

# Source Localization with MEG

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MEG Short Course

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# Overview – SAM Workflow

- Things you need:
  - ✓ Structural MRI including the nose and ears
  - ✓ MEG data
  - ✓ Research hypotheses
- Process the MRI – mark the fiducials, create head conductivity model
- Process the MEG – add event markers, filter, add MRI-based head model
- `sam_cov` – create covariance matrices
- `sam_wts` – create beamformer weights
- `sam_3d`, etc. – create source images
- Group stats – AFNI

# MRI Processing

```
#!/bin/sh

if [ "$#" -ne 1 ]; then
    echo usage: $0 hashcode
    exit
fi

hashcode="$1"

cd mri/${hashcode}

orthohull -c # clean up old files
orthohull -t # ortho + tlrc
```

Once the fiducial markers have been placed, the MRI is rotated into *ortho* space, and the skull removed.

From that, we compute a conductivity model.

This is accomplished by the `orthohull` script. It also creates Talairach (or MNI) versions, for group averaging.

Usually a script like `doOrthohull.sh` is used.

# MRI Processing



# MEG Event Markers – doThresh.sh

```
#!/bin/sh

if [ "$#" -ne 1 ]; then
    echo usage: $0 hashcode
    exit
fi

hashcode="$1"
ds=`grep ${hashcode} dslist`
echo $ds
dirname=`dirname $ds`
setname=`basename $ds .ds`

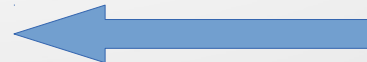
# threshold the ADCs for the ASVEF task

thresholdDetect2 -np -i -m leftpuff -a .5 -d .5 -dt 1 -c UADC001 $ds
thresholdDetect2 -np -i -m rightpuff -a .5 -d .5 -dt 1 -c UADC002 $ds
thresholdDetect2 -np -i -m beep -a .25 -d .25 -dt 1 -c UADC003 $ds
thresholdDetect2 -np -i -m check -a .5 -d .5 -dt 1 -c UADC016 $ds

newDs -f -filter processing.cfg $ds ${dirname}/${setname}-f.ds
```

Along with the MEG data, various information is collected in separate digital and analog channels. Analog channels store things like button presses or stimulus triggers. Here, ADCs 1 & 2 store airpuff stimulator triggers. These can be given names using the `thresholdDetect` program, which is a CTF utility.

Next we apply processing filters.



# processing.cfg

```
processing
{
    // balance: order, adapted
    // (adapted=0 -> not adapted)
    // (adapted=1 -> adapted)
    balance:      3,0
    // lowpass: enable, filterOrder, fc
    lowpass:      0,4,0.0000000000000000
    // highpass: enable, filterOrder, fc
    highpass:     1,4,0.5000000000000000
    // bandreject: enable, filterOrder, fc1, fc2
    bandreject:   0,4,0.0000000000000000,0.0000000000000000
    // bandpass: enable, filterOrder, fc1, fc2
    bandpass:    0,4,0.0000000000000000,0.0000000000000000
    // bandreject: enable, filterOrder, fc1, fc2
    bandreject:   1,2,59.5000000000000000,60.5000000000000000
    // bandreject: enable, filterOrder, fc1, fc2
    bandreject:   1,2,119.5000000000000000,120.5000000000000000
    // bandreject: enable, filterOrder, fc1, fc2
    bandreject:   1,2,179.5000000000000000,180.5000000000000000
    // bandreject: enable, filterOrder, fc1, fc2
    bandreject:   1,2,239.5000000000000000,240.5000000000000000
    // offset: enable, baselineSelection, startPt, endPt
    // (baseline=0 --> use pretrigger data)
    // (baseline=1 --> use from startPt to endPt)
    // (baseline=2 --> use whole trial)
    // (baseline+=10 --> do trend removal)
    offset: 1,2,1,1
}
```

