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The relationship between action, social 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.

37 **Keywords:** PPS - IPS - social interactions - multisensory integration - reachability judgment -
38 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
46 has led to the distinction between *peripersonal space* (PPS, *i.e.*, within reach) and *extrapersonal*
47 *space* (*i.e.*, beyond reach)^{1,2}, which would be underpinned by different neural networks^{3,4}. The
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^{2,5,6}. 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
54 avoidance when facing threatening objects^{7,8}. In line with this, neuroimaging studies revealed
55 that the mere observation of an object located in PPS triggered activation in the sensorimotor
56 brain areas, including the reach-related area of the superior parieto-occipital cortex, and the
57 premotor and motor cortical areas^{9,10,11,12}. As a consequence, the transient disruption of the
58 left motor cortex using transcranial magnetic stimulation has been shown to produce an
59 alteration of the perception of objects located in PPS¹³. Likewise, corticospinal activity¹⁴ and
60 μ rhythm desynchronization^{15,16} increased in the presence of objects in the near (vs. far) space,
61 similar to what has been observed during the preparation and execution of objects-directed
62 motor actions^{17,18}. Moreover, modifying the actual reaching-by-hand capabilities (*e.g.*, through
63 tool-use or limb immobilisation), or biasing the spatial consequences of object-directed
64 actions, entailed a congruent increase or decrease of the PPS^{19,20,21,22}. Altogether, these

65 results suggest that PPS is an action space, enabling access to motor-related information
66 similar to those implied in the planning and execution of voluntary motor actions²³.

67 As revealed by monkey electrophysiological studies, most PPS neurons are
68 multisensory in that they respond to stimuli in two or three different sensory modalities, with
69 overlapping receptive fields anchored onto the same body region^{2,5,6}. 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
72 perceived as potentially interacting with our body⁸. Importantly, some of these neurons are
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 fields². 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 segments²³, as reflected by higher-order activations of somatosensory and associative
78 areas²⁴. Such multisensory integration has also been observed in humans, activating a
79 frontoparietal network^{25,26,27}, in relation to PPS²⁸. However, the main line of evidence in
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
83 which such facilitation is observed (as compared to a unisensory control condition) is usually
84 used as a proxy of the PPS extent^{29,30,31,32}. The scientific consensus is that the integration of
85 visual/auditory and tactile information would provide an interface between perception and
86 action allowing appropriate (re)actions towards (either threatening or incentive) objects to be
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
90 accuracy of motor responses^{33,34,35,36}. Furthermore, electric stimulation of the PPS
91 multisensory neurons in the monkey premotor and intraparietal cortex elicits a pattern of

92 movements that is compatible with defensive arm movements³⁷, while the PPS multisensory
93 neurons in the parietal and precuneus cortex have been shown to discharge during arm
94 reaching movements towards the part of space corresponding to their visual receptive field².
95 PPS represents thus a multisensory and sensorimotor interface mediating the physical
96 interactions between the body and the environment³⁸.

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
101 notably on the body region targeted by the tactile stimulation³². Indeed, when considering
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
108 the head; from 25 to 80 for the trunk^{29,30,31,32,39,40,41,42,43,44,45,46,47,48,49,50,51,52}). Hence, multisensory
109 integration does not seem to systematically overlap with the motor action space. In support
110 of this claim, Zanini and colleagues⁵³ found that the space corresponding to hand-centred
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
113 multisensory and reachable spaces are distinct spatial representations. However, it is worth
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
116 stable trunk-centred frame of reference^{54,55}. By contrast, multisensory integration was
117 thoroughly tested using a hand-centred or head-centred frame of reference, requiring, for
118 motor action, to refer to a more global representation of the body constituting the

