Tracking components of bilingual language control in speech production: an fMRI study using functional localizers.

Abbreviated title: **Tracking components of bilingual language control in speech production**

Agata Wolna[1], Jakub Szewczyk[2], Michele Diaz[3], Aleksandra Domagalik[4],

Marcin Szwed[1], Zofia Wodniecka[1]

[1] Institute of Psychology, Jagiellonian University, Kraków, Poland

[2] Donders Institute for Brain Cognition and Behaviour, Radboud University, Nijmegen, The Netherlands

[3] Social, Life, and Engineering Sciences Imaging Center, the Pennsylvania State University, Pennsylvania, United States

[4] Centre for Brain Research, Jagiellonian University, Kraków, Poland

Corresponding authors:

zofia.wodniecka@uj.edu.pl

agata.wolna@doctoral.uj.edu.pl

Language and Bilingualism Lab

Instytut Psychologii UJ

ul. Ingardena 6

30-060 Kraków

Acknowledgements

The authors would like to thank all members of our Psychology of Language and Bilingualism Laboratory LangUsta who contributed to the research project by collecting and coding the data, all participants who took part in the study, as well as Mike Timberlake who proofread the text. We are very grateful to Evelina Fedorenko for inspiration and insightful comments on the design and results.

Funding

This research was possible thanks to an OPUS grant from the National Science Center Poland, awarded to Zofia Wodniecka (2017/27/B/HS6/00959).

Declaration of Competing Interests

Authors declare no competing interests.

Abstract

When bilingual speakers switch back to speaking in their native language (L1) after having used their second language (L2), they often experience difficulty in retrieving words in their L1: this phenomenon is referred to as the *L2 after-effect*. We used the L2 after-effect as a lens to explore the neural bases of bilingual language control mechanisms. Our goal was twofold: first, to explore whether bilingual language control draws on domain-general or language-specific mechanisms; second, to investigate the precise mechanism(s) that drive the L2 after-effect. We used a precision fMRI approach based on *functional localizers* to measure the extent to which the brain activity that reflects the L2 after-effect overlaps with the language network (Fedorenko et al., 2010) and the domain-general Multiple Demand network (Duncan et al., 2010), as well as three task-specific networks that tap into interference resolution, lexical retrieval, and articulation. Forty-two Polish-English bilinguals participated in the study. Our results show that the L2 after-effect reflects increased engagement of domain-general but not language-specific resources. Furthermore, contrary to previously proposed interpretations, we did not find evidence that the effect reflects increased difficulty related to lexical access, articulation, and the resolution of lexical interference. We propose that difficulty of speech production in the picture naming paradigm – manifested as the L2 after-effect – reflects interference at a non-linguistic level of task schemas or a general increase of cognitive control engagement during speech production in L1 after L2.

Keywords: fMRI, language production, bilingualism, language control, cognitive control

Introduction

In their daily lives, bilinguals must face a constant challenge related to the non-selectivity or constant co-activation of their two languages (Costa et al., 1999; Kroll et al., 2006). To cope with this problem, bilinguals need an efficient system of control that resolves interference between the competing languages (Green, 1998), or a selection mechanism (Blanco-Elorrieta & Caramazza, 2021) that allows them to efficiently use the intended language. In this study we aimed to characterize the neurocognitive mechanisms of control that are engaged in bilingual speech production by capitalizing on the advantages of the functional localizer approach. To this aim, we explored how the control mechanisms affect bilingual speech production by looking at the *L2 after-effect*: a difficulty in native language (L1) production observed after prior use of a second language (L2).

Bilingual language control: domain-general or domain-specific?

One of the most fundamental questions regarding the complex system that bilinguals use to control their two languages is whether it is based on a set of mechanisms specifically tailored to handle language (so called domain-specific, or language-specific mechanisms), or whether bilingual language control is achieved by domain-general cognitive control mechanisms (Abutalebi & Green, 2013; Wu et al., 2019; for a review see: Calabria et al., 2018).

So far, most evidence for the engagement of domain-general control mechanisms in bilingual language production comes from behavioral studies in which bilinguals were required to switch between their languages. The overall conclusion from these experiments is that switching between two languages incurs additional processing costs (Costa & Santesteban, 2004; Gollan & Ferreira, 2009; Meuter & Allport, 1999; Verhoef et al., 2009; for review see: Bobb & Wodniecka, 2013; Declerck et al., 2020; Declerck & Koch, 2022; Declerck & Philipp, 2015). This additional cognitive effort has also been observed as increased neural activation in structures associated with broadly defined cognitive control mechanisms (e.g., Abutalebi & Green, 2007; Coderre et al., 2016; De Baene et al., 2015; Wu et al., 2019; for reviews see: Abutalebi & Green, 2016; Luk et al., 2012; Sulpizio et al., 2020). It has been argued that bilingual language control is supported by domain-general mechanisms because task switching (which recruits domain-general control) and language switching share a number of computational processes, e.g., context monitoring, reconfiguring task-sets, engaging and disengaging from a task, suppressing the interference between

competing representations and selective response inhibition (Green, 1998; Green &
Abutalebi, 2013; Wu et al., 2019). This is also supported by research showing that the brain
networks that support language switching and task switching overlap (Blanco-Elorrieta &
Pylkkänen, 2016; Coderre et al., 2016; De Baene et al., 2015; Luk et al., 2012; Mendoza et
al., 2021; Nee et al., 2007; Niendam et al., 2012; Weissberger et al., 2015; Wu et al., 2019).

On the other hand, the overlap between brain networks activated by language- and
task-switching may not provide conclusive evidence for the domain-general nature of
bilingual language control. First, language switching also engages brain structures that are not
implicated in non-linguistic switching tasks (De Baene et al., 2015; Mendoza et al., 2021;
Weissberger et al., 2015). Second, the overlap between the brain networks recruited by
language- and task-switching may result from idiosyncratic properties of the switching task
itself rather than bilingual language control. More specifically, it can be argued that language
switching is an "artificial" task in the sense that it requires changing languages very
frequently and depending on external cues, while switching in natural contexts is largely
dictated by the speaker (i.e., internal cues; for a discussion on how cues affect switch costs,
see Blanco-Elorrieta & Pylkkänen, 2017 for natural vs. artificial cues and Gollan et al., 2014
and Gollan & Ferreira, 2009 for voluntary vs. forced cues). As such, a language switching
task may require the use of a very specific task set and keeping both languages co-activated.
Such extra demands may recruit mechanisms engaged in performing tasks that are untrained,
difficult, or not automated (Chein & Schneider, 2005; Crittenden & Duncan, 2014;
Shashidhara et al., 2019), and these types of tasks are known to reliably activate the domain-
general Multiple Demand network. What is more, bilinguals rarely switch between their
languages on a word-by-word basis (Wodniecka et al., 2020; see also: Bobb & Wodniecka,
2013); and forced switching is related to greater engagement of cognitive control than
voluntary switching (*for behavioral evidence see:* Gollan et al., 2014; Gollan & Ferreira,
2009; *for fMRI evidence see:* Reverberi et al., 2018; Zhang et al., 2015).

Digging deeper – specific processes involved in bilingual language control
Besides the more general question of whether bilinguals rely on domain-general or
language-specific networks when using their two languages, another unresolved issue is what
specific processes are involved in bilingual language control. Models of bilingual language
production assume that each language of a bilingual speaker is subserved by a set of different
lexical, phonological, and articulatory representations (e.g., Blanco-Elorrieta & Caramazza,
2021; Costa, 2005; La Heij, 2005). Successful production requires using representations that

belong to the target language but not to the non-target language. In the case of bilingual

speakers (especially L1-dominant speakers), it is typically assumed that using their native

language (L1) does not require additional control over representations of their second

language (L2) because L1 representations are readily available and receive no interference

from L2 representations. However, when an L1-dominant bilingual wants to use their weaker

L2, the representations of L2 are less active and receive interference from strongly activated

L1 representations, so accessing them may require additional control processes. The most

prominent theory of bilingual language control proposes that resolution of between-language

interference is implemented by active inhibition of L1 representations (Green, 1998);

however, it does not specify at which level of word processing (e.g., lexical, phonological,

articulatory) inhibition is implemented. On the other hand, other theoretical models argue that

inhibition is not necessary for the successful selection of the target language (Blanco-Elorrieta

& Caramazza, 2021; Costa, 2005; Koch et al., 2010). For example, the persisting activation

account assumes that when a bilingual wants to use the weaker L2, its representations receive

additional activation, which allows them to surpass the level of activation of the stronger L1

(for a review see: Koch et al., 2010). A recent extension of the activation-based account

proposed that activation of target representations does not operate on a whole-language level

(i.e., it does not apply to all items of L1 or L2) but it is rather a sum of activations driven by

different factors, such as baseline word frequency, recency of use and communicative context

(Blanco-Elorrieta & Caramazza, 2021). As such, both bilingual and monolingual speech

production can be explained by the same selection mechanisms that select the most active

representation at a given moment.

The L2 after-effect as a window to study neurocognitive mechanisms of bilingual
language control

As has been already indicated, the lion's share of evidence on the involvement of

control in bilingual language production comes from the language-switching paradigm (for a

recent review see Sulpizio et al., 2020) which – as we argued above - has several important

limitations. Some of these limitations are circumvented in a blocked language picture naming

task that does not require as frequent switches between languages as the language switching

task and yet allows control mechanisms involved in bilinguals' language production to be

tested. Several studies have observed that completing a task in L2 results in a subsequent

slow-down of naming or an increase of errors committed in L1 (Branzi et al., 2014; Degani et

al., 2019; Gollan & Ferreira, 2009; Van Assche et al., 2013; Wodniecka, Szewczyk, et al.,

2020). This difficulty in using L1 after a short exposure to L2 has been referred to as the L2

after-effect. Any task involving language production in L2 can lead to the L2 after-effect:
simply reading a list of words aloud (Degani et al., 2019), naming a set of pictures (Branzi et
al., 2014, 2016; Gollan & Ferreira, 2009; Guo et al., 2011; Wodniecka et al., 2020) or coming
up with words belonging to a given semantic category (Van Assche et al., 2013).

Available theories of bilingual speech production provide different explanations of the
L2 after-effect. According to inhibition-based accounts, the retrieval difficulty that speakers
experience in L1 after using L2 is a lingering consequence of the strong inhibition that had to
be applied to L1 representations during L2 use (Green, 1998; Guo et al., 2011; Misra et al.,
2012). Under the inhibitory accounts, resolution of the interference that drives the L2 after-
effect relies on a set of domain-general mechanisms (e.g., Abutalebi & Green, 2016; Green,
1998; Green & Abutalebi, 2013). An alternative proposal assumes that the L2 after-effect is
driven by increased interference between the representations of L1 and L2 due to the carry-
over effects of increased L2 activation (Blanco-Elorrieta & Caramazza, 2021; Branzi et al.,
2014, 2016; Koch et al., 2010).

To date, the origin of the L2 after-effect has been explored in three neuroimaging
studies using fMRI. They revealed that the difficulty in word retrieval in L1 after using L2
results in increased engagement of brain regions often linked to domain-general control
mechanisms (Branzi et al., 2016; Guo et al., 2011; Rossi et al., 2018) such as interference
suppression (left middle frontal gyrus, left inferior parietal gyrus; Guo et al., 2010), visual
attention shifts (left parietal cortex; Guo et al., 2010), articulation (right postcentral gyrus;
Guo et al., 2010), response inhibition (right inferior frontal cortex; Branzi et al., 2016 right
frontal pole and right superior parietal lobule; Rossi et al., 2018), response selection (left
prefrontal cortex; Branzi et al., 2016), attention (bilateral inferior parietal lobules; Branzi et
al., 2016), and conflict monitoring and resolution (bilateral anterior cingulate cortex; Rossi et
al., 2018). While most of these structures have been linked to several different domain-
general control mechanisms, some structures are thought to support language-specific
processing as well as articulation. As such, the results of previous studies do not unanimously
identify the mechanisms that drive the L2 after-effect.

