# **Operating Manual for**

# Adaptive Signal Analysis & Processing (ASAP) for ERPs

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Web: <a href="http://www.escience.cn/people/cong/AdvancedSP\_ERP.html">http://www.escience.cn/people/cong/AdvancedSP\_ERP.html</a>

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# Introduction

### Why do we write this code?

This code demo is written for processing and analyzing One-factor, Two-factor and Three-factor ERP/EEG data. The specific flow chart is as below:

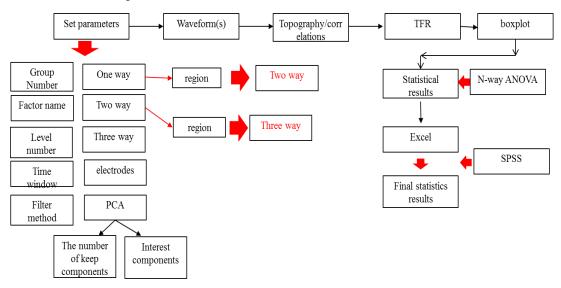


Figure 1.The flow chart of processing and analysis ERP data

#### The information of demo data

In order to best understand how to use this code demo for processing and analysis EEG/ERP data, two demo data can be used.

The first demo data was collected by Pro. Peng Li, from College of Psychology and Sociology, Shenzhen University(within subjects analysis test, two factors [2\*2]. It can also be used to one-factor [4\*1] analysis).

 22 subjects; Epoch Start is -200 millisecond; Epoch End is 1000 millisecond; the sampling rate is 500 Hz.

The second demo data was collected by Dr. Guoliang Chen, 215th Clinical Division, 406th Hospital of PLA, Dalian (between subjects analysis test, two factors [2\*2].

• Two groups: Control (20 subjects); Patient (20 subjects); Epoch Start is -200 millisecond; Epoch End is 1000 millisecond; the sampling rate is 1000 Hz.

# How to use this code demo

#### 1. Forming a struct:D

A struct D should be obtained, including a four-order tensor, epoch start time, epoch end time, sampling rate, and group label. Specific steps are as follows.

• Step1: forming a four-order tensor (channel\*time\*stimulus\*subjects).

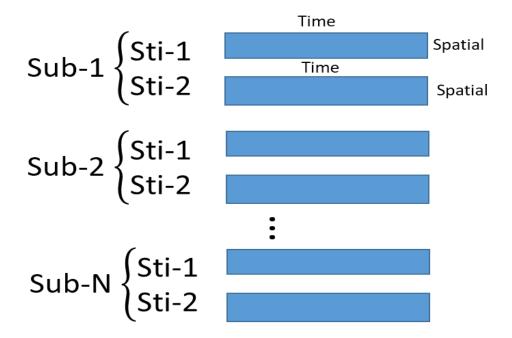
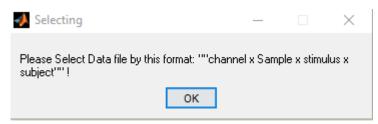
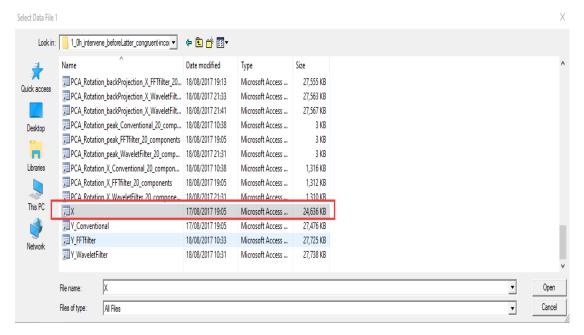


Figure 2. Take one-way test data (two level) forming four-order tensor as example.

- Step2: forming the struct D.
  - run the code:m\_0\_forming\_theStruct\_D, then follow the prompts, as shown below:
  - select the four-order tensor data

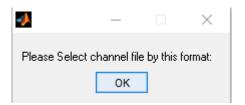


Click 'OK'.

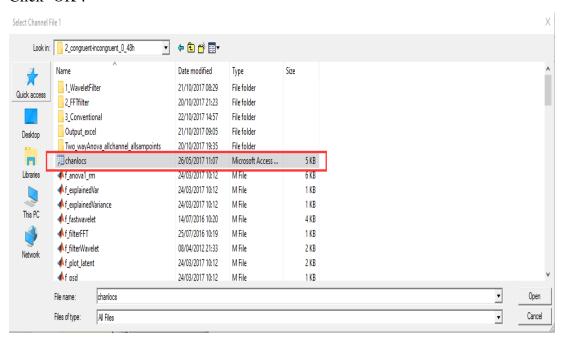


Select the data obtained according to step1.