119 egocentre^{32,38,56,57}. Accordingly, the “trunk-centred” reachable-by-hand space was not
120 expected to coincide exactly with the “hand-centred” multisensory integration space. In line
121 with this claim, Serino and colleagues³² considered that “hand- and face-centred PPS are
122 referenced to the trunk-centred PPS, which [is] a more extended representation of the space
123 surrounding the body”. Hence, multisensory integration might be compatible with the
124 representation of the space that is reachable with the hand when referring to the same frame
125 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 uncomfortable⁵⁸. 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^{59,60,61,62}. 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 interactions⁶³.
136 It is typically assessed with comfort distance judgments requiring the participant to place a
137 confederate at the most comfortable distance to interact with^{64,65}. Not only do these social
138 spaces refer to the space surrounding the body as PPS, but also share common
139 characteristics with PPS. For instance, PPS is modulated by social factors such as the
140 proximity of confederates and the relation that is held with them^{46,66, 67,68}. Furthermore, both
141 PPS and social spaces shrink or enlarge depending on the emotional valence of the facing
142 stimulus^{59,66,69}. They are also both influenced by individual characteristics such as anxiety⁶¹.
143 These observations probably explain why several researchers in the last decades have taken
144 a closer look at the relationship between PPS and social spaces. Until now, studies have
145 mainly focused on the link between PPS and PS. For instance, Iachini and colleagues^{70,60}

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
151 distances^{23,71,72}. In support of this claim, Quesque and colleagues⁶² found that extending arm
152 length's representation through tool-us increased PPS with a concomitant effect on PS. Social
153 spaces seem thus rooted in the same sensorimotor representation as PPS²³. However, the
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
157 of the extent of the different social spaces and their relationship to PPS. Finally, the
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
166 trunk at various times of the approach of the visual stimulus, while ignoring the latter (indexing
167 multisensory integration space, MIS). We expected RS to overlap and correlate with MIS, as
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^{23,71,72}, 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^{60,62,70}. Finally, all spaces should be similarly impacted by the nature of the stimulus,

172 with a preference for the lamp and robot to be kept at a larger distance compared to the
173 human⁷⁰.

174 2. Materials and Methods

175 2.1 Participants

176 Fifty-three participants from the Université of Lille participated in this study, but one
177 participant was excluded because they missed 20% of the tactile stimulations in the
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 \pm standard
181 deviation [*SD*] = 22.6 \pm 4.0). A sample size analysis performed in G*Power indicated that at
182 least 41 participants were required to detect a small effect (Cohen's *f* = 0.15) with a high power
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
187 Lille (CESC Lille, Ref. 2021-515-S95).

188 2.2 Apparatus & stimuli

189 The virtual stimuli were presented through an HTC Vive Pro head-mounted display in a
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
194 the human and robot were aligned with the eye level of the participant. All stimuli had the
195 same height and width. We verified that the visual stimuli were perceived as neutral by
196 requiring the participants to rate the emotional valence of each stimulus on the Self-

197 Assessment Manikin (SAM) scale, a 9-points graphic Likert scale ranging from 1 (extremely
198 negative) to 9 (extremely positive)⁷³. One sample *t*-test to 5 (*i.e.*, neutral emotional valence)
199 indicated that the human, $t(49) = 0.47, p = .643$, robot, $t(49) = 1.85, p = .071$, and lamp, $t(49) =$
200 $-1.24, p = .220$, were similarly judged as neutral.

201 **Figure 1.**



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
210 able to reach the approaching visual stimulus, without actually performing any reaching
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*

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

222 2.3.3 *Discomfort Distance Judgment*

223 The same procedure as in the reachability distance judgment and comfort distance
224 judgment tasks was used, except that participants were required to press the response button
225 whenever the visual stimulus was at a distance that made them feel uncomfortable. This task
226 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
241 visual stimulus while controlling that these effects were not merely due to the expectancy of
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
249 a random delay between 800 and 850 ms following the disappearance of the previous
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
253 in a random order. The trials were divided into 6 blocks of about 6 minutes intermingled with
254 5-minutes breaks. This task was used to assess MIS.

