



HAL
open science

Shared body representations and the "Whose" system

Frédérique de Vignemont

► **To cite this version:**

Frédérique de Vignemont. Shared body representations and the "Whose" system. *Neuropsychologia*, 2013, pp.00-00. ijn_00812272

HAL Id: ijn_00812272

https://hal.science/ijn_00812272v1

Submitted on 11 Apr 2013

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Elsevier Editorial System(tm) for Neuropsychologia
Manuscript Draft

Manuscript Number:

Title: Shared body representations and the "Whose" system

Article Type: Special Issue: Marc Jeannerod

Section/Category: Emotion and Social Neuroscience

Keywords: self; bodily ownership; agency; naked intention; mirroring; imitation; interpersonal body representation; peripersonal space; mirror-tactile synaesthesia; touch; empathy; pain; embodiment

Corresponding Author: Dr. Frederique de Vignemont,

Corresponding Author's Institution: Institut Jean Nicod

First Author: Frederique de Vignemont

Order of Authors: Frederique de Vignemont

Abstract: Mirroring has been almost exclusively analysed in motor terms with no reference to the body that carries the action. According to the standard view, one activates motor representations upon seeing other people moving. However, one does not only see movements, one also sees another individual's body. The following questions then arise. To what extent does one recruit body representations in social context? And does it imply that body representations are shared between self and others? This latter question is all the more legitimate since recent evidence indicates the existence of shared cortical networks for bodily sensations, including pain (e.g., Singer et al., 2004) and touch (e.g., Keysers et al., 2004; Blakemore et al., 2005). But if body representations are shared, then it seems that their activation cannot suffice to discriminate between one's body and other people's bodies. Does one then need a "Whose" system to recognize one's body as one's own, in the same way that Jeannerod argued that one needs a "Who" system to recognize one's actions as one's own?

April 11th, 2013

Dear editor,

Please find attached my manuscript “Shared body representations and the “Whose” system” for submission to *Neuropsychologia*.

Yours sincerely,

Frederique de Vignemont
CNRS – EHESS – ENS
Institut Jean-Nicod, 29 rue d’Ulm, Paris 75005, France

Shared body representations and the “Whose” system

Frédérique de Vignemont

Highlights

Mirroring has been almost exclusively analysed in motor terms with no reference to the body that carries the action. According to the standard view, one activates motor representations upon seeing other people moving. Furthermore, most people have drawn the implications of those shared action representations for 3rd person mindreading (do they allow direct understanding of other people’s intentions?). By contrast, this paper focuses on the existence of shared body representations rather than on shared action representations and on their implications for self-awareness, and in particular on the sense of bodily ownership (how do I recognize this body as my own?).

To what extent does one recruit body representations in social context? And does it imply that body representations are shared between self and others? This latter question is all the more legitimate since recent evidence indicates the existence of shared cortical networks for bodily sensations, including pain (e.g., Singer et al., 2004) and touch (e.g., Keysers et al., 2004; Blakemore et al., 2005). But if body representations are shared, then it seems that their activation cannot suffice to discriminate between one’s body and other people’s bodies. Do we then need a “Whose” system to recognize our body as our own, in the same way that Jeannerod argued that we need a “Who” system to recognize our actions as our own?

Shared body representations and the “Whose” system

Frédérique de Vignemont

Institut Jean Nicod

CNRS-ENS-EHESS

Abstract:

Mirroring has been almost exclusively analysed in motor terms with no reference to the body that carries the action. According to the standard view, one activates motor representations upon seeing other people moving. However, one does not only see movements, one also sees another individual’s body. The following questions then arise. To what extent does one recruit body representations in social context? And does it imply that body representations are shared between self and others? This latter question is all the more legitimate since recent evidence indicates the existence of shared cortical networks for bodily sensations, including pain (e.g., Singer et al., 2004) and touch (e.g., Keysers et al., 2004; Blakemore et al., 2005). But if body representations are shared, then it seems that their activation cannot suffice to discriminate between one’s body and other people’s bodies. Does one then need a “Whose” system to recognize one’s body as one’s own, in the same way that Jeannerod argued that one needs a “Who” system to recognize one’s actions as one’s own?

Keywords: self, bodily ownership, agency, naked intention, mirroring, imitation, interpersonal body representation, peripersonal space, mirror-tactile synaesthesia, vicarious touch, empathy, pain, embodiment

Contact information:

Frédérique de Vignemont, Institut Jean-Nicod, 29 rue d'Ulm 75005 Paris France

Email: frederique.de.vignemont@ens.fr

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1. Introduction

1
2
3 A recent trend in the neuroscientific and philosophical literature suggests that we
4
5 have representations that are shared between self and others (Gallese, 2001; Goldman,
6
7 2006). It all started in 1992 when researchers in Parma found that the same neurons
8
9 fired both when a monkey was grasping a peanut and when it was watching the
10
11 experimenter grasping it (di Pellegrino et al., 1992; Rizzolatti et al., 1995; Grezes &
12
13 Decety, 2001). More generally, the motor system is activated not only when an agent
14
15 executes a goal-directed action, but also when an observer perceives the same action
16
17 performed by another agent. What makes action mirroring special is that it goes
18
19 beyond mere conceptual sharing. In Jeannerod (1994)'s terms, one does not share
20
21 only semantic knowledge about the action; one shares the agent's pragmatic
22
23 perspective. As such, action mirroring allows the observer to internalize another
24
25 individual's actions as if she were the agent by a kind of motor simulation.
26
27
28
29
30
31

32
33 Most people have drawn the implications of those shared action representations for
34
35 3rd person mindreading (do they allow direct understanding of other people's
36
37 intentions?). By contrast, Marc Jeannerod focuses his interest on their implications for
38
39 self-awareness and more specifically, for the sense of agency (how do I recognize my
40
41 actions as my own?). With Elisabeth Pacherie, he argues that shared motor
42
43 representations consist in what they call naked intentions, that is, intentions that are
44
45 neutral relative to the agent (Jeannerod and Pacherie, 2004). The naked content can
46
47 take two forms: either impersonal of the type <action, goal> or personal of the type
48
49 <x, action, goal>, with x as the parameter of the agent still to be determined (de
50
51 Vignemont, 2004; Jeannerod and Pacherie, 2004). In any case, the activation of
52
53 shared motor representations cannot suffice to differentiate whether it is the agent
54
55 moving or another person moving. Further processes based on extra information are
56
57
58
59
60
61
62
63
64
65

1 needed: “Given the existence of shared representations, something more than the sole
2 awareness of a naked intention is needed to determine its author.” (Jeannerod and
3 Pacherie, 2004, p. 140). The lack of differentiation between self and others seems
4 indeed to indicate that one needs a specific mechanism to take apart one’s actions and
5 other people’s actions. Georgieff and Jeannerod (1998) call this mechanism the
6 “Who” system.
7
8
9
10
11
12
13

14 Jeannerod analyses mirroring exclusively in motor terms, with no reference to the
15 body that carries the action. However, it seems that in some situations, we need to
16 establish a correspondence between the representation of our body and the
17 representation of other people’s bodies. How do we achieve such correspondence?
18 Are there shared body representations? These questions are all the more legitimate
19 since recent evidence indicates the existence of shared cortical networks for bodily
20 sensations. Brain activity partially overlaps when one experiences touch or pain and
21 when one observes another individual receiving a tactile or a painful stimulus (e.g.,
22 Singer et al., 2004; Keysers et al., 2004). Strikingly, people with mirror-tactile
23 synaesthesia even consciously feel tactile sensations on their face upon perceiving
24 another person touched on the face (Blakemore et al., 2005). Like for action, shared
25 cortical networks for bodily sensations have been systematically considered from the
26 perspective of the understanding of other people, and especially in relation to empathy
27 (Goldman, 2011; Banissy and Ward, 2007; Keysers et al., 2010; Ebish et al., 2009,
28 Wood et al., 2010). But what consequences do they have for the sense of bodily
29 ownership? All together, these findings may be taken as evidence of shared body
30 representations. But if body representations are shared, then it seems that their
31 activation cannot suffice to discriminate between one’s body and other people’s
32 bodies. Should we then extend Jeannerod’s conclusion to the sense of bodily
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

ownership? In other words, do we need a “Whose” system to recognize our body as our own?

2. An embodied approach to the correspondence problem

A classic problem in the imitation literature is known as the intersubjective correspondence problem: how does one map another person’s movements to one’s own movements (Goldenberg, 1995; Heyes, 2001)? The classic way to approach the correspondence problem has been in sensorimotor terms: how does one map visual information to motor command (e.g., Brass and Heyes, 2005)? Alternatively, one can consider that the main challenge is a problem of intermodal correspondence: how does one map visual information to somatosensory information (e.g., Meltzoff and Moore, 1995)? This latter problem can be found not only in imitation (and especially in the case of opaque movements like tongue protrusion for which one has no visual feedback), but also in vicarious bodily sensations experienced when observing other people receiving tactile (or painful) stimulation. However, the question that interests us here is whether there is a further approach to the correspondence problem. In other words, is there a bodily correspondence problem? In imitation, one maps the movement of another body onto one’s body, and in mirror-tactile synaesthesia, one maps the location of touch on another body to one’s own body. For all that, is intersubjective correspondence mediated by bodily correspondence? Shared motor, affective and somatosensory representations have recently been taken as evidence in favour of embodied social cognition (Gallese, 2007; Gallagher, 2005). However, the sense in which those states are embodied remains often obscure. We shall see now that they are two distinct types of shared representations, and only one of them involves body representations.