> select the channel file

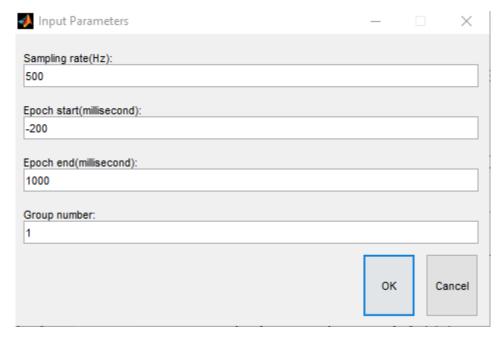


#### Click 'OK'.

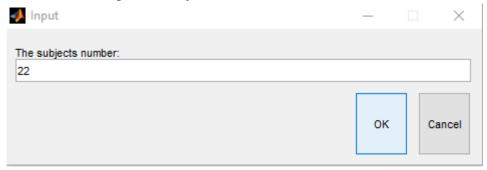


Select the channel file

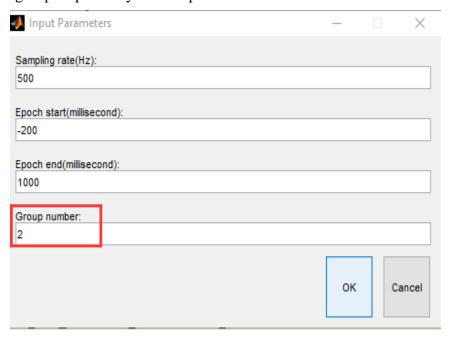
- set the parameters
- One group



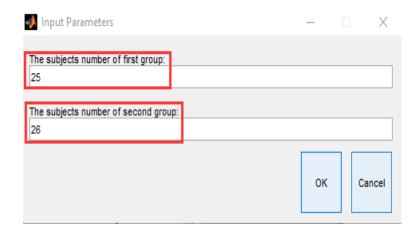
click 'OK', then input the subjects.



➤ If the group number is more than 1, you should input the subjects number of each group respectively. The steps are as follow.



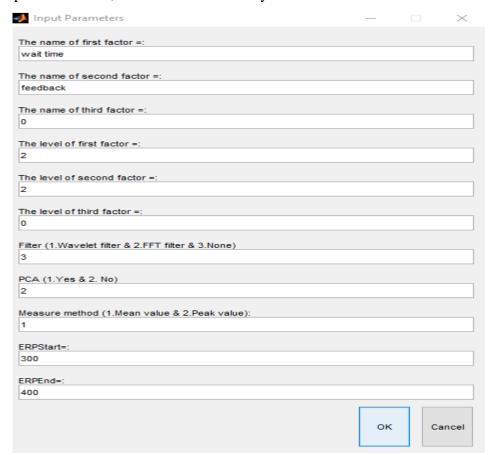
Click 'OK'.



# 2. Within subjects analysis

#### 2.1 Conventional method

• Firstly, you should run the code: m\_1\_ERP\_Analysis, then set the parameters, the grand average waveform of interest electrodes, the topography, the topo the topography correlation between subjects, time frequency representation, box plot of the data, and the statistical analysis results obtained.



Set the parameters: the name of first factor is wait time, the name of second factor is feedback; the name of third factor is 0; the level of first factor is 2; the level of second factor is 2; the level of third factor is 0; filter method is none; PCA is not used, measure method is mean value; the start time of time window is 300 millisecond; the end time of time window is 400 millisecond.

Then, input the name of interest electrodes according to channel file.

