# SUPPLEMENTARY MATERIAL FOR

# **The morphospace of the brain-cognition organisation**

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**Supplementary Figure 1. Replication of the morphospace in four additional parcellation approaches.** Pearson's r of the comparison between each additional parcellation and the morphospace is indicated for the 100 (top left), 400 (top right), 800 (bottom left) and a control random parcellation (i.e. null model, bottom right).



**Supplementary Figure 2**. **Representation of the morphospace embedding with different n neighbours and minimum distance values.** Higher values average the manifold largely across the data and vastly distribute the data in the low embedding. Extremely low values result in spurious clusters of connected neighbours. Given that the neuron-shaped architecture is stable across low to medium values and that there is no goal standard in the finer manifold/global structure trade-off, we opted to rely on the low default metrics that reveal the data manifolds maintaining the global structure without spurious neighbouring connections. Points represent the 506 meta-analytic maps.



**Supplementary Figure S3. Comparison between versions of the morphospace built in 2, 3, 4, and 5 dimensions.** (a) The distribution of the MAE between measured and predicted maps indicates that the 3D predictions are more precise than the 2D ( $z = 4.37$ ; p<.001) and 5D ( $z = 5.18$ ; p<.001). (b) The difference in predictability index between the different dimensions indicates that the 3D predictability index is higher than the 4D ( $z = 3.29$ ; p<.001), and the 2D ( $z = 9.63$ ; p<.001). \*\*\*: p<.001. Lines, boxes, whiskers and dots represent the median, quartiles, distribution, and observations (506 meta-analytc maps).



**Supplementary Figure 4.** The association of the predictability index the number of terms aggregated in each out-of-sample meta-analytic map. The low Pearson's correlation  $(r = 0.15)$  between the predictability index (x-axis) and the number of terms aggregated in each map (y-axis) reveals that the index is not associated with the magnitude of data used for each meta-analysis. Datapoints representing the 888 meta-analytic maps are represented in blue, and the regression line is indicated in red. r: Pearson's regression coefficient.



**Supplementary Figure S5. Robustness of results against a spatial autocorrelation-preserving null model.** a) Pearson's correlation of the comparison between the 400 parcels resolution of the Schaefer and colleagues atlas (2018) applied to the maps' surface projection and the original morphospace. b)The plot shows the distribution similarity between the surface-projected empirical and SA-preserving null versions an example meta-analytic map (auditory). c) Pearson's r of the comparison of the Euclidean distances between the surface-projected empirical and SA-preserving null maps, both parcellated via Schaefer and colleagues' atlas (2018). d) Pearson's r of the comparison between predictability indices obtained from the predicted 506 meta-analytic maps and the predicted SA-preserving null maps. Datapoints representing the 506 meta-analytic maps are represented in blue, and the regression line is indicated in red. The axial slices next to the x and y axes represent the example version of the auditory map predicted from the original morphospace and the SA-preserving null space, respectively. Dim: dimension. SA: spatial autocorrelation.



**Supplementary Figure S6.** predictability index of 888 Neuroquery meta-analytic maps. Topic modelling allowed for (a) 12 broader and (b) 55 finer clusters summary of the predictability index of (c) the meta-analytic maps colocalised onto the morphospace. Bar colours indicate the predictability index of their five nearest neighbours (5nn) in the morphospace. Triangles indicate each new map's coordinate in the morphospace, while transparent circles indicate the morphospace meta-analytic maps' location.



