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Sense of Agency for intracortical brain machine interfaces

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43 Abstract

Intracortical brain machine interfaces decode motor commands from neural signals and translate them into actions, enabling movement for paralyzed individuals. The subjective sense of agency associated to actions generated via intracortical brain machine interfaces, the involved neural mechanisms and its clinical relevance are currently unknown. By experimentally manipulating the coherence between decoded motor commands and sensory feedback in a tetraplegic individual using a brain machine interface, we provide evidence that primary motor cortex processes sensory feedback, sensorimotor conflicts and subjective states of actions generated via the brain machine interface. Neural signals processing the sense of agency affected the proficiency of the brain machine interface, underlining the clinical potential of the present approach. These findings show that primary motor cortex encodes information related to action and sensing, but also sensorimotor and subjective agency signals, which in turn are relevant for clinical applications of brain machine interfaces.

75 Main text

76

77 Introduction

When performing a voluntary movement, motor commands from the brain activate body 78 effectors, which produce a cascade of reafferent sensory (proprioceptive, tactile, visual) 79 cues. Motor commands are also associated with prediction signals about the sensory 80 consequences of the movement. The congruency between motor commands, reafferent 81 sensory feedback, and sensory predictions is at the basis of the sense of agency, our 82 feeling of being in control of our actions¹⁻³. In case of damage to the motor system, 83 motor commands that would trigger actions do not reach body effectors, leading to 84 different types of paralysis, depending on the location and severity of damage. 85 Intracortical brain machine interfaces (BMI) bypass such brain-body disconnection by 86 decoding brain signals from different regions (i.e., primary motor cortex (M1), parietal or 87 premotor cortex) and translating them into motor commands for the control of robots, 88 exoskeletons^{4,5}, neuromuscular functional electrical stimulation^{6,7} or other devices⁸, 89 enabling different actions (BMI actions) for patients with severe neuromotor 90 impairments⁹. 91

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Here we study how it feels to generate movements with an intracortical-BMI, that is what is the sense of agency for BMI actions ((see ^{10,11} for recent studies with noninvasive brain computer interfaces (BCI)) and search for a potential neural mechanism. In particular, we asked whether motor neurons in human M1 encode not only motor commands, but also sensory feedback and whether these signals covary with agency for BMI actions Finally we tested whether agency also affects the efficiency of the BMI system - i.e. whether agency has a potential therapeutic benefit.

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We applied approaches psychophysics, neurophysiology, 101 classic from neuroengineering and virtual reality (VR) to ask these questions in a patient suffering 102 from tetraplegia (caused by severe cervical spinal cord injury; C5/C6), who had been a 103 BMI expert for two years before the start of the present study⁶. The patient had no 104 preserved motor function below the C5 level. His sensory functions were extremely 105

limited and only showed partially preserved function at the C6 level on the left side and at C5 on the right side (there was also residual sensation for pressure on his right thumb). Concerning proprioception, he had preserved perception for shoulder, elbow and wrist joint position, but no proprioception for digits joint position (see Material and methods for more details).

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The BMI consisted of a 96-channel array implanted in the hand area of left M1 and 112 actuated a transcutaneous forearm neuromuscular electrical stimulation (NMES) system 113 (see ⁶ for a full description of the system) to translate decoded cortical signals into right 114 forearm and hand movements. In order to study the sense of agency for BMI actions 115 and to evaluate its clinical impact, we experimentally manipulated the congruency 116 between the decoded actions and the actions actuated by the BMI-NMES system. As 117 illustrated in Figure 1, the participant was instructed to realize a cued action with the 118 BMI and was provided with movement-related sensory feedback using visual (via VR) 119 and/or somatosensory (via NMES) stimulation. Critically, this feedback was either 120 congruent or incongruent with respect to the motor commands decoded from M1: half of 121 122 the trials, in which the decoded action corresponded to the cued action (e.g., open hand), were associated with congruent feedback (e.g., open hand), while the other half 123 were associated with incongruent feedback (e.g. the opposite action: close hand). For 124 each BMI action, we asked the participant whether he felt in control of that action and to 125 rate his confidence about this judgement, allowing us to (1) gauge the sense of agency 126 127 for BMI actions and how this was modulated by the congruency between motor commands and sensory feedback. Next, neural data from the M1 implant were analyzed 128 to measure how (2) the sense of agency and (3) sensory feedback were encoded in the 129 activity of M1 neurons, quantified as multi-unit (MU) firing rates and local field potentials 130 (LFP). Finally, we investigated (4) how visual and somatosensory feedback, and the 131 associated sense of agency, affected the performance of the BMI system by changing 132 the pattern of response of M1 neurons. By investigating what it feels like to control 133 actions mediated by an intracortical BMI, our data show neural patterns in M1 activity 134 (MU and LFP) reflecting the processing of agency for BMI actions, as generated by the 135 congruency between intention and sensory feedback. Importantly, we show that the 136 nature of somatosensory feedback (and the related sense of agency) affected the 137

efficiency of the BMI system by modulating the response properties of M1 neurons,
underlining the clinical relevance of sensory feedback and agency for the BMI field.

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During the experiment, the participant was cued to execute one of four target actions 141 (hand opening, hand closing, thumb extension, thumb flexion) using a validated BMI 142 143 neuroprosthesis. Neural activity corresponding to each target movement was recorded via a 96-channel microelectrode array in M1 and a nonlinear support vector machine 144 classifier was applied to decode the participant's chosen action from MU activity (see ⁶ 145 for full description). On each trial, the classifier provided the likelihood of each target 146 action (on a -1 to +1 range, in 100 ms bins), thus decoding one of the four target actions 147 from the participant's M1 activity. In three different experiments, visual, somatosensory, 148 or visual-somatosensory feedback about the BMI action was provided (Figure 1). In 149 Experiment 1, VR was used to provide visual feedback, consisting of a life-size virtual 150 arm on a monitor superimposed over the participant's right arm, matching the location 151 and dimensions of the participant's real arm, which was occluded from view. In 152 Experiment 2, NMES was used to provide 'somatosensory' feedback: the patient's 153 upper limb muscles were electrically stimulated so he could feel, but not see the 154 selected movement. Experiment 3 combined VR and NMES to provide 'visual-155 somatosensory feedback' (see below). In half of the trials, sensory feedback was 156 congruent with the cued action, while in the other half it was incongruent (i.e., the 157 opposite, action was executed) (see Figure 1B). At the end of each trial, we gauged the 158 159 participant's sense of agency (0 or 1; Q1) and confidence (rating between 0 and 100; Q2). Importantly, the amount of sensory information was kept constant across 160 experiments, by providing non-informative sensorimotor feedback in Experiment 1 (i.e., 161 a pattern of NMES triggering no BMI action) and non-informative visual feedback in 162 Experiment 2 (i.e., a static visual hand performing no action). 163

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166 **Results**

Sensory feedback determines agency and confidence. Agency ratings were collected in a total of 844 trials (155, 243 and 448 trials for Experiments 1, 2 and 3, respectively; for Experiment 3 see below and Supplementary Information) and

compared across feedback conditions using permutation tests. A null distribution of the 170 mean agency rating was created by shuffling the condition labels over 10'000 iterations. 171 P-values (2-sided) were estimated by counting the proportion of shuffled samples 172 exceeding the observed average difference across conditions. As expected, and as 173 shown in Figure 2A, we were able to manipulate agency and confidence for BMI 174 actions. Thus, congruent visual (Experiment 1, 93.8% (bootstrapped 95% confidence 175 interval 93.4% - 94.2%) and 5.2% (4.8% - 5.6%) of positive responses to Q1 for 176 congruent and incongruent trials, respectively, p<.0001) and congruent somatosensory 177 (Experiment 2, 97.5% (97.3% - 97.6%) and 8.8% (8.4% - 9.1%) of positive responses 178 for congruent and incongruent trials respectively, p<.0001) feedback resulted in more 179 180 frequent agency responses versus incongruent conditions. Analyzing the role of feedback for confidence ratings (irrespective of the agency ratings), we found that 181 confidence was modulated by somatosensory congruency (Experiment 2, Q2 ratings 182 were higher for somatosensory congruent [M = 74.0 (73.9 - 74.2)] than incongruent [M = 183 65.0 (64.8 - 65.2)] feedback; p < 0.001). The effect of visual congruency on confidence 184 ratings was not significant (Experiment 1, mean Q2 rating = 70.9 (70.6 - 71.1) for 185 186 congruent, 73.4 (73.1 - 73.6) for incongruent trials; p = 0.28).

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In order to disentangle the role of visual and somatosensory cues for agency and 188 confidence, Experiment 3 combined VR and NMES including combinations of congruent 189 and incongruent visual and somatosensory feedback (Figure 1). Most relevant are the 190 191 comparisons between feedback conditions in which visual (V) and somatosensory (S) signals were both congruent (+) or both incongruent (-) (V+/S+; V-/S-) or when feedback 192 was congruent in one modality and incongruent in the other modality (V+/S-; V- /S+). 193 Results revealed that somatosensory congruency was more effective in driving the 194 sense of agency and the associated confidence: ratings were higher not only when both 195 feedback signals were congruent (Q1 = 100% "Yes", mean Q2 = 83.98 [83.7 - 84.0]) as 196 compared to both being incongruent (Q1 = 7.6% "Yes" [7.2% - 7.9%], mean Q2 = 72.4197 [72.2 - 72.6)], both p-values < 0.001), but also in the V- /S+ (Q1 = 68.9% "Yes" [68.4% -198 69.4%], mean Q2 = 59.4 [59.3 - 59.6] as compared to the V+/S- condition (Q1 = 52.2%) 199 "Yes" [51.5% - 52.8%], mean Q2 = 54.6 [54.4 - 54.8]; p = 0.0035 and p = 0.036, for 200 agency and confidence respectively) (Figure 2). Collectively, these data from 201

Experiments 1-3 show that the congruency between decoded actions and sensory feedback, especially for the somatosensory modality, alters the sense of agency and confidence for actions mediated by an intracortical BMI.

