

# The relationship between action, social and multisensory spaces

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1	The relationship between action, social
2	and multisensory spaces
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### 19 Abstract

20 Several spaces around the body have been described, contributing to interactions with 21 objects (peripersonal) or people (interpersonal and personal). The sensorimotor and 22 multisensory properties of action peripersonal space are assumed to be involved in the 23 regulation of social personal and interpersonal spaces, but experimental evidence is tenuous. 24 Hence, the present study investigated the relationship between multisensory integration and 25 action and social spaces. Participants indicated when an approaching social or non-social 26 stimulus was reachable by hand (reachable space), at a comfortable distance to interact with 27 (interpersonal space), or at a distance beginning to cause discomfort (personal space). They 28 also responded to a tactile stimulation delivered on the trunk during the approach of the visual 29 stimulus (multisensory integration space). Results showed that participants were mostly 30 comfortable with stimuli outside reachable space, and felt uncomfortable with stimuli well 31 inside it. Furthermore, reachable, personal and interpersonal spaces were all positively 32 correlated. Multisensory integration space extended beyond all spaces and correlated only 33 with personal space when facing a social stimulus. Considered together, these data confirm 34 that action peripersonal space contributes to the regulation of the social spaces, and that 35 multisensory integration is not specifically constrained by the spaces underlying motor action 36 and social interactions.

Keywords: PPS - IPS - social interactions - multisensory integration - reachability judgment comfort distance judgment

### 39 1. Introduction

40 The space immediately surrounding the body is of foremost importance for any living 41 being as it is the space in which physical interactions with the environment take place. During 42 the last decades, countless studies in cognitive neurosciences have fortified the idea that the 43 representation of space is functional, *i.e.*, the space offering information on the possibilities 44 of acting on objects must be processed differently by the brain than the space offering 45 information on the mere presence of objects with no possibilities to act on them. This view has led to the distinction between peripersonal space (PPS, i.e., within reach) and extrapersonal 46 space (i.e., beyond reach)<sup>1,2</sup>, which would be underpinned by different neural networks<sup>3,4</sup>. The 47 48 concept of PPS originates from single-unit electrophysiological studies in monkeys showing 49 that a number of neurons within the ventral premotor cortex, the parietal cortex and the 50 putamen responded more to objects presented in the near reachable space than objects 51 presented in the far unreachable space<sup>2,5,6</sup>. Thus, PPS has been conceived as an interface 52 between the body and the environment, contributing to the organisation of object-directed 53 motor actions, either in terms of approach when facing incentive objects or in terms of avoidance when facing threatening objects<sup>7,8</sup>. In line with this, neuroimaging studies revealed 54 55 that the mere observation of an object located in PPS triggered activation in the sensorimotor brain areas, including the reach-related area of the superior parieto-occipital cortex, and the 56 premotor and motor cortical areas<sup>9,10,11,12</sup>. As a consequence, the transient disruption of the 57 58 left motor cortex using transcranial magnetic stimulation has been shown to produce an alteration of the perception of objects located in PPS<sup>13</sup>. Likewise, corticospinal activity<sup>14</sup> and 59  $\mu$  rhythm desynchronization<sup>15,16</sup> increased in the presence of objects in the near (vs. far) space, 60 61 similar to what has been observed during the preparation and execution of objects-directed motor actions<sup>17,18</sup>. Moreover, modifying the actual reaching-by-hand capabilities (e.g., through 62 63 tool-use or limb immobilisation), or biasing the spatial consequences of object-directed actions, entailed a congruent increase or decrease of the PPS<sup>19,20,21,22</sup>. Altogether, these 64

results suggest that PPS is an action space, enabling access to motor-related information
 similar to those implied in the planning and execution of voluntary motor actions<sup>23</sup>.

As revealed by monkey electrophysiological studies, most PPS neurons are 67 multisensory in that they respond to stimuli in two or three different sensory modalities, with 68 69 overlapping receptive fields anchored onto the same body region<sup>2,5,6</sup>. In addition, neural and 70 behavioural investigations have consistently shown that stimuli in one sensory modality 71 enhance the processing of stimuli in another modality, especially when those stimuli are perceived as potentially interacting with our body<sup>8</sup>. Importantly, some of these neurons are 72 73 particularly responsive to a tactile stimulation delivered in co-occurrence with an approaching 74 visual stimulus, provided the two stimuli fall in the neuron's receptive fields<sup>2</sup>. This 75 multisensory integration is of particular relevance for interactions with the environment, which 76 require the position of external stimuli to be combined with information about different body 77 segments<sup>23</sup>, as reflected by higher-order activations of somatosensory and associative 78 areas<sup>24</sup>. Such multisensory integration has also been observed in humans, activating a frontoparietal network<sup>25,26,27</sup>, in relation to PPS<sup>28</sup>. However, the main line of evidence in 79 80 humans supporting multisensory integration in relation to PPS comes from behavioural 81 studies showing that the proximity of a visual/auditory stimulus from a certain body region 82 fastens the detection of a tactile stimulation on that body region, and the maximal distance at which such facilitation is observed (as compared to a unisensory control condition) is usually 83 used as a proxy of the PPS extent<sup>29,30,31,32</sup>. The scientific consensus is that the integration of 84 visual/auditory and tactile information would provide an interface between perception and 85 action allowing appropriate (re)actions towards (either threatening or incentive) objects to be 86 87 generated. The relevance of multisensory integration to action preparation and execution is 88 indeed supported by the studies on the effect of permanent or temporary damage to the 89 monkey's cortex showing a direct relationship between the PPS multisensory network and the accuracy of motor responses<sup>33,34,35,36</sup>. Furthermore, electric stimulation of the PPS 90 91 multisensory neurons in the monkey premotor and intraparietal cortex elicits a pattern of

movements that is compatible with defensive arm movements<sup>37</sup>, while the PPS multisensory
neurons in the parietal and precuneus cortex have been shown to discharge during arm
reaching movements towards the part of space corresponding to their visual receptive field<sup>2</sup>.
PPS represents thus a multisensory and sensorimotor interface mediating the physical
interactions between the body and the environment<sup>38</sup>.

97 Hence, if PPS consists in a multisensory interface dedicated to physical interactions 98 with the environment, the reachable and multisensory integration spaces are expected to 99 overlap. However, the wealth of studies on behavioural multisensory facilitation in humans 100 has highlighted a high degree of lability of the multisensory integration space, depending notably on the body region targeted by the tactile stimulation<sup>32</sup>. Indeed, when considering 101 102 similar experimental conditions (*i.e.*, the detection of a tactile stimulus in the presence of a 103 looming auditory stimulus), the extent of the multisensory integration space tended to be 104 shorter when the tactile stimulus was delivered on the hand (around 40 cm), than on the face 105 (around 50 cm) or trunk (around 55 cm). Moreover, it is worth noting that the range of 106 distances leading to multisensory integration varied considerably across studies, even when 107 using the same experimental conditions (from 20 to 66 cm for the hand, from 17 to 86 cm for the head; from 25 to 80 for the trunk<sup>29,30,31,32,39,40,41,42,43,44,45,46,47,48,49,50,51,52</sup>. Hence, multisensory 108 integration does not seem to systematically overlap with the motor action space. In support 109 of this claim, Zanini and colleagues<sup>53</sup> found that the space corresponding to hand-centred 110 111 visuotactile integration was shorter than the space reachable with the hand, and moved with 112 the hand, while reachable space was insensitive to hand position. They concluded that multisensory and reachable spaces are distinct spatial representations. However, it is worth 113 114 underlying that the observed dissociation might also arise from the different frames of 115 reference involved in the two tasks. It is indeed known that object-directed action involves a stable trunk-centred frame of reference<sup>54,55</sup>. By contrast, multisensory integration was 116 117 thoroughly tested using a hand-centred or head-centred frame of reference, requiring, for motor action, to refer to a more global representation of the body constituting the 118

egocentre<sup>32,38,56,57</sup>. Accordingly, the "trunk-centred" reachable-by-hand space was not expected to coincide exactly with the "hand-centred" multisensory integration space. In line with this claim, Serino and colleagues<sup>32</sup> considered that "hand- and face-centred PPS are referenced to the trunk-centred PPS, which [is] a more extended representation of the space surrounding the body". Hence, multisensory integration might be compatible with the representation of the space that is reachable with the hand when referring to the same frame of reference, *i.e.*, a trunk-centred frame of reference, which has never been truly tested.