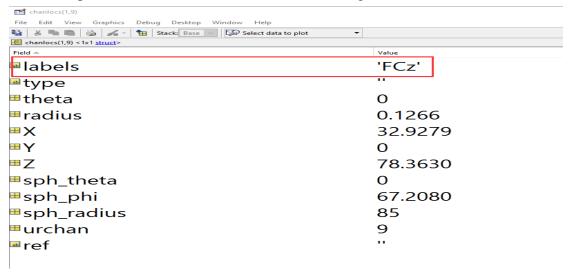
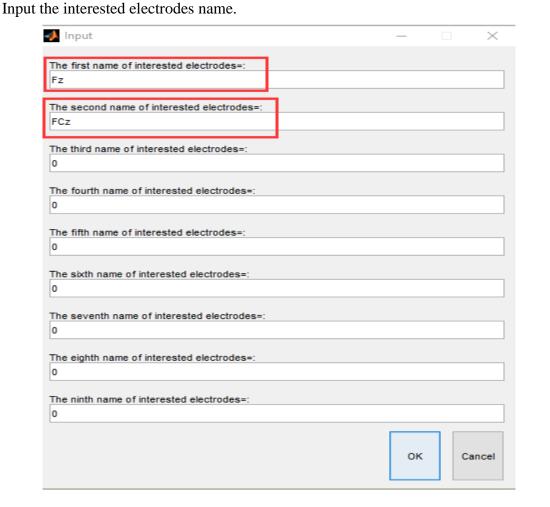


Figure 3.An example of one electrode name



# Click 'OK'. Those results will be obtained.

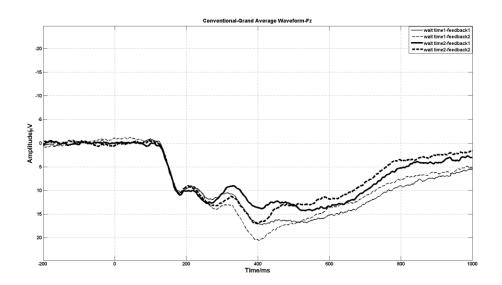


Figure 4.The grand average waveform at Fz

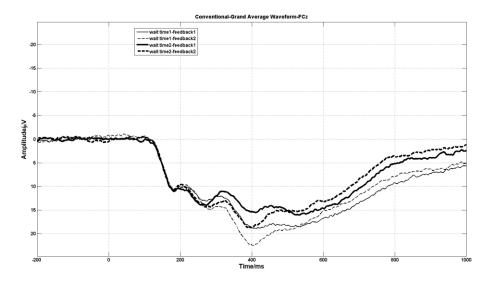


Figure 5. The grand average waveform at FCz  $\,$ 

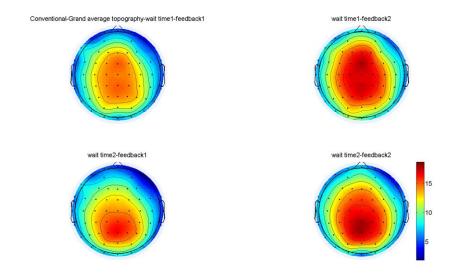


Figure 6.The grand average topography

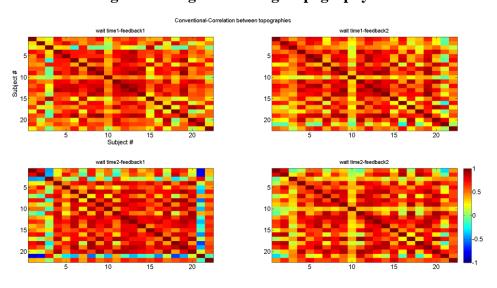
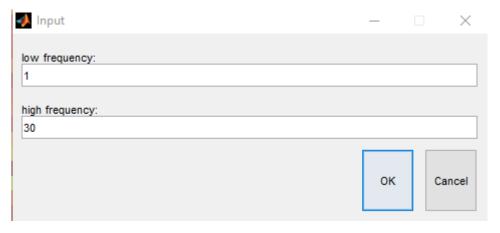


Figure 7.The correlation between topographies

Set the parameters for time-frequency analysis, low frequency value and high frequency value of frequency range.



Then, click 'OK'.

