Evaluating Computer-based Treatment of Anomia: Results of Phase I Trials

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Todays Talk

- 1. Present a model for facilitating programmatic research to advance the state of evidence on a computer-assisted treatment (MTW).
- 2. Present and summarize the data that emerged from this project.
- 3. Discuss clinical implications

Learner Outcomes

Participants will be able to:

- Describe MossTalk's two main treatment modules and rationale for using each
- Summarize the evidence on efficacy including:
 - impact of therapy intensity
 - effectiveness when self administered
 - characteristics of patients who may benefit from MTW
- Identify factors to consider (e.g., barriers and facilitators) before using CAT

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What treatment approaches are empirically supported?

What is the level of evidence?

Phases of Research

- Pre-efficacy studies (Phase 1 and 2)
- Efficacy studies (Phase 3)
- Effectiveness (Phase 4 and 5)

Pre- efficacy Studies

Phase I

examines new treatments tests for therapeutic effect small, single subject designs

Phase II

optimizes procedures determines appropriate candidates dosage (intensity) further explores potential efficacy

Efficacy studies

Phase III: Clinical trial

 Controlled large group design
 Tests the efficacy of the treatment under ideal conditions

Effectiveness studies

- Phase IV
 - Potency under typical clinical conditions
- Phase V
 - Practical considerations(e.g., Cost-benefit analysis, consumer satisfaction)

Computer-Assisted Treatments: a popular movement

Computer-assisted treatments have potential to:

- Increase the intensity of therapy
- Improve outcome and efficiency of therapy
- Extend the period of rehabilitation

State of the evidence

A growing body of experimental literature attests to the benefits of this approach, for example:

- Lingraphica: Aftonomos, Steele, & Wertz, 1997
- ORLA: Cherney, Halper, Holland, & Cole, 2008
- Sentactics: Choy, Holland, Cole, & Thompson, 2009
- MossTalk Words: Fink, Brecher, Schwartz, & Robey, 2002

Large-scale (Phase 3) clinical trials, a level of evidence critical for establishing treatment efficacy are lacking

- Preliminary research (Phase I and II trials) needed to shape factors (patient selection criteria, intensity of administration, etc.) that are prerequisite to a Phase 3 clinical trial.
- Important to inform clinicians about the evidence available for treatment technology they may recommend.

What is MossTalk Words[®] (MTW)

- A computerized therapy system for aphasic adults with word retrieval deficits
- Provides extensive practice in word comprehension and production using multimodality cues and feedback
- Treatment modules
 - Theoretically motivated
 - based on effective treatments
 - routinely employed by clinicians

Two Modules

Cued Naming (CN):

Provides visual and auditory cues that can be systematically applied in a hierarchy to promote retrieval (Linebaugh & Lehner, 1977)

Multimodality Matching (MMM):

Encourages semantic processing to strengthening the association between words and pictures (Howard, Patterson, Franklin, Orchard-Lisle, & Morton, 1985a,b)



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Cued Naming Exercise Settings									
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U	Fill In 🔽 Spoken 🔽 Written								
Е	Word 🔽 Spoken 🔽 Written								
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Additional features

- Customize vocabulary
- Create homework assignments
- Track results

Fink, Brecher, Schwartz & Robey (2002) Phase 1 Study

- Investigated effects of CN Module: a hierarchical phonological cueing procedure
- Two conditions of instruction:
 - clinician guided (CG) condition
 - Partially self-guided (PSG)
- 6 subjects with primarily phonologically based deficits, 3 in each instruction condition

Conditions of instruction

- Clinician guided (CG)
 - worked on computer exercises with clinician 3 times/week
- Partially self-guided (PSG)
 - Worked on computer exercises 3 times/week
 - 1 day with clinician
 - 2 days independently

Prior Studies

Our study draws on prior studies without replicating any of them.

- From Linebaugh and Lehner we took the idea of individuating the cueing hierarchy- moving up and down hierarchy on each trial.
- From Howard et al., Raymer et al. and Thompson et al., we limited cues to phonological type.
- To provide maximum support for all severity levels, we included both written and spoken cues.

Study Aims

To assess acquisition, generalization and maintenance effects associated with computer-assisted hierarchical cueing.