Capturing the neural basis of the L2 after-effect: methodological challenges

Importantly, three studies that explored the L2 after-effect were not free of
methodological challenges. Two out of three studies report neural contrasts that involved
different samples of participants (Guo et al., 2011; Rossi et al., 2018). This could be

problematic as the engagement of language control mechanisms may be dependent on some
aspects of language experience (see Casado et al., 2022 for evidence of the influence of
language balance on the magnitude of engagement of global language control). What is more,
a study by Rossi et al. (2018) compared a sample of bilinguals to monolinguals, but this does
not make it possible to disentangle the effects of previous use of L2 from more general
differences between bilingual and monolingual language processing. Branzi and colleagues
(2016) addressed these limitations by comparing preceding context effects within the same
bilingual participants; however, in this study, the baseline L1 block, which was compared to
L1 after L2 in order to estimate the L2 after-effect, was in some participants preceded by a
task in L2. Since the L2 after-effect is relatively long-lasting (at least 5 minutes, see Branzi et
al., 2014; Casado et al., 2021; Wodniecka, Szewczyk, et al., 2020; for review see: Wodniecka,
et al., 2020), a baseline for comparison (i.e., L1 naming) should not be preceded by a task in
L2. It is therefore possible that the observed L2 after-effect was confounded by factors such
as training, fatigue, or accumulating semantic interference (Runnqvist et al., 2012; for a
discussion of how trial-related factors can affect the measurement of L2 after-effect see:
Casado et al., 2022).

Moreover, the previous studies relied on group-level fMRI analysis. This approach
focuses on the commonalities in activations between participants but is less sensitive to
individual variability in brain organization, which is especially important among bilinguals
whose variable language experience (related to a number of factors, such as immersion, age of
acquisition, proficiency, daily use of language), may translate into changes in the functional
and structural organization of the brain (e.g., DeLuca et al., 2019, 2020; Fedeli et al., 2021).
What is more, while previous studies undoubtedly identified brain structures engaged in the
L2 after-effect, the interpretations regarding the precise mechanisms represented by the
observed brain response were not directly tested and were proposed based on the function
assigned to a given anatomical structure (the so-called *reverse inference* problem, Poldrack,
2006; for a discussion of issues related to this approach in studying language see Fedorenko,
2021). It has been shown that macroanatomical cortical areas are characterized by vast inter-
individual variability, which means that the link between anatomical structure and functional
organization is not straightforward, especially for high-level cognitive processes such as
language (Frost & Goebel, 2012; Tahmasebi et al., 2012; for a discussion see Fedorenko,
2021; Michon et al., 2022).

Current study

In the current study, we explored the nature of bilingual language control by looking at

the brain basis of the L2 after-effect. To address the limitations of experimental designs

previously used to measure the L2 after-effect, we used a within-participant design that

carefully controlled for the effects of the preceding language context as well as for possible

effects of task order and fatigue. We had two specific goals: first, to determine to what extent

the control mechanisms engaged in bilingual language control are domain-general vs.

language-specific; second, to identify the specific process that may underlie difficulties in L1

production following L2 use, namely lexical retrieval difficulty (Casado et al., 2022),

articulatory difficulty (Guo et al., 2011), and increased interference between languages

(Branzi et al., 2014). We addressed these questions by using fMRI and a set of functional

localizers, which - to the best of our knowledge - is an approach that has not been used before

in studies exploring the mechanisms of language control in bilinguals. A crucial advantage of

functional localizers is that they allow the modeling of brain responses at a subject-specific

level; this increases the sensitivity of the statistical analyses and models inter-individual

variability better than typical group level analyses.

To explore the domain-specificity or domain-generality of bilingual language control

mechanisms engaged in the L2 after-effect, we used two well-established functional localizers

which make it possible to track down subject-specific networks that correspond to the

language system (Malik-Moraleda, Ayyash et al., 2022; Fedorenko et al., 2010) and the

Multiple Demand network (Duncan, 2010; Fedorenko et al., 2013). The language localizer is

a well-established paradigm that allows the identification of a robust network in the brain

which spans the frontal, temporal, and parietal regions, which support the processing of

language (e.g., Malik-Moraleda, Ayyash et al., 2022; Fedorenko, 2021; Fedorenko et al.,

2010, 2011). This network has a robust specificity for language: it is engaged during reading

(e.g. Fedorenko et al., 2010), listening (e.g. Pallier et al., 2003; Scott et al., 2017) and

production (Hu et al., 2022; Liljeström et al., 2015), but it shows little engagement in non-

linguistic tasks (Benn et al., 2021; Chen et al., 2021; Fedorenko et al., 2011; Ivanova et al.,

2021) or social reasoning (Shain et al., 2022). Importantly, even though the language network

is usually localized using a comprehension-based task it is equally sensitive to language

production (Hu et al., 2020).

The Multiple-Demand localizer, on the other hand, allows the identification of another

robust network in the brain which responds to increased task demands across many different

types of tasks (Assem, Blank, et al., 2020; Braver et al., 2003; Cole & Schneider, 2007;
Dosenbach et al., 2006; Duncan, 2010; Fedorenko et al., 2011, 2013; Miller & Cohen, 2001).
Activity within this network has been shown to decrease with the increasing automatization of
task performance (Chein & Schneider, 2005) and increase as a function of task complexity
(Crittenden & Duncan, 2014; Fedorenko et al., 2013; Shashidhara et al., 2019), time pressure
and reward (Shashidhara et al., 2019). Importantly, the frontoparietal Multiple Demand
network shows little or no domain-specificity (Assem, Glasser, et al., 2020): it is robustly
activated in a range of different linguistic and non-linguistic cognitive tasks (Fedorenko et al.,
2013; Dosenbach et al., 2006; Shashidhara et al., 2019), including low-level perceptual tasks
(Cole & Schneider, 2007).

While both the Language and the Multiple Demand networks show vast cross-subject
variability (Fedorenko et al., 2010; Smith et al., 2021), they show a clear functional and
topographic dissociation on a single-subject level (e.g., Blank et al., 2014; Diachek et al.,
2020). As such, if brain activity corresponding to the L2 after-effect is found within one of
these two networks, it will provide straightforward evidence for the engagement of domain-
general or language-specific mechanisms.

To investigate the specific mechanisms that drive the L2 after-effect, we used 3
additional localizers that focused on lexical access (a localizer based on the Verbal Fluency
task), articulation (a localizer based on a simple articularion task), and lexical interference (a
localizer based on the Stroop task). Each of them tapped into one of the previously described
mechanisms that possibly underlies the L2 after-effect. The Stroop task (Stroop, 1935) was
used to capture brain structures engaged in the resolution of interference and inhibition of
non-target responses in speech production (Diamond, 2013). An overlap between these
structures and those involved in the L2 after-effect would provide an argument that the L2
after-effect is driven by increased interference between the two languages.

The verbal fluency task was used to identify brain regions that are sensitive to the
difficulty of lexical access. As previously discussed, difficulties in lexical access seem to be
one of the most parsimonious explanations of the L2 after-effect (Wodniecka, Szewczyk, et
al., 2020; Casado et al., 2022). The verbal fluency task is thought to engage cognitive
mechanisms that support word retrieval (Birn et al., 2010; Friesen et al., 2021; Shao et al.,
2014). Specifically, the verbal fluency task was shown to tap into the efficiency of the lexical
search and selection processes (Shao et al., 2014) as well as executive function mechanisms
engaged in word retrieval (Friesen et al., 2021; Martin et al., 1994). Therefore, if L1 retrieval

difficulty after L2 is due to L1 lexical access being hampered, then the brain activity
corresponding to the L2 after-effect should overlap with the network identified using a verbal
fluency localizer. Finally, verbal fluency performance has been shown to depend either on
individual differences between subjects that are related to vocabulary size and working
memory capacity, or on the mere speed of information processing (Unsworth et al., 2011).
Therefore, a subject-specific localizer based on a verbal fluency task allows us to better
capture the inter-individual variability in lexical retrieval between subjects.

Finally, to test the extent to which the L2 after-effect is driven by the articulatory
difficulty of word production in L1, we used a localizer that allows us to identify brain
regions specifically linked to articulation (Basilakos et al., 2018). This localizer was based on
a contrast between articulating syllables and performing a simple motor sequence (finger
tapping). An fMRI experiment which examined the neural basis of the L2 after-effect
suggested that speaking in L1 after being exposed to L2 entails the engagement of additional
articulatory resources (Guo et al., 2011). Moreover, more-recent behavioral evidence (Broos
et al., 2018) suggests that greater difficulty in speaking L2 compared to L1 only affects the
late articulatory stage; if so, it is also plausible that exposure to L2 results in articulatory
difficulties for subsequent L1 use.

Method

Participants

Forty-two Polish-English late bilinguals took part in the study. One participant was
excluded due to excessive head motion (>2mm) in the main fMRI task, resulting in a final
sample of forty-one participants (31 females, 10 males, mean age = 23.29, SD = 3.24, range:
19-32). All participants were Polish native speakers who learned English as their second
language. To participate in the experiment, they had to declare sufficient proficiency in
English (B2 or higher) and obtain at least 20/25 points on the General English Test by
Cambridge Assessment (mean score = 21.32, SD = 1.94). In the behavioral session, the
participants' proficiency in English was assessed again using LexTale (mean score = 71.58%,
SD = 9.91%; Lemhöfer & Broersma, 2012). In addition, participants completed a
questionnaire in which they rated their proficiency in all known languages with regards to
reading, writing, listening, speaking, and accent; participants also gave information on their
daily use of all languages and the age at which they started to learn each of them. Detailed
information on the participants' proficiency, age of acquisition, and daily use of languages
can be found in Table 1. All participants gave written consent for participation in the study.
The study was approved by the Ethics Committee of the Institute of Psychology of the
Jagiellonian University concerning experimental studies with human subjects.

Table 1. Language experience of participants. Information on self-rated proficiency, language learning and daily use of languages is given for all languages that participants declared to know. L1 always refers to Polish and L2 to English. L3 and L4 were a variety of different languages (incl. German, French, Italian, Spanish, Russian, Czech, Japanese, Korean, Norwegian, Latin and Esperanto). Not all the participants reported knowing an L3 or L4.

Language	**L1** (Polish)	**L2** (English)	**L3** (various)	**L4** (various)
n	41	41	36	17
Self-rated proficiency (1-10)				
reading	9.72	4.34	2.76	1.79
listening	9.62	4.10	2.45	1.84
writing	9.26	3.81	2.21	1.48
speaking	9.49	3.89	2.48	1.68
accent	9.56	3.45	2.45	1.65
Language learning				
Age of acquisition onset of learning	0.00	6.44	14.34	15.24
Age of acquisition: onset of using	3.17	9.15	15.41	16.88
Years of formal education	14.72	11.56	4.03	4.18
Daily use of languages				
passive	63.26%	31.79%	4.03%	3.25%
active	81.31%	16.31%	2.06%	1.31%
Verbal Fluency (mean number of words produced)				
category	20.04	13.36	-	-
letter	12.64	10.62	-	-

Study Design

The experiment consisted of 3 testing sessions: one behavioral and two MRI scanning

sessions. The design of the behavioral and MRI sessions is presented in detail in Figure 1, and

the details of each task are discussed in the Design section.