126 Another important aspect of the body-environment interactions concerns the nature of 127 the stimulus under consideration. Studies in social psychology have focused on interactions 128 with conspecifics instead of physical objects, and have typically divided the space around the 129 body in a series of bubbles that serve to maintain proper spacing between individuals. The 130 smallest bubble is the personal space (PS), which is defined as the space in which social 131 intrusion is felt to be threatening or uncomfortable<sup>,58</sup>. It is assumed to serve as a margin of 132 safety around the body and is typically assessed with discomfort distance judgments 133 requiring the participants to judge at which distance a confederate makes them 134 uncomfortable<sup>59,60,61,62</sup>. A second and larger bubble is the *interpersonal space* (IPS), which is 135 defined as the space one maintains between oneself and others during social interactions<sup>63</sup>. 136 It is typically assessed with comfort distance judgments requiring the participant to place a confederate at the most comfortable distance to interact with<sup>64,65</sup>. Not only do these social 137 spaces refer to the space surrounding the body as PPS, but also share common 138 139 characteristics with PPS. For instance, PPS is modulated by social factors such as the proximity of confederates and the relation that is held with them<sup>46,66, 67,68</sup>. Furthermore, both 140 141 PPS and social spaces shrink or enlarge depending on the emotional valence of the facing stimulus<sup>59,66,69</sup>. They are also both influenced by individual characteristics such as anxiety<sup>61</sup>. 142 These observations probably explain why several researchers in the last decades have taken 143 a closer look at the relationship between PPS and social spaces. Until now, studies have 144 mainly focused on the link between PPS and PS. For instance, lachini and colleagues<sup>70,60</sup> 145

146 reported that both spaces have a similar size (around 50cm) and are similarly affected by the 147 nature, age, and gender of the stimulus. They reported that both PPS and PS reduce with 148 humans as compared to robots and cylinders, with females as compared to males, and with 149 children as compared to adults. It has therefore been proposed that PPS, and more particularly 150 its sensorimotor and multisensory properties, serves as a spatial anchor to calibrate social distances<sup>23,71,72</sup>. In support of this claim, Quesque and colleagues<sup>62</sup> found that extending arm 151 length's representation through tool-us increased PPS with a concomitant effect on PS. Social 152 spaces seem thus rooted in the same sensorimotor representation as PPS<sup>23</sup>. However, the 153 154 above studies have mainly focused on the relative impact of different factors on the PPS and 155 social spaces, which provides little information about their relationship. Moreover, these 156 studies did not include a measure of IPS and thus failed to provide a comprehensive picture of the extent of the different social spaces and their relationship to PPS. Finally, the 157 158 involvement of multisensory integration in social spaces has not yet been studied in depth.

159 In this context, the present study investigated the relationship between the different 160 action and social spaces anchored on the body and multisensory integration. Participants had 161 to indicate when an approaching neutral visual stimulus (human, robot or lamp) was reachable 162 with the arm (indexing reachable space, RS), at the most comfortable distance to interact with 163 (indexing interpersonal space, IPS), or started to generate discomfort due to too much 164 proximity (indexing personal space, PS). We also included a visuotactile integration task that 165 required participants to respond as fast as possible to a tactile stimulation delivered on the trunk at various times of the approach of the visual stimulus, while ignoring the latter (indexing 166 multisensory integration space, MIS). We expected RS to overlap and correlate with MIS, as 167 168 being two representative measures of the trunk-centred PPS. Also, along with the idea that 169 the regulation of social distances is based on PPS representation<sup>23,71,72</sup>, we expected RS and 170 MIS to correlate with PS and IPS, although IPS should be larger and PS should be smaller than 171 RS and MIS<sup>60,62,70</sup>. Finally, all spaces should be similarly impacted by the nature of the stimulus,

with a preference for the lamp and robot to be kept at a larger distance compared to thehuman<sup>70</sup>.

### 174 2. Materials and Methods

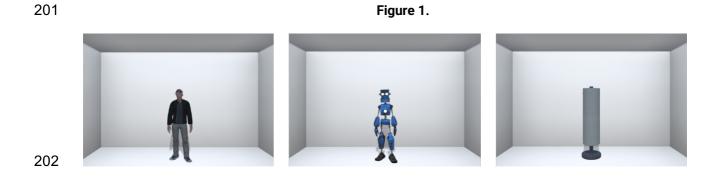
#### 175 2.1 Participants

176 Fifty-three participants from the Université of Lille participated in this study, but one participant was excluded because they missed 20% of the tactile stimulations in the 177 178 multisensory integration task, and two others were excluded because they showed no 179 multisensory facilitation effect, making it impossible to compute its location in space. The 180 final sample was thus composed of 50 participants (12 males, mean [M] age ± standard 181 deviation  $[SD] = 22.6 \pm 4.0$ ). A sample size analysis performed in G\*Power indicated that at least 41 participants were required to detect a small effect (Cohen's f = 0.15) with a high power 182 183 criterion (0.9) in a 4 x 3 repeated-measure ANOVA. All participants were right-handed and had 184 a normal or corrected-to-normal vision. They all gave written informed consent prior to the 185 experiment. The study was performed in accordance with the ethical standards of the 186 Declaration of Helsinki and was approved by the Research Ethics Board of the University of Lille (CESC Lille, Ref. 2021-515-S95). 187

### 188 2.2 Apparatus & stimuli

The virtual stimuli were presented through an HTC Vive Pro head-mounted display in a 189 190 virtual room measuring 6 x 5 x 3 m, and consisting of a white floor, a grey ceiling and grey 191 walls. The stimuli consisted of a human male avatar aged about 30 years, an anthropomorphic 192 robot and a cylindrical lamp. The man and robot looked straight ahead and showed a neutral 193 facial expression (Figure 1). The height of the stimuli was calibrated so that the eye level of the human and robot were aligned with the eye level of the participant. All stimuli had the 194 195 same height and width. We verified that the visual stimuli were perceived as neutral by requiring the participants to rate the emotional valence of each stimulus on the Self-196

Assessment Manikin (SAM) scale, a 9-points graphic Likert scale ranging from 1 (extremely negative) to 9 (extremely positive)<sup>73</sup>. One sample *t*-test to 5 (*i.e.*, neutral emotional valence) indicated that the human, t(49) = 0.47, p = .643, robot, t(49) = 1.85, p = .071, and lamp, t(49) =-1.24, p = .220, were similarly judged as neutral.



#### 203 2.3 Tasks & procedure

204 Participants were standing while holding a response button in their right hand, and 205 wearing the head-mounted display. A vibrotactile stimulator (DRV2605 Haptic Driver, Texas 206 Instruments) was fixed to their sternum with an elastic band. They performed the four 207 following tasks in a counterbalanced order:

#### 208 2.3.1 Reachability Distance Judgment

209 Participants were required to press the response button as soon as they judged being able to reach the approaching visual stimulus, without actually performing any reaching 210 211 movement. Each trial started with the appearance of a visual stimulus at 300 cm in front of 212 the participant for a duration of 500 ms, which then approached the participant at a velocity 213 of 0.75m/sec. Whenever the participant pressed the response button, the visual stimulus 214 stopped moving and remained still for 1000 ms before disappearing. The next trial started at 215 a random delay between 800 and 850 ms following the disappearance of the previous 216 stimulus. The task consisted of 18 trials (3 stimuli x 6 repetitions), lasted about 2 minutes, 217 and was used to assess RS.