255 2.4 Data analyses

256 The data were analysed using *R* (version 4.1.0) and *R Studio* software (version 1.3.1093).
257 We first verified that our multisensory integration task succeeded in showing the typical
258 effects of the tactile stimulation delay on reaction times (RT) in each of the three stimuli used
259 (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
263 reachability judgement task, and in the discomfort and comfort distance judgement tasks,
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 visual 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 colleagues⁷⁴. 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
276 a virtual robot and a lamp^{70,60} when using stimuli controlled for their (neutral) emotional
277 valence. The paired comparisons were performed using the ART⁷⁴ or ART-C⁷⁵ alignment
278 procedure, as appropriate to the requested contrast, and with Bonferroni correction.

279 *2.4.2 Relationship between the different spaces.*

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

284 2.4.3 Bayesian analyses

285 We also conducted the corresponding Bayesian analyses in JASP (with default values)
286 in order to quantify the evidence in favour of an effect (H1) compared to an absence of effect.
287 These analyses provided Bayes Factors (BF_{10}) varying between 0 and ∞ , where values below
288 1 provide increasing evidence in favour of the null hypothesis and values above 1 provide
289 increasing evidence for the alternative hypothesis (H1/H0)⁷⁶. A BF above 3 is typically
290 considered sufficient evidence for the alternative hypothesis, while a BF below $\frac{1}{3}$ is typically
291 considered sufficient evidence for the null hypothesis⁷⁷.

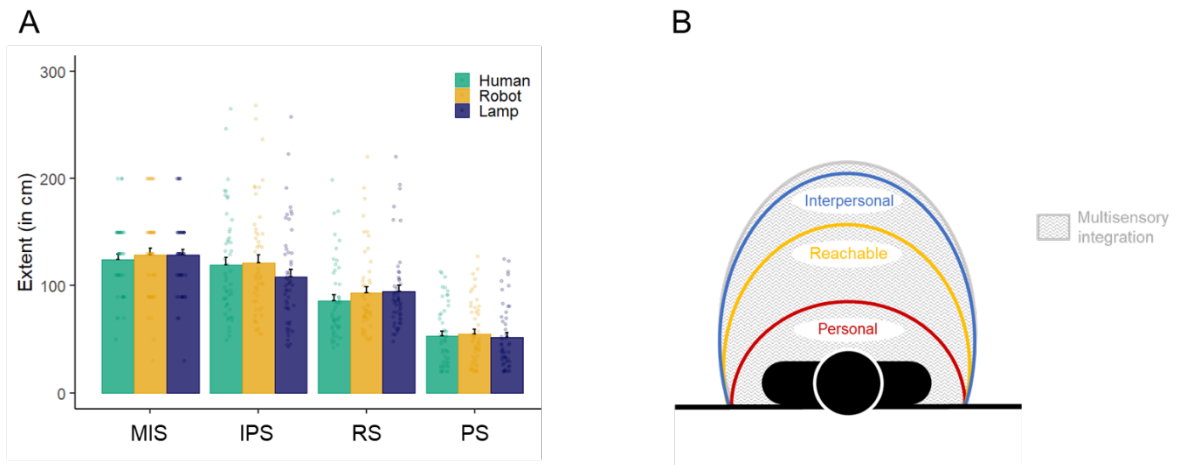
292 3. Results

293 3.1 Extent of the different spaces

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

309 (121.34 ± 7.34 cm), $t(98) = -3.42, p = .003, BF_{10} = 14.78$, and the human (119.27 ± 6.98 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, $F(2, 98) = 0.28, p = .752, \eta_p^2 = .005, BF_{10} = 0.089$.