Figure 1 . Summary of the design of the behavioral and MRI sessions. In the behavioral sessions,

all participants completed the tasks in a fixed order. In the MRI sessions, the order of conditions

completed in the first and the second session (experimental manipulation with L1 or L2) was

independently counterbalanced between participants; the set of localizer tasks assigned to each session

was also counterbalanced between participants.

In the behavioral session, participants completed a series of tasks, including semantic

and phonetic verbal fluency in L1 and L2 (see details below), LexTale in English (Lemhöfer

& Broersma, 2012), a language questionnaire, and a short version of Raven's Progressive

Matrices test (only uneven items) as a measure of a non-verbal intelligence. The behavioral

session also included a short training session on the tasks to be performed in the MRI

sessions.

During the MRI sessions, participants completed the main task, namely picture

naming, and a series of localizer tasks. The main task was split between two sessions, with

one condition (L1 or L2 in the first block) completed in the first session and the other

condition completed in the second session, with the order counterbalanced across sessions.

Localizer tasks included an auditory language localizer task (Malik-Moraleda, Ayyash et al.,

2022; Scott et al., 2017), an articulation localizer, a Stroop task, a visual working memory

task corresponding to the Multiple Demand localizer (Fedorenko et al., 2013), and a verbal
fluency task (order counterbalanced, see Figure 1 for details). In tasks requiring overt spoken
responses (i.e., picture naming, Stroop task, articulation task, and verbal fluency) participants'
responses were recorded with FOMRI III, which is an MR-compatible, fiber-optic
microphone system (OptoacousticsLtd., Or-Yehuda, Israel). Participants were instructed to
minimize jaw movements while giving the spoken responses, which they had practiced during
the behavioral session.

Materials and Procedure

Behavioral session

Verbal Fluency task. In this task, participants were asked to produce as many words
belonging to a given category (fruits or animals) or beginning with a given letter (B or M) as
they could within a 1-minute time limit. Cues were counterbalanced between languages. Each
participant first completed the task in Polish (with one category and one letter) and then in
English (also with one category and one letter). Each cue (letter or category) was displayed on
the screen for 1 minute, during which participants provided verbal responses that were
recorded with the microphone. This task was used as a complementary measure of relative
proficiency in L1 and L2; the mean number of words produced for letter and category tasks in
each language is reported in Table 1.

LexTale. In the LexTale task, participants were instructed to say whether the string of letters
presented on a screen corresponded to an existing word in English. The response (yes/no) was
provided via a keyboard. Each word or nonword appeared in the center of the screen and
remained there until the participant responded. The task used the same words and nonwords
as the original LexTale task (Lemhöfer & Broersma, 2012).

Language questionnaire. In the language questionnaire, participants were asked to respond to
four categories of questions. The first category concerned proficiency in the native and
foreign languages that the participants knew. It included a self-rating scale on the five
language skills (reading, writing, listening, speaking, and accent) and questions concerning
the age of acquisition. The second category included questions on patterns of language use:
the participants were asked to estimate how much time during a typical day they actively (i.e.,
speaking, writing) and passively (i.e., watching movies, reading, listening to
music/radio/podcasts) use the languages they know. In the third category, participants were

asked to provide information about language use contexts: home, work, school, and free time.

Specifically, they were asked to estimate the number of hours that they use each language

they know in each of these contexts. Finally, the fourth category of questions referred to

demographic information, such as age, gender, and education. Filling out the entire

questionnaire took approximately 15-20 minutes.

MRI task training. In this part of the behavioral session, participants completed short

versions of the tasks that they were going to perform during the MRI session (for details of

the tasks see the MRI session design). Each task lasted approximately 1-2 minutes and was

repeated if a participant asked for more practice. In tasks requiring overt responses,

participants were asked to practice minimizing jaw movements while speaking.

Raven test. In this task, participants completed a shortened version of Raven's Advanced

Progressive Matrices test (only odd-numbered items). They had 20 minutes to complete as

many matrices as they could. The test was used in a paper-pencil format and was aimed to

provide a measure of fluid intelligence.

MRI sessions – the main task

In the main task, participants were asked to overtly name pictures of objects in four

task blocks: two blocks in each session. One task block corresponded to one functional run.

The first block of each session corresponded to one of the experimental manipulations:

exposure to L1 (which required naming pictures in L1) or exposure to L2 (which required

naming pictures in L2). The order of experimental manipulation blocks was counterbalanced

between participants, i.e., some of them started with the L2 exposure conditions in the first

session and completed the L1 exposure condition in the second session; some of them started

with the L1 exposure condition in the first session and completed the L2 exposure condition

in the second session. In the second block of each session, participants were asked to name

pictures in L1. Each block contained 55 colored pictures representing objects (from the CLT

database: Haman et al., 2015; Wolna et al., 2022). Pictures were not repeated between blocks.

Each block started with a fixation cross ("+") displayed on a screen for 7000ms.

Subsequently, 55 pictures were presented following a rapid event-related design. Each picture

was displayed on a white background for 2000ms and was followed by a fixation cross,

jittered at an interstimulus interval ranging from 607ms to 10144ms (M = 3091ms, SD =

2225ms). The order of picture presentation and interstimulus intervals was optimized using

optseq2 (Dale, 1999). Finally, at the end of each block, a fixation cross was presented for

8000ms. Overall, each block of the Picture Naming task lasted 295s.

MRI sessions – localizer tasks

After completing the main task, in each session participants completed two or three

localizer tasks. Each localizer consisted of two functional runs.

Language localizer. Participants listened to fragments of *Alice in Wonderland* in L1 and

distorted speech recordings in which it was impossible to understand what the speaker was

saying. The task followed the design introduced by Scott et al. (2017) and Malik-Moraleda,

Ayyash et al. (2022), in which the localizer contrast was based on the intact > distorted speech

condition contrast. The intact condition consisted of short passages from Alice in Wonderland

read by the native speaker. The degraded speech condition was created based on the intact

condition but, it sounded like poor, indecipherable radio reception of speech with preserved

prosody but not decipherable words or phonemes. In each functional run, participants listened

to 6 short passages of Alice in Wonderland in their native language (18s each) and 6 passages

of distorted speech (18s each). Additionally, 4 fixation blocks (12s each) were included in

each run. The total duration of each run was 264s. Each participant completed 2 runs of the

task.

Articulation localizer. Participants performed two tasks: an articulation task in which they

were asked to repeat out loud a string of previously memorized syllables ("bła" [bwa], "sza"

[a], "tra" [tra] – all frequently occurring in Polish as word constituents, but meaningless in

isolation) and a motor task in which they were asked to perform a simple motor sequence with

the fingers of both hands (touching the thumb with their index, ring, and little finger,

respectively). The localizer was based on the articulation > motor sequence contrast. In each

functional run, participants completed 6 articulation and 6 finger tapping blocks (18s each).

Additionally, 4 fixation blocks (12s each) were included in each run. Each participant

completed 2 runs of the task.

Stroop localizer. Participants were asked to overtly name the color of the presented word's

font. The design of the task followed the one used by Fedorenko et al. (2011), in which

participants completed blocks of the easy and hard versions of the task. In the easy condition,

the words corresponded to non-color adjectives [examples in the original language mased for

review]; in the hard condition, the words corresponded to color adjectives [examples in the

original language mased for review]. In the hard condition, half of the trials were conflict

trials in which the word's font color did not match its meaning; the other half were congruent trials. The order of trials within the hard condition was randomized. Each run started with a fixation cross ("+") displayed on a gray background for 8000ms. Following that, participants completed 16 task blocks (8 per condition, 18s each). Task blocks contained 12 trials which started with a fixation cross displayed for 500ms followed by a word displayed for 1000ms. Additionally, five fixation blocks (18s) were included in each run. The order of experimental and fixation blocks was counterbalanced between participants and runs (four counterbalance lists were created and each participant completed two of them). The localizer was based on the hard > easy condition contrast. The total duration of one run was 394s. Each participant completed 2 runs of the task.

Multiple Demand localizer. Participants were asked to perform a spatial working memory task (e.g., Fedorenko et al., 2011, 2013). The design of this task followed the one used by Fedorenko et al. (2011), in which the localizer contrast was based on the hard > easy condition contrast. In each trial, participants saw four 3 x 4 grids and were asked to memorize the locations of the fields that were marked as black on each of the grids. In the easy condition, they had to memorize one location per grid (four in total); in the hard condition, they had to keep track of two locations per grid (eight in total). After memorizing the grids, participants had to choose a grid that contained all the memorized locations from among two grids presented on the screen. Each trial started with a fixation cross displayed on the screen for 500ms; this was followed by four grids, each displayed for 1000ms. After that, the choice task appeared on the screen until a response was given (for a max of 3750ms); this was followed by feedback (right or wrong) displayed for 250ms and a fixation cross, which was displayed for 3750ms minus the reaction time in the choice task. In each of the conditions (easy and hard) participants completed 4 trials per block (34s in total). Each run contained five easy condition blocks and five hard condition blocks. Additionally, six fixation blocks (16s) were included in each run. The order of experimental and fixation blocks was counterbalanced between participants and runs (four counterbalance lists were created and each participant completed two of them). The total duration of one run was 438s.

Verbal fluency localizer. Within 10 s blocks, participants were asked to produce as many words as they could that either pertained to a given category (semantic fluency) or started with a given letter (phonemic fluency). The entire task was completed in L1. In each run, participants completed six semantic fluency and six phonemic fluency blocks, each focusing on a separate letter or semantic category. Each letter and category were presented only once,

and they did not repeat between blocks and runs. A full list of stimuli used in this task is

available in the Supplementary Materials. In addition, each run included five baseline blocks,

in which participants were asked to enumerate the months starting from January (Birn et al.,

2010). The localizer contrast was based on task (semantic or phonemic fluency) > baseline

condition contrast. Following Birn et al. (2010), each block was followed by a fixation cross

displayed for 7s–13 s. Each run lasted 348s. Each participant completed 2 runs of the task.

MRI Data Acquisition

MRI data were acquired using a 3T scanner (Magnetom Skyra, Siemens) with a 64-

channel head coil. High-resolution, whole-brain anatomical images were acquired using a T1-

weighted MPRAGE sequence (208 sagittal slices; voxel size $0.9 \times 0.9 \times 0.9$ mm^3; TR = 1800ms,

TE = 2.32ms, flip angle = 8°). A gradient field map (magnitude and phase images) was

acquired with a dual-echo gradient-echo sequence, matched spatially with fMRI scans (TE1 =

4.92ms, TE2 = 7.3ms, TR = 503ms, flip angle = 60°). Functional T2*-weighted images were

acquired using a whole-brain echo-planar (EPI) pulse sequence (50 axial slices, 3x3x3 mm

isotropic voxels; TR = 1400ms; TE = 27ms; flip angle = 70°; FOV 192 mm; MB acceleration

factor 2; in-plane GRAPPA acceleration factor 2 and phase encoding A>>P) using 2D

multiband echo-planar imaging (EPI) sequences from the Center for Magnetic Resonance

Research (CMRR), University of Minnesota (Xu et al, 2013). In order to account for magnetic

saturation effects, the first four volumes of each sequence were not included in the analysis.

