



Using neurostimulation to understand the impact of pre-morbid individual differences on post-lesion outcomes

Anna M. Woollams^{a,1}, Gaston Madrid^a, and Matthew A. Lambon Ralph^a

^aNeuroscience and Aphasia Research Unit, Division of Neuroscience and Experimental Psychology, School of Biological Sciences, University of Manchester, Manchester, M13 9PL, England

Edited by Robert D. Rafal, University of Delaware, Newark, DE, and accepted by Editorial Board Member Michael S. Gazzaniga September 6, 2017 (received for review May 4, 2017)

Neuropsychological data have proven invaluable in advancing our understanding of higher cognition. The interpretation of such data is, however, complicated by the fact that post-lesion behavioral abnormalities could reflect pre-morbid individual differences in the cognitive domain of interest. Here we exploited the virtual lesion methodology offered by transcranial magnetic stimulation (TMS) to explore the impact of pre-morbid individual differences on post-lesion performance. We applied this approach to the domain of reading, a crucial ability in which there are known to be considerable individual differences in the normal population. As predicted by neuropsychological studies of surface dyslexia in semantic dementia and the connectionist triangle model of reading, previous empirical work has shown that healthy participants vary in their reliance on meaning for reading words with atypical correspondences between spelling and sound. We therefore selected participants who varied along this dimension and applied a virtual lesion to the left anterior temporal lobe. As expected, we observed a significant three-way interaction between “pre-morbid” reading status, stimulation, and word type, such that TMS increased the disadvantage for spelling-sound atypical words more for the individuals with stronger semantic reliance. This successful test-case study provides an approach to understanding the impact of pre-morbid individual variation on post-lesion outcomes that could be fruitfully applied to a variety of cognitive domains.

transcranial magnetic stimulation | anterior temporal lobe | surface dyslexia | semantic dementia | reading

The study of neuropsychological patients to inform models of normal cognitive function has proven highly productive. A major issue, however, that arises in this research is the possibility of systematic individual differences in normal pre-morbid performance impacting significantly on post-damage neuropsychological profiles. Such differences can complicate the interpretation of behavior in terms of its implications for both cognitive and neural models (1), particularly because neuropsychological conclusions are often based on single cases. Estimation of these differences is challenging because they are conflated with the impact of brain damage; hence, one cannot retrospectively assess pre-morbid performance. A three-pronged approach is needed to establish and understand the impact of these individual differences. Firstly, detailed data on the full spectrum of patients’ performance is needed from large-scale case-series studies. Secondly, computational models that implement the core cognitive framework and incorporate an account of individual variation are required. Finally, new approaches, such as neurostimulation techniques that can transiently mimic patient performance, are necessary to relate pre-morbid individual differences directly to post-damage performance. In this paper, we illustrate the key challenges and potential solutions to the issue of pre-morbid individual differences within the domain of reading aloud, where there is already extensive neuropsychological data linking semantic deficits to surface dyslexia and an implemented computational model of individual differences in semantic

reliance (SR) in normal reading and its post-damage consequences. Until now, however, there has not been a direct test of the model’s hypothesis, and here we provide this by exploiting the “virtual lesion” approach offered by repetitive transcranial magnetic stimulation (rTMS).

Reading is a particularly mature and sophisticated area of research because competing alternative accounts have been implemented as computational models, allowing direct quantitative simulation of behavior. It is also a key ability in which there are considerable individual differences among the normal population (2). Reading aloud in English constitutes a quasi-regular domain (3), such that most words have predictable pronunciations according to the most typical subword correspondences between spelling and sound (e.g., *pink*, *scribe*), but there are nevertheless many words that contain atypical correspondences that render their pronunciations unpredictable (e.g., *pint*, *scarce*). Current models of reading vary in how they propose the human reading systems deals with this quasi-regularity (3, 4).

The connectionist triangle model proposes a direct subword pathway between spelling and sound that contains representations of orthography and phonology distributed over a set of units (3, 5, 6). These units are linked by connections, with the weights on these learned through probabilistic exposure on a representative corpus. When trained in isolation, this pathway can successfully translate

Significance

Data from patients with brain damage have provided unique insights into the neural bases of cognitive function. Yet interpretation of patient data is complicated by the possible influence of pre-morbid individual differences on performance. We addressed this issue by considering the impact of transcranial magnetic stimulation (TMS)-induced “virtual lesions” of the left anterior temporal lobe upon reading performance in healthy individuals who vary in their degree of semantic reliance during reading. TMS only disrupted performance in higher semantic reliance readers. These results establish a direct link between pre-morbid individual differences and post-damage outcomes. Our virtual lesion approach provides a methodology through which the impact of pre-morbid individual differences can be examined and their theoretical implications understood.

Author contributions: A.M.W. and M.A.L.R. designed research; A.M.W. and G.M. performed research; A.M.W. and G.M. analyzed data; and A.M.W. and M.A.L.R. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission. R.D.R. is a guest editor invited by the Editorial Board.

This is an open access article distributed under the PNAS license.

¹To whom correspondence should be addressed. Email: anna.woollams@manchester.ac.uk.

