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Can we decode phonetic features in inner speech using surface electromyography?

Ladislas Nalborczyk 1 , 2 *, Romain Grandchamp 1 , Ernst H. W. Koster 2 , Marcela Perrone-Bertolotti 1 , 3 , Hélène Lœvenbruck 1

1 Univ. Grenoble Alpes, CNRS, LPNC, 38000 Grenoble, France

2 Department of Experimental Clinical and Health Psychology, Ghent University

3 Institut Universitaire de France (IUF), France

* Corresponding author: ladislas.nalborczyk@univ-grenoble-alpes.fr

Abstract

Although having a long history of scrutiny in experimental psychology, it is still controversial whether wilful inner speech (covert speech) production is accompanied by specific activity in speech muscles. We present the results of a preregistered experiment looking at the electromyographic correlates of both overt speech and inner speech production of two phonetic classes of nonwords. An automatic classification approach was undertaken to discriminate between two articulatory features contained in nonwords uttered in both overt and covert speech. Although this approach led to reasonable accuracy rates during overt speech production, it failed to discriminate inner speech phonetic content based on surface electromyography signals alone. However, exploratory analyses conducted at the individual level revealed that it seemed possible to distinguish between rounded and spread nonwords covertly produced, in two participants. We discuss these results in relation to the existing literature and suggest alternative ways of testing the engagement of the speech motor system during wilful inner speech production.

Introduction

As you read these words, you may be experiencing the presence of a familiar speechlike companion. This internal speech production may accompany daily activities such as reading (see [1-4], but see [5,6]), writing ([7]), memorising ([8,9]), future planning [8], problem solving [9,10] or musing (for reviews see [11–14]). Several studies using experience sampling or questionnaires (e.g.,[15,16]) have shown that by deliberately paying attention to this internal speech, one can examine its phenomenological properties such as identity (whose voice is it?) or other high-level characteristics (e.g., is it gendered?). Moreover, it is often possible to examine lower-level features like the tone of the inner speech, its pitch or its tempo. This set of basic observations leads to some important insights about the nature of inner speech. The simple fact that we can make sensory

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judgements about our inner speech tautologically reveals that inner speech is accompanied by sensory percepts (e.g., speech sounds, kinaesthetic feelings). Some of these introspective accounts have been examined, tested and complemented using empirical methods from cognitive neuroscience. As summarised in [17], behavioural and neuroimaging data reveal that some variants of inner speech are associated with auditory and/or somatosensory sensations that are reflected by auditory and/or somatosensory cortex activations. Visual representations may also be at play, typically for inner language in the deaf population. Inner verbalising therefore involves the reception of imaginary multisensory signals. This leads to other fundamental questions: where do these percepts come from? Why do they sound and feel like the ones we experience when we actually (overtly) speak?

Two main classes of explanatory theories have been offered to answer these questions. A first class of theories, that derives from Vygotky's views on language and thought, and that we describe as the abstraction view [18], suggest that inner speech is profoundly internalised, abbreviated and condensed in form. Vygotsky suggested that inner speech evolved from so-called egocentric speech (i.e., self-addressed overt speech or private speech), via a gradual process of internalisation during childhood [19]. According to him, the properties of speech are transformed during this internalisation, and inner speech cannot be merely be described as a weakened form of overt speech (as claimed for instance by [20]). This has led some scholars to conceive of inner speech as predominantly pertaining to semantics, excluding any phonological, phonetic, articulatory or even auditory properties (e.g., [18,21,22]). The property of abbreviation and condensation is supported by several psycholinguistic experiments on the qualitative and quantitative differences between overt and covert speech, as concerns rate and error biases (e.g., [18,22-24], but see [25]). Such condensation implies that the auditory gualities mentioned above would only rarely be observed during introspection and would merely be the result of learned associations between abstract linguistic representations and auditory percepts. A second class of theories is described under the umbrella term of motor simulation view. These theories suggest that inner speech can be conceived as a kind of action on its own [26,27], produced in the same way as overt speech is, except that the last stage of articulatory execution is only simulated. Most theories under this view share the postulate that the speech motor system is involved (to some extent) during inner speech production and that the auditory and somatosensory consequences of the simulated articulatory movements constitute the inner speech percepts referred to in subjective studies.

As explained in the ConDialInt model [28], these two views can be reconciled if various degrees of unfolding of inner speech are considered. Fully condensed forms of inner speech only involve semantics, and are deprived of the acoustic, phonological and syntactic qualities of overt speech. Expanded forms inner speech, on the other hand, presumably engage prosodic and morpho-syntactic formulation as well as phonological specification, articulatory simulation and the perception of an inner voice. Between the fully condensed abstract forms and the expanded articulation-ready form, it can be assumed that various semi-condensed forms may exist, with morphosyntactic properties and perhaps even phonological features, depending on the stage at which the speech production process is truncated. Such a view was also taken by [29] who has

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suggested that inner speech varies with cognitive demands and emotional conditions on a continuum between extremely condensed and expanded forms (see also [11,27]). Therefore, the two views of inner speech (abstraction vs. simulation) can be construed as descriptions of two opposite poles on the condensation dimension. On the most expanded side of the continuum, inner speech entails full phonetic specification and articulatory simulation. It might therefore be expected that speech motor activity could be detectable. If the motor simulation view is correct, then motor activity could be recorded during expanded forms of inner speech. If, on the other hand, the abstraction view applies to all forms of inner speech, then no motor activity should be present, even in phonologically-expanded forms.

Previous research has demonstrated that it is possible to record muscle-specific electromyographic correlates of inner speech (e.g., [30,31]). However, these studies mostly focused on small samples of participants and sometimes used invasive intramuscular electromyography. In contrast, more recent research studies using surface electromyography lead to mixed results (e.g.,[32]). Building upon previous work, we describe an experimental set-up using surface electromyography with the aim of testing the involvement of specific speech muscle groups during the covert production of phonologically expanded speech forms.

Inner speech as motor imagery of speech

Speech production is a complex motor action, involving the fine-grained coordination of more than 100 muscles in the upper part of the body [33]. In adult humans, its covert counterpart (referred to as *inner speech* or *verbal imagery*) has developed to support a myriad of different functions. In the same way as visual imagery permits to mentally examine visual scenes, *verbal imagery* can be used as an internal tool, allowing –amongst other things- to rehearse or to prepare past or future conversations [11,14]. Because speech production results from sequences of motor commands which are assembled to reach a given goal, it belongs to the broader category of motor actions [34]. Therefore, a parallel can be drawn between verbal imagery and other forms of motor imagery (e.g., imagined walking or imagined writing). Accordingly, studies on the nature of inner speech might benefit from insights gained from the study of motor imagery and the field of motor cognition [34,35].

Motor imagery can be defined as the mental process by which one 101 rehearses a given action, without engaging in the physical movements 102 involved in this particular action. One of the most influential theoretical 103 accounts of this phenomenon is the motor simulation theory [34,36,37]. In 104 this framework, the concept of simulation refers to the "offline rehearsal of 105 neural networks involved in specific operations such as perceiving or acting" 106 [34]. The MST shares some similarities with the theories of embodied and 107 grounded cognition [38] in that both account for motor imagery by appealing 108 to a simulation mechanism. However, the concept of simulation in grounded 109 theories is assumed to operate in order to acquire specific conceptual 110 knowledge [39], which is not the concern of the MST. In other words, we 111 should make a distinction between embodiment of content, which concerns 112 the semantic content of language, and *embodiment of form*, which concerns 113 "the vehicle of thought", that is, proper verbal production [40]. 114

A second class of explanatory models of motor imagery are concerned

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with the phenomenon of *emulation* and with *internal models* [41]. Internal 116 model theories share the postulate that action control uses internal models, 117 that is, systems that simulate the behaviour of the motor apparatus [42,43]. 118 The function of internal models is to estimate and anticipate the outcome of 119 a motor command. Among the internal model theories, motor control 120 models based on robotic principles [44,45] assume two kinds of internal 121 models (that are supposed to be coupled and regulated): a forward model 122 (or simulator) that predicts the sensory consequences of motor commands 123 from efference copies of the issued motor commands, and an inverse model 124 (or controller) that calculates the feedforward motor commands from the 125 desired sensory states [17,41]. 126

Emulation theories [46,47] borrow from both simulation theories and internal model theories and provide operational details of the simulation 128 mechanism. In the emulation model proposed by [46], the *emulator* is a 129 device that implements the same input-output function as the body (i.e., the 130 musculoskeletal system and relevant sensory systems). When the emulator 131 receives a copy of the control signal (which is also sent to the body), it 132 produces an output signal (the emulator feedback), identical or similar to the 133 feedback signal produced by the body, yielding mock sensory percepts (e.g., 134 visual, auditory, kinaesthetic) during motor imagery. 135

By building upon models of speech motor control [45,48], a recent model describes wilful (voluntary) expanded inner speech as "multimodal acts with multisensory percepts stemming from coarse multisensory goals" [17]. In other words, in this model the auditory and kinaesthetic sensations perceived during inner speech are assumed to be the predicted sensory consequences of simulated speech motor acts, emulated by internal forward models that use the efference copies of motor commands issued from an inverse model [46]. In this framework, the peripheral muscular activity recorder during inner speech production is assumed to be the result of *partially* inhibited motor commands. It should be noted that both simulation, emulation, and motor control frameworks can be grouped under the *motor simulation view* and altogether predict that the motor system should be involved to some extent during motor imagery, and by extension, during inner speech production. We now turn to a discussion of findings related to peripheral muscular activity during motor imagery and inner speech.