2.1 Disembodied shared representations

According to a narrow interpretation of the notion of embodiment, shared representations qualify as embodied in virtue of their bodily content (Goldman and Vignemont, 2009). They carry information about the body, and in particular about its spatial properties. In this sense, embodied representations are representations of the body. But there is another interpretation of the notion of embodiment, according to which shared representations qualify as embodied in virtue of their specific bodily format (Goldman and de Vignemont, 2009; Gallese and Sinigaglia, 2011). Bodily format is a class of representational code, which includes somatosensory, interoceptive, affective, and motor codes. It is then one thing to be endowed with a bodily format and it is another thing to have a bodily spatial content. For example, the visual representation of one's hand has a spatial bodily content, but it is not encoded in a bodily format. Conversely, affective states are endowed with a bodily format (because they are associated to - or caused by - certain physiological bodily states), but they do not have a spatial bodily content.

Let us now apply this distinction between content and format to social cognition. With Alvin Goldman, we argued that shared representations – motor, somatosensory and affective – are endowed with a bodily format. In this sense, they are embodied. For all that they do not necessarily represent the properties of the body, and in particular its spatial properties. In this narrower sense, they are not embodied. In other words, the existence of shared representations in bodily format does not necessarily imply the existence of shared body representations. Here I will focus exclusively on the notion of bodily content. The question then is whether action mirroring, vicarious pain and vicarious touch are embodied or disembodied in the narrow sense.

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33

Let us consider first the case of action mirroring. It is useful here to draw the parallel with the distinction between imitation and emulation. Emulation involves the copy of the goal of a perceived action. For instance, upon seeing someone taking an umbrella, one takes an umbrella as well, but the bodily movement can be completely different. In contrast, in imitation not only is the goal copied, but also the bodily movements used to reach the goal. For instance, upon seeing someone taking an umbrella by lifting it with the foot, one performs the same movement instead of simply grasping it with the hand. The difference between imitation and emulation is thus a matter of motor hierarchy: at what level the visually perceived action is analysed and copied. The low level is strongly embodied (imitation), whereas the more abstract levels are weakly embodied (emulation). The question then is at what level action mirroring occurs. To what extent are bodily parameters represented in mirror systems? Does one use body representations to covertly replicate actions that one perceives?

34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

A series of brain imaging studies show that mirroring is closer to emulation than imitation insofar as it encodes the motor goal instead of the specific bodily movement to achieve the goal (Csibra, 2007). On the one hand, the same bodily movement performed for two distinct intentions (grasping a mug for drinking or for cleaning for instance, cf. Iacoboni et al., 2005) activates two distinct brain areas. On the other hand, different bodily movements performed for the same intention (grasping with the mouth and grasping with the hand, cf. Gallese et al., 1996; grasping food with one's hand and with a stick, cf. Ferrari et al., 2005) activate the same brain area. Interestingly, Csibra (2007) notes that the majority of mirror neurons are only broadly congruent. What is shared is not the representation of the specific limb to move, but rather some motor representation higher in the motor hierarchy. Intersubjective

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

correspondence is then achieved exclusively in motor terms, and not in bodily terms. These findings show how weakly embodied (in the content sense, not in the format sense) the mirror systems can be.

The question of bodily correspondence may then seem more legitimate in the case of vicarious bodily sensations than in the case of action mirroring. At first sight, one may expect the involvement of body representation more systematic, or even necessary, in vicarious pain and in vicarious touch insofar they are vicarious versions of bodily sensations. Yet, this is not the case, as we shall see now.

Let us consider the case of pain first. The pain matrix includes two functionally specialized networks. The sensory-discriminative component involves the experience of the intensity of pain and its bodily location. It recruits primary and secondary somatosensory areas (SI and SII) as well as the posterior insula. Since SI has a somatotopic organization, different regions of SI are activated according to whether one experiences standard pain in one's hand, in one's foot, or in one's mouth. In addition, the sensory-discriminative component has a motor counterpart: its activity underlies specific automatic localized motor responses whose function is to avoid (or decrease) the pain. For instance, the muscles adjacent to the location of the painful stimulus freeze, so to speak. The affective component involves the experience of the unpleasantness of the painful experience. It recruits the anterior insula, the anterior cingulate cortex, the thalamus, and the brain stem. It lacks somatotopic organization.

Now if one experiences vicarious pain, then what happens? Are both components active? More particularly, is vicarious pain localized in a part of one's body that matches the body part that is seen injured? Interestingly, when participants observe cues indicating that another individual is receiving a painful stimulus, most brain imaging studies report a selective activation of the affective component only, with no

1 associated activation of SI (e.g., Singer et al., 2004). For instance, the same brain
2 activation was found whether the hand or the foot was injured (Jackson et al., 2005).
3
4 Vicarious affective pain is thus indifferent to the bodily location of pain. This is why
5 one can vicariously experience pain upon either perceiving another's facial
6 expression, which does not reveal the bodily location of the other's pain (Botvinick et
7 al., 2005) or imagining another's non-located painful experience (Jackson et al.,
8 2006). One may conceive vicarious affective pain as a vicarious version of a non-
9 localized bodily feeling of the type "It hurts" (de Vignemont and Jacob, 2012). In this
10 sense, it is disembodied.
11
12
13
14
15
16
17
18
19
20
21

22 One may expect vicarious tactile sensations to be more strongly embodied, since
23 they are generally devoid of affectivity. Yet, this is far from being so. Touch is also
24 characterized by its dual nature, although of a different kind from pain. Touch can be
25 active (touchant) or passive (touché), or both at the same time in self-touch. One can
26 then distinguish between vicarious touchant experience and vicarious touché
27 experience, although they are confounded in most experiments. Several brain imaging
28 studies found activity only in SII, and not in SI, when participants observed another
29 person being touched (Keysers et al., 2004; Ebisch et al., 2008). Interestingly, SII was
30 activated even when participants watched an object being touched, rolls of paper for
31 instance. It thus seems that most of the time vicarious touch takes into account
32 primarily the body part that is touching (vicarious touchant experience): "What is
33 being touched does not matter as long as touch occurs" (Keysers et al., 2004, p. 339).
34 According to Keyser and coll. (2010), vicarious touch corresponds to the mirroring of
35 tactile consequences of action.
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54

55 To conclude, there is evidence of shared action, shared touch and shared pain, but
56 so far there is no evidence of shared body representations. The intersubjective
57
58
59
60
61
62
63
64
65

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

correspondence problem is not necessarily encoded in bodily terms. One can map the other onto oneself while bracketting the spatial properties of the body. Shared representations are then disembodied insofar as they do not have a bodily spatial content. Yet, as we shall see now, it can be useful, or even necessary in some situations, to analyse other people's movements and sensations in terms of the body parts that move and that are stimulated.

2.2 Embodied shared representations in imitation

“Body part coding reduces the visual appearance of the demonstrated gestures to simple spatial relationships between a limited set of discrete body parts. Body part coding facilitates imitation because it produces equivalence between demonstration and imitation that is independent of the different modalities and perspectives of perceiving one's own and other persons' bodies, and because it reduces the load on working memory in which the shape of the gesture must be held until motor execution is completed.” (Goldenberg, 2009, p. 1455)

Goldenberg (2009) defends an embodied approach to imitation, according to which one uses knowledge about the body to decode the other's movement and copy it. The special role of body representation in imitation is especially salient for some specific types of actions. This is the case for instance when one imitates meaningless gestures. Upon seeing a military salute, one can recognize it as such and use a stored motor schema of the salute to replicate it. But when one sees a person putting her thumb above her eyebrow, the only solution is to encode the movement in terms of the body parts that are seen and their spatial relations. Hence, it is classically assumed that

1 there are two routes for imitation: one that involves the recognition of the seen action
2 and the other that does not, and only the latter uses ‘body part coding’ (for review, see
3 Rumiati et al., 2009). Deficits in imitation are then sometimes explained in terms of
4 deficits of body representation. This is the case for instance in patients with ideomotor
5 apraxia who are most affected in the imitation of meaningless gestures (Goldenberg,
6
7 1995; Buxbaum et al., 2000; Schwoebel & Coslett, 2005).
8
9
10
11
12
13
14