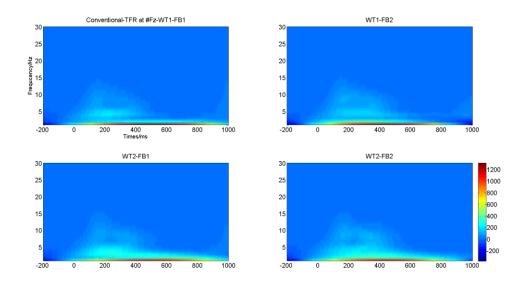


Figure 8.The time-frequency representation at Fz

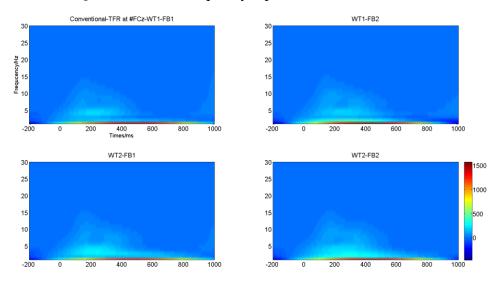


Figure 9. The time-frequency representation at FCz

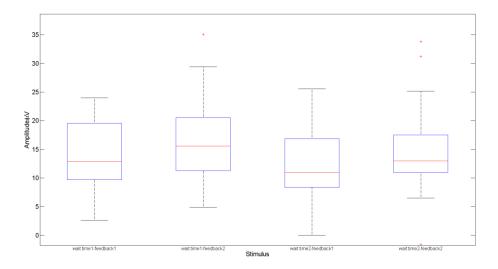


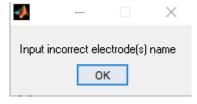
Figure 10.The boxplot of the data (the window is from 300 to 400 millisecond, the interest electrodes are Fz,FCz)

1.43 571.15	1 1 1	Mean Sq.  1.428  571.153  447.587	0.03 13.42	0.8551 0.0004	
1.43 571.15 447.59 3574.3	1 1 1	571.153 447.587	13.42	0.0004	
447.59 3574.3	1	447.587			
3574.3			10.52	0.0017	
	84	40 554		0.001/	
4594.47		42.551			
	87				

Constrained (Type III) sums of squares.

Figure 11.The results of statistical analysis (the window is from 300 to 400 millisecond; the interest electrodes are Fz,FCz)

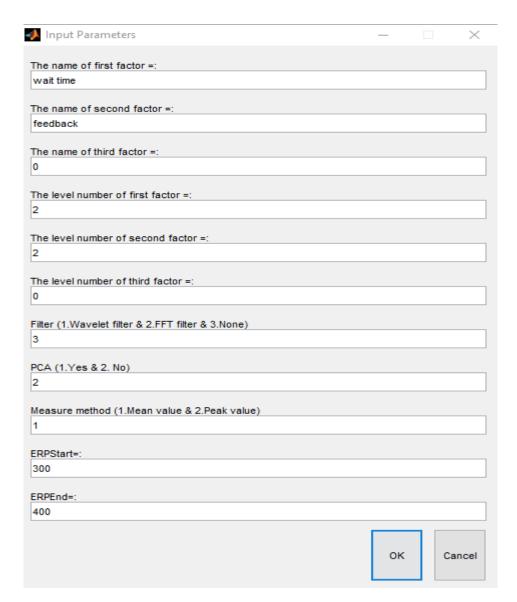
If you input incorrect electrode(s) name or do not input anything, the following tips will be given:



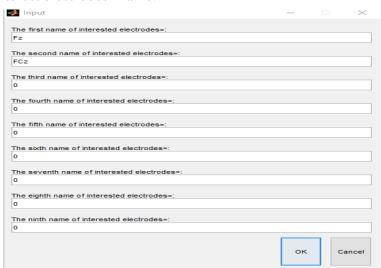
Please, click 'OK'. Other input window will appear. Input interested electrode(s) name again.

#### • Output the data

Run the code: m\_4\_outputdata\_excel. Output the data of interested component at interested electrodes, and then use SPSS for further analysis. The steps are as follow: Set the parameters:



#### Input interest electrodes' name:



#### Click 'OK'.

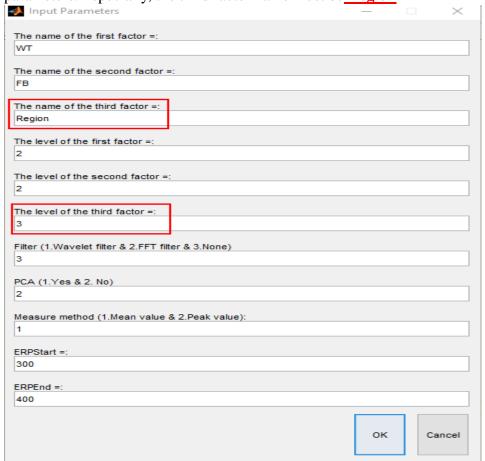
Then, the data of the time window 300-400ms and interest electrodes were

imported into excel as shown in figure 12.

