

# EEG/MEG 1:

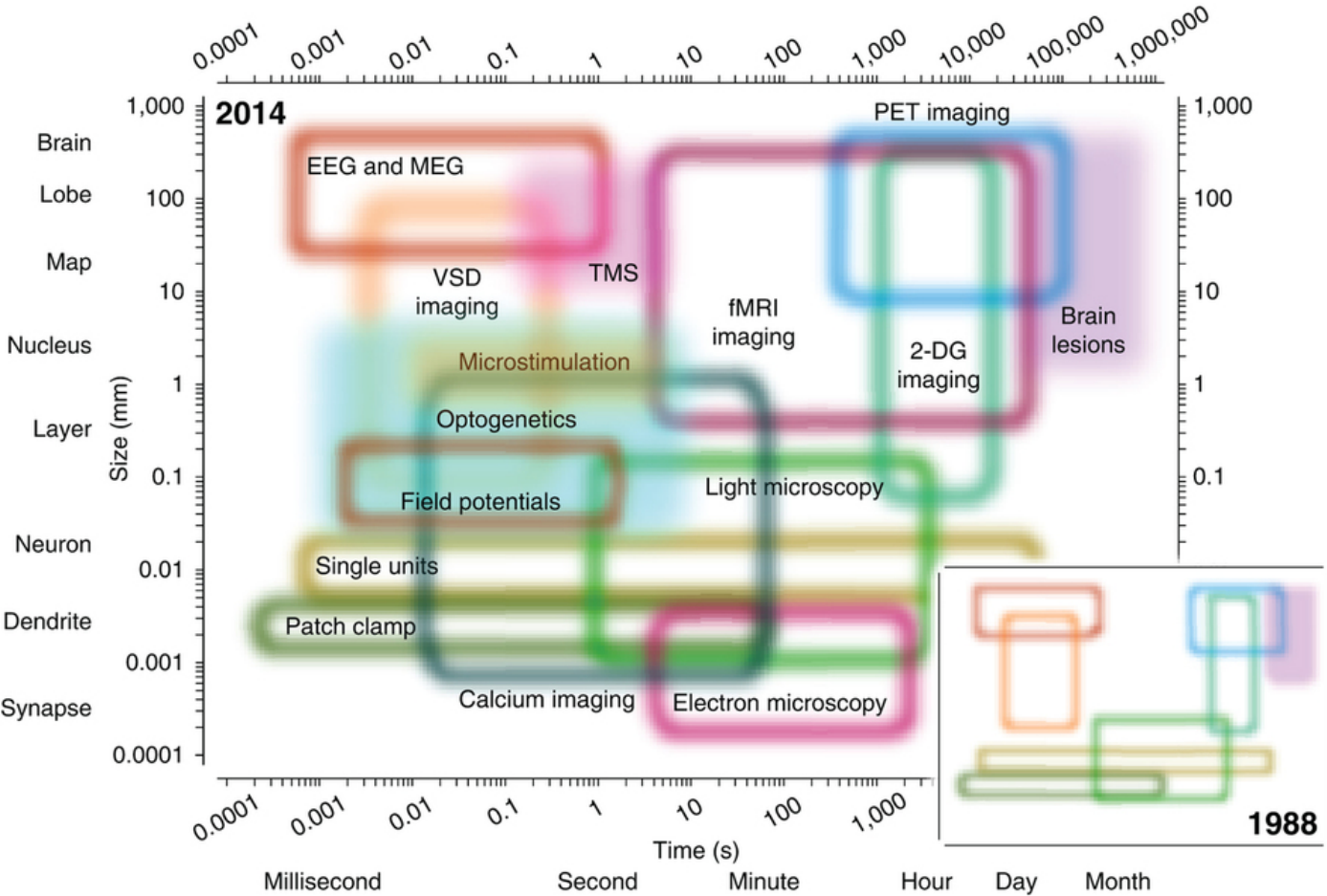
## Measurement, Pre-Processing and Data Reviewing

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Introduction to Neuroimaging Methods, 19.2.2018

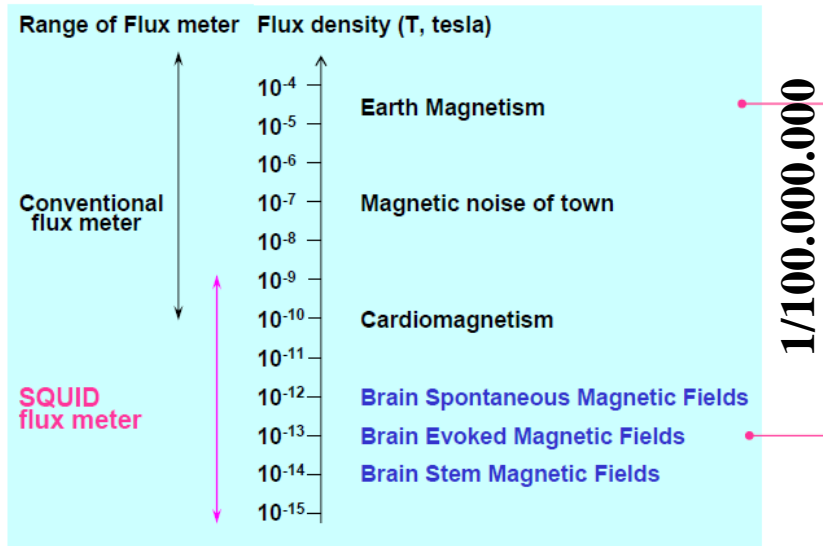
# A Big Picture: Spatial vs Temporal Resolution



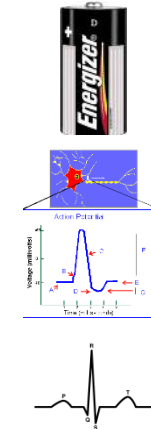
Sejnowski, Churchland, Movshon, Nat Nsc 2014

# What We are Measuring

## Magnetoencephalography (MEG)



## Electroencephalography (EEG)

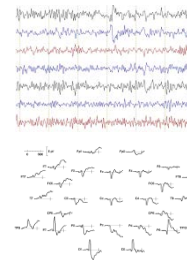


Household Batteries  
~ 1-12 V

Cell Membrane Potentials  
~ 70 mV

ECG:  
~ 1mV

Raw EEG: ~ 30  $\mu$ V  
Eye blinks: > 100  $\mu$ V

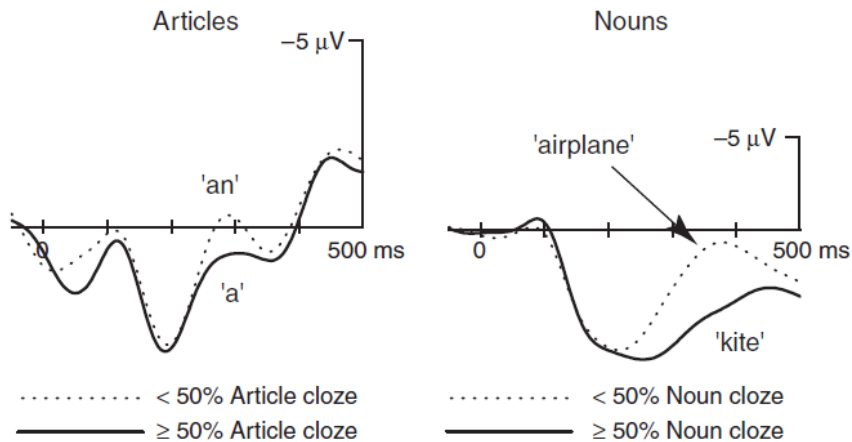


ERPs: ~ 0-10  $\mu$ V

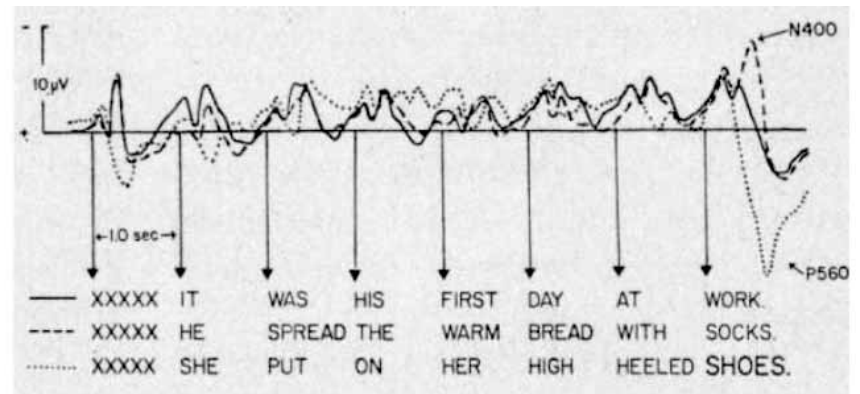


# When Timing Is Of The Essence

Vertex ERPs by median split on cloze probability,  
*e.g.*, 'The day was breezy so the boy went outside to fly ...'



deLong, Urbach, Kutas, Nat Nsc 2005



Kutas&Hillyard, Science 1980

# EEG/MEG Introductory Literature

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<http://imaging.mrc-cbu.cam.ac.uk/meg/MEGpapers>

## **Books:**

Hansen, Kringelbach, Salmelin: “MEG: An Introduction to Methods”, OUP 2010  
SJ Luck: “An Introduction to The Event-Related Potential Technique”, MIT 2005  
TC Handy: “Event-Related Potentials”, MIT 2004

## **Guidelines for MEG and EEG research:**

Gross et al., “Good practice for conducting and reporting MEG research.“, Neuroimage  
2013  
Picton et al., “Guidelines for using human event-related potentials to study cognition:  
recording standards and publication criteria.“, Psychophysiology 2000

# A Brief History Of Bioelectromagnetism

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## **Ancient Egypt, 2750 BC:**

Electric Fish (“Thunderer of the Nile”)  
Some Roman writers mention electric shocks as an ailment for headaches (~ 0 AC)...