15 Imitation is often considered as the key mechanism for acquisition of new skills.
16
17 But the use of imitation is not restricted to explicit learning context. Rather, we have a
18 permanent tendency to automatically imitate other people’s movements, which we
19 need to inhibit (Brass et al., 2001). This can be shown if subjects simultaneously
20 perform a movement. For example, participants are faster in moving their index finger
21 when observing an index finger movement than a middle finger movement. Motor
22 facilitation effect is driven by the bodily congruency between the effectors and cannot
23 be reduced to a mere spatial congruency effect. For instance, Brass and coll. (2001)
24 asked participants to observe an index finger tapping and lifting, but the image was
25 inverted (upward motion for tapping and downward motion for lifting). Yet,
26 participants were faster in tapping when observing tapping in spite of the fact that
27 tapping responses (downward motion) were spatially incompatible with tapping
28 stimuli (upward motion) and were spatially compatible with lifting stimuli (downward
29 motion). This study is only one among many other experiments that indicate that how
30 parts of the body are located and move relative to one another is mirrored when
31 observing other people’s actions (e.g. Gillmeister et al., 2008). Interestingly, it was
32 found that the mere observation of coloured patches on another person’s static body
33 parts (head, hand or foot) sufficed to prime action with the same body parts (Bach et
34 al., 2007). The authors concluded in favour of a “body schema that represents
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 locations on the observer's body and on the bodies of others in a common format" (p.
2 515). Hence, in some cases, what is shared is not only the representation of the goal or
3 the movement, but the representation of the effector itself that performs the
4 movement. This is confirmed at the neural level. For example, observation of hand,
5 foot and mouth actions selectively activates distinct regions of human ventral
6 premotor and parietal cortex (Buccino et al., 2001). Importantly, Wheaton et al.
7 (2004) showed this somatotopic pattern of activation even when movements were
8 held constant across effectors (opening and closing movements of a hand and a
9 mouth, respectively). Hence, correspondence between self and others can be encoded
10 in bodily terms in some situations.
11
12
13
14
15
16
17
18
19
20
21
22
23

24 There are several versions of the bodily approach to imitation. For example,
25 Goldenberg (1995) argues that imitation recruits abstract conceptual knowledge of the
26 human body. By contrast, Chaminade and coll. (2005) as well as Buxbaum and coll.
27 (2000) consider the primitive body schema as the most likely candidate in virtue of its
28 sensorimotor content. Finally, the body image is sometimes also mentioned in virtue
29 of its visuo-spatial content (Gallagher and Meltzoff, 1996). Given the confusion
30 around these various notions (de Vignemont, 2010), I shall leave them aside and
31 instead specify what bodily information needs to be encoded in imitation. A first point
32 is that it must be quite rough-grained. There are many differences between bodies.
33 The body whose movements I copy is never identical to my body: not only is our
34 posture rarely the same, but our gender can also be different, as well as the size of our
35 limbs, the strength of our muscles, the flexibility of our joints, and so forth. Yet this
36 does not prevent intersubjective correspondence. Typically a child can imitate an
37 adult. The brain must thus abstract from major bodily differences. It then seems that
38 what remains in common between all bodies is the configuration of the various body
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 parts, i.e. the fact that we have two hands and two feet and they are respectively
2 located at the end of our arms and legs. More precisely, what needs to be represented
3 is the functional arrangement of bodily effectors. It was indeed found that the
4 movements that are seen are perceived in terms of functional units of effectors that
5 move together. Functional body parts consist in parts of the body that are regrouped
6 by their role in action. Fingers and hands can constitute grasping units for instance.
7 The foot and the leg can constitute kicking units or walking units. The fact that it is
8 visually shaped like a hand or like a foot and a leg does not matter. For example,
9 observing hand, foot or mouth actions selectively activate brain areas for hand, foot,
10 and mouth even if the movement is performed by non-conspecifics such as monkeys
11 or dogs (Buccino et al., 2004), or robots (Gazzola et al., 2007a; Press et al., 2005).
12 What matters to mirroring is mouth-action, not whether the mouth that is moving is
13 yours or your dog's. The visual appearance and exact shape of the perceived body
14 part are of little importance. What matters is the type of movements it can afford. This
15 is well illustrated by the following finding in aplasic individuals who were born
16 without hands and who perform with their feet and their mouth actions normally
17 performed with hands. It was found that they activated regions of the mirror system
18 recruited in mouth and foot movement execution when observing hand movements
19 (Gazzola et al., 2007b). In mirroring, one matches functional bodily units dedicated to
20 specific types of movements (grasping for instance), no matter whether those
21 functional bodily units are instantiated by a hand or a mouth. Arguably, this level of
22 segmentation of the body into parts is the most relevant for action in general, and for
23 imitation and mirroring in particular. For most gestures one can dispense with more
24 detailed bodily specification. All that needs to be represented is the dynamic position
25 of functionally defined body parts with respect to one another (Buxbaum et al., 2000).
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

2.3 Embodied shared representation in vicarious sensations

Intersubjective correspondence recruits body representations not only in imitation, but also for some vicarious bodily sensations. Vicarious pain can be of two distinct kinds. It includes not only vicarious affective pain, which is disembodied, but also vicarious sensory pain, which is embodied (de Vignemont and Jacob, 2012). For instance, it was found activation of the somatotopically organized SI upon seeing the back of another's hand being deeply penetrated by a needle (Bufalari et al., 2007). Motor responses to vicarious sensory pain are even muscle-specific, similar to those found when one is injured: when one sees another's hand being hurt, one automatically freezes one's own hand, as if one's own hand were injured (Avenanti et al., 2005; 2009). These experimental results strongly suggest that in experiencing vicarious sensory pain, one responds to the perception of another's bodily part subjected to painful stimulation by expecting specific sensorimotor consequences of pain at the same location on one's own body.

Likewise individuals with mirror-touch synaesthesia consciously feel tactile sensations on their own body when they see another person's being touched (Blakemore et al., 2005; Banissy and Ward, 2007). Upon seeing another being touched on the left cheek for example, they experience a sensation of touch on their own cheek. If at the same time they are touched on the right cheek, they either report feeling touch on both sides or they make mistakes (they report feeling touch on the left). This is not a mere attentional effect. They do not make mistakes if they see a visual flash on the left of the face rather than receive a tactile stimulus (Banissy and Ward, 2007). When compared to normal subjects, it was found a more intense activity of the somatosensory cortex when subjects with mirror-tactile synaesthesia saw

1 people being touched (Blakemore et al., 2005). In addition to individuals with mirror-
2 tactile synaesthesia, it was found that some patients with phantom limbs reported
3 feeling tactile sensations on their phantom hand when observing another person's
4 hand being touched (Ramachandran and Brang, 2009).
5
6
7
8
9

10 The interpersonal dimension of body representations in the sensory domain is
11 confirmed by results both in the multisensory literature and in the neuropsychological
12 literature. It is well known that viewing the body part that is touched (without viewing
13 the touching object) enhances tactile acuity so that one's judgements about tactile
14 sensations are both faster (Tipper et al., 1998) and more accurate (Kennett et al.,
15 2001). Importantly, the effect of visual enhancement of touch is as effective when
16 seeing one's own body part as when seeing another person's body part (Haggard,
17 2006). Likewise, Thomas and coll. (2006) found that participants were faster in
18 detecting touch when they saw before a non-predictive visual cue on another person's
19 body at the corresponding location. The authors concluded: "We believe that our
20 results provide the first behavioural evidence in normal subjects for interpersonal
21 body representation based on a somatotopic spatial map, at the purely sensory level."
22 (Thomas et al., 2006, p. 328). Findings in neuropsychology also indicate the existence
23 of a similar body representation for perception that is 'interpersonal'. Most – but not
24 all – patients with autotopagnosia fail to locate body parts not only on their own body,
25 but also on other people's bodies, on mannequins and on drawings of a human body.
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48

49 Hence, one exploits body representations in order to map the location of the tactile
50 or painful stimulation of another's body onto one's own body. One can speculate that
51 like in action, bodily coding in vicarious sensations must go beyond the differences
52 between bodies and carry information only about the configuration of the body. But
53 the spatial organization of the body is not similarly represented in imitation and in
54
55
56
57
58
59
60
61
62
63
64
65

1 vicarious bodily sensations. Whereas the representation of the body that is used in
2 imitation represents functionally defined body parts, the representation of the body
3 that is used in vicarious bodily sensations represents anatomically defined body part.
4 One may for instance note that the disruption of the representation of the
5 configuration of the body in patients with autotopagnosia does not affect their actions.
6 Unfortunately, there is little evidence on the spatial organization of body
7 representation in the case of vicarious bodily sensations. The distinction between a
8 functional and an anatomical representation of the body, however, is in line with a
9 classic distinction between two types of body representation, one oriented towards
10 action and the other towards body perception (Dijkerman & de Haan, 2007;
11 Gallagher, 2005; Head & Holmes, 1911; Paillard, 1980; de Vignemont, 2010). The
12 two types of body representation follow distinct principles of spatial organization (de
13 Vignemont et al., 2005; de Vignemont et al., 2009; Tsakiris et al., 2006; Hach et al.,
14 2011). In particular, action requires a more unified map of the body than perception:
15 when I move my forearm, my hand and fingers follow. By contrast, if someone
16 touches my forearm, this usually does not tell me anything about the sensation in my
17 hand and fingers. On the basis of the results described above, it seems that both types
18 of body representation can be recruited in social context.