Design

Single Subject (replicated)

➤Multiple Baseline Across Behaviors

Two conditions:
 Partially self-guided (PSG)
 Clinician-guided (CG).

Participants

- 6 chronic aphasic subjects
 - 5 M; 1 F
 - 54-64 yrs (mn= 60 yrs)
 - 2.3-7.5 yrs post onset (mn=4 yrs)
- Moderate-severe naming deficits
 - Naming severity: 17.8 77.4 % (PNT)
 - Aphasia Severity: 2 4 (BNT)
- Primarily phonological in nature
 - Phonological retrieval and/or
 - Phonological encoding
- Patients with central semantic deficits excluded
 - Mild semantic (2)

	Clini	cian Guid	ed	Partially Self Guided		
	GM	AS	BM	EL	EG	RH
Age-(rounded year)	54	64	60	59	63	63
Gender	М	М	М	F	Μ	М
Handedness	R	R	L	R	R	L
Time Post Onset (mos.)	92	40	28	34	40	61
Prior Language Therapy	12	9	9	5	8	11
Aphasia Subtype	Conduction	Broca	Anomic	Conduction C	Conduction	n Anomic
BDAE severity level	4	2	2	3	2	2

Table 1. Demographic information and language classification.

Training Procedure

- The Cued Naming module of MTW software delivered the picture stimuli, cues and feedback
- 6 of the 8 cues were used and presented in a hierarchy, individually determined for each subject

Multimodality Cues

Auditory cues

Written cues

- Initial phoneme
- Sent. completion
- Word repetition

- First letter
- Sent. completion
- Oral reading

Training conditions

 Clinician guided condition (CG) 3 participants

 Partially self-guided condition (PSG) 3 participants

Duration of Treatment

Subjects were treated 3 times a week

 Treatment continued until criterion was reached or for a maximum of 4 weeks

Outcome measures: naming

- Big Naming test- pre and post
- Daily naming probes of trained and untrained items during baseline, training, maintenance and follow-up phases
- Follow-up naming probes were administered after an average of 4 weeks

Outcome measures

Philadelphia Repetition Test (PRT)
Philadelphia Oral Reading Test (PORT)

Results

01 Set 1 • 12 Treatment- Set 1 Maintenance follow-up Baseline (6 weeks) • 18 • 19 01 Set 2 follow-up Treatment- Set 2 Baseline (6 weeks)

Figure 1. Subject GM (CG)

Figure 2. Subject AS (CG)







Figure 4. Subject EL (PSG)



sessions

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Figure 5. Subject EG (PSG)



Figure 6. Subject RH (PSG)



sessions

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		Se	t 1		Set 2		
		Baseline vs. Treatment	Baseline vs. Maintenance	4-week Follow-up	Baseline vs. Treatment	4-week Follow-up	
Group	Subject						
	GM	10.74	13.02	7.82	11.74	13.18	
CG	AS	7.78	8.37	8.49	3.91	3.91	
	BM	8.9	10.83	12.99	2.3	5.26	
	EL	5.5	4.86	6.05	2.67	2.73	
PSG	EG	9.76	10.81	12.92	13.51	16.05	
	RH	8.14	6	10.5	1.67	-0.35	
Benchma	rks (Beeso	on & Robey, 200	6): 4.0 (S); 7.	0 (M); 10.1(L)			

Study Results

- Training specific acquisition was demonstrated in both conditions for all subjects
 - 2 of 3 subjects in each group showed moderatestrong gains
 - 1 subject in each group showed weaker gains
 - Set 1 performance higher for 4 of 6 participants (2 from each group)
 - Gains were maintained when treatment was withdrawn
 - Small advantage for Clinician-guided group

Results: Generalization

- Limited and variable generalization patterns were noted in:
 - Naming of untrained items during training (EL and AS)
 - Oral Reading and Repetition
 - 339 item pre-post Naming test
 - All showed improved scores on trained items
 - GM and AS also showed significant improvement on untrained items

Conclusions

- Chronic aphasic subjects with moderate to severe phonologically-based naming impairments can benefit from a computerized cued naming protocol.
- Independent work on the computer can be an effective adjunct to therapy.