MRI Data preprocessing

All data were visually inspected for artifacts. The non-brain tissue was removed using

the FSL Brain Extraction Tool (Smith, 2002). For the preprocessing of functional data, we

used FSL FEAT (version 6.0.0; Woolrich et al., 2004). The preprocessing steps included

high-pass temporal filtering (100s), spatial smoothing (FWHM = 5 mm), distortion correction

of functional images using fieldmap with FUGUE, co-registration and normalization using

FLIRT (Jenkinson et al., 2002; Jenkinson & Smith, 2001) and motion correction using

MCFLIRT (Jenkinson et al., 2002) with 6 rigid-body transformations. Functional images were

co-registered with their anatomical images; subsequently, they were registered to MNI

standard space (FSL's 2mm standard brain template).

MRI Data analysis

In the 1st-level statistical analysis, a double-gamma hemodynamic response function

was used to model the BOLD signal corresponding to each event (trial in the main task, as it

followed the event-related design; and blocks in the localizer tasks which followed a blocked

design). The estimates of motion obtained with MCFLIRT (Jenkinson et al., 2002) were

included in the 1st-level GLM as nuisance covariates. For the localizer tasks, the two

functional runs of each localizer task were combined in a second-level analysis for each

participant using an FSL fixed-effect model. To ensure better comparability of our findings

with previous studies, for language, Multiple Demand, and articulation localizers we used sets

of functional parcels established and validated by previous studies instead of creating group-

level parcels based on our data (twelve language parcels: Fedorenko et al., 2010; twenty

multiple demand parcels: Fedorenko et al., 2013; eleven articulation parcels: Basilakos et al.,

2018). It is important to note that the group-level parcels from previous studies showed a high

overlap with parcels for the language, Multiple Demand, and articulation networks defined

based on our data. For the remaining Stroop and Verbal Fluency localizers, group-constrained

subject-specific ROIs were defined for each participant following the approach proposed by

Fedorenko et al. (2010). In the first step, the individual subject activation maps thresholded at

5% of most active voxels. In most cases this threshold was slightly more conservative than z

> 3.1, uncorrected at the whole brain level. We chose to use the 5% of most active voxels

instead of a standard threshold because it yielded maps with comparable number of voxels

between subjects that consequently produced more coherent group-level parcels.

Subsequently, the individual parcels were binarized and overlaid on top of each other. The

probabilistic map obtained this way was then thresholded so that only voxels active in more

than 5 participants (~20%) would be included. In the second step, the probabilistic maps were

smoothed using an 8x8x8 Gaussian kernel and subsequently, group-level partitions were

created using a watershed algorithm for segmentation. This procedure yielded nineteen group-

level parcels for the Stroop task and fifteen group-level parcels for the Verbal Fluency task.

Before further analyses, we excluded very small parcels (mean size across subjects smaller

than 10 voxels). This yielded a final set of thirteen parcels within the Stroop network and

thirteen parcels within the verbal fluency network. In each of these parcels significant

activation to the localizer contrast, i.e.: color words > neutral words for the Stroop task and

fluency > automated speech for verbal fluency (thresholded at z>2, corrected for multiple

comparison using a cluster-wise correction at p<0.05) was found in at least 51% of

participants for the Stroop parcels (mean = 70.36%) and in at least 73.17% of participants for

the Verbal Fluency parcels (mean = 84.99%). In the last step, group-level partitions were

intersected with subject-specific data, thus yielding a set of subject-specific group-constrained

ROIs. Finally, we selected the top 10% most responsive voxels within each parcel based on

the *z*-value map for a given localizer contrast (Language: speech > degraded speech, MD:

hard > easy visual working memory task, Articulation: syllables > motor sequence, Stroop:

color words > neutral words, and Verbal Fluency: fluency > automated speech) and we

binarized the obtained ROIs. We then used the ROIs corresponding to all localizer tasks to

extract parameter estimates corresponding to % signal change in the two critical conditions of

the main task: naming in L1 after L1, and naming in L1 after L2. Parameter estimates were

extracted from the 1st-level analyses for each participant separately. For the analysis of the

functional localizer scope, we used an across-run cross-validation procedure to extract the %

signal change corresponding to each localizer task condition (i.e., we defined the functional

parcels based on the first functional run and estimated responses to the localizer contrast on

the second run and repeated this procedure using the second run to define the functional

parcels and the first run to extract response estimates; see Fedorenko et al., 2010). This was

done using FSL's FEATquery tool (http://www.FMRIb.ox.ac.uk/fsl/feat5/featquery.html).

Further analyses were performed on the extracted parameter estimates in R (version: 4.0.2; R

Core Team, 2020). Following an approach that proposes modeling activity within a functional

network on the network level instead of relying on separate analyses for each ROI (Malik-

Moraleda et al., 2021), we fitted one linear mixed-effect model for each functional network

according to the following formula:

554   % signal change ~ PrecedingLanguage +

555   (1 + PrecedingLanguage | Subject) +

556   (1 + PrecedingLanguage | ROI)

In each model, the fixed effect of PrecedingLanguage corresponded to a predictor with

two levels: *L1 after L1* and *L1 after L2*. Before the analysis, this categorical predictor was

deviation-coded (L1 after L1 = -0.5; L1 after L2 = 0.5). The analysis was performed using the

lmer() function from the lmerTest package (Kuznetsova et al., 2017)For each functional

network, we first fitted a maximal model and then identified the best random-effects structure

following the recommendations of Bates et al. (2018). The reason for modeling the activity

within each network with ROI as a random effect instead of a fixed effect relies on the

assumption that the different brain regions engaged within a given functional network exhibit

a significant degree of functional integration. To further explore the fine-grained differences

between our critical conditions, which might only affect a subset of functional parcels within

each network, we also fitted separate models for each ROI in each of the functional networks.

The results of the ROI-specific analyses are reported in the *Supplementary Materials*. Raw

neuroimaging data used in this study are freely available at the OpenNeuro repository

https://openneuro.org/datasets/ds004456. Data and code necessary to reproduce the ROI and

behavioral   analyses   are   freely   available   at

https://osf.io/59za8/?view_only=e23f3500d12e4109b75ca18fdbd320e5.

Results

Functional localizers' scope

Functional networks identified with a set of localizer tasks were based on robust contrasts that

yielded strong effects in each network (language localizer: intact vs. degraded speech: β =

0.520, $t(40) = 5.684$, $p < .001$; Multiple Demand localizer: hard vs. easy visual working

memory task: β = 0.580, $t(40) = 13.12$, $p < .001$; Articulation localizer: articulation vs. motor

task: β = 1.337, $t(40) = 20.84$, $p < .001$; Stroop localizer: Stroop task blocks vs. congruent

color naming: β = 0.435, $t(40) = 11.75$; $p < .001$; Verbal Fluency localizer: fluency vs.

enumerating months: β = 1.155, $t(40) = 16.73$, $p < .001$). The localizer tasks yielded five sets

of parcels covering a wide range of brain regions. Language and Multiple Demand parcels

covered two robust non-overlapping networks in the brain (see Figure 2A, although small

parts of language and Multiple Demand network show some overlap, this was only the case

on for the group-level parcels, not the individual subject-specific parcels). The Articulation

parcels included a set of regions partially overlapping with the language network in the

temporal ROIs but it also included parcels that selectively responded to articulation but not

language (see figure 2B). As expected, the Stroop network largely overlapped with the

Multiple Demand network (for similar results see Fedorenko et al., 2013), however, the

Stroop parcels were considerably smaller than the Multiple Demand network (see figure 2C).

The differences in size between the Stroop and the Multiple Demand networks are partially

driven by the fact that the Stroop parcels were created based on single-subject activation maps

thresholded at 5% of the most active voxels (for details see Methods), which puts an a priori

constraint on the size of the activation maps and the parcels that are created based on these

maps. However, the Stroop network was also considerably smaller than the Multiple Demand

network when other, less stringent threshold were used (e.g., z>2). As such, the Stroop

network can be considered a functional sub-system of the larger and more general Multiple

Demand network. Finally, the Verbal Fluency localizer yielded a set of strongly left-

lateralized parcels partially overlapping both with the Language and the Multiple Demand

networks (for Verbal Fluency parcels see Figure 4).

To further characterize the functional networks as well as the scope of the localizer tasks used

in this study, we ran additional analyses showing (1) responses to all localizer tasks used in

this study within the language and the Multiple Demand networks; and (2) responses to the

Language and Multiple Demand localizer tasks (i.e., listening to degraded vs. intact speech

and hard vs. easy working memory task) in all five functional networks used in this study.

Results of these analyses can be found in the Supplementary Materials.

Figure 2. Brain masks corresponding to group-level functional networks. The figure presents a

comparison of brain networks identified using different functional localizers (A) Language (red) and

Multiple Demand (blue) networks; (B) Language (red) and articulation (yellow) networks; (C)

Multiple Demand (blue) and Stroop (turquoise) networks. All networks presented on the graph

correspond to a combination of group-level parcels.

Bilingual language control: domain-general or language-specific

To address the question of whether the bilingual language control mechanisms driving

the L2 after-effect rely on domain-general or domain-specific mechanisms, we compared

brain activity corresponding to naming pictures in *L1 after L1* and *L1 after L2* within two

functional networks: the language network and the Multiple Demand network. The results

revealed no significant effect of the preceding language within the language network (β =

0.026, *t(40)*= 0.89, *p* = .379, effect size = 0.28); however, a significant effect of the preceding

language was found within the Multiple Demand network (β = 0.072, *t(40)* = 2.679, p = .011,

effect size = 0.85). The results of the comparison between *L1 after L1* and *L1 after L2* within

the language and Multiple Demand networks are presented in **Figure 3.**

Figure 3. Neural activation corresponding to the L2 after-effect within the Multiple Demand and
Language networks. Plots represent % BOLD signal change in response to naming pictures in L1 after
L1 (light hues) and L1 after L2 (dark hues). Rendered maps present sets of group-level parcels used in the
subject-specific analysis (LH = left hemisphere; RH = right hemisphere).

Separate analyses on each individual ROI showed significant differences between *L1 after L1*

and *L1 after L2* in four ROIs in the Multiple Demand network (left and right angular gyrus /

superior parietal lobule, right superior and middle frontal gyrus, and right frontal pole), but no

significant results were found in the language network. Full results showing comparisons

between conditions for each ROI and each network are presented in **Tables S1 and S2** in the

Supplementary Materials.

Digging deeper: specific mechanisms of bilingual language control

To explore which specific mechanisms drive the L2 after-effect, we compared the

brain activity corresponding to naming pictures in *L1 after L1* and *L1 after L2* within three

functional networks identified with the Articulation, Stroop, and Verbal Fluency tasks. We

did not find a significant effect of the preceding language in any of the three networks

(Articulation: $\beta = 0.005$, $t(40) = 0.09$, $p = .926$, effect size = 0.03; Stroop: $\beta = 0.034$, $t(40) =$

1.02, $p = .314$, effect size = 0.32; Verbal Fluency: $\beta = 0.003$, $t(40) = 0.07$, $p = .941$, effect size

= 0.02; see Figure 4). Separate analyses on each individual ROI for each of these networks are

presented in **Tables S3-5** in the *Supplementary Materials*.

Figure 4. Neural activation corresponding to the L2 after-effect within the Articulation, Verbal Fluency and Stroop networks. Plots represent % BOLD signal change in response to naming pictures in L1 after L1 (bright hues) and L1 after L2 (dark hues). Rendered maps present sets of group-level parcels used in the subject-specific analysis (LH = left hemisphere; RH = right hemisphere).

