



Predictors of Poststroke Aphasia Recovery

A Systematic Review-Informed Individual Participant Data Meta-Analysis

The REhabilitation and recovery of peopLE with Aphasia after StrokE (RELEASE) Collaborators*

BACKGROUND AND PURPOSE: The factors associated with recovery of language domains after stroke remain uncertain. We described recovery of overall-language-ability, auditory comprehension, naming, and functional-communication across participants' age, sex, and aphasia chronicity in a large, multilingual, international aphasia dataset.

METHODS: Individual participant data meta-analysis of systematically sourced aphasia datasets described overall-language ability using the Western Aphasia Battery Aphasia-Quotient; auditory comprehension by Aachen Aphasia Test (AAT) Token Test; naming by Boston Naming Test and functional-communication by AAT Spontaneous-Speech Communication subscale. Multivariable analyses regressed absolute score-changes from baseline across language domains onto covariates identified a priori in randomized controlled trials and all study types. Change-from-baseline scores were presented as estimates of means and 95% CIs. Heterogeneity was described using relative variance. Risk of bias was considered at dataset and meta-analysis level.

RESULTS: Assessments at baseline (median=43.6 weeks poststroke; interquartile range [4–165.1]) and first-follow-up (median=10 weeks from baseline; interquartile range [3–26]) were available for n=943 on overall-language ability, n=1056 on auditory comprehension, n=791 on naming and n=974 on functional-communication. Younger age (<55 years, +15.4 Western Aphasia Battery Aphasia-Quotient points [CI, 10.0–20.9], +6.1 correct on AAT Token Test [CI, 3.2–8.9]; +9.3 Boston Naming Test points [CI, 4.7–13.9]; +0.8 AAT Spontaneous-Speech Communication subscale points [CI, 0.5–1.0]) and enrollment <1 month post-onset (+19.1 Western Aphasia Battery Aphasia-Quotient points [CI, 13.9–24.4]; +5.3 correct on AAT Token Test [CI, 1.7–8.8]; +11.1 Boston Naming Test points [CI, 5.7–16.5]; and +1.1 AAT Spontaneous-Speech Communication subscale point [CI, 0.7–1.4]) conferred the greatest absolute change-from-baseline across each language domain. Improvements in language scores from baseline diminished with increasing age and aphasia chronicity. Data exhibited no significant statistical heterogeneity. Risk-of-bias was low to moderate-low.

CONCLUSIONS: Earlier intervention for poststroke aphasia was crucial to maximize language recovery across a range of language domains, although recovery continued to be observed to a lesser extent beyond 6 months poststroke.

Key Words: aphasia ■ comprehension ■ demography ■ language ■ survivor

Approximately one-third of the 25.7 million stroke survivors¹ worldwide experience aphasia,² affecting spoken language, auditory and reading comprehension, writing, and everyday communication. Aphasia is associated with poorer performance on functional recovery³ activities of daily living,⁴ and emotional well-being after stroke.⁵

Aphasia also affects hospital discharge destination⁶ and the likelihood of successful return to work. Long-term language impairment affects 61% of stroke survivors' communication at 1 year after onset.⁷ While spontaneous recovery seems limited after that point^{8,9} focused therapeutic intervention may benefit people with chronic aphasia.¹⁰ Evidence is

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Nonstandard Abbreviations and Acronyms

AAT-TT	Aachen Aphasia Test Token Test
IPD	individual participant data
IQR	interquartile range
RCT	randomized controlled trial
SLT	speech and language therapy

needed to clarify the nature and role of demography, stroke, and aphasia profiles on language recovery after stroke.

Clinical guidelines¹¹ recommend communication of realistic prognoses for recovery. However, accurate prognostication based on existing evidence is challenging,¹² due to use of single-center studies offering small sample sizes¹³ and investigating single language domains,¹⁴ thereby limiting information on the recovery potential across the spectrum of impaired language domains and demography for people with aphasia.

Severity of aphasia at onset has been linked with recovery^{15,16} but is often categorized in broad terms (eg, mild/moderate/severe)⁷ or as an item within a global stroke severity measure,¹⁷ lacking sensitivity to provide clinically meaningful indications of recovery potential across the range of affected language domains. Previous studies examining aphasia profiles and recovery have been limited to literature reviews and aggregated summary data,¹⁸ retrospective review of hospital records,^{19,20} nonstandardized determination of aphasia, or aggregation within systematic reviews,¹³ often using English-speaking datasets, uncontrolled and nonrandomized studies.²¹ Associations between age, sex, and language recovery have been suggested, but the nature of that interaction remains unclear.^{7,22} The degree of recovery from aphasia across the poststroke trajectory is also uncertain, despite variable evidence for early intervention,²³ with some suggesting intervention should be delayed.²⁴

The recovery potential across the spectrum of people with aphasia requires analysis of large samples reflecting a range of demographic variables, across multiple geographic centers, with detailed assessment of language domains, using standardized methods of data collection, within a clinically relevant time-frame. A pragmatic approach using existing datasets may inform clinical insights and direct future research. Systematic review-based Individual Participant Data (IPD) meta-analysis allows for individual representation and exploration of missing data, as well as individual adjustment for prognostic factors on a larger scale. Examining IPD from several studies facilitates greater participant representation,²⁵ enables subgroup analyses, and can synthesize smaller and disparate datasets at IPD level without bias that might be introduced through the use of single-center meta-analyses.

We conducted an IPD meta-analysis using a rigorous, systematically collated international aphasia research

database to identify demographic, stroke- and language-related factors associated with aphasia recovery.

METHODS

Ethics, Protocol, Registration, and Guidelines

The Rehabilitation and Recovery of People With Aphasia After Stroke Project collated data from completed aphasia studies (University ethical approval: HLS/NCH/15/09; PROSPERO; CRD42018110947; IRAS; database ID 179505). The protocol for development of this database including search strategy and eligibility criteria was published elsewhere²⁶; methods and findings are reported according to the PRISMA-IPD guidelines.