2.3 Embodied shared representations of peripersonal space

19 There is a last series of evidence that argue in favour of shared body
20 representations. We know that the space immediately surrounding one's body, namely
21 peripersonal space, is processed differently than extrapersonal space (for review,
22 Brozzoli et al., 2012). A large number of studies with monkeys have now found
23 bimodal neurons in several brain structures (putamen, parietal and premotor area),
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 which are activated both by touch and by vision (or audition). Interestingly, they are
2 activated even when the visual stimulus is not on the body part itself, but up to 30-
3 50cm off the body part (e.g., Cooke and Graziano, 2003). Moreover, the visual (or
4
5 auditory) activity follows the position of the tactile receptive field associated to a
6
7 specific part of the body when the part is moved. It is thus body-centered. Several
8
9 studies support the existence of a similar multimodal representation of peripersonal
10
11 space in humans, which results in the interaction between visual (or auditory)
12
13 stimulus near the body and tactile processing. The best known evidence in favour of
14
15 such multimodal interaction is the cross-modal congruency effect. Participants are
16
17 asked to report when they feel a vibrotactile stimulation and to judge its location. It
18
19 was shown that neutral visual stimuli interfere with tactile detection and tactile
20
21 localization, if the location of the visual stimuli is incongruent with the location of the
22
23 tactile stimuli (Spence et al., 2004). This effect works only if the visual stimuli are
24
25 close to a part of the body. For such interference to happen, both visual and tactile
26
27 experiences must share a common spatial frame of reference. It is not the visual frame
28
29 because the interference is not sensitive to bodily posture. If the flash of light is close
30
31 to your left hand and the touch is applied to your right hand, your performance
32
33 decreases even if your hands are crossed. Hence, even if the flash is visually located
34
35 close to the hand that is touched, they are not perceived as being congruent. As
36
37 Graziano and Gross (1993, p. 107) described it, the peripersonal space is like “a
38
39 gelatinous medium surrounding the body that deforms whenever the head rotates or
40
41 the limbs move”.

52
53
54 Objects in peripersonal space are endowed with a special significance for the body.
55
56 They may be perceived as potential threats or obstacles. The function of information
57
58 processing in peripersonal space is not uniquely to avoid snakes, and other life-risking
59
60
61
62
63
64
65

1 situations. One needs to protect one's body in everyday life as well, as when one walks
2 through a room without hitting the furniture, one brushes away an insect, one reaches
3 safely around a prickly object, or one sits at a desk without bruising one's elbows and
4 arms as one works. Objects in peripersonal space may also be perceived as potential
5 targets for action. One may question whether there is a single peripersonal space or two,
6 one defined as margin of safety and the other defined as reaching space, but I shall leave
7 this question aside and focus instead on the representation of other people's peripersonal
8 space. A series of evidence indeed indicates that it recruits the same resource as the
9 representation of one's peripersonal space.
10
11
12
13
14
15
16
17
18
19
20
21

22 Ishida and coll. (2010) found in monkey parietal areas that some visuo-tactile
23 neurons fired when a visual stimulus was close both to a part of the monkey's body
24 and to the experimenter's equivalent body part. The effect was body part specific.
25 Likewise in humans, it was found that SI was activated when the experimenter
26 stroked with a paintbrush the region of space close to a hand visually presented from a
27 third-person perspective (Schaefer et al., 2012). Surprisingly, however, the activity
28 was stronger than when the experimenter stroked the peripersonal space of a hand
29 visually presented from a first-person perspective. In addition, it was shown that one
30 represents other people's reaching space in the same way as one represents one's
31 reaching space. For example, it is known that the motor system is activated by the
32 mere vision of a graspable object. Furthermore, it has been shown that the spatial
33 alignment of the object with the effector affects the motor response. For example, if
34 the handle of a mug is aligned with the left hand, one is quicker if using the left hand
35 than the left hand to grasp it. This effect works, however, only if the mug is in the
36 agent's reaching space (Costantini et al., 2010). Now a recent study showed that the
37 effect worked as well if the mug was in a virtual individual's reaching space, although
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

it was not in the participants' own reaching space. There was no effect, on the other hand, if the mug was close to a cylinder of the same size as the virtual avatar. Neither was there any effect if there was a near-transparent panel between the avatar and the mug (Costantini et al., 2011). Cardellicchio and coll. (2012) further showed that motor-evoked potentials were higher when the mug was close either to the participants or to the avatar. The authors concluded: "Our proposal is that such sensitivity can be explained by means of an interpersonal bodily space representation allowing one to map the body of other people in terms of their actual motor possibilities." (p. 4).

To conclude, there are several ways to solve the intersubjective correspondence problem. In the domain of mirroring, it can involve representations of the motor goal. In the domain of vicarious bodily sensations, it can involve affective representations. Those representations qualify as embodied insofar their mental format is concerned. But they do not qualify as embodied in a narrower sense: they do not represent bodily states. Yet, there are cases in which the correspondence problem is solved thanks to representations of the body. Some cases of imitation as well as some vicarious bodily sensations respect bodily congruency. In order to do so, they need to exploit a representation of the body to map other people's bodies to one's own body. The body is then the common "currency" between self and others. And this is so both in the motor and in the perceptual domains.

But what consequences does it have for the sense of bodily ownership? Jeannerod and Pacherie's argument for the sense of agency can be articulated into four steps: (i) there are representations shared between self and others; (ii) thus, they must be endowed with a naked content (neutral relative to the agent); (iii) thus, they can lead to confusion between self and others; (iv) thus, one needs a "Who" system. We shall

1 now ask whether this argument can apply to the sense of bodily ownership and
2 whether the evidence reviewed so far raises a challenge for theories of bodily
3 ownership, in the same way that the existence of shared action representations raises a
4 challenge for theories of agency. I will now consider two ways to escape Jeannerod
5 and Pacherie's argument. First, one may object to the very first step: there are no
6 representations shared between self and others, but only a coupling of self-specific
7 and non self-specific representations. Second, one may object to the transition from
8 the second step to the third step: naked content does not necessarily open the door to
9 confusion.
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24

25 **3. A matter of coupling?**

26
27
28 Intersubjective correspondence can be encoded in bodily terms. So far I have
29 assumed that it involves shared body representation, that is, a single representation
30 exploited for both one's body and other bodies. There is, however, an alternative
31 model that can account for the evidence reviewed above. Intersubjective
32 correspondence can also be achieved by the automatic association of distinct
33 representations. This latter view has been defended among others by Celia Heyes
34 (2001). According to the Associative Sequence Learning (ASL) model, imitation is
35 based on past experiences of the systematic coupling between the action one performs
36 and its sensory consequences. Typically, when I wave my hand, I see it waving. I can
37 thus learn the sensorimotor association so that when I see another person waving, the
38 sensory input can automatically elicit the associated motor output. Another example
39 can be found in Gallagher and Meltzoff (1993). They argue that imitation depends on
40 the systematic association between the body schema, which is primarily
41 proprioceptive and motor, and the body image, which is primarily visual. One can
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 also interpret some of the multisensory results along these lines. On the basis of their
2 findings that the vision of non-predictive cues of another person's body could
3 interfere with tactile processing on one's own body, Thomas and coll. (2006) defined
4 shared body representations in terms of "a special, automatic mechanism for
5 associating sensory body events" (p. 327).
6
7
8
9
10

11
12 These different conceptions do not directly address the question of the specific
13 content of the representations that are associated. In particular, are those
14 representations naked? Are they neutral relative to the owner of the body? Let us
15 consider in more detail the ASL model. The two terms of the association are (i) a
16 motor representation, which consists in the activity of the mirror system at the neural
17 level, and (ii) a visual representation of the movement. The visual representation can
18 represent either one's own movement or another person's movement. In this sense, it
19 can be said to be naked. The seen movement is unattributed. The motor
20 representation, on the other hand, is always used for one's own motor system. It plays
21 a role for one's own actions, and one's own actions only. Hence, one may argue that
22 it is first-personal (although not necessarily reflexively). The association can thus be
23 reformulated as follows: an intention <I raise my finger> associated to a visual
24 representation <a finger raising>. The association was built on the basis of past
25 experiences of seeing my finger raising when I have the intention to raise my finger.
26
27 But it has generalized to any raising finger. Hence, when I see you raising your finger,
28 this activates my mirror system and I have an automatic tendency to imitate the
29 movement. On this view, the motor representation is activated by another person's
30 actions, but for all that it does not represent intentions other than one's own. Only the
31 visual representation has a naked content, not the motor representation. We can apply
32 the same analysis to mirror-tactile synaesthesia. Then the association is between a
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 tactile representation <touch on my cheek> and a visual representation <tactile stimuli
2 on a cheek>. Again, only the visual representation has a naked content, even if the
3 tactile representation is activated by another person being touched.
4
5
6

7 If this is the right interpretation, then it is misleading to talk of ‘shared
8 representation’. The results described in the first section are merely a matter of
9 coupling, rather than sharing. They have no implication for the content of the motor
10 and the somatosensory representations. All they imply is that visual representations
11 have a naked content, but this is hardly surprising. Discriminating one’s body from
12 other bodies is then not an issue. As Gallagher and Meltzoff (1996, p. 225-226) noted,
13 “Thus there is a coupling between self and other, and this coupling does not involve a
14 confused experience”. The body representation associated to the visual representation
15 of other people’s bodies can remain self-specific. One can thus dispense with a
16 “Whose” system. The possibility of bodily correspondence has no consequence for
17 the sense of bodily ownership.
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34