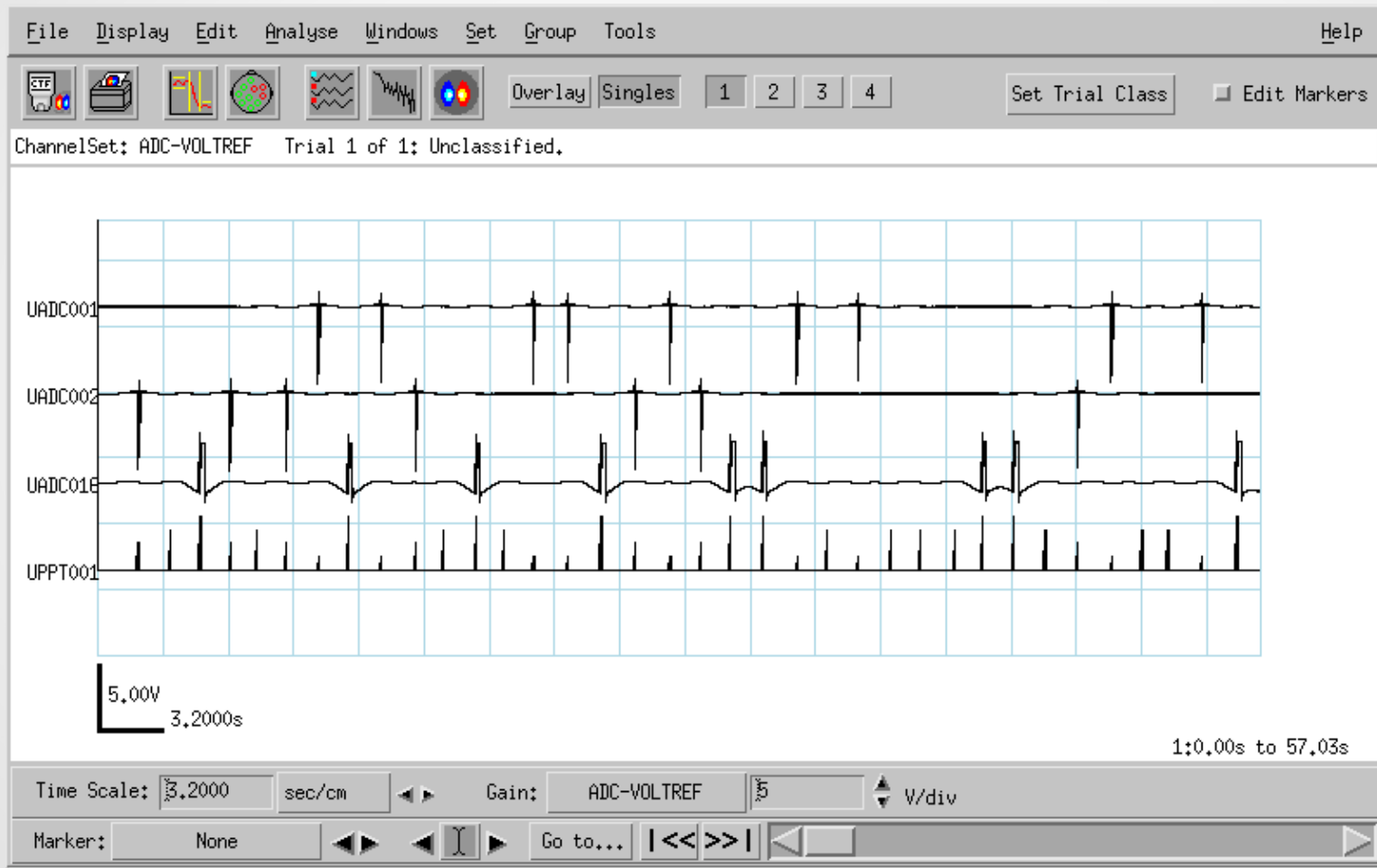
3rd gradient (default)

High pass filter (optional)

