Visceromotor roots of aesthetic evaluation of pain in art: an fMRI study

Martina Ardizzi,1,2 Francesca Ferroni,1,2 Maria Alessandra Umiltà,2,3,4 Chiara Pinardi,5 Antonino Errante,1 Francesca Ferri,6 Elisabetta Fadda,2,7 and Vittorio Gallese1,2,4

1Department of Medicine and Surgery, University of Parma, 43126, Parma, Italy, 2Neuroscience & Humanities Lab, University of Parma, 43125, Parma, Italy, 3Department of Food and Drug, University of Parma, 43124, Parma, Italy, 4Department of Art History Columbia University, Italian Academy for Advanced Studies, Columbia University, 10027, New York, NY, USA, 5Department of Neuroradiology, Fondazione IRCCS Istituto Neurologico Carlo Besta, 20133, Milan, Italy, 6Department of Neuroscience, Imaging and Clinical Science, University G. d’Annunzio, 66100, Chieti, Italy, and 7Department of Humanities, Social Sciences and Cultural Industries, University of Parma, 43125, Parma, Italy

Correspondence should be addressed to Martina Ardizzi, Department of Medicine and Surgery - Unit of Neuroscience, University of Parma, Via Volturno, 39/E, Parma, 43121, Italy. E-mail: martina.ardizzi@unipr.it, ardizzi.martina@gmail.com.

Abstract

Empathy for pain involves sensory and visceromotor brain regions relevant also in the first-person pain experience. Focusing on brain activations associated with vicarious experiences of pain triggered by artistic or non-artistic images, the present study aims to investigate common and distinct brain activation patterns associated with these two vicarious experiences of pain and to assess whether empathy for pain brain regions contributes to the formation of an aesthetic judgement (AJ) in non-art expert observers. Artistic and non-artistic facial expressions (painful and neutral) were shown to participants inside the scanner and then aesthetically rated in a subsequent behavioural session. Results showed that empathy for pain brain regions (i.e. bilateral insular cortex, posterior sector of the anterior cingulate cortex and the anterior portion of the middle cingulate cortex) and bilateral inferior frontal gyrus are commonly activated by artistic and non-artistic painful facial expressions. For the artistic representation of pain, the activity recorded in these regions directly correlated with participants’ AJ. Results also showed the distinct activation of a large cluster located in the posterior cingulate cortex/precuneus for non-artistic stimuli. This study suggests that non-beauty-specific mechanisms such as empathy for pain are crucial components of the aesthetic experience of artworks.

Key words: art; anterior insula; cingulate cortex; inferior frontal gyrus; pain

Introduction

Neuroimaging studies demonstrate that the observation of other individuals’ somatosensory experiences or emotional facial expressions recruits sensory, premotor and visceromotor brain areas involved also in the first-person experience of the same state (Carr et al., 2003; Wicker et al., 2003; Ebisch et al., 2008, 2011; Caruana et al., 2017). Our hypothesis is that this common neural ground for experiencing and observing emotions...
also serves the formation of an aesthetic judgement (AJ) when the emotional content of the artistic work is represented at a figurative level. To address our hypothesis, we focus on empathy for pain triggered by artistic and non-artistic painful facial expressions. Among the numerous cerebral regions active both during the first-person experience of pain and the mere observation of others’ facial expressions of pain, the bilateral anterior insula (AI)/fronto-insular cortices and the cingulate cortex (CC)—especially in its anterior medial (mACC) and posterior anterior (pACC) sectors—are, indeed, consistently identified (Botvinick et al., 2005; Lamm et al., 2011). This functional overlap suggests that empathy for pain is underpinned by visceromotor and viscerosensitive neural structures that are also involved in the direct experience of pain. Indeed, the insular cortex is characterized by a clear anatomical and functional caudal-to-rostral gradient of increasingly complex integration of bodily signals. Whereas the direct experience of pain is somatotopically mapped only in the posterior insular subdivision associated with the sensory components of nociception (Ostrowsky et al., 2002; Kurth et al., 2010), visceromotor and viscerosensitive responses are related to the activation of the AI, induced both during the actual experience of pain and by the observation of others’ pain facial expressions (Ostrowsky et al., 2000; Lamm et al., 2011; Benuzzi et al., 2018). Interestingly, although far from being a pain-specific region (Kurth et al., 2010), when the AI is damaged by brain lesions, patients worsen their recognition of another person’s pain experience, suggesting its causal involvement in other’s pain detection (Gu et al., 2012). At the same time, mounting evidence supports the role of the pACC and mACC in autonomic and motor control. A recent study (Caruana et al., 2018) demonstrates that the intracerebral high-frequency electrical stimulation of the mACC in a large cohort of drug-resistant epileptic patients triggers a variety of goal-oriented and defensive behaviours involving the upper limbs or the entire body. Differently, the adjacent pACC appears to be involved in the production of facial emotional displays and autonomic responses identified by the patients as fear and anxiety (Caruana et al., 2018). Coherently, a study conducted on rats’ homologous mesial regions found neurons responding both when rats experience pain, as triggered by a laser, and while they witness another rat receiving shocks (Carrillo et al., 2019). These authors also demonstrate that the deactivation of this region reduces rats’ nociceptive behaviours (i.e. freezing) while observing a conspecific experiencing painful shocks.

