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NEGATION OF ABSTRACT-CONCRETE CONCEPTS

Sentential negation of abstract and concrete conceptual categories: a brain decoding MVPA study

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Summary

We rarely use abstract and concrete concepts in isolation but rather embedded within a linguistic context. To examine the modulatory impact of the linguistic context on conceptual processing, we isolated the case of sentential negation polarity, in which an interaction occurs between the syntactic operator *not* and conceptual information in the negation's scope. Previous studies suggested that sentential negation of concrete action-related concepts modulates activation in the fronto-parieto-temporal action representation network. In this functional magnetic resonance imaging study, we examined the influence of negation on a wider spectrum of meanings, by factorially manipulating sentence polarity (affirmative, negative) and fine-grained abstract (mental state, emotion, mathematics) and concrete (related to mouth, hand, leg actions) conceptual categories. We adopted a multivariate pattern analysis approach, and tested the accuracy of a machine learning classifier in discriminating brain activation patterns associated to the factorial manipulation. Searchlight analysis was used to localize the discriminating patterns. Overall, the neural processing of affirmative and negative sentences with either an abstract or concrete content could be accurately predicted by means of multivariate classification. We suggest that sentential negation polarity modulates brain activation in distributed representational semantic networks, through the functional mediation of syntactic and cognitive control systems.

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1. Introduction

Research on conceptual knowledge has provided consistent neuropsychological and neuroimaging evidence that abstract and concrete concepts are represented in distributed brain networks, which are best differentiated at the level of structural and functional connectivity [1]. In so-called “hubs and spokes” models of conceptual knowledge, modality-specific nodes (i.e., spokes), for example in sensory-motor or limbic areas, are linked together with modality-invariant and graded hubs (though it is debated whether there are several hubs in the fronto-parieto-temporal cortices [2] or a single hub in the anterior temporal lobe [3]). According to multiple representation accounts developed within the grounded cognition framework, the differential involvement of these distributed networks for processing abstract and concrete concepts, and their more fine-grained sub-categories, reflects the type of experiential information that is more relevant for either concepts during acquisition [4–5]. Concrete concepts mostly draw on sensory-motor experience [6], whereas abstract concepts mostly draw on emotional, introspective, social, and linguistic experience [5, 7–10]. Whether and to which extent experiential information also contributes to a context-dependent appropriate use of concepts is debated. Previous work revealed differential contextual effects on abstract and concrete word processing. These were explained in terms of either lower context availability [11] or higher context diversity [12] for abstract versus concrete words. However, the neural mechanisms through which linguistic contextual information modulates multiple grounded representations are still poorly understood [13].

In the present study, we isolated the case of sentential negation as a benchmark for testing the role of linguistic context on neural conceptual representations, for two reasons. First, by reversing the truth value of conceptual information, sentential negation has clear-cut, discrete effects on its semantic interpretation, applying equally to concrete (e.g., *She doesn't kick the ball*) and abstract (e.g., *She doesn't remember the past*) concepts. Second, there is evidence that the sentential negation of concrete, action-related concepts reduces the access to experience-related sensory-motor brain networks. In a previous fMRI study [14], we showed that sentential negation of

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action-related concepts reduces activation and effective connectivity within the left-hemispheric premotor-parieto-temporal action representation system. Concordant evidence of motor cortex activity reduction in the presence of sentential negation was obtained in other fMRI [15], EEG [16], and TMS [17] studies. This effect has been interpreted as a reduced grounding in the sensory-motor system for the semantic processing of negative action-related sentences [18], possibly due to action inhibition mechanisms [16].

Building on this evidence, in the present study we tested not only concrete but also abstract concepts in order to address the question whether processing of specific semantic categories entails activation of brain areas coding for the corresponding experiential information and whether this activation is modulated by negation. We applied multivariate pattern analysis (MVPA) to fMRI data obtained while participants processed fine-grained abstract (mental state, emotion, mathematics) and concrete (related to mouth, hand, leg actions) concepts, presented either in the affirmative or negative polarity context (Table S1). Compared to univariate methods, MVPA can attain increased sensitivity, by considering activation across multiple voxels that may be even distributed in non-adjacent anatomical regions, rather than accepting or rejecting individual voxels based on a given significance cut-off [19-20]. Based on machine learning classification algorithms, MVPA aims at solving a classification problem (CP), by predicting the stimuli associated with a given activation pattern. Sentential negation for abstract and concrete conceptual categories was examined by specifying the following classification problems:

The first classification problem (CP1) tested whether sentential negation differentially modulates abstract and concrete semantic representations. According to multiple representation accounts and to “hubs and spokes” neuroanatomical models, abstract and concrete concepts are represented in partially distinct and distributed brain networks. Sentential negation may therefore operate on abstract versus concrete networks independently, yielding in principle four distinct neural representation levels: affirmative abstract, negative abstract, affirmative concrete, and negative concrete. By means of CP1, we tested whether our fMRI data contained sufficient

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information to discriminate between these four sentence types. Previous evidence demonstrated the feasibility of decoding abstract versus concrete semantic content processing, with classification sensitive regions widely distributed across the cortex [21]. To the best of our knowledge, this study provides the first decoding attempt applied to concreteness categories and negation polarities jointly, as reflected by their 2x2 factorial interaction.