Data Availability

Where contributors have given permission, fully anonymized datasets will be made available to the wider research community through the Collaboration of Aphasia Trialists (<https://www.aphasiatrials.org/aphasia-dataset/>) from December 2020.

Procedures

We systematically identified datasets with at least 10 people with poststroke aphasia, documented language assessments and time since index stroke from MEDLINE, EMBASE, CINAHL, LLBA, and SpeechBITE (inception to September 2015) supplemented by a search of trial registrations to identify emerging datasets beyond this period. Building upon previous aggregated trial data syntheses,²⁷ we extracted anonymized IPD on demography (age, sex, handedness, language of assessment, education level, socioeconomic status), time from stroke onset to inclusion in the primary research study, type of stroke, study design, language outcome data (overall-language-ability, auditory comprehension, naming, other spoken language, reading comprehension, writing, and functional-communication), and timing of language assessment. Assessment instruments were categorized by the language domain measured; categorizations were reviewed, discussed, and accepted by the Rehabilitation and Recovery of People With Aphasia After Stroke Collaborators a priori. We retained complete IPD on language domain assessments where available at both baseline and first follow-up (Figure 1).

Standardization of Outcomes

We converted IPD gathered on multiple assessment instruments into one standardized measurement for each language domain (see Table I in the [Data Supplement](#)) as previously reported.²⁶ Data integrity was assessed by performing checks for ranges, missingness, and expected formats. Selection, detection, attrition, and reporting biases were assessed at study and database level.

Primary analysis was based on randomized controlled trial (RCT) IPD; secondary analyses included all study designs (RCT plus non-RCT group comparisons, cohort, case series and registry datasets). Analyses were specified a priori and comprised at least 2 source datasets. Our primary outcome was mean absolute change in language domain score from baseline to first follow-up. Secondary outcomes were absolute and relative proportions of change in language scores from baseline.

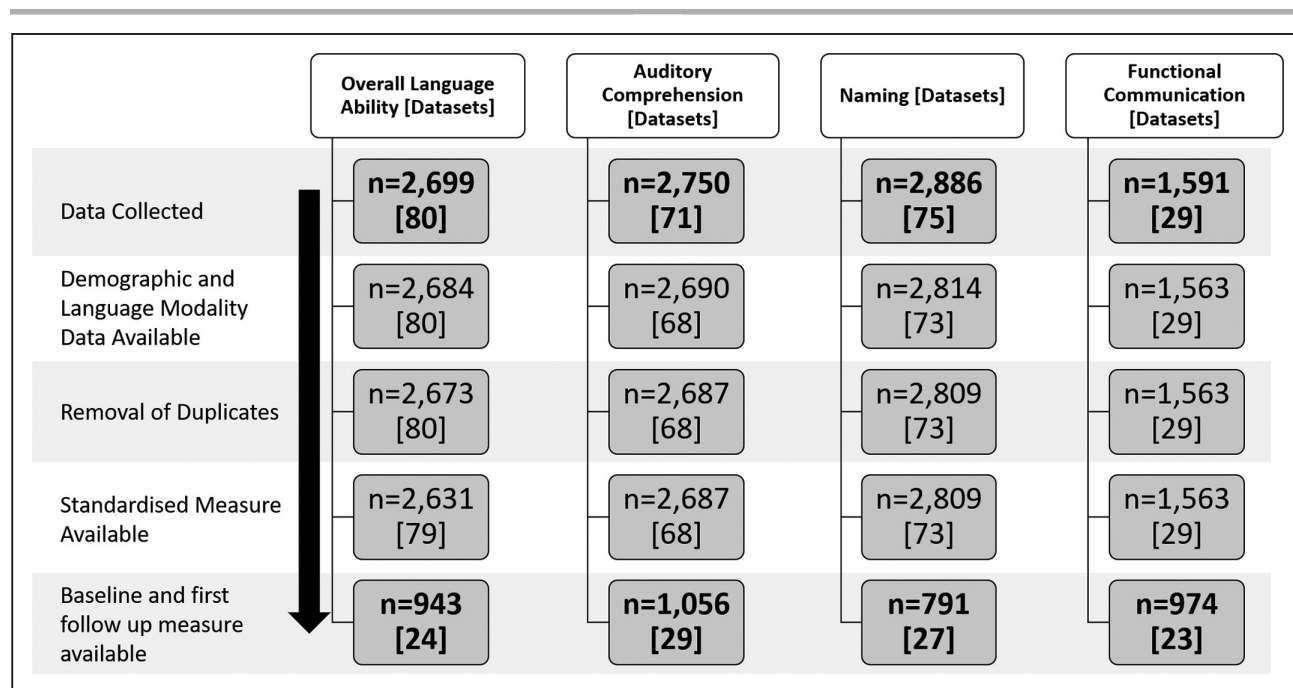


Figure 1. Flowchart of individual participant data and dataset inclusion.

n=individual participant data.

Statistical Analysis

A one-stage meta-analysis approach combined available IPD from eligible datasets with analyses preserving the clustering of participants within each study and were generated using SAS/STAT software, Version 9.4 of the SAS System for Windows.

Where possible, we presented data according to whether the participant had access to speech and language therapy (SLT). People with aphasia following a first stroke, allocated to receive no study-mediated SLT, no standard care SLT, and who were enrolled within 15 days of stroke onset (reducing the possibility of unreported historical SLT exposure) were described as the No SLT group. People with aphasia who were exposed to any form of SLT (eg, as part of a study or standard SLT care during the study period) were the SLT group. A historical SLT group comprised people who received no study-mediated SLT intervention, no standard SLT care during the study period but who were enrolled after 15 days of onset and, unless otherwise stated, were therefore likely to have received SLT as part of standard care in the past.

