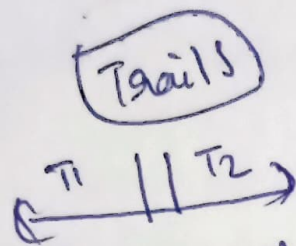


Quiz-1

- Lecture slides
- Notes
- Highlighted points in TB
- Any previous year papers.
- Understanding Matlab Code.
(whatever is shared).
- Read ch-7, 8, 9 & write
Notes

Rolep

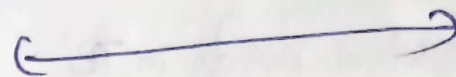


timing b/w
the end of
one trail &
start of next
trail

⇒ need of Base line
period.

⇒ for TFD,
Baseline should be
before trail onset
eg: -500 to -200ms

Typical Inter-trail interval
↓
at least 1000ms = 1sec



① Trail

↓
timing b/w events
within a
trial

1. Pilot test
2. Trail onset
3. alpha power/
activity
4. Edge Artifacts
5. Avg, Bipolar
Reference

Fixed ⇒ participants know
exactly when next
trial starts
variable
↳ can't predict
exactly

→ Brain Responses linger

→ Baseline in TFA should end before stimulus, not at
stimulus onset

→

① Preprocessing

P → Q
 R → X
 Q → R
 S → Q

only deleting
bad segments/
electrodes

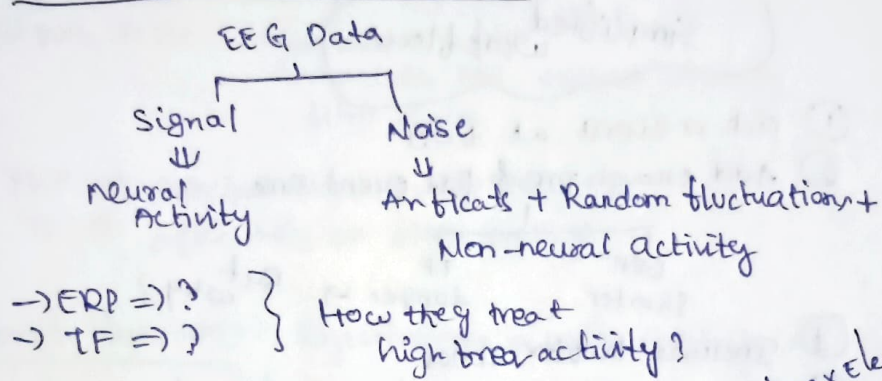
- organizing data — cutting continuous data into epochs
- Removing bad data — rejecting artifact trials
- Transforming clean data — changes data itself
eg. filtering, re-referencing, ICA, spatial filters.

no change to signal

→ Avoid introducing biases
↓
do same preprocessing steps.

② Balance b/w signal & Noise

$U \times G = 100$
 $U \rightarrow 10$
 $G \rightarrow 10$

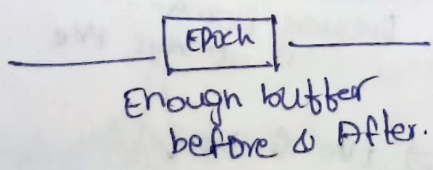


EEG → time × electrodes (2D)

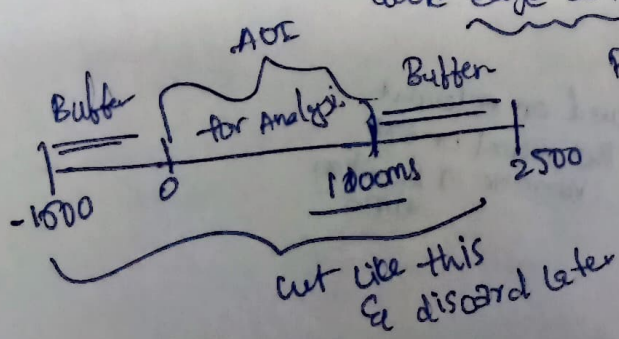
③ creating Epochs :

↓
For studying brain activity around specific events (stimulus, response...)

⇒ cut into epochs
EEG: electrodes × time × trials (3D)



ERP → relatively short buffer (-200ms → 800ms)
TF → need longer epochs with buffer zones to avoid edge artifacts



Rule of thumb

↓
they last about 3 cycles of FOI

Special cases → overlapping epochs? Happens when trials were close
 Problematic for ICA.
 ↓
 Shouldn't see same time points

→ If epochs are too short? (Already cut)

1) use Reflection Method → flip data Before vs After
 create Artificial buffer → Analyze & trim off

→ Tapering out → fading in/out
 → distorts baseline & power estimates

Simplified workflow

① pick a event at $t=0$

② Add enough pre & post event time

ERP Shorter TF longer → But why?

