

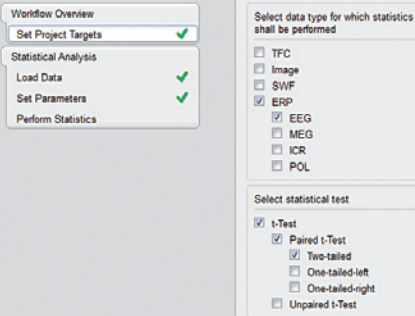


BESA Statistics

**CE-certified software for cross-subject
statistics of EEG / MEG data**



Intuitive Workflow



Optimized for data type

Preliminary Statistics



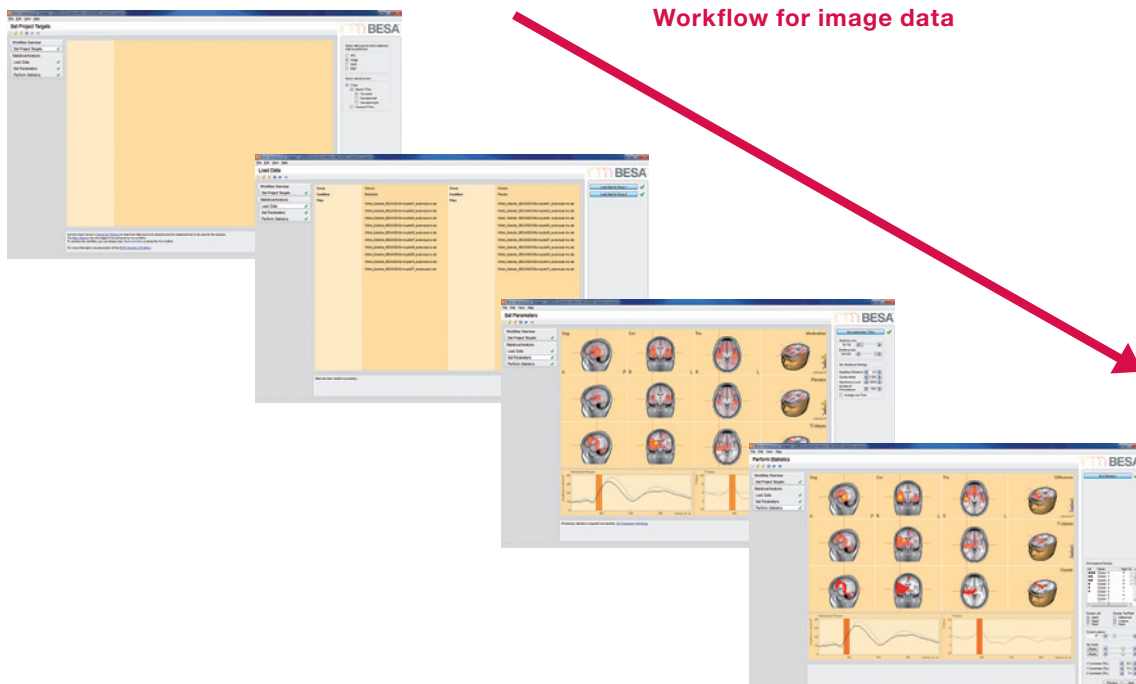
For defining regions of interest

Data Input

- Event-related potentials / fields (ERP/F)
- Image data, e.g. LORETA, beamforming
- Time-frequency data, e.g. temporal-spectral evolution, coherence, inter-trial phase-locking
- Source waveforms

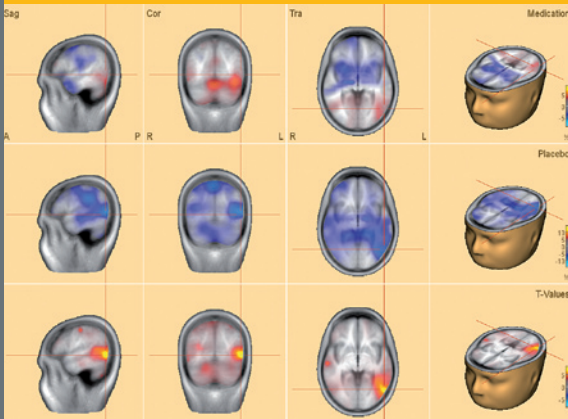
Preliminary Statistics

- Two groups (e.g. patients, controls) or conditions within the same group of subjects (e.g. time 1, time 2) can be compared
- Paired or unpaired Student's T-tests are calculated and visualized
- Based on these preliminary T-tests regions of interest can be defined that are passed on to permutation testing