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The sense of agency has been traditionally studied by presenting participants with 206 different visuo-motor couplings^{2,12–15}. In comparison, the role of somatosensory signals 207 remains poorly understood¹⁶, notably because it is normally impossible to decouple 208 motor commands, somatosensory feedback and visual feedback, with extremely rare 209 exceptions as in deafferented patients. Here we were able to contrast feedback cues 210 that were congruent in one modality (e.g., visual) and incongruent in the other modality 211 (e.g., somatosensory; and vice versa) with respect to the motor command and 212 demonstrate that somatosensory cues dominate the sense of agency and the 213 associated confidence for BMI-NMES actions. Of note, this effect cannot be due to the 214 presence of somatosensory cues alone, as BMI actions in the visual condition were 215 always associated with non-informative NMES stimulation producing somatosensory 216 sensations without generating any actions (i.e., pseudo random somatosensory 217 feedback, see Supplementary Information). Collectively these psychophysical data in a 218 BMI expert reveal that agency for BMI actions depends on visual and somatosensory 219 feedback (tactile and proprioceptive input) with somatosensory cues being more 220 relevant. 221

222

Cortical signatures of sensory feedback in M1. We next investigated how such 223 sensory feedback, that modulated the sense of agency, was encoded in M1 activity. We 224 first analyzed the LFP amplitude in the different feedback conditions across the three 225 experiments, using a regularized generalized linear model (ridge regression) and input 226 signals from each individual channel at every time point (see Supplementary 227 Information). As shown in Figure 3A (left), the analysis distinguished congruent vs. 228 incongruent visual feedback (maximum Cohen's Kappa K = 0.40, t-sum = 219.4, p < 229 0.001) within a single period of a positive potential that lasted from ~700-1200 ms after 230 the BMI action classification onset (Experiment 1). We could also distinguish congruent 231 vs. incongruent somatosensory feedback (maximum Cohen's Kappa K = 0.64, t-sum = 232 233 979.8, p < 0.001) during two time periods: an early period characterized by a negative

potential (stronger for incongruent feedback), starting at ~200 ms after BMI 234 classification onset, followed by a later persistent differentiation lasting almost until the 235 end of the trial. These results were further corroborated by data from Experiment 3: 236 congruent trials in both modalities were clearly distinguished from incongruent trials in 237 both modalities, lasting from ~250-1900 ms after BMI classification onset (maximum K = 238 0.66, t-sum = 958.8, p < 0.001). In addition, V+/S- trials were different from V-/S+ trials 239 from ~300-1400 ms from BMI classification onset (maximum K = 0.31, t-sum = 256.7, p 240 < 0.001) (Figure 3B left). These findings show that visual and somatosensory feedback 241 were both encoded by LFPs in human M1 and that such M1-LFP coding started earlier 242 and was more stable over time for somatosensory feedback. 243

244

Applying the same decoding algorithm as for LFPs, we next determined if sensory 245 feedback was also encoded by the spiking rate of MU in M1 (for methods see 246 Supplementary Information). As shown in Figure 3A (right), in Experiment 1, MU activity 247 distinguished between congruent and incongruent visual feedback from ~400-900 ms 248 from BMI classification onset (max K value = 0.41, t-sum = 56.2, p < .001). Extending 249 LFP findings, an earlier and more stable differentiation between congruent and 250 incongruent somatosensory feedback was found in MU activity in Experiment 2, with an 251 effect as early as ~200 ms from the BMI classification onset (max K value = 0.66, t-sum 252 = 390.7, p = < .001) and then persisted from 800 to 2000 ms. Similar results were found 253 in Experiment 3 (Figure 3B, right), where MU activity distinguished between trials 254 congruent and incongruent in both modalities (max K value = 0.68, t-sum = 271.8, p = < 255 .001) and between V+/S- and V-/S+ trials from ~160 ms from BMI classification onset 256 (max K value = 0.55, t-sum = 151.2, p = < .001). These data show that LFP and MU 257 activity reflects visual and somatosensory feedback during actions driven by a BMI 258 neuroprosthesis, with M1 activity early reflecting somatosensory feedback starting ~200 259 ms after NMES activation (~150 ms after BMI classification onset, ~200 ms before M1 260 activity encoding visual feedback) and persisting for a longer period. 261

262

The role of somatosensory and visual information is an important topic in motor control, with robust evidence showing how perturbations of sensory feedback impact motor

execution and adaptation¹⁷. The present data show that the congruency between an 265 intended action and somatosensory/visual feedback is encoded by M1 neurons at 266 different latencies. Previous studies in non-human primates described responses in M1 267 related to tactile and visual input^{18,19}, during active and passive movements²⁰ and during 268 visual feedback of a pre-recorded movement^{21,22}. The present results are consistent 269 with proposals that suggest that M1 activity codes both for movement types and their 270 sensory consequences, in line with recent works describing how M1 neurons encode 271 different movement parameters (see ^{19,23,24} for reviews). Here we report that, at the 272 population level, human M1 activity in addition discriminates between arm movements 273 that were congruent or incongruent with the motor command, as defined by 274 somatosensory and visual feedback, with higher accuracy, earlier and more consistent 275 processing for the former type of sensory information. Thus, neural coding in M1 276 contains, at the population level, information not only about the movement itself, but 277 also about sensory consequences of actions, involving somatosensory-motor and visuo-278 motor loops. These results are important to explain how sensory feedback affects the 279 proficiency of the BMI system as described below. 280

281

Cortical signatures of the sense of agency in M1. It is known that sensory-motor 282 congruency is a key mechanism of agency for able-bodied actions^{2,3}; here we have 283 shown that this also applies to agency and confidence for BMI-mediated actions and 284 that LFPs and MU activity in human M1 distinguishes congruent vs. incongruent BMI 285 actions. Next, we investigated to what extent LFP and MU activity in M1 also 286 discriminate actions with and without an accompanying sense of agency. For each trial, 287 we sorted LFP responses as a function of whether the participant reported agency or 288 not. As seen in Figure 4A (left), LFP activity starting ~270 ms after BMI classification 289 onset was found to code for agency and reached a maximum information value (Max K 290 = 0.41, t-sum = 1258.3, p < 0.001) at ~1000 ms after BMI movement onset. Thus, BMI 291 actions for which the participant felt to be the agent were characterized by a different 292 LPF pattern compared to BMI actions for which he did not. This was corroborated by 293 MU activity analysis (Figure 4A, right). The MU firing rate was higher for trials with 294 versus no agency; this discrimination started at ~300ms after BMI classification onset, 295

until 500 ms, and peaked at ~400 ms (max K = 0.45, t-sum = 232.5, p < 0.001). Later on, MU activity also differentiated for agency, with higher firing rate for trials with no agency (800-1600 ms after BMI classification). The same decoding was also able to discriminate trials with high vs. low confidence, based on a median split of Q2, from LFPs (max K = 0.296 at ~1200 ms, t-sum = 8449.3, p < 0.001) and MU (max K = 0.225 at ~400 ms, t-sum = 214.9, p < 0.001).

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In the experimental design, sensory feedback congruency was used to modulate the 303 sense of agency and this may have influenced these agency findings. Accordingly, we 304 next tested whether LFP and MU contained information related to the sense of agency 305 per se, after controlling for the effect of sensory feedback. For this we built a continuous 306 measure of sense of agency and confidence allowing us to regress out the effect of 307 sensory feedback. This new index was computed by recoding confidence ratings (Q2) 308 309 as -Q2, for trails with no agency (as indicated in Q1) and +Q2 for trials with agency (from Q1). This index was then orthogonalized with respect to congruency in order to 310 regress out this effect from the agency scores. As M1 signals also varied as a function 311 of the different cued actions (see SI), the index was also orthogonalized for the type of 312 action. We then used the same decoder to predict orthogonalized agency scores from 313 LFP and MU activity over time. This analysis shows that LFPs predicted the sense of 314 agency starting at ~450 ms after BMI classification onset (max $R^2 = 0.03$, t-sum = 69.8, 315 p = 0.017) (see Figure 4B left). A similar pattern was found when considering MU 316 activity, although the peak failed to reach significance after cluster-based correction for 317 multiple comparisons (Figure 4B right). These data show that M1 activity encodes the 318 sense of agency and associated confidence level and was modulated by the 319 congruency between motor commands and sensory feedback. Thus, subjective mental 320 states associated with BMI actions and control are encoded by M1 activity at the LFP 321 level (and to a minor extent at MU), independent of the neural processing associated 322 with sensory feedback (see Supplementary Information for single channel analyses). 323