#### 218 2.3.2 Comfort Distance Judgment

The same procedure as in the reachability distance judgment task was used, except that participants were required to press the response button as soon as the visual stimulus was judged at the most comfortable distance to interact with it. This task was used to assess IPS.

222 2.3.3 Discomfort Distance Judgment

The same procedure as in the reachability distance judgment and comfort distance judgment tasks was used, except that participants were required to press the response button whenever the visual stimulus was at a distance that made them feel uncomfortable. This task was used to assess PS.

227 2.3.4 Multisensory Integration Task

228 Participants were required to respond as quickly as possible to a tactile stimulation (60 229 ms, 3.6 V, 250 Hz) delivered well above the detection threshold on their sternum while ignoring 230 the visual stimulus facing them. The task included 4 types of trials: bimodal visuotactile, 231 unimodal tactile, bimodal catch and unimodal catch trials. In all types of trials, the visual 232 stimulus appeared at 300 cm in front of the participants for 500 ms. In the bimodal visuotactile 233 trials, the stimulus moved towards the participants at a velocity of 0.75m/sec. A tactile 234 stimulation was delivered at one of the 8 following delays: 1333, 2000, 2267, 2533, 2800, 3067, 235 3333 or 3600 ms after the setting in motion of the visual stimulus. This means that the visual 236 stimulus was respectively at 200, 150, 130, 110, 90, 70, 50 and 30 cm from the participant at 237 the time the tactile stimulation occurred. Hence, the longer the delay, the closer the stimulus 238 from the participants. In the unimodal tactile trials, the tactile stimulation was provided after 239 1333, 2800 or 3600 ms, but the visual stimulus remained still. These trials served as baseline 240 and allowed us to investigate the facilitation effects induced by the spatial proximity of the visual stimulus while controlling that these effects were not merely due to the expectancy of 241 242 tactile stimulation or attention varying with temporal delay. In the bimodal catch trials, the 243 visual stimulus moved toward the participant until being at a distance of 20 cm, but no tactile 244 stimulation was delivered. In the unimodal catch trials, the visual stimulus remained still, but 245 no tactile stimulation was delivered. These catch trials were included to avoid automatic 246 motor responses and make sure that the participants were attentive to the task all along the 247 experiment. Whenever the participant pressed the response button, the visual stimulus 248 stopped moving and remained still for 1000 ms before disappearing. The next trial started at a random delay between 800 and 850 ms following the disappearance of the previous 249 250 stimulus. The whole task consisted of 414 trials, including 240 visuotactile bimodal (3 stimuli 251 x 8 delays x 10 repetitions), 90 unimodal (3 stimuli x 3 delays x 10 repetitions), 42 bimodal 252 catch (3 stimuli x 14 repetitions) and 42 unimodal catch (3 stimuli x 14 repetitions) presented in a random order. The trials were divided into 6 blocks of about 6 minutes intermingled with 253 254 5-minutes breaks. This task was used to assess MIS.

#### 255 2.4 Data analyses

The data were analysed using *R* (version 4.1.0) and *R Studio* software (version 1.3.1093). We first verified that our multisensory integration task succeeded in showing the typical effects of the tactile stimulation delay on reaction times (RT) in each of the three stimuli used (see Supplemental Materials for procedure and results).

#### 260 2.4.1 Extent of the different spaces.

261 To determine the individual extent of RS, PS and IPS, we averaged for each participant 262 and each stimulus the distance of the visual stimulus at the time of the response in the reachability judgement task, and in the discomfort and comfort distance judgement tasks, 263 264 respectively. The extent of MIS was determined by identifying the farthest distance at which 265 the bimodal trials induced facilitation effects as compared to the unimodal trials in the 266 visuotactile integration task (see Supplemental Materials for detailed procedure). We then 267 compared the different spaces in terms of their average extent and their sensitivity to the 268 nature of the visuals stimulus by entering the computed extents in a repeated-measures 269 ANOVA with the Space (RS, MIS, IPS, PS) and type of Stimulus (human, robot, lamp) as within-270 subject variables. Since the extent of MIS was an ordinal variable and the extent of the 271 different spaces, as well as the residuals of the model, did not follow a normal distribution, we 272 used an Aligned Rank Transform (ART) for nonparametric factorial ANOVAs as described by 273 Wobbrock and colleagues<sup>74</sup>. We planned to conduct pairwise comparisons on the significant 274 effects, but also on the effect of the Stimulus on each task, to investigate whether we replicate 275 the observation of expanded PPS and PS in the presence of a virtual human as compared to a virtual robot and a lamp<sup>70,60</sup> when using stimuli controlled for their (neutral) emotional 276 valence. The paired comparisons were performed using the ART<sup>74</sup> or ART-C<sup>75</sup> alignment 277 278 procedure, as appropriate to the requested contrast, and with Bonferroni correction.

279 2.4.2 Relationship between the different spaces.

We then further investigated the relationship between the different spaces with pairwise correlation analyses. We computed the correlation coefficients for each stimulus separately. In particular, we computed Pearson *r* coefficients, except when correlation included MIS, in which case we computed the Spearman *r* correlation coefficient for ordinal variables.

#### 284 2.4.3 Bayesian analyses

We also conducted the corresponding Bayesian analyses in JASP (with default values) in order to quantify the evidence in favour of an effect (H1) compared to an absence of effect. These analyses provided Bayes Factors ( $BF_{10}$ ) varying between 0 and  $\infty$ , where values below 1 provide increasing evidence in favour of the null hypothesis and values above 1 provide increasing evidence for the alternative hypothesis (H1/H0)<sup>76</sup>. A BF above 3 is typically considered sufficient evidence for the alternative hypothesis, while a BF below <sup>1</sup>/<sub>3</sub> is typically considered sufficient evidence for the null hypothesis<sup>77</sup>.

### 292 3. Results

### 293 3.1 Extent of the different spaces

294 The ANOVA comparing the extent of the different spaces and the sensitivity to the different visual stimuli showed a significant effect of Space, F(3,539) = 181.31, p < .001,  $\eta_p^2 =$ 295 0.502,  $BF_{10} = 6.94^{+66}$ . The average extent ± standard error [SE] was 127.40 ± 2.98 cm for MIS, 296 297 116.35 ± 4.05 cm for IPS, 91.36 ± 3.08 cm for RS and 53.47 ± 2.37 cm for PS. Post hoc pairwise 298 comparisons showed that all spaces were significantly different from each other (all p-values < .001; Figure 2A). There was no significant effect of the Stimulus, F(2,539) = 0.62, p = .536, 299  $\eta_p^2$  = .002, *BF*<sub>10</sub> = 0.04, or Space by Stimulus interaction, *F*(6,539) = 0.73, *p* = .626,  $\eta_p^2$  = .008, 300  $BF_{10} = 0.02$ . The planned comparisons, however, showed a significant effect of the Stimulus 301 on RS, F(2, 98) = 13.84, p < .001,  $\eta_p^2 = .220$ ,  $BF_{10} = 2018.02$ , with participants judging the human 302 303 as reachable at shorter distances (M ± SE = 86.24 ± 5.04 cm) than the robot (93.16 ± 5.55 cm), t(98) = -3.43, p = .002,  $BF_{10} = 3835.21$ , and the lamp (94.66 ± 5.41 cm), t(98) = -5.17, p < .001, 304  $BF_{10}$  = 355.93, while RS for the robot and lamp did not significantly differ from each other, t(98)305 = -1.74, p = .195,  $BF_{10} = 0.205$ . The effect of Stimulus was also significant for IPS, F(2, 98) =306 6.88, p = .002,  $\eta_p^2 = .123$ ,  $BF_{10} = 36.62$ . Post-hoc pairwise comparisons further indicated that 307 participants preferred to place the lamp at shorter distances (108.43 ± 6.65 cm) than the robot 308