## **Ancient Greece, 600 BC:**

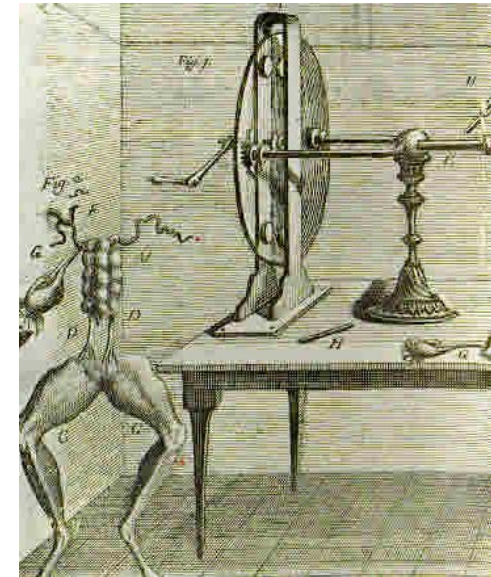
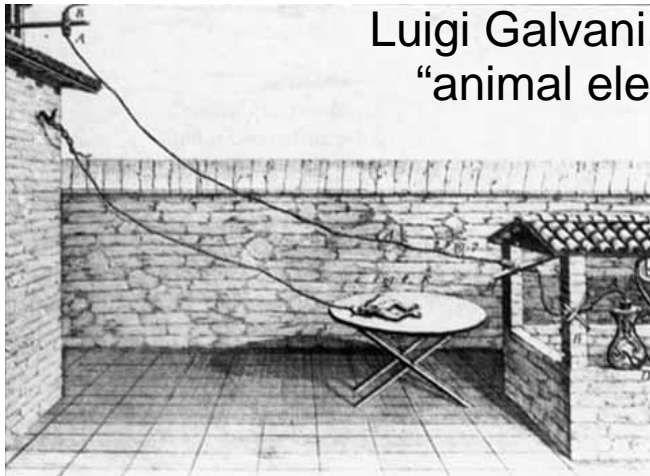
Thales describes static electricity  
“electron”



# Early Science

1771

Luigi Galvani, Bologna  
“animal electricity”



In 1803:

“On the first application of the process to the face, the jaws of the deceased criminal began to quiver, and the adjoining muscles were horribly contorted, and one eye was actually opened. ...

Mr Pass, the beadle of the Surgeons' Company, who was officially present during this experiment, was so alarmed that he died of fright soon after his return home.”

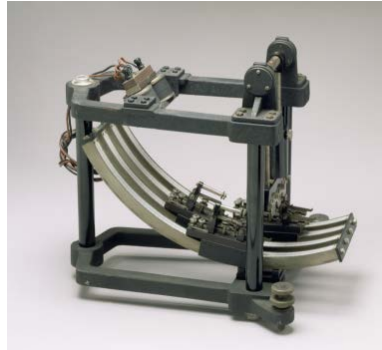
<http://www.executedtoday.com/2009/01/18/1803-george-foster-giovanni-aldini-galvanic-reanimation/>



# Early Electrophysiology

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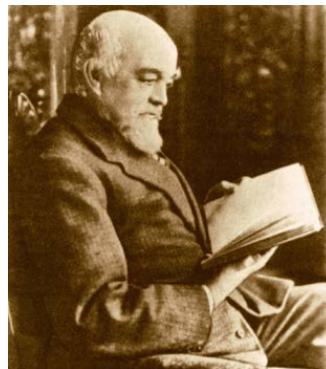
1842: Du Bois-Reymond, Berlin  
nerve action potentials neurons



1852: Helmholtz, Berlin  
speed of action potentials in frogs neurons

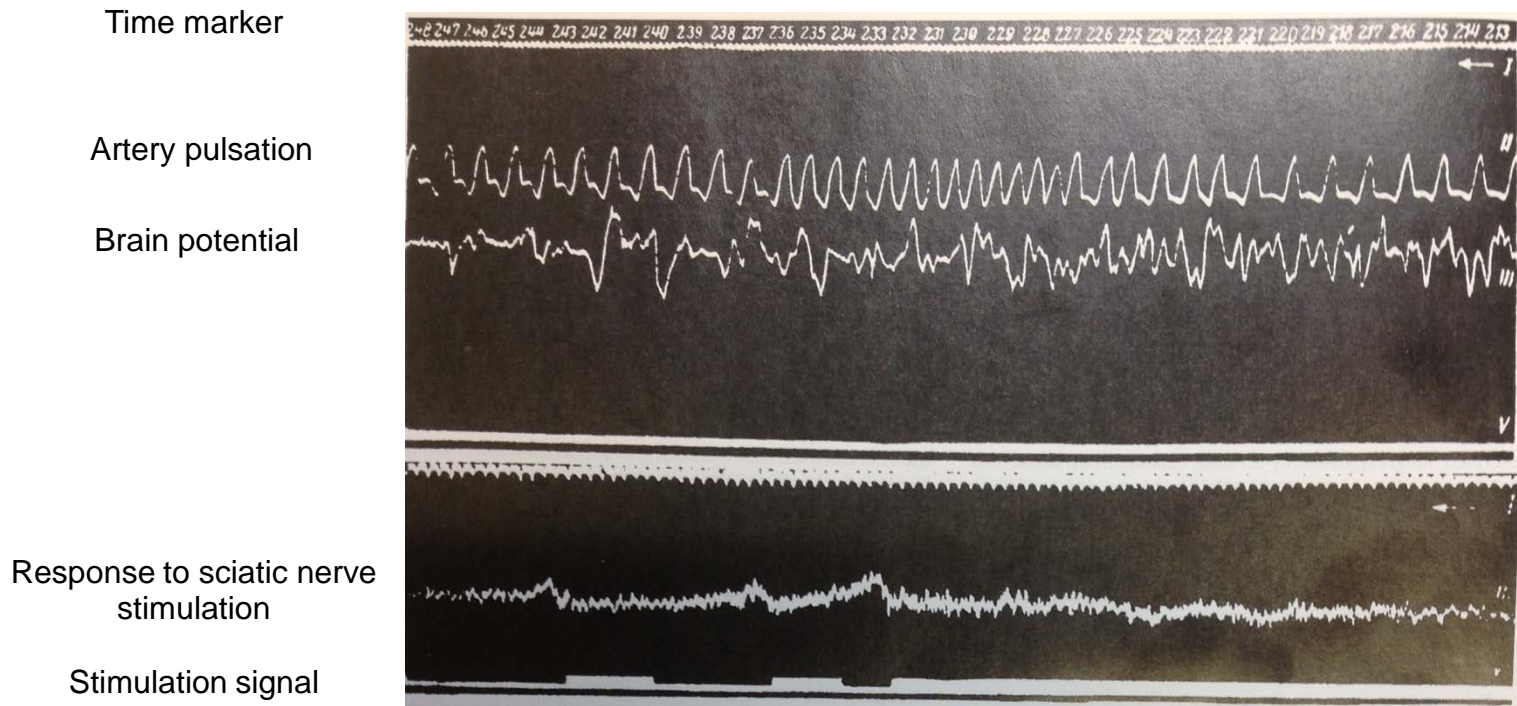


1875: Richard Caton, Liverpool  
first “ECoG” from animals



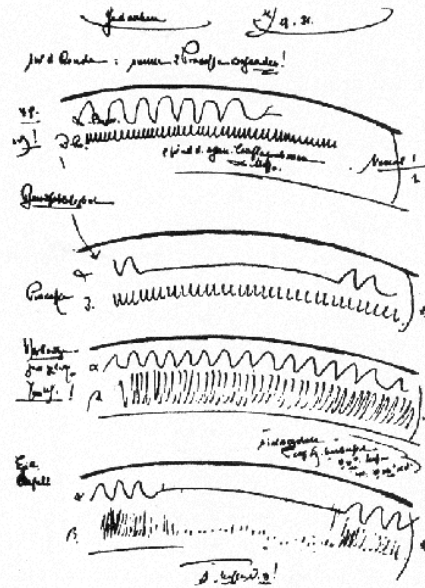


# Early EEG

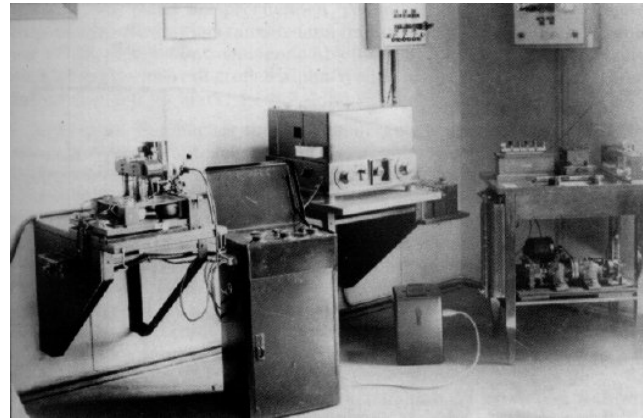


“Danilevsky (1852-1939) ... finished his thesis entitled “Investigations into the Physiology of the Brain (1877). ... He published an extensive textbook of human physiology in 1915. ... He saw his high hopes unfulfilled as far as the spontaneous electrical activity of the brain was concerned. ... He was not the only EEG researcher with shattered hopes in the field of psychophysiology”.

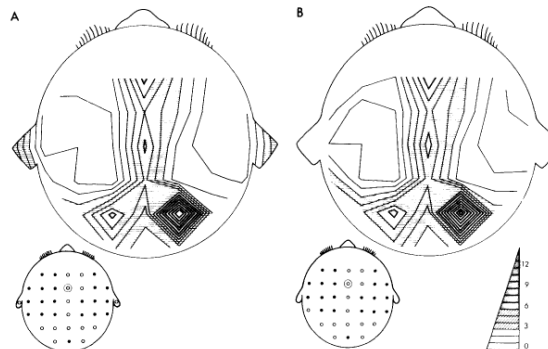
# Early EEG



Hans Berger, Jena 1924  
First Fourier Analysis of EEG: Berger&Dietsch 1931

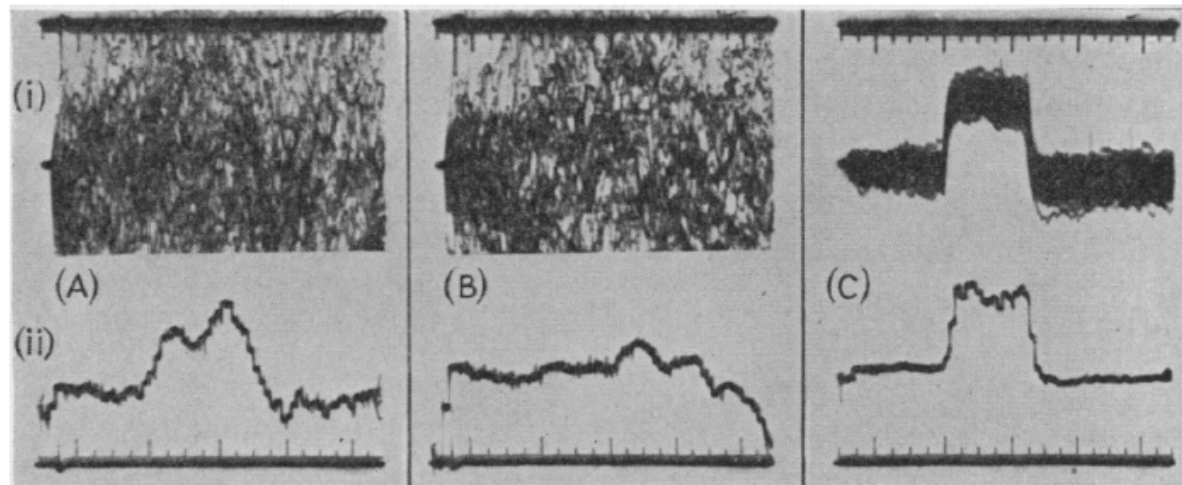


1969/70: 32/48-channel EEG, "generators"