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325 **Somatosensory feedback modulates BMI classifier accuracy.** Given the strong role 326 of sensory congruency in determining agency and its coding in M1, we finally asked

whether sensory feedback has any impact on the BMI classifier. To this aim, we tested 327 whether the congruency between the decoded motor commands and sensory feedback 328 (visual, somatosensory) affected the accuracy of the BMI classifier, defined as the 329 summed suprathreshold activation values across a 4s window. In Experiments 1 and 2 330 we found that congruent somatosensory feedback improved classifier accuracy (t(241) 331 = 9.29, Cohen's d = 1.238, p < 0.001) (Figure 5B right). There was no effect due to 332 visual feedback (t(153) = 1.523, Cohen's d = -0.245, p = 0.14) (Figure 5A). Moreover, 333 incongruent somatosensory feedback was associated with lower classifier accuracy for 334 the cued movement (Figure 5B left), and even increased classifier accuracy for the 335 opposite movement (Figure 5B left). Thus, only somatosensory feedback congruency 336 affected BMI accuracy in the present participant. This was extended by the results of 337 Experiment 3, where we found a significant main effect of sensory feedback condition 338 $(F(3,444) = 15.83, \eta^2 = 0.097, p < 0.001;$ Figure 5C). Further post-hoc corrected tests 339 showed that the BMI classifier's accuracy was higher when feedback was congruent, 340 than incongruent, in either modality (Tukey corrected t = 4.966, Cohen's d = 0.666, $p < 10^{-1}$ 341 0.001). More interestingly, when feedback was congruent for the somatosensory 342 modality and incongruent for the visual modality (V-/S+) BMI accuracy was higher than 343 in the opposite feedback condition (S-/V+) (t = 4.558, Cohen's d = 0.612, p < 0.001). 344 Figure 5 also shows the modulation of the BMI decoder as function of sensory feedback 345 over time during the trial. Significant change of the decoder's output is visible from 430 346 ms from somatosensory feedback. 347

348 These data from Experiments 1-3 show that BMI performance is affected by the congruency between the decoded motor commands and the somatosensory feedback 349 induced by the action actuated by NMES. This finding is also coherent with the more 350 reliable (i.e., earlier, more long-lasting and better decoded) processing of 351 somatosensory feedback from M1 activity (LFP, MU). The fact that the same action as 352 actuated by NMES (e.g., open hand) increased or decreased the BMI classifier 353 performance, depending on whether somatosensory feedback was congruent (open 354 hand) or incongruent (close hand) with the cued action, excludes that this effect was a 355 generic artifact of NMES stimulation affecting the input to the BMI classifier 356 independently from sensory information. 357

In order to better understand how somatosensory feedback affected the accuracy of the 358 BMI classifier, we analyzed time point by time point changes in multiunit activity for the 359 whole array. We computed the average Euclidean distance between firing patterns of 360 trials with a given cued movement and either congruent or incongruent somatosensory 361 feedback. For a given cued movement (e.g., movement hand open) at congruent 362 feedback (hand open), we computed its distance either with the same movement (cued: 363 hand open) at incongruent feedback (hand close) or with the opposite movement (hand 364 close), at its relative incongruent feedback (hand open). This way, we compared cases 365 with the same motor intention, but opposite sensory feedback, and trials with the 366 opposite motor intention, but the same sensory feedback. As shown in Figure 6, trials 367 from Experiment 2 with opposite somatosensory feedback, but the same motor 368 intention, diverge after sensory feedback, whereas trials with opposite motor intention, 369 but the same sensory feedback, seem to even Otheconverge slightly with respect to 370 371 baseline. This shows that M1 activity after feedback reflects the movement implemented via NMES more than the intended movement, thus explaining the modulation of 372 somatosensory feedback in BMI proficiency (see Figure 6B). As a control, we also 373 analyzed trials with opposite motor intention and congruent somatosensory feedback. 374 We found the activity patterns to differ only slightly with respect to trials with same 375 somatosensory feedback, but opposite motor intention, further showing that 376 somatosensory feedback is prevailing over motor intention after movement onset. In the 377 case of visual feedback (Experiment 1), instead, there was no divergence of activity 378 379 patterns after the feedback, while trails with different motor intention clearly diverged before the movement onset (see Figure 6A). 380

381

In order to better display the effect of somatosensory feedback on M1 activity for each 382 type of movements, we computed a 2D multidimensional scaling of neural activity as a 383 function of intended movement and congruency of somatosensory feedback. This 384 technique aims at representing the high dimensional spatio-temporal pattern of neural 385 activity in 2D plane, while maximising the fraction of retained variance. As shown in 386 Figure 6C, both for hand (opening/closing) and thumb (flection/extension) movements, 387 before sensory feedback (in the window between -650 and -150 ms before sensory 388 389 feedback onset), M1 neural activity is clustered solely as a function of the intended

390 movement. After somatosensory feedback (between 0 and 600 ms from sensory 391 feedback onset, Figure 6D), trials with congruent somatosensory feedback and a given 392 intended movement are clustered more with trials coding for the opposite movement, 393 but receiving the same sensory feedback rather than with trials coding for the same 394 movement.

No prior study in humans and only few studies in monkeys directly tested the effects of 395 sensory feedback on BMI performance^{21,25}. Here we show, for the first time, an effect of 396 feedback congruency on BMI performance, and the underlying role of M1 in this 397 process. Our findings indicate that the recorded M1 units processed motor signals for 398 the trained BMI actions, for sensory and sensory-motor signals reflecting the type and 399 congruency of the sensory feedback. Importantly, these processes were found to 400 change across time, as a function of the sensory feedback provided. In particular, our 401 results show that, after somatosensory feedback, the pattern of neural activity from M1 402 reflected more closely the type of movement realized by the NMES (i.e., the pattern of 403 somatosensory feedback) rather than the intended and decoded movement. This re-404 writing of the encoded M1 movement as a function of the NMES-implemented 405 406 movement directly relates to the improvement of BMI efficiency based on congruent somatosensory feedback that we observed and was absent in visual feedback trials. 407

This effect might be mediated by mutual connections between the primary motor and 408 the primary somatosensory cortices, which have been extensively documented in non-409 humans primates²⁶ and in humans²⁷. In addition, this effect might also depend on direct 410 411 somatosensory inputs reaching M1 neurons likely from the dorsal columns via the ventrolateral thalamic nucleus²⁸. This is an important finding, considering that original 412 BMI approaches for severely motor-impaired patients generally provide visual feedback 413 only^{5,29} or somatosensory feedback by directly stimulating primary somatosensory 414 cortex^{30–32} (see ³¹ for a review). Although from a single tetraplegic participant, the 415 present data show that non-invasive somatosensory feedback via NMES not only 416 enables higher subjective feeling of being in control (agency and confidence), but also 417 leads to better actual control of the patient's BMI actions. 418

419

Agency covaries with BMI classifier. We finally investigated whether agency has an 420 impact on BMI efficiency and thus tested whether the sense of agency covaried with 421 BMI classifier accuracy. We found that trials with agency versus trials without agency 422 were associated with higher classifier accuracy when somatosensory feedback was 423 modulated (Experiment 2; t(241) = 8.91, Cohen's d = 1.199, p < 0.001), as confirmed 424 from analysis of data from Experiment 3 (t(446) = 6.256, Cohen's d = 0.601, p < 0.001). 425 We found no statistically significant difference between trials with agency and trials 426 without agency in classifier accuracy when visual feedback was modulated (Experiment 427 1; t(153) = 0.690, Cohen's d = 0.111, p = 0.49). In addition, there was a significant 428 correlation across all three experiments between BMI classifier accuracy and 429 confidence (See Supplementary Table 1 for multiple regression analyses). Thus, 430 431 agency and confidence were both directly related to the performance of the present BMI system, but only when somatosensory feedback was involved. In order to confirm the 432 role of agency on BMI performance, while controlling for other potential factors, we 433 compared the BMI performance between trials in which the BMI user reported high and 434 low agency, within conditions at equivalent sensory feedback, that is V-/S+ and V+/S-435 from Experiment 3 (which resulted in a balanced and sufficient numbers of trials with 436 "Yes" and "No" responses to Q1). As shown in Figure 5D, BMI accuracy varied as a 437 function of subjective agency judgments, in conditions of equivalent sensory feedback. 438 BMI accuracy was significantly higher in trials with high agency as compared to trials 439 with low agency from 300 ms in the V-/S+ condition. The same pattern is visible in the 440 V-/S+ condition, although the comparison was not significant (i.e., did not survive to 441 correction for multiple comparisons). The same analysis run on confidence ratings (by 442 sorting high and low confidence ratings by means of a median split) did not show any 443 significant difference in BMI accuracy due to confidence at equivalent conditions of 444 sensory feedback (see Supplementary Figure S6). These results suggest that the sense 445 of agency, and not confidence (see Supplementary Table 1 for further analyses), has an 446 447 effect on BMI accuracy beyond the prominent role of sensory feedback, and impacts BMI accuracy at a later time point. Since agency judgments and confidence ratings 448 reflect two different processes of subjective experience, the present data suggest that 449

pre-reflexive, rather than post-decisional agency components more strongly affect the
 proficiency of a BMI decoder in M1.