Within each domain, we performed meta-analysis with fixed demographic and recovery effects and included study as a random effect; variance was also evaluated. The absolute change in language domain scores from baseline to first follow-up was regressed onto stroke, aphasia, and demographic covariates specified a priori (baseline language impairment score, age, sex, lesioned hemisphere, stroke type, handedness, language of data collection, and time since stroke). Only variables that were significant at univariable level ($P < 0.1$) were included in the multivariable analysis. Each multivariable analysis included the study as a random covariate to account for possible variability in the outcome measure between studies, with the demographic variables modeled jointly. This allowed us to present each demographic variable as the independent effect following the inclusion of the other variables in the model and allowed us to avoid the effect of confounding. Results were presented as

estimate mean change in absolute scores from baseline and 95% confidence limits.

We described recovery in the context of absolute and relative proportions of change from baseline. Absolute proportion of change was described as the difference between scores at baseline and first follow-up, expressed as a proportion of the maximum possible assessment instrument score $[(\text{individual change score} + \text{maximum possible score}) \times 100]$. Relative proportion of change since baseline was described as the proportion of change in score at first follow-up, relative to baseline score $[(\text{individual change score} + \text{baseline score}) \times 100]$. Absolute proportion has a ceiling of 100% and assumes that recovery scales are linear; that is, an improvement of 20% anywhere on the scale from a starting point of 0 to 80 is equivalent. Relative proportion is unbounded and emphasizes gains at the more severe end of the scale (eg, a patient recovering from 10% to 30% will improve by 200% on this measure, while a person recovering from 70% to 90% will improve by 29%).

We examined chronicity since stroke for each language domain across the following study entry points: 0 to 1, 1 to 3, 3 to 6, and >6 months. We accounted for the skewness of data for each language domain by presenting medians and interquartile ranges (IQRs).

RESULTS

We screened 5256 records. From 698 potentially eligible datasets (592 resulted in no data contribution: 193/592 were trial registration records, 318 received no responses, 78 had no available data and 3 declined to participate), 174 research studies were included (IPD=5928; 24.9% of potentially eligible datasets). We included IPD from 47 RCTs, 18 non-RCTs, 5 registries, and 104 case-series/cohort studies and extracted data

on demography and language outcomes for analysis; median age was 63 (IQR, 53–72), 3407/5550 (61.4%) were male and median aphasia chronicity was 321 (IQR, 30–1156) days. Table II in the [Data Supplement](#) describes participant demographics across each language domain. Characteristics of included studies are reported elsewhere.²⁸

Data checks revealed no clustering of transformed language domain scores that were generated from any single assessment instrument (Figure I in the [Data Supplement](#)). The transformed values were therefore considered to be accurate and valid. Data at baseline (median=43.6 weeks since stroke; IQR [4–165.1]) and first follow-up (median=10 weeks since baseline; IQR [3–26]) were available for overall-language-ability ($n=943$; 24 datasets, represented by the Western Aphasia Battery-AQ), auditory comprehension ($n=1056$; 29 datasets, represented by the Aachen Aphasia Test [AAT] Token Test), naming ($n=791$, 27 datasets, by Boston Naming Test), and functional-communication ($n=974$, 23 datasets, represented by the AAT Spontaneous-Speech Communication subscale, Figure 1). Data were extracted on other spoken language production ($n=231$, represented by the Porch Index of Communicative Ability), reading comprehension ($n=219$, represented by the Reading subtest of the Comprehensive Aphasia Test), and writing ($n=253$, represented by the Writing subtest of the Comprehensive Aphasia Test). Results for language production, reading comprehension, and writing are not presented as they did not meet the threshold for analysis defined a priori (at least 2 datasets), did not include RCT data, or contained too few subgroup data points for reliable analyses.

There were inadequate data on handedness (left-handed $n=133$; ambidextrous $n=27$) in combination with other key demographic variables to permit informative covariate-adjusted analyses. Socioeconomic status ($n=175$) and education level ($n=3125$) were reported using a range of nonaggregable formats (eg, multiple deprivation index, years of education and occupation). In combination with available outcome measurements, these resulted in very small strata within analyses and hindered meaningful covariate adjustment. Education, handedness, and socioeconomic status were therefore excluded as covariates from our analyses.

There were also insufficient language data on participants where we could be confident that had No SLT before or during the primary research (IPD=0 for overall-language-ability; IPD=15 for auditory comprehension; IPD=29 for naming). Data on participants in the historical SLT only group were limited to 2 RCTs (IPD=22) and 3 datasets in all study types (IPD=34); corresponding postintervention follow-up values were unavailable or did not meet our minimum eligibility criteria for analysis. Data are therefore presented on those who had access to SLT during the study intervention period.

Predictors of Language Recovery

Overall-Language-Ability

Within the RCT datasets, the largest mean absolute change in overall-language-ability (presented as Western Aphasia Battery-AQ points) was seen in those aged <55 years (+15.4 points CI [10–20.9] IPD=136, 11 RCTs). Gains were also observed in those aged 56 to 65 years (+12.4 points; IPD=141, 11 RCTs), 66 to 75 years (+11.5 points; IPD=96, 10 RCTs), and >75 years (+13.8 points; IPD=109; 7 RCTs; see Table III in the [Data Supplement](#)).

When examining aphasia chronicity, enrollment within 1 month of stroke was associated with greatest mean absolute change in overall-language-ability (+19.1 points on the, CI [13.9–24.4]; IPD=260, 8 RCTs). Gains were also significant for participants enrolled at later time points (1–3 months: +16.2 points [IPD=64, 6 RCTs]; 3–6 months: +9.6 points [IPD=16, 3 RCTs] and >6 months +8.2 points [IPD=142, 4 RCTs]).