The second classification problem (CP2) examined the sentential negation of abstract and concrete concepts at a fine-grained semantic level, based on grounded, multiple representation hypotheses that concepts representing different types of experience are represented by distinct modality-specific networks, and that negation operates on each of these networks independently. By applying a MVPA decoding approach, we aimed at predicting the brain activation patterns of each fine-grained conceptual category (mental state, emotion, mathematics, mouth, hand, leg) presented either in the affirmative or negative polarity context (6x2 factorial interaction).

We leveraged MVPA classification to also investigate the main effects nested in our factorial manipulation. Namely, sentential negation polarity (CP3: affirmative vs. negative); concreteness (CP4: abstract vs. concrete), and fine-grained conceptual category (CP5: mental state, emotion, mathematics, mouth, hand, leg). CP3 sought confirmation for a neural correlate of negation at the syntactic level (i.e. independently of the semantic meanings on which negation operates), an aspect for which limited evidence obtained by means of univariate analysis techniques is available [14,22]. CP4 and CP5 tackled the important question of replicability of previous results for concreteness [21] and fine-grained conceptual categories [23–24].

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2. Methods**(a) Participants**

Fifty subjects (25 females, M age = 23.02 years, SD age = 4.88) volunteered in the study. All subjects were Italian native speakers with a comparable level of education and with no reported history of neurological or psychiatric disorders. All subjects were right-handed (Edinburgh Inventory score: $M = .94$, $SD = .05$). Two subjects (1 female) were excluded due to brain structural anomalies.

(b) Experimental design

We applied a within-subject factorial combination of Conceptual category (3 abstract: mental state (Ms), emotion (Em), mathematics (Ma); 3 concrete: hand (Ha), mouth (Mo), leg (Le) action-related), and Polarity (affirmative (A), negative (N)). This resulted in 12 experimental conditions: AMs, NMs, AEm, NEm, AMa, NMa, AMo, NMo, AHa, NHa, ALe, NLe.

(c) Experimental stimuli

The set of stimuli consisted of 35 Italian sentences for each of the six conceptual categories, each sentence presented in the affirmative and negative form (total = 420 sentences) (Table S1). The set of affirmative sentences was validated in a previous rating study [9] with respect to linguistic variables, namely number of words (all 4-word sentences), number of syllables (Chi-square (25) = 36.37, $p = .07$); number of letters ($F(5,204) = 1.25$, $p = .29$); lexical frequency for, respectively, nouns ($F(5,204) = 1.86$, $p = .10$), verbs ($F(5,204) = 1.72$, $p = .13$), and noun-verb combinations ($F(5,204) = 1.82$, $p = .11$). In addition, sentences were rated with respect to semantic variables, including category-specific association, body-part association, concreteness, context availability and familiarity (see Tables 2, 3, 4 in [9]).

All affirmative and negative sentences were read aloud by a female native Italian speaker and recorded in a sound-proof room. Intensity was normalized for all sentences at 70 dB. Pitch was balanced across the six conceptual categories, both for affirmative, $F(5, 204) = 1.43$, $p = .21$, and negative sentences, $F(5, 204) = 0.65$, $p = .66$.

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In order to avoid repetition effects that could arise if participants were exposed to the same sentences in both polarities, the pool of 420 sentences was split in two lists of 210 experimental stimuli, each including only one version of each sentence, either with affirmative or negative polarity. Each list was assigned to one of two groups of subjects (Group A: 24 subjects, 12 females, M age = 23.71 years, SD = 6.14, Edinburgh score M = 0.94, SD = 0.06); Group B: 24 subjects, 12 females, M age = 22.42 years, SD = 3.61, Edinburgh score M = 0.95, SD = 0.05).

(d) Experimental Procedure

Presentation 14.9 (Neurobehavioral Systems) was used for stimulus presentation. Sentences were presented in an event-related design. The acquisition session for each participant comprised four runs of 12 minutes and 40 seconds each. Each run consisted of 67 randomized trials: 51 experimental, 9 catch, and 7 null trials. Experimental trials began with the auditory presentation of one sentence, followed by a 2000 ms interval, after which a visual fixation cross appeared for 500 ms. In catch trials, the fixation cross was replaced by a question mark, followed by a written sentence presented for 1000 ms, and participants were instructed to blink their eyelids once in case the written sentence matched the auditory sentence, or twice in case of mismatch (for full details, see [24], where the same procedure was applied).

(e) Data Acquisition

MRI scans were acquired with a 3 Tesla Intera Philips body scanner (Philips Medical Systems), equipped with an 8-channels head coil (SENSE factor = 2). In order to prevent scanner noise from affecting auditory sentence presentation, fMRI sparse sampling was employed [25–26]. Whole brain T2* BOLD images were acquired with a gradient-echo, EPI pulse sequence (TR = 2915 ms acquisition time + 7585 ms silent period = 10500 ms; TE = 30 ms). Each functional image comprised 35 axial slices (3.2 mm thick, .8 mm gap) acquired in ascending order (FOV: 240x240 mm; matrix size: 128x128). Each participant underwent four fMRI scanning sessions, each comprising 71 scans, plus two initial dummy scans, which were discarded prior to data analysis.