Discussion

Bilingual language control: domain-general or language-specific?

We explored the neurocognitive mechanisms underlying the L2 after-effect: a difficulty in producing words in the native language (L1) after having used the second language (L2). We compared brain activity corresponding to speaking in L1 after L2 with speaking in L1 after L1 within functionally defined brain networks. Our goal was to test (1) whether the L2 after-effect is related to activity in domain-general or domain-specific brain networks, and (2) whether the L2 after-effect reflects lexical retrieval difficulty, interference resolution, or increased difficulty at the phonological and articulatory stages of speech production. Unlike previous work exploring the neural basis of bilingual speech production, we used individual-level functional localizers (Fedorenko et al., 2010) to provide functional interpretations of brain responses associated with the L2 after-effect.

To address the question of whether the L2 after-effect is associated with increased

engagement of language-specific and/or domain-general brain networks, we compared brain

responses to speaking in L1 after L2 and L1 after L1 with two well-established, robust

functional networks: the language network (Malik-Moraleda, Ayyash et al., 2022; Fedorenko

et al., 2010; Scott et al., 2017) and the Multiple Demand network (Duncan, 2010; Fedorenko

et al., 2013). We found that the difficulty of naming pictures in L1 after L2 is related to

increased BOLD responses within the domain-general Multiple Demand network, but not

within the domain-specific language network. Furthermore, to explore whether the L2 after-

effect is driven by lexical access difficulty, increased interference between language-specific

representations, or articulatory difficulty, we used three additional functional tasks as

localizers: verbal fluency, the Stroop task, and an articulation task. We found that the L2

after-effect does not overlap with any of these process-specific localizers. We have also

replicated the results of a behavioral study (Casado et al., 2022) showing that the magnitude

of the L2 after-effect depends on the balance between languages, defined as a difference

between mean naming latencies in L2 and L1 (see *Supplementary Materials* for a detailed

report). This is an important note as our study is the first to record naming latencies

corresponding to the L2 after-effect in the scanner. As such, our data provides straightforward

evidence that the L2 after-effect can be successfully tested in the MRI scanner, despite the

noisy and challenging environment.

The L2 after-effect reflects the increased engagement of domain-general cognitive

control

We found evidence for increased response for speech production in L1 after L2

compared to L1 after L1 in the Multiple Demand network but not within the language

network (the interaction between condition and network was, however, not significant (β =

0.046, $t(2470)$ = 1.89, p = .058)). On the most general level, linking the L2 after-effect to

increased activation in the Multiple Demand system, supports models that posit domain-

general bilingual language control (Abutalebi & Green, 2016; Green & Abutalebi, 2013; Wu

et al., 2019).

The Multiple Demand network has been claimed to support cognitive control by

integrating different types of information and binding them together to support the current

cognitive operation (Assem et al., 2020, 2022; Cole et al., 2013; Koechlin & Summerfield,

2007). Activity within the Multiple Demand network has been linked to mechanisms

responsible for the implementation and reconfiguration of task sets (Dosenbach et al., 2006,
2007), voluntary attentional control (goal-directed attention; Asplund et al., 2010; Corbetta &
Shulman, 2002), and reorienting attention to novel tasks (Corbetta et al., 2008). An important
feature of the Multiple Demand network is that it is more engaged in less-automatized and
untrained tasks (Shashidhara et al., 2019), thus implying that it implements a domain-general
mechanism that supports any difficult or not sufficiently automatized activity that requires
additional attentional support. Seen from this perspective, speaking in L1 after L2 may recruit
additional domain-general resources to support the execution of a language task. However,
this leaves open the question of why exactly speaking in L1 after L2 is so much more difficult
than speaking in L1 after L1 that it requires additional support from the Multiple Demand
network.

The L2 after-effect reflects neither word retrieval difficulties nor increased
interference between language-specific representations

The L2 after-effect has been previously explained within different theoretical
frameworks (see Introduction). While these accounts provide plausible explanations of this
effect, no univocal evidence in favor or against these interpretations has been provided
(Casado et al., 2022; Wodniecka, Szewczyk, et al., 2020; Declerck & Koch, 2022). All
available accounts of the L2 after-effect build on a common theoretical assumption that the
difficulty in producing words in L1 after using L2 is a consequence of a control mechanism
that changes the balance between the relative activation of L1 and L2 representations (Casado
et al., 2022; for a discussion see Declerck & Koch, 2022). Importantly, according to these
accounts the change in L1-L2 balance results in increased interference between the two
languages that persists for some time after the speaker switches back to speaking in L1
(Casado et al., 2022), which, in turn, increases the difficulty of access and selection of (a)
lexico-semantic representations (e.g., Casado et al., 2022) and/or (b) phonological
representations and articulatory programs of L1 (e.g., Guo et al., 2011).

The results of the current study do not provide evidence supporting the notion that the
L2 after-effect is related to increased interference at the lexico-semantic, phonological, or
articulatory levels. We found that the L2 after-effect was not linked to an increased BOLD
response within the core language network. Since lexical access and selection mechanisms
pertain to core language computations (Hu et al., 2022), difficulties in lexical access should be
reflected by an increased BOLD response within the language network. This was, however,

not the case in the current study. What is more, we did not find increased brain response to

speech production in the L1 after L2 condition in either of the functional networks that were

identified using verbal fluency and articulation tasks. Even though both localizer tasks

produced robust responses and allowed us to identify a number of functional ROIs, they did

not overlap with the neural correlates of the L2 after-effect. Furthermore, we did not find any

overlap between the neural correlates of the L2 after-effect and the functional network that

was identified using the Stroop task; thus, we found no evidence for increased interference

between language representations during speech production in L1 after L2.

The results of the localizer-based analyses provide a rather unexpected

characterization of the neural correlates of the L2 after-effect. Contrary to the previous

interpretations (Branzi et al., 2014, 2016; Guo et al., 2011; Misra et al., 2012; Declerck &

Koch, 2022), the current results indicate that the L2 after-effect reflects neither lexical access

difficulty, nor articulatory difficulty, nor the additional engagement of language-specific

mechanisms that support word retrieval. In other words, our results suggest that the L2 after-

effect does not arise from within the language system itself (or from within language-specific

representations). So, how can we then account for the increased difficulty in speaking in the

native language after using L2? In the following paragraphs, we propose two putative

explanations of the difficulty in production in L1 following the use of L2.

Putative explanations of the L2 after-effect

The L2 after-effect as interference between task schemas

As described in the Introduction, most of the evidence for the involvement of

cognitive control in bilingual processing comes from the language-switching paradigm and,

more specifically, from the asymmetrical language switch cost (i.e., the finding that the cost

of switching to the dominant (L1) language is bigger than to the weaker (L2) language) and

language mixing cost effects (i.e., the stronger language (L1) suffers more when both

languages are mixed within one block of naming) (Costa & Santesteban, 2004; Gollan &

Ferreira, 2009; Meuter & Allport, 1999; Verhoef et al., 2009; for review see: Bobb &

Wodniecka, 2013; Declerck et al., 2020; Declerck & Koch, 2022; Declerck & Philipp, 2015).

Perhaps the most classic explanation of these effects assumes that they result from the

proactive language control that is recruited whenever participants have to name a picture in

the weaker L2. This control is recruited to "protect" the weaker L2 (less activated or less

automatized) from interference with the stronger L1, either by inhibiting L1 (e.g., Green,

1998; Green & Abutalebi, 2013) or by bolstering the activation of L2 (e.g., Blanco-Elloreta &

Caramazza, 2021). When, after using L2, a bilingual speaker switches to her L1, naming in

L1 is hampered by the consequences of prior control mechanisms: L1 is now either inhibited,

or there is an increased interference with L2, which is still strongly activated.

Explanations that assume the involvement of proactive control have been previously

used to account for the L2 after-effect (Wodniecka et al., 2020; Declerck & Koch, 2022), and

our current findings are generally consistent with this framework. However, they also point to

an important caveat regarding the level of the targeted representations. While models of

bilingual language processing typically assume that proactive control targets language-

specific representations, such as specific lexical units (see, e.g., BIA+ model, Dijkstra & van

Heuven 2002 for bilingual language comprehension; Blanco-Elorrieta & Caramazza, 2021 for

bilingual language production), the findings of the current study are incompatible with such

views. If proactive control operated on language-specific representations, we should see

modulations of brain activity in the after-L1 vs after-L2 conditions in regions directly

operating on these language-specific representations, namely in the language network, which

was not the case.

Nevertheless, our results support accounts that assume proactive control if one

assumes that the control does not target language-specific representations but rather more-

abstract ones. Previous literature claims the existence of such representations and refers to

them as control representations (Braver et al., 2021) or hierarchical task-schemas (Badre &

Nee, 2018). Such representations could be conceived as sets of rules for orchestrating access

to language-specific representations – the program we execute when we intend to speak in a

given language and which guides all subprocesses involved in subsequent episodes of

language production. This may perhaps be best understood in light of the cascade model of

executive control (Koechlin & Summerfield, 2007), which model proposes a hierarchy of

increasingly abstract control processes. At the sensorimotor control level, a stimulus is simply

associated with a reaction (e.g., picking up the phone when it rings). At the contextual control

level, the stimulus-response link depends on the current context (not picking up the phone

when it rings at your friend's home). Episodic and branching control levels enable exceptions

to the rules under which the current contextual rules can be temporarily and conditionally

withheld or reinstated (not picking up the phone at a friend's place unless he asked you to).

Koechlin and Summerfield note that all dual-task situations require such high-level

reconfigurations of rules (i.e., task 2 requires temporarily withholding the rulesets of task 1).

Thus, when conceived from the perspective of this framework, the interference between L1

and L2 can occur at the level of rulesets that are managed by these more-abstract levels of

cognitive control. The interference occurs between the sets of task rules for speaking in L1

and the sets of task rules for speaking in L2, both of which compete for selection to guide

language production. If the interference indeed occurs at the language task-schema level, it

would explain why language representations, supported and processed by the language

network, are not affected by a prior use of L2, yet it usually slows down the naming process

(Branzi et al., 2014; Wodniecka et al., 2020, Casado et al., 2022; Degani et al., 2019). We

therefore propose that the activation within the Multiple Demand network may reflect the

costs of resolving the interference between L1 and L2 production task schemas.

This idea is in line with the Inhibitory Control Hypothesis (Green, 1998), which

proposed that language-switching effects are a consequence of competition between language

schemas. However, the Inhibitory Control Hypothesis did not clearly define "language

schemas": they are sometimes described at a more abstract level as sets of rules that guide

speech production in different languages, similar to what we propose here; and at other times,

they are framed as "regulating the outputs from the lexico-semantic system by altering the

activation levels of representations within that system and by inhibiting outputs from that

system" (Green, 1998), which implies that they engage in resolution of interference between

language-specific representations.

Our proposal is also relevant to a distinction made in the literature regarding bilingual

cognitive control, which differentiates between local and global language control. According

to this distinction, bilingual language control mechanisms (for example, inhibition) can target

not only specific lexical representations (local control) but also the entire language system

(global control: Branzi et al., 2014; Van Assche et al., 2013; Wodniecka et al., 2020). The

idea that difficulty in bilingual speech production can be driven by increased interference

between sets of rules that guide speech production in a given language aligns with the idea of

global language control as mechanisms that operate not on language-specific representations

(e.g., lemmas, c.f. Van Assche et al., 2013; or nodes linked to a common language node;

Dijkstra & van Heuven, 2001; Blanco-Elorrieta & Caramazza, 2021) but on sets of schemas

for speaking in a given language.