35 However, one may challenge Gallagher and Meltzoff’s conclusion: coupling can
36 lead to confusion if the coupling is mandatory. If the activation of one representation
37 automatically and systematically induces the activation of the other, and vice-versa,
38 then the coupled representations behave as if they constitute a unique representation.¹
39 It would be like marriage: after a while, you can never invite one spouse without the
40 other; they have lost their individuality. More than coupling, one should then talk of
41 fusion, which can lead to confusion. And if this is the case, then we are back to our
42 original problem. If the couple is activated both by seeing one’s body and other
43 people’s bodies, then the activation of the couple can no longer suffice to discriminate
44 among bodies.
45
46
47
48
49
50
51
52
53
54
55
56
57
58

59
60 ¹ I would like to thank Wolfgang Prinz for this objection.
61
62
63
64
65

1 The question is thus whether the coupling is mandatory or not. If it is, then it does
2 not make much functional difference with actual sharing. To settle the debate, one
3 must look at the empirical data. However, the evidence is not straightforward. On the
4 one hand, some results indicate that the association can allow for some flexibility and
5 plasticity. For example, it has been shown that automatic imitation is sensitive to
6 sensorimotor learning (Cook et al., 2010). Body part priming in imitation was reduced
7 following the repeated exposure to incongruent sensorimotor associations such as
8 observing a hand action while performing a movement with the foot. On the other
9 hand, other findings indicate a tight coupling, so tight that there are bidirectional
10 relations between the two terms of the association.
11
12
13
14
15
16
17
18
19
20
21
22
23
24

25 Most studies have investigated the effect of the observation of others upon oneself,
26 but a few studies showed that the representation of one's body influences the
27 perception of other people's bodies. For example, when participants observe two
28 photographs that differ only in the position of one limb that alternates, they see an
29 illusory movement of the limb. If the photographs are flashed in rapid succession,
30 they perceive the limb traversing the shortest possible path of visual apparent motion,
31 although it is biologically impossible because of joint constraints. But if the
32 presentation rate slows down, they perceive paths of apparent limb movement that
33 follow natural human limb trajectories (Shiffrar and Freyd, 1990). This result
34 indicates that knowledge of one's joint constraints determines how one perceives
35 other people's bodily movements. This effect is not driven by mere visual familiarity
36 of other people's movements, as shown by a study with two aplasic patients born with
37 no arm (Funk et al., 2005). They shared the same visual familiarity of the movements
38 that are biologically possible, but only one of the patients experienced phantom arms,
39 and thus, had bodily familiarity with joint constraints. Interestingly, it was found that
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 only this patient showed the same perceptual pattern as normal subjects. Hence, the
2 representation of one's bodily constraints can affect the perception of another
3 individual's bodily movements. Another study showed that healthy participants were
4 more efficient in detecting changes in a model's leg posture than in the model's arm
5 posture if they were moving their legs, and conversely that they were more efficient in
6 detecting changes in a model's arm posture if they were moving their arms (Reed and
7 Farah 1995). Hence, not only does the perception of another person's movement
8 affect one's own movements (other-to-self), but one's own movements can also
9 facilitate the perception of another's body (self-to-other). These results do not offer a
10 definite reply about the strength of the association, if association there is. Still they
11 argue in favour of a strong coupling, or even of actual sharing. Let us now consider
12 the implications of the hypothesis that body representations can be shared between
13 self and others.

34 **4. A "Whose" system?**

35
36
37
38 If there are shared body representations, then the same representations are
39 activated both for one's body and for other people's bodies. In order for body
40 representations to fulfil their intersubjective function, they must thus represent what
41 one's body and other bodies have in common, and only that. In Jeannerod and
42 Pacherie's terms, they must have a naked content. Their content is neutral relative to
43 whose body it is. They leave the body unattributed and represent indifferently one's
44 body and other people's bodies. They can be either impersonal representations of the
45 type <body part, bodily property> or personal representations with the owner x left
46 unspecified of the type <x, body part, bodily property>. Thanks to their naked content
47 they enable the perceiver to imitate another individual's actions and to map her
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 sensations onto the perceiver's body. What can then be the content of naked body
2 representations? On the basis of his findings on ideomotor apraxia, Goldenberg (1995,
3 p.71) proposes the following characterization: a "general concept of the human body
4 which applies regardless of whether one's own body is concerned or not". But if this
5 is the right way to characterize shared body representations, then they can hardly
6 ground the sense of bodily ownership. It then seems that one cannot dispense with a
7 "Whose" system.
8
9
10
11
12
13
14
15
16
17
18
19
20

21 4.1 The Comparator models of ownership 22

23 According to Jeannerod and Pacherie (2004, p. 139-140), the existence of shared
24 action representations implies that they cannot ground the sense of agency: "If,
25 however, we can be aware of both our intentions and those of others in the same way,
26 namely as unattributed or 'naked' intentions, the problem of self-other discrimination
27 does indeed arise (...) this cortical network provides the basis for the conscious
28 experience of goal-directedness—the primary awareness of intentions—but does not
29 by itself provide us with a conscious experience of self- or other-agency." Likewise
30 one may argue that because of their naked content, shared body representations
31 cannot suffice to distinguish between one's body and other people's bodies. The
32 impossibility lies in the fact that it cannot solve two opposite problems: the
33 correspondence problem in intersubjective situations and the ownership problem. If at
34 some level the representation of one's body is similar to the representation of other
35 people's bodies, then how could it ground the sense of ownership? It seems that the
36 naked body representation would have to play two incompatible roles: grounding self-
37 awareness and grounding other-awareness. It thus follows from the existence of naked
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 body representations that further processes are needed to discriminate between one's
2 body and other people's bodies. But what is the exact nature of the "Whose" system?
3
4

5 In the recent philosophical and empirical literature, the prominent view is that the
6 sense of agency should be understood as the result of the comparison between the
7 prediction of the sensory consequences of one's actions and their actual
8 consequences. Can one generalize the comparator model to the sense of body
9 ownership? But then what are the different types of information to compare? In the
10 case of the sense of agency, there is efferent information that can be compared to
11 sensory information, but this is not true in the case of the sense of ownership.
12 Arguably, one can feel one's body as one's own although one is not moving. What
13 type of information can help differentiate one's body from other bodies?
14
15
16
17
18
19
20
21
22
23
24
25
26

27 The dominant model of the "Whose" system is entirely drawn from the Rubber
28 Hand Illusion (RHI): synchronous stroking of one's own occluded hand and an
29 anatomically congruent visible rubber hand leads to a sense of ownership over the
30 rubber hand. On the basis of the RHI, it has been suggested that intermodal matching
31 plays a key role for ownership (Botvinick and Cohen, 1998, Makin et al., 2008, but
32 also Rochat 1998 for developmental evidence): "It has been proposed that the body is
33 distinguished from other objects as belonging to the self by its participation in specific
34 forms of intermodal perceptual correlation" (Botvinick and Cohen, 1998, p. 756). In
35 support of this view, it was found that the ownership rating in questionnaires was
36 correlated with the activity in brain regions that are classically involved in
37 multisensory processes: "the detection of correlated multisensory signals by these
38 regions [premotor cortex and posterior parietal area] is the mechanism for body
39 ownership." (Ehrsson et al., 2005, p. 10571).
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

However, the hypothesis of intermodal matching leaves many questions unanswered. As pointed out by Botvinick and Cohen (1998), one still needs to determine the “special ingredient” for ownership. Indeed, the sense of body ownership cannot derive from any kind of intermodal correlation. Imagine that you see and hear two hands clapping. Despite visuo-auditory correlation, you do not feel these hands as your own. You also need the visuo-auditory information to correlate with proprioceptive and tactile information indicating that you, and nobody else, are clapping your hands. In other words, there must be information that is self-specific (e.g. somatosensory information) for intermodal correlation to play a role for ownership. Furthermore, there are some constraints that lay upon the RHI. For instance, visuo-somatosensory correlation fails to elicit an ownership illusion when the rubber hand is replaced by a wooden spoon (Tsakiris and Haggard, 2005). Hence, Tsakiris (2010) argues in favour of a multi-layered “Whose” system that involves three distinct comparators: between the visual form of the viewed object and a pre-existing body model, between the current state of the body and the postural and anatomical features of the body-part that is to be experienced as one’s own, and between the vision of touch and the felt touch and their respective reference frames.