Women experienced slightly greater gains in overall-language-ability (+14.3 points from baseline; 95% CI [9–19.5], IPD=206, 11 RCTs) compared with men (+12.3 points 95% CI [7.2–17.4]; IPD=276, 11 RCTs); however, score differences were not clinically meaningful. These observations were also consistent when analyzing data from all study types (see Table III in the [Data Supplement](#)).

Auditory Comprehension

Based on the analysis of RCT datasets, younger people experienced greater gains in auditory comprehension (presented as correct items on the AAT Token Test; <55 years: +6.1 correct; CI [3.2–8.9]; IPD=178, 16 RCTs) than older people. More correct responses were observed for enrollment within 1 month of stroke (+5.3 correct, 95% CI [1.7–8.8], IPD=139, 6 RCTs); gains were also evident >6 months post aphasia onset (+1.4 correct, 95% CI [–1.9 to 4.7] IPD=243, 9 RCTs). These observations were also consistent when analyzing data from all study types (Table IV in the [Data Supplement](#)).

Naming

Younger people (<55 years; IPD=103, 13 RCTs) experienced a gain of +9.3 points on the Boston Naming Test (CI [4.7–13.9]). Gains of +6.2 and +4.4 points were evident for people aged 66 to 75 years (IPD=97, 12 RCTs) and >75 years (IPD=61, 11 RCTs), respectively. When examining aphasia chronicity, the greatest gain was observed for enrollment within 1 month of stroke (+11.1 points, CI [5.7–16.5], IPD=129, 5 RCTs). Gains were also apparent in those enrolled at 1 to 3 months (+7.7 points; IPD=93, 8 RCTs), 3 to 6 months (+4.3 points; IPD=70, 6 RCTs), and >6 months (+4.1 points; IPD=93, 7 RCTs). These observations were also evident when analyzing all study types (Table V in the [Data Supplement](#)).

Functional Communication

Younger people (<55 years, IPD=147, 14 RCTs) experienced an absolute gain of +0.75 points on the AAT Spontaneous-Speech Communication subscale (CI [0.5–1]). Gains were also seen for those aged 56 to 65 years (+0.7 points; IPD=145, 13 RCTs), 66 to 75 years (+0.55 points; IPD=121, 14 RCTs), and >75 years (+0.65 points; IPD=119, 12 RCTs).

When examining chronicity, the greatest absolute gain was observed for enrollment within 1 month of stroke (+1-point, 95% CI [0.7–1.4]; IPD=232, 6 RCTs). Gains were also evident at later time points (1–3 months=+0.87 points [IPD=68, 5 RCTs]; 3–6 months=+0.4 points [IPD=62, 4 RCTs]; >6 months=+0.33 points [IPD=170, 7 RCTs]).

Women (IPD=236, 14 RCTs) experienced greater gains in functional-communication (+0.76 points [95% CI, 0.5–1]), compared with men (+0.57 points [95% CI, 0.3–0.8]; IPD=296, 14 RCTs). These observations were consistent with analysis of all study types (Table VI in the Data Supplement).

Proportion of Recovery Across Each Language Domain

RCT Population

We considered the absolute and relative proportions of change on each language domain score from baseline. The largest absolute proportion of change was observed on measures of functional-communication (median=10%; IQR [0%–26.6%]; 16 RCTs, 608 IPD) and overall-language-ability (median=8.4% IQR [1.3%–22%], 11 RCT, 418 IPD, Figure 2A), while the largest relative proportion of change was in naming (median=35.7%; IQR [4.2%–133.3%]; 14 RCTs, 293 IPD) and functional-communication (median=25% IQR [0%–98.9%], 16 RCTs, 595 IPD; Figure 2B).

We observed a greater absolute (16.7% for overall-language-ability; IQR [4.3–35.1]; 12.4% for auditory comprehension; IQR [4–23.9]; 16.7% for naming; IQR [4.5–40]; and 18.9% for functional-communication; IQR [5.6–42.6] Figure 3A) and relative proportion of change (29.6% for overall-language-ability; IQR [11.3–73.9]; 72.7% for naming; IQR [20–310]; and 52.9% for functional-communication; [9.8–152.9] Figure 3B) for enrollment within 1 month of onset compared with other time windows poststroke.

All Study Designs

Data were available on overall-language-ability (IPD=788, 21 datasets); auditory comprehension (IPD=921, 27 datasets); naming (IPD=646, 24 datasets); and functional-communication (IPD=802, 22 datasets). We observed the largest absolute proportion of change in scores since baseline for functional-communication (9.5% IQR [0%–26.6%]) and overall-language-ability

(6.8% IQR [0.3%–19.3%], Figure II in the Data Supplement), and the largest relative proportion of change for naming (IPD=582, 24 datasets; median=24.3% IQR [0–115.4], see Figure II in the Data Supplement).

Heterogeneity and Bias

We found no evidence of significant statistical heterogeneity. Within all study types, most participant groups were comparable with respect to demographic, stroke, and aphasia severity variables at baseline, with baseline significant differences for subgroups only evident for age in 3/47 RCTs, sex in 1/47 RCTs, time since stroke in 2/47 RCTs, and language impairment in 1/47 RCTs. RCT datasets within Rehabilitation and Recovery of People With Aphasia After Stroke had a low risk of several sources of bias; most RCTs reported outcome assessor blinding (n=34/47; 75%). Attrition bias was low, and there was a moderate to low risk of selection bias, despite the use of predominantly research datasets. We considered the level of evidence available to be moderate because of potentially eligible datasets being identified in our systematic search but were not confirmed as eligible or contributed to Rehabilitation and Recovery of People With Aphasia After Stroke.²⁸ Our analysis dataset showed no evidence of publication bias by sample size ($P=0.77$). Comparisons of missingness with the main demographic variables did not reveal any systematic relationships.