309  $(121.34 \pm 7.34 \text{ cm}), t(98) = -3.42, p = .003, BF_{10} = 14.78, and the human (119.27 \pm 6.98 \text{ cm}),$ 310  $t(98) = -2.95, p = .011, BF_{10} = 3.02$ , while the preferred distance for the human and robot did 311 not significantly differ from each other,  $t(98) = -0.47, p = .884, BF_{10} = 0.271$ . By contrast, the 312 effect of the Stimulus was marginal (or null, according to Bayesian analyses) on PS, F(2, 98)313 = 3.01,  $p = .054, \eta_p^2 = .057, BF_{10} = 0.290$ , with only the lamp being tolerated closer than the 314 robot,  $t(98) = -2.45, p = .047, BF_{10} = 0.56$ . Finally, the effect of Stimulus on multisensory space 315 was not significant,  $F(2, 98) = 0.28, p = .752, \eta_p^2 = .005, BF_{10} = 0.089$ .

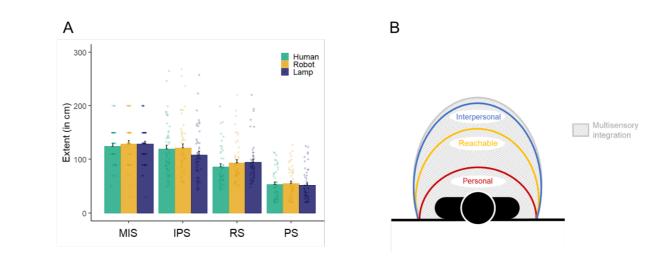


Figure 2.

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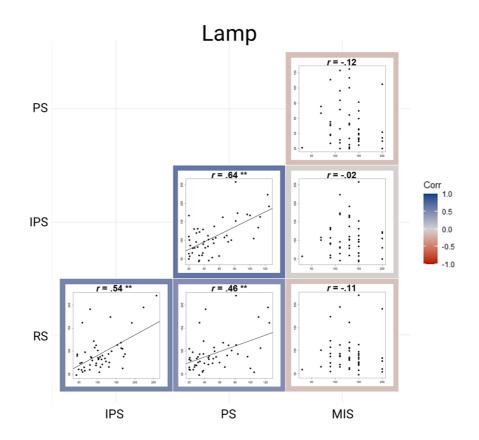
#### 318 3.2 Relationship between the different spaces

319 Regarding the lamp, a significant positive correlation was found between RS and IPS, r = .54, p < .001,  $BF_{10} = 594.58$ , between RS and PS, r = .46, p < .001, BF = 42.33, as well as 320 between IPS and PS, r = .64, p < .001,  $BF_{10} = 31768.77$ . The correlation between RS and MIS 321 was not significant, r = -.11, p = .443,  $BF_{10} = 0.28$ , so as the other correlations including MIS (all 322 *p*-values > .407, all  $BF_{10}$ -values < 0.27; Figure 3). Regarding the robot, we also found a 323 324 significant positive correlation between RS and IPS, r = .41, p = .003,  $BF_{10} = 11.23$ , RS and PS, r = .35, p = .014, BF<sub>10</sub> = 3.38, as well as between IPS and PS, r = .53, p < .001, BF<sub>10</sub> = 335.11. In 325 addition, there was a significant negative correlation between PS and MIS, r = -.45, p < .001, 326

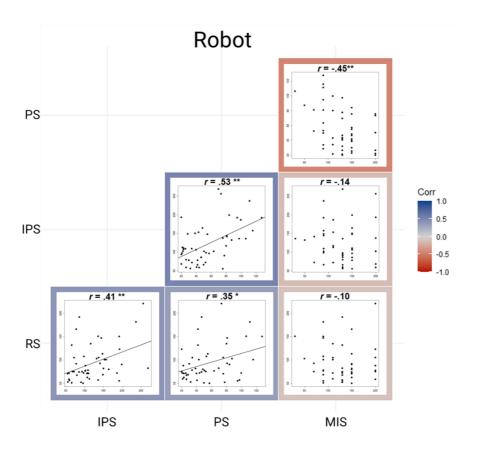
327  $BF_{10} = 63.84$ . No other correlation was significant (all *p*-values > .938, all  $BF_{10} < 0.31$ ), including 328 the correlation between RS and MIS, r = -.10, p = .499,  $BF_{10} = 0.23$  (Figure 4). Regarding the 329 human, we found the same significant correlations as in the robot: a positive correlation 330 between RS and IPS, r = .42, p = .003,  $BF_{10} = 13.92$ , and between RS and PS, r = .37, p = .007, 331  $BF_{10} = 5.40$ , PS and IPS, r = .59, p < .001,  $BF_{10} = 4169.65$ , as well as a negative relation between 332 PS and MIS, r = -.38, p = .006,  $BF_{10} = 8.94$ . There was no other significant correlation (*p*-values 333 >.210,  $BF_{10}$ -values < 0.439; Figure 5).

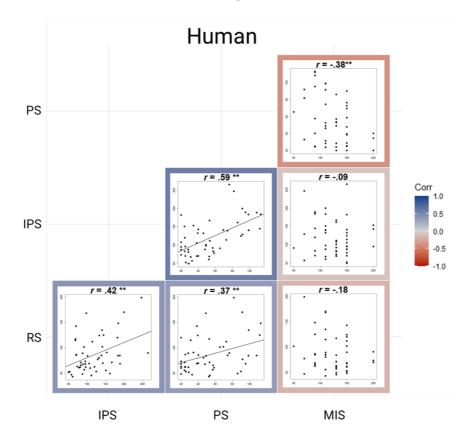












340 4. Discussion

341 The aim of the present study was to assess the relationship between the reachable and 342 multisensory spaces, two representative measures of the trunk-centred PPS, and how the latter related to the social spaces (interpersonal and personal). To do so, we required 343 344 participants to indicate when an approaching neutral visual stimulus (human, robot or lamp) 345 was reachable with the arm (RS), at the most comfortable distance to interact with (IPS), or 346 too close so that it generated a feeling of discomfort (PS). We also included a visuotactile integration task (MIS) that required participants to respond as fast as possible to tactile 347 stimulation delivered on the trunk at various times of the approach of the visual stimulus. 348 349 Based on the idea that PPS is an action space characterised by sensorimotor and 350 multisensory properties, we expected the extent of RS and MIS not only to overlap but also to 351 correlate. Moreover, along with the idea that PPS contributes to the regulation of the social spaces, we expected all spaces to correlate and to be similarly impacted by the nature of thestimulus, although PS should have the smallest extent and IPD the largest one.