316 **Figure 2.**



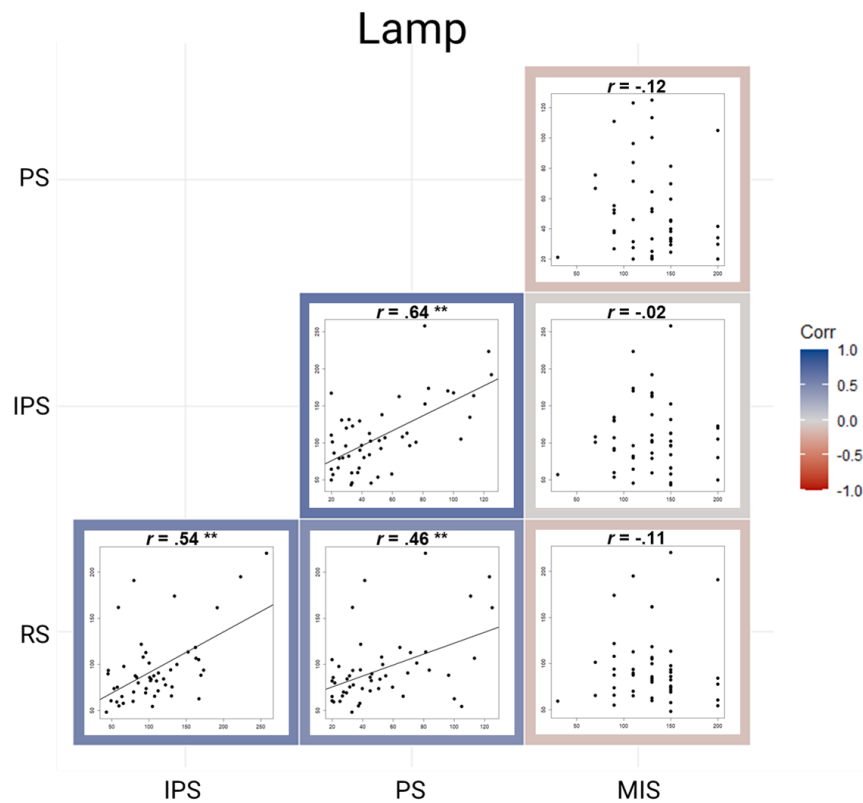
317

318 3.2 Relationship between the different spaces

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

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).