③ Include buffer zones

④ Discard buffer zones before Analysis (plotting)

④ Unequal Trail count = Biased Results? Yes (said in class)

Analysis type

① Phase Based → more sensitive → small No. of trials → tve Bias

② Power Based $\xrightarrow{\text{Noise will increase}}$ Because power is always tve

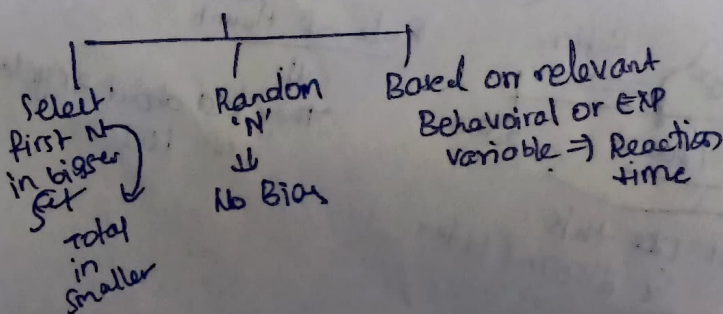
③ ERP → Not much Bias

Voltages → tve & -ve ✓

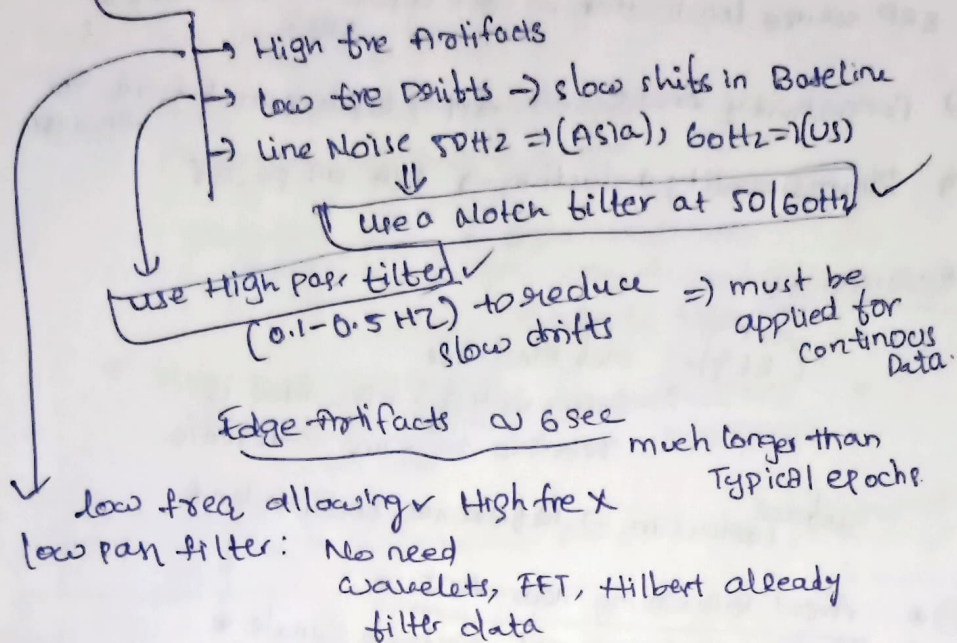
But Effects "SNR"

Soln:

Match trail count



⑤ Filtering: helps remove noise from EEG Data



1) ERP ⇒ careful Bandpass filtering

2) TF ⇒ often rely on other methods

⑥ Trial Rejection: Rejecting trials with Artifacts ✓

Approaches

Manual

- visual inspection
(Better) ⇒ Recommended
Sia said in cls
- Time consuming

Automatic

- Algo / Tool box / An based
- Risk of FP ⇒ May reject too much
- Diff thresholds
↓
diff results

sharp in data ⇒ less prominent in ERP ⇒ smoothing while averaging

⑦ Spatial Filtering:

Localization of sources

- which part of Brain?

separating overlapping brain processes

- using surface laplacian

Preprocessing for connectivity analyses

- same signal appearing at multiple electrodes

- when to apply spatial filters?

i) ERP source localisation \rightarrow fit dipole on grand averaged ERP's

ii) Connectivity Analyses \rightarrow Apply Laplacian before TF Analyses

iii) Dimensionality reduction \rightarrow PCA on power

⑧ Referencing:

EEG \rightarrow ref electrode
 \rightarrow every voltage recorded is relative to some reference.

Surface Laplacian \rightarrow Reference Independent

* Avoid Referencing near AOI *

* Always clean your ref electrode signals *

* Using one of the scalp electrode as ref is suboptimal *

Electrode
 \rightarrow Bipolar Ref \rightarrow Measures diff b/w 2 electrodes directly (EEG or EcG)
 \rightarrow Avg reference (all electrodes) > 100
 \rightarrow Mastoids (behind ears) \rightarrow picks up less Brain activity

⑨ Interpolation of Bad Electrodes:

Process by which data from missing electrodes are estimated based on activity & locations of other electrodes

Determine whether a noisy electrode?

\Downarrow

Apply a low pass filter to data at 30Hz

\Downarrow

Compare the activity with surrounding electrodes

Ch 8

Independent Component Analysis

→ ICA for Artifact Removal:

- source separation technique
- preprocessing, data reduction
 - ↓
 - subtract them
- Analyze component time series instead of electrode time series

* Max no. of components that can be isolated in EEG data = Max No. of Electrodes *

→ Removing trails because of ...