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454 **Discussion**

By combining techniques from neurophysiology, neuroengineering, and VR with 455 psychophysics of agency, we were able to study the sense of agency for actions 456 enabled by a BMI-based intracortical neuroprosthesis and found that congruent sensory 457 feedback boosted agency and confidence when controlling BMI actions. Moreover, we 458 showed that human M1 processes not only motor and sensory information, but also 459 different levels of congruency between sensory and motor signals and the resulting 460 sense of agency. The present data are also of clinical relevance, because our NMES-461 462 based BMI approach, by providing congruent somatosensory feedback (without direct S1 stimulation) to a tetraplegic patient, improved the ability of the BMI classifier in 463 decoding the patient's motor commands. Interestingly, such higher BMI proficiency was 464 associated with a stronger sense of agency, suggesting that, beyond supporting close-465 loop systems and M1 feedback in general, somatosensory feedback and signals related 466 to subjective aspects of motor control (i.e., agency) are important input for improving 467 BMI proficiency. Quantifying subjective action-related mental states and including 468 controlled motor and sensory feedback may therefore provide new levels of comfort and 469 personalization and should be considered for the design of future BMIs. 470

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The present data demonstrate that M1 activity contains information specifically linked to 472 subjective aspects of motor control, in particular the sense of agency and confidence 473 that our participant associated with his BMI actions. It is known that agency likely 474 involves a network of multiple brain areas from which we did not record in the present 475 study (e.g., posterior parietal cortex³³ and angular gyrus; anterior insula^{34,35}; 476 supplementary motor cortex³⁶; premotor cortex³⁷; for reviews see ^{3,38}). However, our 477 findings – even if coming from a single tetraplegic patient - directly demonstrate that M1 478 activity contains sufficient information to decode actions for which a human participant 479 feels to be in control. The present BMI findings extend previous research that 480 investigated the sense of agency for non-invasive BCI, as based on scalp 481

electroencephalography (EEG). They add important information about the underlying 482 neural underpinnings based on M1 multiunit activity of the sense of agency in humans. 483 In keeping with a prominent line of research on the role of visuo-motor (and visuo-484 tactile) cues in boosting or modulating body ownership for artificial and real limbs³⁹⁻⁴². 485 previous BCI studies also demonstrated that coherent visual feedback results in a 486 higher sense of agency for BCI actions¹⁴. This effect is associated with stronger 487 activations in a cortical-subcortical network, recruited during motor imagery used to 488 control the BCI, consisting of posterior parietal cortex, insula, lateral occipital cortex, 489 and basal ganglia¹³. Another study¹⁰ further demonstrated that a stronger sense of 490 agency for BCI-mediated actions is associated with stronger activity in sensorimotor 491 areas during motor imagery based BCI. The present data on the sense of agency when 492 using an intracortical BMI, although from a single, highly proficient BMI user (see 493 below), demonstrate that this relationship can be tracked down even at the level of 494 multi-unit activity from M1 neurons, and it is further associated with higher BMI 495 proficiency. 496

497

498 Moreover, the present findings offer a mechanistic explanation for the relationship between sensorimotor activity, sensory feedback and the resulting sense of agency, by 499 showing that M1 activity before movement execution codes for the intended movement, 500 while activity after movement execution encodes the sensory feedback associated with 501 the implemented movement. By showing that somatosensory feedback in particular 502 503 affects the performance of the BMI classifier, these analyses provide insights into the sensorimotor mechanisms of BMI proficiency. We note that this last finding was 504 possible only due to the combination of a SCI lesion and the present intracortical BMI, 505 which allowed us to decode efferent signals and manipulate afferent signals, not only as 506 visual reproductions of body movements (via VR, as in previous studies), but also as 507 physical movements of the real body (via NMES). In order to highlight the dynamic, 508 multiscale brain mechanisms underlying the sense of agency in humans, future studies 509 should combine insights that can be gained from invasive BMI - with ultra-high spatial 510 resolution, but limited coverage in a handful of subjects - and non-invasive BCI - with 511 limited resolution, but recording from the entire brain in larger subject samples. 512

Finally, our results can be of interest not only within the field of neuroprosthetics and its clinical application, but also for basic neuroscience as well as current ethical and legal debates about the subjective sense of agency and responsibility when applying neurotechnology solutions for human repair or enhancement^{38,43,44}.

518 519

520 Limitations of the study

Because of the uniqueness of the present experimental setup, generalization of the 521 present findings to the general population should be done carefully. First, we tested a 522 single participant, who is an extremely trained BMI user, who could have developed an 523 extraordinary capacity of controlling his BMI system. This could have in turn impacted 524 the associated sense of agency and the discovered links with BMI proficiency. Second, 525 in order to enable movements of his upper limb, we used an NMES system that 526 provides a series of somatosensory cues, which are only partially comparable to those 527 associated with natural movements. For example, the intensity and temporal activation 528 of somatosensory fibers as well as of the recruited motor fibers (antidromic) differs from 529 sensorimotor activation during natural movements. We also note that although our 530 participant suffered severe somatosensory loss (following damage at the C5-C6 level), 531 he may have "learned" to associate some patterns of cutaneous sensations with the 532 specific type of NMES stimulation used to enable specific movements. Indeed, outside 533 of the experimental tasks described here, he was able to identify NMES-implemented 534 movements even without seeing his arm. Finally, given the long-term spinal cord lesion 535 suffered by this participant, we cannot exclude that some plastic changes have occurred 536 in his motor representations in M1, his somatosensory representations in S1, or the 537 connectivity between the two. There is still no consensus about plasticity following 538 spinal cord injury, with some evidence of preserved network organization, some 539 possible changes in grey matter density^{45,46} or activation in the sensorimotor 540 cortices^{47,48}. It is also not clear how these results at the population level of SCI patients 541 are predictive of changes at the single patient level. Thus, at the moment, it is not 542 possible to exclude that some of the present results are idiosyncratic to this particular 543 clinical case. 544

548 **References**

- 1. Blakemore, S. J., Wolpert, D. M. & Frith, C. D. Abnormalities in the awareness of action. *Trends Cogn. Sci.* **6**, 237–242 (2002).
- 551 2. Jeannerod, M. *Motor cognition : what actions tell the self*. (Oxford University 552 Press, 2006).
- 553 **3**. Haggard, P. Sense of agency in the human brain. *Nat. Rev. Neurosci.* **18**, 196– 554 207 (2017).
- 4. Hochberg, L. R. *et al.* Reach and grasp by people with tetraplegia using a neurally controlled robotic arm. *Nature* **485**, 372–375 (2012).
- 557 5. Collinger, J. L. *et al.* High-performance neuroprosthetic control by an individual 558 with tetraplegia. *Lancet* **381**, 557–564 (2013).
- 559 6. Bouton, C. E. *et al.* Restoring cortical control of functional movement in a human 560 with quadriplegia. *Nature* **533**, 247–250 (2016).
- Ajiboye, A. B. *et al.* Restoration of reaching and grasping movements through
 brain-controlled muscle stimulation in a person with tetraplegia: a proof-of-concept
 demonstration. *Lancet* 389, 1821–1830 (2017).
- Lebedev, M. A. & Nicolelis, M. A. L. Brain-Machine Interfaces: From Basic
 Science to Neuroprostheses and Neurorehabilitation. *Physiol. Rev.* 97, 767–837
 (2017).
- 567 9. Donoghue, J. P. Connecting cortex to machines: recent advances in brain 568 interfaces. *Nat. Neurosci.* **5**, 1085–1088 (2002).
- Nierula, B. *et al.* Agency and responsibility over virtual movements controlled
 through different paradigms of brain-computer interface. *J. Physiol.* 0, 1–16
 (2019).
- Nierula, B. & Sanchez-Vives, M. V. Can BCI Paradigms Induce Feelings of
 Agency and Responsibility Over Movements? in (2019). doi:10.1007/978-3-030 05668-1_10.
- 575 **12.** Sato, A. & Yasuda, A. Illusion of sense of self-agency: Discrepancy between the 576 predicted and actual sensory consequences of actions modulates the sense of 577 self-agency, but not the sense of self-ownership. *Cognition* **94**, 241–255 (2005).
- 13. Marchesotti, S. *et al.* Cortical and subcortical mechanisms of brain-machine interfaces. *Hum. Brain Mapp.* **38**, 2971–2989 (2017).
- Evans, N., Gale, S., Schurger, A. & Blanke, O. Visual Feedback Dominates the
 Sense of Agency for Brain-Machine Actions. *PLoS One* **10**, e0130019 (2015).
- 582 15. Knoblich, G. & Sebanz, N. Agency in the face of error. *Trends Cogn. Sci.* 9, 259–
 583 261 (2005).
- Tsakiris, M., Prabhu, G. & Haggard, P. Having a body versus moving your body:
 How agency structures body-ownership. *Conscious. Cogn.* 15, 423–432 (2006).
- Scott, S. H., Cluff, T., Lowrey, C. R. & Takei, T. Feedback control during voluntary
 motor actions. *Curr. Opin. Neurobiol.* 33, 85–94 (2015).

- Shokur, S. *et al.* Expanding the primate body schema in sensorimotor cortex by
 virtual touches of an avatar. *Proc. Natl. Acad. Sci.* **110**, 15121–15126 (2013).
- Hatsopoulos, N. G. & Suminski, A. J. Sensing with the motor cortex. *Neuron* 72, 477–487 (2011).
- 592 20. Hatsopoulos, N. G., Xu, Q. & Amit, Y. Encoding of movement fragments in the 593 motor cortex. *J. Neurosci.* **27**, 5105–14 (2007).
- Suminski, A. J., Tkach, D. C., Fagg, A. H. & Hatsopoulos, N. G. Incorporating
 Feedback from Multiple Sensory Modalities Enhances Brain-Machine Interface
 Control. *J. Neurosci.* **30**, 16777–16787 (2010).
- 597 22. Tkach, D., Reimer, J. & Hatsopoulos, N. G. Congruent activity during action and 598 action observation in motor cortex. *J. Neurosci.* **27**, 13241–50 (2007).
- Single-Neuron Activity in Premotor and Motor Cortex. *J. Neurophysiol.* 97, 4235–4257 (2007).
- Schwartz, A. B. Movement: How the Brain Communicates with the World. *Cell* 164, 1122–1135 (2016).
- 604 25. O'Doherty, J. E. *et al.* Active tactile exploration using a brain–machine–brain 605 interface. *Nature* **479**, 228–231 (2011).
- Stepniewska, I., Preuss, T. M. & Kaas, J. H. Architectionis, somatotopic
 organization, and ipsilateral cortical connections of the primary motor area (M1) of
 owl monkeys. *J. Comp. Neurol.* 330, (1993).
- Eickhoff, S. B. *et al.* Anatomical and functional connectivity of cytoarchitectonic areas within the human parietal operculum. *J. Neurosci.* **30**, (2010).
- Fetz, E. E., Finocchio, D. V., Baker, M. A. & Soso, M. J. Sensory and motor
 responses of precentral cortex cells during comparable passive and active joint
 movements. *J. Neurophysiol.* 43, (1980).
- 614 29. Hochberg, L. R. *et al.* Neuronal ensemble control of prosthetic devices by a 615 human with tetraplegia. *Nature* **442**, 164–171 (2006).
- 61630.Tabot, G. A. *et al.* Restoring the sense of touch with a prosthetic hand through a617brain interface. *Proc. Natl. Acad. Sci. U. S. A.* **110**, 18279–84 (2013).
- Flesher, S. N. *et al.* Intracortical microstimulation of human somatosensory cortex.
 Sci. Transl. Med. 8, 361ra141 LP-361ra141 (2016).
- Bensmaia, S. J. & Miller, L. E. Restoring sensorimotor function through
 intracortical interfaces: progress and looming challenges. *Nat. Rev. Neurosci.* 15, 313–325 (2014).
- 33. Desmurget, M. *et al.* Movement intention after parietal cortex stimulation in humans. *Science* 324, 811–3 (2009).
- Farrer, C. & Frith, C. D. Experiencing Oneself vs Another Person as Being the
 Cause of an Action: The Neural Correlates of the Experience of Agency.
 Neuroimage 15, 596–603 (2002).