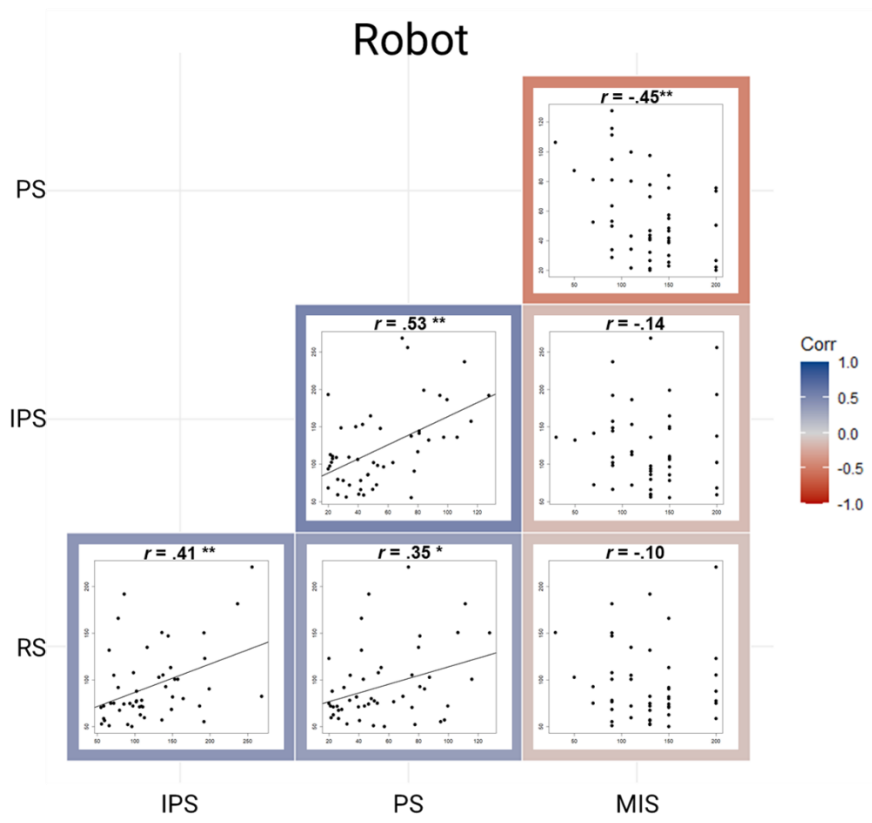
334 **Figure 3.**



335

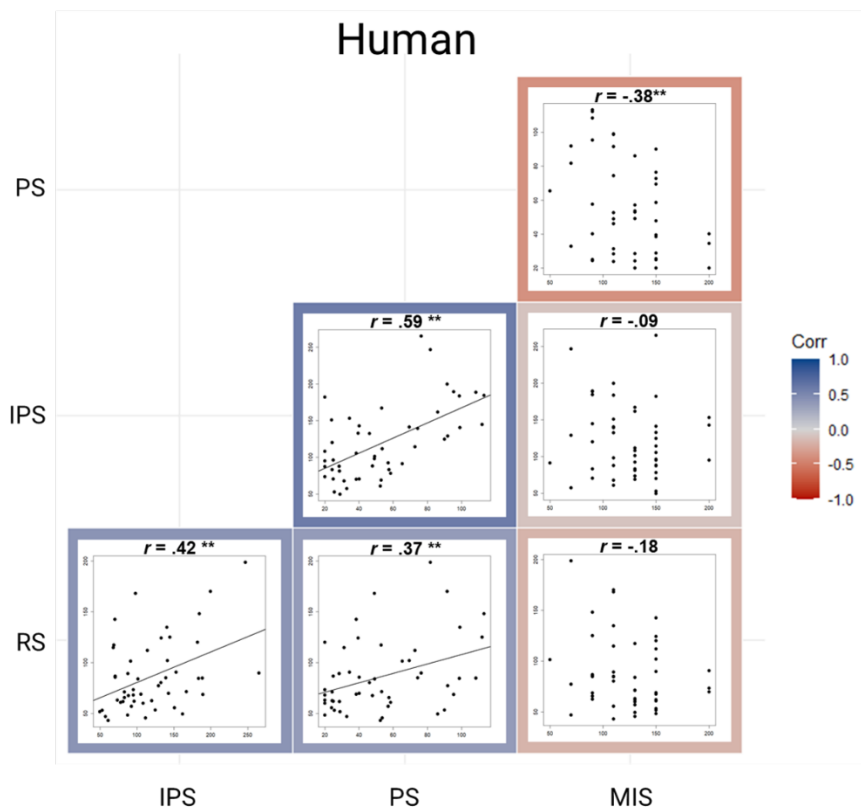
336

Figure 4.



337

Figure 5.



339

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
 343 latter related to the social spaces (interpersonal and personal). To do so, we required
 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
 347 integration task (MIS) that required participants to respond as fast as possible to tactile
 348 stimulation delivered on the trunk at various times of the approach of the visual stimulus.
 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

352 spaces, we expected all spaces to correlate and to be similarly impacted by the nature of the
353 stimulus, 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
358 known to slightly overestimate arm length⁷⁸ (here 73.2 ± 5.6 cm, corresponding for RS to an
359 overestimation of 29%), in particular in virtual environments⁷⁹. Moreover, the relative extents
360 of the reachable and social (PS and IPS) spaces are in line with previous observations showing
361 that IPS extent is typically between 80 and 140 cm⁶⁵, while RS and PS extents are typically
362 smaller, i.e., between the range of 50-70 cm^{20,70,62}. The data analysis conducted in the present
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^{80,81}. 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
367 shown^{60,70}, RS was significantly shorter in the presence of a virtual human than in the presence
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
372 touching them and thus be at shorter distances than people for which interactions might be
373 primarily conceived as a verbal exchange, especially when the situation involves a stranger⁶³.
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)
377 emotional valence should be positioned at the same IPS^{59,71,80}. A complementary
378 interpretation could be that the human stimulus used in the present study was a male who

379 was shown to trigger larger IPS than a female human stimulus^{60,70}. Further experiments would
380 be required to disentangle these different interpretations.