Electromyography

Blinks ①

- correction Methods

i) ICA

ii) Regression Based

- Only Missed Stimulus → remove

Spatial filtering & ICA

oculomotor Activity ②

↓
Eye movements

- use central fixation points

- use short stimulus duration

- Ref choice

Nose Earlobe
move contamination less

- Research Focus

③ Based on EMG in EEG Channels

- High freq. 20-40 Hz Burst like

- Subject sneezed, coughed, moved their jaw

- Reject trails, (15 Hz) localized

- Baseline Normalisation, ICA

④ Based on task performance

- Error trails, No resp. trails, Too many response

- unrealistic response

* Not always visible in EEG *

Frontal → Big problem
Central → Not an issue

⑤ Based on Response Hand EMG

Why Response EMG? → ideally

- Detects partial errors But How?

① record

② preprocess

i) take derivative of EMG → Z-transf → rectify

ii) Normalise

③ criteria:-

- Zscore of incorrect hand > 2 b/w stimulus & Press button
- Peak = 2x larger than baseline

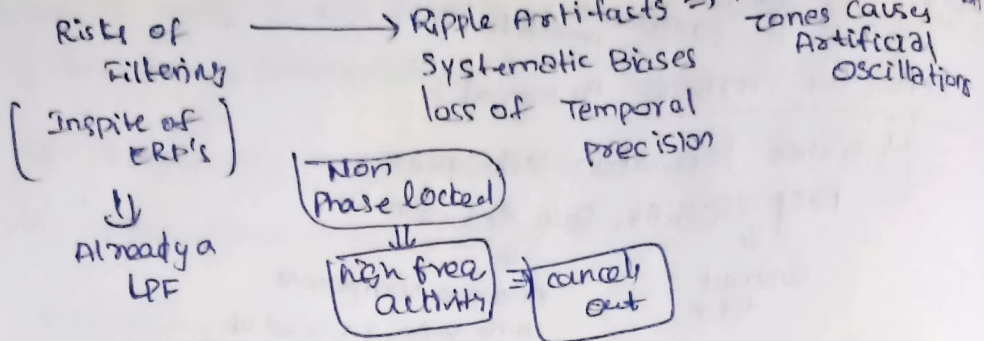
⑥ Train Subjects to minimize Artifacts

Explain them clearly

Awareness helps a lot!!

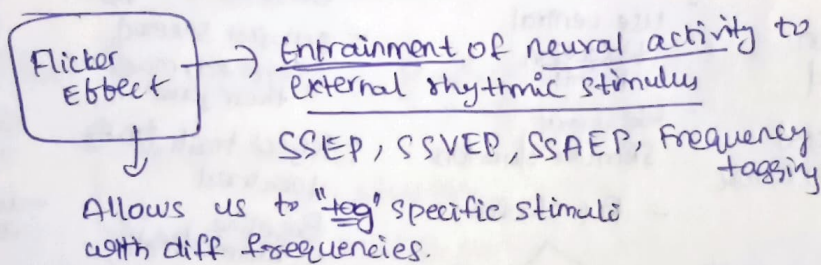
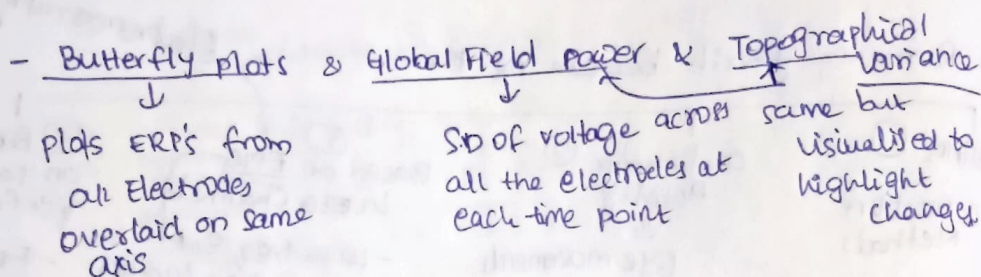
⑦ minimize during data collection

↓
our own Responsibility
No other choice!!



* Filtering ERP's \Rightarrow optional *

It done :- choose the range of filter properly



\rightarrow Is it oscillations? or Repeated ERP's?

Analysis

i) compare flicker freq power before vs during flicker or against neighbouring non flicker freq

\rightarrow Topographical Maps \rightarrow Shows spatial distribution of EEG/ERP activity across scalp

- Constructed by Interpolating voltages b/w electrodes
- identify Bad electrodes
- Easy to interpret

Micro States

EEG scalp topography remains stable \Rightarrow "quasi stable"

for ~70-130ms then

Rapidly shifts



\Rightarrow Each stable period is microstate

\Rightarrow linked to cognitive fns :- perception, memory, language

Global map Dissimilarity

\Downarrow

Diff b/w Scalp map at

$t=t_0$ & $t=t_1$

Low = stable state

Sudden Increase = Transition

\rightarrow

ERP Images

- 2D representation of single trial EEG at one electrode

-