This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1707162114/-DCSupplemental.

all known words and also novel letter strings (3). However, when the model is trained with an additional semantic pathway to emulate access to word meaning, then a division of labor emerges such that this whole-word information comes to support pronunciation of atypical words, allowing the direct pathway to partially specialize to typical subword mappings (3, 5, 7, 8). Evidence for this account is provided by reports of stronger semantic effects for reading aloud atypical than typical words (9–13), which has been simulated in the connectionist triangle model (5). In this model then, semantic damage results in deficits in reading atypical words, a pattern called surface dyslexia (3, 7).

The connectionist triangle model account therefore makes the strong prediction that semantic deficits should compromise reading of words with atypical spelling–sound mappings (3). This prediction is supported by the observation that the vast majority of patients with semantic dementia, a progressive and selective deterioration of semantic memory due to atrophy and hypometabolism of the anterior temporal lobes (ATLs) (14, 15), show a reading profile of surface dyslexia (16, 17). The causes of this association have been hotly debated, however, because there have been a small minority of cases of semantic dementia who show preserved atypical word reading (18–21).

The connectionist triangle model account proposes that, in addition to degree of semantic damage, a key factor in explaining variation in the degree of surface dyslexia seen in semantic dementia is pre-morbid individual differences in the degree of semantic involvement in reading aloud (7, 8). Previous work on individual differences in reading styles had been framed in terms of differential reliance on whole-word and subword strategies (22, 23), rather than the involvement of semantic information, and had yielded inconsistent cognitive correlates (2). More recently, Graves et al. (21) have considered individual variation in degree of SR during reading and found this to be related to white matter connectivity within the left hemisphere reading network. The connectionist triangle model account of variation in surface dyslexia in semantic dementia focuses particularly on individual differences in SR specifically for words with atypical spelling–sound mappings.

As noted earlier, to explore the proposal derived from the connectionist triangle model, large-scale case-series neuropsychological data are needed to quantify the strength of the relationship between semantic and reading impairments and the incidence of exceptional cases and more broadly to establish the scale of variation around the central tendency of the association. This was provided by the largest case-series study of this issue performance to date, based on 100 observations of reading behavior from 51 semantic dementia patients (8). This study found an extremely strong association between semantic deficits and surface dyslexia, with the degree of semantic impairment accounting for half the variance in patients’ reading of atypical words. While there were three cases showing preserved atypical word reading at initial testing, all progressed to a surface dyslexic reading pattern at follow-up. However, despite the strength of the association between semantics and atypical word reading in this study, there was nevertheless considerable variation in reading performance between patients with the same degree of semantic impairment.

To further our understanding of the role of individual differences, we also need a computational model that implements the role of semantic information in normal reading and incorporates a formal mechanism and simulation of how pre-morbid variation in SR can impact upon post-damage performance. Woollams et al. (8) simulated their case-series data using a sample of different instantiations of the connectionist triangle model that varied in the degree of semantic support provided during training of the direct pathway. For models trained with strong semantic support, lesioning in the semantic pathway had a marked negative impact on atypical word reading. For models trained with weak semantic support, lesioning had a much milder effect on atypical word reading. In other words, for the more semantic reading models,

little damage was needed before surface dyslexia emerged, while for the less semantic reading models, greater damage was required to produce a surface dyslexic reading pattern. Incorporating individual differences in this manner allowed the model to capture over 90% of the variance in the patient data (8).

Although this individual differences account is a plausible one, validation is needed. Specifically, by definition, it is not possible to ascertain the *pre-morbid* reading performance of the semantic dementia patients, yet the account predicts that (i) there are individual differences in SR for atypical word reading among healthy participants and (ii) these will have differential consequences if the neural system for semantic representation is compromised. The virtual lesion methodology offered by rTMS (24) provides a unique way to explore the impact of transient disruption to specific neural regions in healthy participants. Across multiple studies, performance on various verbal and nonverbal semantic tasks has been shown to be disrupted by offline rTMS to a specific area of the left ATL, the anterior middle temporal gyrus (MTG), mimicking in reaction time the deficits observed in semantic dementia (25–27).

In this study, we exploit the virtual lesion methodology to understand the impact of pre-morbid individual differences on post-damage performance. This approach allows us to circumvent concerns about variation in the location/extent of damage and also in adaptation to damage, both of which are inherent in neuropsychological research. We applied rTMS to the left ATL of normal readers to determine how variation in SR for atypical word reading impacts upon deficits after disruption. We selected normal readers for their low or high SR using a behavioral measure previously shown to predict the size of imageability and semantic priming effects in reading (2). We therefore used rTMS to target the anterior MTG of low and high SR readers for virtual lesioning. The individual differences hypothesis predicts that the disadvantage for atypical words should be increased by ATL rTMS more for the high SR readers than the low SR readers and that this pattern should be particularly apparent for low-frequency words.

Results

An ANOVA on reaction time data for low-frequency words (Table 1) with reader type (low SR/high SR) as a between-participants variable and rTMS (pre/post) and typicality (typical/atypical) as within-participant variables revealed that the critical expected three-way interaction between reader type, stimulation, and typicality was significant, $F(1, 16) = 0.23, P = 0.036$. There was also a two-way interaction between typicality and reader type, $F(1, 16) = 4.90, P = 0.042$, a significant main effect of typicality, $F(1, 16) = 35.15, P < 0.0005$, and a marginally significant main effect of reader type, $F(1, 16) = 4.20, P = 0.057$.