42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

However, although the RHI has been used as the experimental paradigm to investigate the sense of ownership, one can question whether the same type of mechanisms are at stake for the ownership of the rubber hand, which is merely seen, and for the ownership of a biological hand, which can be moved and experienced from the inside independently of any visual feedback. A recent lesion study showed dissociation between patients who failed to experience the RHI and patients who denied ownership of their own hand (Zeller et al., 2011). The authors concluded that the RHI recruits different brain regions than those involved for the sense of ownership

1 of one's biological body. One can also question whether there are alternative accounts
2 of ownership that do not appeal to a "Whose" system despite the existence of shared
3 body representations. I will now argue that the existence of naked body content does
4 not necessarily lead to confusion between self and others.
5
6
7
8
9

10 11 12 13 4.2 The Janus head hypothesis 14 15

16 The debate here is far broader than about some putative "Whose" system. What is
17 at stake is whether the exploitation of common resources prevents or reduces the
18 modularity of the mind. According to the modular view, the brain is organized in
19 specialized separately modifiable cognitive abilities that use specific, dedicated neural
20 resources. A modularist may for example claim that some are dedicated to the self and
21 others are dedicated to social cognition. However, more and more evidence indicates
22 that brain regions are not recruited by a single task. Rather, they are recycled to
23 support numerous cognitive functions (e.g., Anderson, 2010, Dehaene and Cohen,
24 2007; Goldman, 2012). In other words, they are originally established for one purpose
25 and reused for a different cognitive purpose. Recycling makes sense from an
26 evolutionary perspective insofar as it is more parsimonious than developing new
27 neural systems. But does it challenge the modular conception of the mind?
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44

45 The fact that there are modules does not preclude those modules from sharing parts
46 wherever possible (Carruthers, 2006, p. 23): "As a result, what we should predict is
47 that while there will be many modules, those modules should share parts wherever
48 this can be achieved without losing too much processing efficiency". This may work
49 along the "time-sharing model" offered by Jungé and Dennett (2010, p. 278): "(1) At
50 any given time, one high-level process uses the "workings" of multiple lower-level
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

1 areas, and (2) numerous high-level processes are hypothesized to alternately access a
2 common pool of specialized lower-level resources”. The point is that modules are not
3 exhausted by the parts that are shared. If the parts that are not shared are disrupted or
4 modified, then it alters only one module and not the other. Likewise, shared body
5 representations do not mean the end of the boundary between self and others. What is
6 shared is the representation of the rough structure of the body, either for action or for
7 perception. But this rough structure needs to be filled in for a full-fleshed spatial
8 representation of one’s body, including information about body metrics for example,
9 which is highly specific and can hardly be shared. Hence, some processing of bodily
10 information constitutes a common resource between self and other people, which can
11 in turn be used for one or the other. But what is shared, and thus naked, is only one
12 component of more complex representations of the body that are not shared. Those
13 more complex body representations may be called “superficial schema” (Head and
14 Holmes, 1911), “body structural description” (Schwoebel and Coslett, 2005) or “long-
15 term body image” (O’Shaughnessy, 1980). All refer to more or less the same notion,
16 that is, the representation of the spatial configuration and the dimension of the body,
17 what I call myself a body blueprint. One peculiarity of the body blueprint is that its
18 spatial organization differs whether it is used to shape bodily experiences or to guide
19 action.

20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47 To recap, some components of the body blueprint are shared between self and
48 others, but some components only. This is confirmed by brain imaging studies, which
49 never show perfect identity between activation for the self and activation for others.
50
51 For instance, vicarious touch can activate SI, but this hardly exhausts the neural basis
52 of the representation of the spatial properties of one’s body, which includes brain
53 areas at a higher level, including the parietal area. There are differences even in visual
54
55
56
57
58
59
60
61
62
63
64
65

1 representations of the body. For example, it was found that in the extrastriate body
2 area in the visual cortex, different brain regions selectively responded to images of
3 one's own body parts or other people's body parts (Myers and Sowden, 2008).
4
5 Furthermore, it was found an implicit self-advantage in body visual recognition
6
7 (Frassinetti et al., 2008, 2012). Participants were asked to match pictures of body
8
9 parts together. Their performance improved when the pictures displayed their own
10
11 body parts from an egocentric perspective compared to when they display other
12
13 people's body parts from the same perspective. Hence, they were better in visually
14
15 processing their own body that any other bodies.
16
17
18
19
20
21

22 The fact that self and others are not fully confounded is also confirmed by
23
24 neuropsychological dissociations. For example, patients with anorexia nervosa were
25
26 asked to imagine walking through a door-like aperture and then to judge whether or
27
28 not they would be able to walk at a normal speed without turning sideways (Guardia
29
30 et al., 2012). Alternatively they were asked to imagine another person of the same size
31
32 walking through the aperture and to judge whether she could pass. It was found that
33
34 the patients mistakenly judged that they could not pass in apertures in which they
35
36 accurately judged other people could pass. One way to interpret the results is that the
37
38 blueprint of their own body was impaired with no consequence on the representation
39
40 of other people's bodies. Conversely, patients with heterotopagnosia have selective
41
42 difficulties in locating another individual's body parts on her body, but no difficulty
43
44 on their own body. Rather than pointing on another's body, they indicate the location
45
46 of the named body part on their own body (Felician et al., 2003). Consequently,
47
48 deficits of the blueprint of one's body do not necessarily lead to intersubjective
49
50 impairments.
51
52
53
54
55
56
57
58

59 We can then propose the following hypothesis:
60
61
62
63
64
65

Janus head hypothesis: Intersubjective correspondence is achieved by time-sharing processing of bodily information that is common to the blueprint of one's body and the blueprint of other people's bodies.

On this view, neural resources that were originally designed to represent one's bodily states and were later shaped to represent other people's states are Janus-faced. They face inward as representation of one's body and they face outward as representation of other people's bodies. They can thus bridge the gap between one's body and other people's bodies, but without losing the distinction between self and others. At the level of the body blueprint, representations of one's body still differ from representations of other people's bodies. They are not shared, and thus, they cannot lead to confusion. It then seems that one can dispense with a "Whose" system despite the existence of shared body representations. The content is not naked at all levels. The blueprint of one's body is not exhausted by what is shared. It can thus keep its self-specificity, and thus, suffice to ground the sense of body ownership.

Conclusion

Do we need a "Whose" system like we have a "Who" system? Here I have addressed these questions by analysing the implications of intersubjective correspondence for the content of body representations. I have defended a limited embodied approach to the problem, according to which intersubjective correspondence is encoded in bodily terms in some – but not all – situations. There are shared body representations endowed with naked content, both in the motor and in the perceptual domains, but that they do not exhaust the blueprint of the body, that is, the representation of the spatial properties of the body. Some components of the body

1 blueprints are naked, but others are not. Hence, body blueprints do not leave the body
2 unattributed. They are thus susceptible to ground the sense of body ownership. It thus
3
4 seems that one can dispense with a WHOSE system despite the existence of shared
5
6
7 body representations.
8
9

10 11 12 **Acknowledgement**

13
14
15
16 This paper is dedicated to Professor Marc Jeannerod, who inspired all my work. I
17
18 would also like to thank Pierre Jacob for his very helpful comments.
19
20
21
22
23

24 **References**

- 25
26
27 Anderson, M. (2010). Neural reuse: A fundamental organizational principle of the
28
29 brain. *Behavioral and Brain Sciences*, 33, 4, 245-313.
30
31
32 Avenanti A, Buetti D, Galati G, Aglioti SM (2005) Transcranial magnetic stimulation
33
34 highlights the sensorimotor side of empathy for pain. *Nat Neurosci* 8(7):955–
35
36 960
37
38
39 Avenanti, A., Minio-Paluello, I., Sforza, A. and Aglioti, S.. (2009). Freezing or
40
41 Escaping? Opposite Modulations of Empathic Reactivity to the Pain of Others.
42
43 *Cortex* 45 (9): 1072–77.
44
45
46
47 Bach P, Peatfield NA, Tipper SP. (2007). Focusing on body sites: the role of spatial
48
49 attention in action perception. *Exp Brain Res*. 178(4):509-17.
50
51
52
53 Banissy MJ, Ward J (2007). Mirror-touch synaesthesia is linked with empathy. *Nat*
54
55 *Neurosci* 10:815–816
56
57
58
59
60
61
62
63
64
65

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Blakemore S-J, Bristow D, Bird G, Frith C, Ward J (2005) Somatosensory activations during the observation of touch and a case of vision-touch synaesthesia. *Brain* 128:1571–1583
- Botvinick, M., and Cohen, J. (1998). Rubber hands ‘feel’ touch that eyes see. *Nature*, 391, 756.
- Botvinick, M., Jhab, A., Bylsmaa, L., Fabian, S., Solomon, P. and Prkachin, K. (2005). “Viewing Facial Expressions of Pain Engages Cortical Areas Involved in the Direct Experience of Pain.” *Neuroimage* 251:312–19.
- Brass M, Bekkering H, Prinz W. (2001). Movement observation affects movement execution in a simple response task. *Acta Psychol (Amst)*. 106(1-2):3-22.
- Brass, M., & Heyes, C. (2005). Imitation: Is cognitive neuroscience solving the correspondence problem? *Trends in Cognitive Sciences*, 9, 489-495.
- Brozzoli, C., Makin, T.R., Cardinali, L., Holmes, N.P., Farnè, A. (2012) Peripersonal Space: A Multisensory Interface for Body–Object Interactions. In MM Murray, MT Wallace (eds). *The Neural Bases of Multisensory Processes*. Boca Raton (FL): CRC Press
- Buccino, G., Binkofski, F., Fink, G. R., Fadiga, L., Fogassi, L., Gallese, V., Seitz, R. J., Zilles, K., Rizzolatti, G. and Freund, H.-J. (2001). Action observation activates premotor and parietal areas in a somatotopic manner: An fMRI study. *European Journal of Neuroscience*, 13, 400-404.
- Buccino, G., Lui, F., Canessa, N., Patteri, I., Lagravinese, G., Benuzzi, F., Porro, C. A., & Rizzolatti, G. (2004). Neural circuits involved in the recognition of actions performed by nonconspecifics: An FMRI study. *Journal of Cognitive Neuroscience*, 16(1), 114-126.