DISCUSSION

Our findings improve our understanding of poststroke language recovery, the domains that recover most, and the degree of recovery reported (in relative and absolute terms), and in which poststroke time windows the greatest recovery takes place. We observed the greatest improvement for enrollment within 1-month poststroke across all language domains. Improvements in mean absolute scores from baseline diminished with increasing time since stroke, yet still exceeded established group-level benchmarks of significant change for overall-language-ability (5.03 Western Aphasia Battery-AQ points) and naming (3.3 Boston Naming Test points).²⁹ Relative and absolute proportions of change in scores across each language domain were typically the greatest within 1 month of stroke onset.

Our findings have important implications for the timing of SLT delivery; earlier intervention was associated with the greatest improvement across language domains. However, should early intervention be infeasible, for example, due to concurrent illness or inability to engage in rehabilitation, significant improvements were still observed beyond the acute period. We demonstrated clinical gains in the cohort who were enrolled beyond 6 months poststroke, which included those with a chronicity beyond 2 years. This population typically received usual care before study enrollment and

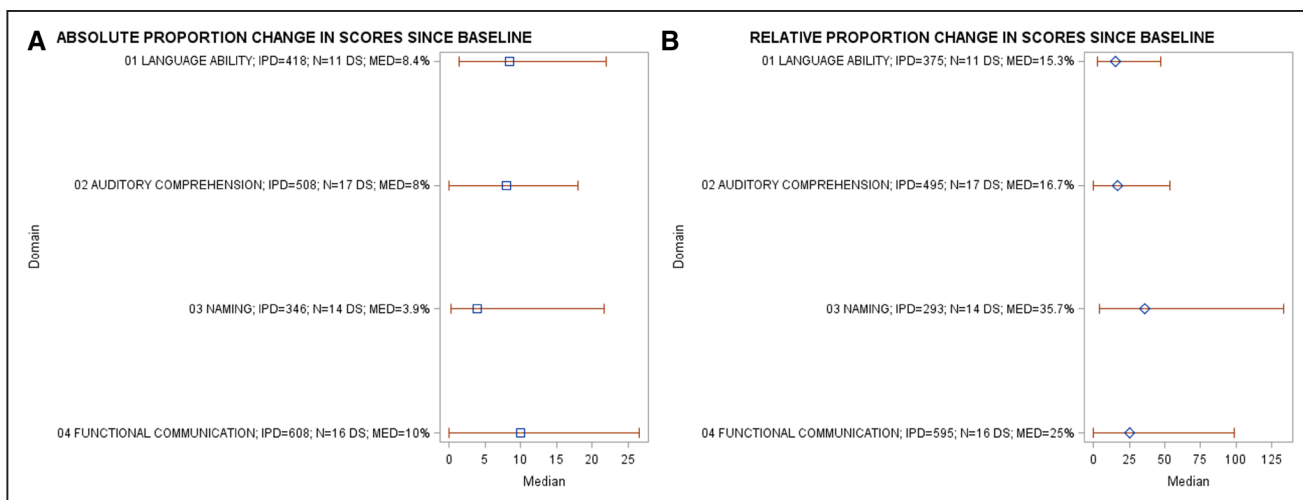


Figure 2. Absolute and relative proportions of recovery across all language domains in the randomized controlled trial population. IPD indicates individual participant data; MED, median proportion of recovery; and N, number of datasets.

still made clinical gains in language outcomes. In addition to greater improvements observed in the youngest population (<55 years) across all domains, we also

observed substantial rehabilitation potential for elderly participants (>75 years), where gains of +13.8 Western Aphasia Battery-AQ and +4.4 Boston Naming