The existence of a common neural ground for experiencing and observing emotions has been mainly investigated as the functional mechanism underpinning the recognition of others’ emotions (Gallese et al., 2004; Gallese and Sinigaglia, 2011; Gallese, 2014). However, this functional mechanism has been proposed to also play a role in other contexts. For example, when reading single words with threat connotation, not only proposed to also play a role in other contexts. For example, the faculty of beauty seems to recruit the medial orbitofrontal (Ishizu and Zeki, 2011; Ishizu, 2014; Zeki et al., 2014) and the dorsolateral prefrontal cortices (Cela-Conde et al., 2004; Cattaneo et al., 2014). This brain-based approach to the study of aesthetics suggests that all works of art that appear beautiful to a subject affect the activity of specific brain regions (Ishizu and Zeki, 2011). Another approach is to investigate whether neural circuits that we know are linked to functions not associated with the experience of beauty (e.g. empathy for pain brain regions) can also play a role in the formation of aesthetic experience. Among these non-beauty-specific neural mechanisms possibly underpinning the aesthetic power of images, convergent studies propose the observer’s sensory and visceromotor engagement with images as a valid candidate. Behavioural studies show that observer’s simulation of artists’ creative gestures increases the aesthetic evaluation of paintings made with congruent hand movements (Leder et al., 2012; Ticini et al., 2014; McLean et al., 2015). Direct demonstration of the link between activation in premotor cortices and AJ can be found in studies investigating the neural correlates of dance enjoyment (for a review, see Kirsch et al., 2016). In a pioneering study, Calvo-Merino and colleagues (2008) found a significant activation in the premotor cortex during passive viewing of dance stimuli that was related to the subsequent aesthetic evaluation of the same stimuli. More recently, using stimuli depicting static or dynamic representational paintings of human figures or landscapes, a link—mediated by dynamism impression—between the amplitude of observers’ motor evoked potentials and their liking judgements has been demonstrated (Fiori et al., 2020). Coherently, the AJ of landscape paintings involves the posterior and central sectors of the insular cortex, in relation to the intrinsic dynamism of the artwork (Di Dio et al., 2016). Also, observers’ sensorimotor engagement with portrayals of painful facial expressions influences their explicit AJs (Ardivizi et al., 2020a). Specifically, it has been found that the overt contraction of the corrugator supercilii facial muscle increased the aesthetic rating of artistic facial expressions of pain, where the contraction of the same facial muscle was visible. This latter result seems to support Schott’s claim about the potential engagement of empathy for pain brain regions (i.e. sensory, premotor and visceromotor brain areas) during the aesthetic appreciation of pictorial representation of pain (Schott, 2015). Until now, no studies have directly explored this hypothesis. Only two neuroimaging studies not specifically interested in aesthetic appreciation but using pictorial representation of injured bodies (De Gelder et al., 2018) or mourning scenes (Labek et al., 2017) offered mixed results in support of the recruitment of empathy for pain brain regions during the enjoyment of such artistic stimuli.