1	wait timel-feedback1	wait time1-feedback2	wait time2-feedback1	wait time2-feedback2
2	5. 559942592	4.892975658	-0.034007352	-1.604017693
3	13.59754728	15. 11314073	8.35611875	13.03624674
4	18. 12222851	13. 24727698	10.12511959	11.94992744
5	8. 229799446	11.98082454	11.13041595	12.96222205
6	12.74391173	11.91424574	7. 289813658	10.96095858
7	5.616828777	8.836184928	8.18290422	7. 231415237
8	17.87280772	20. 34327672	16.90528599	13. 35448377
9	12.64478801	13.73555039	9.872050005	15.60136149
10	11.10733538	17.05754501	13.82617256	15. 94076294
11	2. 630242538	6.061902244	2. 966013157	6.503776074
12	23. 74954989	27. 84338496	19.06123768	25.16335193
13	22.10258569	26. 72755104	25. 54351495	33.80483879
14	22. 33392649	35.08289627	12. 29419849	20. 2554984
15	15.15969405	20.50756067	10.0314783	11.27086617
16	19.54930569	17.18631152	13.58982647	12.96389031
17	13.02535008	16.09033757	16.88000233	10.99830198
18	20. 5452699	29. 411 46674	20.61972324	31.17620688
19	11.99591296	9. 9258263	8.666429602	7. 924248386
20	10. 45313239	19.69905225	11.2762523	14.39468026
21	23. 95938855	27. 17721218	19. 41782923	21.10729015
22	9. 63054688	10.18664747	4.802277119	8. 767951759
23	9. 701472041	11. 27713333	10.81494372	17. 54484471

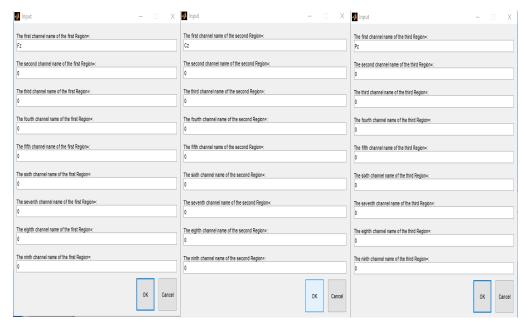
Figure 12.An example for importing data into excel\

• The operations for different brain regions:run **m\_1\_ERP\_Analysis**. Set the parameters. Especially, the third factor name must be **Region**.

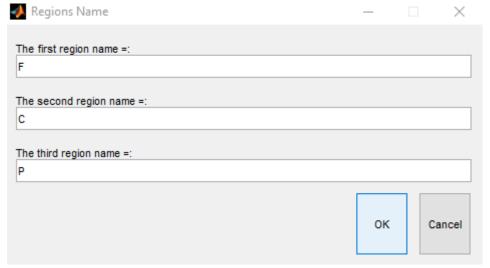


Click 'OK', then input the name of interest channel, the grand average waveform of interest electrodes, the topography, the topography correlation between subjects, and time frequency representation obtained.

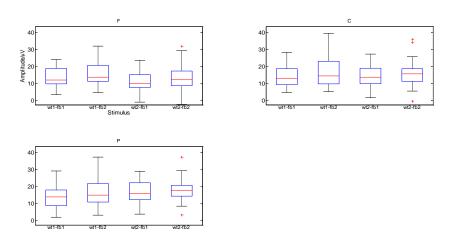
Next, three user interfaces will appear respectively for inputting the name of the first region/second region/third region.

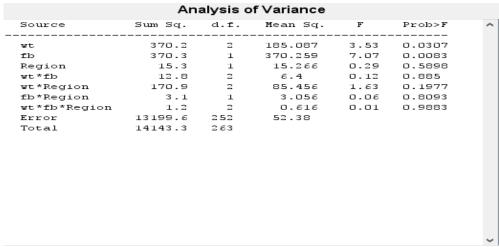


Input the name of each region.



Click 'OK'.





Constrained (Type III) sums of squares.

#### 2.2 FFT filter & Wavelet filter

• Step1, Run the code: m\_2\_Filter\_ERP -to filter the average ERP data.