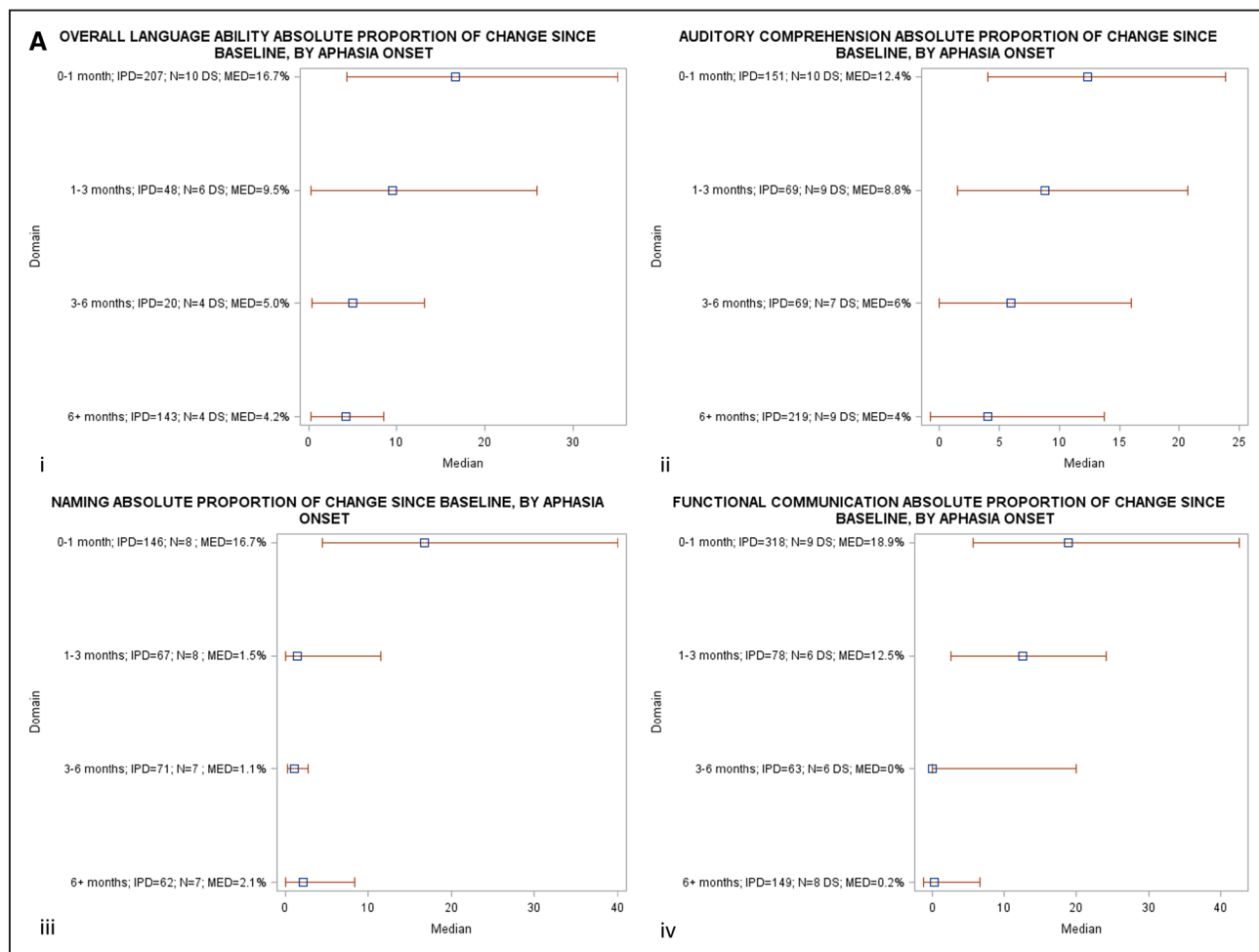


Figure 3. A, Absolute proportion of recovery across all language domains, stratified by time since index stroke, in randomized controlled trial (RCT) populations.

B, Relative proportion of recovery across all language domains, stratified by time since index stroke, in RCT populations. DS indicates datasets; IPD, individual participant data; MED, median recovery; and N, number of datasets. (Continued)

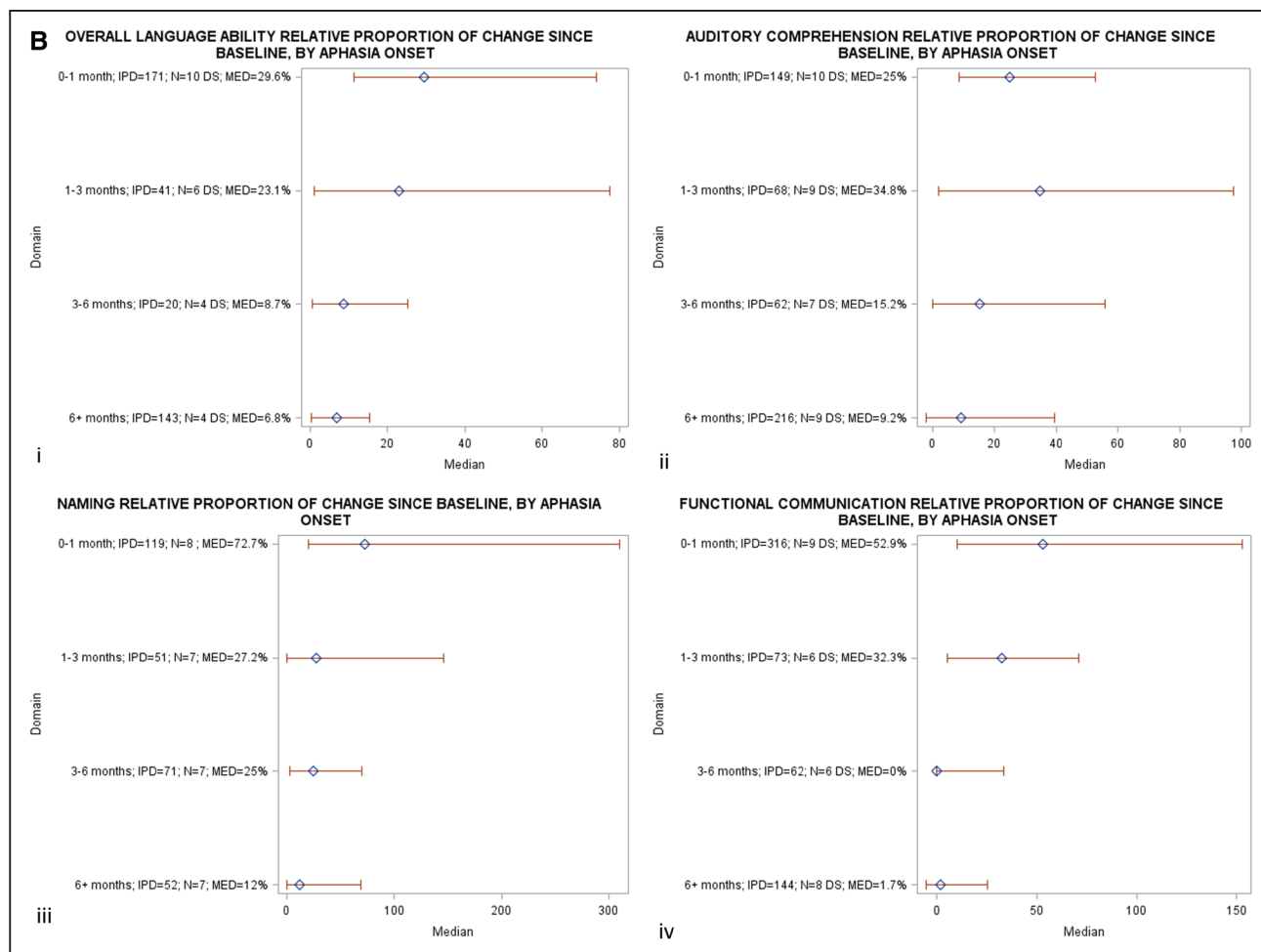


Figure 3 Continued.