In sum, while our first tentative explanation of the L2 after-effect builds on a

mechanism that has been proposed before (proactive interference during L2 naming), it also

points to a critical difference. The key novelty of the proposed interpretation is that the

affected representations are not language-specific (lexical units, morphosyntactic units,

phonological representations, motor programs) but more general and abstract task-schemas

that determine the general programs of using L1 or L2.

The L2-after effect as sustained activation of cognitive control

Another possibility that, to the best of our knowledge, has not been proposed in the

bilingual language processing literature attributes the after-effects in blocked designs entirely

to cognitive control mechanisms without involving any representations at the level of either

language or task-schema. This proposal is similar to the first explanation, because it also

assumes that a prior speech production task in L2 requires intensified proactive cognitive

control to resolve the interference between stronger L1 and weaker L2. However, unlike the

first explanation, here we propose that the prior use of L2 leads to increased activation of

cognitive control without ever impacting language-specific representations. Instead, the

difficult task of speaking in L2 engages cognitive control so much that the control system

remains alert even after the task in L2 concludes. We propose that this increased alertness

might be achieved by increased engagement of the performance or error-monitoring systems

in the brain. As such, the execution of any task (even a non-linguistic one) would proceed in a

more controlled way leading to an increase in reaction times. Support for the explanation

assuming that the L2 after-effect reflects persisting increased engagement of cognitive control

may be found in the Dual Mechanisms of Control framework (Braver, 2021). According to

this framework, proactive control recruits the lateral prefrontal cortex, which, unlike the

anterior cingulate cortex, is sensitive not only to errors or conflicts in the previous trial but

also to control demands specific to the entire task, or the current communicative context

(Blanco-Elorrieta & Caramazza, 2021, Casado et al., 2022). Since we did not observe

modulations between the L1-after and L2-after conditions in the anterior cingulate cortex

activity (see *Supplementary Materials*) but in the lateral prefrontal cortex, which is part of the

Multiple Demand system (by-ROI results in the Multiple Demand network are available in

Supplementary Materials), it can be argued that the L2 after-effect is a manifestation of

proactive control.

In sum, the second tentative explanation of the increased brain response related to the

L2 after-effect proposes that it reflects the increased engagement of cognitive control

triggered by the previous use of L2 and that this engagement lingers during a subsequent task

in L1. However, this hypothesis has never been directly tested and, as such, further
experiments are needed to falsify it.

While the increased effort related to the L2 after-effect observed within the Multiple
Demand network can be putatively explained by increased engagement of proactive
mechanisms engaged in performance or error-monitoring systems, other explanations of the
observed effect cannot be ruled out. One alternative explanation of the increased engagement
of the Multiple Demand network is related to the specificity of the task used in the current
study. We used a picture naming task, which requires producing single words. Previous
studies on monolingual speakers have shown that the Multiple Demand network responds
more strongly to naming objects (i.e., producing isolated words) than to describing events
presented with pictures (i.e., producing full sentences; Hu et al., 2022). Similar effects have
also been found for language comprehension: the Multiple Demand network was shown to
respond more strongly to listening to lists of unconnected words than full sentences (Diachek
et al., 2020). Altogether these results suggests that the Multiple Demand network does not
support the core aspects of language production. This does not rule out the possibility that the
engagement of the domain-general network reflects increased engagement of non-linguistic
control processes in the production of words in L1 after L2 in contrast to L1 after L1;
however, it may suggest that the L2 after-effect measured in a picture naming task may be
"inflated" by the mere demands related to producing single, isolated words. As such, the
effect might reflect a laboratory phenomenon which not necessarily exemplifies cognitive
control mechanisms that bilinguals engage in more natural conversations of everyday
language use. As such, further studies implementing more naturalistic paradigms that require
bilinguals to produce more complex outputs are needed to separate the components driven by
task-related demands from the actual engagement of cognitive control in the L2 after-effect.

Conclusions

The novelty of our contribution is twofold: First, to identify the cognitive mechanisms
that correspond to the neural signature of the L2 after-effect, we used the functional localizer
approach (Fedorenko et al., 2010), which to the best of our knowledge has never been used in
experiments looking at the brain basis of bilingual speech production. Second, unlike previous
studies, our conclusions are based on a design that was optimized to measure the index of
language control, i.e., the L2 after-effect, while reducing many possible confounds. We
believe that the combination of precise analytical tools, carefully designed tasks, as well as

the relatively large number of participants tested (n=42) lead to new insights into the

neurocognitive mechanisms of language control in bilinguals.

In contrast to the dominating theories and views in the field, we found that the L2

after-effect is unlikely to arise as a result of interference between language-specific

representations in L1 and L2 because it does not overlap with the language network, or with

any task-specific localizer that targets specific language processes (lexical access, articulation,

language interference, and inhibition). Instead, we found that the L2 after-effect overlaps with

the Multiple Demand network and thus reflects a domain-general mechanism. We propose

that the L2 after-effect may manifest as either increased interference between high-level task

schemas used to orchestrate language use in L1 and L2, or as a lingering increase in the

engagement of domain-general cognitive control mechanisms in speech production.

Data/code availability statement

The raw neuroimaging data collected for this study along with the data on participants' language experience is available at the OpenNeuro repository https://openneuro.org/datasets/ds004456.

Data and code necessary to reproduce the presented ROI and behavioral analyses is available at the https://osf.io/59za8/?view_only=e23f3500d12e4109b75ca18fdbd320e5.

References

Abutalebi, J., & Green, D. (2007). Bilingual language production: The neurocognition of language representation and control. *Journal of Neurolinguistics*, *20*(3), 242–275. https://doi.org/10.1016/j.jneuroling.2006.10.003

Abutalebi, J., & Green, D. W. (2016). Neuroimaging of language control in bilinguals: Neural adaptation and reserve. *Bilingualism: Language and Cognition*, *19*(4), 689–698. https://doi.org/10.1017/S1366728916000225

Asplund, C. L., Todd, J. J., Snyder, A. P., & Marois, R. (2010). A central role for the lateral prefrontal cortex in goal-directed and stimulus-driven attention. *Nature neuroscience*, *13*(4), 507-512.

Assem, M., Shashidhara, S., Glasser, M. F., & Duncan, J. (2022). Precise topology of adjacent domain-general and sensory-biased regions in the human brain. *Cerebral Cortex*, *32*(12), 2521-2537. https://doi.org/10.1093/cercor/bhab362

Assem, M., Blank, I. A., Mineroff, Z., Ademoğlu, A., & Fedorenko, E. (2020). Activity in the fronto-parietal multiple-demand network is robustly associated with individual differences in working memory and fluid intelligence. *Cortex*, *131*, 1–16. https://doi.org/10.1016/j.cortex.2020.06.013

Assem, M., Glasser, M. F., Van Essen, D. C., & Duncan, J. (2020). A Domain-General Cognitive Core Defined in Multimodally Parcellated Human Cortex. *Cerebral Cortex*, *30*(8), 4361–4380. https://doi.org/10.1093/cercor/bhaa023

Badre, D., & Nee, D. E. (2018). Frontal cortex and the hierarchical control of behavior. *Trends in cognitive sciences*, *22*(2), 170-188.

Basilakos, A., Smith, K. G., Fillmore, P., Fridriksson, J., & Fedorenko, E. (2018). Functional Characterization of the Human Speech Articulation Network. *Cerebral Cortex*, *28*(5), 1816–1830. https://doi.org/10.1093/cercor/bhx100

Bates, D., Kliegl, R., Vasishth, S., & Baayen, H. (2018). Parsimonious mixed models. *arXiv preprint arXiv:1506.04967*. https://doi.org/10.48550/arXiv.1506.04967

Benn, Y., Ivanova, A. A., Clark, O., Mineroff, Z., Seikus, C., Silva, J. S., Varley, R., & Fedorenko, E. (2021). *No evidence for a special role of language in feature-based categorization* (p. 2021.03.18.436075). bioRxiv. https://doi.org/10.1101/2021.03.18.436075

Birn, R. M., Kenworthy, L., Case, L., Caravella, R., Jones, T. B., Bandettini, P. A., & Martin, A. (2010). Neural systems supporting lexical search guided by letter and semantic category cues: A self-paced overt response fMRI study of verbal fluency. *NeuroImage*, *49*(1), 1099–1107. https://doi.org/10.1016/j.neuroimage.2009.07.036

Blanco-Elorrieta, E., & Caramazza, A. (2021). A common selection mechanism at each linguistic level in bilingual and monolingual language production. *Cognition*, 104625. https://doi.org/10.1016/j.cognition.2021.104625

Blanco-Elorrieta, E., & Pylkkänen, L. (2017). Bilingual Language Switching in the Laboratory versus in the Wild: The Spatiotemporal Dynamics of Adaptive Language Control.

Journal of Neuroscience, *37*(37), 9022–9036. https://doi.org/10.1523/JNEUROSCI.0553-
17.2017

Blank, I., Kanwisher, N., & Fedorenko, E. (2014). A functional dissociation between
language and multiple-demand systems revealed in patterns of BOLD signal fluctuations.
Journal of Neurophysiology, *112*(5), 1105–1118. https://doi.org/10.1152/jn.00884.2013

Bobb, S. C., & Wodniecka, Z. (2013). Language switching in picture naming: What
asymmetric switch costs (do not) tell us about inhibition in bilingual speech planning. *Journal*
of Cognitive Psychology, *25*(5), 568–585. https://doi.org/10.1080/20445911.2013.792822

Branzi, F. M., Della Rosa, P. A., Canini, M., Costa, A., & Abutalebi, J. (2016). Language
Control in Bilinguals: Monitoring and Response Selection. *Cerebral Cortex*, *26*(6), 2367–
2380. https://doi.org/10.1093/cercor/bhv052

Branzi, F. M., Martin, C. D., Abutalebi, J., & Costa, A. (2014). The after-effects of bilingual
language production. *Neuropsychologia*, *52*, 102–116.
https://doi.org/10.1016/j.neuropsychologia.2013.09.022

Braver, T. S., Kizhner, A., Tang, R., Freund, M. C., & Etzel, J. A. (2021). The dual
mechanisms of cognitive control project. *Journal of Cognitive Neuroscience*, *33*(9), 1990-
2015.

Braver, T. S., Reynolds, J. R., & Donaldson, D. I. (2003). Neural Mechanisms of Transient
and Sustained Cognitive Control during Task Switching. *Neuron*, *39*(4), 713–726.
https://doi.org/10.1016/S0896-6273(03)00466-5

Broos, W. P. J., Duyck, W., & Hartsuiker, R. J. (2018). Are higher-level processes delayed in
second language word production? Evidence from picture naming and phoneme monitoring.
Language, Cognition and Neuroscience, *33*(10), 1219–1234.
https://doi.org/10.1080/23273798.2018.1457168

Calabria, M., Costa, A., Green, D. W., & Abutalebi, J. (2018). Neural basis of bilingual
language control. *Annals of the New York Academy of Sciences*, *1426*(1), 221–235.
https://doi.org/10.1111/nyas.13879

Casado, A., Szewczyk, J., Wolna, A., & Wodniecka, Z. (2022). The relative balance between
languages predicts the degree of engagement of global language control. *Cognition*, *226*,
105169.