# Early ERPs

**A summation technique for detecting small signals in a large irregular background.** By G. D. DAWSON. *Neurological Research Unit, Medical Research Council, National Hospital, Queen Square, London, W.C. 1*



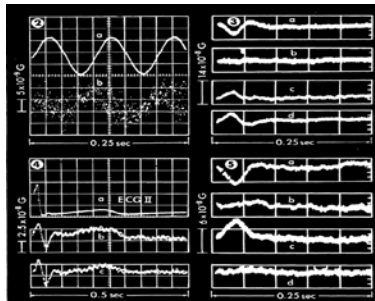
**Fig. 1.** An experiment to detect cerebral responses when the left ulnar nerve was stimulated at the wrist once per second. The upper line of traces shows sets of 55 records superimposed and the lower line the averages of these given by the machine. In A, from the contralateral scalp, there was one electrode on the midline and one over the right central sulcus. In B, from the ipsilateral scalp, the record was taken from the same midline electrode and one over the left central sulcus. In C is shown the result of making the electrode over the central sulcus positive to that on the midline by  $5 \mu\text{V}$ . The largest spikes in the time scales show intervals of 20 msec., and the stimulus was applied 5 msec. after the start of each sweep.

# First MEG: Pre-SQUID age

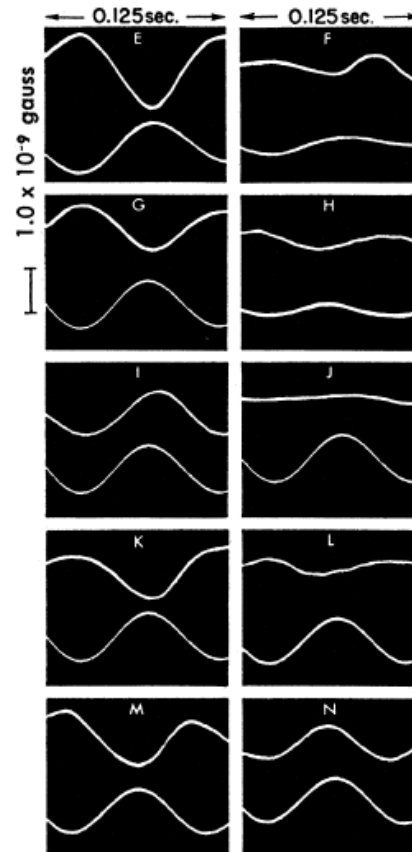
MEG pioneers MIT



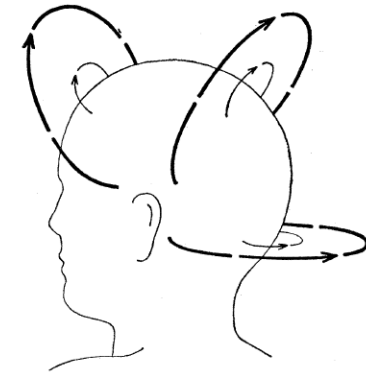
MCG, 1967/(63)



MEG, 1968



Alpha Rhythm



Cohen, Science 1968

# The Fast Evolution of MEG



1983  
by HUT  
4 channels  
30 mm in  
diameter  
(coverage:  
7 cm<sup>2</sup>)  
Axial



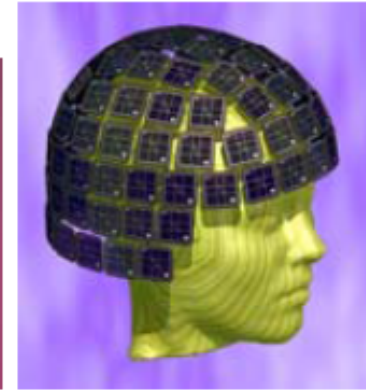
1986  
by HUT  
7  
channels  
93 mm in  
diameter  
(coverage:  
68 cm<sup>2</sup>)  
Axial



1989  
by HUT  
24 channels  
125 mm in  
diameter  
(coverage:  
123 cm<sup>2</sup>)  
Planar



1991  
by Neuromag  
122 channels  
whole head  
(coverage:  
1100 cm<sup>2</sup>)  
Planar  
12 Deliveries

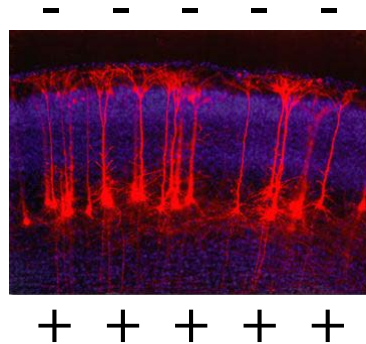
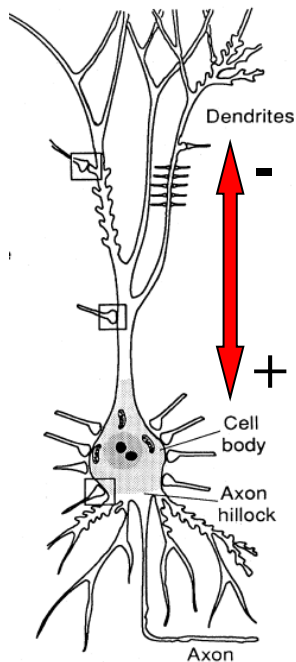


1997  
by Neuromag  
306 channels  
whole head  
(coverage:  
1220 cm<sup>2</sup>)  
Planar &  
Magnetometers



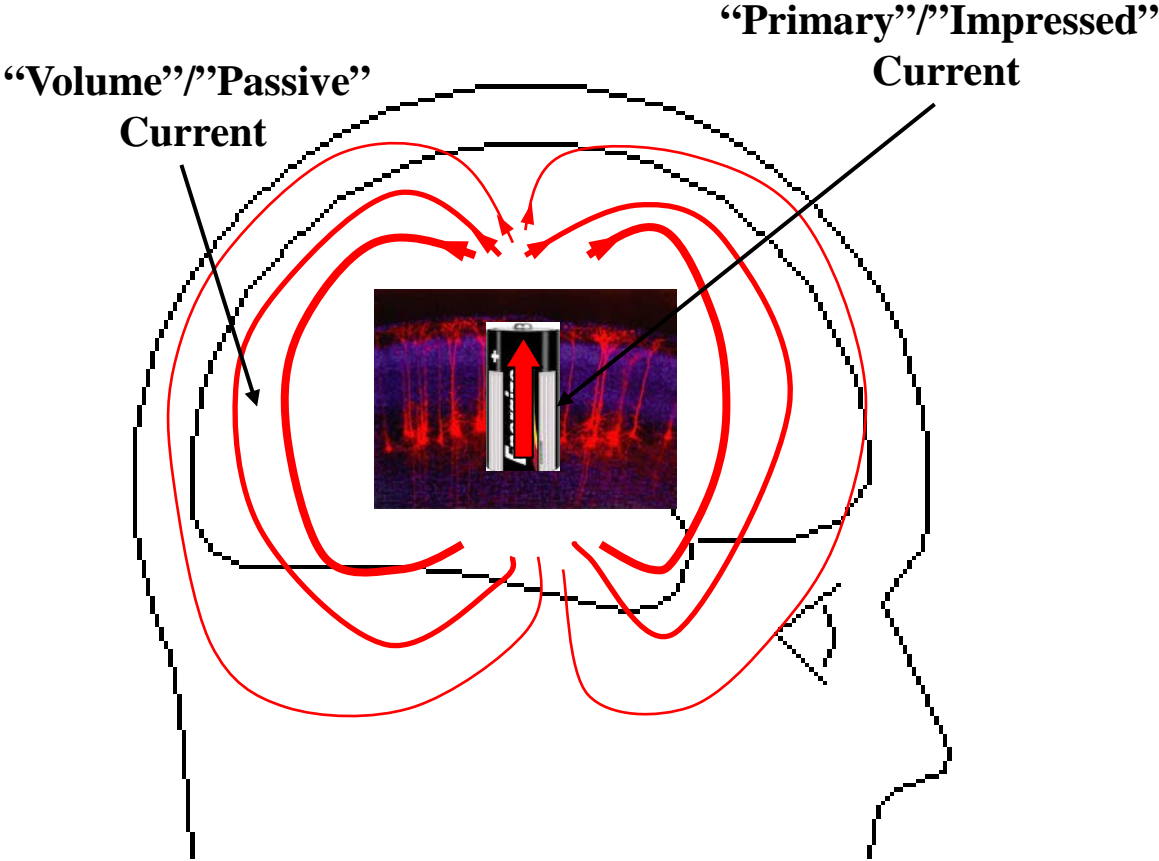
# Main Generators of Electrical Activity in the Brain

- Apical dendrites of pyramidal cells
- **NOT action potentials** (too short-lived and quadrupolar)
- **EEG/MEG: same generators, different sensitivity**



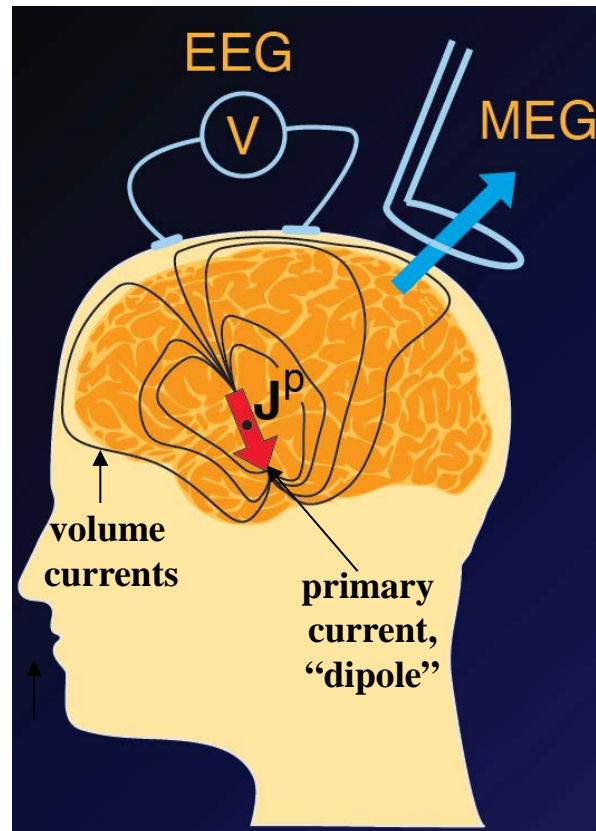
- ~ 1 Million synapses needed to activate simultaneously
- Luckily: ~10000 cells per  $\text{mm}^2$ , ~ 1000 synapses per cell  
=> several  $\text{mm}^2$  can produce measurable signal