381 The correlational analyses revealed that RS, IPS and PS, although they were
382 characterised by different extents, were positively correlated to each other, whatever the
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
386 on the representation of PPS⁶², although the outcome of the present study extends the
387 contribution of PPS also to IPS. The observed pattern of results, therefore, provides an
388 additional argument for the involvement of PPS in the calibration of social spaces^{60,62,70,71}, and
389 corroborates brain imaging studies showing that the frontoparietal network involved in the
390 representation of PPS also supports social interactions^{82,83}. Overall, these findings comfort
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 interactions⁷¹. 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
398 buffer zone whose intrusion produces discomfort^{80,81} and triggers defensive behaviour⁸⁴. As
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,
406 which is in contradiction with our initial hypothesis. Indeed, MIS was 11.05 cm larger than IPS,
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)³². One potential
410 explanation could be that multisensory integration extended more when facing meaningful
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
413 virtual human characters (up to 127 and 150 cm, respectively)^{85,86} than when facing looming
414 pink noise (up to 66 and 75 cm, respectively)^{42,49}. However, even when centred on the same
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 RS². However, one may hypothesise that the sensory
423 facilitation reported in the behavioural studies and the neural mechanisms highlighted in the
424 single-cell studies do not refer to the exact same multisensory integration process⁸⁷. While
425 the link between the two has been strongly advocated²⁹, it is apparent that the behavioural
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
428 be altered by the valence or meaning of the visual/auditory stimulus⁶⁶, individual traits such
429 as anxiety/phobia⁴⁵, interoceptive traits⁴⁰, bodily changes such as pregnancy⁴¹ or limb
430 immobilisation⁸⁸, and even lockdown experience⁸⁵. Moreover, a number of studies indicated

431 that the visual/auditory stimulus does not have to target the same body part as the tactile
432 stimulation to trigger multisensory facilitation⁸⁷. 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 representation⁸⁹ and object-
435 directed action control⁷. 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
437 condition. Thus, despite their pioneering role, single-cell studies might represent only a small
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
441 revealed a link between multisensory integration and arm RS^{2,5}. From a behavioural
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 purpose⁸.
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.

451 Another implication of MIS encompassing all other spaces is that multisensory
452 processes, usually related to the action space, extend also to the social space. This is not that
453 surprising since the need to combine several sensory cues is not restricted to interactions
454 with objects but also applies to social stimuli. For instance, emotions are expressed through
455 facial expressions but also voice such that visual and auditory cues integration is an essential
456 part of emotion reading and more globally of social interactions⁹⁰. Moreover, multisensory
457 integration is assumed to allow the preparation of the body for action, either for the purpose

458 of defensive or approaching behaviour^{7,8,70,71}. 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
464 others and programme appropriate actions and responses – for example, to avoid harmful
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
468 interaction envisaged, and which thus seems not constitutive of the spaces underlying object-
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
472 confirms previous studies reporting that RS and PS are related^{60,62,70,80}, but extends this
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 interactions⁷¹. 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).
- 485 2. Rizzolatti, G., Scandolara, C., Matelli, M. & Gentilucci, M. Afferent properties of periarculate
486 neurons in macaque monkeys. II. Visual responses. *Behavioural Brain Research* **2**, 147–
487 163 (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).
- 505 10. Ferri, F. *et al.* Intertrial Variability in the Premotor Cortex Accounts for Individual
506 Differences 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 Suberved
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).
- 550 31. Noel, J.-P., Pfeiffer, C., Blanke, O. & Serino, A. Peripersonal space as the space of the bodily
551 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).
- 554 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*
556 *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).