354 The analyses of the extent of the different spaces showed that MIS was larger (127 cm) 355 than IPS (121 cm), which was, in turn, larger than RS (95 cm) and PS (58 cm, Figure 2B). This 356 indicates that, as expected, both objects and humanoids were preferentially placed outside 357 RS to interact with, and generated discomfort when present well inside it. The extent of RS is known to slightly overestimate arm length<sup>78</sup> (here 73.2  $\pm$  5.6 cm, corresponding for RS to an 358 overestimation of 29%), in particular in virtual environments<sup>79</sup>. Moreover, the relative extents 359 360 of the reachable and social (PS and IPS) spaces are in line with previous observations showing that IPS extent is typically between 80 and 140 cm<sup>65</sup>, while RS and PS extents are typically 361 smaller, i.e., between the range of 50-70 cm<sup>20,70,62</sup>. The data analysis conducted in the present 362 363 study further showed that the extent of PS is smaller than that of RS. This confirms the 364 previous findings highlighting that one feels progressively uncomfortable whenever RS is 365 violated<sup>80,81</sup>. It is also worth noting that the extent of RS and IPS were both affected by the 366 type of stimulus presented, even though the latter were all rated as neutral. As already shown<sup>60,70</sup>, RS was significantly shorter in the presence of a virtual human than in the presence 367 368 of a lamp or robot (7.7 cm in the present study). This confirms that PPS representation 369 expanded with virtual objects and reduced with virtual humans. Conversely, the extent of IPS 370 was not different between the robot and human, and significantly shorter with the lamp (11.9 371 cm in the present study). This might reflect the fact that interactions with objects require touching them and thus be at shorter distances than people for which interactions might be 372 primarily conceived as a verbal exchange, especially when the situation involves a stranger<sup>63</sup>. 373 374 The lack of difference between the robot and human stimulus might suggest that the 375 anthropomorphic aspect of the robot used in the present study was sufficient to consider 376 social interaction with it. It is indeed expected that human-like stimuli with the same (neutral) emotional valence should be positioned at the same IPS<sup>59,71,80</sup>. A complementary 377 interpretation could be that the human stimulus used in the present study was a male who 378

was shown to trigger larger IPS than a female human stimulus<sup>60,70</sup>. Further experiments would
be required to disentangle these different interpretations.

381 The correlational analyses revealed that RS, IPS and PS, although they were characterised by different extents, were positively correlated to each other, whatever the 382 383 stimulus presented. This means that the participants with a larger RS were also those who 384 had a larger IPS and PS, and conversely, whichever the stimulus presented. These data 385 confirm previous studies that highlighted that the regulation of PS depends in some respect on the representation of PPS<sup>62</sup>, although the outcome of the present study extends the 386 387 contribution of PPS also to IPS. The observed pattern of results, therefore, provides an additional argument for the involvement of PPS in the calibration of social spaces<sup>60,62,70,71</sup>, and 388 389 corroborates brain imaging studies showing that the frontoparietal network involved in the representation of PPS also supports social interactions<sup>82,83</sup>. Overall, these findings comfort 390 391 the idea that action and social spaces are related but more specifically that the sensorimotor 392 properties of PPS serve as a spatial reference to specify the appropriate social distances, as 393 suggested by the homeostatic theory of social interactions<sup>71</sup>. According to this theory, the 394 appropriate inter-individual distance corresponds to PPS plus an extra margin of safety, that 395 adapts according to the valence or level of threat endowed on conspecifics. This theory, 396 therefore, accounts for the observation that IPS correlates with RS but has a larger extent. In 397 its original form, the theory did not take into account PS and assumed that PPS is a protective buffer zone whose intrusion produces discomfort<sup>80,81</sup> and triggers defensive behaviour<sup>84</sup>. As 398 399 discussed above, the present study rather underlines that discomfort is experienced when 400 stimuli are well inside RS. PS is therefore a better candidate if we consider the priority space 401 dedicated to the protection of the body, although it seems calibrated from PPS representation. 402 This spatial relationship between PPS and PS would allow for PPS intrusion, at least to some 403 extent, which is often required during interactions both with objects and living beings.

404 The striking result of the present study is however the observation that trunk-centred 405 multisensory integration extended much further away than both reachable and social spaces, which is in contradiction with our initial hypothesis. Indeed, MIS was 11.05 cm larger than IPS, 406 407 36.04 cm larger than RS and 73.93 cm larger than PS. MIS extent is furthermore much larger 408 in the present study than what was previously observed with auditory stimuli when also using 409 a trunk-centred frame of reference (*i.e.*, around 55 cm, from 25 to 80)<sup>32</sup>. One potential explanation could be that multisensory integration extended more when facing meaningful 410 411 visual stimuli. A careful inspection of previous studies supports this hypothesis: hand-centred 412 and face-centred multisensory integration were found to be both more extended when facing virtual human characters (up to 127 and 150 cm, respectively)<sup>85,86</sup> than when facing looming 413 pink noise (up to 66 and 75 cm, respectively)<sup>42,49</sup>. However, even when centred on the same 414 415 trunk-centred frame of reference as the reachability task, MIS did not correspond to RS. This 416 result has two consequences. First, it indicates that multisensory integration is not specifically 417 related to the motor action space. Second, the fact that MIS encompasses both action space 418 and social spaces may suggest that multisensory integration contributes to the overall 419 interactions with objects and people in the environment, without specifically contributing to 420 the specification of the spaces where these interactions occur. These findings contrast with 421 the single-cell recording studies in monkeys showing that the receptive fields of the 422 multisensory neurons are within RS<sup>2</sup>. However, one may hypothesise that the sensory 423 facilitation reported in the behavioural studies and the neural mechanisms highlighted in the single-cell studies do not refer to the exact same multisensory integration process<sup>87</sup>. While 424 the link between the two has been strongly advocated<sup>29</sup>, it is apparent that the behavioural 425 426 multisensory facilitation effect in humans is more flexible than what was reported in single-427 cell studies. As evidence, multisensory facilitation in behavioural studies has been found to be altered by the valence or meaning of the visual/auditory stimulus<sup>66</sup>, individual traits such 428 as anxiety/phobia<sup>45</sup>, interoceptive traits<sup>40</sup>, bodily changes such as pregnancy<sup>41</sup> or limb 429 immobilisation<sup>88</sup>, and even lockdown experience<sup>85</sup>. Moreover, a number of studies indicated 430

431 that the visual/auditory stimulus does not have to target the same body part as the tactile 432 stimulation to trigger multisensory facilitation<sup>87</sup>. This might be because the behavioural 433 effects evidenced arose not only from the multisensory brain areas but also from their 434 interaction with other brain areas such as those involved in body representation<sup>89</sup> and object-435 directed action control<sup>7</sup>. Another aspect of the behavioural studies on humans is that they 436 implied a task-dependent motor response, while monkeys were generally studied in a passive condition. Thus, despite their pioneering role, single-cell studies might represent only a small 437 438 window onto the network underpinning multisensory integration in the context of goal-439 directed motor action and social interaction. This may explain the lack of correlation that we 440 found between MIS and RS, corroborated by the Bayesian analysis, albeit single-cell studies revealed a link between multisensory integration and arm RS<sup>2,5</sup>. From a behavioural 441 442 perspective, it seems thus that RS refers to a different spatial representation than MIS despite 443 being tested with a typical looming task and using the same spatial frame of reference. PPS, 444 as an action space, must thus be viewed as a sensorimotor interface anchored on the body 445 that involves, but does not depend on, multisensory integration. Moreover, the negative 446 correlation found between MIS and PS, although specific to the humanoid stimuli (human and 447 robot), could suggest that multisensory integration serves mostly a defensive purpose<sup>8</sup>. 448 People characterised by a larger MIS were also characterised by a shorter PS, which may 449 reveal an adaptive link between anticipation of physical contact with social stimuli and 450 acceptance of the proximity of these stimuli.