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A high-resolution T1 anatomical image was acquired for each participant (TR = 7.2 ms; TE = 3.5 ms), comprising 200 axial slices (1 mm slice thickness, 1x1 mm in-plane resolution).

(f) Data analysis

(i) Data preprocessing

Data were preprocessed with SPM8 (www.fil.ion.ucl.ac.uk/spm), including slice time correction, realignment, and normalization to the MNI space. Smoothing was not performed to provide optimal sensitivity for high-frequency multi-voxel patterns in MVPA [27]. The time series of each subject were high-pass filtered at 128 sec. No pre-whitening and no global normalization were applied. For each subject, we modelled a 6x2 factorial design (Conceptual category x Polarity) with four separate sessions and one regressor for each experimental condition. Additional regressors modelled the catch trials and movement parameters. For each subject, we defined a set of Student's t-contrasts, with a weight of +1 for one of the experimental condition regressors and a weight of zero for all the other regressors. The resulting 12 t-contrast images (AMs, NMs, AEm, NEm, AMa, NMa, AMo, NMo, AHa, NHa, ALe, and NLe) of each participant were used for the MVPA [28].

(ii) Multivariate Pattern Analyses

PyMVPA 2.5.0 (www.pympva.org; [29]) running under Python 2.7.9 (www.python.org) was used for MVPA. The t-contrast images were inclusively masked by an image defining the set of non-zero voxels shared by all subjects with grey matter tissue probability > .1 (for an equivalent procedure, see [24]). Subject-wise z-scoring normalization was applied to correct for noise-related inhomogeneities in voxel intensities. The t-contrast images were averaged subject-wise and condition-wise.

A linear SVM algorithm was used for all CPs (CP1: 2x2 factorial interaction, and post-hoc classification of affirmative versus negative sentences, separately for the abstract and concrete conditions, see Results 3.a; CP2: 6x2 factorial interaction; CP3: main effect of polarity; CP4: main effect of concreteness; CP5: main effect of fine-grained conceptual category). Classifications were performed between subjects in order to examine whether brain activation patterns were consistent

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across subjects, by means of a leave-one-subject-out cross-validation [24]. We report the mean cross-individual classification accuracies across all inclusive mask voxels (i.e. at the whole brain level).

In addition, we used searchlight analysis [30] with 4-mm radius spheres and a Gaussian Naive Bayes classifier [31] to localize anatomically the brain regions that significantly contributed to accurate discrimination of the different classification problems, as determined through a Monte Carlo permutation testing procedure. The condition labels were permuted in each sphere 1000 times [32], and the actual classification accuracy was then compared against the random permutation distribution with a declared $P < .001$ threshold. We report the mean classification accuracies across leave-one-subject-out cross-validations and the corresponding confusion matrices for the significant searchlight spheres.

For CP5, in order to gain a deeper insight in the brain coding of semantic information for the target conceptual categories, we adapted the procedure described in [24], which is based on recursive feature elimination and the sensitivity weights it provides. Sensitivity weights reflect the contribution of each voxel to the discrimination of one category from the others [33]. For each category, we calculated the spatial intersection of all pairwise sensitivity maps involving that category (e.g., for Ms: Ms-Em, Ms-Ma, Ms-Mo, Ms-Ha, Ms-Le). Before calculation of the category-specific intersections, the pairwise sensitivity maps were filtered for a minimum cluster extension of 20 voxels, only the clusters with sensitivity weights in the 9.5 highest percentile were retained, and smoothing with a 3-mm FWHM Gaussian kernel was applied. The sensitivity intersection maps were then inspected anatomically and projected to the AAL ROI atlas [34], in order to identify four distinct levels of semantic coding, derived from “hubs and spokes” models [2–3]: i) category-specific spokes: brain regions that were specific for just one particular category; ii) multi-category spokes: AAL ROIs presenting distinct and non-overlapping clusters for two or more category-specific intersection maps; iii) category-invariant hubs: AAL ROIs presenting clusters with spatial overlap from all the category-specific intersection maps; iv) graded hubs: AAL ROIs

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presenting both category-specific and overlapping clusters from all the category-specific intersection maps.

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3. Results

(a) CP1: 2x2 concreteness by polarity interaction

The four concreteness by polarity factorial combinations were classified with a mean accuracy of 35.42% across participants (chance level: $100\% / 4 = 25\%$) at the whole brain level. Searchlight permutation testing indicated that there were 135 searchlight spheres that significantly discriminated between the four factorial combinations ($P < .001$ against 1000 permutations), with a mean classification accuracy of 38.41% (Figure 1A). The significant searchlights were localized over extensive bilateral regions of the brain, including the medial and lateral frontal and parietal cortices, the anterior temporal lobes, the anterior and middle cingulate cortex, the caudate, and the cerebellum (Figure 1B, Table S2A).

Given successful discrimination of the 2x2 concreteness by polarity interaction, we further examined our experimental hypothesis that sentential negation modulates the modality-specific brain networks not only for concrete but also for abstract concepts, by post-hoc classification of affirmative versus negative sentences, separately for the abstract and concrete conditions.