- 561 36. Rizzolatti, G., Matelli, M. & Pavesi, G. Deficits in attention and movement following removal
562 of postarcuate (area 6) and prearcuate (area 8) cortex in macaque monkeys. *Brain* **106**,
563 655–673 (1983).
- 564 37. Cooke, D. F. & Graziano, M. S. A. Defensive Movements Evoked by Air Puff in Monkeys.
565 *Journal of Neurophysiology* **90**, 3317–3329 (2003).
- 566 38. Serino, A. Peripersonal space (PPS) as a multisensory interface between the individual
567 and the environment, defining the space of the self. *Neuroscience & Biobehavioral Reviews*
568 **99**, 138–159 (2019).
- 569 39. Amemiya, T., Ikei, Y. & Kitazaki, M. Remapping Peripersonal Space by Using Foot-Sole
570 Vibrations Without Any Body Movement. *Psychol Sci* **30**, 1522–1532 (2019).
- 571 40. Ardizzi, M. & Ferri, F. Interoceptive influences on peripersonal space boundary. *Cognition*
572 **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).
- 579 44. Galli, G., Noel, J. P., Canzoneri, E., Blanke, O. & Serino, A. The wheelchair as a full-body tool
580 extending the peripersonal space. *Front. Psychol.* **6**, (2015).
- 581 45. Taffou, M. & Viaud-Delmon, I. Cynophobic Fear Adaptively Extends Peri-Personal Space.
582 *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).
- 585 47. Hobeika, L., Taffou, M. & Viaud-Delmon, I. Social coding of the multisensory space around
586 us. *R. Soc. open sci.* **6**, 181878 (2019).

- 587 48. Hobeika, L., Taffou, M., Carpentier, T., Warusfel, O. & Viaud-Delmon, I. Capturing the
588 dynamics of peripersonal space by integrating expectancy effects and sound propagation
589 properties. *Journal of Neuroscience Methods* **332**, 108534 (2020).
- 590 49. Noel, J.-P., Blanke, O., Magosso, E. & Serino, A. Neural adaptation accounts for the
591 dynamic resizing of peripersonal space: evidence from a psychophysical-computational
592 approach. *Journal of Neurophysiology* **119**, 2307–2333 (2018).
- 593 50. Maister, L., Cardini, F., Zamariola, G., Serino, A. & Tsakiris, M. Your place or mine: Shared
594 sensory experiences elicit a remapping of peripersonal space. *Neuropsychologia* **70**, 455–
595 461 (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).
- 598 52. Pfeiffer, C., Noel, J., Serino, A. & Blanke, O. Vestibular modulation of peripersonal space
599 boundaries. *Eur J Neurosci* **47**, 800–811 (2018).
- 600 53. Zanini, A. et al. Peripersonal and reaching space differ: Evidence from their spatial extent
601 and multisensory facilitation pattern. *Psychon Bull Rev* **28**, 1894–1905 (2021).
- 602 54. Lacquaniti & Caminiti. Visuo-motor transformations for arm reaching. *European Journal of*
603 *Neuroscience* **10**, 195–203 (1998).
- 604 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).
- 607 56. Alsmith, A. J. T. & Longo, M. R. Where exactly am I? Self-location judgements distribute
608 between head and torso. *Consciousness and Cognition* **24**, 70–74 (2014).
- 609 57. Bertossa, F., Besa, M., Ferrari, R. & Ferri, F. Point Zero: A Phenomenological Inquiry into
610 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).
- 621 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).
- 627 65. Sorokowska, A. *et al.* Preferred Interpersonal Distances: A Global Comparison. *Journal of*
628 *Cross-Cultural Psychology* **48**, 577–592 (2017).
- 629 66. Pellencin, E., Paladino, M. P., Herbelin, B. & Serino, A. Social perception of others shapes
630 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
632 of peripersonal space depends on the success of one's own motor actions, but also the
633 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. Iachini, 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).
- 641 71. Coello, Y. & Cartaud, A. The Interrelation Between Peripersonal Action Space and
642 Interpersonal Social Space: Psychophysiological Evidence and Clinical Implications.
643 *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
647 semantic differential. *Journal of Behavior Therapy and Experimental Psychiatry* **25**, 49–59
648 (1994).
- 649 74. Wobbrock, J. O., Findlater, L., Gergle, D. & Higgins, J. J. The aligned rank transform for
650 nonparametric factorial analyses using only anova procedures. in *Proceedings of the*
651 *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*
654 *Software and Technology* 754–768 (ACM, 2021).
- 655 76. Dienes, Z. Using Bayes to get the most out of non-significant results. *Front. Psychol.* **5**,
656 (2014).
- 657 77. Jeffreys, H. *The theory of probability.* (1988).
- 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
661 Using Virtual Reality Technology and Manual Measurement. in *2020 IEEE 7th International*
662 *Conference on Industrial Engineering and Applications (ICIEA)* 390–393 (IEEE, 2020).
- 663 80. Cartaud, A., Ruggiero, G., Ott, L., Iachini, T. & Coello, Y. Physiological Response to Facial
664 Expressions in Peripersonal Space Determines Interpersonal Distance in a Social
665 Interaction Context. *Front. Psychol.* **9**, 657 (2018).