An ANOVA on the data for low-frequency words before TMS with reader type (low SR/high SR) as a between-participants variable and typicality (typical/atypical) as a within-participant

Table 1. Millisecond RTs of low and high SR readers on the frequency by typicality reading list completed pre-TMS and post-TMS

Reader type	Frequency	Typicality	Pre-TMS		Post-TMS	
			Mean	SD	Mean	SD
Low SR	High	Typical	502	63	507	67
		Atypical	508	63	508	67
	Low	Typical	519	67	527	79
		Atypical	545	82	540	87
High SR	High	Typical	555	55	543	61
		Atypical	565	62	552	54
	Low	Typical	587	73	578	74
		Atypical	622	67	629	83

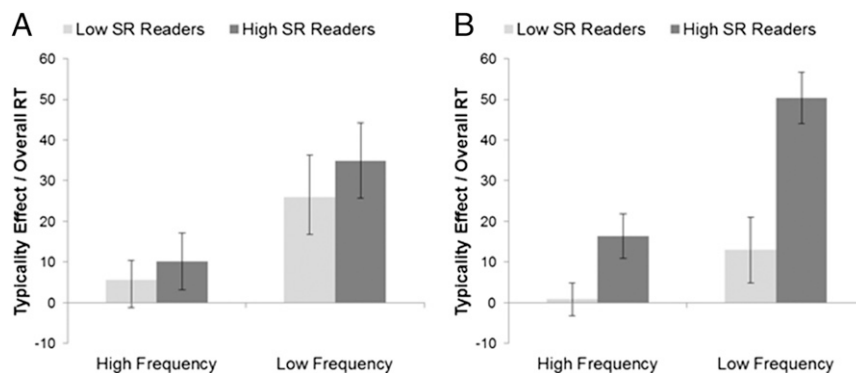


Fig. 1. Performance of low and high SR readers on the frequency by typicality reading list before and after rTMS to the left ATL: (A) pre-stimulation typicality effects and (B) post-stimulation typicality effects. Error bars are \pm SE. Values represent effects adjusted for overall level of performance (difference in values across two conditions divided by the mean over those two conditions).

variable revealed only significant main effects of typicality, $F(1, 16) = 19.31$, $P < 0.0005$, and reader group, $F(1, 16) = 4.72$, $P = 0.045$, with no interaction, $F(1, 16) = 0.42$, $P = 0.528$. After rTMS, there was a significant interaction between typicality and reader group, $F(1, 16) = 13.36$, $P = 0.002$, indicating that the typicality effect for the high SR readers was now significantly larger than that for the low SR readers, as can be seen in Fig. 1. There was also a significant main effect of typicality, $F(1, 16) = 38.21$, $P < 0.0005$, and a trend toward an effect of reader group, $F(1, 16) = 3.41$, $P = 0.083$. Follow-up paired t tests demonstrated that the typicality effect for low-frequency words after TMS was not significant for the low SR readers, $t(1, 8) = 1.6$, $P = 0.148$, but was highly significant for the high SR readers, $t(1, 8) = 8.0$, $P < 0.0005$.

This pattern of results was confirmed by using regression to consider the extent to which degree of SR, as a continuous variable, predicted the size of typicality effects for low-frequency words. Before rTMS, there was no significant relationship between SR and the typicality effect for low-frequency words, $F(1, 16) = 0.37$, $P = 0.554$. In contrast, after rTMS, there was a significant relationship between SR and the typicality effect for low-frequency words, $F(1, 16) = 7.14$, $P = 0.017$, $R^2 = 31\%$. Essentially, the two groups were reading atypical words in different ways before rTMS, and disruption of semantic processing only affected the high SR readers, producing a relationship between the SR measure and the size of the typicality effect after stimulation.

An ANOVA on reaction time data for high-frequency words (Table 1) with reader type (low SR/high SR) as a between-participants variable and rTMS (pre/post) and typicality (typical/atypical) as within-participant variables revealed only a main effect of typicality, $F(1, 16) = 5.95$, $P = 0.027$.

Results for high- and low-frequency words were compared using an omnibus ANOVA on reaction times, with reader type (low SR/high SR) as a between-participants variable and rTMS (pre/post), typicality (typical/atypical), and frequency (high/low) as within-participant variables, and revealed a marginally significant four-way interaction, $F(1, 16) = 3.83$, $P = 0.068$. This result supports the previous analysis and indicates that, as predicted, the disruptive effect of TMS on atypical word reading for the high SR readers was most apparent for low-frequency words.

An omnibus ANOVA on error rates, presented in Table 2, with reader type (low SR/high SR) as a between-participants variable and TMS (pre/post), typicality (typical/atypical), and frequency (high/low) as within-participant variables revealed only significant effects of typicality, $F(1, 16) = 50.59$, $P < 0.0005$, frequency, $F(1, 16) = 64.54$, $P < 0.0005$, and an interaction between them, $F(1, 16) = 36.18$, $P < 0.0005$. No other effects approached significance.