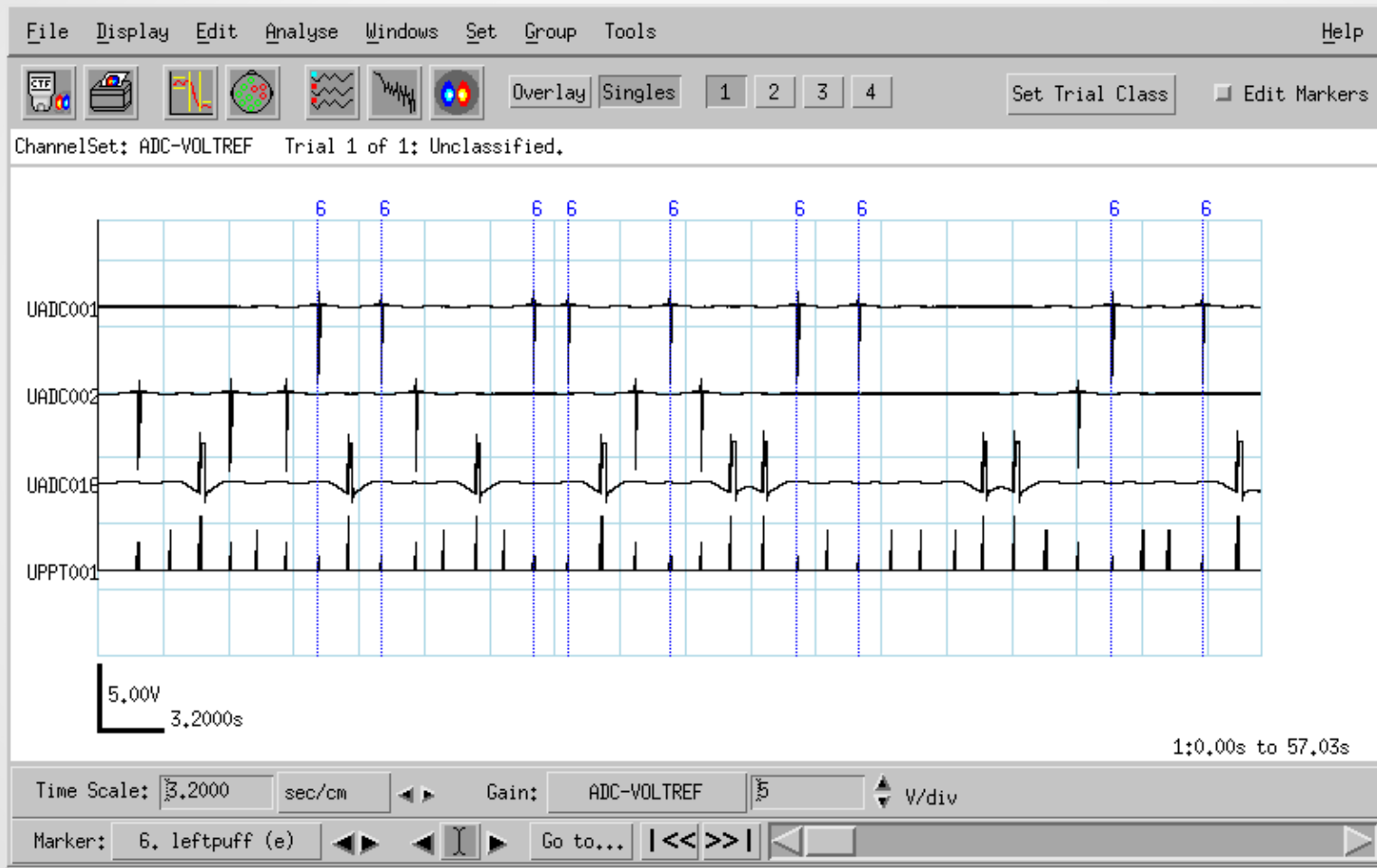
60 Hz line-noise notch filters  
including harmonics

Baseline removal

# Visually Check the Markers



# Visually Check the Markers





# doLocalSpheres.sh

```
#!/bin/sh
```

```
if [ "$#" -ne 1 ]; then
```

```
    echo usage: $0 dataset
```

```
    exit
```

```
fi
```

```
ds="$1"
```

```
hashcode=`basename $ds | cut -d_ -f1`
```

```
localSpheres -s mri/${hashcode}/multisphere.shape -d $ds -r 9
```

The `multisphere.shape` file is the output of `orthohull.localSpheres` uses that to create the final MultiSphere model used for magnetic field calculations. It is stored inside the MEG dataset folder.

# doSam.sh

```
#!/bin/sh

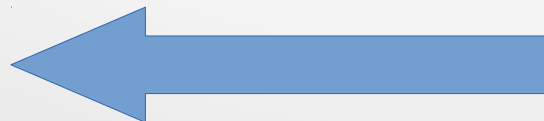
dosam() {
  ds="$1"
  m="$2"

  sam_cov -r $ds -m $m
  sam_wts -r $ds -m $m
  sam_4d -r $ds -m $m
}

for dsname in `cat dslist`; do
  dirname=`dirname $dsname`           # directory
  setname=`basename $dsname .ds`     # setname without .ds
  ds=${dirname}/${setname}-f.ds      # full pathname

  dosam $ds sef
done
```

Usually we use a script like this to run SAM. First `sam_cov` creates the covariance matrices (which are stored in the dataset's SAM subdirectory). Then, `sam_wts` creates the beamformers for the ROI. Finally, other programs such as `sam_3d` or `sam_4d` use the weights to create 3d (or 3d+time) volumes.



All SAM analysis is controlled by a *parameter* file (specified with `-m`).