- 666 81. Kennedy, D. P., Gläscher, J., Tyszka, J. M. & Adolphs, R. Personal space regulation by the
667 human amygdala. *Nat Neurosci* **12**, 1226–1227 (2009).
- 668 82. Holt, D. J. *et al.* Neural Correlates of Personal Space Intrusion. *Journal of Neuroscience* **34**,
669 4123–4134 (2014).
- 670 83. Vieira, J. B., Pierzchajlo, S. R. & Mitchell, D. G. V. Neural correlates of social and non-social
671 personal space intrusions: Role of defensive and peripersonal space systems in
672 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).
- 675 85. Serino, S. *et al.* Sharpening of peripersonal space during the COVID-19 pandemic. *Current*
676 *Biology* **31**, R889–R890 (2021).
- 677 86. Buck, L. E., Park, S. & Bodenheimer, B. Determining Peripersonal Space Boundaries and
678 Their Plasticity in Relation to Object and Agent Characteristics in an Immersive Virtual
679 Environment. in *2020 IEEE Conference on Virtual Reality and 3D User Interfaces (VR)* 332–
680 342 (IEEE, 2020).
- 681 87. Beccherle, M., Facchetti, S., Villani, F., Zanini, M. & Scandola, M. Peripersonal Space from
682 a multisensory perspective: the distinct effect of the visual and tactile components of
683 Visuo-Tactile stimuli. *Exp Brain Res* **240**, 1205–1217 (2022).
- 684 88. Bassolino, M., Finisguerra, A., Canzoneri, E., Serino, A. & Pozzo, T. Dissociating effect of
685 upper limb non-use and overuse on space and body representations. *Neuropsychologia*
686 **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).

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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

700 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

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

706 10. Figure Captions

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

710 **Figure 2.** (A) The extent of the different spaces (MIS, IPS, RS, PS) expressed in centimetres as a function
711 of the stimulus (human, robot, lamp). The bars represent the average extent (error bars represent the

712 SE), while the dots represent the individual performances. (B) Schematic representation of the
713 organisation of the different spaces.

714 **Figure 3.** Correlation matrix plot showing the relation between RS, IPS, PS and MIS when facing the
715 virtual lamp. The r refers to the Spearman coefficient when the correlation includes MIS and to the
716 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
721 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.