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Bufalari, I., Aprile, T., Avenanti, A., Di Russo, F., Aglioti, S. (2007). Empathy for Pain and Touch in the Human Somatosensory Cortex. *Cerebral Cortex* 17 (11): 2553–61.
- Buxbaum LJ, Giovannetti T, Libon D. (2000). The role of the dynamic body schema in praxis: evidence from primary progressive apraxia. *Brain Cogn.* 44(2):166-91.
- Cardellicchio P, Sinigaglia C, Costantini M. (2012). Grasping affordances with the other's hand: A TMS study. *Soc Cogn Affect Neurosci.*
- Carruthers, P. (2006). *The architecture of the mind: Massive modularity and the flexibility of thought.* Clarendon Press.
- Chaminade T, Meltzoff AN, Decety J. (2005). An fMRI study of imitation: action representation and body schema. *Neuropsychologia.* 43(1):115-27.
- Costantini M, Ambrosini E, Tieri G, Sinigaglia C, Committeri G (2010) Where does an object trigger an action? An investigation about affordances in space. *Experimental Brain Research* 207: 95.
- Costantini M, Committeri G, Sinigaglia C. (2011). Ready both to your and to my hands: mapping the action space of others. *PLoS One.* 6(4):e17923.
- Cook R, Press C, Dickinson A, Heyes C. (2010). Acquisition of automatic imitation is sensitive to sensorimotor contingency. *J Exp Psychol Hum Percept Perform.* 36(4):840-52
- Cooke, D. F., and Graziano, M. S. A. (2003). Defensive movements evoked by air puff in monkeys. *Journal of Neurophysiology*, 90, 3317–3329.

- 1 Csibra G (2007) Action mirroring and action interpretation: An alternative account. In
2 Haggard P, Rosetti Y, Kawato M (eds) Sensorimotor foundations of higher
3 cognition. Attention and performance XXII. Oxford University Press, Oxford
4
5
6
7 Dehaene, S. & Cohen, L. (2007) Cultural recycling of cortical maps. *Neuron* 56:384–
8 98.
9
10
11
12 de Vignemont F (2004) The co-consciousness hypothesis. *Phenomenol Cogn Sci*
13 3:97–114
14
15
16
17
18 de Vignemont, F. (2010) Body schema and body image: pros and cons.
19 *Neuropsychologia*, 48(3), 669–80.
20
21
22
23
24 de Vignemont, F., Tsakiris, M., Haggard, P. (2005), Body mereology. In *Human*
25 *Body Perception from the Inside Out*, edited by G. Knoblich, I.M. Thornton, M.
26 Grosjean, M. Shiffrar. New York: Oxford University Press, 147-170.
27
28
29
30
31 de Vignemont, F., Majid, A., Jolla, C., and Haggard, P. (2009). Segmenting the body
32 into parts, evidence from biases in tactile perception. *Quarterly Journal of*
33 *Experimental Psychology*, 3, 1–13.
34
35
36
37
38
39 de Vignemont, F. and Jacob, P. (2012), What it's like to feel another's pain.
40 *Philosophy of Science*, 79, 2, 295-316.
41
42
43
44
45 Dijkerman, H. C., and de Haan, E. H. (2007). Somatosensory processes subserving
46 perception and action. *The Behavioral and Brain Sciences*, 30, 189–201.
47
48
49
50 Di Pellegrino, G., Fadiga, L., Fogassi, L., Gallese, V., & Rizzolatti, G. (1992).
51 Understanding motor events: a neurophysiological study. *Experimental Brain*
52 *Research*, 91, 176-180.
53
54
55
56
57
58
59
60
61
62
63
64
65

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Ebisch SJ, Perrucci MG, Ferretti A, Del Gratta C, Romani GL, Gallese V. (2008). The sense of touch: embodied simulation in a visuotactile mirroring mechanism for observed animate or inanimate touch. *J Cogn Neurosci*. 20(9):1611-23.
- Ehrsson HH, Holmes NP, Passingham RE. (2005). Touching a rubber hand: feeling of body ownership is associated with activity in multisensory brain areas. *J Neurosci*. 25(45):10564-73.
- Felician O, Ceccaldi M, Didic M, Thinus-Blanc C, Poncet M. (2003). Pointing to body parts: a double dissociation study. *Neuropsychologia*. 41(10):1307-16.
- Ferrari PF, Rozzi S, Fogassi L (2005) Mirror neurons responding to observation of actions made with tools in monkey ventral premotor cortex. *J Cogn Neurosci* 17:212–226
- Frassinetti F, Maini M, Romualdi S, Galante E, Avanzi S. (2008). Is it mine? Hemispheric asymmetries in corporeal self-recognition. *J Cogn Neurosci*. 20(8):1507-16.
- Frassinetti F, Fiori S, D'Angelo V, Magnani B, Guzzetta A, Brizzolara D, Cioni G. (2012). Body knowledge in brain-damaged children: a double-dissociation in self and other's body processing. *Neuropsychologia*. 50(1):181-8.
- Funk, M., Shiffrar, M., & Brugger, P. (2005). Hand movement observation by individuals born without hands: Phantom limb experience constrains visual limb perception. *Experimental Brain Research*, 164(3), 341-346.
- Gallagher, S. (2005). *How the body shapes the mind*. New York: Oxford University Press.

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Gallagher, S., & Meltzoff, A. N. (1996). The earliest sense of self and others: Merleau-Ponty and recent developmental studies. *Philosophical Psychology*, 9, 211-233
- Gallese V (2001) The 'Shared Manifold' hypothesis: from mirror neurons to empathy. *J Conscious Stud* 8(5-7):33-50
- Gallese V (2007) Embodied simulation: from mirror neuron systems to interpersonal relations. *Novartis Found Symp* 278:3-12
- Gallese, V., Fadiga, L., Fogassi, L., & Rizzolatti, G. (1996). Action recognition in the premotor cortex. *Brain*, 119, 593-609.
- Gallese V, Sinigaglia C. (2011). What is so special about embodied simulation? *Trends Cogn Sci*. 15(11):512-9.
- Gazzola V, Rizzolatti G, Wicker B, Keysers C. (2007a). The anthropomorphic brain: the mirror neuron system responds to human and robotic actions. *Neuroimage*. 35(4):1674-84.
- Gazzola, V., van der Worp, H., Mulder, T., Wicker, B., Rizzolatti, G., & Keysers, C. (2007b). Aphasics born without hands mirror the goal of hand actions with their feet. *Current Biology*, 17(14), 1235-1240.
- Georgieff, N. and Jeannerod, M. (1998) Beyond consciousness of external reality: a "who" system for consciousness of action and self-consciousness. *Conscious Cogn*. 7(3):465-77.
- Gillmeister H, Catmur C, Liepelt R, Brass M, Heyes C. (2008). Experience-based priming of body parts: a study of action imitation. *Brain Res*. 1217:157-70

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Goldenberg G. (1995) Imitating gestures and manipulating a mannikin--the representation of the human body in ideomotor apraxia. *Neuropsychologia*. 33(1):63-72.
- Goldenberg G. (2009). Apraxia and the parietal lobes. *Neuropsychologia*. 47(6):1449-59
- Goldman, A.I. (2006). *Simulating minds*. Oxford: Oxford University Press
- Goldman, A.I. (2011) Two Routes to Empathy: Insights from Cognitive Neuroscience. In A. Coplan and P. Goldie (eds.), *Empathy: Philosophical and Psychological Perspectives*, pp. 31-44. Oxford University Press (2011).
- Goldman, A.I. (2012). A Moderate Approach to Embodied Cognitive Science. In A. Alsmith, & de F. de Vignemont (eds), *The body represented/Embodied representation*. *Review of Philosophy and Psychology*, Special issue, 3 (1).
- Goldman, A.I. and de Vignemont, F. (2009). Is social cognition embodied? *Trends in Cognitive Sciences*, 13(4), 154–9.
- Graziano, M.S.A. and Gross, C.G. (1993). A bimodal map of space: somatosensory receptive fields in the macaque putamen with corresponding visual receptive fields. *Exp Brain Res*, 97:96-109.
- Grezes J, Decety J (2001) Functional anatomy of execution, mental simulation, observation, and verb generation of actions: a meta-analysis. *Hum Brain Mapp* 12(1):1–19
- Guardia D, Conversy L, Jardri R, Lafargue G, Thomas P, Dodin V, Cottencin O, Luyat M. (2012). Imagining one's own and someone else's body actions: dissociation in anorexia nervosa. *PLoS One*. 7(8):e43241