In the present functional magnetic resonance imaging (fMRI) study, we investigate the brain activations related to the vicarious pain experience triggered by artistic and non-artistic stimuli with the following aims: (i) to elucidate whether the observation of artistic facial expressions of pain is able to activate the brain regions normally associated with empathy for pain; (ii) to understand whether this specific activation pattern could be involved in the AJ of the same images and (iii) to verify whether the two vicarious experiences of pain, one induced by art and the other aroused by non-artistic stimuli, evoke different brain activation patterns.

**Materials and methods**

**Participants**

Twenty healthy right-handed volunteers with no training in art or art history [11 females; mean age = 25.15, Standard Error (SE) = 0.68, mean schooling = 15.25, SE = 0.41; mean Art
Experience Questionnaire (Chatterjee et al., 2010) score = 12.5, SE 1.89] participated in the study. Handedness was assessed by means of the Edinburgh Inventory (Oldfield, 1971). All participants had normal or corrected-to-normal visual acuity. No participant had a history of neurologic, general medical or psychiatric conditions. The experimental protocol was approved by the Ethics Committee of the University of Parma, and it was in line with the Declaration of Helsinki 2013. Written informed consents were collected from all participants.

Stimuli
Twenty-four high-resolution digital versions of neutral (N = 12) and painful (N = 12) facial expressions were used as experimental stimuli. Half of the stimuli were selected from Renaissance and Baroque paintings [Art Pain (AP) stimuli, N = 6; Art Neutral (AN) stimuli, N = 6], whereas the other half derived from non-artistic digital photographs of models’ facial expressions [non-Art Pain (nAP) stimuli, N = 6; non-Art Neutral (nAN) stimuli, N = 6]. Stimuli selection followed recent guidelines for the use of artworks as stimuli in empirical research (Hayn-Leichsenring, 2017). Please see Supplementary Material for a detailed description of the procedure followed to select images and validate the final set of stimuli.

Experimental design
The experimental protocol consisted of two sessions (see Figure 1):

fMRI session. Participants lay in the scanner in a dimly lit environment. The stimuli were viewed via digital visors (VisuaStim) with a 500,000 pixel x 0.25 square inch resolution and horizontal eye field of 30°. The digital transmission of the signal to the scanner occurred via optic fibre. The software E-Prime 2 Professional (Psychology Software Tools, Inc., Pittsburgh, USA, http://www.pstnet.com) was used both for stimuli presentation and the recording of participants’ answers.

Participants were instructed to indicate, on the appearance of the task question, if the face depicted showed an expression of pain or not. Responses were given using the index or medium fingers of their right hands. Options’ order was balanced across participants congruently with the screen that appeared in the scanner (i.e. ‘Pain: Yes or No’; ‘Pain: No or Yes’).

The experiment consisted of six runs lasting 7 minutes each. The total duration of the entire experiment was approximately 50 minutes. Each run consisted of 24 randomized trials, six for each condition (i.e. AP, AN, nAP and nAN), constituting the event-related fMRI design. Each stimulus was presented six times across the six runs. Each trial began with a central fixation cross (ranging from 10 to 15 sec; i.e. implicit baseline of the subsequent functional analyses) followed by stimuli presentation lasting 2.5 sec. After stimulus presentation, the task question (i.e. ‘Pain: Yes or No’ or ‘Pain: No or Yes’) lasting 2.5 sec appeared. Overall, the experiment consisted of 144 trials, 36 for each condition. Before the beginning of the fMRI session, an out-of-scanner training of eight stimuli (two for each condition), different from those showed in the following scan session, was administered to ensure that participants understood the instructions and became familiar with timing and also with the use of the dial.