Discussion

The influence of pre-morbid individual differences on the interpretation of post-damage neuropsychological profiles is a critical issue. Here we addressed this challenge in the domain of reading, where there is already a large body of case-series evidence showing the distribution of reading deficits observed after semantic damage and an implemented computational model that explains the variation in these deficits (7, 8). We employed rTMS to the left ATL to test the hypothesis that individual differences in degree of SR when reading atypical words produces appreciable variation in the impact of a virtual lesion. Before ATL stimulation, the two groups did not differ in their reading performance, whereas after stimulation, the typicality effect for low-frequency words was significantly greater for the high than low SR readers. In addition, the size of the typicality effect for low-frequency words was predicted by the individual differences measure.

The finding that stimulation had no appreciable impact on high-frequency words for any participants illustrates that left ATL rTMS does not interfere with reading in general. It also argues against an account of variation in stimulation effects according to SR in terms of individual differences in global susceptibility to ATL stimulation. The reading deficit we observed for the high SR readers was manifested in reaction times, as is the case for most cognitive TMS studies, including those involving ATL stimulation (25, 27, 28). Although this contrasts with the accuracy effects seen in semantic dementia patients, this is to be expected given that the neural impact of rTMS is more anatomically and temporally punctate than that of any neurodegenerative condition.

The results of the present study are consistent with the neuropsychological data concerning atypical word reading deficits in semantic dementia (8, 29). Structural neuroimaging shows reading deficits to be associated with damage to the left ATL (ref. 30 and

Table 2. Percentage error rates of low and high SR readers on the frequency by typicality reading list completed pre-TMS and post-TMS

Reader type	Frequency	Typicality	Pre-TMS		Post-TMS	
			Mean	SD	Mean	SD
Low SR	High	Typical	0.53	1.68	0.53	1.68
		Atypical	0.53	1.68	1.06	2.20
	Low	Typical	1.59	2.46	1.59	2.46
		Atypical	11.64	7.09	10.58	8.02
High SR	High	Typical	0.53	1.68	1.59	2.46
		Atypical	3.70	3.37	3.17	2.20
	Low	Typical	2.65	2.55	1.59	2.46
		Atypical	11.64	6.23	10.58	5.36

Fig. 2A). Our stimulation target was squarely within the peak region of atrophy in these patients (Fig. 2B). Our results also extend previous work using functional imaging to understand reading in healthy participants, which revealed areas of the anterior MTG and inferior temporal gyrus that were more strongly activated during atypical than typical word reading (31). Strikingly, these areas overlapped with areas more strongly activated for high than low SR readers (Fig. 2C) and intersected with the area we targeted for stimulation in this study. Our work therefore provides crucial evidence that higher left ATL activation in high SR readers plays an appreciable functional role in supporting efficient reading of atypical words. Hence our neurostimulation results provide an important direct link between neuropsychological studies and neuroimaging data concerning individual differences in SR during reading, thereby validating the connectionist triangle model's account of surface dyslexia in semantic dementia.

This study assessed the impact of rTMS of the ATL on reading aloud. Other researchers have explored the effect of online TMS on reading aloud, however they have stimulated different sites within the neural reading network (32–36). The effect of stimulation of left ventral occipito-temporal cortex is of particular relevance given the attribution of reading deficits in semantic dementia to progressive atrophy of this region (37, 38). Pattamadilok et al. (39) reported that online TMS 140 ms after stimulus onset disrupted reading of both typical and atypical words and nonwords when delivered to left ventral occipito-temporal cortex and also disrupted both typical and atypical word reading when delivered to either the left posterior MTG or left supramarginal gyrus (see also ref. 36). To the extent that the impact of online and offline TMS are comparable (cf. refs. 25 and 40), these studies demonstrate that the disruption of left ventral occipito-temporal cortex causes different reading deficits to those reported here after stimulation of the left ATL. This agrees with results of comparative case series contrasting the reading performance of patients with pure alexia from posterior cerebral artery stroke with that of patients with surface dyslexia and semantic dementia (41, 42). Indeed, there have been a number of semantic dementia cases reported who have had marked surface dyslexia without any evidence of hypometabolism extending into the left ventral occipito-temporal cortex (29). Furthermore, recent lesion-symptom mapping work has demonstrated a significant relationship between the regularization errors in reading and the degree of left ATL atrophy among semantic dementia patients (43). Taken together, these results from neurostimulation and neuropsychology combine to support a specific role of the ATL in atypical word reading.

One reason that the necessity of ATL semantic processing in atypical word reading has been a topic of debate is because it is a dimension on which competing computational models of word reading differ markedly (3, 4). The connectionist triangle model proposes that semantic information is necessary for reading aloud atypical words, particularly those low in frequency. In this model, it is the semantic pathway that provides the whole-word information that is particularly important for atypical word reading (3, 5). This

account therefore predicts the very strong association observed between semantic dementia and surface dyslexia. Variation around this central tendency, including cases of dissociation at the extremes, is accounted for individual differences in the degree of pre-morbid SR for atypical word reading (7, 8). Here, we exploited virtual lesioning of the left ATL using rTMS to provide support for this individual differences hypothesis. Our results support the connectionist triangle model as it is this framework that incorporates both (i) a need for semantic activation for atypical word reading and (ii) some mechanism of variation in the strength of this necessity due to individual differences. Our results make it clear that reading models must incorporate some account of individual variation in SR if they are to capture the full range of normal and disordered performance in behavioral and brain data. Indeed, Dilkina et al. (44) used a connectionist model to explore the extent to which individual differences arise from variation in reading practice, direct pathway resources, and lesion distribution. Their results allowed them to fit data from both semantic dementia patients who showed associations and who showed dissociations between performance on reading aloud and semantic tests.