- 1 Hach S, Ishihara M, Keller PE, Schütz-Bosbach S. (2011). Hard and fast rules about
2 the body: contributions of the action stream to judging body space. *Exp Brain*
3
4
5 Res. 212(4):563-74.
6
- 7 Haggard P. (2006). Just seeing you makes me feel better: interpersonal enhancement
8 of touch. *Soc Neurosci.* 1(2):104-10.
9
- 10
11
12
13 Head, H., and Holmes, G. (1911). Sensory disturbances from cerebral lesions. *Brain*,
14
15 34, 102–254.
16
- 17
18 Heyes, C. (2001). Causes and consequences of imitation. *Trends Cogn Sci.* 5(6):253-
19
20 261.
21
22
- 23
24 Iacoboni M, Molnar-Szakacs I, Gallese V, Buccino G, Mazziotta JC, Rizzolatti G
25
26 (2005) Grasping the intentions of others with one's own mirror neuron system.
27
28 *PLoS Biol* 3(3):529–535
29
- 30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- 66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99
100
- 101
102
103
104
105
106
107
108
109
110
111
112
113
114
115
116
117
118
119
120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
- 166
167
168
169
170
171
172
173
174
175
176
177
178
179
180
181
182
183
184
185
186
187
188
189
190
191
192
193
194
195
196
197
198
199
200
- 201
202
203
204
205
206
207
208
209
210
211
212
213
214
215
216
217
218
219
220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240
241
242
243
244
245
246
247
248
249
250
251
252
253
254
255
256
257
258
259
260
261
262
263
264
265
266
267
268
269
270
271
272
273
274
275
276
277
278
279
280
281
282
283
284
285
286
287
288
289
290
291
292
293
294
295
296
297
298
299
300
- 301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340
341
342
343
344
345
346
347
348
349
350
351
352
353
354
355
356
357
358
359
360
361
362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
- 401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445
446
447
448
449
450
451
452
453
454
455
456
457
458
459
460
461
462
463
464
465
466
467
468
469
470
471
472
473
474
475
476
477
478
479
480
481
482
483
484
485
486
487
488
489
490
491
492
493
494
495
496
497
498
499
500
- 501
502
503
504
505
506
507
508
509
510
511
512
513
514
515
516
517
518
519
520
521
522
523
524
525
526
527
528
529
530
531
532
533
534
535
536
537
538
539
540
541
542
543
544
545
546
547
548
549
550
551
552
553
554
555
556
557
558
559
560
561
562
563
564
565
566
567
568
569
570
571
572
573
574
575
576
577
578
579
580
581
582
583
584
585
586
587
588
589
590
591
592
593
594
595
596
597
598
599
600
- 601
602
603
604
605
606
607
608
609
610
611
612
613
614
615
616
617
618
619
620
621
622
623
624
625
626
627
628
629
630
631
632
633
634
635
636
637
638
639
640
641
642
643
644
645
646
647
648
649
650
651
652
653
654
655
656
657
658
659
660
661
662
663
664
665
666
667
668
669
670
671
672
673
674
675
676
677
678
679
680
681
682
683
684
685
686
687
688
689
690
691
692
693
694
695
696
697
698
699
700
- 701
702
703
704
705
706
707
708
709
710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727
728
729
730
731
732
733
734
735
736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
- 801
802
803
804
805
806
807
808
809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828
829
830
831
832
833
834
835
836
837
838
839
840
841
842
843
844
845
846
847
848
849
850
851
852
853
854
855
856
857
858
859
860
861
862
863
864
865
866
867
868
869
870
871
872
873
874
875
876
877
878
879
880
881
882
883
884
885
886
887
888
889
890
891
892
893
894
895
896
897
898
899
900
- 901
902
903
904
905
906
907
908
909
910
911
912
913
914
915
916
917
918
919
920
921
922
923
924
925
926
927
928
929
930
931
932
933
934
935
936
937
938
939
940
941
942
943
944
945
946
947
948
949
950
951
952
953
954
955
956
957
958
959
960
961
962
963
964
965
966
967
968
969
970
971
972
973
974
975
976
977
978
979
980
981
982
983
984
985
986
987
988
989
990
991
992
993
994
995
996
997
998
999
1000

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Jeannerod, M. & Pacherie, E. (2004). Agency, Simulation and Self-identification. *Mind & Language*, 19, 2: 113-146
- Jungé, J.A and Dennett, D.C. (2010) Multi-use and constraints from original use. *Behavioral and Brain Sciences*, 33, 4.
- Kennett, S., Taylor-Clarke, M., Haggard, P. 2001. Noninformative vision improves the spatial resolution of touch in humans. *Curr Biol*. 11(15), 1188–91.
- Keysers, C., Wicker, B., Gazzola, V., Anton, J.L., Fogassi, L. and Gallese, V. (2004). A Touching Sight: SII/PV Activation during the Observation and Experience of Touch. *Neuron* 42:335–46.
- Keysers C, Kaas JH, Gazzola V. (2010) Somatosensation in social perception. *Nat Rev Neurosci*. 11(6):417-28.
- Makin, T.R., Holmes, N.P., Ehrsson, H.H. (2008). On the other hand: dummy hands and peripersonal space. *Behav Brain Res*. 191(1), 1–10.
- Meltzoff, A. N., & Moore, M. K. (1995). Infants' Understanding of People and Things: From Body Imitation to Folk Psychology. In J.L. Bermudez, A. Marcel, & N. Eilan (Eds.), *The Body and Self* (pp. 43-69), MIT Press: Bradford
- Myers A, Sowden PT. (2008). Your hand or mine? The extrastriate body area. *Neuroimage*. 42(4):1669-77.
- O'Shaughnessy, B. (1980). *The Will*. Cambridge: Cambridge University Press.
- Paillard, J. (1980). Le corps situé et le corps identifié. Une approche psychophysiologique de la notion de schéma corporel. *Revue Medicale de la Suisse Romande*, 100, 129–141.

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
- Press C, Bird G, Flach R, Heyes C. (2005). Robotic movement elicits automatic imitation. *Brain Res Cogn Brain Res.* 25(3):632-40.
- Ramachandran VS, Brang D. (2009). Sensations evoked in patients with amputation from watching an individual whose corresponding intact limb is being touched. *Arch Neurol.* 66(10):1281-4.
- Reed CL, Farah MJ (1995) The psychological reality of the body schema: a test with normal participants. *J Exp Psychol Hum Percept Perform* 21:334–343
- Rizzolatti, G., Fadiga, L., Gallese, V., & Fogassi, L. (1995). Premotor cortex and the recognition of motor actions. *Cognitive Brain Research*, 3, 131-141.
- Rochat, P. (1998). Self perception and action in infancy. *Experimental Brain Research*, 123, 102–109.
- Rumiati RI, Carmo JC, Corradi-Dell'Acqua C. (2009). Neuropsychological perspectives on the mechanisms of imitation. *Philos Trans R Soc Lond B Biol Sci.*364(1528):2337-47.
- Schaefer M, Heinze HJ, Rotte M. (2012) Close to you: embodied simulation for peripersonal space in primary somatosensory cortex. *PLoS One.* 7(8):e42308.
- Schwoebel, J., and Coslett, H. B. (2005). Evidence for multiple, distinct representations of the human body. *Journal of Cognitive Neuroscience*, 17, 543–553.
- Shiffrar M, Freyd JJ (1990) Apparent motion of the human body. *Psychol Sci* 1:257–264

- 1 Singer, T., Seymour, B., O'Doherty, J., Kaube, H., Dolan, R. and Frith, C. (2004).
2 Empathy for Pain Involves the Affective but Not Sensory Components of Pain.
3 Science 303:1157–62.
4
5
6
7
8 Spence, C., Pavani, F., Driver, J. (2004). Spatial constraints on visual-tactile cross-
9 modal distractor congruency effects. *Cogn Affect Behav Neurosci.* 4(2), 148–69.
10
11
12
13 Thomas R, Press C, Haggard P. (2006). Shared representations in body perception.
14 Acta Psychol (Amst). 121(3):317-30.
15
16
17
18 Tipper, S.P., Phillips, N., Dancer, C., Lloyd, D., Howard, L.A., McGlone, F. (2001).
19 Vision influences tactile perception at body sites that cannot be viewed directly.
20 Exp Brain Res. 139(2), 160–7.
21
22
23
24
25
26 Tsakiris, M. (2010). My body in the brain: a neurocognitive model of body-
27 ownership. *Neuropsychologia.* 48(3), 703-12.
28
29
30
31 Tsakiris, M., Haggard, P. (2005). The rubber hand illusion revisited: visuotactile
32 integration and self-attribution. *J Exp Psychol Hum Percept Perform.* 31(1), 80–
33 91
34
35
36
37
38
39 Tsakiris, M., Prabhu, G., Haggard, P. (2006). Having a body versus moving your
40 body: How agency structures body-ownership. *Conscious Cogn.* 15(2):423-32.
41
42
43
44
45 Wheaton KJ, Thompson JC, Syngeniotis A, Abbott DF, Puce A. (2004). Viewing the
46 motion of human body parts activates different regions of premotor, temporal,
47 and parietal cortex. *Neuroimage.* 22(1):277-88.
48
49
50
51
52
53 Wood R, Gallese V, Cattaneo L. (2010). Visuotactile empathy within the primary
54 somatosensory cortex revealed by short-latency afferent inhibition. *Neurosci*
55 *Lett.* 473(1):28-31.
56
57
58
59
60
61
62
63
64
65

1 Zeller D, Gross C, Bartsch A, Johansen-Berg H, Classen J. (2011). Ventral premotor
2 cortex may be required for dynamic changes in the feeling of limb ownership: a
3
4
5 lesion study. J Neurosci. 31(13):4852-7.
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24

25 Action mirroring is considered as evidence of shared action representation, but can it
26
27 be considered as evidence as well of shared body representation?
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65