Electromyographic correlates of covert actions

Across both simulationist and emulationist frameworks, motor imagery has 152 consistently been defined as the mental rehearsal of a motor action without 153 any overt movement. One consequence of this claim is that, in order to 154 prevent execution, the neural commands for muscular contractions should 155 be blocked at some level of the motor system by active inhibitory 156 mechanisms [49]. Despite these inhibitory mechanisms, there is abundant 157 evidence for peripheral muscular activation during motor imagery [49–51]. 158 As suggested by [36], the incomplete inhibition of the motor commands 159 would provide a valid explanation to account for the peripheral muscular 160 activity observed during motor imagery. This idea has been corroborated by 161 studies of changes in the excitability of the motor pathways during motor 162 imagery tasks [52]. For instance, [53] measured spinal reflexes while 163 participants were instructed to either press a pedal with the foot or to 164 simulate the same action mentally. They observed that both H-reflexes and 165

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T-reflexes increased during motor imagery, and that these increases correlated with the force of the simulated pressure. Moreover, the pattern of results observed during motor imagery was similar (albeit weaker in amplitude) to that observed during execution, supporting the *motor simulation view* of motor imagery. Using transcranial magnetic stimulation, several investigators observed muscle-specific increases of motor evoked potentials during various motor imagery tasks, whereas no such increase could be observed in antagonist muscles [54,55].

When considered as a form of motor imagery, inner speech production is 174 also expected to be accompanied with peripheral muscular activity in the 175 speech muscles. This idea is supported by many studies showing peripheral 176 muscular activation during inner speech production [10,30,31,56–58], during 177 auditory verbal hallucinations in patients with schizophrenia [59], or during 178 induced mental rumination [60]. Some authors also recently demonstrated 179 that it is possible to discriminate inner speech content based on surface 180 electromyography (EMG) signals with a median 92% accuracy [61]. However, 181 other teams failed to obtain such results [32]. 182

Many of these EMG studies concluded on the involvement of the speech motor system based on a difference in EMG amplitude by contrasting a period of inner speech production to a period of rest. However, as highlighted by [62], it is usually not enough to show an increase of speech muscle activity during inner speech to conclude that this activation is related to inner speech production. Indeed, three sorts of inference can be made based on the studies of electromyographic correlates of inner speech production, depending on the stringency of the control procedure. The stronger sort of inference is permitted by highlighting a discriminative pattern during covert speech production, as for instance when demonstrating a dissociation between different speech muscles during the production of speech sounds of different phonemic class (e.g. contrasting labial versus non-labial words). According to [62], other (weaker) types of control procedures include i) comparing the EMG activity during covert speech production to a baseline period (without contrasting phonemic classes in covert speech utterances), or ii) comparing the activity of speech-related and non-speech related (e.g., forearm) muscle activity. Ideally, these controls can be combined by recording and contrasting speech and non-speech related muscles in different conditions (e.g., rest, covert speech, overt speech) of pronunciation of different speech sounds classes (e.g., labial versus non-labial).

Previous research studies carried out using the preferred procedure recommended by [62] suggest a discriminative patterns of electromyographic correlates according to the phonemic class of the words being covertly uttered [30,31], which would corroborate the *motor simulation view* of inner speech. However, these studies used limited sample sizes (often less than ten participants) and worked mostly with children. These factors limit the generalisability of the above findings because i) low-powered experiments provide biased estimates of effects, ii) following the natural internalisation process, inner speech muscular correlates are expected to weaken with age and iii) a higher sensitivity could be attained by using modern sensors and signal processing methods.

The present study intends to bring new information to the debate215between the motor simulation view and the abstraction view of inner speech,216by focusing on an expanded form of inner speech, wilful nonword covert217

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production. This work can be seen as a replication and extension of previous 218 works carried out by McGuigan and collaborators [30,31]. We aimed to 219 demonstrate similar dissociations by using surface electromyography 220 recorded over the lip (orbicularis oris inferior, OOI) and the zygomaticus 221 *major* (ZYG) muscles. More precisely, given that rounded phonemes (such as 222 /u/) are articulated with orbicular labial contraction, whereas spread 223 phonemes (such as /i/) are produced with zygomaticus contraction, if the 224 motor simulation view is correct, we should observe a higher average EMG 225 amplitude recorded over the OOI during both the overt and inner production 226 of rounded nonwords in comparison to spread nonwords. Conversely, we 227 would expect a lower average EMG amplitude recorded over the ZYG during 228 both the inner and overt production of rounded nonwords in comparison to 229 spread nonwords. In addition, we would not expect to observe 230 content-specific differences in EMG amplitude concerning the non 231 speech-related muscles (i.e., forehead and forearm muscles). 232

Methods

In the *Methods* and *Data analysis* sections, we report how we determined our sample size, all data exclusions, all manipulations, and all measures in the study [63]. A pre-registered version of our protocol can be found on OSF: https://osf.io/czer4/. 237

Participants

As previous studies of the electromyographic correlates of inner speech were 239 mostly carried out with samples of children or young adults, used different 240 kinds of EMG measures (surface EMG or needle EMG), and different kinds of 241 signal processing methods, it was impractical to determine the effect size of 242 interest for the current study. Therefore, we used sequential testing as our 243 sampling procedure, based on the method described in [64] and [65]. We 244 fixed a statistical threshold to $BF_{10} = 10$ and $BF_{10} = 1/10$ (i.e., $BF_{01} = 10$), 245 testing the difference between the inner production of labial items versus 246 the inner production of non-labial items on the standardised EMG amplitude 247 of the lower lip (orbicularis oris inferior). In order to prevent potential 248 experimenter and demand biases during sequential testing, the 249 experimenter was blind to BFs computed on previous participants [66]. All 250 statistical analyses have been automatised and a single instruction was 251 returned to the experimenter (i.e., "keep recruiting participants" or "stop the 252 recruitment"). We fixed the maximum sample size to 100 participants. 253

As a result of the above sampling procedure, a total of 25 254 French-speaking female undergraduate students in Psychology from the Univ. 255 Grenoble Alpes (mean age = 19.57, SD = 1.1). took part in this experiment, 256 in exchange for course credits. It should be noted that this procedure did not 257 work optimally because we later spotted an error in the EMG signal 258 processing workflow. Thus, the sequential testing stopped earlier than it 259 should have. These participants were recruited via mailing list, online 260 student groups, and posters. Each participant provided a written consent 261 and the present study was approved by the local ethics committee (Grenoble 262 CERNI agreement #2016-05-31-9). 263

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Material

EMG recordings

EMG activity was recorded using TrignoTM Mini sensors (Delsys Inc.) with a sampling rate of 1926 samples/s, a band pass of 20 Hz (12 dB/ oct) to 450 Hz (24 dB/oct) and were amplified by a TrignoTM 16-channel wireless EMG system (Delsys Inc.). These sensors consist of two 5 mm long, 1 mm wide parallel bars, spaced by 10 mm, which were attached to the skin using double-sided adhesive interfaces. The skin was cleaned by scrubbing it with 70% isopropynol alcohol. EMG signals were synchronised using the PowerLab 16/35 (ADInstrument, PL3516). Raw data from the EMG sensors were then resampled at a rate of 1 kHz and stored in digital format using Labchart 8 software (ADInstrument, MLU60/8).

EMG sensors were positioned over five muscles: the corrugator supercilii 276 (COR), the frontalis (FRO), the zygomaticus major (ZYG), the orbicularis oris 277 inferior (OOI), and the flexor carpi radialis (FCR).1 The two speech-related 278 muscles (OOI and ZYG) were chosen to show speech-specific EMG correlates, 279 whereas the two non-speech related facial muscles (ZYG and FRO) were 280 chosen to control for overall facial muscular activity. We also recorded the 281 activity of the FCR of the non-dominant forearm to control for overall (body) 282 muscular activity. 283

As reviewed in [67], the dominant side of the face displays larger movements than the left side during speech production, whereas the non-dominant side is more emotionally expressive. Therefore, we recorded the activity of control and emotion-linked muscles (i.e., COR and FRO) that were positioned on the non-dominant side of the face (i.e., the left side for right-handed participants), while sensors recording the activity of the speech muscles (i.e., ZYG and OOI) were positioned on the dominant side of the face.

The experiment was video-monitored using a Sony HDR-CX240E video camera to track any visible facial movements. A microphone was placed 20–30 cm away from the participant's lips to record any faint vocal production during the inner speech and listening conditions. Stimuli were displayed using the OpenSesame software [68] on a 19-inch colour monitor.

Linguistic material

We selected ten rounded and ten spread bi-syllabic nonwords (cf. Table 1). Each class of nonwords was specifically designed to either induce a greater 298 activation of the lip muscle (rounded items) or a greater activation of the 200 zygomaticus muscle (spread items). These stimuli were selected based on 300 phonetic theoretical constraints, with the aim of maximising the differences 301 between the two classes of non-words in their involvement of either the OOI 302 or the ZYG muscle. More precisely, rounded items consisted in the repetition 303 of a syllable containing a bilabial consonant followed by a rounded vowel, 304 whereas spread items consisted in the repetition of a syllable containing a 305 lingual consonant followed by a spread vowel. 306

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¹Given that the activity of the *orbicularis oris inferior* and *orbicularis oris superior* muscles has previously been observed to be strongly correlated and that the activity of the OOI was more strongly affected by the experimental manipulation [59,60], we decided to record only the activity of the OOI in this study.

session.	
rounded items	spread items
/mumu/	/gigi/
/pupu/	/sese/
/fofo/	/lele/
/mymy/	/sisi
/pypy/	/didi/
/byby/	/nini/
/vøvø/	/ʒiʒi/*
/pøpø/	/lili/
/bøbø/	\RIRI\
/mɔ̃mɔ̃/	/aeae/

Table 1. List of bisyllabicnonwords used in the test

Note. *Because the production of the French palato-alveolar fricative in /ʒiʒi/ may involve a protrusion of the lips, this item theoretically slightly deviates from other items of this class.