In terms of the sources of the individual differences we have considered, our previous large-scale behavioral study indicated that stronger SR is associated with weaker performance on phonological processing tasks, in the form of slower nonword reading and rhyme judgment and lower scores on rhyme fluency (2). This aligns well with the Hoffman et al. (31) finding that high SR readers showed lower activation in the left pre-central gyrus during reading, as this is a region critical for phonological processing (45). It is possible that a mild disadvantage in phonological processing might provide the impetus for increased reliance on semantic activation during reading of atypical words, which are phonologically challenging by virtue of the competing pronunciations that they generate (46). Connectionist modeling of developmental dyslexia has shown that pre-reading phonological damage can undermine reading of both nonwords and atypical words (47). Such damage would undermine the competence of the direct pathway, and connectionist simulations have shown that this can be compensated for, in the case of atypical word reading, by increased SR (7). This working hypothesis could be pursued in future developmental research, while future imaging research could consider how individual differences in SR may manifest in other areas of the left hemisphere language network (21, 48) and also how the right ATL interacts with this network to support fluent reading.

This study has provided evidence to suggest that systematic individual differences in the healthy normal population can have appreciable consequences for the pattern of impairment observed after neural disruption (see also ref. 49). Such a demonstration has implications for the interpretation of neuropsychological data, as this has been heavily reliant upon the notion that dissociations indicate functional independence. Shallice (1) noted that pre-morbid individual differences could modulate the extent to which a dissociation is manifest, and that would certainly seem to be the case for the normal variation in SR for atypical word reading that

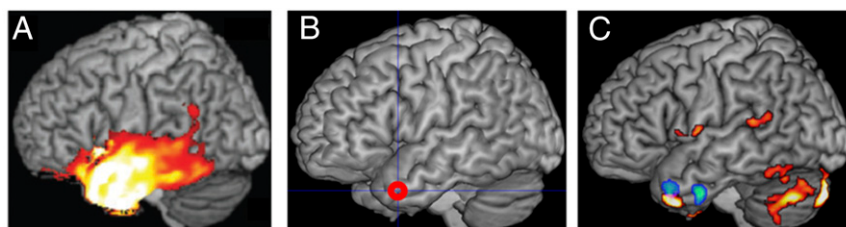


Fig. 2. A comparison of (A) areas of left ATL atrophy in semantic dementia patients with surface dyslexia (reproduced with permission by Oxford University Press from ref. 30); (B) the area of rTMS stimulation previously shown to disrupt semantic processing (27) and targeted in the current study; and (C) the areas activated by atypical word reading (warm colors) and showing a positive relationship with degree of SR (cool colors) based on data from Hoffman et al. (31).

we report here. Specifically, lower SR readers may initially show a dissociation between impaired semantic processing and relatively intact reading performance (18–20), while higher SR readers may initially show a dissociation between impaired reading performance and relatively intact semantic processing (50, 51). Taken together, these cases provide a double dissociation between reading and meaning, the gold standard of evidence for functional independence. However, the account proposed and supported here is instead of a graded relationship between reading and meaning in the healthy population that affects the degree of association between the two capacities seen after brain damage.

While the individual differences we have reported here do pose some challenges to the traditional interpretation of neuropsychological data, these merely emphasize the importance of considering the full range of performance across a case series of patients (52). Although Shallice (1) was skeptical that individual differences could speak directly to the functional architecture, we would suggest that the individual differences in reading that we have reported are extremely theoretically informative in showing a key relationship between atypical word reading and semantic processing. This study and its linked papers (2, 8, 31) indicate that individual differences can speak directly to issues of functional architecture as long as the relevant theory has a formal way to consider individual variation. The exploration of individual differences within the implemented triangle model of reading provides a tangible example of how this can be achieved (7, 8, 44). The underlying causes for this variation have yet to be determined (2), but the connectionist triangle model is well placed to explore this issue given its focus on learnt representations. Our results suggest any neurocognitive model needs to incorporate mechanisms that permit substantial systematic individual differences in the dynamics of the reading network. Given that reading is a late-acquired skill both phylogenetically and ontogenetically (53, 54), it is a prime example of a domain in which individual differences in the normal healthy population may be at their most marked, behaviorally (2) and neurally (21, 31, 48). However, substantial normal variation is apparent in a variety of higher cognitive capacities, and the neurostimulation approach we present could be harnessed to explore the nature of individual differences in other domains.