- 628 35. Chambon, V., Wenke, D., Fleming, S. M., Prinz, W. & Haggard, P. An online 629 neural substrate for sense of agency. *Cereb. Cortex* **23**, 1031–1037 (2013).
- 36. Fried, I., Mukamel, R. & Kreiman, G. Internally Generated Preactivation of Single
 Neurons in Human Medial Frontal Cortex Predicts Volition. *Neuron* 69, 548–562
 (2011).
- Fornia, L. *et al.* Direct electrical stimulation of the premotor cortex shuts down awareness of voluntary actions. *Nat. Commun.* **11**, 1–11 (2020).
- 38. Sperduti, M., Delaveau, P., Fossati, P. & Nadel, J. Different brain structures
 related to self- and external-agency attribution: A brief review and meta-analysis. *Brain Struct. Funct.* 216, 151–157 (2011).
- Blanke, O., Slater, M. & Serino, A. Behavioral, Neural, and Computational
 Principles of Bodily Self-Consciousness. *Neuron* 88, 145–166 (2015).
- 40. Makin, T. R., Holmes, N. P. & Ehrsson, H. H. On the other hand: Dummy hands and peripersonal space. *Behav. Brain Res.* **191**, 1–10 (2008).
- Rognini, G. *et al.* Multisensory bionic limb to achieve prosthesis embodiment and
 reduce distorted phantom limb perceptions. *J. Neurol. Neurosurg. Psychiatry* 90,
 833–836 (2019).
- 42. Sanchez-Vives, M. V., Spanlang, B., Frisoli, A., Bergamasco, M. & Slater, M.
 Virtual hand illusion induced by visuomotor correlations. *PLoS One* 5, e10381
 (2010).
- 43. Yuste, R. *et al.* Four ethical priorities for neurotechnologies and Al. *Nature* **551**, 159–163 (2017).
- 44. Fried, I., Haggard, P., He, B. J. & Schurger, A. Volition and Action in the Human
 Brain: Processes, Pathologies, and Reasons. *J. Neurosci.* 37, 10842–10847
 (2017).
- 45. Wang, W. *et al.* Specific Brain Morphometric Changes in Spinal Cord Injury: A
 Voxel-Based Meta-Analysis of White and Gray Matter Volume. *J. Neurotrauma* 36, (2019).
- 46. Melo, M. C., Macedo, D. R. & Soares, A. B. Divergent Findings in Brain
 Reorganization After Spinal Cord Injury: A Review. *Journal of Neuroimaging* vol.
 30 (2020).
- Freund, P. *et al.* MRI investigation of the sensorimotor cortex and the
 corticospinal tract after acute spinal cord injury: A prospective longitudinal study.
 Lancet Neurol. 12, (2013).
- 48. Henderson, L. A., Gustin, S. M., Macey, P. M., Wrigley, P. J. & Siddall, P. J.
 Functional reorganization of the brain in humans following spinal cord injury:
 Evidence for underlying changes in cortical anatomy. *J. Neurosci.* **31**, (2011).
- Mallat, S. A Wavelet Tour of Signal Processing. A Wavelet Tour of Signal Processing (2009). doi:10.1016/B978-0-12-374370-1.X0001-8.
- 667 50. Humber, C., Ito, K. & Bouton, C. Nonsmooth Formulation of the Support Vector 668 Machine for a Neural Decoding Problem. *arXiv* (2010).

- 51. Colachis, S. C. *et al.* Dexterous control of seven functional hand movements
 using cortically-controlled transcutaneous muscle stimulation in a person with
 tetraplegia. *Front. Neurosci.* **12**, 1–14 (2018).
- 672 52. Colachis, S. C. I. Optimizing the Brain-Computer Interface for Spinal Cord Injury
 673 Rehabilitation. (2018).
- 53. Zhang, M. *et al.* Extracting wavelet based neural features from human intracortical recordings for neuroprosthetics applications. *Bioelectron. Med.* **4**, 11 (2018).
- 54. Quiroga, R. Q., Nadasdy, Z. & Ben-Shaul, Y. Unsupervised Spike Detection and
 Sorting with Wavelets and Superparamagnetic Clustering. *Neural Comput.* 16,
 1661–1687 (2004).
- 679 55. Hoerl, A. E. & Kennard, R. W. Ridge Regression: Biased Estimation for 680 Nonorthogonal Problems. *Technometrics* **12**, 55–67 (1970).
- 681 56. Kuhn, M. R Package: caret, Ver. 6.0-80. CRAN (2018).
- 682 57. Maris, E. & Oostenveld, R. Nonparametric statistical testing of EEG- and MEG-683 data. *J. Neurosci. Methods* **164**, 177–190 (2007).
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Competing interests: AS, MS, TB, NF, PS, and OB declare no competing interests.
CD, KE, PG, GS, NA, and DF hold patents for the BMI system.

710 Figure legends

711

Figure 1. Experimental setup. A. Events during trials. One (out of four possible 712 movements) was cued, following a "Go" signal to initiate the movement. The BMI 713 classifier decoded the movement from M1 activity and sensory feedback was given. The 714 715 patient answered two questions: Q1. "Are you the one who generated the movement?", by saying "Yes" or "No"; and Q2. "How confident are you?", by indicating a number 716 ranging from 0 (absolutely unsure) to 100 (absolutely sure). B: Example of sensory 717 feedback for one type of movement. The chosen movement was realized as a visual 718 feedback, via virtual reality (VR - Experiment 1), as a somatosensory feedback, via 719 NEMS (Experiment 2) or both (Experiment 3). In different congruency conditions, either 720 the cued and correctly decoded movement (Congruent) or the opposite movement 721 (Incongruent) was realized for the different modalities. The location of the electrodes 722 array in the M1, with respect to the pattern of activity for upper limb attempted 723 movement from fMRI is also shown (from ⁶). 724

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Figure 2. Agency judgements and confidence depends on sensory feedback. A. Proportion of "Yes" and "No" answers (Q1) to congruent and incongruent trials for the visual (Experiment 1) somatosensory (Experiment 2) and the combination of the two modalities (Experiment 3). B-C-D: Confidence about agency judgments. Distribution of Q2 responses as a function of the congruency of visual (B), somatosensory (C) or both (D) sensory feedback.

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Figure 3. M1 activity depends on sensory feedback. Sensory feedback as encoded 733 by Local field potentials (LFP; left panel) and Multiunit firing rates (MU; right panel). LFP 734 and MU modulation for congruent and incongruent visual (Experiment 1) and 735 somatosensory (Experiment 2) feedback (A) and for the combination of the two 736 (Experiment 3, B). Colored lines represent averaged signal across all channels (shaded 737 areas indicate SEMs); black lines report the time-related k-values of the multivariate 738 decoder distinguishing between congruent and incongruent feedback; the underlying 739 thick segments indicate k-values significantly higher than chance level from cluster-740 based permutation analyses. 741

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Figure 4. Sense of agency in M1. Sense of agency as coded by LFP (left) and Multi-743 unit firing rates (right). A. Left and right panels respectively show averaged LFP and 744 Multi-unit modulation for high (green) and no (grey) agency response to Q1 (shaded 745 areas indicate SEMs); black lines report the time-related k-values of the multivariate 746 747 decoder distinguishing the two conditions; the underlying thick segments indicate kvalues significantly higher than chance level from cluster-based permutation analyses. 748 B: Results of the decoder discriminating between high vs. low orthogonalized agency 749 scores from LFP (left) and MU (right) after regressing out for the effects of the 750 congruency of sensory feedback and type of movements. 751

752

Figure 5. Performance of BMI classifier as a function of sensory feedback and 753 sense of agency. A-B-C. The left panels (A, B, C) show the modulation in time of the 754 performance of the BMI classifier for the 4 types of movements indicated for the cued 755 movement (filled line) and the opposite (dashed line), as a function of feedback (i.e. 756 black dots indicate time points with significant difference). The right panels show the 757 area under the curve taken as an index of global performance of the BMI. The 758 performance of the BMI classifier does vary not as a function of visual feedback 759 (Experiment 1, A), but it is significantly better when somatosensory feedback is 760 congruent both in Experiment 2 (B) and in Experiment 3 (C). D. BMI accuracy in time as 761 a function of sense of agency. Blue/red curves represent the BMI classifier output for 762 763 the cued movement as a function of agency judgements (Q1: 1=high agency; 0=low agency) in conditions of equal sensory feedback. 764

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Figure 6. Somatosensory feedback changes firing rates of M1 neurons. A-B. Euclidean distance in time between trials with same motor intention and opposite feedback (red), same feedback and opposite intention (blue), or opposite congruent feedback and intention (green), for experiment 1 (A) and 2 (B). In Experiment 1, neural activity diverged as a function of motor intention before the movement, as shown by the increase in Euclidean distances between the green and blue curves. In Experiment 2, neural activity diverged as function of sensory feedback after NMES activation. C-D. Multidimensional scaling of neural activity before (-650/150 ms; C) and after (0/500 ms; D) sensory feedback. The plots show a 2D dimensionality reduction of population activity in the target period, in order to represent it on a plane. As in a principal component analysis, Dimensions 1 and 2 can be seen as the two abstract coordinates explaining most variance in the data. Movements are separated by classes of hand (open/close; right) and thumb (extension / flection; left) movements.