For abstract concepts, affirmative and negative sentences were discriminated with a mean classification accuracy of 41.66% (chance level: $100\% / 2 = 50\%$). Classification accuracy in individual searchlights was more successful. Searchlight permutation testing indicated that there were 23 spheres that significantly discriminated between affirmative and negative sentences ($P < .001$ against 1000 permutations), with a mean classification accuracy of 76.32%, and the mean confusion matrix [34.35 13.65; 13.65 34.35].

For concrete concepts, affirmative and negative sentences were discriminated with a mean classification accuracy of 56.25% (chance level: $100\% / 2 = 50\%$). Again, classification accuracy in individual searchlights was more successful. Searchlight permutation testing indicated that there were 44 spheres that significantly discriminated between affirmative and negative sentences ($P < .001$ against 1000 permutations), with a mean classification accuracy of 76.91%, and the mean confusion matrix [34.82 13.18; 13.18 34.82].

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The abstract and concrete conditions partially differed with respect to the macro-anatomical distribution of the searchlights significantly discriminating between affirmative and negative sentences (Figure S1). For the abstract conditions, there was a unique involvement of the left temporal pole, right medial temporal lobe (amygdala, hippocampus, and parahippocampal gyrus), right precuneus, and cerebellum (Table S2B). Only the concrete conditions, in turn, involved the left posterior middle temporal gyrus, the left angular gyrus, the pars opercularis of the right inferior frontal gyrus, the right superior frontal gyrus, the calcarine and lingual gyri, and, bilaterally, the postcentral gyrus and the putamen (Table S2C).

(b) CP2: 6x2 fine-grained conceptual category by polarity interaction

The mean whole-brain accuracy for the classification of the 12 classes of sentences was 14.93% (chance level: $100\% / 12 = 8.33\%$). However, the confusion matrix showed a meaningless structure, that is, an inconsistently populated leading diagonal, and a disproportionately high rate of densely populated off-the-diagonal cells, representing incorrect predictions-to-target correspondences (Figure S2). Therefore, no further analysis of CP2 was carried out.

(c) CP3: Main effect of polarity

Affirmative and negative sentences were discriminated with a mean classification accuracy of 62.50% (chance level: $100\% / 2 = 50\%$). Classification accuracy in individual searchlights was more successful. Searchlight permutation testing indicated that there were 47 spheres that significantly discriminated between affirmative and negative sentences ($P < .001$ against 1000 permutations), with a mean classification accuracy of 75.66%, and the mean confusion matrix [33.28 14.72; 14.72 33.28]. These searchlights were localized in the left dorsolateral and medial frontal cortex, anterior and middle cingulate gyrus, precuneus and calcarine gyri, left caudate nucleus, right putamen and left thalamus, and cerebellar hemispheres (Figure 1B, Table S3).

(d) CP4: Main effect of concreteness

Abstract and concrete sentences were discriminated with a mean classification accuracy of 83.33% (chance level: $100\% / 2 = 50\%$) at the whole brain level. Searchlight permutation testing

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indicated that there were 464 spheres that significantly discriminated between abstract and concrete sentences ($P < .001$ against 1000 permutations), with a mean classification accuracy of 77.82%, and the mean confusion matrix [34.9 13.1; 13.1 34.9]. These searchlights were localized in the left dorsolateral frontal cortex and, bilaterally, in the anterior, posterior, and medial temporal lobe, the angular gyrus, and the cerebellum (Figure 1B, Table S4).

(e) CP5: Main effect of fine-grained conceptual category

The mean whole-brain accuracy for the classification of the fine-grained conceptual categories was 52.78 % (chance level: $100\% / 6 = 16.67\%$) (Figure 2A). Searchlight permutation testing yielded a much lower mean classification accuracy (18.17%), indicating that the to-be-discriminated patterns distinguishing between the six categories are sparse and distributed over broad neural territories.

To further investigate this broadly distributed category-specificity we applied an alternative whole-brain approach based on sensitivity weights, yielded by recursive feature elimination. We calculated all pairwise classifications among the six categories. For each and every pair, the mean classification accuracy was well above the 50% chance level, with a significant rate of correct predictions-to-targets correspondences ($P < .001$ for all pairs) (Table S5). Based on these successful pairwise classifications and on the corresponding sensitivity weights' anatomical distribution, we identified brain regions which might be ascribed to four distinct levels of semantic coding, reflecting multiple representations (Figure 2B, Table S6, see also Methods 2.f.ii): i) brain regions that were specific for just one particular category ("category-specific spokes"), and ii) regions presenting distinct and non-overlapping clusters for two or more categories ("multi-category spokes"): both these coding levels were sparsely distributed over large portions of the heteromodal cortices in both hemispheres, including the cerebellum; iii) regions of spatial overlap of all categories ("category-invariant hubs"): these were identified in the left superior temporal gyrus / Heschl's gyrus, right fusiform gyrus, and the cerebellar vermis; iv) regions presenting both spatial

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overlap and specific clusters for all six categories (“graded hubs”): these were identified in the left insula, left anterior middle temporal gyrus, and right precuneus.