Behavioural session. Immediately after the fMRI session, the stimuli were shown again in the AJ task administered outside the scanner. Participants were asked to answer the question ‘How artistically beautiful do you think this image is?’ using a 5-point ordinal scale ranging from ‘not at all’ (1) to ‘extremely beautiful’ (5). AJ task consisted of 144 randomized trials, 36 for each condition. Each trial began with a central fixation cross lasting 0.5 sec followed by stimulus presentation lasting 2.5 sec. After this period, task question and ordinal scale appeared. The next trial began after participants’ no-time-limit answers. The entire duration of AJ task was approximately 12 minutes, depending on participants’ response time. Lastly, participants were required to respond in a yes/no forced choice task whether they had seen the stimuli before the study. All participants reported that they had not seen any of the images before (100% unfamiliarity rating).

fMRI data acquisition
Anatomical T1-weighted and functional T2*-weighted MR images were acquired with a 3-Tesla General Electric scanner equipped with an 8-channel receiver head coil. Functional images were acquired using a T2*-weighted gradient-echo, echoplanar (EPI) pulse sequence (acceleration factor asset = 2, 40 sequential transverse slices covering the whole brain, with a Repetition Time (TR) time of 2.5 sec, Echo Time (TE) = 30 msec, flip angle = 90°, Field of View (FOV) = 205 x 205 mm², inter-slice gap = 0.5 mm, slice thickness = 3 mm, in-plane resolution = 2.5 x 2.5 x 2.5 mm³). At the end of the six functional runs, a T1-weighted anatomical scan (acceleration factor arc = 2, 156 sagittal slices, matrix 256 x 256, isotropic resolution 1 x 1 x 1 mm³, Time to Invert (TI) = 450 msec, TR = 8100 msec, TE = 3.2 msec, flip angle = 12°) was acquired for each participant.

Fig. 1. Experimental design and stimuli. In the fMRI session, participants judged if the face depicted in the stimulus showed an expression of pain or not. In the behavioural session, performed outside the scanner, participants were asked to express an AJ on a 5-point ordinal scale. Four exemplificative stimuli are displayed in the right panel of the figure.
Statistical analyses

Data analysis was performed with SPM12 (Statistical Parametric Mapping software; The Wellcome Department of Imaging Neuroscience, London, UK; http://www.fil.ion.ucl.ac.uk) running on MATLAB R2017b (The Mathworks, Inc., Natick, MA) and IBM SPSS statistics 24. The first four volumes of each run were discarded to allow for T1 equilibration effects. For each participant, all volumes were corrected for slice timing using the middle slice as reference. Then, all volumes were spatially realigned to the first one of the first session and un-warped to correct for between-scanner motion, and a mean image from the realigned volumes was created. T1-weighted images were co-registered to the mean fMRI volume; then segmented into grey, white and cerebrospinal fluid; and finally spatially normalized to the Montreal Neurological Institute (MNI) coordinates system. The thereby derived spatial transformation by co-registered T1 normalization was applied to the realigned EPI volumes, which after normalization were re-sampled in 1 × 1 × 1 mm³ voxels using trilinear interpolation in space. All functional volumes were then spatially smoothed with a 6 mm full-width half-maximum isotropic Gaussian kernel for the group analysis.

Data were analysed using a random-effects model (Friston et al., 1999), implemented in a two-level procedure. In the first level, single-subject fMRI responses were modelled in a General Linear Model by a design matrix comprising the onsets and durations of each event for each functional run (AP, AN, nAP, nAN and Response). Trials erroneously identified by participants were regressed separately. No participant exceeded the 16% of incorrect trials identified. This analysis employed event-related convolution models using the hemodynamic response function provided by SPM12. The presentation of the stimuli for each trial condition was modelled as mini-epoch lasting 2.5 sec, whereas the motor response was modelled as one punctual single event. Motion regressors were included to control for possible artefacts related to head motion. For all participants, head motion never exceeded 3 mm.

In the second-level analysis (group analysis), corresponding contrast images from the first level for each participant were entered into flexible analysis of variance (ANOVA) with sphericity correction for repeated measures (Friston et al., 2002). This model considered the patterns of activation obtained for the main effects—Emotion (Pain, Neutral) and Content (Art, Non-Art)—as well as the interaction between the two factors (Emotion × Content). All results were thresholded at $P < 0.05$ family wise error (FWE) corrected. The main effect of Emotion revealed a quite extensive activation in a midline cluster encompassing the cingulate cortices and the insular cortices (ROI rAI: $t_{19} = 0.7$, $P = 0.001$; ROI IAI: $t_{19} = 0.6$, $P = 0.006$; ROI CC: $t_{19} = 0.48$, $P = 0.04$) (Figure 4). Differently, Pearson’s correlation analysis performed between $\Delta A_{IA}$ and the beta weights extracted from the Fusiform Face Area was not significant (control ROI FFA: $t_{19} = −0.009$, $P = 0.97$).