Chein, J. M., & Schneider, W. (2005). Neuroimaging studies of practice-related change:
FMRI and meta-analytic evidence of a domain-general control network for learning.
Cognitive Brain Research, *25*(3), 607–623. https://doi.org/10.1016/j.cogbrainres.2005.08.013

Chen, X., Affourtit, J., Ryskin, R., Regev, T. I., Norman-Haignere, S., Jouravlev, O., Malik-
Moraleda, S., Kean, H., Varley, R., & Fedorenko, E. (2021). *The human language system*
does not support music processing (p. 2021.06.01.446439). bioRxiv.
https://doi.org/10.1101/2021.06.01.446439

Coderre, E. L., Smith, J. F., Heuven, W. J. B. V., & Horwitz, B. (2016). The functional
overlap of executive control and language processing in bilinguals. *Bilingualism: Language*
and Cognition, *19*(3), 471–488. https://doi.org/10.1017/S1366728915000188

Cole, M. W., Reynolds, J. R., Power, J. D., Repovs, G., Anticevic, A., & Braver, T. S. (2013). Multi-task connectivity reveals flexible hubs for adaptive task control. *Nature neuroscience*, *16*(9), 1348-1355.

Cole, M. W., & Schneider, W. (2007). The cognitive control network: Integrated cortical regions with dissociable functions. *NeuroImage*, *37*(1), 343–360. https://doi.org/10.1016/j.neuroimage.2007.03.071

Corbetta, M., Patel, G., & Shulman, G. L. (2008). The reorienting system of the human brain: from environment to theory of mind. *Neuron*, *58*(3), 306-324.

Corbetta, M., & Shulman, G. L. (2002). Control of goal-directed and stimulus-driven attention in the brain. *Nature reviews neuroscience*, *3*(3), 201-215.

Costa, A. (2005). Lexical access in bilingual production. In J. F. Kroll & A. M. B. De Groot (Eds.), *Handbook of bilingualism: Psycholinguistic approaches* (pp. 308–325). Oxford University Press.

Costa, A., Miozzo, M., & Caramazza, A. (1999). Lexical Selection in Bilinguals: Do Words in the Bilingual's Two Lexicons Compete for Selection? *Journal of Memory and Language*, *41*(3), 365–397. https://doi.org/10.1006/jmla.1999.2651

Costa, A., & Santesteban, M. (2004). Lexical access in bilingual speech production: Evidence from language switching in highly proficient bilinguals and L2 learners. *Journal of Memory and Language*, *50*(4), 491–511. https://doi.org/10.1016/j.jml.2004.02.002

Crittenden, B. M., & Duncan, J. (2014). Task Difficulty Manipulation Reveals Multiple Demand Activity but no Frontal Lobe Hierarchy. *Cerebral Cortex*, *24*(2), 532–540. https://doi.org/10.1093/cercor/bhs333

Dale, A. M. (1999). Optimal experimental design for event‐related fMRI. *Human brain mapping*, *8*(2‐3), 109-114.

De Baene, W., Duyck, W., Brass, M., & Carreiras, M. (2015). Brain Circuit for Cognitive Control Is Shared by Task and Language Switching. *Journal of Cognitive Neuroscience*, *27*(9), 1752–1765. https://doi.org/10.1162/jocn_a_00817

Declerck, M., & Koch, I. (2022). The concept of inhibition in bilingual control. *Psychological Review*. https://doi.org/10.1037/rev0000367

Declerck, M., Kleinman, D., & Gollan, T. H. (2020). Which bilinguals reverse language dominance and why? *Cognition*, *204*, 104384. https://doi.org/10.1016/j.cognition.2020.104384

Declerck, M., & Philipp, A. M. (2015). A review of control processes and their locus in language switching. *Psychonomic Bulletin & Review*, *22*(6), 1630–1645. https://doi.org/10.3758/s13423-015-0836-1

Degani, T., Kreiner, H., Ataria, H., & Khateeb, F. (2019). The impact of brief exposure to the second language on native language production: Global or item specific? *Applied Psycholinguistics*, 1–31. https://doi.org/10.1017/S0142716419000444

DeLuca, V., Rothman, J., Bialystok, E., & Pliatsikas, C. (2019). Redefining bilingualism as a spectrum of experiences that differentially affects brain structure and function. *Proceedings of the National Academy of Sciences*, *116*(15), 7565–7574. https://doi.org/10.1073/pnas.1811513116

DeLuca, V., Rothman, J., Bialystok, E., & Pliatsikas, C. (2020). Duration and extent of bilingual experience modulate neurocognitive outcomes. *NeuroImage*, *204*, 116222. https://doi.org/10.1016/j.neuroimage.2019.116222

Diachek, E., Blank, I., Siegelman, M., Affourtit, J., & Fedorenko, E. (2020). The Domain-General Multiple Demand (MD) Network Does Not Support Core Aspects of Language Comprehension: A Large-Scale fMRI Investigation. *The Journal of Neuroscience*, *40*(23), 4536–4550. https://doi.org/10.1523/JNEUROSCI.2036-19.2020

Diamond, A. (2013). Executive functions. *Annual review of psychology*, *64*, 135-168. https://doi.org/10.1146/annurev-psych-113011-143750

Dijkstra, T., & Van Heuven, W. J. (2002). The architecture of the bilingual word recognition system: From identification to decision. *Bilingualism: Language and cognition*, *5*(3), 175-197.

Dosenbach, N. U. F., Fair, D. A., Miezin, F. M., Cohen, A. L., Wenger, K. K., Dosenbach, R. A. T., Fox, M. D., Snyder, A. Z., Vincent, J. L., Raichle, M. E., Schlaggar, B. L., & Petersen, S. E. (2007). Distinct brain networks for adaptive and stable task control in humans. *Proceedings of the National Academy of Sciences*, *104*(26), 11073–11078. https://doi.org/10.1073/pnas.0704320104

Dosenbach, N. U. F., Visscher, K. M., Palmer, E. D., Miezin, F. M., Wenger, K. K., Kang, H. C., Burgund, E. D., Grimes, A. L., Schlaggar, B. L., & Petersen, S. E. (2006). A Core System for the Implementation of Task Sets. *Neuron*, *50*(5), 799–812. https://doi.org/10.1016/j.neuron.2006.04.031

Duncan, J. (2010). The multiple-demand (MD) system of the primate brain: Mental programs for intelligent behaviour. *Trends in Cognitive Sciences*, *14*(4), 172–179. https://doi.org/10.1016/j.tics.2010.01.004

Fedeli, D., Del Maschio, N., Sulpizio, S., Rothman, J., & Abutalebi, J. (2021). The bilingual structural connectome: Dual-language experiential factors modulate distinct cerebral networks. *Brain and Language*, *220*, 104978. https://doi.org/10.1016/j.bandl.2021.104978

Fedorenko, E. (2021). The early origins and the growing popularity of the individual-subject analytic approach in human neuroscience. *Current Opinion in Behavioral Sciences*, *40*, 105–112. https://doi.org/10.1016/j.cobeha.2021.02.023

Fedorenko, E., Behr, M. K., & Kanwisher, N. (2011). Functional specificity for high-level linguistic processing in the human brain. *Proceedings of the National Academy of Sciences*, *108*(39), 16428–16433. https://doi.org/10.1073/pnas.1112937108

Fedorenko, E., Duncan, J., & Kanwisher, N. (2012). Language-Selective and Domain-General Regions Lie Side by Side within Broca's Area. *Current Biology*, *22*(21), 2059–2062. https://doi.org/10.1016/j.cub.2012.09.011

Fedorenko, E., Duncan, J., & Kanwisher, N. (2013). Broad domain generality in focal regions of frontal and parietal cortex. *Proceedings of the National Academy of Sciences*, *110*(41), 16616–16621. https://doi.org/10.1073/pnas.1315235110

Fedorenko, E., Hsieh, P.-J., Nieto-Castañón, A., Whitfield-Gabrieli, S., & Kanwisher, N. (2010). New Method for fMRI Investigations of Language: Defining ROIs Functionally in Individual Subjects. *Journal of Neurophysiology*, *104*(2), 1177–1194. https://doi.org/10.1152/jn.00032.2010

Friesen, D. C., Edwards, K., & Lamoureux, C. (2021). Predictors of verbal fluency performance in monolingual and bilingual children: The interactive role of English receptive vocabulary and fluid intelligence. *Journal of Communication Disorders*, *89*, 106074. https://doi.org/10.1016/j.jcomdis.2020.106074

Frost, M. A., & Goebel, R. (2012). Measuring structural–functional correspondence: Spatial variability of specialised brain regions after macro-anatomical alignment. *NeuroImage*, *59*(2), 1369–1381. https://doi.org/10.1016/j.neuroimage.2011.08.035

Gollan, T. H., & Ferreira, V. S. (2009). Should I stay or should I switch? A cost–benefit analysis of voluntary language switching in young and aging bilinguals. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *35*(3), 640–665. https://doi.org/10.1037/a0014981

Gollan, T. H., Kleinman, D., & Wierenga, C. E. (2014). What's easier: Doing what you want, or being told what to do? Cued versus voluntary language and task switching. *Journal of Experimental Psychology: General*, *143*(6), 2167–2195. https://doi.org/10.1037/a0038006

Green, D. W. (1998). Mental control of the bilingual lexico-semantic system. *Bilingualism: Language and Cognition*, *1*(02), 67. https://doi.org/10.1017/S1366728998000133

Green, D. W., & Abutalebi, J. (2013). Language control in bilinguals: The adaptive control hypothesis. *Journal of Cognitive Psychology*, *25*(5), 515–530. https://doi.org/10.1080/20445911.2013.796377

Guo, T., Liu, H., Misra, M., & Kroll, J. F. (2011). Local and global inhibition in bilingual word production: FMRI evidence from Chinese–English bilinguals. *NeuroImage*, *56*, 2300–2309.

Haman, E., Łuniewska, M., & Pomiechowska, B. (2015). Designing Cross-Linguistic Lexical Tasks (CLTs) for Bilingual Preschool Children. In *Assessing Multilingual Children* (pp. 196–240). Multilingual Matters. https://www.degruyter.com/document/doi/10.21832/9781783093137-010/html

Hu, J., Small, H., Kean, H., Takahashi, A., Zekelman, L., Kleinman, D., Ryan, E., Ferreira, V., & Fedorenko, E. (2022). Precision fMRI reveals that the language-selective network supports both phrase-structure building and lexical access during language production. *Cerebral Cortex*, *1*, 21.

Ivanova, A. A., Mineroff, Z., Zimmerer, V., Kanwisher, N., Varley, R., & Fedorenko, E. (2021). The Language Network Is Recruited but Not Required for Nonverbal Event Semantics. *Neurobiology of Language*, *2*(2), 176–201. https://doi.org/10.1162/nol_a_00030

Jenkinson, M., Bannister, P., Brady, M., & Smith, S. (2002). Improved optimization for the
robust and accurate linear registration and motion correction of brain
images. *Neuroimage*, *17*(2), 825-841.

Jenkinson, M., & Smith, S. (2001). A global optimisation method for robust affine
registration of brain images. *Medical image analysis*, *5*(2), 143-156.

Koch, I., Gade, M., Schuch, S., & Philipp, A. M. (2010). The role of inhibition in task
switching: A review. *Psychonomic Bulletin & Review*, *17*(1), 1–14.
https://doi.org/10.3758/PBR.17.1.1

Koechlin, E., & Summerfield, C. (2007). An information theoretical approach to prefrontal
executive function. *Trends in cognitive sciences*, *11*(6), 229-235.