# Current Flow in the Head





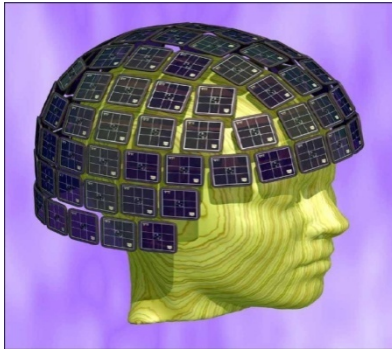
# EEG/MEG Measurements



Volume currents affect both EEG and MEG –  
but EEG more than MEG

# The Neuromag Vectorview System

306 channels in 102 locations



1 magnetometer and 2 planar gradiometers at each location

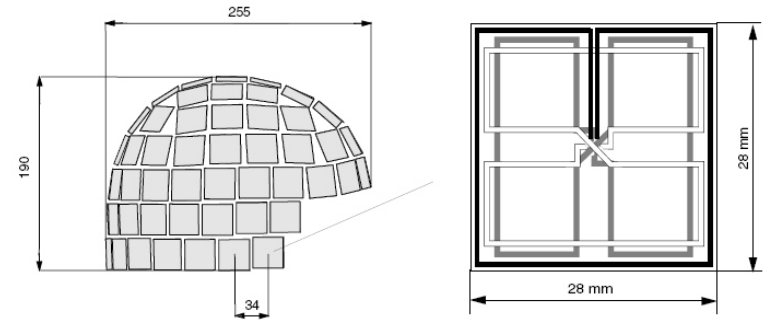
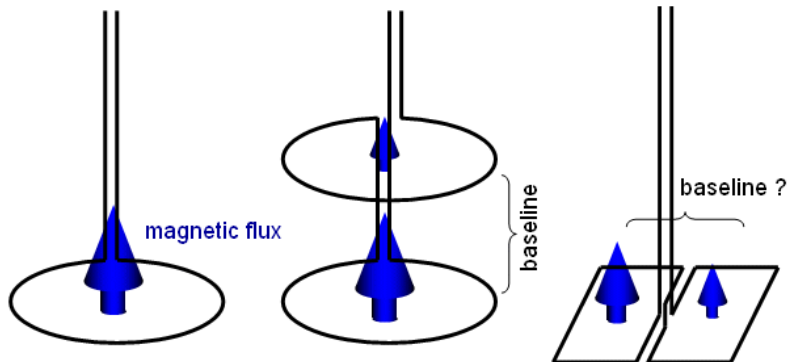


Figure 1.6. (left) Detector array, side view. Average distance between sensor elements : 34,6 mm. (right) Triple sensor detector unit.

## MEG sensor types

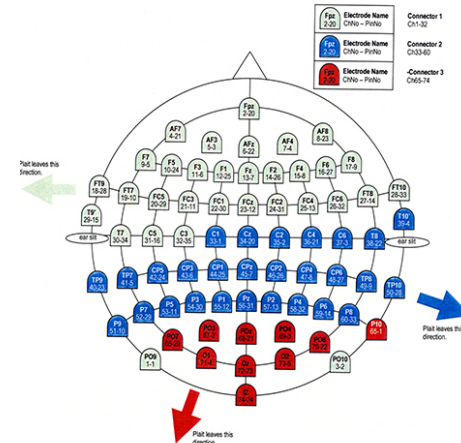
magnetometer    axial gradiometer    planar gradiometer



<http://meg.aalip.jp/scilab/CoilType.html>

MRC Cognition and Brain Sciences Unit

Up to 120 EEG electrodes  
(we typically use 70, plus EOG/ECG)



<http://imaging.mrc-cbu.cam.ac.uk/meg/VectorviewDescription>

# Leadfields

**Leadfields are sensitivity profiles of individual sensors.**

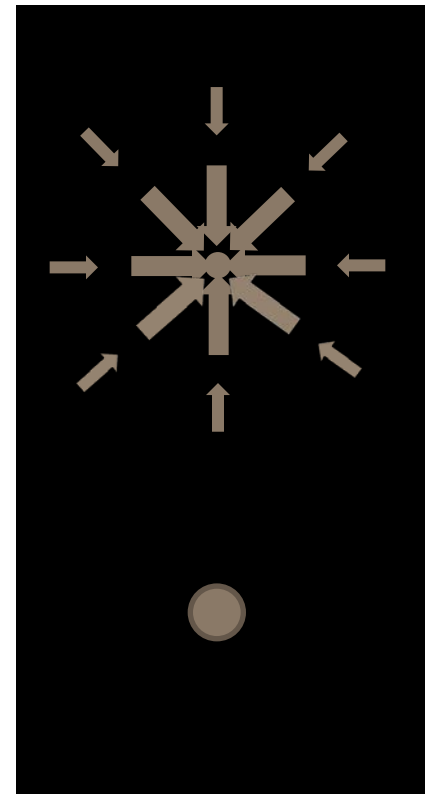
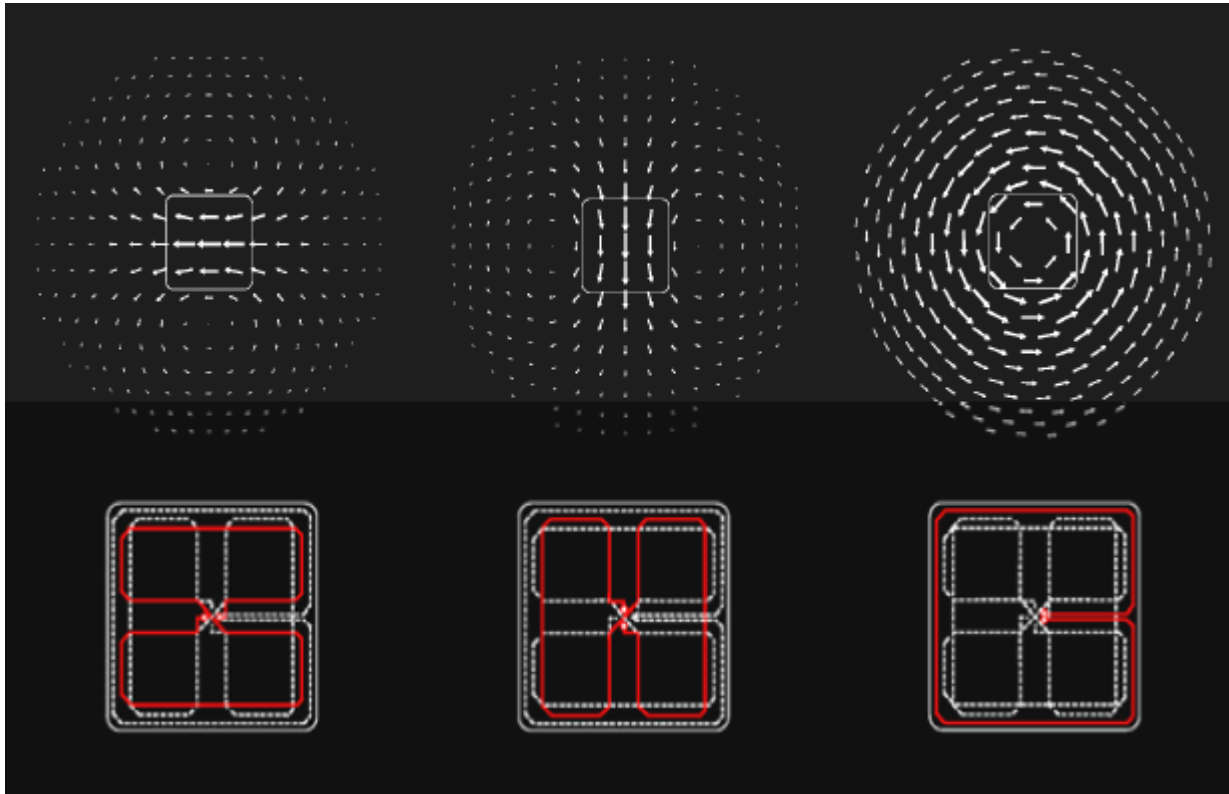
Each sensor is maximally sensitive to sources oriented along the arrows, and insensitive to sources perpendicular to the arrows.