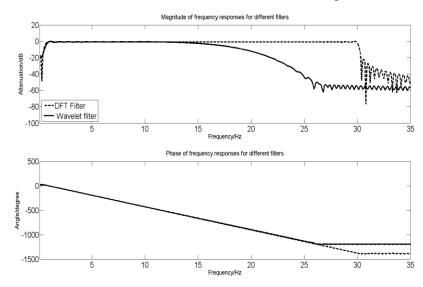
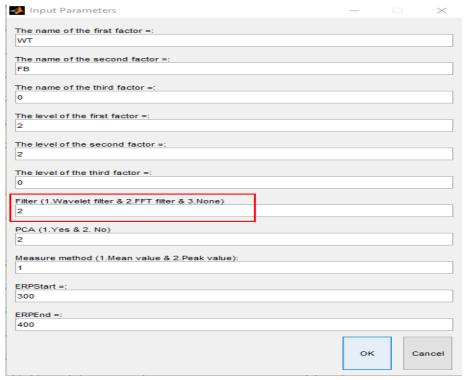


Figure 2.1 frequency responses for different filters

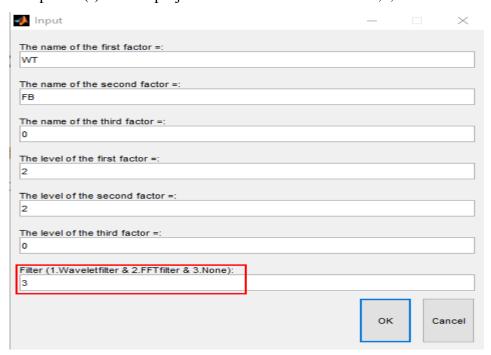
• Step2, Run the code: **m\_1\_ERP\_Analysis**, the filter method is 1 or 2 and the rest parameters setting are similar to the conventional method, shown as below.



• Step 3, run the code: m\_4\_outputdata\_excel,output the data of interested component, then use SPSS to further analysis.

#### 2.3 Conventional method/FFT/Wavelet Filter + PCA

• Step1, Run the code: m\_3\_PCA\_rotation\_ERP\_multipleGroups, determine how many components should be retained, then select the interested component(s) to back projection. Filter method can use 1,2, and 3.



Next, select the reserved components according to explained variance of 17/25

accumulated lambda.

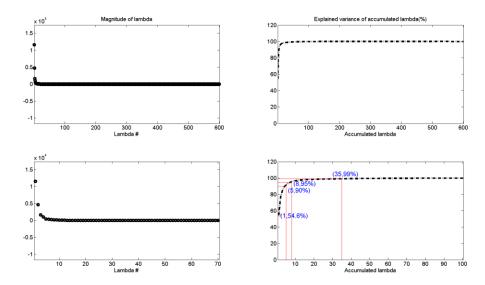


Figure 2.3.1 explained variance of accumulated lambda of conventional method Here, 10 was reserved, then select the interested component(s) to back projection.

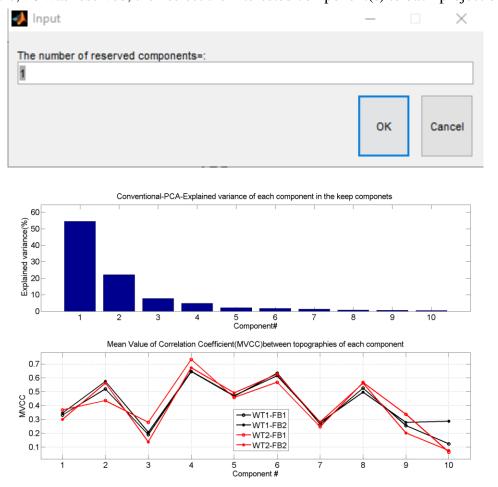


Figure 2.3.2 Explained variance of accumulated lambda of conventional method and Mean Value of Correlation Coefficient (MVCC) between topographies of each component