**Table S1** Features of the new task-related activation maps.

![](_page_8_Picture_398.jpeg)

![](_page_9_Picture_520.jpeg)

![](_page_10_Picture_455.jpeg)

### *Relationship between cognitive domains branches*

The neuron-shaped architecture of the morphospace clusters cognitive domains within each branch, with the position of each branch reflecting the relationship between the domains. For instance, the close position of the vision and attention branches with regard to others reflects the anatomical overlap between activations related to vision paradigms, from simple stimuli observation to eye-tracking paradigms, and with attentional networks<sup>12</sup>. Vision and action activations are also closely located in the morphospace, and their interaction is known to manifest as embodiment mechanisms (e.g. rubber hand illusion 13 ). Motor cognition and somatosensory mechanisms are jointly recruited during a movement to ensure online control and the successful outcome of the performance<sup>14</sup>. Further, the clusterisation of the domains within the morphospace shows that the emotion and somatosensory domains are adjacents, reflecting bodily signal generation and processing of emotional responses<sup>15-17</sup>. Emotions and somatic responses guide decision-making<sup>18</sup> as confirmed by the proximity of Emotion and Decision-making within the morphospace. The joint contribution of learning and memory allows humans to orient in social experiences<sup>19</sup>, thus their close clusterisation in the space. Contextualisation of memories occurs by assigning meaning and words to encoded items<sup>20</sup>, and language has been ascribed as part of the working memory as the phonological loop component<sup>21</sup>. Accordingly, memory, language, and working memory have interrelated aspects and follow one another in the morphospace. The auditory cognition clusters far away from the other domains. The striking difference in the anatomical pattern of auditory-modality fMRI task opens further queries on the possible influence of stimuli modality in activation studies.

## *Brain structures of the* predictability *map*

The cerebral regions associated with high predictability indices exhibit a strong correlation with the gradients that explain overall brain activity in particular areas of auditory and motor processing<sup>22</sup>. The superior temporal cortex has been shown to contribute to auditory cognition<sup>23</sup> and processing of the object's spatial features<sup>24</sup>, while medial temporal cortices such as the rhinal cortex, hippocampus, and amygdala play a role in memory<sup>25-28</sup> and stimuli representation (e.g. objects, faces, and scenes)<sup>29</sup>. The premotor cortex contributes to action planning<sup>30</sup> and speech<sup>31</sup>, while the FEF and PEF are involved in visual target detection<sup>32,33</sup>. An extensive range of functions for the implementation of voluntary action, such as timing, sensory predictions, sequence implementation, and inhibition of concurrent movements, involve the SMA and pre-SMA areas<sup>34-36</sup>. Finally, the involvement of Broca's area as a hub of the language network in the brain has been extensively confirmed by the literature since its first description by Broca in 1861<sup>31,37,38</sup>. Prediction, learning and reward mechanisms emerge from the activity of subcortical structures such as the basal ganglia and its connections<sup>39,40</sup> as well as the medial temporal lobe structures 41 .

### *Replication of the* morphospace *space architecture in 2021 dataset*

In order to assess the morphospace reliability in terms of clustering cognitive domains and their spatial positioning, the Euclidean distances between the morphospace maps were compared to the distances in a three-dimensional space built using the updated 2021 version of the 2017 meta-analytic maps. A total of 506 maps were obtained from the Neurosynth repository and matched with the 2017 dataset terms. No thresholding was applied to the 2021 meta-analytic maps as the newer version of Neurosynth automatically corrects for multiple comparisons by applying a threshold of  $z \geq 3.4$ . The maps underwent parcellation using the Glasser and colleagues <sup>42</sup> and AAL3 43,44 atlases delineated by our group.

The Uniform Manifold Approximation and Projection (UMAP<sup>45</sup>) algorithm was applied to reduce the dimensionality of the parcelled 2021 meta-analytic dataset in a three-dimensional space, and UMAP default parameter values were used. Specifically, the algorithm used the information of 15 local neighbours to learn the manifold structure of the data points; 0.1 minimum distance was allowed by the algorithm to pack the data; the Euclidean metric was used for the data embedding.

Using Python (https://github.com/vale-pak/BCS.git), the Euclidean distances between the 2021 three-dimensional space maps were computed and subsequently compared with those of the 2017 morphospace meta-analytic maps. Pearson's correlations revealed a positive correlation ( $r = 0.53$ ), affirming that the cognitive domain clusterisation and positioning observed in 2017 can be reliably replicated in later versions of the dataset.

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