Three regions of interest (ROIs) were created, based on a previous meta-analysis study (Lamm et al., 2011), on the left and right AI (IAI and rAI) and on the CC using MarsBaR Toolbox for SPM (release 0.44). All ROIs were defined centring the sphere (radius = 10 mm) around the maxima of these clusters (ROI rAI: $x = 39$, $y = 23$, $z = −4$; ROI IAI: $x = −40$, $y = 22$, $z = 0$; ROI CC: $x = −2$, $y = 23$, $z = 40$). These regions were reported as consistently activated across the 32 studies included both in the coordinate- and image-based meta-analysis (Lamm et al., 2011). Furthermore, the selected ROIs are also close to the activation foci associated with the main effect of Emotion in the present study. Mean beta weights associated with the contrast images, AP vs AN, of each participant were extracted using REX Toolbox (Duff et al., 2007). Coherent with this investigated contrast, ROIs’ mean beta weights were correlated with the change score between the mean AI attributed to AP and the same assigned to AN ($\Delta A_{IA} = A_{IA} − A_{AN}$). Three Pearson’s two-tailed correlation analyses were then performed between ROIs mean beta weights and $\Delta A_{IA}$.

The specificity of this effect was investigated by performing an additional correlation analysis between $\Delta A_{IA}$ and mean beta weights extracted from a control brain region not directly involved in other’s pain detection but related to face perceptual analysis, the Fusiform Face Area (FFA) (ROI FFA: $x = 42$, $y = −50$, $z = −19$; Cohen et al., 2019). One participant was removed from ROIs analyses due to technical problems in the recording of the responses in the behavioural session. Consequently, ROIs analyses were performed on 19 participants.

See Supplementary Material for the results of the AJ task.

Results

In the fMRI session, brain activity was measured as a function of Emotion (Pain, Neutral), Content (Art, Non-Art) and their interaction (Emotion × Content). fMRI results are listed in Table 1.

The main effect of Emotion revealed a quite extensive activation for painful facial expressions, irrespective of Content, in the pACC and aMCC partially extending to the medial frontal gyrus (MFG). Additional activations were present bilaterally in large clusters including the AI and inferior frontal gyrus (IFG); the latter is more extended into the right hemisphere (Figure 2, panel A).

The main effect of Content revealed significant activation of the posterior CC (PCC)/precuneus mainly evoked by non-artistic stimuli (Figure 2, panel B).

The interaction Emotion × Content showed increased activation in a cluster located in the right inferior occipital gyrus (rIOG) extending to the calcarine gyrus, with AP and nAN producing greater activations than AN and nAP (Figure 2, panel C).

The results of the conjunction analysis (AP ∩ nAP; Figure 3; Table 2) showed that both in AP and nAP conditions there were significant activations in a midline cluster encompassing the pACC, aMCC and MFG, as well as, in two bilateral clusters including the AI and the IFG.

Pearson’s correlation analyses performed between $\Delta A_{IA}$ and the beta weights extracted from the cingulum and the insular cortices were significant (ROI rAI: $r_{19} = 0.7$, $P = 0.001$; ROI IAI: $r_{19} = 0.6$, $P = 0.006$; ROI CC: $r_{19} = 0.48$, $P = 0.04$) (Figure 4). Differently, Pearson’s correlation analysis performed between $\Delta A_{IA}$ and the beta weights extracted from the Fusiform Face Area was not significant (control ROI FFA: $r_{19} = −0.009$, $P = 0.97$).