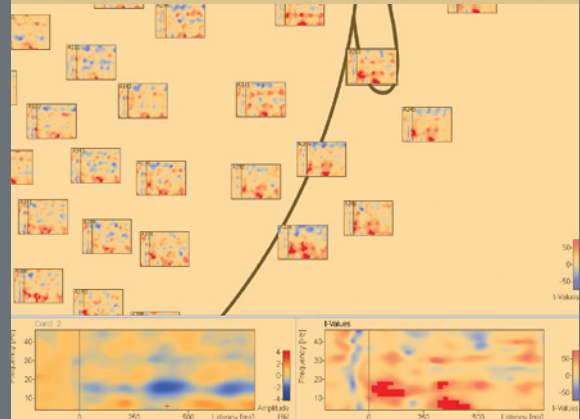
Workflow for image data

Cross-Subject Statistics based on Permutation Tests



Significant clusters in time, space and frequency are determined

Visualization of Significant Data Clusters



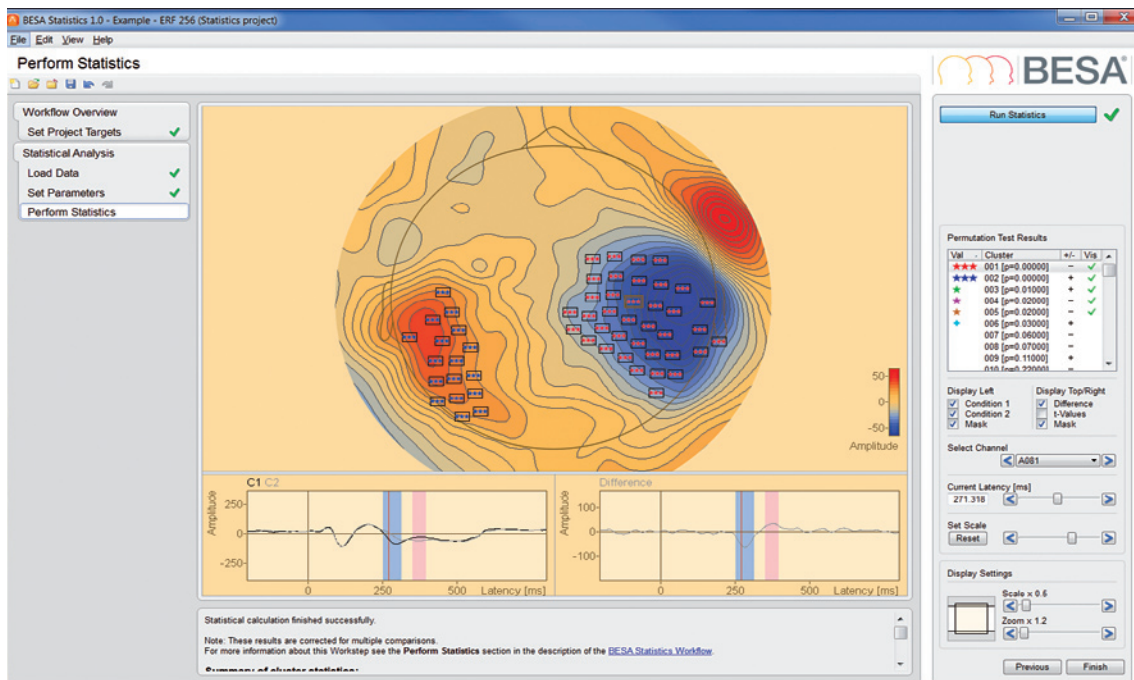
Suitable for publications

Cluster Permutation Statistics

- BESA® Statistics will determine significant clusters in time and if applicable space and frequency that show robust differences between groups / conditions
- Clusters will be probed for significance by permutation testing
- Significant data clusters are visualized in categories (highly significant, significant, trend)
- Results are corrected for multiple comparisons

Exporting Data

- Statistical values can be directly used in scientific reports without further analysis
- NEW** - All statistical results can be exported
- NEW** - Detailed cluster information per person can be exported
- NEW** - All images can be saved as vector graphics (eps)





For sales and product enquiries, please contact:

Rogue Resolutions Ltd
The Creative Quarter
8a Morgan Arcade
CARDIFF
CF101AF

info@rogue-resolutions.com
www.rogue-resolutions.com

BESA GmbH
Freihamer Str. 18
82166 Gräfelfing – Germany

Phone + 49.89.89 80 99 66
Email info@besa.de
Web www.besa.de



The CE marking certifies that this product fulfills the basic requirements of the Medical Devices Directive MDD 93/42/EEC. The number represents the identification number of the Notified Body which carried out testing and certification.

© Copyright 2015