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4. Discussion

This study investigated how the neural processing of abstract and concrete concepts expressed at the sentence level is modulated by negation polarity. Sentential negation polarity is thought to operate at the syntax-semantic interface [35–36], thus representing a case of interplay between a linguistic contextual operator and conceptual representations. Our main intent was to provide a proof of concept for the hypothesis that the neural networks supporting semantic representations are flexibly modulated by the linguistic sentential context [18]. We expanded on two different lines of research. A first line indicating that abstract and concrete concepts, and their respective fine-grained sub-categories, are distinctively encoded in distributed brain networks including category-invariant and category-specific nodes [3–4]. A second line, so far limited to concrete action-related concepts, indicating that sentential negation modulates neural activity of category-specific conceptual representation nodes [14–17]. Our sentence processing fMRI study sought generalization evidence for the modulatory effects of sentential negation by factorial combination of fine-grained abstract and concrete conceptual categories and affirmative/negative polarity.

By applying MVPA across participants, we demonstrated that the fMRI data contained sufficient information to discriminate between affirmative abstract, negative abstract, affirmative concrete, and negative concrete sentences resulting from the 2x2 concreteness by polarity interaction (CP1). Post-hoc classifications showed that affirmative and negative sentences were discriminable also when abstract and concrete conditions were analysed separately. However, when considering a 6x2 factorial combination between polarity (affirmative vs. negative) and fine-grained abstract (Ms, Em, Ma) and concrete (Mo, Ha, Le) categories (CP2), MVPA failed to accurately discriminate the associated brain activation patterns. Lack of sentential negation modulation on fine-grained semantic categories might be due to methodological aspects. In fact, analyses related to CP2 required separately modelling each of the 12 experimental conditions. This was not the case for CP1, where data were averaged across multiple experimental conditions. It is possible that the

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number of trials for each experimental condition in our study was not sufficient to ensure full-fledged category by polarity MVPA separation in CP2. However, this result might also constitute true negative evidence, weakening the conclusions drawn in the present study, and this important limitation needs to be considered. To sum up, the expectations of our proof of concept on negative versus affirmative linguistic contexts were fully met at a level of a simple but conceptually relevant distinction between concrete and abstract sentences, suggesting that the effect of sentential negation is not only limited to concrete action-related concepts but also extends to abstract ones. Furthermore, successful classifications were observed when considering the main effects nested in our factorial manipulation. Concerning sentential negation polarity, the MVPA yielded accurate whole-brain classification of the neural patterns associated with affirmative versus negative sentences although classification accuracy in individual searchlights was more successful (CP3). More robust classification accuracy was observed for concreteness (CP4), and fine-grained conceptual category (CP5) main effects.

Anatomical localization of condition-specific fMRI activation patterns is not straightforward in MVPA, since decoding is blind to the spatial organization of these patterns [37]. Nevertheless, MVPA localization techniques such as searchlight analysis [30], in combination with independent meta-analytic evidence on the brain functional organization of semantic processing, can provide useful information on the brain regions where contextual sentential negation modulations may occur. In CP1, CP3, and CP4 the searchlight analysis revealed an involvement of regions broadly distributed over both hemispheres, with a high degree of spatial overlap across these three classification problems (Figure 1B, Tables S2A, S3, S4). The classification problems CP1 and CP3, which are united by the manipulation of syntactic polarity, showed overlap in the pars triangularis of the inferior frontal gyrus, the basal ganglia (notably, the left caudate nucleus), and the anterior and middle cingulate cortex. Both the pars triangularis of the inferior frontal gyrus [38–39] and the left caudate nucleus [40–41] have been consistently implicated in syntactic structure processing, that is, word order computation at the sentence level above and beyond the specific issue of

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sentential negation. Of even greater relevance are previous univariate fMRI studies specifically investigating negation at the syntactic level, independently of the meanings on which it operates, which found an involvement of the basal ganglia [14] and of the left pars triangularis [22], among a set of other brain regions not identified in the present study. However, the inclusion of the anterior and middle cingulate cortex as a region of overlap between CP1 and CP3 prompts another possible functional interpretation. The anterior/middle cingulate and the left pars triangularis are known to be key components of the cognitive control network, also referred to as multiple-demand system, which is recruited by language, including syntactic tasks [42]. The left caudate nucleus, in turn, is known to be crucially involved in language monitoring and control [43]. It is therefore possible that these three brain regions jointly contribute to a cognitive control system that may help regulating the contextually-driven modulatory effects of negation on conceptual representations. The syntactic and cognitive control interpretation need not be mutually exclusive: the neural circuit activations underlying syntactic and cognitive control functions may be spatially and temporally distinct, but display the observed spatio-temporal overlap due to the relatively low spatial and temporal resolution of the fMRI data, and to the multivariate analysis technique. Our results may therefore indicate that both syntactic computation and cognitive control are involved in processing negation polarity at the sentence level.

