')
fig3.tight_layout()
fig3
```



## Time Filter Around Lick Bout Epocs

Note that we are using dFF of the full time series, not peri-event dFF where  $f_0$  is taken from a pre-event baseline period.

```

PRE_TIME = 5 # five seconds before event onset
POST_TIME = 10 # ten seconds after
fs = data.streams[GCAMP].fs/N #recall we downsampled by N = 10 earlier

# time span for peri-event filtering, PRE and POST, in samples
TRANGE = [-PRE_TIME*np.floor(fs), POST_TIME*np.floor(fs)]

dFF_snips = []
array_ind = []
pre_stim = []
post_stim = []

for on in data.epocs[LICK_EVENT].onset:
    # If the bout cannot include pre-time seconds before event, make zero
    if on < PRE_TIME:
        dFF_snips.append(np.zeros(TRANGE[1]-TRANGE[0]))
    else:
        # find first time index after bout onset
        array_ind.append(np.where(time > on)[0][0])
        # find index corresponding to pre and post stim durations
        pre_stim.append(array_ind[-1] + TRANGE[0])
        post_stim.append(array_ind[-1] + TRANGE[1])
        dFF_snips.append(dFF[int(pre_stim[-1]):int(post_stim[-1])])

# Make all snippets the same size based on min snippet length
min1 = np.min([np.size(x) for x in dFF_snips])
dFF_snips = [x[1:min1] for x in dFF_snips]

mean_dFF_snips = np.mean(dFF_snips, axis=0)
std_dFF_snips = np.std(mean_dFF_snips, axis=0)

peri_time = np.linspace(1, len(mean_dFF_snips), len(mean_dFF_snips))/fs -
PRE_TIME

```

## Make a Peri-Event Stimulus Plot and Heat Map

```

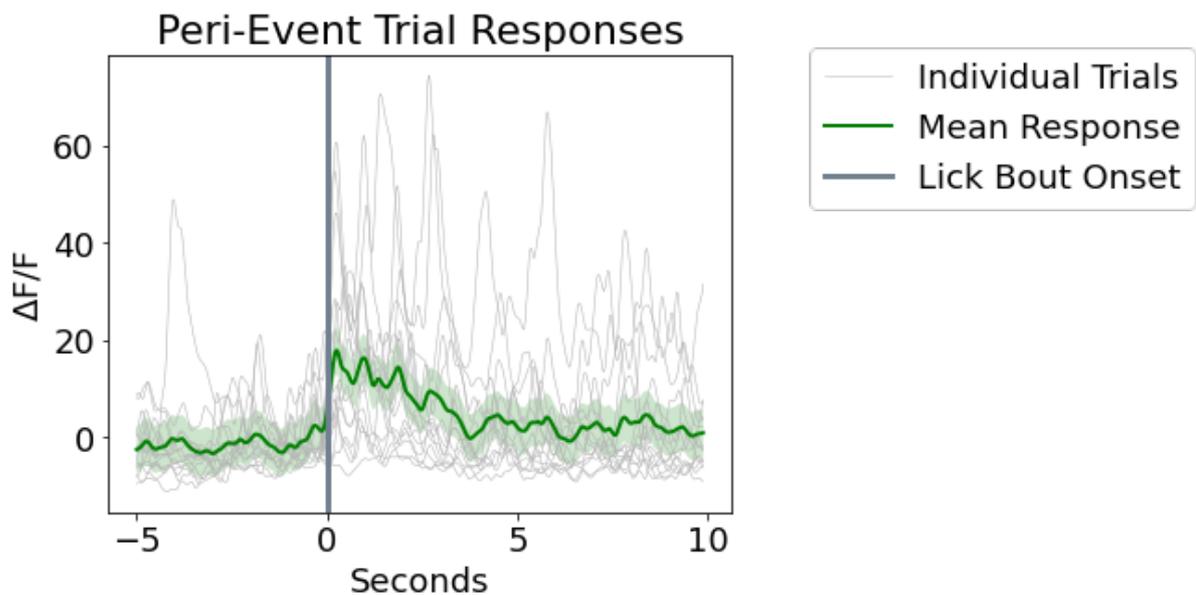
fig4 = plt.figure(figsize=(6,10))
ax5 = fig4.add_subplot(211)

for snip in dFF_snips:
    p1, = ax5.plot(peri_time, snip, linewidth=.5, color=[.7, .7, .7],
label='Individual Trials')
    p2, = ax5.plot(peri_time, mean_dFF_snips, linewidth=2, color='green',
label='Mean Response')

# Plotting standard error bands
p3 = ax5.fill_between(peri_time, mean_dFF_snips+std_dFF_snips,
                    mean_dFF_snips-std_dFF_snips, facecolor='green',
alpha=0.2)
p4 = ax5.axvline(x=0, linewidth=3, color='slategray', label='Lick Bout Onset')

ax5.axis('tight')
ax5.set_xlabel('Seconds')
ax5.set_ylabel(r'$\Delta F/F$')
ax5.set_title('Peri-Event Trial Responses')
ax5.legend(handles=[p1, p2, p4], bbox_to_anchor=(1.1, 1.05));

```



```

ax6 = fig4.add_subplot(212)
cs = ax6.imshow(dFF_snips, cmap=plt.cm.Greys,
               interpolation='none', extent=[-
PRE_TIME, POST_TIME, len(dFF_snips), 0],)
ax6.set_ylabel('Trial Number')
ax6.set_yticks(np.arange(.5, len(dFF_snips), 2))
ax6.set_yticklabels(np.arange(0, len(dFF_snips), 2))
fig4.colorbar(cs)
fig4

```