Materials and Methods

Participants. An initial pool of 129 individuals completed a reading-aloud task designed to assess their degree of SR for atypical word reading, as described in *Computation of the Individual Differences Index*. Stimulus properties are provided in Table S1. All participants were University of Manchester undergraduate or postgraduate students. We recruited as many TMS eligible and willing participants as possible from the upper and lower ends of the SR distribution to return to undergo a structural MRI scan and participate in the main TMS experiment. We obtained 18 participants in total. Nine of these from the lower end of the distribution (ranks 3, 8, 17, 18, 23, 29, 52, 54, 58, mean SR = 24.51, SD = 22.04) were classed as the low SR readers, while nine from the higher end of the distribution (ranks 60, 62, 83, 94, 97, 112, 114, 117, 118, mean SR = 78.97, SD = 23.20) were classed as the high SR readers. Their results on the SR index reading aloud task can be seen in Fig. S1 and Tables S2 and S3. All 18 had normal or corrected-to-normal vision; five participants were male. All participants spoke English as a first language, with one participant in the high SR group also speaking Finnish. The age of the low SR readers (mean = 22.00, SD = 3.00) did not differ significantly from that of the high SR readers (mean = 21.89, SD = 3.30). Scores on the Edinburgh Handedness

Inventory for the low SR readers (mean = 24.44, SD = 85.75) did not differ significantly from that of the high SR readers (mean = 21.11, SD = 75.57). None of the participants were taking medication, and all were free from any history of neurological disease or mental illness. The study was approved by the UK National Research Ethics Service Greater Manchester West research ethics committee, and informed consent was obtained from all participants.

Stimuli. The stimuli used in the TMS experiment were the 168 words from the Surface List (17). This list contains a factorial manipulation of frequency and spelling–sound regularity, with 48 items per cell, with properties of the stimuli provided in Table S4. This list is the same as that (i) used to quantify surface dyslexia in semantic dementia (17), (ii) to simulate this pattern within the connectionist triangle model (8), and also (iii) to determine neural activation among normal readers varying in degree of SR using fMRI (31). Accordingly, our TMS study can be directly related to these three previous, key studies. To prevent item repetition, the words were divided into two matched lists of 84 items (21 words per cell), one of which was presented before TMS and one of which was presented afterward. The order of lists was balanced within group across participants.

Procedure. The DMDX experimental software package (55) was used to record RTs and vocal responses and to display instructions and stimuli. Responses were collected by a voice-key plus headset connected to an IBM compatible Pentium III computer with a 60 Hz refresh rate at 1,280 × 1,024 pixel screen resolution. Vocal responses were recorded from the beginning of the trial for a period of 1,000 ms after the voice key triggered. Trials from all conditions were presented mixed together in a pre-stimulation and post-stimulation block, with the order of trial presentation within each block randomized anew for each participant, and stimuli were presented in white on a black background. The assignment of item sets to pre- or post-stimulation was counterbalanced across participants. Mispronunciations and measurement errors were recorded by hand. Participants were instructed to name the centrally presented words as rapidly and accurately as possible. Trials began with a 500-ms fixation cross followed by the word that disappeared from the screen upon response or after 2,000 ms.

The study used the virtual lesion method in which there was (i) a reading-aloud task (baseline), then (ii) rTMS stimulation, and immediately after (iii) an analogous reading-aloud task (probe). This meant that rTMS was delivered without a concurrent task and that the probe task was performed during the rTMS refractory period, which has been estimated to last for approximately 20 min (27). Focal magnetic stimulation was delivered using a 70-mm figure-of-eight coil attached to a MagStim Rapid2 stimulator (Magstim). Before experimental stimulation, motor threshold (MT) was determined for every participant as a visible twitch in the relaxed contralateral abductor pollicis brevis muscle in three out of six trials. Stimulation intensity for the experiment was then set at 120% of MT for each participant and consisted of 10 min of 1-Hz stimulation. A structural T1-weighted MRI scan was acquired for each participant to guide rTMS stimulation. The ATL site was defined as the region 10 mm posterior from the tip of the left temporal lobe along the MTG, consistent with previous studies where left ATL stimulation has been shown to disrupt semantic processing (25–27). The average Montreal Neurological Institute coordinates for the stimulated site were –53, 13, –32 (Fig. 2B). For stimulation, this site was determined by coregistering the cortical surface with 11 anatomical landmarks (inion, tip of the nose, left/right ear canals, and left/right ear projections), some of which were marked before the scan with oil capsules (vertex, nasion, left/right ear tragus, and beneath lip in chin indentation). Coregistration was made using Ascension miniBIRD magnetic tracking system and MRIreg software (www.cabiatl.com/micro/micro/mrireg/index.html).

ACKNOWLEDGMENTS. This work was supported by Leverhulme Trust Research Fellowship RF2/RFG/2008/0298 (to A.M.W.), MRC Programme Grant MR/J004146/1 (to M.A.L.R.), and ERC Advanced Grant GAP: 670428 - BRAIN2MIND_NEUROCOMP (to M.A.L.R.).