Procedure

Participants were seated in front of a computer screen while audio stimuli 308 (when applicable) were presented through speakers on both sides of the 309 screen. A video camera was positioned on one side of the screen to monitor 310 facial movements. A microphone was positioned at approximately 10cm of 311 the participant to record possible speech sounds. After positioning of the 312 EMG sensors, each participant underwent a relaxation session aiming to 313 minimise pre-existing inter-individual variability on facial muscle contraction 314 (approximate duration was 330s). This relaxation session was recorded by a 315 trained professional sophrology therapist. Baseline EMG measurements were 316 performed during the last minute of this relaxation session, resulting in 60s 317 of EMG signal at baseline. By using this relaxation period as a baseline, we 318 made sure that participants were all in a comparable relaxed state. In 319 addition, several previous EMG studies have argued for the use of a 320 relaxation period as a baseline, since mere resting periods may include some 321 inner speech production (e.g., [69,70], for a review). 322

Subsequently, participants went through a training session, during which 323 they could get familiar with the main task. They trained with 8 stimuli in 324 total (4 rounded nonwords and 4 spread nonwords, cf. supplementary 325 materials). Each training stimulus appeared in three conditions (for all 326 participants): overt speech, inner speech and listening. Nonwords to be produced (covertly or overtly) were visually presented on the screen. Then, 328 a central fixation appeared on the screen, indicating to the participant that 329 s•he should utter the nonword (either overtly or covertly). This aimed to 330 ensure that participants were actually producing a nonword, not just simply 331 visually scanning it. In the overt speech condition, participants were asked 332 to produce the nonword "just after the word disappeared from the screen", 333

with "the most neutral intonation possible". In the inner speech condition, participants were asked to "innerly produce the nonword" (cf. the supplementary materials for precise instructions in French), with "the most neutral intonation possible" and while remaining as still as possible. In the listening condition, the order of these two screens was reversed. A fixation dot was first presented (for 1 second), followed by a blank screen (for 1 second). The audio stimulus was presented when the blank screen appeared, while participants were asked to remain as still as possible.

After the training, participants moved to the experimental part, that included a novel list of 20 nonwords (cf. Table 1). Each nonword was presented 6 times in each condition for each participant. The EMG activity was recorded during the entire experiment. The periods of interest consisted in one-second portions, after each stimulus presentation and during either production or listening. This resulted in 60 observations (60 periods of 1 second) for both classes of nonword in each test condition. The total duration of the experiment ranged between 30 min and 40 min.

EMG signal processing

EMG signal pre-processing was carried out using Matlab r2014a (Version 8.3.0.532, www.mathworks.fr). We first applied a 50Hz frequency comb filter to eliminate power noise. Then, we applied a 20 Hz – 450 Hz bandpass filter to the EMG signals, in order to focus on the 20–450 Hz frequency band, following current recommendations for facial EMG studies [71,72].

Although participants were explicitly asked to remain still during inner 356 speech production or listening, small facial movements (such as swallowing 357 movements) sometimes occurred. Such periods were excluded from the final 358 sample of EMG signals. To remove these signals, we first divided the portions 359 of signals of interest into periods of 1 second. The baseline condition was 360 therefore composed of 60 trials of 1-second. The periods of interest in all the 361 speech conditions consisted of the 1 second interval during which the 362 participants either produced speech or listened to speech. It is possible that 363 the nonword took less than 1 second to be produced, but since there was no 364 way to track when production started and ended in the inner speech 365 condition, the entire 1-second period was kept. Therefore, the overt speech 366 condition was composed of 6 repetitions of each nonword, that is 6x20 trials 367 of 1 second. The "inner speech" and "listening" conditions were similarly 368 composed of 6x20 trials of 1 second. Then, we visually inspected the EMG 369 signals recorded during each trial and listened to the audio signal 370 simultaneously recorded. In all conditions, any time a non-speech noise 371 (such as coughing or yawning) was audible in a trial, the trial was discarded 372 (i.e., we did not include this trial in the final analysis, for any of the recorded 373 muscles). In the listening and overt speech conditions, if a burst of EMG 374 activity was present after the relevant audio speech signal, then the trial was 375 discarded. In the overt speech condition, if the participant started too early 376 or too late and only part of the nonword was recorded in the audio signal, 377 then the corresponding trial was discarded. In the inner speech and listening 378 condition, if a large EMG burst of activity was present, potentially associated 379 with irrelevant non-speech activity, we excluded the trial. The fact that the 380 artefact rejection procedures slightly differ in the various conditions is not an 381 issue since we do not directly compare between conditions. Instead, we 382 compare the EMG correlates of the two classes of nonwords within each 383

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condition.

This inspection was realised independently by two judges. Subsequently, we only kept the trials that were not rejected by any of these two judges (i.e., we removed a trial as soon as it was rejected by at least one judge). The agreement rate between the two judges was of 87.82% (with a moderate Cohen's κ of approximately 0.48). The overall procedure led to an average (averaged over participants) rejection rate of 22.96% (SD = 6.49) in the baseline condition and 18.49% (SD = 6.48) in the other conditions.

After pre-processing and artefact rejection, we computed the by-trial 392 average amplitude of the centred and rectified EMG signal. This provided a 393 score for each muscle of interest (OOI, ZYG, FRO, COR, FCR) in each 394 condition (Baseline, Overt Speech, Inner Speech, Listening) and for each participant. Absolute EMG values are not meaningful as muscle activation is 396 never null, even in resting conditions, due in part to physiological noise. In 397 addition, there are inter-individual variations in the amount of EMG activity in 398 the baseline. To normalise and standardise for baseline activity across 399 participants, we thus expressed the EMG amplitude as a z-score from 400 baseline activity (i.e., we subtracted the mean amplitude of the centred and 401 rectified baseline signal and divided the result by the standard deviation of 402 the centred and rectified baseline signal), thereafter referred to as δ . 403

Data analysis

Statistical analyses were conducted using R version 3.5.3 [73], and are reported with the papaja [74] and knitr [75] packages. To assess the effects of the condition and the class of nonwords on the standardised EMG amplitude, we analysed these data using *Condition* (3 modalities: speech, inner speech, and listening) and *Class* of nonwords (2 modalities, rounded and spread, contrast-coded) as within-subject categorical predictors, and the standardised EMG amplitude as a dependent variable in a multivariate (i.e., with multiple outcomes) Bayesian multilevel linear model (BMLM). An introduction to Bayesian statistics is outside the scope of this paper. However, the interested reader is referred to [76] for an introduction to Bayesian multilevel modelling using the brms package.

In order to take into account the dependencies between repeated 416 observations by participant, we also included in this model a varying 417 intercept by participant. Contrary to what we pre-registered, we used a 418 multivariate model (instead of separate models by muscle). This allowed us 419 to estimate the correlation between each pair of muscles. Models were fitted 420 with the brms package [77] and using weakly informative priors (see the 421 supplementary materials for code details). Two Markov Chain Monte-Carlo 422 (MCMC) were run for each model to approximate the posterior distribution, 423 including each 5.000 iterations and a warmup of 2.000 iterations. Posterior convergence was assessed examining trace plots as well as the 425 Gelman-Rubin statistic R. Constant effect estimates were summarised via 426 their posterior mean and 95% credible interval (CrI), where a credible 427 interval can be considered as the Bayesian analogue of a classical 428 confidence interval. When applicable, we also report Bayes factors (BFs), 429 computed using the Savage-Dickey method, which consists in taking the 430 ratio of the posterior density at the point of interest divided by the prior 431 density at that point. These BFs can be interpreted as an updating factor, 432 from prior knowledge (what we knew before seeing the data) to posterior 433

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knowledge (what we know after seeing the data).

Results

The *Results* section is divided into two parts. First, we present results from 436 confirmatory (preregistered) analyses, aiming to test whether it is possible to 437 dissociate the activity of the OOI and the ZYG during inner speech 438 production, according to the content of inner speech (here, the class of 439 nonword). More precisely, we expected an increased EMG activity of the OOI 440 during the inner production of rounded nonwords in comparison to spread 441 nonwords. Conversely, we expected elevated EMG activity of the ZYG during 442 the inner production of spread nonwords in comparison to rounded nonwords. 443 Second, we present results from exploratory (non-preregistered) analyses. 444

To foreshadow the results, we did not observe such a clear dissociation between the EMG activity of the OOI and the ZYG muscles, neither in the inner speech condition nor in the overt speech condition. Contrary to theoretical expectations based on phonetics and speech production theory [78–81], the activity of both muscles was of higher amplitude for the pronunciation of rounded nonwords (as compared to spread nonwords) during overt speech production. Additionally, the EMG amplitude on both muscles of interest was similar during the inner production (or listening) of the two classes of nonwords. However, in the exploratory analyses section, we report results from supervised machine learning algorithms (classification using random forests), showing a reasonable accuracy to classify EMG signals according to the class of nonwords during overt speech production. This strategy was however unsuccessful for the inner speech and the listening conditions.

Before moving to the statistical results, we represent the distribution of the whole dataset, by class, by condition and by muscle for the two main muscles of interest (OOI and ZYG) in Fig 1. More precisely, the first row of this figure represents the distribution of the standardised EMG scores in the inner speech condition, the second row depicts the distribution of these scores in the listening condition, whereas the third row depicts the distribution of the standardised EMG scores in the overt speech condition. The first column depicts the distribution of the standardised EMG scores recorded over the OOI muscle whereas the second one represents the distribution of the standardised EMG scores recorded over the ZYG muscle. Each individual data point is represented as a vertical bar along the x-axis of each panel whereas the vertical coloured line represents the class-specific median. Additionally, a vertical dashed line is plotted at zero, which represents the baseline level. Thus, a positive value on the x-axis represents EMG standardised scores that are higher than baseline.

Fig 1. Distribution of standardised EMG scores by class and by muscle. The first row corresponds to the inner speech condition, the second one to the listening condition, and the third one to the overt speech condition. The first column depicts the EMG amplitude recorded over the OOI muscle while the second column represents the EMG amplitude recorded over the ZYG muscle. Each individual data point is represented as a vertical bar along the x-axis. The vertical coloured line represents the by-class median. Please note that the scale of the x-axis may differ considerably between panels.

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In Table 2, we report the mean standardised EMG amplitude of all recorded muscles in each condition. Given the skewness of the distribution of these scores, the mean and the standard deviation (SD) are not the best indicators of the central tendency and dispersion of these distributions. Therefore, we also report the median, the median absolute deviation (MAD), and the inter-quartile range (IQR).