In turn, the classification problems CP1 and CP4, which are united by the manipulation of semantic concreteness, showed overlap in an extended set of regions which is more consistent with a semantic functional role. It included the bilateral anterior temporal lobe, which has been suggested to serve as the main brain’s semantic hub [3,44]. But it also included, bilaterally, areas in the ventral posterior and medial temporal cortex (fusiform and parahippocampal gyri), parietal cortex (angular and supramarginal gyri, precuneus), and frontal lobe (inferior frontal gyrus, ventro- and dorso-medial prefrontal cortex), which, together with the anterior temporal lobe, have been postulated to form an extended semantic hub network based on the results of extant metaanalyses of fMRI studies [13,45–47].

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The broad involvement of brain regions revealed by CP1, CP3, and CP4 searchlight analysis also comprised modality-specific areas, which are thought to contribute to the discrimination of concrete and abstract concepts in embodied and grounded cognition accounts [4–5]. These included areas in sensory-motor systems, such as the visual and auditory cortices, the post-central gyrus, and supplementary motor area, and in the limbic system, particularly the amygdala. These and additional modality-specific brain regions also emerged from the post-hoc classification of affirmative and negative sentences, separately for abstract and concrete concepts. For abstract concepts, the discrimination of affirmative versus negative sentences mainly occurred in regions in the limbic system, including the amygdala. A specific modulation of this system by processing abstract sentences is in line with previous evidence showing that abstract concepts do have an emotional connotation. It is possible that sentential negation reduced access to this emotional content, dampening amygdala activation [48]. For concrete concepts, in turn, modality-specific brain regions that contributed most to the discrimination of affirmative versus negative sentences included the post-central gyrus, and the visual cortices (calcarine and lingual gyrus), along with portions of the left posterior middle temporal gyrus previously ascribed to the processing of concrete concepts, such as action-related sentences [49], manipulable objects [50], or manipulation and visual motion features [51]. The modulation of the left posterior middle temporal gyrus by affirmative versus negative concrete sentences confirms previous results on negation of action-related sentences [14], although only partially since in that study polarity modulations extended to the left premotor-parietal action representation system. Altogether, the distinct effects of negation polarity on abstract versus concrete sentences in the present study are largely compatible with our main hypothesis that sentential negation modulates modality-specific semantic brain regions.

The “hubs and spokes” organization of conceptual knowledge also emerged from the pattern of results obtained in the analysis of the fine-grained conceptual categories as a main effect (CP5). CP5 replicated previous studies [23–24] with respect to both the successful discrimination between fine-grained conceptual categories, and the anatomical sparseness of the involved brain regions. In

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particular, we replicated the findings in [24] by showing that category-specific and multi-category spoke regions were sparsely distributed over sensory-motor, limbic, and heteromodal cortices. As in [24], however, we didn't observe consistent correspondences between the conceptual categories and the experiential systems in which their representation is thought to be grounded according to grounded cognition accounts (e.g., somatotopically organized motor areas for Mo, Ha, and Le action-related sentences). The sensitivity analysis also identified a number of candidate hub brain regions, albeit with the notable exclusion of the ventral anterior temporal lobe, for which the conventional acquisition protocol used in the present study may have lacked signal detection sensitivity, as compared to other studies (e.g., [52] using an optimised fMRI acquisition protocol [53]. Some of the identified candidate hub regions (i.e. the right fusiform gyrus, the left superior temporal gyrus / Heschl's gyrus, and the cerebellar vermis) presented complete spatial overlap in the voxels that contributed to discriminate between the fine-grained conceptual categories. Based on this characteristic, the "category-invariant hubs" as we called them (see Methods 2.f.ii), may be assimilated to the class of semantic hubs. This appears quite straightforward in the case of the fusiform gyrus, since it has been identified as part of the "conceptual hub" network in a meta-analysis of neuroimaging studies of semantic processing [13,45]. Less straightforward, however, is the case of the superior temporal gyrus/Heschl's gyrus, which has been usually considered to play a role in speech perception and phonological rather than semantic processing (but see [45]). As for the cerebellum, there is increasing evidence of its contribution to language functions and, more specifically, to semantic processing tasks, such as predictive processing during language comprehension, lexical-semantic associations, and mapping of novel words onto existing concepts [54–55]. However, there is no consensus so far as to what the specific cerebellar contribution to semantic processing might be [55].

Some other candidate hub brain regions (i.e. the left insula, the left anterior middle temporal gyrus, and the right precuneus) contained, in addition to voxels with complete spatial overlap, specific voxels for each of the six categories, thus contributing to discriminate all the categories

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from each other. These hubs rather conform to the class of “graded hubs”, namely brain regions that present a functionally graded organization, possibly reflecting the varying pattern of connectivity links with category-specific spoke layers [3,24], such as the graded organization for abstract and concrete concepts in the anterior temporal lobe [52].