- Shallice T (1988) *From Neuropsychology to Mental Structure* (Cambridge Univ Press, New York).
- Woollams AM, Lambon Ralph MA, Madrid G, Patterson KE (2016) Do you read how I read? Systematic individual differences in semantic reliance amongst normal readers. *Front Psychol* 7:1757.
- Plaut DC, McClelland JL, Seidenberg MS, Patterson K (1996) Understanding normal and impaired word reading: Computational principles in quasi-regular domains. *Psychol Rev* 103:56–115.
- Coltheart M, Rastle K, Perry C, Langdon R, Ziegler J (2001) DRC: A dual route cascaded model of visual word recognition and reading aloud. *Psychol Rev* 108: 204–256.
- Harm MW, Seidenberg MS (2004) Computing the meanings of words in reading: Cooperative division of labor between visual and phonological processes. *Psychol Rev* 111:662–720.
- Seidenberg MS, McClelland JL (1989) A distributed, developmental model of word recognition and naming. *Psychol Rev* 96:523–568.
- Plaut DC (1997) Structure and function in the lexical system: Insights from distributed models of word reading and lexical decision. *Lang Cogn Process* 12:765–805.
- Woollams AM, Ralph MA, Plaut DC, Patterson K (2007) SD-squared: On the association between semantic dementia and surface dyslexia. *Psychol Rev* 114:316–339.
- Cortese MJ, Simpson GB, Woolsey S (1997) Effects of association and imageability on phonological mapping. *Psychon Bull Rev* 4:226–231.

