

# fctSPM: Factorial ANOVA and post-hoc tests for Statistical Parametric Mapping in MATLAB

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### Summary

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Statistical Parametric Mapping (SPM) is a statistical method originally used in neuroimaging developed in the early 90's in biomedical imaging, allowing to determine which brain zones were solicited during a functional MRI (Friston et al., 1995). Originally developed for a three dimensional analysis, the application of this method to the analysis of vectors or matrices was made possible thanks to Pataky (2010) whom allows to perform statistical inference on curves (vectors - 1D) or maps (matrices - 2D).

As in "classical" statistics on scalar values (0D), there is a parametric and a non-parametric approach to the SPM method. While the parametric method is based on random gaussian fields, the non-parametric method is based on label permutation tests (Nichols & Holmes, 2002), and thus, on re-sampling and randomness to make statistical inference. The main advantage of the non-parametric approach is that a gaussian distribution of the data is not required, making possible it to work with both curves and maps.

## Statement of need

Most of physiological data measured during human movement are continuous and expressed 20 in function of time. However, researchers predominantly analyze extracted scalar values from 21 the continuous measurement, as the mean, the maximum, the amplitude, or the integrated 22 value over the time. Analyzing continuous values (i.e., time series) can provide more infor-23 mation than extracted indicators, as the later discards one dimension of the data among the 24 magnitude and localization in time. In addition, oscillatory signals such as muscle vibrations 25 and electromyograms contain information in the temporal and frequency domains. However, 26 scalar analysis reduces the information at only one dimension by discarding two dimensions 27 among the magnitude and the localization in the time and/or frequency domain. 28 To analyze all the dimensions of a signal without losing information, the analysis of curves

29 or maps was proposed, coded, and put online by Pataky. However, the use of the proposed 30 functions does not allow the analysis of 2D data automatically. Moreover, a rather frequent 31 error is to consider only the significance of the last statistical test performed and not the 32 intersection between the post-hoc tests and the ANOVA. Indeed, a difference between two 33 samples can be significant if, and only if the ANOVA is significant in the same areas. This 34 package, integrated in the fctSPM repository at ./fctSPM/src/spm1d\_Pataky is published 35

elsewhere, and thus is not part of this JOSS review. 36

### DOI: 10.21105/joss.0XXXX

#### Software

- Review C<sup>\*</sup>
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#### Editor: Editor Name C

Submitted: 01 January XXXX 11 Published: 01 January XXXX 12

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# <sup>37</sup> fctSPM

 $_{38}$  The function we propose meets two objectives. 1/ to allow statistical inferences on curves

 $_{
m 39}$  and maps with a standardized format and 2/ to simplify analyses by comparing means while

 $_{\rm 40}$   $\,$  considering intersections with tests performed upstream (ANOVA and post-hoc of main ef-  $_{\rm 41}$   $\,$  fects).

The statistical tests are performed taking into account the independent and repeated measure effects provided in the obligatory function inputs. ANOVA, up to three-way ANOVA with

<sup>43</sup> effects provided in the obligatory function inputs. ANOVA, up to three-way ANOVA with <sup>44</sup> three repeated measures, is performed if required, and followed by post-hoc tests as paired or

<sup>45</sup> independent Student t-tests. By default, the ANOVA is performed with an alpha risk of 5%,

<sup>46</sup> while post-hoc tests alpha risk is adjusted with Bonferronni correction. A number of 10/alpha

47 iterations (200 for a 5% risk) is defined for each test. Statistical parameters are customizable

via optional inputs, like multiIT which can be used to increase the number of iterations and

<sup>49</sup> achieve numerical stability and reliable analysis (Nichols & Holmes, 2002). A Matlab (.mat)

 $_{50}$  file containing the number of permutations, the significant clusters, the statistical thresholds,

<sup>51</sup> and the raw data used in the analysis is also generated for each test family.

To interpret the results, figures directly usable for presentations and/or articles are available. In one dimension, the main figure contains the mean and standard deviations for each group of the analyzed condition, and the results of the post-hoc tests corrected with the result of the ANOVA. In two dimensions, the mean maps and the standard deviations are on two separate figures, and the result of the statistical analysis is displayed on the map of differences between two modalities. Therewith, other figures that display absolute and relative differences, effect sizes, and the raw value of the statistical test and its threshold are available.

<sup>59</sup> To personalize the figures, Matlab (.fig) files are implemented to perform a posteriori modifi-

<sup>60</sup> cations, and optional inputs are available to a priori customize the figures. These parameters

 $_{61}$  are gathered in three categories: 1/ general plot parameters working identically for one and

two dimensions, acting on axis labels and limits, image resolution and size 2/ one dimension parameters that act on the characteristics of the curves like color and transparency, or the

parameters that act on the characteristics of the curves like color and transparency, or the position of the statistical analysis relatively to curves 3/ two dimensions parameters acting

<sup>64</sup> position of the statistical analysis relatively to curves 3/ two dimensions parameters acting <sup>65</sup> on the colormap and its limits, as well as the color of the statistical test displayed on map of <sup>66</sup> differences

66 differences

This function was already used in Trama et al. (2021) to compare soft-tissue and muscle vibrations of the *vastus lateralis*. It is currently used to assess modifications of soft-tissue vibrations caused by mountain ultra-marathons, the effect of the pedaling phase on *quadriceps* soft-tissue vibrations, and differences in isokinetic torque after ACL operation and rehabilitation.

# 71 References

Friston, KJ., Holmes, AP., Worsley, KJ., Poline, JB., Frith, CD., & Frackowiak, RSJ. (1995).
 Statistical parametric maps in functional imaging: a general linear approach. *Human Brain*

74 Mapping, 2, 189–210. https://doi.org/10.1002/hbm.460020402

Nichols, TE., & Holmes, AP. (2002). Nonparametric permutation tests for functional neuroimaging: a primer with examples. *Human Brain Mapping*, *15*, 1–25. https://doi.org/
 10.1016/j.jbiomech.2010.03.008

78 Pataky, TC. (2010). Generalized n-dimensional biomechanical field analysis using statistical

parametric mapping. Journal of Biomechanics, 43, 1976–1982. https://doi.org/10.1016/
 j.jbiomech.2010.03.008

Trama, R., Hautier, C., Souron, R., Lapole, T., Fouré, A., & Blache, Y. (2021). Is accelerometry an effective method to assess muscle vibrations in comparison to ultrafast



- ultrasonography. *IEEE Transactions on Biomedical Engineering*, *68*, 1409–1416. https://doi.org/10.1109/TBME.2020.3035838 83
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