Condition	Item	Muscle	Mean	Median	SD	MAD	IQR
innerspeech	rounded	COR	0.34	0.17	0.54	0.42	0.65
innerspeech	spread	COR	0.32	0.16	0.54	0.40	0.63
listening	rounded	COR	0.22	0.09	0.46	0.36	0.57
listening	spread	COR	0.24	0.11	0.46	0.37	0.58
speech	rounded	COR	0.29	0.17	0.52	0.40	0.56
speech	spread	COR	0.25	0.13	0.52	0.37	0.52
innerspeech	rounded	FCR	-0.01	-0.04	0.18	0.09	0.13
innerspeech	spread	FCR	0.00	-0.04	0.19	0.09	0.12
listening	rounded	FCR	-0.01	-0.04	0.15	0.09	0.12
listening	spread	FCR	0.00	-0.03	0.15	0.09	0.12
speech	rounded	FCR	-0.01	-0.03	0.14	0.08	0.11
speech	spread	FCR	-0.01	-0.03	0.13	0.09	0.12
innerspeech	rounded	FRO	0.63	0.47	0.68	0.66	0.95
innerspeech	spread	FRO	0.62	0.47	0.67	0.64	0.95
listening	rounded	FRO	0.59	0.40	0.67	0.57	0.82
listening	spread	FRO	0.60	0.41	0.67	0.58	0.85
speech	rounded	FRO	0.66	0.48	0.71	0.62	0.88
speech	spread	FRO	0.62	0.46	0.72	0.60	0.81
innerspeech	rounded	001	1.31	0.72	1.63	0.94	1.80
innerspeech	spread	001	1.24	0.64	1.67	0.85	1.61
listening	rounded	001	1.11	0.62	1.40	0.85	1.44
listening	spread	001	1.11	0.62	1.42	0.83	1.40
speech	rounded	001	12.01	10.99	6.44	6.22	8.75
speech	spread	001	11.71	8.88	9.13	6.56	10.25
innerspeech	rounded	ZYG	0.01	-0.04	0.17	0.11	0.17
innerspeech	spread	ZYG	0.00	-0.03	0.18	0.11	0.16
listening	rounded	ZYG	0.01	-0.02	0.17	0.12	0.17
listening	spread	ZYG	0.00	-0.02	0.16	0.12	0.18
speech	rounded	ZYG	1.57	1.33	1.10	0.92	1.29
speech	spread	ZYG	0.69	0.56	0.78	0.51	0.70

Table 2. Descriptive statistics of the standardised EMG amplitude foreach muscle in each condition.

We also created a shiny application [82] allowing for further visual exploration of the data by muscle, by condition, and by participant, in the 3D space formed by three (to be chosen) muscles. This application is available online (https://barelysignificant.shinyapps.io/3d_plotly/) and the associated code is available in the OSF repository (https://osf.io/czer4).

Confirmatory (preregistered) analyses

Bayesian multivariate multilevel Gaussian model

We then compared the standardised EMG amplitude δ for each muscle in each condition (Overt Speech, Inner Speech, Listening) by fitting a

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multivariate multilevel Gaussian model (as detailed previously in the Methods section). We predicted a higher increase of OOI activity during the production of rounded items in comparison to spread items and conversely, a higher increase of ZYG activity during the inner production of rounded items in comparison to spread items. These predictions should also apply to the overt speech condition (and to the listening condition). We should not observe any by-class differences of FRO and COR activity in any condition.

Response	Term	Estimate	SE	Lower	Upper	Rhat	BF01
001	Inner Speech	1.21	0.27	0.67	1.75	1.00	0.04
001	Listening	1.09	0.23	0.65	1.53	1.00	<0.001
001	Overt Speech	11.59	1.28	9.08	14.18	1.00	<0.001
001	Inner Speech x Class	0.07	0.14	-0.20	0.34	1.00	64.45
001	Listening x Class	-0.08	0.20	-0.47	0.32	1.00	47.05
001	Overt Speech x Class	0.02	0.19	-0.35	0.39	1.00	52.11
ZYG	Inner Speech	0.00	0.03	-0.05	0.06	1.00	379.5
ZYG	Listening	0.01	0.02	-0.04	0.05	1.00	388.4
ZYG	Overt Speech	1.15	0.15	0.86	1.43	1.00	<0.001
ZYG	Inner Speech x Class	0.01	0.02	-0.03	0.04	1.00	532.81
ZYG	Listening x Class	0.00	0.03	-0.06	0.05	1.00	389.12
ZYG	Overt Speech x Class	0.86	0.03	0.81	0.91	1.00	<0.001

Table 3. Estimates from the Gaussian BMLM concerning the OOI and the ZYG.

Note. For each muscle (response), the first three lines represent the estimated average amplitude in each condition, and its standard error (SE). The three subsequent rows represent the estimated average difference between the two classes of nonwords in each condition (i.e., the interaction effect). The 'Lower' and 'Upper' columns contain the lower and upper bounds of the 95% CrI, whereas the 'Rhat' column reports the Gelman-Rubin statistic. The last column reports the Bayes factor in favour of the null hypothesis (BF01).

The results of the Bayesian Gaussian multivariate model are summarised in Table 3. This table reports the estimated average EMG amplitude in each 497 condition and the corresponding BF. As they are not the main focus of 498 interest here and for the sake of clarity, descriptive results for the other two 499 facial muscles and for the forearm muscle are reported in the supplementary 500 materials. This analysis revealed that the EMG amplitude of the OOI was 501 higher than baseline (the standardised score was above zero) in every 502 condition whereas it was only the case in the overt speech condition for the 503 ZYG. Moreover, in all conditions, the EMG amplitude of the ZYG was lower 504 than that of the OOI. Crucially, we did not observe the hypothesised 505 difference according to the class of nonwords on the OOI during inner speech 506 production ($\beta = 0.071$, 95% CrI [-0.204, 0.342], BF₀₁ = 64.447) nor on the 507 ZYG (β = 0.005, 95% Crl [-0.031, 0.041], BF₀₁ = 532.811). 508

Fig 2 depicts these results by representing the distribution of the raw data 509 (coloured dots) along with the predictions from this model. The black dots 510 and vertical intervals represent the predicted mean and associated 95% 511 credible interval for each class of non-word, each condition and for the OOI 512 and the ZYG. Coherently with Table 3, this figure shows that the fitted model 513 predicts no noticeable differences between the two classes of non-words in 514 any condition for the OOI muscle. However, it predicts a higher average EMG 515 amplitude associated with the rounded item as compared to the spread 516

items in the overt speech condition for the ZYG muscle.

Fig 2. Raw data along with posterior predictions of the first model for the OOI and the ZYG muscles. Dots represent the mean prediction of this model by condition, whereas the vertical error bars represent the 95% credible intervals around the mean.

Before proceeding further with the interpretation of the results, it is 518 essential to check the quality of this first model. A useful diagnostic of the 519 model's predictive abilities is known as *posterior predictive checking* (PPC) 520 and consists in comparing observed data to data simulated from the 521 posterior distribution [83]. The idea behind PPC is that a good model should 522 be able to generate data that resemble the observed data [84]. In this vein, 523 Fig 3 represents the distribution of the whole dataset (across all participants 524 and conditions) by muscle (the dark blue line) along with the distribution of 525 hypothetical datasets generated from the posterior distribution of the model 526 (the light blue lines). As can be seen from this figure, the distributions of the 527 data generated from the model differ considerably from the distribution of 528 the observed data. Therefore, in the next section, we turn to a more 529 appropriate model for these data. 530

Fig 3. Posterior predictive checking for the first model concerning the OOI and ZYG muscles. The dark blue line represents the distribution of the raw data (across all conditions) whereas light blue lines are dataset generated from the posterior distribution.

Bayesian multivariate multilevel distributional Skew-Normal model 531

Fig 3 reveals an important failure of the first model, as it fails to generate 532 data that look like the data we have collected. More precisely, the collected 533 data look right-skewed, as it usually happens with physiological 534 measurements. To improve on the Gaussian model, we then assumed a 535 Skew-normal distribution for the response variable (the standardised EMG 536 amplitude δ). The Skew-normal distribution is a generalisation of the 537 Gaussian distribution with three parameters ξ (xi), ω (omega), and α (alpha) 538 for location, scale, and shape (skewness), respectively (note that the 539 Gaussian distribution can be considered a special case of the Skew-normal 540 distribution when $\alpha = 1$). In addition, we also improved the first model by 541 turning it into a *distributional model*, that is, a model in which we can specify 542 predictor terms for all parameters of the assumed response distribution [85]. 543 More precisely, we used this approach to predict both the location, the scale, 544 and the skewness of the Skew-Normal distribution (whereas the first model 545 only allowed predicting the mean of a Gaussian distribution). As can been 546 seen in Fig 4, this second model seems better than the first one at 547 generating data that fit the observed data. 548

Fig 4. Posterior predictive checking for the Skew-Normal model concerning the OOI and ZYG muscles. The dark blue line represents the distribution of the raw data whereas light blue lines are dataset generated from the posterior distribution.

Response	Term	Estimate	SE	Lower	Upper	Rhat	BF01
001	Inner Speech	1.47	0.03	1.41	1.53	1.00	0.04
001	Listening	1.24	0.02	1.19	1.29	1.00	< 0.001
001	Overt Speech	12.15	0.14	11.87	12.43	1.00	< 0.001
001	Inner Speech x Class	0.03	0.02	-0.01	0.06	1.00	64.45
001	Listening x Class	0.00	0.02	-0.05	0.05	1.00	47.05
001	Overt Speech x Class	1.42	0.18	1.05	1.78	1.00	52.11
ZYG	Inner Speech	0.02	0.00	0.01	0.02	1.00	379.5
ZYG	Listening	0.01	0.00	0.00	0.02	1.00	388.4
ZYG	Overt Speech	1.21	0.02	1.18	1.24	1.00	< 0.001
ZYG	Inner Speech x Class	0.00	0.01	-0.01	0.01	1.00	532.81
ZYG	Listening x Class	0.00	0.01	-0.02	0.02	1.00	389.12
ZYG	Overt Speech x Class	0.39	0.02	0.35	0.43	1.00	< 0.001

Table 4. Estimates from the distributional Skew-Normal model concerning the OOI and the ZYG.

Note. For each muscle (response), the first three lines represent the estimated average amplitude in each condition, and its standard error (SE). The three subsequent rows represent the estimated average difference between the two classes of nonwords in each condition (i.e., the interaction effect). The 'Lower' and 'Upper' columns contain the lower and upper bounds of the 95% Crl, whereas the 'Rhat' column reports the Gelman-Rubin statistic. The last column reports the Bayes factor in favour of the null hypothesis (BF01).