A model for facilitating research

Identify intervention (e.g., MossTalk Words)

Organize collaborative network

Site A Site B Site C

Evaluate results

Plan Phase 3 Clinical Trials

Organizing Collaborative network

Letters of invitation were sent to researchers and clinicians who work with individuals with aphasia.

Collaborators agreed to:

- * Participate in a brief training program
- * Complete a set of evaluation forms
- * Execute a controlled experiment of their design (research sites)
- * Use MTW in clinical setting (clinical sites)

Host provided ongoing training, technical assistance and support

Results of Dissemination

End of Year 1

* 3 Research groups had preliminary data on clinically relevant factors

- * Effectiveness for various etiologies and language impairements
- * Effectiveness when self administered
- * Impact of therapy intensity on outcomes

Subsequently

* Researchers presented and published several articles on clinically relevant aspects of MTW

Overview of Research Studies

Research Group	Module Used	Number of participants	Clinically Relevant Factors Studied					
			Etiologies & language impairment	Effectiveness when self- administered	Impact of therapy intensity			
Jokel & Rochon (2009)	CN	2	NPA (P)		\checkmark			
Jokel & Rochon (2010)	CN	1	SD (P)					
Raymer, et al (2006)	MMM	5	CVA 2(S) 3(P)		\checkmark			
Raymer, et al (2009)	MMM	4	CVA (S)	\checkmark				
Ramsberger & Marie (2007)	CN	4	CVA 1(S) 3 (P)	\checkmark	\checkmark			
Fink, et al (2002)	CN	6	CVA (P)	\checkmark				
CN=Cued Naming: MMM=Multimodality Matching:								

CN=Cued Naming; MMM=Multimodality Matching;

NPA=Nonfluent Progressive Aphasia; SD=Semantic Dementia;

S=semantic impairments; P=phonological impairments

Panel Presentation

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Elizabeth Rochon, PhD

 University of Toronto and Toronto Rehabilitation Institute

Anastasia Raymer Old Dominion University, Norfolk, VA.

THE END OF PART 1

PART 3: Summary and Discussion

Acquisition, Maintenance & Generalization

Acquisition

Most participants demonstrated measurable acquisition of trained items, though they varied in degree of improvement

Maintenance

Most maintained gains above baseline levels when treatment was withdrawn

maintenance phase

1 month follow-up

Generalization

Some evidence -but limited and variable

Does Intensity Matter?

- Significant improvement noted with intensive and non-intensive schedules
- Some advantage for greater intensity (Ramsberger, Raymer)
- When asked, participants preferred non-intense condition (Ramsberger)

Effectiveness when selfadministered

- Participants able to use computer independently
- Improvement noted when treatment was
 - Clinician guided
 - Partially self-guided
 - Completely self guided
- Effect sizes somewhat favor clinician- guided group

Who benefits?

- Adults with stroke related aphasia (15 studied); NPA (2 studied) and SD (1 studied)
- Moderate-severe production deficits
- Moderate-severe comprehension deficits
- Varied aphasia subtypes (Broca, Anomia, Conduction, Wernicke*)
 *limited # of Wernicke aphasia studied)

Modules/cues used

• CN

- Ramsberger (all written and spoken cues, individually determined)
- Fink et al (all but description cues, individually determined)
- Jokel & Rochon (printed and spoken cues (Study 1); written and spoken description (study 2)

• MMM

Raymer and colleagues

Barriers

- No computer in home or support
- Cognitive deficits
- Severe apraxia
 - need to be able to repeat or
 - respond to one of the cues provided by the computer)

Conclusions

- Findings confirm and extend Fink et al data:
- CN and MMM modules were effective in improving naming of **trained words** (acquisition and maintenance) for individuals with moderate severe naming impairments.
- Software effective with varied population (NPA, Semantic Dementia, and moderate-severe chronic aphasia)
- INTENSITY
 - Some advantage for greater intensity, but significant improvement noted with either intensive and non-intensive schedules.
- IINDEPENDENT work on computer can be an effective adjunct to clinician guided treatment
- BUT
 - Limited and variable generalization to untrained words or tasks

Future Directions

- Assess effects with new speech recognition component
- Incorporate more functional outcome measures

 Generalization to untrained production tasks (e.g., picture description, conversational sample)
- Prepare for Phase III Trial (RCT)

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