**Gradiometer |**

**Gradiometer**

**Magnetometer**

**EEG**

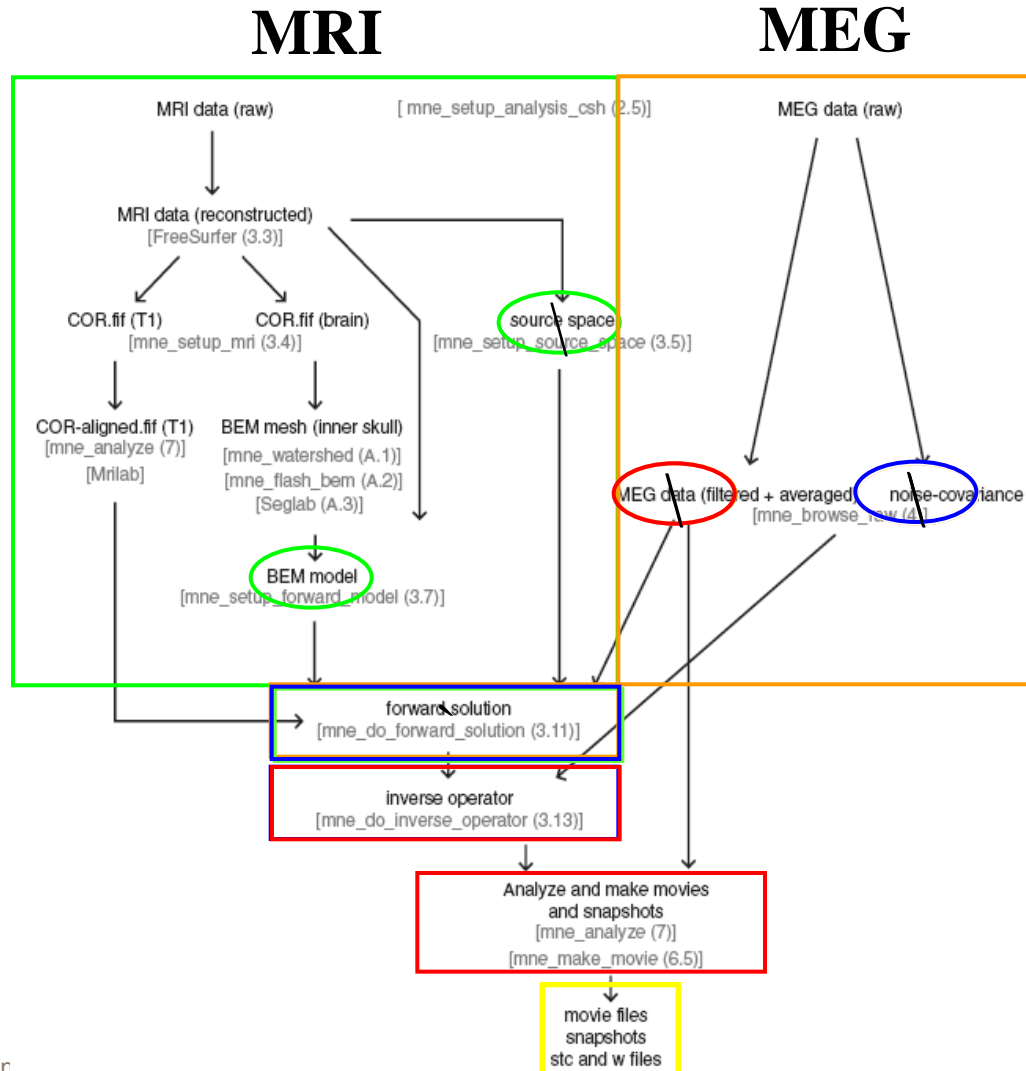


The “right-hand-rule” comes in handy here.

This bit I made up.



# Typical EEG/MEG Analysis Pipeline



# Artefacts

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## Artefacts can be

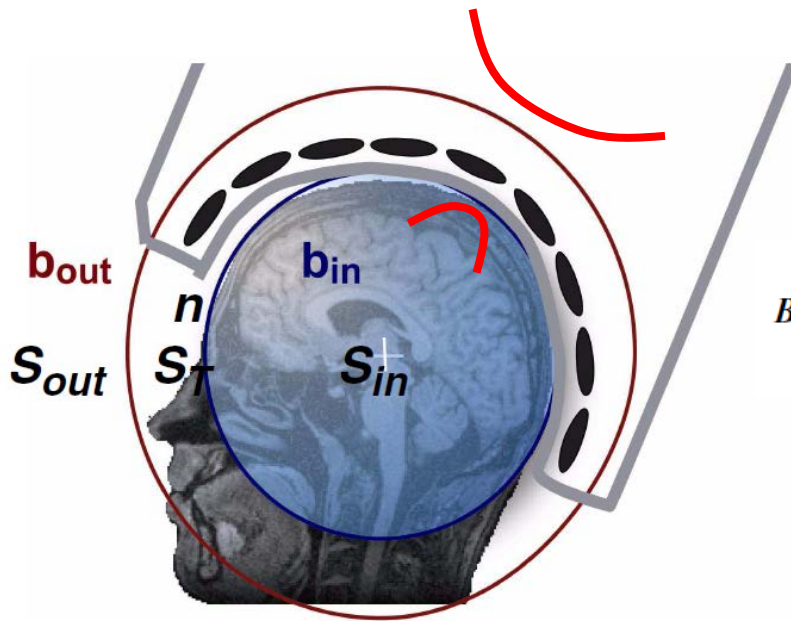
- **non-physiological**, i.e. from outside the body (sensor-intrinsic noise, line noise, moving objects, vibrations)  
=> Maxfilter (SSS), Frequency-Filtering, SSP, PCA/ICA
- **Physiological but non-brain**, e.g. eye movements, muscles  
=> SSP, PCA/ICA, H/L-Filtering
- **Physiological from the brain**, i.e. brain sources that are not of interest or not included in your source model  
=> choose appropriate source estimation, regularisation

### Wisdoms:

“Some people’s signal is other people’s noise.”

It’s always better to avoid artefacts than to correct them.

# Maxfilter



$$\mathbf{b} = \mathbf{b}_{in} + \mathbf{b}_{out} + \mathbf{n}$$

**Maxmagic** (spherical harmonics):

$$B(r) = -\mu_o \sum_{n=1}^{\infty} \sum_{m=-n}^n \alpha_{nm} \frac{v_{nm}(\theta, \varphi)}{r^{n+2}} - \mu_o \sum_{n=1}^{\infty} \sum_{m=-n}^n \beta_{nm} r^{n-1} \omega_{nm}(\theta, \varphi).$$

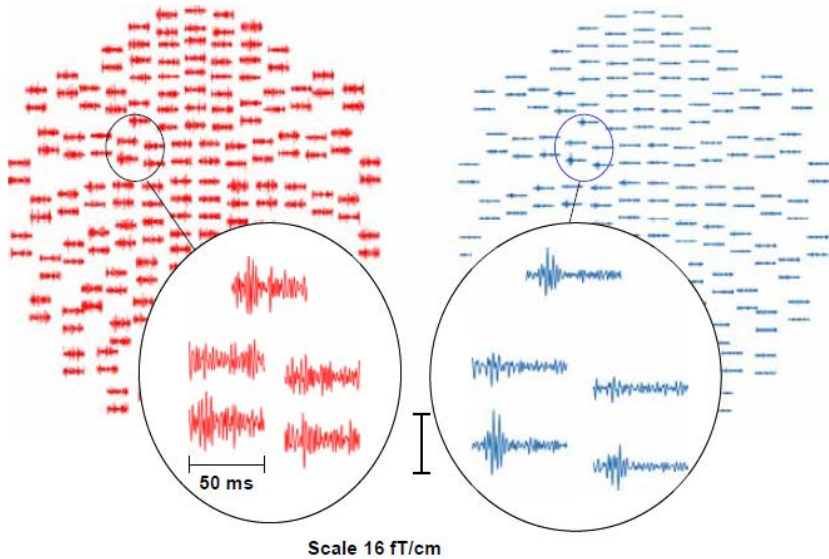
$$v_{nm}(\theta, \varphi) = -(n+1)Y_{nm}e_r + \frac{\partial Y_{nm}}{\partial \theta}e_{\theta} + \frac{imY_{nm}}{\sin \theta}e_{\varphi},$$

$$\omega_{nm}(\theta, \varphi) = nY_{nm}e_r + \frac{\partial Y_{nm}}{\partial \theta}e_{\theta} + \frac{imY_{nm}}{\sin \theta}e_{\varphi},$$

# Maxfilter

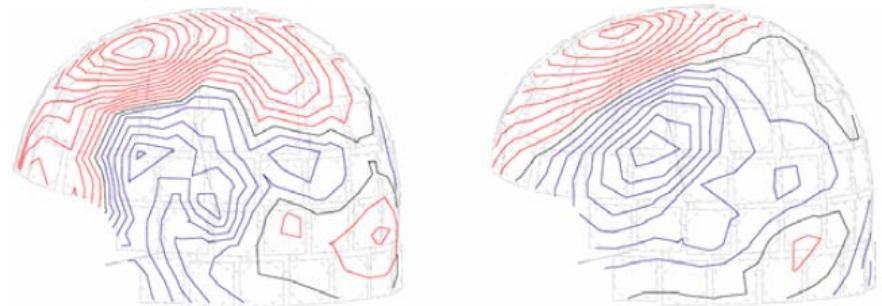
Without

With



Without

With



Original Field Map

SSS Reconstructed  
Field Map

Latency 20 ms  
Q = 2 nAm



# Maxfilter

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## **Software shielding (Signal Space Separation, SSS)**

By subtracting the outer SSS components from measured signals, the program suppresses artifacts from distance sources.

## **Automated detection of bad channels**

By comparing the reconstructed sum with measured signals, the program can automatically detect if there are MEG channels with bad data that need to be excluded from Maxwell-filtering.

## **Spatio-temporal suppression of artifacts (“-st”)**

By correlation the time courses of SSS artefact components with the cleaned signal, the program can identify and suppress further artefacts that arise close to the sensor array.

**Notch Filter** to remove 50Hz line noise

## **Transformation of MEG data between different head positions (“-trans”)**

By transforming the inner components into harmonic amplitudes (i.e. virtual channels), MEG signals in a different head position can be estimated easily.

## **Compensation of disturbances caused by head movements (“-movecomp”)**

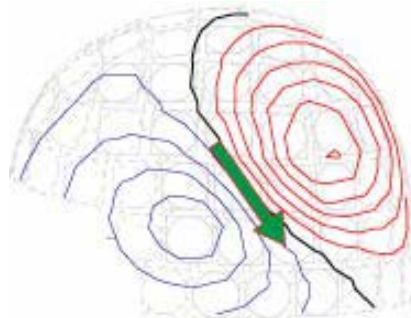
By extracting head position indicator (HPI) signals applied continuously during a measurement, the data transformation capability is utilized to estimate the corresponding MEG signals in a static reference head position.

# Maxfilter – Movement Compensation

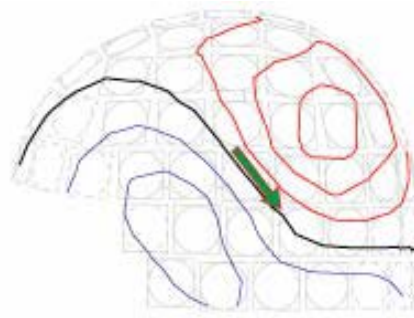
Head movement is tracked continuously (well, every 200 ms) via HPI (Head Position Indicator) coils.

We can take Maxfilter parameters from any time point  $t$ , and estimate the MEG signals at sensor positions of time point  $t_0$ .

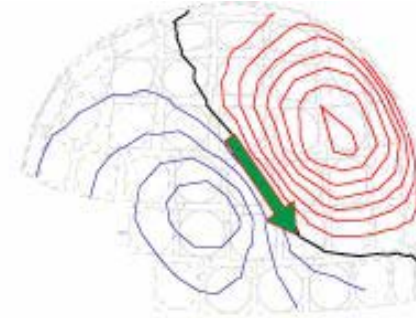
This compensates – to some degree – for spatial variation caused by head movements.