The estimates of this second model are summarised in Table 4 and Fig 5. 549 According to this model, the EMG amplitude of the OOI was higher than 550 baseline (the estimated standardised score was above zero) in every 551 condition whereas, for the ZYG, it was only the case in the overt speech 552 condition. We did not observe the hypothesised difference according to the 553 class of nonwords during inner speech production, neither on the OOI (β = 554 0.025, 95% Crl [-0.012, 0.064], BF₀₁ = 64.447) nor on the ZYG (β = 0.004, 555 95% Crl [-0.007, 0.014], BF₀₁ = 532.811). 556

Fig 5. Raw data along with posterior predictions of the third model for the OOI and the ZYG muscles. Dots represent the mean prediction of this model by condition (concerning the location parameter) whereas the vertical error bars represent the 95% credible intervals.

Predictions from this model are visually represented in Fig 5. This figure differs from Fig 2 (showing the predictions of the Gaussian model) in that the second model (the Skew-normal model) predicts shifts in location for both the OOI and the ZYG muscles according to the class of non-word in overt speech prediction. In contrast, the first model (the Gaussian model) predicted a by-class difference only for the ZYG muscle.

Exploratory (non-preregistered) analyses

In the previous section, we tried to predict the average EMG amplitude by condition on each single muscle. Although this approach was appropriate to tackle our initial research question (i.e., can we distinguish muscle-specific EMG correlates of inner speech production?), it is not optimal to answer more general questions such as "can we predict the content of inner speech

based on the available EMG data?". In Fig 6, we depict the distribution of the 569 by-word averaged EMG scores in the 2D space formed by the OOI and the 570 ZYG muscles. This figure reveals that although different nonwords produced 571 in overt speech seem difficult to discriminate on the basis of a single muscle 572 (cf. Fig 1), it seems easier to discriminate them in the space formed by two 573 muscles (here OOI and ZYG). More precisely, the two classes of nonwords 574 seem to form two separate clusters in the overt speech condition, but these 575 clusters do not seem discriminable in the inner speech or in the listening 576 condition. 577

Fig 6. Average standardised EMG amplitude for each nonword in each condition, in the 2D space formed by the OOI and the ZYG. Ellipses represent 95% data ellipses, that is, the 95% contours of a bivariate normal distribution.

In other words, it is easier to discriminate these signals in the 578 multidimensional space of all speech muscles, rather than by considering 579 each muscle independently. Thus, we used a supervised machine learning 580 algorithm aiming to classify speech signals according to the class of 581 nonwords. Broadly, the machine learning approach seeks to find a 582 relationship between an input X (e.g., EMG recordings over the four facial 583 muscles) and an output Y (e.g., the class of nonwords). Once trained, it 584 allows predicting a value of the output based on some input values, whose 585 prediction can be evaluated against new observations. 586

We used a random forest algorithm, as implemented in the caret package [86]. Random forests (RFs) represent an ensemble of many decision trees (a forest), which allow predictions to be made based on a series of decision rules (e.g., is the score on predictor x_1 higher or lower than z? If yes, then ..., if not, then ...). The specificity of RFs is to combine a large number of trees (usually above 100 trees), and to base the final conclusion on the average of these trees, thus preventing overfitting. We used three separate RFs to classify EMG signals in each condition (Overt Speech, Inner Speech, and Listening).

To evaluate the performance of this approach, we report the raw accuracy 596 (along with its resampling-based 95% confidence interval), or the proportion 597 of data points in the test dataset for which the RF algorithm predicted the 598 correct class of nonwords. First, we randomly split the entire dataset into a 599 training (80%) and a test set (20%). The training set was used for the 600 learning whereas the test set was used to evaluate the predictions of the 601 algorithm. To prevent overfitting, we used repeated 10-fold cross-validation 602 during the learning phase. 603

Predicting the class of nonwords during overt speech production

We first tried to predict the class of nonwords produced in overt speech, based on the activity of the four facial muscles (OOI, ZYG, COR, FRO). Each predictor was centred to its mean and standardised before the analysis.

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Table 5. Confusion matrix with by-class error for the overt speech condition.

Prediction	rounded	spread	class.error
rounded spread	917 198	163 898	0.151 0.181

This analysis revealed an overall classification accuracy of 0.847, 95% Cl608[0.814, 0.876] (cf. confusion matrix in Table 5). Examining the relative609importance of each feature (i.e., each muscle) for prediction revealed that610the muscles containing most information to discriminate the two classes of611nonwords were the ZYG and the OOI, whereas, as predicted, forehead612muscles did not seem to strongly contribute to predictive accuracy in the613overt speech condition.614

Predicting the class of nonwords during inner speech production and listening

We then applied the same strategy (the same algorithm) to the signals recorded in the inner speech and listening conditions. The results of these analyses are reported in Table 6 and Table 7.

condition.			•
	Refere	ence	
Prediction	rounded	spread	class.error
rounded spread	386 473	502 454	0.565 0.510

Table 6. Confusion matrix with by-classclassification error for the inner speechcondition.

This analysis revealed an overall classification accuracy of 0.472, 95% CI 620 [0.426, 0.52] in the inner speech condition, which indicates that the RF 621 algorithm did not allow discriminating the two classes of nonwords better 622 than random guessing. As the classification accuracy in the inner speech 623 and listening conditions was not better than chance, we do not report the 624 relative importance of the predictors. Indeed, it would be difficult to interpret 625 the importance of predictors for a classification task at which they do not 626 perform better than chance. 627

Table 7. Confusion matrix with by-class

 classification error for the listening condition.

Prediction	rounded	spread	class.error
rounded spread	426 508	499 406	0.539 0.556

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This analysis similarly revealed an overall classification accuracy of 0.46, 628 95% CI [0.413, 0.507] in the listening condition. 629

Discussion

In the present study we aimed to replicate and extend previous findings showing that facial electromyography can be used to discriminate expanded inner speech content [30,31]. As these studies used small samples of children, our study aimed to examine whether such results can be reproduced using surface electromyography and modern signal processing methods in an adult sample.

To this end, it was crucial to first show that the EMG correlates of our two 637 classes of nonwords were discriminable during overt speech production. 638 Surprisingly, the data we collected during overt speech production do not 639 corroborate the hypothesis according to which the average EMG amplitude 640 of the OOI should be higher during the production of "rounded" nonwords as 641 compared to "spread" nonwords. For both orofacial speech muscles (OOI 642 and ZYG), the average EMG amplitude was higher for rounded nonwords 643 than for spread nonwords during overt speech production. Moreover, while 644 the average EMG amplitude recorded over speech muscles was higher than 645 baseline in both the inner speech and listening conditions, we did not find 646 differences of activation according to the content of the material (the class 647 of nonword). An automatic classification approach, using the four facial 648 muscles (OOI, ZYG, COR, FRO), revealed that although it was possible to 649 discriminate EMG signals related to the two classes of nonwords with a 650 reasonable accuracy during overt speech production, this approach failed in 651 discriminating these two classes during inner speech production or during 652 listening. We also observed a higher EMG amplitude recorded over the facial 653 (both orofacial and non-orofacial) muscles during inner speech production 654 and during the listening of speech production than during rest. However, as 655 pinpointed by [62], this observation is not sufficient to conclude that these 656 activations were actually related to inner speech production, because i) both orofacial speech-related muscles and forehead non-speech related muscles 658 showed similar EMG amplitude changes from baseline and ii) we did not 659 observe different changes in EMG amplitude depending on the content of 660 inner speech (i.e., depending on the class of nonword to be uttered). 661

Before discussing the theoretical implications of these results, two main issues are worth discussing. First, how can we explain that rounded nonwords were associated with higher EMG amplitude during overt speech on both OOI and ZYG muscles? Second, how can we explain the indiscriminability of inner speech content, which seems to contradict classic as well as recent findings in the field [61]? We turn to each of these questions in the following.

To answer the first question, we began by comparing our results to results 669 obtained by another group [87]. The authors of this study recorded surface 670 EMG activity from five participants while they were producing seven facial 671 expressions and five isolated vowel sounds (/a/, /e/, /i/, /o/, /u/), repeated five 672 times each. They recorded EMG activity over eight facial muscles (the 673 zygomaticus major (ZYG), the risorius (RIS), the orbicularis oris superior 674 (OOS) and inferior (OOI), the mentalis (MEN), the depressor anguli oris (DAO), 675 the levator labii superioris (LLS) muscles, and the digastric muscle (DIG)). 676

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We divided these vowels in two classes to fit our own classes of nonwords. More precisely, we have created the following two classes: a *rounded* class, composed of the vowels /o/ and /u/, and a *spread* class, composed of the vowels /e/ and /i/ (note that we did not include the vowel /a/ because it theoretically does not fit in one of these two categories). We present the average EMG amplitude recorded over the OOI and the ZYG according to the vowel class in Table 8.

Table 8. Standardised EMG amplitude recorded over the OOI and the ZYG during overt speech production of rounded versus spread vowels in Eskes et al. (2017).

Muscle	Item	Observations	Mean	SD	Median	Histogram
00I 00I ZYG	rounded spread rounded	50 50 50	59.70 22.15 7.39	60.09 11.92 3.78	42.03 20.65 6.27	
ZYG	spread	50	10.15	6.20	7.99	

Note. The number of observations is given by the number of vowels to be pronunced in each category (2) times the number of repetitions (5) times the number of participants (5), for a total of 50 observations per cell.