Test points also exceeded the established group-level benchmarks of significant change for overall-language-ability and naming, respectively.²⁹

We employed novel data transformation methods to standardize the language outcome data, and robust data synthesis methods including a systematic search for international and multilingual aphasia studies, including IPD from not only RCTs but also case-series/ cohort studies, registries and non-RCTs. Inclusion of international, clinical datasets alongside research datasets enhanced the relevance of our data. We provided evidence primarily based on RCT sources and validated these observations in data from all study types. Importantly, collation and standardization of data from these sources permitted analysis of a much larger sample size and included a broader range of language domains than previously available. Our analyses preserved the clustering of participants within each trial.³⁰ We reduced the risk of bias and increased applicability by preventing analyses to be undertaken on single datasets. Our operational definition of recovery was based on absolute proportion of change in scores from baseline, consistent with existing definitions.¹⁵

While some previous studies have described a beneficial impact of intervention within 3 months of onset, these analyses were based on data from English-language only, uncontrolled and nonrandomized studies.²¹ Other studies recommended delayed interventions^{24,31} or reported on SLT initiation after 12 weeks post-onset.³² Our study extends these findings, describing marked improvements in language outcomes for earlier initiation of therapy, compared with later initiation in a large, multilingual, international sample, and provides data across a range of affected language domains.

We were unable to account for spontaneous recovery in our study. Many potentially important covariates such as language stimulation in the living environment, other concomitant rehabilitation interventions, education level, initial stroke severity, mood disorders, co-existing cognitive impairments, and socioeconomic status were inconsistently available across the datasets, and therefore could not inform our planned analyses. Additionally, there were inadequate data on reading and writing assessments to permit examination of these factors. While we collated data from almost 6000 IPD, presence of single assessment time points, availability of

language domain assessments, demographic and clinical factors only permitted analyses of samples ranging between 943 and 1056 IPD.

Nevertheless, our estimates and predictors of recovery are robust, use clinically meaningful assessment instruments, adjusting for critical participant confounders and are based on analyses of much larger sample sizes than previously examined.³³ Our findings indicate a need for further investigation of therapy-associated recovery effects, and subgroup differences to ascertain which subpopulations may respond better to different types of intervention. Planned analyses will explore the associations between treatments, intensities, durations, dosages, and language outcomes.