10. Shibahara N, Zorzi M, Hill MP, Wydell T, Butterworth B (2003) Semantic effects in word naming: Evidence from English and Japanese Kanji. *Q J Exp Psychol A* 56: 263–286.
11. Strain E, Patterson K, Seidenberg MS (1995) Semantic effects in single-word naming. *J Exp Psychol Learn Mem Cogn* 21:1140–1154.
12. Strain E, Patterson K, Seidenberg MS (2002) Theories of word naming interact with spelling-sound consistency. *J Exp Psychol Learn Mem Cogn* 28:207–214, discussion 215–220.
13. Woollams AM (2005) Imageability and ambiguity effects in speeded naming: Convergence and divergence. *J Exp Psychol Learn Mem Cogn* 31:878–890.
14. Hodges JR, Patterson K, Oxbury S, Funnell E (1992) Semantic dementia. Progressive fluent aphasia with temporal lobe atrophy. *Brain* 115:1783–1806.
15. Nestor PJ, Fryer TD, Hodges JR (2006) Declarative memory impairments in Alzheimer's disease and semantic dementia. *Neuroimage* 30:1010–1020.
16. Graham KS, Hodges JR, Patterson K (1994) The relationship between comprehension and oral reading in progressive fluent aphasia. *Neuropsychologia* 32:299–316.
17. Patterson K, Hodges JR (1992) Deterioration of word meaning: Implications for reading. *Neuropsychologia* 30:1025–1040.
18. Blazely AM, Coltheart M, Casey BJ (2005) Semantic impairment with and without surface dyslexia: Implications for models of reading. *Cogn Neuropsychol* 22:695–717.
19. Cipolotti L, Warrington EK (1995) Semantic memory and reading abilities: A case report. *J Int Neuropsychol Soc* 1:104–110.
20. Schwartz MF, Marin OSM, Saffran EM (1979) Dissociations of language function in dementia: A case study. *Brain Lang* 7:277–306.
21. Graves WW, et al. (2014) Anatomy is strategy: Skilled reading differences associated with structural connectivity differences in the reading network. *Brain Lang* 133:1–13.
22. Baron J, Strawson C (1976) Use of orthographic and word-specific knowledge in reading words aloud. *J Exp Psychol Hum Percept Perform* 2:386–393.
23. Brown P, Lupker SJ, Colombo L (1994) Interacting sources of information in word naming: A study of individual differences. *J Exp Psychol Hum Percept Perform* 20: 537–554.
24. Walsh V, Cowey A (1998) Magnetic stimulation studies of visual cognition. *Trends Cogn Sci* 2:103–110.
25. Lambon Ralph MA, Pobric G, Jefferies E (2009) Conceptual knowledge is underpinned by the temporal pole bilaterally: Convergent evidence from rTMS. *Cereb Cortex* 19: 832–838.
26. Pobric G, Jefferies E, Ralph MA (2010) Amodal semantic representations depend on both anterior temporal lobes: Evidence from repetitive transcranial magnetic stimulation. *Neuropsychologia* 48:1336–1342.
27. Pobric G, Jefferies E, Ralph MA (2007) Anterior temporal lobes mediate semantic representation: Mimicking semantic dementia by using rTMS in normal participants. *Proc Natl Acad Sci USA* 104:20137–20141.
28. Woollams AM (2012) Apples are not the only fruit: The effects of concept typicality on semantic representation in the anterior temporal lobe. *Front Hum Neurosci* 6:85.
29. Woollams AM, Lambon Ralph MA, Plaut DC, Patterson K (2010) SD-squared revisited: Reply to Coltheart, Tree, and Saunders (2010). *Psychol Rev* 117:273–281, discussion 282–283.
30. Wilson SM, et al. (2009) The neural basis of surface dyslexia in semantic dementia. *Brain* 132:71–86.
31. Hoffman P, Lambon Ralph MA, Woollams AM (2015) Triangulation of the neuro-computational architecture underpinning reading aloud. *Proc Natl Acad Sci USA* 112: E3719–E3728.
32. Braet W, Humphreys GW (2006) Case mixing and the right parietal cortex: Evidence from rTMS. *Exp Brain Res* 168:265–271.
33. Costanzo F, Menghini D, Caltagirone C, Oliveri M, Vicari S (2012) High frequency rTMS over the left parietal lobule increases non-word reading accuracy. *Neuropsychologia* 50:2645–2651.
34. Nakamura K, et al. (2006) Task-guided selection of the dual neural pathways for reading. *Neuron* 52:557–564.
35. Skarratt PA, Lavidor M (2006) Magnetic stimulation of the left visual cortex impairs expert word recognition. *J Cogn Neurosci* 18:1749–1758.
36. Duncan KJ, Pattamadilok C, Devlin JT (2010) Investigating occipito-temporal contributions to reading with TMS. *J Cogn Neurosci* 22:739–750.
37. Coltheart M, Tree JJ, Saunders SJ (2010) Computational modeling of reading in semantic dementia: Comment on Woollams, Lambon Ralph, Plaut, and Patterson (2007). *Psychol Rev* 117:256–271, discussion 271–272.
38. Noble K, Glosser G, Grossman M (2000) Oral reading in dementia. *Brain Lang* 74: 48–69.
39. Pattamadilok C, et al. (2015) How early does the brain distinguish between regular words, irregular words, and pseudowords during the reading process? Evidence from neurochronometric TMS. *J Cogn Neurosci* 27:1259–1274.
40. Jackson RL, Lambon Ralph MA, Pobric G (2015) The timing of anterior temporal lobe involvement in semantic processing. *J Cogn Neurosci* 27:1388–1396.
41. Cumming TB, Patterson K, Verfaellie M, Graham KS (2006) One bird with two stones: Abnormal word length effects in pure alexia and semantic dementia. *Cogn Neuropsychol* 23:1130–1161.
42. Woollams AM, Hoffman P, Roberts DJ, Lambon Ralph MA, Patterson KE (2014) What lies beneath: A comparison of reading aloud in pure alexia and semantic dementia. *Cogn Neuropsychol* 31:461–481.
43. Joyal M, et al. (2017) The role of the left anterior temporal lobe for unpredictable and complex mappings in word reading. *Front Psychol* 8:517.
44. Dilkina K, McClelland JL, Plaut DC (2008) A single-system account of semantic and lexical deficits in five semantic dementia patients. *Cogn Neuropsychol* 25:136–164.
45. Vigneau M, et al. (2006) Meta-analyzing left hemisphere language areas: Phonology, semantics, and sentence processing. *Neuroimage* 30:1414–1432.
46. Woollams AM, Patterson K (2012) The consequences of progressive phonological impairment for reading aloud. *Neuropsychologia* 50:3469–3477.
47. Harm MW, Seidenberg MS (1999) Phonology, reading acquisition, and dyslexia: Insights from connectionist models. *Psychol Rev* 106:491–528.
48. Blackmon K, et al. (2010) Phonetically irregular word pronunciation and cortical thickness in the adult brain. *Neuroimage* 51:1453–1458.
49. Jefferies E, Rogers TT, Ralph MA (2011) Premorbid expertise produces category-specific impairment in a domain-general semantic disorder. *Neuropsychologia* 49: 3213–3223.
50. Mendez MF (2002) Slowly progressive alexia. *J Neuropsychiatry Clin Neurosci* 14:84.
51. Weekes B, Coltheart M (1996) Surface dyslexia and surface dysgraphia: Treatment studies and their theoretical implications. *Cogn Neuropsychol* 13:277–315.
52. Lambon Ralph MA, Patterson K, Plaut DC (2011) Finite case series or infinite single-case studies? Comments on "Case series investigations in cognitive neuropsychology" by Schwartz and Dell (2010). *Cogn Neuropsychol* 28:466–474, discussion 515–520.
53. Patterson K, Ralph MA (1999) Selective disorders of reading? *Curr Opin Neurobiol* 9: 235–239.
54. Woollams AM (2013) Connectionist neuropsychology: Uncovering ultimate causes of acquired dyslexia. *Philos Trans R Soc Lond B Biol Sci* 369:20120398.
55. Forster KI, Forster JC (2003) DMDX: A windows display program with millisecond accuracy. *Behav Res Methods Instrum Comput* 35:116–124.
56. Jared D (1997) Spelling-sound consistency affect the naming of high-frequency words. *J Mem Lang* 36:505–529.
57. Jared D (2002) Spelling-sound consistency and regularity effects in word naming. *J Mem Lang* 46:723–750.
58. Ziegler JC, Stone GO, Jacobs AM (1997) What is the pronunciation for -ough and the spelling for /u/? A database for computing feedforward and feedback consistency in English. *Behav Res Methods Instrum Comput* 29:600–618.
59. Cortese MJ, Simpson GB (2000) Regularity effects in word naming: What are they? *Mem Cognit* 28:1269–1276.
60. Townsend J, Ashby F (1978) Methods of modeling capacity in simple processing systems. *Cognitive Theory*, eds Castellan JN, Restle F (Erlbaum, Hillsdale, NJ), Vol 3, pp 200–239.