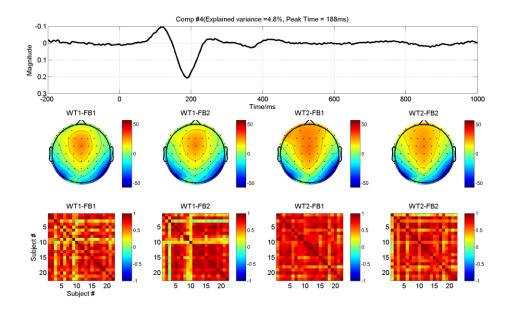


Figure 2.3.3 The 4<sup>th</sup> component

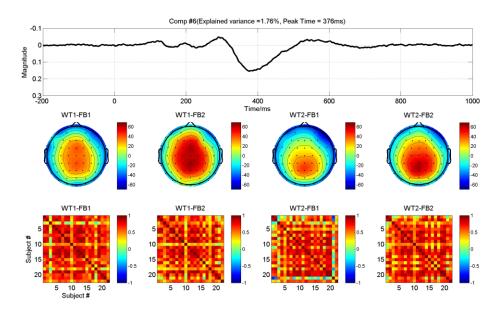


Figure 2.3.4 The 6<sup>th</sup> component

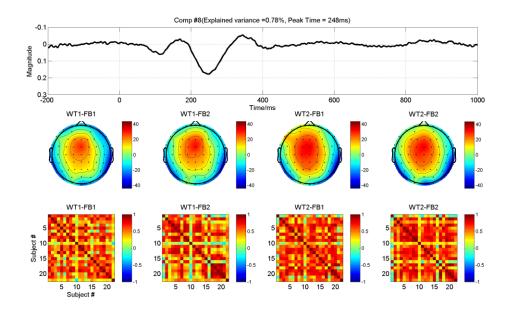


Figure 2.3.5 The 8<sup>th</sup> component

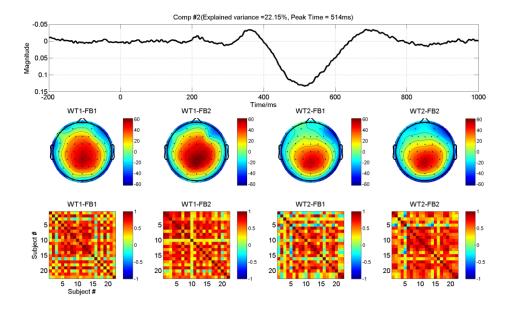


Figure 2.3.6 The 2<sup>th</sup> component

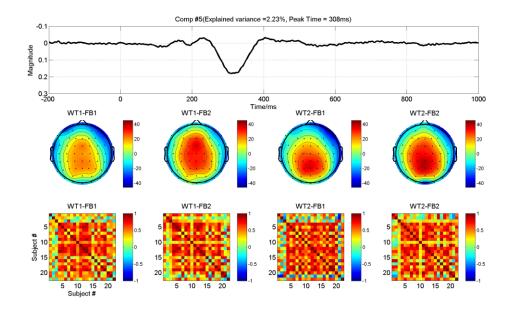


Figure 2.3.7 The 5<sup>th</sup> component

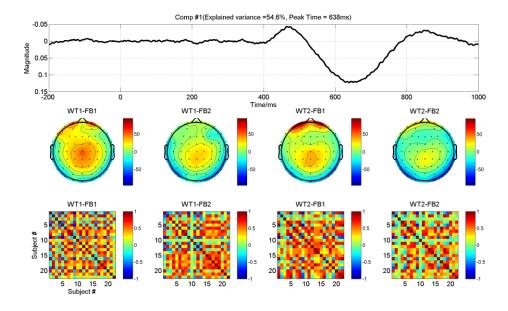


Figure 2.3.8 The 1<sup>th</sup> component

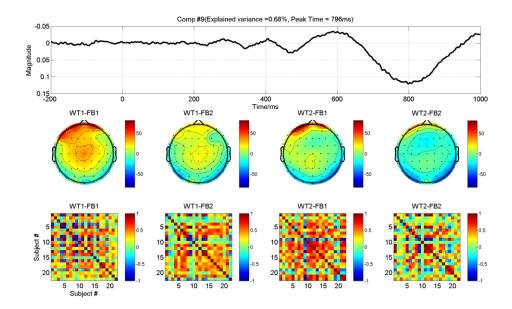


Figure 2.3.9 The 9<sup>th</sup> component

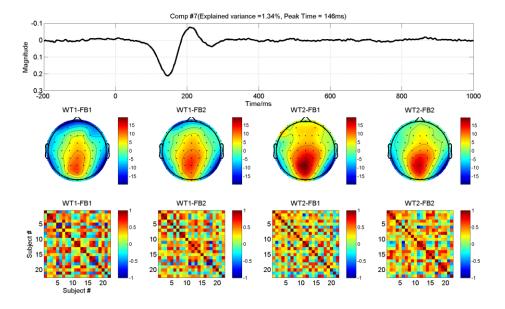


Figure 2.3.10 The 7<sup>th</sup> component

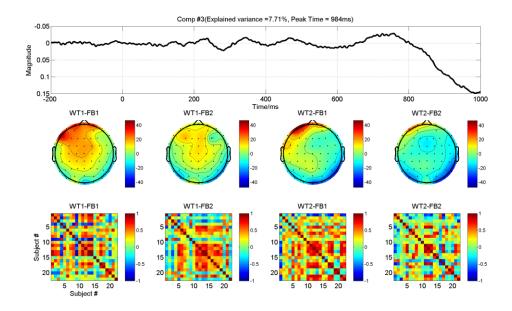


Figure 2.3.11 The 3<sup>th</sup> component

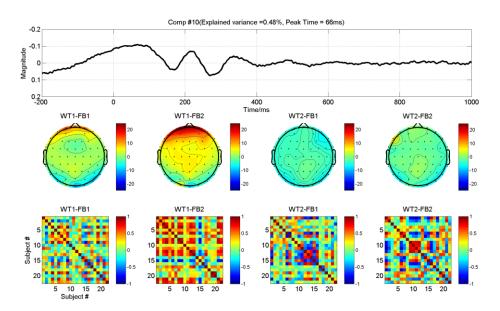
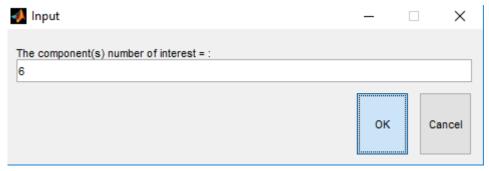
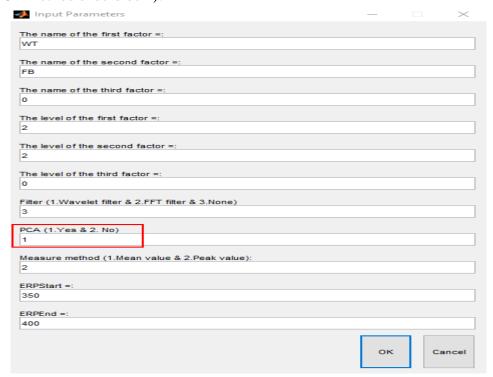


Figure 2.3.12 The 10<sup>th</sup> component

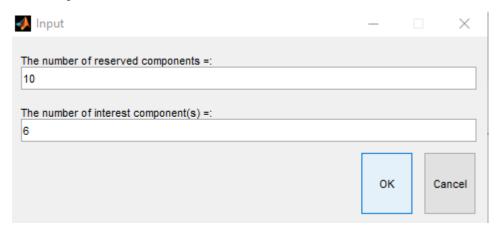
According to the properties of interest component in the time domain and spatial domain and the correlation coefficient between topographies, the  $6^{th}$  component was selected to back projection.