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Discussion

Empirical evidence has consistently demonstrated that the recognition of others’ facial expression of emotions and sensations, like pain, is underpinned by the activation of the brain regions that are active during the subjective experience of the same emotions and sensations. However, new evidence (Leder et al., 2012; Ticini et al., 2014; Kirsch et al., 2016; Flori et al., 2020; Ardizzi et al., 2020a) suggests that this basic access to others’ emotions may also feed other high-level processes, such as the aesthetic experience of works of art and the formation of AJs. In the present study, we better explored this hypothesis investigating the brain activations associated with vicarious experiences of pain triggered by artistic or non-artistic images depicting facial expressions of pain. First, our aim was to understand whether artistic pain representations engage observers’ empathy for pain brain regions. Second, we wanted to understand whether the possible recruitment of this common brain network for experiencing and observing pain helps the formation of the AJ of non-art expert observers. Third, we are totally aware that witnessing others’ pain looking at the picture of a real face or at the pictorial representation of the same facial expression are two different experiences. For this reason, we expected to also find distinctive brain activations for artistic and non-artistic facial expressions of pain.

Considering the first aim of the present study, activations derived from the main effect of Emotion showed the expected empathy for pain brain regions’ activations (i.e. bilateral AI and pACC and aMCC). Interestingly, the bilateral activations in the AI significantly extend to the IFG, especially in the right hemisphere. IFG plays an important role in the motor mirror-neuron mechanism that likely supports recognition and imitation of actions (Rizzolatti and Sinigaglia, 2016), but it is also specifically related to the observation/evaluation and execution of facial expressions of emotions (Carr et al., 2003; Hennenlotter et al., 2005; Van Der Gaag et al., 2007; Jabbi and Keysers, 2008). Indeed, patients with localized damages limited to IFG are selectively compromised in the recognition of facial expressions of emotions, but not in second-order false belief attribution (Shamay-Tsoory et al., 2009). A more recent meta-analysis investigating the specific brain areas subserving specific sub-processes of mindreading found convergent activations in the IFG when others’ internal states are inferred from their faces or eyes (Schurz et al., 2014). The authors concluded that IFG supports a particular form of mindreading made possible by a ‘common coding’ mechanism for action and perception, that is, the mirror mechanism. Coherently, in the context of painful facial expressions, IFG is involved in the detection of the emotional components of pain (Budeli et al., 2010, 2013).

In agreement with the results of the main effect of Emotion, the conjunction analysis performed between the whole-brain activations for AP and nAP stimuli revealed the activation of pACC/aMCC, together with bilateral AI/IFG clusters. The activation patterns revealed by the present study demonstrate that the activation of empathy for pain brain regions (i.e. AI, pACC and aMCC) and IFG can also be elicited by both artistic and non-artistic representations of facial pain. Other studies demonstrated that the observation of actions depicted in figurative works of art (Battaglia et al., 2011; Thakral et al., 2012) stimulates the responses of sensorimotor circuits also involved in actual motor control. However, for the first time, the present results provide evidence that the common neural network for actual and vicarious emotional experiences can also be elicited by art emotional content. When observers try to decode others’ painful facial expressions, be they produced by an expressive natural behaviour or created by intentional artistic practice, both lead to the vicarious activation of a set of brain areas relevant to the direct experience of the same emotional state.

We moved a step forward in the attempt to meet our second aim and link this neural activation pattern to the formation of AJ. Results show that the activations found in empathy for pain brain regions are also positively correlated with participants’ AJs attributed to artistic facial expressions of pain with respect to artistic neutral faces. In other words, the higher the response of these areas, the higher the AJ of artistic beauty of painful facial expressions. As expected, this relation is present only for empathy for pain brain regions and not for brain areas involved in face perceptual analysis but not in others’ pain detection. These findings support the idea that the activation of empathy for pain brain circuit and of IFG is involved in the AJ of beauty when the source of the emotional content is artistic. Even in the context of contemporary dance enjoyment, the activations of IFG and other mentalizing areas were associated with spectators’ grasping of dance coherence (Bachrach et al., 2016). Interestingly, Ishizu and

| Brain structure | Side | Cluster extent size in voxel (Ke) | P FWE corrected at the cluster level | Z | x | y | z |
|-----------------|------|----------------------------------|-----------------------------------|---|---|---|---|
| Main effect of Emotion | pACC/aMCC/MFG | R/L | 9623 | <0.001 | 5.58 | -1 | 28 | 39 |
| | AI/IFG | R | 8003 | <0.001 | 4.52 | 4 | 36 | 28 |
| | | | | | 4.04 | 9 | 20 | 56 |
| | AI/IFG | L | 5061 | <0.001 | 5.10 | 44 | 22 | -6 |
| | | | | | 4.55 | 39 | 11 | 45 |
| | | | | | 4.37 | 54 | 30 | 25 |
| Main effect of Content | PCC/precuneus | R/L | 1086 | 0.027 | 4.51 | 3 | -60 | 28 |
| Emotion × Content | IOG | R | 1357 | 0.009 | 5.36 | 31 | -93 | -5 |
| | | | | | 3.26 | 35 | -92 | -12 |