We notice that [87] have indeed observed the dissociation we initially 684 predicted, that is, that the EMG amplitude recorded over the OOI was higher 685 during the pronunciation of rounded vowels than during pronunciation of 686 spread vowels, whereas the reverse pattern was observed concerning the 687 ZYG. Paired-samples Wilcoxon signed rank tests revealed a shift in location 688 (pseudomedian) between rounded and spread items for the OOI ($\beta = 24.12$, 689 95% CI [15.19, 40.77], V = 1184, p < .001) with rounded items being 690 associated with a higher location than spread items. This analysis also 691 revealed a shift in the inverse direction concerning the ZYG ($\beta = -1.51, 95\%$ 692 CI [-2.94, -0.48], V = 275, p < .001). However, one crucial difference between [87] design and ours is the complexity of the linguistic material. 694 Whereas [87] used single phonemes, we chose to use bisyllabic nonwords to 695 increase the ecological validity of the paradigm. Although these nonwords 696 were specifically created to theoretically increase the engagement of either 697 the OOI or the ZYG (cf. the "Linguistic material" section), it is reasonable to 698 expect differences in the average EMG patterns between isolated phonemes 699 and nonwords. More precisely, we expect the average EMG amplitude 700 associated with the production of a given phoneme (e.g., /y/) to be impacted 701 by the production of the consonant (e.g., /b/) it is paired with, due to 702 coarticulation. More generally, we could hypothesise that the difference 703 between the average EMG amplitude recorded during the production of the 704 phoneme i and during the production of the phoneme y could be reduced 705 when these phonemes are coarticulated in CV or CVCV sequences like /byby/ 706 or /didi/ (as in our study). In other words, we might expect an interaction 707 effect between the structure of the to-be produced speech sequence (either 708 a single vowel or a CV/CVCV sequence) and the class of the vowel. This is 709 coherent with previous findings showing that the muscular activity 710 associated with vowel production is strongly influenced by the surrounding 711 consonants in CVC sequences [78]. Thus, further investigations should focus 712 on how the average EMG amplitude is impacted by coarticulation during the 713 production of CVCV sequences.

With regards to inner speech, our results do not support theoretical 715 predictions of the *motor simulation view*, according to which it should be 716 possible to discriminate classes of nonwords produced in inner speech based 717 on EMG signals. Whereas this outcome is consistent with some recent 718 results [32], it also stands in sharp contrast with classical results in the field 719 [30,31] as well as more recent developments. For instance, [61] developed a 720 wearable device composed of seven surface EMG sensors that can attain a 721 92% median classification accuracy in discriminating internally vocalised 722 digits. There are a few crucial differences between [61]'s work and ours that 723 stand as good candidates to explain the discrepancies between our results. 724 First, the strategy adopted to position the sensors was radically different. Following guidelines from the field of psychophysiology, our strategy was to 726 position sensors precisely over the facial muscles of interest, aligned with the 727 direction of the muscle fibers and in theoretically optimal positions to record 728 activity of this muscle while reducing cross-talk. However, precisely because 729 of pervasive cross-talk in facial surface EMG recordings, this strategy, 730 whereas maximising the probability of recording activity from a given single 731 muscle, was also (as a result) reducing the probability of recording activity 732 from potentially speech-relevant neighbour muscles. Therefore, this strategy 733 might work sub-optimally when the goal of the experiment is to extract the 734 maximum amount of (relevant) EMG information to discriminate inner 735 speech content. However, this problem might be mitigated by using more sensors and a more lenient sensor-positioning approach. Whereas we 737 recorded the EMG amplitude over only two lower facial muscles (OOI and 738 ZIG), [61] analysed EMG data from seven different sensors, whose position 739 and number was defined iteratively in order to maximise the classification 740 accuracy. In other words, the parameters of the experiment were iteratively 741 optimised to maximise a certain outcome (classification accuracy). This 742 strategy is radically different from the classical approach in experimental 743 and cognitive psychology where experimental conditions are defined to test 744 theoretically derived hypotheses. Whereas the first approach is arguably 745 more efficient at solving a particular problem at hand, the second approach 746 might be more efficient in tackling theoretical questions. For instance, a 747 recent study reported a greater EMG amplitude of laryngeal and lip muscles 748 during auditory verbal tasks (covert singing) than during visual imagery 749 tasks [88]. By coupling EMG recording with demographic and psychological 750 measures, they were able to show that these correlates were related to the 751 level of accuracy in singing, thus shedding light upon the nature and 752 functions of peripheral muscular activity during covert singing. However, 753 adding more sensors (e.g., on the risorius), or better optimising sensor 754 placement, could improve the sensitivity of the present approach. 755

Putting aside considerations related to methodological aspects of the 756 present study, these results do not corroborate the motor simulation view of 757 inner speech production. Instead, it seems to support the abstraction view, 758 which postulates that inner speech results from the activation of abstract 759 linguistic representations and does not engage the articulatory apparatus. 760 However, individual differences in discriminability highlight that the 761 abstractness of inner speech might be flexible, as suggested by [22]. Indeed, 762 although for most participants it was not possible to decode the phonetic 763 content of inner speech, rounded and spread nonwords were in fact 764 distinguishable based on OOI and ZYG information only (by visual inspection 765

of the 2D plot), for two of them (S 15 and S 17, cf. supplementary materials). 766 This suggests either that the extent to which inner speech production 767 recruits the speech motor system might vary between individuals or that it 768 might vary within individual depending on the properties of the ongoing task 769 (these two suggestions are not mutually exclusive). For instance, we know 770 from early research on the EMG correlates of inner speech that the average 771 amplitude of these correlates tend to be higher when the task is more 772 difficult [10]. As such, the extent to which inner speech production recruits 773 the speech motor system could be moderated by manipulating the difficulty 774 of the ongoing task. In addition, the electromyographic activity recorded 775 during motor imagery could be modulated by the perspective taken in motor 776 imagery. A distinction is made between first-person perspective or internal 777 *imagery* (i.e., imagining an action as we would execute it) and third-person 778 perspective or external imagery (i.e., imagining an action as an observer of 779 this action), that may involve different neural processes [89]. It has been 780 shown that a first-person perspective may result in greater EMG activity than 781 motor imagery in a third-person perspective [90,91]. Therefore, we 782 hypothesise that the involvement of the speech motor system during inner 783 speech production may be modulated by the specific instructions given to 784 the participants. For instance, by instructing participants to focus on *inner* 785 speaking (imagining speaking), instead of inner hearing (imagining hearing), 786 and by asking them to focus on the kinaesthetic feelings related to speech 787 acts (rather than on auditory percepts), we could expect to find a higher average EMG amplitude recorded over the speech muscles. In addition, by 789 specifically asking the participants to mentally articulate the nonwords, as if 790 they were dictating them to someone, rather than just read and visually scan 791 them, we may expect stronger articulatory involvement. 792

Of course, the current study and the above discussion should be interpreted with a few words of caution in mind. For each class of nonwords, we collected around $6 \times 10 = 60$ observations by condition and by participant. For 25 participants and two classes of nonwords, this results in 25 (participants) x 120 (individual trials) x 3 (conditions) = 9000observations. However, after rejecting trials with movement artefacts, we had 7285 observations in total. Although the number of observations reported in the present study is reasonable, the sensitivity of the experiment could be improved by increasing the number of observations and/or by reducing two important sources of variation. More precisely, one could reduce the variance related to the item (the specific nonword being uttered) by selecting nonwords that are more similar to each other in the way they are uttered, by selecting less items or simpler items. Similarly, particular attention should be devoted to reducing inter-participant variability, which could be done by using more guided and specific instructions, as well as a longer training phase to familiarise the participant with the task.

In summary, we have demonstrated that whereas surface 809 electromyography may lead to reasonable accuracy in discriminating classes 810 of nonwords during overt speech production (using signals recorded over 811 only two speech-related muscles), it did not permit to discriminate these two 812 classes during inner speech production across all participants (only for two 813 participants). These results, in comparison with results obtained by other 814 teams [61], highlight that depending on the aim of the research, different 815 strategies might be more or less successfully pursued. More precisely, if the 816 goal is to attain high classification accuracy (problem-solving approach), 817

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then the parameters of the experiment (e.g., number of repetitions, number 818 of sensors, position of the sensors, parameters of the signal processing 819 workflow) should be optimised based on the desired outcome (i.e., 820 classification accuracy). However, the classical laboratory strategy used in 821 experimental and cognitive psychology, aiming to compare specific 822 conditions (or muscles) to each other in a controlled environment, is deemed 823 to be more appropriate when the aim of the research is to sharpen our 824 understanding of the psychological phenomenon under study. 825

Supplementary materials

Pre-registered protocol, open data, supplementary materials, as well as reproducible code and figures are available at https://osf.io/czer4.

Aside from previously cited packages, several other packages have been used for the writing of this paper, among which the ggrepel and ggplot2 packages for plotting [92,93] as well as the tidyverse, sjstats, here, skimr, and glue packages for code writing and formatting [94–98].

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References

1. Filik R, Barber E. Inner Speech during Silent Reading Reflects the Reader's Regional Accent. PLOS ONE. Public Library of Science; 2011;6: e25782. doi:10.1371/journal.pone.0025782

2. Lœvenbruck H, Baciu M, Segebarth C, Abry C. The left inferior frontal gyrus under focus: An fMRI study of the production of deixis via syntactic extraction and prosodic focus. Journal of Neurolinguistics. 2005;18: 237–258. doi:10.1016/j.jneuroling.2004.12.002

3. Perrone-Bertolotti M, Kujala J, Vidal JR, Hamame CM, Ossandon T, Bertrand O, et al. How Silent Is Silent Reading? Intracerebral Evidence for Top-Down Activation of Temporal Voice Areas during Reading. J Neurosci. Society for Neuroscience; 2012;32: 17554–17562. doi:10.1523/INEUROSCI.2982-12.2012

4. Yao B, Belin P, Scheepers C. Silent reading of direct versus indirect speech activates voice-selective areas in the auditory cortex. J Cogn Neurosci. 2011;23: 3146-3152. doi:10.1162/jocn_a_00022

5. Brouwers VP, Heavey CL, Lapping-Carr L, Moynihan S, Kelsey J, Hurlburt RT. Pristine Inner Experience: While Silent Reading It's Not Silent Speaking of the Text. Journal of Consciousness Studies. 2018;25: 29–54.