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785 Materials and Methods

786 **Participant**

The participant in this study was enrolled in a pilot clinical trial (NCT01997125) of a 787 custom neural bridging system (Battelle Memorial Institute) to reanimate paralyzed 788 upper limbs after C4-6 spinal cord injury. The system consisted of a Neuroport data 789 acquisition system (Blackrock Micro, Salt Lake, Utah), custom signal processing and 790 decoding algorithms (Battelle), and a NeuroLife Neuromuscular Stimulation System 791 (Battelle). The trial received investigational device exemption (IDE) approval by the US 792 Food and Drug Administration and Institutional Review Board approval through the Ohio 793 794 State University (Columbus, Ohio). The study conformed to institutional research requirements for the conduct of human subjects. The site of the experiments was the 795 Ohio State University NeuroRehabLab (Bockbrader, PI) and data was analyzed at Ohio 796 797 State (Columbus, Ohio) and École polytechnique fédérale de Lausanne (EPFL, Switzerland). The participant provided informed consent at the time of enrollment and 798 also provided written permission for photographs and videos. 799

The study participant was a 22 year-old male at the time of study enrollment. He had 800 complete C5 ASIA A, non-spastic tetraparesis from cervical spinal cord injury 801 802 associated with a diving accident 3 years prior. On neurological exam, he had full motor function bilaterally for C5 level muscles (e.g., biceps and shoulder girdle muscles), but 803 no motor function below the C6 level. He had 1/5 strength on the right and 2/5 strength 804 on the left for wrist extension (C6 level) on manual muscle testing. His sensory level 805 was C6 on the left and C5 on the right, although he had sensation for pressure on his 806 right thumb. He had preserved proprioception for shoulder, elbow and wrist joint 807 position, but was at chance level for distinguishing digit joint positions 808 (flexion/neutral/extension) for the thumb and fingers. He had mild finger flexor 809 contractures bilaterally, limiting finger extension at the proximal and distal 810 interphalangeal joints of digits 2-5. 811

He was implanted with a 4.4 x 4.2mm intracortical silicon Neuroport microelectrode array (Blackrock Microsystems) in the dominant hand/arm area of his motor cortex on 4/22/2014, as previously described⁶. The implant site was determined by preoperative functional neuroimaging obtained while the participant visualized movements of his right hand and forearm. He began using cortically-controlled transcutaneous neuromuscular
electrical stimulation (NMES) on his right forearm on 5/23/14. Data for this study was
collected over 13 sessions (45 hours) from 11/16/2016 - 2/20/2017, corresponding to
post-implant days 939-1035. One session with visual and NMES feedback was used for
practice (5 blocks of 32 trials on post-implant day 939). At the time of data collection,
the participant was an expert brain-machine interface (BMI) user with over 800 hours of
study participation.

- The participant underwent cognitive testing approximately one year after Utah array implantation (January – July, 2015). He scored with superior verbal abilities, attention, and working memory (ranging between 92nd - 99th percentile for his age).
- 826

827 Cortical Signal Acquisition and Classification

Neural data (96 channels) were acquired from the left motor cortex Utah array through 828 the Neuroport data acquisition system (Blackrock Micro). Raw data were processed 829 using analog hardware with 0.3Hz 1st order high-pass and 7.5kHz 3rd order Butterworth 830 low-pass filters, then digitized at 30,000 Hz. Data were divided into 100ms bins and 831 passed into Matlab (version 2014b), where signal artifact was removed by blanking over 832 3.5ms around artifacts (defined as signal amplitude >500µV at the same time on 4 of 12 833 randomly-selected channels). Signals were decomposed into mean wavelet power 834 (MWP) using the 'db4' wavelet over 100ms⁴⁹. Coefficients within the multiunit frequency 835 bands (234–3,750Hz, coefficients of scales 3, 4, 5, 6) were averaged across the 100ms 836 window and normalized by channel (by subtracting the mean and dividing by the 837 standard deviation of each channel and scale, respectively). Normalized coefficients for 838 each channel were averaged across scales 3-6, creating 96 MWP values (one for each 839 channel) per each 100ms. MWP values were fed as features into a real-time, nonlinear 840 support vector machine (SVM) classifier⁵⁰ with five classes (hand open, hand closed, 841 thumb extension, thumb flexion, and rest). Classifier activation values were computed 842 for each 100ms bin and ranged from -1 to 1. Classifier output represented the 843 movement pattern (hand open, hand closed, thumb extension, thumb flexion) with the 844 highest activation greater than threshold (zero). If no movement classes had activation 845

greater than zero, the classifier was in the "rest" state. If multiple output classes exceeded threshold, only the one with the highest score was used to provide feedback.

Signal quality was stable⁵¹ during the interval of data collection; but represented about a 30% decline in MWP normalized to post-implant 87^{52} . (See below for single unit statistics.) Average impedance was approximately 200 k Ω , a decline of 40% of its initial value. Average signal-to-noise was approximately 17.5dB, a decrease of about 10% of its initial value⁵³. Most of the decline in signal quality occurred in the first 400 days postimplantation.

854

855 Classifier Training and Neurally Controlled Hand Movements

Before each session, the SVM classifier was trained in an adaptive manner over 5 856 blocks. Each block consisted of 3 repetitions of 4 movements (hand open, hand closed, 857 thumb extension, thumb flexion) presented in a random order. Movements were cued 858 for 3-4s (4-5s inter-cue interval) using a small, animated hand in the corner of the video 859 display. Feedback was given with both NMES and the feedback hand on the video 860 screen. During the first training block, scripted feedback was provided simultaneously 861 with the cued movements. In subsequent blocks, appropriate movements were 862 activated when an output class for a given movement exceeded threshold (>0). 863 Training took approximately 10-15 minutes per session. 864

865

866 **Neuromuscular Electrical Stimulation**

The NMES system was used to evoke hand and finger movements by stimulating 867 forearm muscles. The system consisted of a multi-channel stimulator and a flexible, 868 130-electrode, circumferential forearm cuff. Coated copper electrodes with hydrogel 869 interfaces (Axelgaard, Fallbrook, CA) were 12mm in diameter, spaced at regular 870 intervals in an array (22mm longitudinally X 15mm transversely), and delivered current 871 in monophasic, rectangular pulses at 50Hz (pulse width 500µs, amplitude 0-20mA). 872 Desired hand/finger movements were calibrated at the beginning of each session by 873 determining/confirming the intensity and pattern of electrodes required to stimulate 874 intended movements. This took 5-10 minutes per session. 875

During the experiment, the participant's view of NMES-evoked movements was obscured from view by the video display. During Experiment 1, non-informative NMES feedback was given (current at an intensity equivalent to what was used for movement calibration patterns, but that did not evoke movement). During Experiments 2 and 3, NMES feedback was provided that evoked hand and finger movements.

881

882 Virtual Reality Animation

A non-immersive virtual reality system (i.e. without a head-mounted display or head-883 tracking) was used to provide visual feedback. This was done in order to adopt a 884 previous setup that the participant was already familiar with to the present experiments 885 and also facilitated the calibration procedure to train the BMI classifier. A physics-based 886 animated hand was used to provide visual feedback of classifier activation. During 887 training, two animated hands were displayed, a small cue hand at the bottom left and a 888 larger centrally-placed feedback hand (Figure 1 main text). During the experiment, the 889 display was oriented over the participant's forearm, a single, centrally-placed feedback 890 hand was displayed to match the size and location of the participant's right hand (the 891 cue hand was not displayed). During Experiments 1 and 3, feedback was provided 892 using the virtual hand. During Experiment 2, non-informative visual feedback was given 893 (the feedback hand remained in a neutral, rest position). 894

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896 **Feedback Congruency**

In half of the trials across Experiments, the visual and/or somatosensory feedback was covertly manipulated to be incongruent with the cue. In incongruent trials, when the participant correctly activated the classifier associated with the cue, he received feedback opposite to the cue (i.e., hand closed for "hand open", thumb extension for "thumb flex", etc.). In congruent trials, he received feedback consistent with the cue (i.e., hand open for "hand open", thumb flexion for "thumb flex", etc.).

903

904 Agency Assessment

All experimental trials began with a verbal cue ("hand open", "hand closed", "thumb extend", "thumb flex"), followed by a 2 second delay, then a verbal cue ("go"). During the next 4s, the participant was given feedback based on classifier activation levels, and then was told to "stop"). Over the next 5-5.5s, the participant reported whether he felt in control of the movement ("yes" or "no") and his degree of certainty (0-100). The next trial began at the end of this 5-5.5s interval. There were 32 trials per block in Experiments 1 and 2 and 26 trials per block in Experiment 3.

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913 Trial Selection and Time-locking

To ensure that the participant was successfully activating the classifier for the cued 914 movement, and the signal can be meaningfully time-locked to movement onset, we 915 applied the following selection criteria on the trials. We consider it as a correct imagined 916 movement when the participant is able to maintain the classifier of the cued movement 917 above the threshold for at least 600 ms (6 classifier output bins). We retain trials in 918 which at least one correct movement happens between the GO cue and 1.5 seconds 919 before the STOP cue. Epochs are then constructed by time-locking every trial with 920 respect to the onset of such imagined movements. In case several correct movements 921 occurred during the same trial, the time-locking is relative to the first movement. 922 Furthermore, we excluded 128 trials from the session on which the participant 923 systematically reported problems with controlling the BMI system and absent subjective 924 agency. Globally, we retained 846 out of 1408 trials (60%). 925

Note that, since we define the onset as the beginning of the 100 ms bin of neural activity that is fed to the classifier, and around 50 ms are required to compute the output, the corresponding feedback is received about 150 ms after the onset of the imagined movement.