Results are thresholded at P < 0.05 FWE corrected at the cluster level. Local maxima are given in MNI standard brain coordinates. Most probable anatomical regions are derived from Anatomy Toolbox 1.7 (Eickhoff et al., 2005) and listed in ‘Brain structure’ column.
Zeki (Ishizu and Zeki, 2017) found that when people aesthetically rate sorrowful works of art, the brain areas involved in the empathic experience of other people’s sadness are functionally connected to regions implicated in the judgement of beauty, suggesting how empathic engagement and aesthetic experience are two interrelated phenomena.

In a similar vein, the present results on the aesthetic involvement of visceromotor and premotor brain regions related to the experience of pain suggest that the empathic engagement with works of art concerns a bodily-based direct access to art emotional content. Indeed, motor and physiological responses coherent with the artistic emotional climax are widely demonstrated across many forms of art (Lundqvist et al., 2009; Koelsch, 2014; Wassiliwizky et al., 2017; Siri et al., 2018; Kaltwasser et al., 2019; Ardizzi et al., 2020b). The absence of a significant correlation between FFA activation and AJ for artistic images outlines the specificity of such relation for empathy for pain brain regions and stimulates some considerations about the qualification of this functional mechanism in aesthetic appreciation. For example, Cattaneo and collaborators (Cattaneo et al., 2015) interfering with extrastriate area V5 suppressed both the perceived sense of motion and the liking of abstract but not of representational paintings. As in Cattaneo et al. (2015), where the dynamism perceived in abstract paintings drove their aesthetic appreciation, our results show that the direct access to the observed pain experience concurs with the AJ of painful facial expressions as represented in paintings. Overall, these results suggest that non-beauty-specific sensory and visceromotor engagement with images can contribute, under specific circumstances, to the formation of AJ, linking together the emotion-evaluation, sensorimotor and meaning-knowledge processes composing the aesthetic experience (Chatterjee and Vartanian, 2016; Ardizzi, 2020).

As stated before, the decoding of artistic facial expression and the decoding of non-artistic facial expressions of pain are not two perfectly overlapping processes. The results of the main effect of Content showed the activation of a large cluster located in the PCC/precuneus for non-artistic stimuli. These regions turned out to be responsive to a wide range of highly integrated tasks, including visuospatial imagery, visual information processing, episodic memory retrieval and self-processing operations, such as first-person perspective taking and the experience of agency (Cavanna and Trimble, 2006). Thanks to these functional activations, it is not surprising that the precuneus was seen to be consistently activated in mentalizing tasks (Molenberghs et al., 2016). The activation of these regions suggests the preferential recruitment of visual and self-referential processing when decoding others’ emotional states, as when portrayed by non-artistic facial expressions. The results of the interaction Emotion x Content showed interesting distinct activation. Artistic facial expressions of pain and non-artistic neutral facial expressions trigger the response of the rIOG. This region belongs to the distributed neural system devoted to face perception (Haxby et al., 2002; Rossion et al., 2012). The cluster here identified includes the occipital face area, a functionally defined face-selective area usually located in the lateral surface of the occipital lobe either in or in the vicinity of the IOG (Molenberghs et al., 2016). The activation of the rIOG in response to artistic facial expressions of pain and non-artistic neutral faces may suggest that the decoding of such stimuli takes advantage from a detailed analysis of facial visual proprieties.
Fig. 3. Brain activation map resulting from the conjunction analysis. The map is obtained from the conjunction between the contrasts AP vs baseline and nAP vs baseline, masked using an inclusive contrast image derived from the main effect of Emotion. Group-averaged statistical parametric maps are rendered into a standard MNI brain template and in three representative slices ($P < 0.05$ FWE corrected at the cluster level).