6. Hurlburt RT. On investigating inner experience: Contrasting Moore & Schwitzgebel and Brouwers et al. Consciousness and Cognition. 2018;63: 146–150. doi:10.1016/j.concog.2018.06.018

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852

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856

857

7. Frith U. Reading by Eye and Writing by Ear. In: Kolers PA, Wrolstad ME, 859 Bouma H, editors. Processing of Visible Language. Boston, MA: Springer US; 860 1979. pp. 379-390. doi:10.1007/978-1-4684-0994-9 23 861

8. D'Argembeau A, Renaud O, Linden MV der. Frequency, characteristics and functions of future-oriented thoughts in daily life. Applied Cognitive Psychology. 2011;25: 96-103. doi:10.1002/acp.1647

Baldo JV, Dronkers NF, Wilkins D, Ludy C, Raskin P, Kim J. Is problem solving dependent on language? Brain Lang. 2005;92: 240-250. doi:10.1016/j.bandl.2004.06.103

10. Sokolov A. Inner speech and thought. New York: Springer-Verlag; 1972.

11. Alderson-Day B, Fernyhough C. Inner speech: Development, cognitive 870 functions, phenomenology, and neurobiology. Psychological Bulletin. 871 2015;141: 931-965. doi:10.1037/bul0000021 872

12. Lœvenbruck H. What the neurocognitive study of inner language reveals about our inner space. In: Smadja S, Patoine P-L, editors. Épistémocritique, n° 18 : Langage intérieur - Espaces intérieurs / Inner Speech - Inner Space. 2018. Available: http://epistemocritique.org/ what-the-neurocognitive-study-of-inner-language-reveals-about-our-inner-space.

13. Morin A, Uttl B, Hamper B. Self-Reported Frequency, Content, and 878 Functions of Inner Speech. Procedia - Social and Behavioral Sciences. 879 2011;30: 1714-1718. doi:10.1016/j.sbspro.2011.10.331 880

14. Perrone-Bertolotti M, Rapin L, Lachaux JP, Baciu M, Lœvenbruck H. What is that little voice inside my head? Inner speech phenomenology, its role in cognitive performance, and its relation to self-monitoring. Behavioural brain research. Elsevier B.V. 2014;261: 220-39. doi:10.1016/j.bbr.2013.12.034

15. Hurlburt RT, Heavey CL. Investigating pristine inner experience: Implications for experience sampling and guestionnaires. Consciousness and Cognition. 2015;31: 148-159. doi:10.1016/j.concog.2014.11.002

888 16. McCarthy-Jones S, Fernyhough C. The varieties of inner speech: Links 889 between guality of inner speech and psychopathological variables in a 890 sample of young adults. Consciousness and cognition. Elsevier Inc. 2011;20: 891 1586-93. doi:10.1016/j.concog.2011.08.005 892

17. Lœvenbruck H, Grandchamp R, Rapin L, Nalborczyk L, Dohen M, Perrier P, et al. A cognitive neuroscience view of inner language: To predict and to hear, see, feel. In: Langland-Hassan P, Vicente A, editors. Inner speech: New voices. Oxford University Press; 2018. p. 37.

18. Oppenheim GM, Dell GS. Inner speech slips exhibit lexical bias, but not the phonemic similarity effect. Cognition. 2008;106: 528-537. doi:10.1016/j.cognition.2007.02.006

19. Vygotsky LS. Thought and language, revised and expanded edition. The MIT Press; 1934–2012.

20. Watson JB. Psychology from the standpoint of a behaviorist.

Philadelphia, PA, US: J B Lippincott Company; 1919. doi:10.1037/10016-000 MacKay DG. Constraints on theories of inner speech. In: Reisberg D,

editor. Auditory imagery. Erlbaum; Hillsdale, N. J. 1992. pp. 121-149. 22. Oppenheim GM, Dell GS. Motor movement matters: The flexible abstractness of inner speech. Memory & Cognition. 2010;38: 1147–1160. doi:10.3758/MC.38.8.1147

23. Korba RJ. The rate of inner speech. Percept Mot Skills. 1990;71: 1043-1052. doi:10.2466/pms.1990.71.3.1043

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863

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885

886

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893

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897

898

899

900

901

902

903

904

905

906

907

908

909

24. Netsell R, Kleinsasser S, Daniel T. The rate of expanded inner speech
during spontaneous sentence productions. Percept Mot Skills. 2016;123:
912
383-393. doi:10.1177/0031512516664992911

25. Corley M, Brocklehurst PH, Moat HS. Error biases in inner and overt speech: Evidence from tongue twisters. Journal of Experimental Psychology: Learning, Memory, and Cognition. 2011;37: 162–175. doi:10.1037/a0021321

26. Jones SR, Fernyhough C. Thought as action: Inner speech, self-monitoring, and auditory verbal hallucinations. Consciousness and cognition. 2007;16: 391–9. doi:10.1016/j.concog.2005.12.003

27. Martínez-Manrique F, Vicente A. The activity view of inner speech. Frontiers in psychology. 2015;6: 232. doi:10.3389/fpsyg.2015.00232

28. Grandchamp R, Rapin L, Perrone-Bertolotti M, Pichat C, Haldin C, Cousin E, et al. The ConDialInt Model: Condensation, Dialogality, and Intentionality Dimensions of Inner Speech Within a Hierarchical Predictive Control Framework. Front Psychol. Frontiers; 2019;10. doi:10.3389/fpsyg.2019.02019

29. Fernyhough C. Alien voices and inner dialogue: Towards a developmental account of auditory verbal hallucinations. New Ideas in Psychology. 2004;22: 49–68. doi:10.1016/j.newideapsych.2004.09.001

30. McGuigan FJ, Dollins AD. Patterns of covert speech behavior and phonetic coding. The Pavlovian Journal of Biological Science. 1989;24: 19–26.

31. McGuigan FJ, Winstead CL. Discriminative relationship between covert oral behavior and the phonemic system in internal information processing. Journal of Experimental Psychology. 1974;103: 885–890. doi:10.1037/h0037379

32. Meltzner GS, Sroka J, Heaton JT, Gilmore LD, Colby G, Roy S, et al. Speech recognition for vocalized and subvocal modes of production using surface EMG signals from the neck and face. INTERSPEECH. Brisbane, Australia; 2008. pp. 2667–2670.

33. Simonyan K, Horwitz B. Laryngeal motor cortex and control of speech in humans. The Neuroscientist. 2011;17: 197–208. doi:10.1177/1073858410386727

34. Jeannerod M. Motor cognition: What actions tell the self. Oxford ; New York: Oxford University Press; 2006.

35. Haggard P. Conscious intention and motor cognition. Trends in Cognitive Sciences. 2005;9: 290–295. doi:10.1016/j.tics.2005.04.012

36. Jeannerod M. The representing brain: Neural correlates of motor intention and imagery. Behavioral and Brain Sciences. 1994;17: 187. doi:10.1017/S0140525X00034026

37. Jeannerod M. Neural simulation of action: A unifying mechanism for motor cognition. NeuroImage. 2001;14: S103–S109. doi:10.1006/nimg.2001.0832

38. Barsalou LW. Grounded cognition. Annual review of psychology. 2008;59: 617-645. doi:10.1146/annurev.psych.59.103006.093639

39. O'Shea H, Moran A. Does motor simulation theory explain the cognitive mechanisms underlying motor imagery? A critical review. Frontiers in Human Neuroscience. 2017;11. doi:10.3389/fnhum.2017.00072

40. Pickering MJ, Garrod S. An integrated theory of language production and comprehension. Behavioral and Brain Sciences. 2013;36: 329–347. doi:10.1017/S0140525X12001495

41. Gentsch A, Weber A, Synofzik M, Vosgerau G, Schütz-Bosbach S.

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954

955

956

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958

959

960

961

Towards	a common framework of grounded action cognition: Relating motor	963
control,	perception and cognition. Cognition. 2016;146: 81–89.	964
doi:10.1	016/j.cognition.2015.09.010	965
42. K	awato M, Furukawa K, Suzuki R. A hierarchical neural-network model	966
for cont	rol and learning of voluntary movement. Biological Cybernetics.	967
1987;57	: 169–185. doi:10.1007/BF00364149	968
43. Jo	ordan MI, Rumelhart DE. Forward models: Supervised learning with a	969
distal te	acher. Cognitive Science. 1992;16: 307–354.	970
doi:10.1	207/s15516709cog1603_1	971
44. K	awato M. Internal models for motor control and trajectory planning.	972
Current	Opinion in Neurobiology. 1999;9: 718–727.	973
doi:10.1	016/S0959-4388(99)00028-8	974
45. V	Volpert D, Ghahramani Z, Jordan MI. An internal model for	975
sensorir	notor integration. Science. 1995;269: 1880–1882.	976
doi:10.1	126/science. /569931	977
46. 0	rush R. The emulation theory of representation: Motor control,	978
imagery	, and perception. Behavioral and Brain Sciences. 2004;27.	979
doi:10.1	017/50140525X04000093	980
47. N	Ioulton ST, Kosslyn SM. Imagining predictions: Mental imagery as	981
mental	emulation. Philosophical Transactions of the Royal Society B:	982
Biologic	al Sciences. 2009;364: 1273-1280. doi:10.1098/rstb.2008.0314	983
48. F	oude JF, Nagarajan SS. Speech production as state feedback control.	984
Frontiers	s in Human Neuroscience. 2011;5. doi:10.3389/fnhum.2011.00082	985
. 49. 6	uillot A, Di Rienzo F, MacIntyre T, Moran A, Collet C. Imagining is not	986
doing bu	It involves specific motor commands: A review of experimental data	987
	to motor inhibition. Frontiers in Human Neuroscience. 2012;6.	988
doi:10.3	389/fnnum.2012.00247	989
50. G	Sufficient A, Collet C. Contribution from neurophysiological and	990
psychol	Digical methods to the study of motor imagery. Brain Research	991
Reviews	. 2005;50: 387-397. doi:10.1016/j.brainresiev.2005.09.004	992
51. G	uniol A, Lebon F, Collet C. Electromyographic activity during motor	993
inagery of mont	al and motor imagony. Oxford University Pross, 2010, pp. 92-04	994
or ment	al anu motor imagery. Oxford Oniversity Press; 2010. pp. 65-94.	995
	195/dcp101:050/9760199540251.005.0000	996
52. 5 A. Collet	C editors. The neurophysiological foundations of montal and motor	997
A, Collet	C, editors. The neurophysiological foundations of mental and motor	998
inagery 52 p	appet M. Decety L. Jeanpared M. Pequin L. Montal simulation of an	999
action m	onnet M, Decety J, Jeannerou M, Requiri J. Mental Simulation of an	1000
Brain Re	bearch 1007.5, 221_228 doi:10.1016/S0026-6/10/06/00072-0	1001
54 F	adiga Buccino G. Craighero I. Fogassi I. Gallese V. Pavesi G	1002
Corticos	ninal excitability is specifically modulated by motor imagery a	1003
magnet	c stimulation study. Neuropsychologia 1999:37: 147-158	1004
doi:10.1	016/c0028.3032/08/00089.x	1005
55 B	ossini PM. Corticospinal excitability modulation to hand muscles	1000
during r	novement imagery. Cerebral Cortex, 1999.9. 161–167	1007
doi:10.1	093/cercor/9 2 161	1000
56.1	ivesay I. Liebke A. Samaras M. Stanley A. Covert speech behavior	1010
during a	silent language recitation task. Percentual and Motor Skills	1010
1996.83	: 1355-1362. doi:10.2466/pms.1996.83.3f1355	1012
57 I	ocke IL. Subvocal speech and speech. ASHA, 1970:12: 7–14	1012
58. L	ocke IL, Fehr FS. Subvocal rehearsal as a form of speech. Journal of	1014
501 2	,	