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931 Experiment 1: Agency Assessment with Virtual Hand Feedback and Non-932 informative NMES

Twelve blocks of 32 trials were collected on post-implant days 953 (4 blocks), 988 (4 blocks), and 1035 (4 blocks). In each trial, the participant received a verbal cue to perform a movement ("hand open", "hand closed", "thumb extend", "thumb flex"). When a classifier crossed threshold during the 4 second feedback window, feedback was given by showing movement of the virtual hand and by activating non-informative NMES (radial wrist electrode activation that did not elicit movement, did not vary from trial to trial, and that the participant could feel and distinguish from real NMES feedback). Feedback on half of the trials was randomly selected to be incongruent with the cue. His subjective sense of agency and level of certainty were recorded for each trial.

A total of 384 trials were collected across three days. After removing trials where the cued action could not be correctly decoded and the session on post-implant day 1035 (see trial selection paragraph), 83 congruent and 72 incongruent trials remained for behavioral and neural activity analysis.

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Experiment 2: Agency Assessment with NMES Feedback and Non-informative Virtual Hand

Twelve blocks of 32 trials were collected on post-implant days 941 (5 blocks), 960 (3 949 blocks), and 967 (4 blocks). In each trial, the participant received a verbal cue to 950 perform a movement ("hand open", "hand closed", "thumb extend", "thumb flex"). When 951 a classifier crossed threshold during the 4 second feedback window, feedback was 952 given by activating movement of the participant's hand and wrist through NMES and 953 showing non-informative visual feedback (non-moving hand). The participant could not 954 see his own hand/wrist, but could distinguish his hand state based what the stimulation 955 patterns felt like to him. Feedback on half of the trials was randomly selected to be 956 incongruent with the cue. His subjective sense of agency and level of certainty were 957 recorded for each trial. 958

A total of 384 trials were collected across three days. After removing trials where the participant did not respond correctly by activating the classifier associated with the cue, 154 congruent and 89 incongruent trials remained for behavioral and neural activity analysis.

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Experiment 3: Agency Assessment with Virtual Hand and NMES Feedback

Twenty blocks of 32 trials were collected on post-implant days 993 (3 blocks), 990 (5 blocks), 1007 (4 blocks), 1014 (3 blocks), and 1021 (5 blocks). In each trial, the participant received a verbal cue to perform a movement ("hand open", "hand closed", "thumb extend", "thumb flex"). When a classifier crossed threshold during the 4 second feedback window, feedback was given by activating movement of the participant's hand and wrist through NMES and showing movement of the virtual hand. The participant could not see his own hand/wrist, but could distinguish his hand state based what the 972 stimulation patterns felt like to him. Congruency with respect to the cue was 973 manipulated independently in the visual and somatosensory modalities such that 25% 974 of the trials were each: congruent for both visual and NMES feedback, incongruent for 975 both visual and NMES feedback, congruent for visual but incongruent for NMES 976 feedback, congruent for NMES but incongruent for visual feedback. His subjective 977 sense of agency and level of certainty were recorded for each trial.

A total of 520 trials were collected across five days. After removing trials where the participant did not respond correctly by activating the classifier associated with the cue, the number of trials that remained for behavioral and neural activity analysis were: 117 congruent for both visual and NMES feedback, 103 incongruent for both visual and NMES feedback, 101 congruent for visual and incongruent for NMES feedback, and 127 congruent for NMES and incongruent for visual feedback.

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985 Firing Rate Calculation and Single Unit Analyses

Single units were identified through offline data processing. For each block, raw voltage 986 recordings at each channel were processed in a series of steps. First, FES stimulation 987 988 artifact was removed using a 500µV threshold and 3.5ms artifact removal time window. The removed window was replaced with an interpolated segment to retain temporal 989 information. Then, the raw signal with FES artifact removed was processed with a 300-990 3000Hz bandpass filter. The filtered data was fed into an automated spike detection and 991 sorting algorithm, wave_clus⁵⁴ using the default optimization settings. A threshold was 992 993 set to four times the standard deviation of the noise and used to detect spike locations. A wavelet decomposition was performed on the spikes to extract features and a 994 superparamagnetic clustering algorithm was used to cluster the spikes into groups, 995 996 representative of individual single units. The superparamagnetic clustering algorithm was used to eliminate spikes that were considered noise to ensure only single units 997 were analyzed. As spike sorting was not performed before data collection, there was no 998 way to match single units across days. Additionally, the number of single units detected 999 at a given channel fluctuated between days, possibly due to micro-movement of the 1000 array and brain state changes. For this reason, all single units detected at a given 1001 channel were considered the same and pooled at the single channel level as multiunit 1002 1003 activity in subsequent analysis.

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1005 Offline neural decoding

Sensory feedback congruency and subjective ratings (Q1 and Q2) were decoded offline both from LFPs and from multiunit activity. For LFP analysis, the signal amplitude for each channel was downsampled to 500 Hz, band-passed between 0.1 and 12 Hz with an IIR filter and smoothed using sliding averaging windows of 250 ms. Following multiunit spike times calculation (see above), multiunit firing rate was estimated at 20 Hz over a 250 ms sliding window.

We fed each channel's signal amplitude (LFP) or firing rate (multiunit) as predictors to a 1012 penalized linear decoder based on ridge regressions⁵⁵. A separate model was trained to 1013 decode congruency (Q1) or confidence (Q2) on each signal timepoint, with a sampling 1014 1015 rate of 20 and 500 Hz for multiunit and LFPs respectively. Decoding performance was evaluated by computing and averaging Cohen's k (logistic regression; Q1) or R2 (linear 1016 1017 regression; Q2) values over 10 independent 10-fold cross validation runs. The regression was performed through the "train" function of the R "caret" package⁵⁶. To 1018 evaluate the statistical significance of the decoding, we generated a null decoding 1019 1020 performance distribution by applying the same decoding methods on the data after randomly shuffling Q1 and Q2 values. 1000 permutations were generated, and the 1021 decoding performance was evaluated for each of them. Then, a t-value was assigned to 1022 every time-point both in real and permuted data, by comparing its decoding 1023 performance to the null distribution of permuted data. Finally, the t-values were used to 1024 define significant decoding time windows based on a cluster-based permutation test on 1025 each epoch's largest cluster⁵⁷. After checking that the t-value threshold used to define 1026 clusters was not significantly affecting the results, its value was set at 2. 1027

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1029 Computation of distance between neural activity patterns

Since the neural activity recorded by the microelectrode array can change significantly between experimental sessions (i.e., days of recording) spanned by our analysis, Euclidean distances per each pair of conditions were computed separately within each day of recording and then averaged to obtain the final results. Confidence intervals were obtained through a bootstrapping technique, again applied within sessions. For each session and condition, we extracted n random trials with replacement, where n is the number of trials for that condition/session, and the final Euclidean distance was obtained by averaging across sessions as described above. The procedure was repeated 100 times, and 95 % confidence intervals were obtained as 1.96 times the standard deviation of the surrogated distribution obtained as explained here.

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1041 Multidimensional scaling

In order to graphically represent the spatio-temporal patterns of neural activity, we 1042 performed a multidimensional scaling (MATLAB function mds) on correlation distances 1043 computed between spatio-temporal patterns of neural activity. Also, in this case, to 1044 1045 avoid including sources of variances due to the change in signal between experimental sessions, the procedure was run within experimental sessions. In order to obtain 1046 correlation distances between trials we started by concatenating, for each trial, data 1047 1048 from all channels and timepoints within the selected temporal window. Then, we computed the correlation coefficient of the resulting vector with the equivalent vector 1049 from all other trials within the same session and subtracted the obtained values to 1 to 1050 1051 obtain values of the correlation distance. The first two dimensions of the multidimensional scaling were then aligned across sessions via the Procrustes analysis 1052 1053 (MATLAB function Procrustes), using the means by conditions (combinations of movement/somatosensory feedback) in the first session as a reference. 1054

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1057 Data availability statement

1059 Behavioral data and processed data necessary to reproduce the figures in the main text 1060 can be found in the OSF repository accessible at:

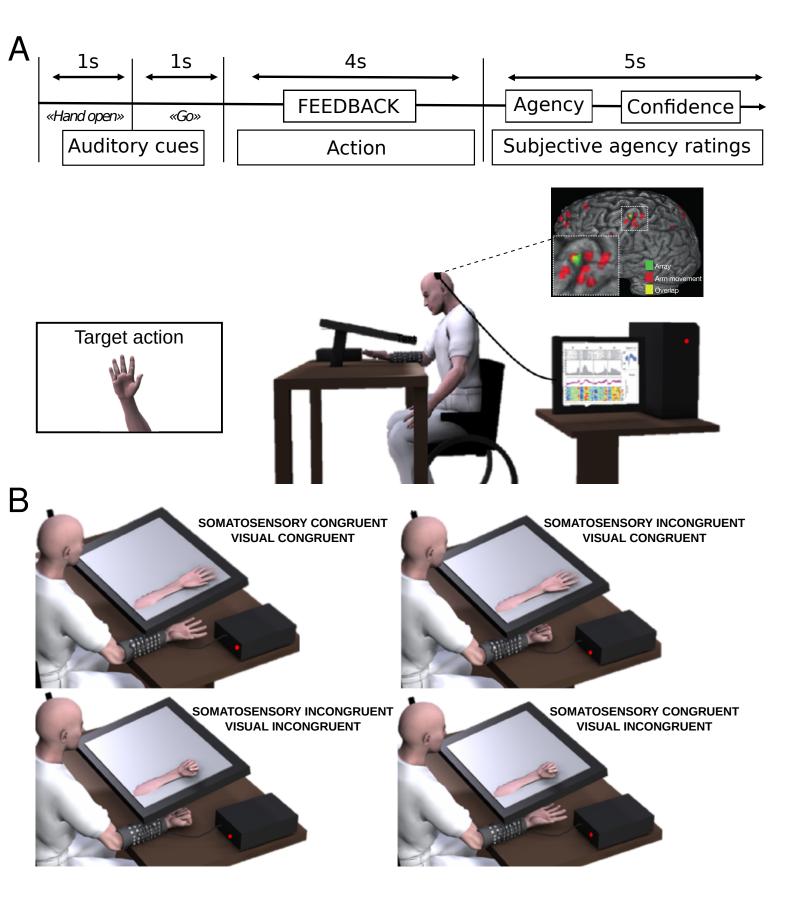
1061 https://osf.io/7rma5/?view_only=9928bd8e32a748828f7ecfdbeb1f8baa.