Table 2. Results of conjunction analysis

| Contrast   | Brain structure | Side | Cluster extent size in voxel (Ke) | P FWE corrected at the cluster level | Z  | x   | y   | z   |
|------------|-----------------|------|----------------------------------|--------------------------------------|----|-----|-----|-----|
| AP ∩ nAP   | pACC/aMCC/MFG   | R/L  | 2569                             | 0.001                                | 7.31| −5  | 13  | 47  |
|            |                 |      |                                  |                                       | 6.99| 7   | 16  | 43  |
|            |                 |      |                                  |                                       | 6.49| 7   | 15  | 52  |
| AI/IFG     |                 | L    | 1298                             | 0.022                                | 6.35| −29 | 27  | 3   |
|            |                 |      |                                  |                                       | 3.26| −37 | 16  | 2   |
| AI/IFG     |                 | R    | 1536                             | 0.01                                 | 5.83| 31  | 27  | 5   |

Results are thresholded at $P < 0.05$ FWE corrected at the cluster level. Local maxima are given in MNI standard brain coordinates. Most probable anatomical regions are derived from Anatomy Toolbox 1.7 (Eickhoff et al., 2005) and listed in ‘Brain structure’ column.

Fig. 4. Pearson’s correlation analyses conducted between mean beta weights and AJs. Mean beta weights were extracted from left (in blue) and right (in green) AI and from the CC (in red). As a control region, mean beta weights were obtained from the FFA (in orange). *$P < 0.05$.

A number of specific methodological choices were made in our paradigm. Here, a balance between a rigorous control over stimuli properties and the ecological power of artistic images was obtained through a meticulous and hypothesis-driven procedure of stimuli selection and validation, potentially limiting the study’s general validity. Despite this rigorous procedure, some differences between artistic and non-artistic stimuli remain (i.e. realism rating). The artistic images used in empirical studies interested in art are not created for experimental purposes and therefore possess a variety of elements that cannot
be fully controlled, which are congenital to the artistic nature of the images themselves. However, the presence of these differences requires caution in the interpretation and generalization of our results. Due to the specificity of our stimuli, we selected a small number of images that consequently required a relatively high number of repetitions (n = 6) across the experimental runs, potentially leading to a decrement in Blood Oxygenation Level Dependent (BOLD) activation. Participants made continuous AJs on stimuli categorized dichotomously as artistic or non-artistic. This procedure constitutes a necessary mismatch between the theoretical formulation and the methodological choices aimed to measure the aesthetic experience (see the ‘Introduction’ section). However, it is important to note that only the judgements offered to artistic images were entered in the ROIs analyses to respond to the hypotheses formulated. With respect to the procedure followed in this study, participants always performed the pain identification task inside the scanner. This procedure was followed in agreement with the extensive literature on empathy for pain adopting the same paradigm, but it prevents us from establishing that the same activation patterns are also possible during the ‘task-free’ observation of pictorial pain expressions. Further studies are needed to investigate the neural activations associated with the AJ of artistic facial expressions of pain, to support and extend the present results. Although observation of painful facial expressions engages pain-related regions (i.e. AI, pACC and aMCC), several arguments have been raised about the pain specificity of those areas, with critical implications for the functional interpretation of the neural overlap triggered by experiencing one’s own pain and observing others’ pain (Iannetti et al., 2013; Zaki et al., 2016). Lastly, whereas we were able to formulate specific predictions about the results expected for our first two aims, our third goal was mostly explorative. In this case, we can only suggest cautious interpretations, accounting for the notion of reverse inference.

In conclusion, our results show that empathy for pain brain regions can also be activated by artistic pain representations. Correlational analyses between functional brain responses and aesthetic ratings suggest that the activity of the insular and cingulate cortices concurs with the formation of an AJ in non-art expert observers. This work supports the necessity to further investigate how non-beauty-specific neural mechanisms could feed the complex phenomena of aesthetic experience and AJ.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Supplementary data

Supplementary data are available at SCAN online.

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