ARTICLE INFORMATION

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Supplemental Materials

Tables I–VI

Figures I and II

APPENDIX

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REFERENCES

1. Feigin VL, Norrving B, Mensah GA. Global burden of stroke. *Circ Res*. 2017;120:439–448. doi: 10.1161/CIRCRESAHA.116.308413
2. van Oers CAMM, van der Worp HB, Kappelle LJ, Raemaekers MAH, Otte WM, Dijkhuizen RM. Etiology of language network changes during recovery of aphasia after stroke. *Sci Rep*. 2018;8:856. doi: 10.1038/s41598-018-19302-4
3. Gialanella B, Bertolinelli M, Lissi M, Prometti P. Predicting outcome after stroke: the role of aphasia. *Disabil Rehabil*. 2011;33:122–129. doi: 10.3109/09638288.2010.488712
4. Paolucci S, Antonucci G, Grasso MG, Morelli D, Troisi E, Coiro P, Bragoni M. Early versus delayed inpatient stroke rehabilitation: a matched comparison conducted in Italy. *Arch Phys Med Rehabil*. 2000;81:695–700. doi: 10.1016/s0003-9993(00)90095-9
5. Thomas SA, Walker MF, Macniven JA, Haworth H, Lincoln NB. Communication and Low Mood (CALM): a randomized controlled trial of behavioural therapy for stroke patients with aphasia. *Clin Rehabil*. 2013;27:398–408. doi: 10.1177/0269215512462227
6. Gialanella B, Prometti P. Rehabilitation length of stay in patients suffering from aphasia after stroke. *Top Stroke Rehabil*. 2009;16:437–444. doi: 10.1310/tsr1606-437
7. Pedersen PM, Vinter K, Olsen TS. Aphasia after stroke: type, severity and prognosis. The Copenhagen aphasia study. *Cerebrovasc Dis*. 2004;17:35–43. doi: 10.1159/000073896
8. Ferro JM, Mariano G, Madureira S. Recovery from aphasia and neglect. *Cerebrovasc Dis*. 1999;9 (Suppl 5):6–22. doi: 10.1159/000047571
9. Prepared by the Intercollegiate Stroke Working Party. *National Clinical Guideline for Stroke*. 4th ed. Royal College of Physicians London; 2012.
10. Meinzer M, Djundja D, Barthel G, Elbert T, Rockstroh B. Long-term stability of improved language functions in chronic aphasia after constraint-induced aphasia therapy. *Stroke*. 2005;36:1462–1466. doi: 10.1161/01.STR.0000169941.29831.2a
11. Royal College of Physicians, National Clinical Guidelines for Stroke, 2016. [https://www.strokeaudit.org/SupportFiles/Documents/Guidelines/2016-National-Clinical-Guideline-for-Stroke-5t-\(1\).aspx](https://www.strokeaudit.org/SupportFiles/Documents/Guidelines/2016-National-Clinical-Guideline-for-Stroke-5t-(1).aspx).
12. Hillis AE, Beh YY, Sebastian R, Breining B, Tippett DC, Wright A, Saxena S, Rorden C, Bonilha L, Basilakos A, et al. Predicting recovery in acute post-stroke aphasia. *Ann Neurol*. 2018;83:612–622. doi: 10.1002/ana.25184
13. Flowers HL, Skoretz SA, Silver FL, Rochon E, Fang J, Flamand-Roze C, Martino R. Poststroke aphasia frequency, recovery, and outcomes: a systematic review and meta-analysis. *Arch Phys Med Rehabil*. 2016;97:2188–2201.e8. doi: 10.1016/j.apmr.2016.03.006
14. El Hachoui H, Lingsma HF, van de Sandt-Koenderman MW, Dippel DW, Koudstaal PJ, Visch-Brink EG. Long-term prognosis of aphasia after stroke. *J Neural Neurosurg Psychiatry*. 2013;84:310–315. doi: 10.1136/jnnp-2012-302596
15. Lazar RM, Minzer B, Antonielli D, Festa JR, Krakauer JW, Marshall RS. Improvement in aphasia scores after stroke is well predicted by initial severity. *Stroke*. 2010;41:1485–1488. doi: 10.1161/STROKEAHA.109.577338
16. Basso A. Prognostic factors in aphasia. *Aphasiology*. 1992;6:337–348.
17. Ali M, Lyden P, Brady M; VISTA Collaboration. Aphasia and dysarthria in acute stroke: recovery and functional outcome. *Int J Stroke*. 2015;10:400–406. doi: 10.1111/ijs.12067
18. Doogan C, Dignam J, Copland D, Leff A. Aphasia recovery: when, how and who to treat? *Curr Neurol Neurosci Rep*. 2018;18:90. doi: 10.1007/s11910-018-0891-x
19. Dickey L, Kagan A, Lindsay MP, Fang J, Rowland A, Black S. Incidence and profile of inpatient stroke-induced aphasia in Ontario, Canada. *Arch Phys Med Rehabil*. 2010;91:196–202. doi: 10.1016/j.apmr.2009.09.020
20. Bersano A, Burgio F, Gattinoni M, Candelise L; PROSIT Study Group. Aphasia burden to hospitalised acute stroke patients: need for an early rehabilitation programme. *Int J Stroke*. 2009;4:443–447. doi: 10.1111/j.1747-4949.2009.00349.x
21. Robey R. A meta-analysis of clinical outcomes in the treatment of aphasia. *J Speech Lang Hear Res*. 1998;41:172–187. doi: 10.1044/jslhr.4101.172
22. Laska AC, Hellblom A, Murray V, Kahan T, Von Arbin M. Aphasia in acute stroke and relation to outcome. *J Intern Med*. 2001;249:413–422. doi: 10.1046/j.1365-2796.2001.00812.x
23. Nouwens F, Visch-Brink EG, Van de Sandt-Koenderman MM, Dippel DW, Koudstaal PJ, de Lau LM. Optimal timing of speech and language therapy for aphasia after stroke: more evidence needed. *Expert Rev Neurother*. 2015;15:885–893. doi: 10.1586/14737175.2015.1058161
24. Bowen A, Hesketh A, Patchick E, Young A, Davies L, Vail A, Long AF, Watkins C, Wilkinson M, Pearl G, et al. Effectiveness of enhanced communication therapy in the first four months after stroke for aphasia and dysarthria: a randomised controlled trial. *BMJ*. 2012;345:e4407. doi: 10.1136/bmj.e4407
25. EFSA NDA Panel. EFSA Panel on dietetic products, nutrition and allergies: scientific opinion on dietary reference values for vitamin D. 2016; <https://efsa.onlinelibrary.wiley.com/doi/epdf/10.2903/j.efsa.2016.4547>. Accessed October 8, 2019.
26. Brady MC, Ali M, VandenBerg K, Williams LJ, Williams LR, Abo M, Becker F, Bowen A, Brandenburg C, Breitenstein C, et al. RELEASE: a protocol for a systematic review based, individual participant data, meta- and network meta-analysis, of complex speech-language therapy interventions for stroke-related aphasia. *Aphasiology*. 2019;34:137–157.
27. Brady MC, Kelly H, Godwin J, Enderby P, Campbell P. Speech and language therapy for aphasia following stroke. *Cochrane Database Syst Rev*. 2016; 6:CD000425. doi: 10.1002/14651858.CD000425.pub4
28. On behalf of the RELEASE Collaborators. RELEASE: An Individual Participant Data Meta-Analysis, Incorporating Systematic Review and Network Meta-Analysis, of Complex Speech-Language Therapy Interventions for

Stroke-Related Aphasia. National Institutes of Health Research: Health Services and Delivery Research; 2021 (in Press).

29. Gilmore N, Dwyer M, Kiran S. Benchmarks of significant change after aphasia rehabilitation. *Arch Phys Med Rehabil*. 2019;100:1131–1139.e87. doi: 10.1016/j.apmr.2018.08.177
30. Riley RD, Lambert PC, Abo-Zaid G. Meta-analysis of individual participant data: rationale, conduct, and reporting. *BMJ*. 2010;340:c221. doi: 10.1136/bmj.c221
31. Nouwens F, de Lau LM, Visch-Brink EG, van de Sandt-Koenderman WM, Lingsma HF, Goosen S, Blom DM, Koudstaal PJ, Dippel DW. Efficacy of early cognitive-linguistic treatment for aphasia due to stroke: a randomised controlled trial (Rotterdam Aphasia Therapy Study-3). *Eur Stroke J*. 2017;2:126–136. doi: 10.1177/2396987317698327
32. Wertz RT, Weiss DG, Aten JL, Brookshire RH, García-Buñuel L, Holland AL, Kurtzke JF, LaPointe LL, Milianti FJ, Brannegan R. Comparison of clinic, home, and deferred language treatment for aphasia. A Veterans Administration Cooperative Study. *Arch Neurol*. 1986;43:653–658. doi: 10.1001/archneur.1986.00520070011008
33. Wilson SM, Eriksson DK, Brandt TH, Schneck SM, Lucanie JM, Burchfield AS, Charney S, Quillen IA, de Riesthal M, Kirshner HS, et al. Patterns of recovery from aphasia in the first 2 weeks after stroke. *J Speech Lang Hear Res*. 2019;62:723–732. doi: 10.1044/2018_JSLHR-L-18-0254