Kroll, J. F., Bobb, S. C., & Wodniecka, Z. (2006). Language selectivity is the exception, not
the rule: Arguments against a fixed locus of language selection in bilingual speech.
Bilingualism: Language and Cognition, *9*(2), 119–135.
https://doi.org/10.1017/S1366728906002483

Kuznetsova, A., Brockhoff, P.B., Christensen, R.H.B. (2017). "lmerTest Package: Tests in
Linear Mixed Effects Models." *Journal of Statistical Software*, 82(13), 1–
26. https://doi:10.18637/jss.v082.i13

La Heij, W. (2005). Selection process in monolingual and bilingual lexical access. In J. F.
Kroll & A. M. B. De Groot (Eds.), *Handbook of bilingualism: Psycholinguistic approaches*
(pp. 289–307). Oxford University Press.

Lemhöfer, K., & Broersma, M. (2012). Introducing LexTALE: A quick and valid lexical test
for advanced learners of English. *Behavior research methods*, *44*, 325-343.

Liljeström, M., Kujala, J., Stevenson, C., & Salmelin, R. (2015). Dynamic reconfiguration of
the language network preceding onset of speech in picture naming. *Human Brain Mapping*,
36(3), 1202–1216. https://doi.org/10.1002/hbm.22697

Luk, G., Green, D. W., Abutalebi, J., & Grady, C. (2012). Cognitive control for language
switching in bilinguals: A quantitative meta-analysis of functional neuroimaging studies.
Language and Cognitive Processes, *27*(10), 1479–1488.
https://doi.org/10.1080/01690965.2011.613209

Malik-Moraleda, S., Ayyash, D., Gallée, J., Affourtit, J., Hoffmann, M., Mineroff, Z.,
Jouravlev, O., & Fedorenko, E. (2022). An investigation across 45 languages and 12 language
families reveals a universal language network. *Nature Neuroscience*, *25*(8), 1014-1019.

Malik-Moraleda, S., Cucu, T., Lipkin, B., & Fedorenko, E. (2021). The Domain-General
Multiple Demand Network Is More Active in Early Balanced Bilinguals Than Monolinguals
During Executive Processing. *Neurobiology of Language*, *2*(4), 647–664.
https://doi.org/10.1162/nol_a_00058

Martin, A., Wiggs, C. L., Lalonde, F., & Mack, C. (1994). Word retrieval to letter and
semantic cues: A double dissociation in normal subjects using interference tasks.
Neuropsychologia, *32*(12), 1487–1494. https://doi.org/10.1016/0028-3932(94)90120-1

Mendoza, M. N., Blumenfeld, H. K., Knight, R. T., & Ries, S. K. (2021). Investigating the Link Between Linguistic and Non-Linguistic Cognitive Control in Bilinguals Using Laplacian-Transformed Event Related Potentials. *Neurobiology of Language*, *2*(4), 605–627. https://doi.org/10.1162/nol_a_00056

Meuter, R. F. I., & Allport, A. (1999). Bilingual Language Switching in Naming: Asymmetrical Costs of Language Selection. *Journal of Memory and Language*, *40*(1), 25–40. https://doi.org/10.1006/jmla.1998.2602

Michon, K. J., Khammash, D., Simmonite, M., Hamlin, A. M., & Polk, T. A. (2022). Person-specific and precision neuroimaging: Current methods and future directions. *NeuroImage*, *263*, 119589. https://doi.org/10.1016/j.neuroimage.2022.119589

Miller, E. K., & Cohen, J. D. (2001). An Integrative Theory of Prefrontal Cortex Function. *Annual Review of Neuroscience*, *24*(1), 167–202. https://doi.org/10.1146/annurev.neuro.24.1.167

Misra, M., Guo, T., Bobb, S. C., & Kroll, J. F. (2012). When bilinguals choose a single word to speak: Electrophysiological evidence for inhibition of the native language. *Journal of Memory and Language*, *67*(1), 224–237. https://doi.org/10.1016/j.jml.2012.05.001

Naylor, L., Stanley, E., & Wicha, N. (2012). Cognitive and Electrophysiological Correlates of the Bilingual Stroop Effect. *Frontiers in Psychology*, *3*. https://www.frontiersin.org/articles/10.3389/fpsyg.2012.00081

Nee, D. E., Jonides, J., & Berman, M. G. (2007). Neural mechanisms of proactive interference-resolution. *NeuroImage*, *38*(4), 740–751. https://doi.org/10.1016/j.neuroimage.2007.07.066

Niendam, T. A., Laird, A. R., Ray, K. L., Dean, Y. M., Glahn, D. C., & Carter, C. S. (2012). Meta-analytic evidence for a superordinate cognitive control network subserving diverse executive functions. *Cognitive, Affective, & Behavioral Neuroscience*, *12*(2), 241–268. https://doi.org/10.3758/s13415-011-0083-5

Pallier, C., Dehaene, S., Poline, J. B., LeBihan, D., Argenti, A. M., Dupoux, E., & Mehler, J. (2003). Brain imaging of language plasticity in adopted adults: Can a second language replace the first?. *Cerebral cortex*, *13*(2), 155-161.

Poldrack, R. A. (2006). Can cognitive processes be inferred from neuroimaging data? *Trends in Cognitive Sciences*, *10*(2), 59–63. https://doi.org/10.1016/j.tics.2005.12.004

R Core Team. (2020). R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL https://www.R-project.org/

Reverberi, C., Kuhlen, A. K., Seyed-Allaei, S., Greulich, R. S., Costa, A., Abutalebi, J., & Haynes, J.-D. (2018). The neural basis of free language choice in bilingual speakers: Disentangling language choice and language execution. *NeuroImage*, *177*, 108–116. https://doi.org/10.1016/j.neuroimage.2018.05.025

Rossi, E., Newman, S., Kroll, J. F., & Diaz, M. T. (2018). Neural signatures of inhibitory control in bilingual spoken production. *Cortex*, *108*, 50–66. https://doi.org/10.1016/j.cortex.2018.07.009

Runnqvist, E., Strijkers, K., Alario, F.-X., & Costa, A. (2012). Cumulative semantic interference is blind to language: Implications for models of bilingual speech production. *Journal of Memory and Language*, *66*(4), 850–869. https://doi.org/10.1016/j.jml.2012.02.007

Scott, T. L., Gallée, J., & Fedorenko, E. (2017). A new fun and robust version of an fMRI localizer for the frontotemporal language system. *Cognitive Neuroscience*, *8*(3), 167–176. https://doi.org/10.1080/17588928.2016.1201466

Shain, C., Paunov, A., Chen, X., Lipkin, B., & Fedorenko, E. (2022). No evidence of theory of mind reasoning in the human language network. *bioRxiv*, 2022-07.

Shao, Z., Janse, E., Visser, K., & Meyer, A. S. (2014). What do verbal fluency tasks measure? Predictors of verbal fluency performance in older adults. *Frontiers in Psychology*, *5*. https://doi.org/10.3389/fpsyg.2014.00772

Shashidhara, S., Mitchell, D. J., Erez, Y., & Duncan, J. (2019). Progressive Recruitment of the Frontoparietal Multiple-demand System with Increased Task Complexity, Time Pressure, and Reward. *Journal of Cognitive Neuroscience*, *31*(11), 1617–1630. https://doi.org/10.1162/jocn_a_01440

Smith, D. M., Perez, D. C., Porter, A., Dworetsky, A., & Gratton, C. (2021). Light through the fog: Using precision fMRI data to disentangle the neural substrates of cognitive control. *Current Opinion in Behavioral Sciences*, *40*, 19–26. https://doi.org/10.1016/j.cobeha.2020.12.004

Stroop, J. R. (1935). Studies of interference in serial verbal reactions. *Journal of experimental psychology*, *18*(6), 643.

Sulpizio, S., Del Maschio, N., Fedeli, D., & Abutalebi, J. (2020). Bilingual language processing: A meta-analysis of functional neuroimaging studies. *Neuroscience & Biobehavioral Reviews*, *108*, 834–853. https://doi.org/10.1016/j.neubiorev.2019.12.014

Tahmasebi, A. M., Davis, M. H., Wild, C. J., Rodd, J. M., Hakyemez, H., Abolmaesumi, P., & Johnsrude, I. S. (2012). Is the Link between Anatomical Structure and Function Equally Strong at All Cognitive Levels of Processing? *Cerebral Cortex*, *22*(7), 1593–1603. https://doi.org/10.1093/cercor/bhr205

Tzelgov, J., Henik, A., & Berger, J. (1992). Controlling Stroop effects by manipulating expectations for color words. *Memory & Cognition*, *20*(6), 727–735. https://doi.org/10.3758/BF03202722

Unsworth, N., Spillers, G. J., & Brewer, G. A. (2011). Variation in verbal fluency: A latent variable analysis of clustering, switching, and overall performance. *Quarterly Journal of Experimental Psychology*, *64*(3), 447–466. https://doi.org/10.1080/17470218.2010.505292

Van Assche, E., Duyck, W., & Gollan, T. H. (2013). Whole-language and item-specific control in bilingual language production. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *39*(6), 1781–1792. https://doi.org/10.1037/a0032859

Verhoef, K., Roelofs, A., & Chwilla, D. J. (2009). Role of inhibition in language switching: Evidence from event-related brain potentials in overt picture naming. *Cognition*, *110*(1), 84–99. https://doi.org/10.1016/j.cognition.2008.10.013

Weissberger, G. H., Gollan, T. H., Bondi, M. W., Clark, L. R., & Wierenga, C. E. (2015).
Language and task switching in the bilingual brain: Bilinguals are staying, not switching,
experts. *Neuropsychologia*, *66*, 193–203.
https://doi.org/10.1016/j.neuropsychologia.2014.10.037

Wodniecka, Z., Casado, A., Kałamała, P., Marecka, M., Timmer, K., & Wolna, A. (2020).
Chapter Eight—The dynamics of language experience and how it affects language and
cognition. In K. D. Federmeier & H.-W. Huang (Eds.), *Psychology of Learning and*
Motivation (Vol. 72, pp. 235–281). Academic Press.
https://doi.org/10.1016/bs.plm.2020.02.005

Wodniecka, Z., Szewczyk, J., Kałamała, P., Mandera, P., & Durlik, J. (2020). When a second
language hits a native language. What ERPs (do and do not) tell us about language retrieval
difficulty in bilingual language production. *Neuropsychologia*, *141*, 107390.
https://doi.org/10.1016/j.neuropsychologia.2020.107390

Wolna, A., Łuniewska, M., Haman, E., & Wodniecka, Z. (2022). Polish norms for a set of
colored drawings of 168 objects and 146 actions with predictors of naming performance.
Behavior Research Methods. https://doi.org/10.3758/s13428-022-01923-3

Wu, J., Yang, J., Chen, M., Li, S., Zhang, Z., Kang, C., Ding, G., & Guo, T. (2019). Brain
network reconfiguration for language and domain-general cognitive control in bilinguals.
NeuroImage, *199*, 454–465. https://doi.org/10.1016/j.neuroimage.2019.06.022

Xu, J., Moeller, S., Auerbach, E. J., Strupp, J., Smith, S. M., Feinberg, D. A., Yacoub, E., &
Uğurbil, K. (2013). Evaluation of slice accelerations using multiband echo planar imaging at
3 T. *Neuroimage, 83,* 991-1001. https://doi.org/10.1016/j.neuroimage.2013.07.055

Zhang, Y., Wang, T., Huang, P., Li, D., Qiu, J., Shen, T., & Xie, P. (2015). Free Language
Selection in the Bilingual Brain: An Event-Related fMRI Study. *Scientific Reports*, *5*(1).
https://doi.org/10.1038/srep11704