• Second, run the code: **m\_1\_ERP\_Analysis**, then set the parameters, the grand average waveform of interest electrodes, the topography, the topo the topography correlation between subjects, time frequency representation, box plot of the data, and the statistical analysis results were obtained(especially, the PCA method should be 1).



Then, input the number of reserved components (here, it is 10) and the number of interest component (here, it is 6).

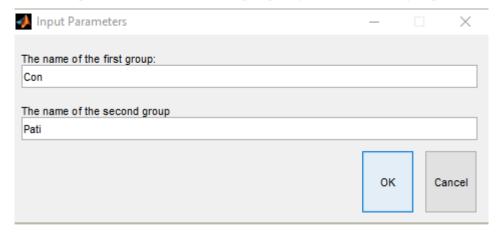


Next, input the name of interest electrode(s) in the user interface.

• Finally, run code: **m\_4\_outputdata\_excel**, the parameters setting are the same as step 2.

## 3. Between subjects analysis

Here, the similar operation should be done except inputing the name of each group.



# 4. Acknowledgement

My acknowledgements are to Prof. Peng Li, Dr.Guoliang Chen for providing ERP data. Meanwhile, express my thanks to Tiantian Yang, Xiaoshuang Wang, Jianrong Li, XiaoyuWang, Jianning Du for helping me to improve this code demo.