# SAM parameter files

```
%include ROI                # standard ROI

NumMarkers 2                 # dataset markers
Marker1 rightpuff -.2 .5 true
Marker2 leftpuff -.2 .5 true

OrientBand 15 30            # beta is recommended because of its stability
CovBand 15 30                # the covariance band can be larger
ImageBand 15 30              # than the image band, for stability
SmoothBand 0 20              # for sam_4d
TimeStep .01

Model MultiSphere           # forward model
Mu 2                         # optional regularization (stability)

MRIDirectory mri            # input and
ImageDir image              # output dirs

ImageMetric Power           # what sort of image to compute
CovType SUM                 # which covariance to use for the image
```

# Running AFNI – .afnirc

```
***ENVIRONMENT
```

```
AFNI_LEFT_IS_LEFT = YES
```

```
AFNI_ALWAYS_LOCK = YES
```

```
AFNI_ORIENT = PRI
```

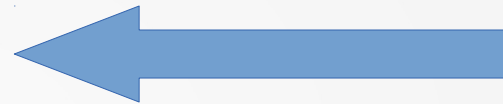
```
AFNI_DATASET_BROWSE = YES
```

```
AFNI_DECONFLICT = OVERWRITE
```

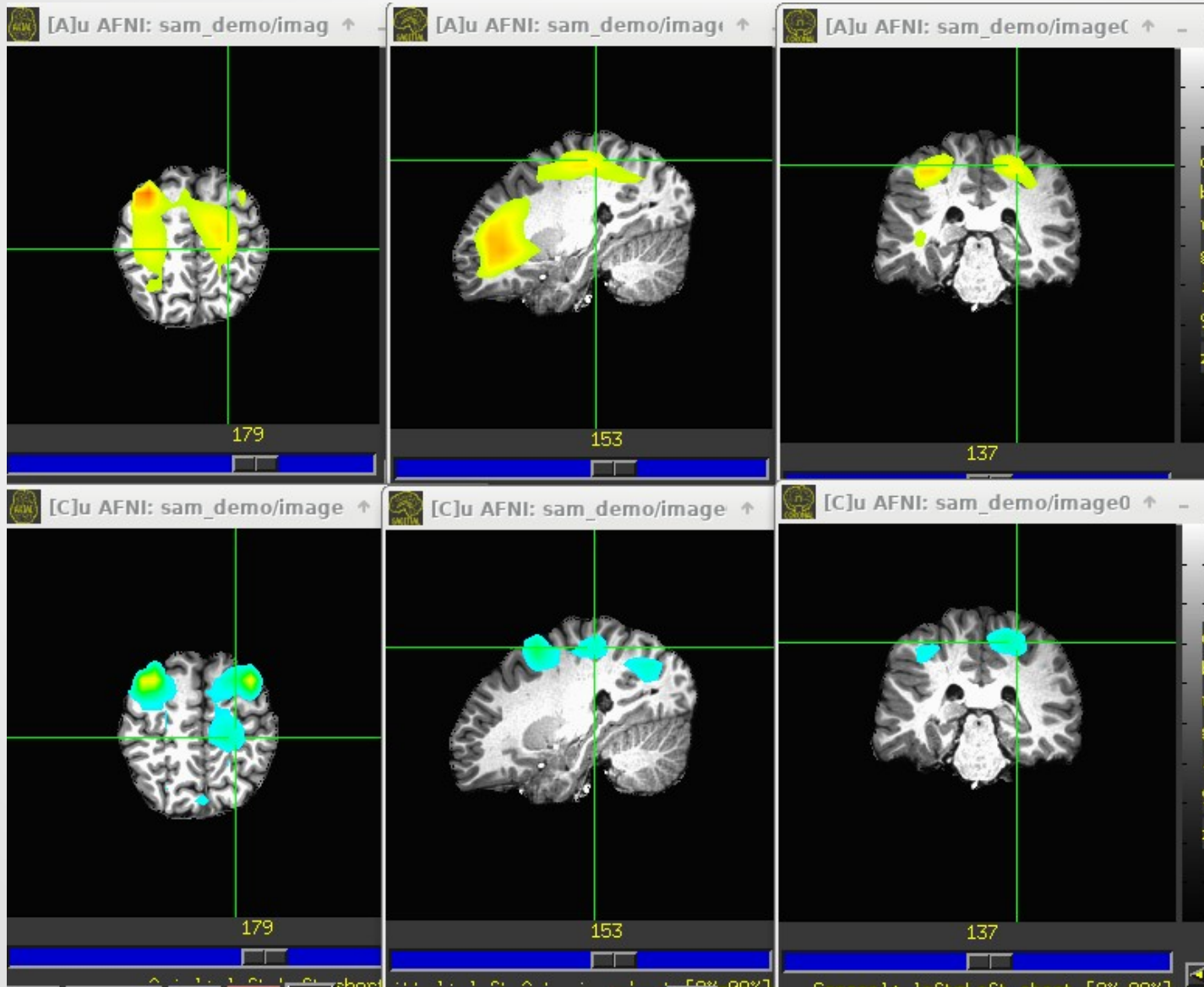
```
AFNI_NIFTI_VIEW = orig
```

```
AFNI_SLAVE_THROLAY = YES
```

CTF ordering is PRI



# sam\_3d – Mean and Variance



# sam\_4d – 3d+time Images and Graph