Stable subject



Moving subject,  
No compensation



Moving subject,  
with compensation



# Filtering and Downsampling

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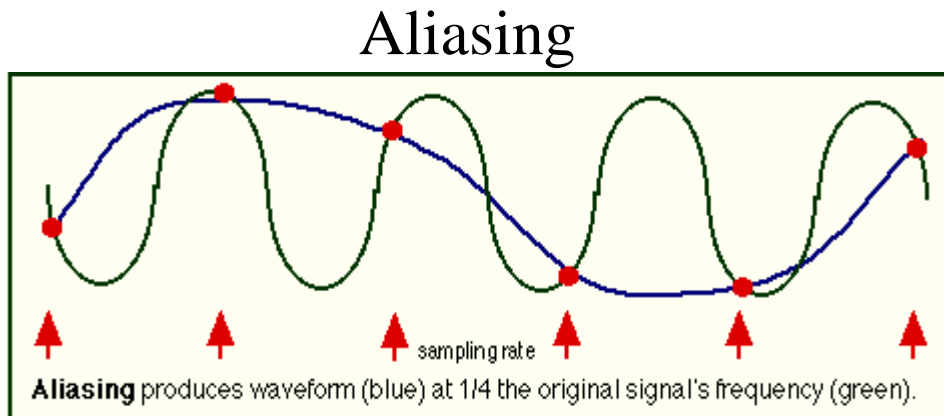
- Choose a “convenient” sampling rate with respect to processing speed and storage (usually 250 Hz to 500 Hz ok).
- We have to sample at 1000 Hz during acquisition because of head position indicator (HPI) signals.
- Downsampling can lead to “aliasing” if the data are not filtered appropriately (Nyquist theorem).
- Filtering can reduce (possibly remove) some artefacts such as sensor noise, muscle artefacts, line noise.

## Further reading:

Widmann et al., “Digital filter design for electrophysiological data – a practical approach”, Journal of Neuroscience Methods 2015.

# Aliasing

- Downsampling can lead to “aliasing” if the data are not filtered appropriately (Nyquist theorem)



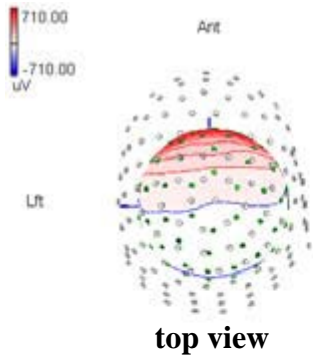
Watch:

<https://www.youtube.com/watch?v=R-IVw8OKjvQ>

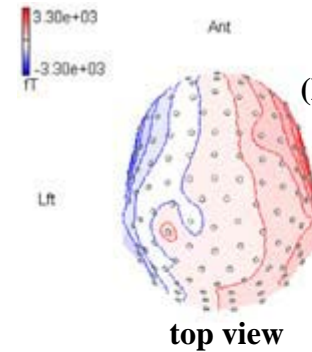
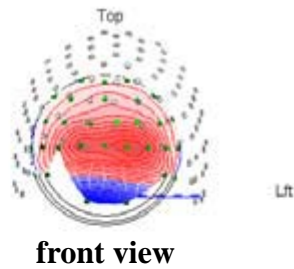
Thanks to Alessandro



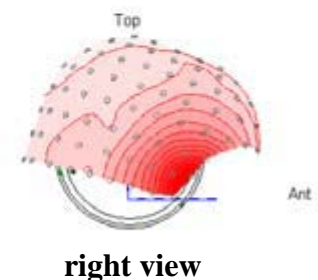
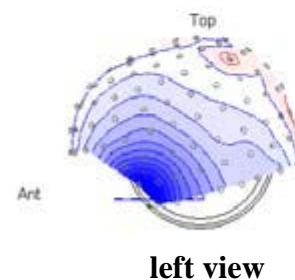
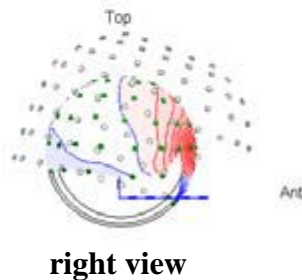
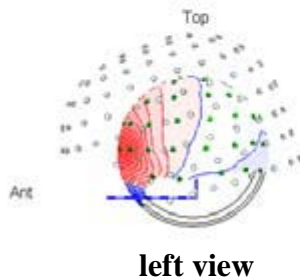
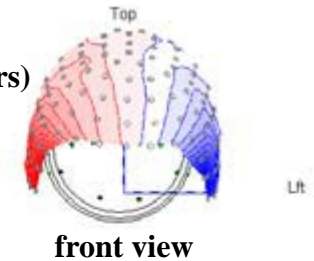
# Common Artefacts: Eye Blink