Verbal Learning and Verbal Behavior. 1970;9: 495–498.	1015
doi:10.1016/s0022-5371(70)80092-5	1016
59. Rapin L, Donen M, Polosan M, Perrier P, Lœvenbruck H. An EMG study	1017
of the lip muscles during covert auditory verbal hallucinations in	1018
Schizophrenia. Journal of Speech, language, and hearing research. 2013;50:	1019
1882-1995. 001:10.1044/1092-4588(2015/12-0210)	1020
00. Nalborczyk L, Perione-Bertololli M, Baeyens C, Grandchamp K,	1021
induced verbal rumination. Dielegical Devebalagy, 2017;127; 52, 62	1022
dai:10.1016/i biopovcho.2017.04.012	1023
Gl. Kapur A. Kapur S. Mass D. Alter Free A. personalized wearable silent	1024
or South Stranger of the 2018 Conference on Human	1025
Information Interaction & Poteculary January ACM Process 2018	1026
nii of nation interaction are ineval - 101 10. 10kyo, japan. ACM FIESS, 2010.	1027
62 Carrity II Electromyography: A roview of the current status of	1028
cubyocal space research. Momory & Cognition 1077:5: 615-622	1029
	1030
63 Simmons Nelson Simonsohn A 21 word solution. The Official	1031
Newsletter of the Society for Personality and Social Psychology 2012:26	1032
doi-10 2139/ssrn 2160588	1033
64 Schönbrodt ED Wagenmakers E-L Bayes factor design analysis	1034
Planning for compelling evidence. Psychonomic Bulletin & Beview 2018:25:	1035
128–142. doi:10.3758/s13423-017-1230-v	1030
65. Schönbrodt FD. Wagenmakers E-I. Zehetleitner M. Perugini M.	1038
Sequential hypothesis testing with Bayes factors: Efficiently testing mean	1039
differences. Psychological Methods. 2017:22: 322–339.	1040
doi:10.1037/met0000061	1041
66. Beffara B, Bret A, Nalborczyk L. A fully automated, transparent,	1042
reproducible and blind protocol for sequential analyses. PsyArXiv. 2019;	1043
doi:10.17605/OSF.IO/MWTVK	1044
67. Everdell IT, Marsh H, Yurick MD, Munhall KG, Paré M. Gaze behaviour	1045
in audiovisual speech perception: Asymmetrical distribution of face-directed	1046
fixations. Perception. 2007;36: 1535-1545. doi:10.1068/p5852	1047
68. Mathôt S, Schreij D, Theeuwes J. OpenSesame: An open-source,	1048
graphical experiment builder for the social sciences. Behavior Research	1049
Methods. 2012;44: 314-324. doi:10.3758/s13428-011-0168-7	1050
69. Jacobson E. Electrical measurements of neuromuscular states during	1051
mental activities. VII. Imagination, recollection, and abstract thinking	1052
involving the speech musculature. American Journal of Physiology. 1931;897:	1053
200–209.	1054
70. Vanderwolf CH. Brain, behavior, and mind: What do we know and	1055
what can we know? Neurosci Biobehav Rev. 1998;22: 125-142.	1056
doi:10.1016/s0149-7634(97)00009-2	1057
71. Boxtel A. Optimal signal bandwidth for the recording of surface EMG	1058
activity of facial, jaw, oral, and neck muscles. Psychophysiology. 2001;38:	1059
22-34. doi:10.1111/1469-8986.3810022	1060
72. De Luca CJ. The use of surface electromyography in biomechanics.	1061
Journal of Applied Biomechanics. 1997;13: 135–163.	1062
doi:10.1123/jab.13.2.135	1063
73. R Core leam. R: A language and environment for statistical	1064
computing [Internet]. Vienna, Austria: R Foundation for Statistical	1065
Computing; 2018. AVailable: https://www.R-project.org/	1066

74. Aust F, Barth M. papaja: Create APA manuscripts with R Markdown 1067 [Internet]. 2018. Available: https://github.com/crsh/papaja 1068 75. Xie Y. Dynamic documents with R and knitr [Internet]. 2nd ed. Boca 1069 Raton, Florida: Chapman; Hall/CRC; 2015. Available: 1070 https://yihui.name/knitr/ 1071 76. Nalborczyk L, Batailler C, Lœvenbruck H, Vilain A, Bürkner P-C. An 1072 introduction to Bayesian multilevel models using brms: A case study of 1073 gender effects on vowel variability in standard indonesian. Journal of Speech 1074 Language and Hearing Research. 2019;62: 1225–1242. 1075 doi:10.1044/2018 JSLHR-S-18-0006 1076 77. Bürkner P-C. brms: An R package for Bayesian multilevel models 1077 using Stan. Journal of Statistical Software. 2017;80: 1–28. 1078 doi:10.18637/jss.v080.i01 1079 78. Fromkin VA. Neuro-muscular specification of linguistic units. Language 1080 and Speech. 1966;9: 170-199. doi:10.1177/002383096600900304 1081 79. Ladefoged P, Johnson K. A Course in Phonetics. Thomson Wadsworth. 1082 2006. 1083 80. Lieberman P, Blumstein SE. Speech Physiology, Speech Perception, 1084 and Acoustic Phonetics [Internet]. Cambridge: Cambridge University Press; 1085 1988. doi:10.1017/CBO9781139165952 1086 81. Zemlin WR. Speech and Hearing Science: Anatomy and Physiology: 1087 United States Edition. Boston: Pearson; 1968. 1088 82. Chang W, Cheng J, Allaire J, Xie Y, McPherson J. shiny: Web application 1089 framework for r [Internet]. 2018. Available: 1090 https://CRAN.R-project.org/package=shiny 1091 83. Gelman A, Carlin JB, Stern H, Dunson DB, Vehtari A, Rubin DB. 1092 Bayesian data analysis, third edition. CRC Press, Taylor & Francis Group; 1093 2013. 1094 84. Gabry J, Simpson D, Vehtari A, Betancourt M, Gelman A. Visualization 1095 in Bayesian workflow. Journal of the Royal Statistical Society: Series A 1096 (Statistics in Society). 2019;182: 389–402. doi:10.1111/rssa.12378 1097 85. Bürkner P-C. Advanced Bayesian multilevel modeling with the R 1098 package brms. The R Journal. 2018;10: 395-411. 1099 86. Jed Wing MKC from, Weston S, Williams A, Keefer C, Engelhardt A, 1100 Cooper T, et al. caret: Classification and regression training [Internet]. 2018. 1101 Available: https://CRAN.R-project.org/package=caret 1102 87. Eskes M, van Alphen MJA, Balm AJM, Smeele LE, Brandsma D, van der 1103 Heijden F. Predicting 3D lip shapes using facial surface EMG. Zhang Y, editor. 1104 PLOS ONE. 2017;12: e0175025. doi:10.1371/journal.pone.0175025 1105 88. Pruitt TA, Halpern AR, Pfordresher PQ. Covert singing in anticipatory 1106 auditory imagery. Psychophysiology. 2018; e13297. doi:10.1111/psyp.13297 1107 89. Ruby P, Decety J. Effect of subjective perspective taking during 1108 simulation of action: A PET investigation of agency. Nat Neurosci. 2001;4: 1109 546-550. doi:10.1038/87510 1110 90. Hale BD. The effects of internal and external imagery on muscular 1111 and ocular concomitants. Journal of Sport Psychology. 1982;4: 379-387. 1112 doi:10.1123/isp.4.4.379 1113 Harris DV, Robinson WJ. The effects of skill level on EMG activity 1114 during internal and external imagery. Journal of Sport Psychology. 1986;8: 1115 105-111. doi:10.1123/jsp.8.2.105 1116 92. Slowikowski K. ggrepel: Automatically position non-overlapping text 1117 labels with 'ggplot2' [Internet]. 2018. Available: 1118

https://CRAN.R-project.org/package=ggrepel	1119
Wickham H. ggplot2: Elegant graphics for data analysis [Internet].	1120
Springer-Verlag New York; 2016. Available: http://ggplot2.org	1121
94. Wickham H. tidyverse: Easily install and load the 'tidyverse' [Internet].	1122
2017. Available: https://CRAN.R-project.org/package=tidyverse	1123
95. Lüdecke D. sjstats: Statistical functions for regression models (version	1124
0.17.1) [Internet]. 2018. doi:10.5281/zenodo.1284472	1125
96. Hester J. glue: Interpreted string literals [Internet]. 2018. Available:	1126
https://CRAN.R-project.org/package=glue	1127
97. Müller K. here: A simpler way to find your files [Internet]. 2017.	1128
Available: https://CRAN.R-project.org/package=here	1129
98. McNamara A, Arino de la Rubia E, Zhu H, Ellis S, Quinn M. Skimr:	1130
Compact and flexible summaries of data [Internet]. 2018. Available:	1131
https://CRAN.R-project.org/package=skimr	1132