Another implication of MIS encompassing all other spaces is that multisensory processes, usually related to the action space, extend also to the social space. This is not that surprising since the need to combine several sensory cues is not restricted to interactions with objects but also applies to social stimuli. For instance, emotions are expressed through facial expressions but also voice such that visual and auditory cues integration is an essential part of emotion reading and more globally of social interactions<sup>90</sup>. Moreover, multisensory integration is assumed to allow the preparation of the body for action, either for the purpose 458 of defensive or approaching behaviour<sup>7,8,70,71</sup>. Physical contact with people, though less 459 frequent than with objects, is also experienced on a daily-base: we shake hands, hug, are 460 tapped on the shoulder to get our attention, or brush against each other in crowded 461 environments, with some of these contacts, for instance when concerning people with bad 462 intentions, being at risk for the body. The functional advantage provided by multisensory 463 integration is thus also relevant for social interactions to anticipate possible contact with others and programme appropriate actions and responses – for example, to avoid harmful 464 465 contact or shake hands properly with our interlocutor. A consequence of this approach is that 466 multisensory integration must be viewed as a process at hand during interactions with either 467 objects or individuals, which is not specific to the nature of the present stimulus or the type of interaction envisaged, and which thus seems not constitutive of the spaces underlying object-468 469 directed actions and social interactions.

470 In conclusion, this first study comparing PPS (RS and MIS) and the social spaces (PS 471 and IPS) showed that only (the action PPS was related to the social spaces. This finding confirms previous studies reporting that RS and PS are related<sup>60,62,70,80</sup>, but extends this 472 473 relationship to IPS. This further underlines the particular role of the sensorimotor aspects of 474 PPS in the regulation of the social spaces, providing new evidence in support of the 475 homeostatic theory of social interactions<sup>71</sup>. Multisensory integration was not restricted to 476 action PPS and social spaces, as it extended beyond all these spaces. This indicates that 477 multisensory integration is involved in interactions with objects and people, in relation to the 478 anticipatory aspects of these interactive behaviours, but does not specifically determine the 479 representation of both action PPS and social spaces. The specific role of multisensory 480 integration in the different interactions with the environment, therefore, remains to be further 481 clarified, paving the way for future research.

### 482 5. References

- 483 1. Previc, F. H. The neuropsychology of 3-D space. *Psychological Bulletin* **124**, 123–164
  484 (1998).
- 2. Rizzolatti, G., Scandolara, C., Matelli, M. & Gentilucci, M. Afferent properties of periarcuate
  neurons in macaque monkeys. II. Visual responses. *Behavioural Brain Research* 2, 147–
  163 (1981).
- 488 3. di Pellegrino, G. & Làdavas, E. Peripersonal space in the brain. *Neuropsychologia* 66, 126–
  489 133 (2015).
- 490 4. Cléry, J., Guipponi, O., Wardak, C. & Ben Hamed, S. Neuronal bases of peripersonal and
  491 extrapersonal spaces, their plasticity and their dynamics: Knowns and unknowns.
  492 *Neuropsychologia* **70**, 313–326 (2015).
- 493 5. Graziano, M. S. A. & Gross, C. G. A bimodal map of space: somatosensory receptive fields
  494 in the macaque putamen with corresponding visual receptive fields. *Exp Brain Res* 97,
  495 (1993).
- 496 6. Graziano, M. S. A., Yap, G. S. & Gross, C. G. Coding of Visual Space by Premotor Neurons.
  497 Science 266, 1054–1057 (1994).
- 498 7. Brozzoli, C., Makin, T. R., Cardinali, L., Holmes, N. P. & Farnè, A. Peripersonal Space: A
  499 Multisensory Interface for Body–Object Interactions. in *The Neural Bases of Multisensory*500 *Processes* (CRC Press/Taylor & Francis, 2012).
- 501 8. Graziano, M. S. A. & Cooke, D. F. Parieto-frontal interactions, personal space, and
  502 defensive behavior. *Neuropsychologia* 44, 845–859 (2006).
- 503 9. Bartolo, A. *et al.* Contribution of the motor system to the perception of reachable space:
  504 an fMRI study. *Eur J Neurosci* 40, 3807–3817 (2014).
- 50510. Ferri, F. et al. Intertrial Variability in the Premotor Cortex Accounts for Individual506Differences in Peripersonal Space. Journal of Neuroscience 35, 16328–16339 (2015).

- 507 11. Grafton, S. T., Fadiga, L., Arbib, M. A. & Rizzolatti, G. Premotor Cortex Activation during
  508 Observation and Naming of Familiar Tools. *NeuroImage* 6, 231–236 (1997).
- 509 12. Quinlan, D. J. & Culham, J. C. fMRI reveals a preference for near viewing in the human
  510 parieto-occipital cortex. *NeuroImage* 36, 167–187 (2007).
- 511 13. Coello, Y. *et al.* Perceiving What Is Reachable Depends on Motor Representations:
  512 Evidence from a Transcranial Magnetic Stimulation Study. *PLoS ONE* **3**, e2862 (2008).
- 513 14. Cardellicchio, P., Sinigaglia, C. & Costantini, M. The space of affordances: A TMS study.
  514 *Neuropsychologia* 49, 1369–1372 (2011).
- 515 15. Proverbio, A. M. Tool perception suppresses 10–12Hz μ rhythm of EEG over the
  516 somatosensory area. *Biological Psychology* 91, 1–7 (2012).
- 517 16. Wamain, Y., Gabrielli, F. & Coello, Y. EEG μ rhythm in virtual reality reveals that motor
   518 coding of visual objects in peripersonal space is task dependent. *Cortex* 74, 20–30 (2016).
- 519 17. Bestmann, S. & Duque, J. Transcranial Magnetic Stimulation: Decomposing the Processes
  520 Underlying Action Preparation. *Neuroscientist* 22, 392–405 (2016).
- 521 18. Pfurtscheller, G. & Neuper, C. Event-related synchronization of mu rhythm in the EEG over
  522 the cortical hand area in man. *Neuroscience Letters* 174, 93–96 (1994).
- 523 19. Bourgeois, J. & Coello, Y. Effect of visuomotor calibration and uncertainty on the 524 perception of peripersonal space. *Atten Percept Psychophys* **74**, 1268–1283 (2012).
- 525 20. Bourgeois, J., Farnè, A. & Coello, Y. Costs and benefits of tool-use on the perception of
  526 reachable space. *Acta Psychologica* 148, 91–95 (2014).
- 527 21. Leclere, N. X., Sarlegna, F. R., Coello, Y. & Bourdin, C. Sensori-motor adaptation to novel
  528 limb dynamics influences the representation of peripersonal space. *Neuropsychologia*529 131, 193–204 (2019).
- 530 22. Toussaint, L., Wamain, Y., Bidet-Ildei, C. & Coello, Y. Short-term upper-limb immobilization
  531 alters peripersonal space representation. *Psychological Research* 84, 907–914 (2020).
- 532 23. Coello, Y. & Fisher M. Foundations of embodied cognition. Volume 1: Perceptual and
- 533 emotional embodiment. (Routledge, Taylor & Francis Group, 2016).

534 24. Guipponi, O., Cléry, J., Odouard, S., Wardak, C. & Ben Hamed, S. Whole brain mapping of
535 visual and tactile convergence in the macaque monkey. *NeuroImage* **117**, 93–102 (2015).

536 25. Bremmer, F. *et al.* Polymodal Motion Processing in Posterior Parietal and Premotor
537 Cortex. *Neuron* 29, 287–296 (2001).