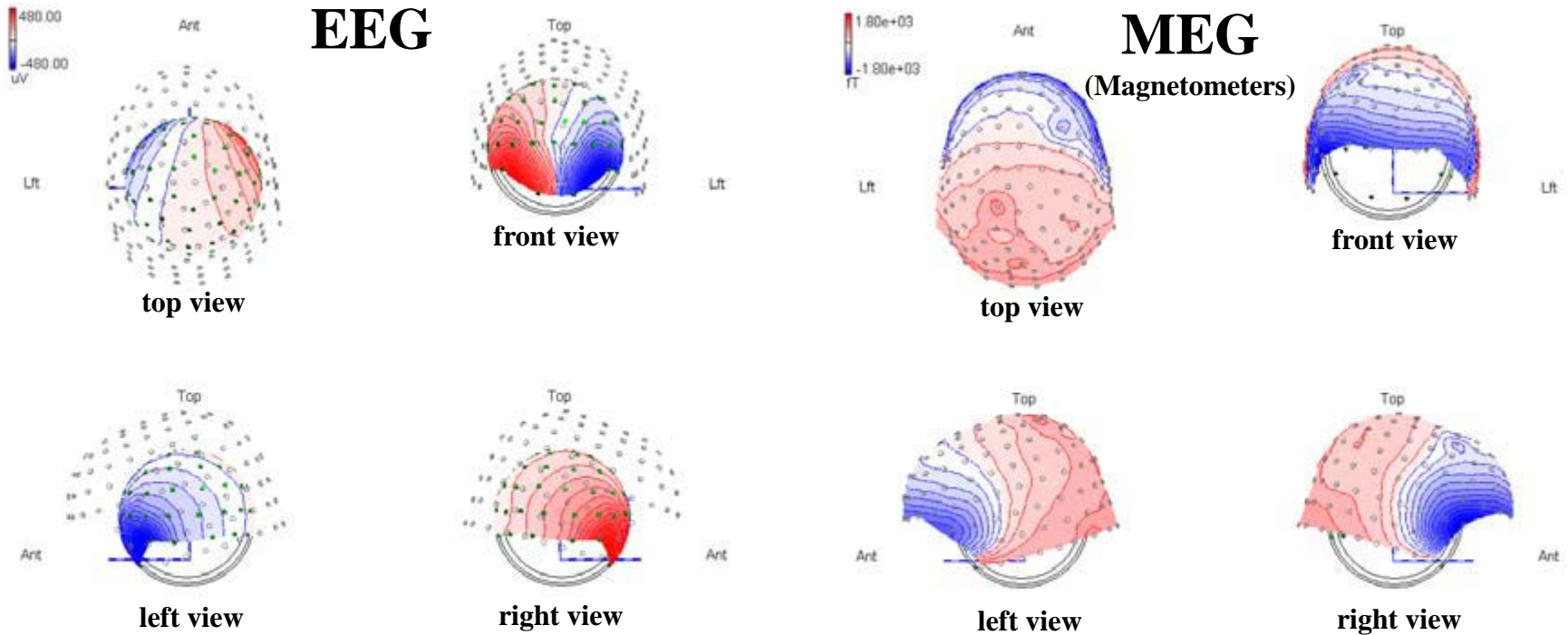
**EEG**



**MEG**  
(Magnetometers)

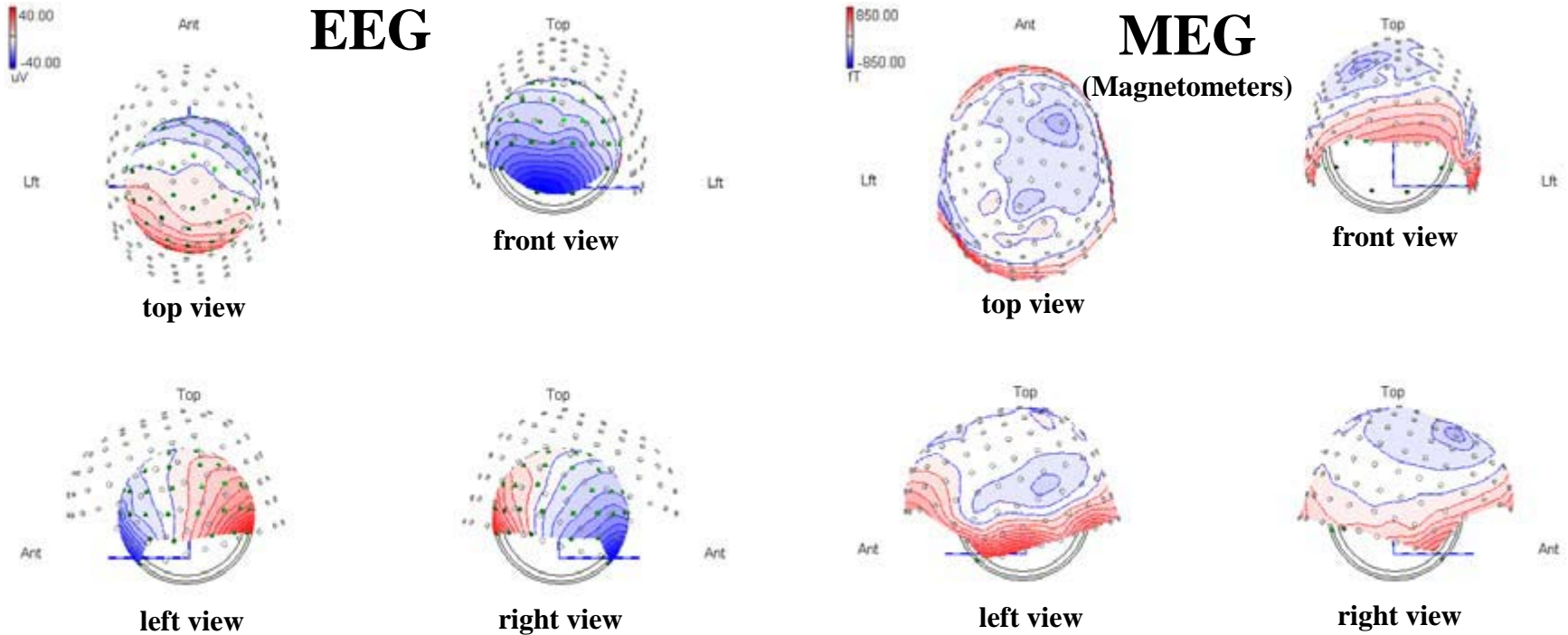


# Common Artefacts: Eye Movement to the Right





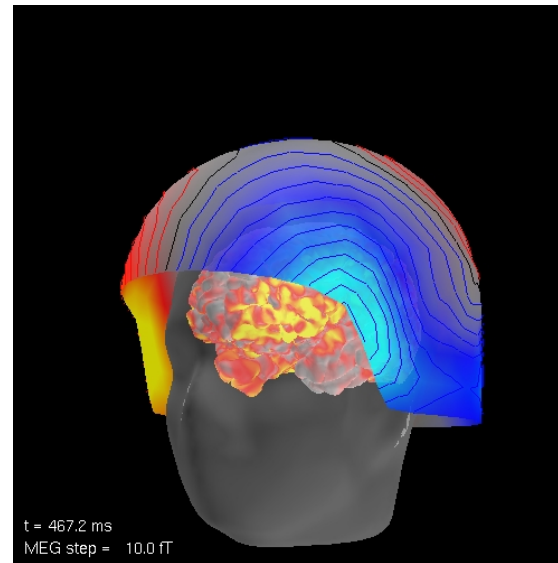
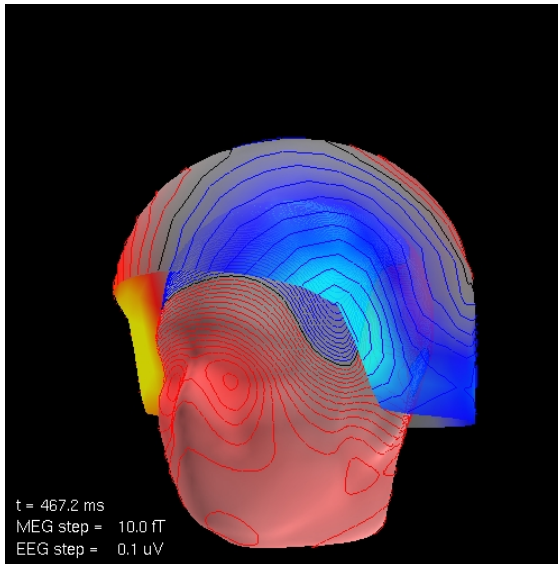
# Common Artefacts: Heart Beat



# Artefacts in EEG and MEG (Can) End Up in Source Space

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## Example: Eye Blink



This is a problem with all source estimation methods –  
get rid of your artefacts beforehand.

# Separating Signal and Noise Components

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If signal and noise have characteristic topographies, several methods can be applied to remove (some) noise or extract signals:

- SSP: Signal Space Separation

The following often go under the term “blind source separation”, because the topographies are not pre-defined, and found by the methods themselves (under certain assumptions):

- PCA: Principal Component Analysis
- SVD: Singular Value Decomposition

# Signal Space Projection (SSP)

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You know the noise topography **N**

You decompose your data **D**, such that

$$\mathbf{D} = \mathbf{a} * \mathbf{N} + \mathbf{Signal}$$

You only analyse **Signal**.

This works well with eye-movement and blink artefacts.

Note:

Brain signals whose topographies are highly correlated with **T** will also be removed or attenuated.

# PCA and SVD

---

- Decompose data into **orthogonal** components  $\mathbf{T}_1, \mathbf{T}_2$ , etc. (topographies or time courses), i.e. data

$$\mathbf{D} = a*\mathbf{T}_1 + b*\mathbf{T}_2 + \dots$$

- Find the components you don't like (e.g. correlate highly with EOG and ECG, or components that explain little variance).

- Reconstitute your data only with the “good” components,

e.g.  $\mathbf{D} = a*\mathbf{T}_1 + c*\mathbf{T}_3 + \dots$  if component 2 reflects eye blinks.

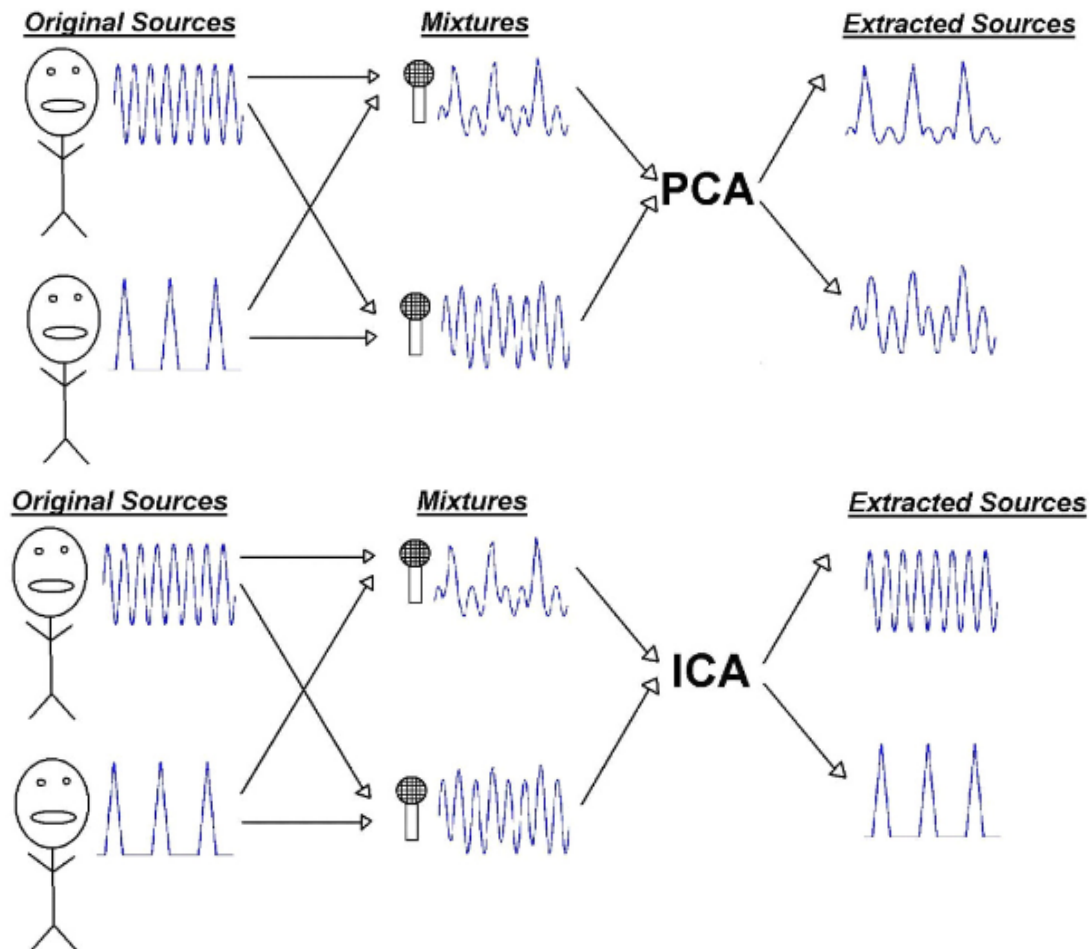
Also:

- Components have an order according to the variance they explain (e.g.  $\text{var}(\mathbf{T}_1) > \text{var}(\mathbf{T}_2) > \dots$ )
  - Can be used to determine the number of independent components (according to specified criteria)
    - Relatively fast (try `svd()` or `princomp()` in Matlab).

**•Unfortunately: Orthogonality and variance ordering not physiologically plausible.**

# Independent Component Analysis

Example: (De-)mixing of sources in the cocktail party effect



# Independent Component Analysis

---

Basic idea is similar to PCA and SVD:

Decompose data into components  $\mathbf{T}_1$ ,  $\mathbf{T}_2$ , etc. (topographies or time courses), i.e.

$$\text{data } \mathbf{D} = a * \mathbf{T}_1 + b * \mathbf{T}_2 + \dots$$

**But:**

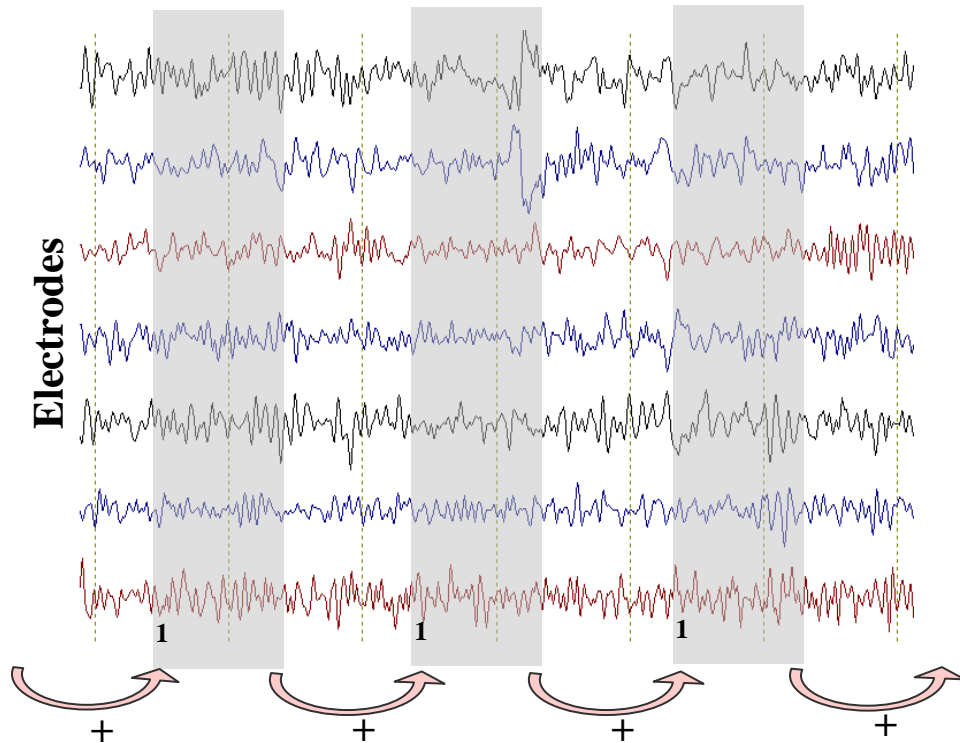
ICA does not produce orthogonal components,  
and does not assume Gaussianity of signals.