As discussed so far, the neural code patterns for the interaction between sentential negation polarity and abstract/concrete semantic categories that emerge from our results appear largely compatible with “hubs and spokes” accounts of semantic processing. Hub regions highly interconnected with primary and secondary sensory-motor spoke regions may be optimally suited to compute semantic representation by integrating information from different modalities [13,51], and also to combine multiple semantic representations, thus producing semantic meanings at the sentence level [2,13]. The interplay between hubs and spokes might therefore account for the distinction between abstract and concrete concepts, which result from the differential integration of information from multiple modalities (e.g., sensory and motor information for concrete concepts, versus emotional, introspective, social, and linguistic information for abstract concepts [5, 7–10]). The linguistic context, such as the one investigated here (i.e., sentential negation), might modulate such interplay between hubs and spokes and therefore modulate the representation of abstract and concrete concepts. Still, the exact information processing and coding dynamics occurring between hub and spoke brain regions remain relatively unclear. In one view, the anterior temporal lobe is put forward as the sole brain’s semantic hub [3], but with multiple subregions that have been suggested to code for different semantic categories or representational modalities, based on cytoarchitectonic, anatomical, and functional connectivity data [3]. In another view, semantic representations are instead served by relatively undifferentiated but multiple hubs [2], including some of the candidate hub regions identified in the present study, such as the medial prefrontal cortex, the anterior and posterior cingulate, the insula, the lateral parietal cortex, and the precuneus. Our findings are suggestive of semantic “hubs and spokes” dynamics that incorporate features from both views. On the one hand, we found multiple hub brain regions located in the frontal, temporal, and parietal

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lobes. On the other hand, while some of the multiple hub regions were undifferentiated (i.e. category-invariant hubs), some other presented with category-specific subregions for all types of abstract and concrete concepts included in our study (i.e. graded hubs). Our results indicate that combinatorial meaning specificities produced by the interaction between linguistic context and conceptual category emerge from activation patterns across multiple voxels distributed in unimodal spokes, as well as in undifferentiated and graded hubs.

Conclusions

The overall picture emerging from this set of results is largely compatible with the view that sentential negation polarity operates by modulating the neural activation patterns coding for concepts, to an extent that is sufficient to make the processing of affirmative and negative sentences with either an abstract or concrete content distinguishable by means of MVPA. We suggest that the negation polarity modulation occurs in distributed representational semantic networks, through the functional mediation of syntactic and cognitive control systems.

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Ethics

The study was approved by the Ethics Committee of the San Raffaele Scientific Institute, Milan, Italy. Subjects gave written consent for their participation in the study.

Data Accessibility

The data and code that support the findings of this study are publicly available at <https://doi.org/10.6084/m9.figshare.5492698.v2>.

Authors' Contributions

MG, MMV and MT conceived and designed the experiments; MG, MMV and MT performed the experiment; MG, KH and MT analysed the data; MG, KH, MMV and MT wrote the manuscript. All authors approved the final version of the manuscript.

Competing Interests

We have no competing interests.