- 538 26. Cléry, J. *et al.* The Prediction of Impact of a Looming Stimulus onto the Body Is Subserved
  539 by Multisensory Integration Mechanisms. *J. Neurosci.* **37**, 10656–10670 (2017).
- 540 27. Serino, A., Canzoneri, E. & Avenanti, A. Fronto-parietal Areas Necessary for a Multisensory
- 541 Representation of Peripersonal Space in Humans: An rTMS Study. *Journal of Cognitive*542 *Neuroscience* 23, 2956–2967 (2011).
- 543 28. Brozzoli, C., Gentile, G., Petkova, V. I. & Ehrsson, H. H. fMRI Adaptation Reveals a Cortical
- 544 Mechanism for the Coding of Space Near the Hand. *Journal of Neuroscience* 31, 9023–
  545 9031 (2011).
- 546 29. Canzoneri, E., Magosso, E. & Serino, A. Dynamic Sounds Capture the Boundaries of 547 Peripersonal Space Representation in Humans. *PLoS ONE* **7**, e44306 (2012).
- 548 30. Canzoneri, E. *et al.* Tool-use reshapes the boundaries of body and peripersonal space
  549 representations. *Exp Brain Res* 228, 25–42 (2013).
- 31. Noel, J.-P., Pfeiffer, C., Blanke, O. & Serino, A. Peripersonal space as the space of the bodily
  self. *Cognition* 144, 49–57 (2015).
- 552 32. Serino, A. *et al.* Body part-centered and full body-centered peripersonal space 553 representations. *Sci Rep* **5**, 18603 (2015).
- 33. Ettlinger, G. & Kalsbeck, J. E. Changes in tactile discrimination and in visual reaching after
- 555 successive and simultaneous bilateral posterior parietal ablations in the monkey. *Journal*
- of Neurology, Neurosurgery & Psychiatry **25**, 256–268 (1962).
- 557 34. Faugier-Grimaud, S., Frenois, C. & Stein, D. G. Effects of posterior parietal lesions on
  558 visually guided behavior in monkeys. *Neuropsychologia* 16, 151–168 (1978).
- 559 35. Gallese, V., Murata, A., Kaseda, M., Niki, N. & Sakata, H. Deficit of hand preshaping after
- 560 muscimol injection in monkey parietal cortex: *NeuroReport* **5**, 1525–1529 (1994).

- 36. Rizzolatti, G., Matelli, M. & Pavesi, G. Deficits in attention and movement following removal
  of postarcuate (area 6) and prearcuate (area 8) cortex in macaque monkeys. *Brain* 106,
  655–673 (1983).
- 37. Cooke, D. F. & Graziano, M. S. A. Defensive Movements Evoked by Air Puff in Monkeys.
  Journal of Neurophysiology 90, 3317–3329 (2003).
- 38. Serino, A. Peripersonal space (PPS) as a multisensory interface between the individual
  and the environment, defining the space of the self. *Neuroscience & Biobehavioral Reviews*99, 138–159 (2019).
- 39. Amemiya, T., Ikei, Y. & Kitazaki, M. Remapping Peripersonal Space by Using Foot-Sole
  Vibrations Without Any Body Movement. *Psychol Sci* **30**, 1522–1532 (2019).
- 40. Ardizzi, M. & Ferri, F. Interoceptive influences on peripersonal space boundary. *Cognition*177, 79–86 (2018).
- 573 41. Cardini, F., Fatemi-Ghomi, N., Gajewska-Knapik, K., Gooch, V. & Aspell, J. E. Enlarged 574 representation of peripersonal space in pregnancy. *Sci Rep* **9**, 8606 (2019).
- 575 42. Di Cosmo, G. *et al.* Peripersonal space boundary in schizotypy and schizophrenia.
  576 Schizophrenia Research 197, 589–590 (2018).
- 577 43. Ferroni, F. *et al.* Schizotypy and individual differences in peripersonal space plasticity.
  578 *Neuropsychologia* 147, 107579 (2020).
- 44. Galli, G., Noel, J. P., Canzoneri, E., Blanke, O. & Serino, A. The wheelchair as a full-body tool
  extending the peripersonal space. *Front. Psychol.* 6, (2015).
- 45. Taffou, M. & Viaud-Delmon, I. Cynophobic Fear Adaptively Extends Peri-Personal Space. *Front. Psychiatry* 5, (2014).
- 583 46. Teneggi, C., Canzoneri, E., di Pellegrino, G. & Serino, A. Social Modulation of Peripersonal
  584 Space Boundaries. *Current Biology* 23, 406–411 (2013).
- 47. Hobeika, L., Taffou, M. & Viaud-Delmon, I. Social coding of the multisensory space around
  us. *R. Soc. open sci.* 6, 181878 (2019).

- 48. Hobeika, L., Taffou, M., Carpentier, T., Warusfel, O. & Viaud-Delmon, I. Capturing the
  dynamics of peripersonal space by integrating expectancy effects and sound propagation
  properties. *Journal of Neuroscience Methods* 332, 108534 (2020).
- 49. Noel, J.-P., Blanke, O., Magosso, E. & Serino, A. Neural adaptation accounts for the
  dynamic resizing of peripersonal space: evidence from a psychophysical-computational
  approach. *Journal of Neurophysiology* **119**, 2307–2333 (2018).
- 50. Maister, L., Cardini, F., Zamariola, G., Serino, A. & Tsakiris, M. Your place or mine: Shared
  sensory experiences elicit a remapping of peripersonal space. *Neuropsychologia* 70, 455–
  461 (2015).
- 596 51. Noel, J.-P. *et al.* Full body action remapping of peripersonal space: The case of walking.
  597 *Neuropsychologia* **70**, 375–384 (2015).
- 52. Pfeiffer, C., Noel, J., Serino, A. & Blanke, O. Vestibular modulation of peripersonal space
  boundaries. *Eur J Neurosci* 47, 800–811 (2018).
- 53. Zanini, A. *et al.* Peripersonal and reaching space differ: Evidence from their spatial extent
- and multisensory facilitation pattern. *Psychon Bull Rev* 28, 1894–1905 (2021).
- 54. Lacquaniti & Caminiti. Visuo-motor transformations for arm reaching. *European Journal of Neuroscience* 10, 195–203 (1998).
- 55. McIntyre, J., Stratta, F. & Lacquaniti, F. Short-Term Memory for Reaching to Visual Targets:
- 605 Psychophysical Evidence for Body-Centered Reference Frames. J. Neurosci. 18, 8423–
  606 8435 (1998).
- 56. Alsmith, A. J. T. & Longo, M. R. Where exactly am I? Self-location judgements distribute
  between head and torso. *Consciousness and Cognition* 24, 70–74 (2014).
- 57. Bertossa, F., Besa, M., Ferrari, R. & Ferri, F. Point Zero: A Phenomenological Inquiry into
  the Seat of Consciousness. *Percept Mot Skills* 107, 323–335 (2008).
- 611 58. Hediger, H. Studies of the psychology and behavior of captive animals in zoos and circuses.
- 612 (Books, Criterion Inc., 1955).

- 613 59. Cartaud, A., Lenglin, V. & Coello, Y. Contrast effect of emotional context on interpersonal
  614 distance with neutral social stimuli. *Cognition* 218, 104913 (2022).
- 615 60. Iachini, T., Coello, Y., Frassinetti, F. & Ruggiero, G. Body Space in Social Interactions: A
- 616 Comparison of Reaching and Comfort Distance in Immersive Virtual Reality. *PLoS ONE* 9,
  617 e111511 (2014).
- 618 61. Iachini, T., Ruggiero, G., Ruotolo, F., Schiano di Cola, A. & Senese, V. P. The influence of
  619 anxiety and personality factors on comfort and reachability space: a correlational study.
  620 Cogn Process 16, 255–258 (2015).
- 62. Quesque, F. *et al.* Keeping you at arm's length: modifying peripersonal space influences
  622 interpersonal distance. *Psychological Research* 81, 709–720 (2017).
- 623 63. Hall, E. T. *The Hidden Dimension*. (NY: Doubleday, 1966).
- 624 64. Perry, A., Nichiporuk, N. & Knight, R. T. Where does one stand: a biological account of
  625 preferred interpersonal distance. *Social Cognitive and Affective Neuroscience* 11, 317–326
  626 (2016).
- 65. Sorokowska, A. *et al.* Preferred Interpersonal Distances: A Global Comparison. *Journal of Cross-Cultural Psychology* 48, 577–592 (2017).
- 66. Pellencin, E., Paladino, M. P., Herbelin, B. & Serino, A. Social perception of others shapes
  one's own multisensory peripersonal space. *Cortex* **104**, 163–179 (2018).
- 631 67. Coello, Y., Quesque, F., Gigliotti, M.-F., Ott, L. & Bruyelle, J.-L. Idiosyncratic representation
- of peripersonal space depends on the success of one's own motor actions, but also the
  successful actions of others! *PLoS ONE* 13, e0196874 (2018).
- 634 68. Gigliotti, M. F., Soares Coelho, P., Coutinho, J. & Coello, Y. Peripersonal space in social
- 635 context is modulated by action reward, but differently in males and females. *Psychological*636 *Research* 85, 181–194 (2021).
- 637 69. Coello, Y., Bourgeois, J. & Iachini, T. Embodied perception of reachable space: how do we
  638 manage threatening objects? *Cogn Process* 13, 131–135 (2012).