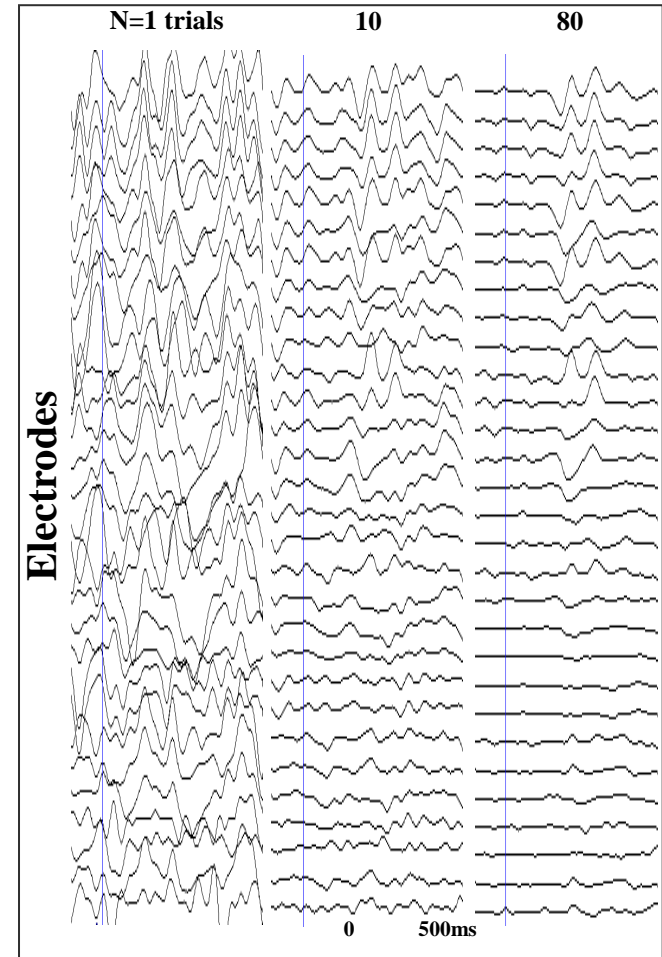


# Data Averaging

Continuous “raw” data:



Averaged data:



# Data Averaging

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The necessary number of trials depends on effect size, noise, variability across participants, your stats etc. –  
the more the better.

For random noise, variance goes down with  $n$ , and standard deviation with  $\sqrt{n}$ .

For “one-off” artefacts, amplitude in the average goes down with  $n$ .

“Robust Averaging” procedures exist (e.g. in SPM) that weigh epochs with an estimate of their reliability (e.g. distance to mean).

# Artefact Rejection

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Usually, epochs are excluded from averaging when they exceed some maximum-minimum criterion.

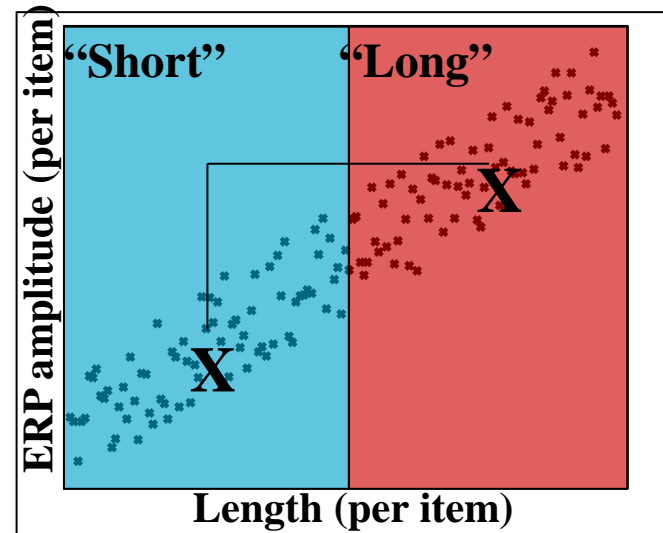
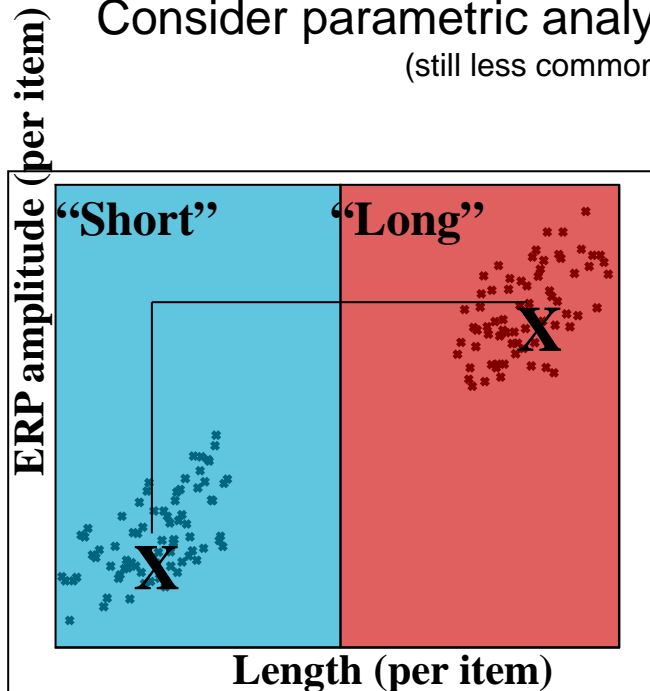
Make sure “chronically bad channels” are excluded from this procedure (or there won’t be any data left to average).

Prior to any procedure that combines signals across channels, such as average reference, SSP or ICA, bad channels should be removed (or signals from bad channels may be projected into the good ones).

Appropriate filtering and artefact correction (e.g. ICA) should be applied beforehand (but don’t feel too safe: artefacts may slip through).

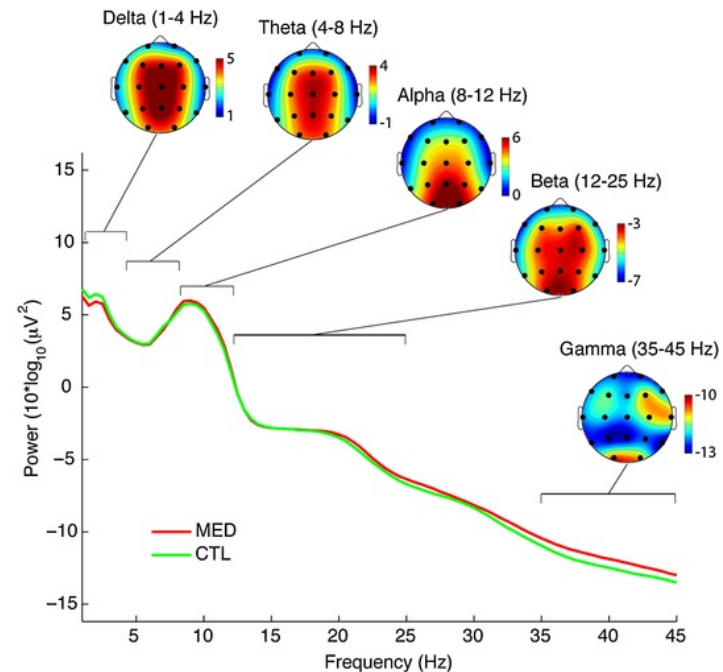
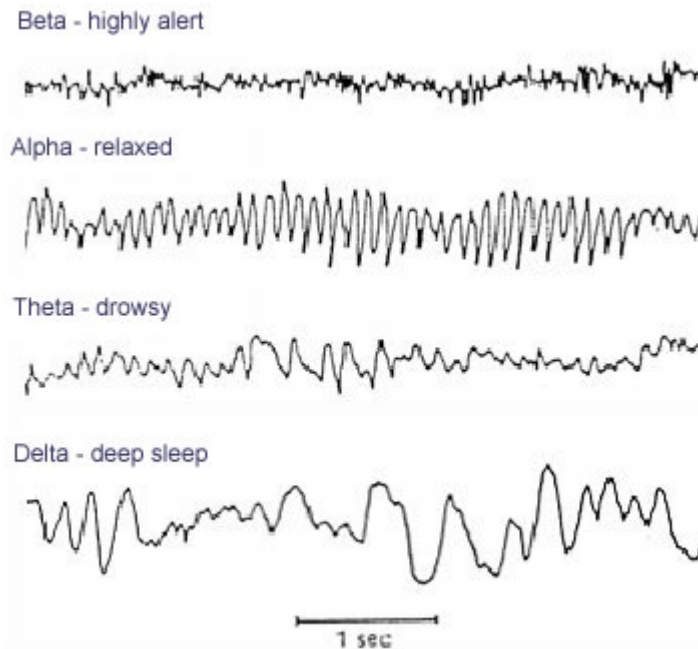
# Parametric vs Factorial Designs

Consider parametric analysis if stimulus variables are continuous  
(still less common in EEG/MEG than in fMRI analysis)



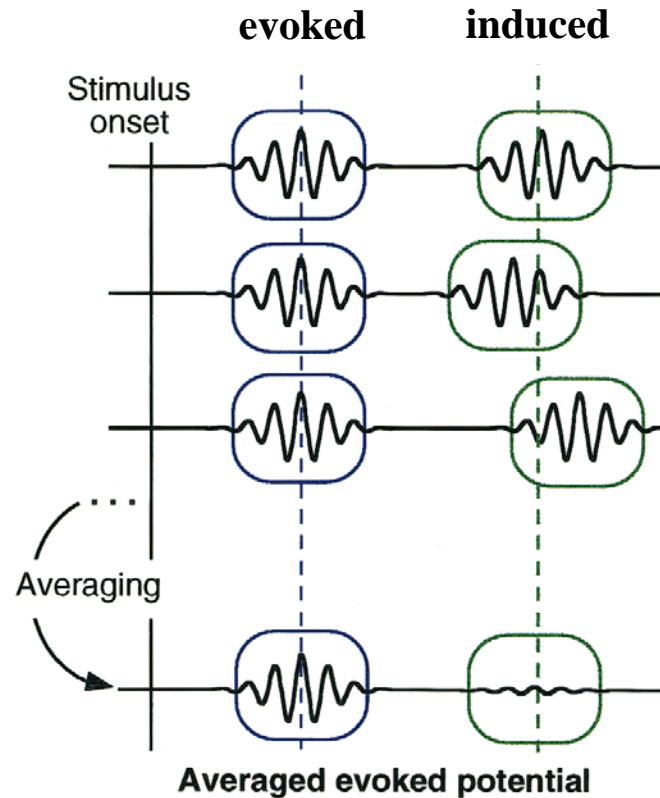
# “Brain Rhythms” and “Oscillations”

**Time course and topography may differ  
among different frequency bands  
(and may depend on task, environment, subject group etc.)**



<http://link.springer.com/article/10.1007%2Fs10339-009-0352-1/>

# Evoked and Induced Activity



Tallon-Baudry & Bertrand, TICS 1999

# The End Of #1

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