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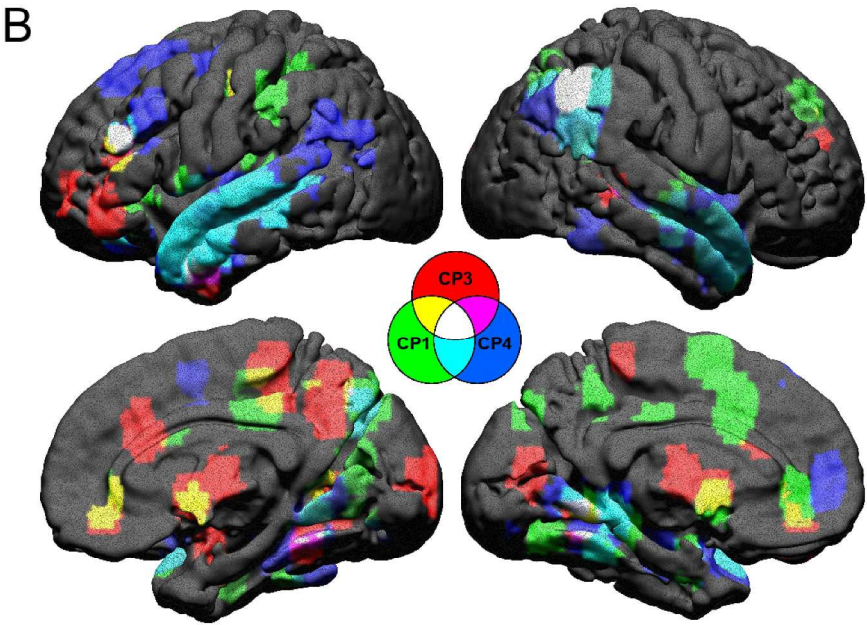
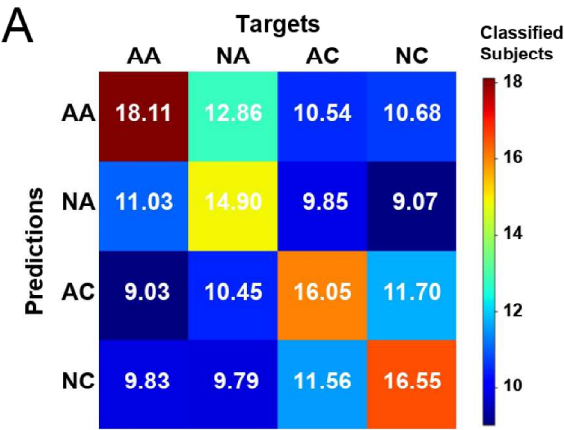
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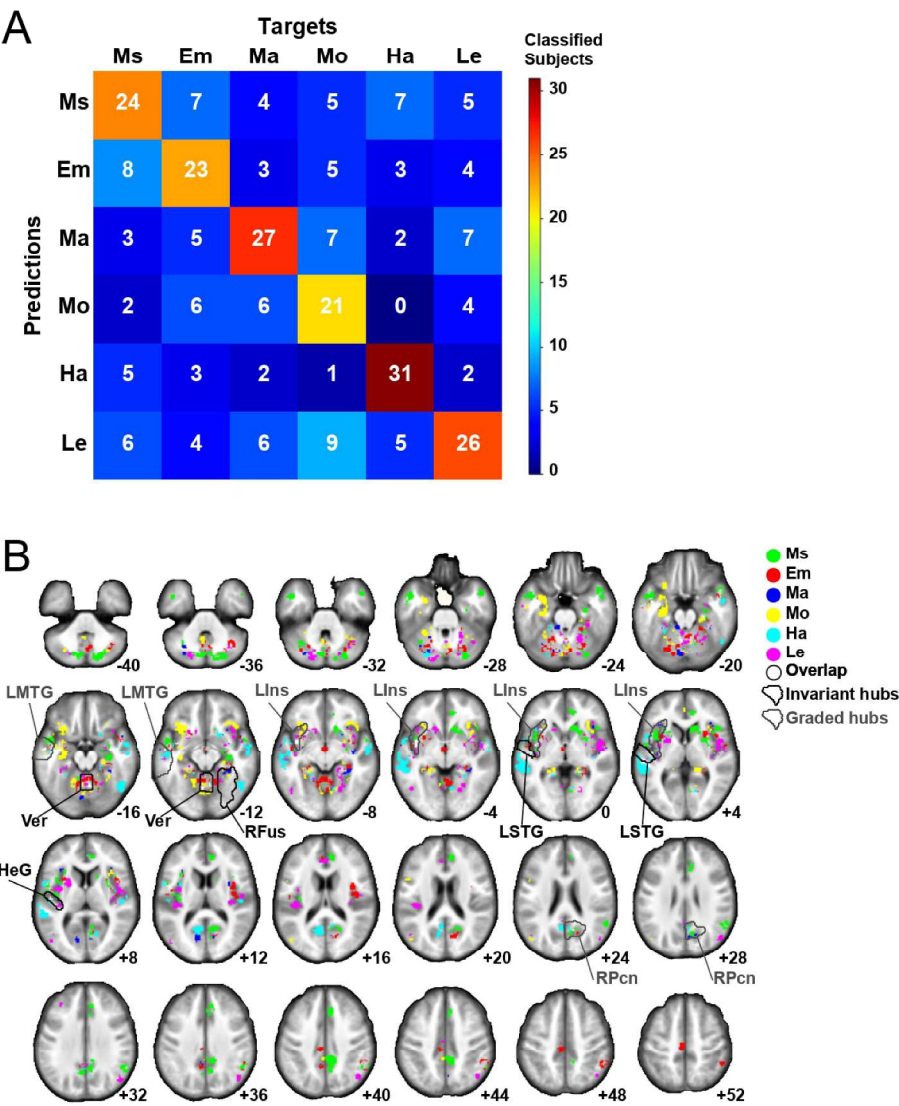
Figure legends

Figure 1. MVPA classification of the main effects of sentential negation polarity, concreteness, and their 2x2 factorial interaction. A) Confusion matrix for CP1, featuring the means of the classified spmT maps across the significant ($P < .001$, against 1000 permutations) searchlight spheres. Cell numbers represent the mean number of subjects that were either classified correctly (diagonal) or incorrectly (off the diagonal), with respect to the four concreteness by polarity factorial combinations (AA: affirmative abstract; NA: negative abstract; AC: affirmative concrete; NC: negative concrete). B) Anatomical localization of the searchlight spheres yielding significant ($P < .001$, against 1000 permutations) classification accuracy for CP1, CP3, and CP4. Colour codes are indicated by the colour palette inset. The effects are displayed on lateral and medial wall surface renderings of the average anatomical image of all participants. Left and right hemispheres are displayed according to the neurological convention.

Figure 2. MVPA classification of the main effect of fine-grained conceptual category. A) Confusion matrix for CP5, with cell numbers featuring the number of subjects that were either classified correctly (diagonal) or incorrectly (off the diagonal) at the whole brain level, with respect to the six fine-grained conceptual categories. B) Category-specific sensitivity intersection maps, representing the anatomical regions that consistently allowed us to correctly discriminate each category from the other five categories, in a pairwise fashion. The category-specific intersections are displayed on axial slices (z coordinate levels indicated in mm) of the average anatomical image of all participants (neurological convention). Anatomical region boundaries are displayed for AAL ROIs classified as “category-invariant hubs” (black colour, LHeG: left Heschl's gyrus; LSTG: left superior temporal gyrus; RFus: right fusiform gyrus; Ver: cerebellar vermis) or “graded hubs” (grey colour, LMTG: left middle temporal gyrus; LIns: left insula; RPcn: right precuneus).



319x392mm (300 x 300 DPI)



310x370mm (300 x 300 DPI)