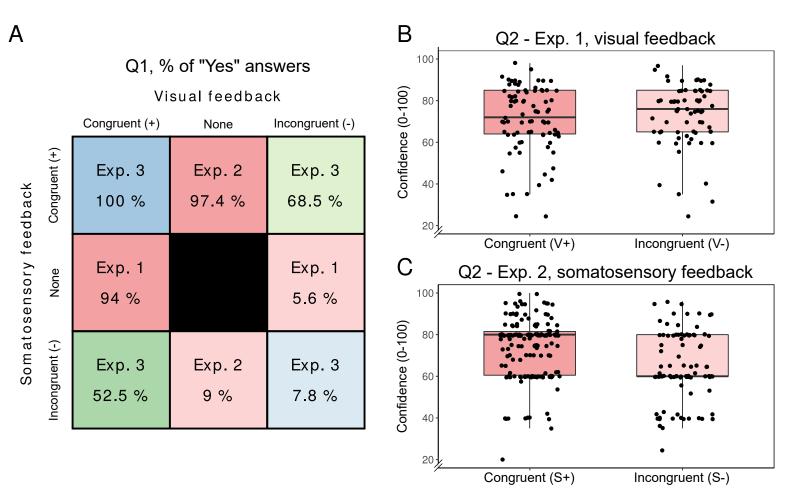
Neural data and code for BMI control can be made available to qualified individuals for
 collaboration via a written agreement between Battelle Memorial Institute and the
 requester's affiliated institution. Such inquiries or requests should be directed to:
 <u>ganzer@battelle.org</u>.

1066

1067 **Code availability statement**

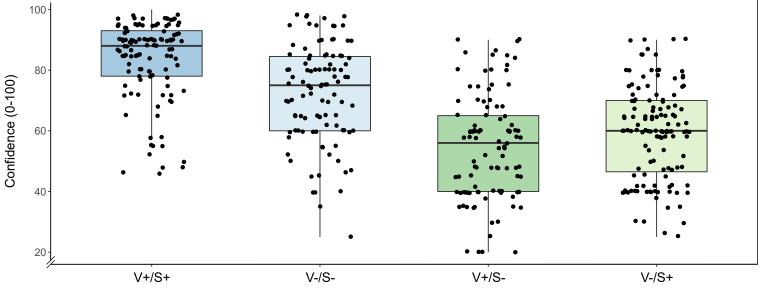
1069 Custom code for neural data analysis and BMI control can be obtained following a 1070 written agreement between Battelle Memorial Institute and the requester's affiliated 1071 institution. Such inquiries or requests should be directed to: <u>ganzer@battelle.org</u>. 1072 Inquiries or requests concerning custom analysis code used for this study should be 1073 directed to AS.

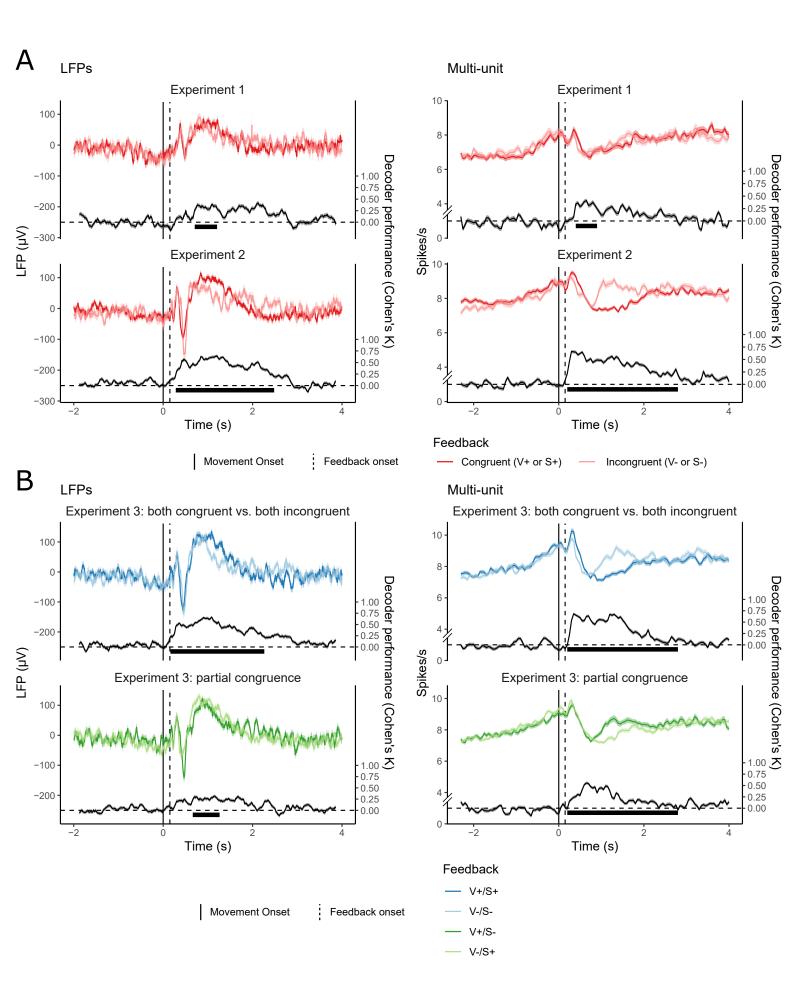


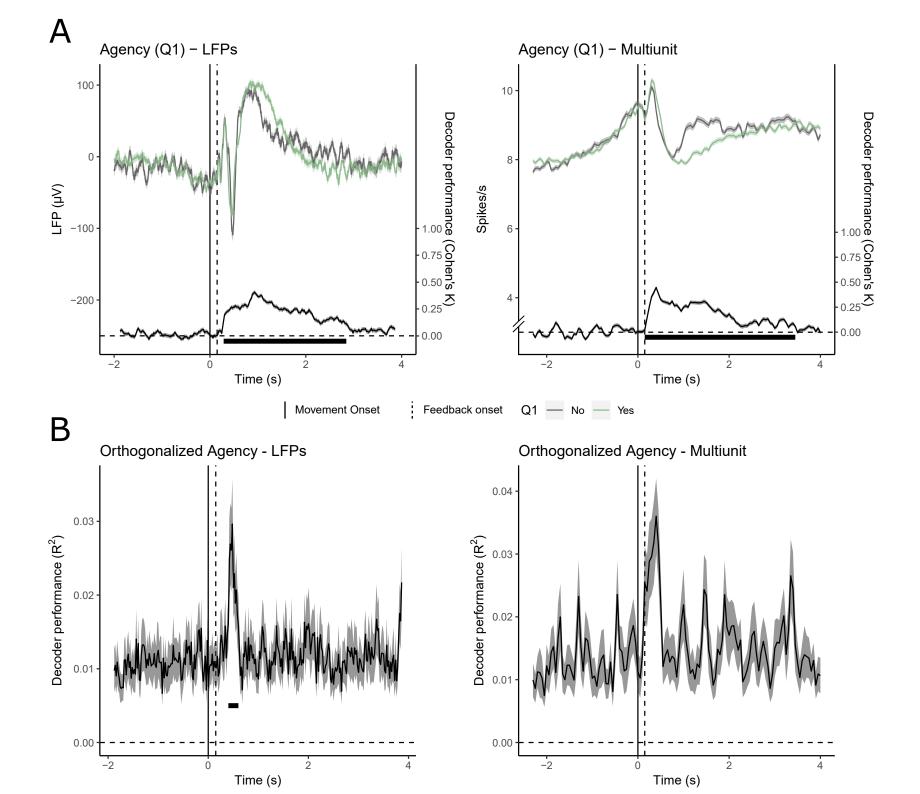


D

Q2 - Exp. 3, visual and somatosensory feedback







Experiment 1 – Visual feedback Α 40 0.5 Classifier AUC 30 0.0 -0.5 Congruent (V+) Incongruent (V-) $\mathbf{B}^{-1.0}$ Experiment 2 – Somatosensory feedback Decoder output (A.U.) 0.5 Classifier AUC 30 0.0 -0.5 10 **C**^{-1.0} Congruent (S+) Incongruent (S-) Experiment 3 - Visual and somatoensory feedback 0.5 30 Classifier AUC 0.0 -0.5 10 **e**. -1.0 V+/S+ V-/S-V+/S-V-/S+ Time (s) Movement type Feedback Cued Cued Congruent (V+ or S+) Movement Onset Feedback onset Incongruent (V- or S-) V+/S-V-/S+