- 639 70. lachini, T. *et al.* Peripersonal and interpersonal space in virtual and real environments:
  640 Effects of gender and age. *Journal of Environmental Psychology* **45**, 154–164 (2016).
- 71. Coello, Y. & Cartaud, A. The Interrelation Between Peripersonal Action Space and
  Interpersonal Social Space: Psychophysiological Evidence and Clinical Implications. *Front. Hum. Neurosci.* **15**, 636124 (2021).
- 644 72. Lloyd, D. M. The space between us: A neurophilosophical framework for the investigation
  645 of human interpersonal space. *Neuroscience & Biobehavioral Reviews* 33, 297–304 (2009).
- 646 73. Bradley, M. M. & Lang, P. J. Measuring emotion: The self-assessment manikin and the
- semantic differential. *Journal of Behavior Therapy and Experimental Psychiatry* 25, 49–59
  (1994).
- 74. Wobbrock, J. O., Findlater, L., Gergle, D. & Higgins, J. J. The aligned rank transform for
  nonparametric factorial analyses using only anova procedures. in *Proceedings of the SIGCHI Conference on Human Factors in Computing Systems* 143–146 (ACM, 2011).
- 652 75. Elkin, L. A., Kay, M., Higgins, J. J. & Wobbrock, J. O. An Aligned Rank Transform Procedure
  653 for Multifactor Contrast Tests. in *The 34th Annual ACM Symposium on User Interface*
- 76. Dienes, Z. Using Bayes to get the most out of non-significant results. *Front. Psychol.* 5, (2014).
- 657 77. Jeffreys, H. The theory of probability. (1988).

Software and Technology 754-768 (ACM, 2021).

- 658 78. Fischer, M. H. Estimating reachability: Whole body engagement or postural stability?
  659 *Human Movement Science* 19, 297–318 (2000).
- 660 79. Xiong, W., Yu, X. & Lee, Y.-C. The Difference in Measuring Reachability Distance between
- Using Virtual Reality Technology and Manual Measurement. in 2020 IEEE 7th International
   Conference on Industrial Engineering and Applications (ICIEA) 390–393 (IEEE, 2020).
- 80. Cartaud, A., Ruggiero, G., Ott, L., Iachini, T. & Coello, Y. Physiological Response to Facial
  Expressions in Peripersonal Space Determines Interpersonal Distance in a Social
  Interaction Context. *Front. Psychol.* 9, 657 (2018).

- 81. Kennedy, D. P., Gläscher, J., Tyszka, J. M. & Adolphs, R. Personal space regulation by the
  human amygdala. *Nat Neurosci* 12, 1226–1227 (2009).
- 82. Holt, D. J. *et al.* Neural Correlates of Personal Space Intrusion. *Journal of Neuroscience* 34,
  4123–4134 (2014).
- 83. Vieira, J. B., Pierzchajlo, S. R. & Mitchell, D. G. V. Neural correlates of social and non-social
  personal space intrusions: Role of defensive and peripersonal space systems in
  interpersonal distance regulation. *Social Neuroscience* **15**, 36–51 (2020).
- 673 84. Evans, G. W. & Wener, R. E. Crowding and personal space invasion on the train: Please
  674 don't make me sit in the middle. *Journal of Environmental Psychology* 27, 90–94 (2007).

85. Serino, S. *et al.* Sharpening of peripersonal space during the COVID-19 pandemic. *Current Biology* **31**, R889–R890 (2021).

- 86. Buck, L. E., Park, S. & Bodenheimer, B. Determining Peripersonal Space Boundaries and
  Their Plasticity in Relation to Object and Agent Characteristics in an Immersive Virtual
  Environment. in 2020 IEEE Conference on Virtual Reality and 3D User Interfaces (VR) 332–
  342 (IEEE, 2020).
- 87. Beccherle, M., Facchetti, S., Villani, F., Zanini, M. & Scandola, M. Peripersonal Space from
  a multisensory perspective: the distinct effect of the visual and tactile components of
  Visuo-Tactile stimuli. *Exp Brain Res* 240, 1205–1217 (2022).
- 88. Bassolino, M., Finisguerra, A., Canzoneri, E., Serino, A. & Pozzo, T. Dissociating effect of
  upper limb non-use and overuse on space and body representations. *Neuropsychologia* **70**, 385–392 (2015).
- 687 89. Grivaz, P., Blanke, O. & Serino, A. Common and distinct brain regions processing
  688 multisensory bodily signals for peripersonal space and body ownership. *NeuroImage* 147,
  689 602–618 (2017).
- 690 90. de Gelder, B. & Vroomen, J. The perception of emotions by ear and by eye. *Cognition &*691 *Emotion* 14, 289–311 (2000).
- 692 6. Acknowledgement

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### 696 **7. Author Contributions**

697 All authors contributed to the design of the study and the writing of the manuscript.698 L.G., in addition, collected and analysed the data.

### 699 8. Data Availability Statement

All data analysed in this study have been made publicly available on Open Science

701 Framework (OSF) via the following link:

702 https://osf.io/xp9r8/?view\_only=ed8daecc5dfa43b8b1a024abdb37bb2f.

## 703 9. 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.

### 706 10. Figure Captions

Figure 1. The virtual environment and stimuli used in the four tasks: a neutral human adult male, an
anthropomorphic robot and a cylindrical lamp appearing at 300 cm in front of the participant in an
undecorated and unequipped room.

**Figure 2.** (A) The extent of the different spaces (MIS, IPS, RS, PS) expressed in centimetres as a function

of the stimulus (human, robot, lamp). The bars represent the average extent (error bars represent the

- SE), while the dots represent the individual performances. (B) Schematic representation of theorganisation of the different spaces.
- **Figure 3.** Correlation matrix plot showing the relation between RS, IPS, PS and MIS when facing the virtual lamp. The *r* refers to the Spearman coefficient when the correlation includes MIS and to the Pearson coefficient when it does not. \*\*p-values < .001, \*p-values < .05.
- 717 Figure 4. Correlation matrix plot showing the relation between RS, IPS, PS and MIS when facing the
- 718 virtual robot. The *r* refers to the Spearman coefficient when the correlation includes MIS and to the
- 719 Pearson coefficient when it does not. \*\**p*-values < .001, \* *p*-values < .05.
- 720 Figure 5. Correlation matrix plot showing the relation between the RS, IPS, PS and MIS when facing the
- virtual human. The *r* refers to the Spearman coefficient when the correlation includes MIS and to the
- 722 Pearson coefficient when it does not. \*\**p*-values < .001, \* *p*-values < .05.