



RESEARCH INFORMATION for PEOPLE WITH APHASIA

Constraint Induced or Multi-Modal aphasia rehabilitation: A Randomised Controlled Trial (RCT) for stroke related chronic aphasia



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Miranda is a **speech pathologist** working in **research** at La Trobe University.

The study has been granted **permission** by the **La Trobe**

University Ethics Committee (Application No. 15-043)





WHAT IS THIS RESEARCH



ABOUT?

We know that living with aphasia is difficult.



Speech pathology can help. There are **lots** of types of treatments.



We don't know **which treatment** is the **best** for each person.



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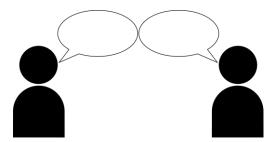


THIS STUDY

We will **compare** different treatments for people with aphasia.

1. Group A - Constraint-Induced Aphasia Therapy (CIAT)

CIAT therapy focuses on **talking**.



Therapy lasts for **30 hours** over **2 weeks**.

| Monday | Tuesday | Wed | Thursday | Thursday Friday Satur | | Sunday | |
|--------|---------|-----|----------|-----------------------|----------|--------|--|
| | | | | | | | |
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |

Therapy is delivered in a **group** of **three people** with aphasia and it is in addition to your usual aphasia therapy.



You **CAN** continue your usual aphasia therapy during this time.





2. Group B - Multi-Modality Aphasia Treatment (M-MAT)

M-MAT focuses on talking, drawing, writing and gestures.



Therapy lasts for **30 hours** over **2 weeks**.

drawing

| Monday | Tuesday | Wed | Thursday | nursday Friday S | | Sunday | |
|--------|---------|-----|----------|------------------|----------|--------|--|
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |

Therapy is delivered in a **group** of **three people** with aphasia and it is in addition to your usual aphasia therapy.



You **CAN** continue your usual aphasia therapy during this time.

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3. Group C - Usual Care

No extra treatment is given. You continue with your **normal** appointments or therapy for two weeks.

| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday |
|--------|---------|-----|----------|--------|----------|--------|
| | | | | | | |
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday |

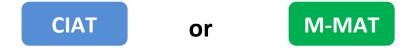
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4. Extra therapy for Group C

Later in the study, we will check that you are still able to participate in therapy. If yes, you may receive CIAT or M-MAT.



Therapy lasts for **30 hours** over **5 weeks**.

| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |
|--------|---------|-----|----------|------------------------|----------|--------|--|
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |
| Monday | Tuesday | Wed | Thursday | ursday Friday Saturday | | Sunday | |
| Monday | Tuesday | Wed | Thursday | Friday | Saturday | Sunday | |

Therapy is delivered in a **group** of **three people** with aphasia.



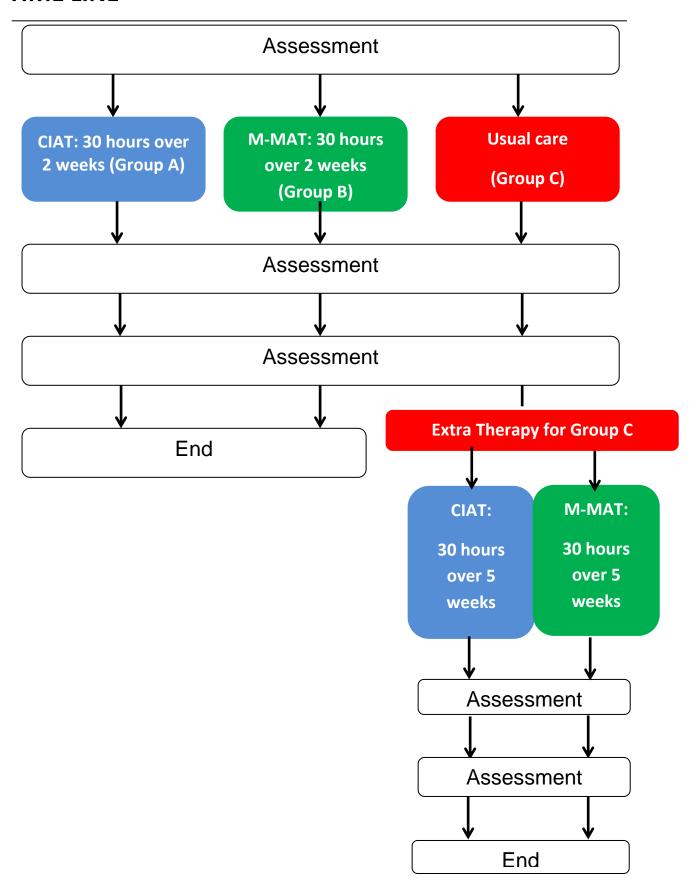
You **CAN** continue your usual aphasia therapy during this time.

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TIME LINE



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SEVEN PARTS IN THE STUDY

1. Assessment



You will complete **tests** of **language**, **communication**, **memory**, and **thinking**.

This will happen at a **clinic** or at your **home**.

You will be **asked to provide** copies of your **brain scans** from **after your stroke** if possible or **allow** one of the **researchers** to follow up a **copy** of these **scans for you**.

There will be $\bf 2$ - $\bf 3$ assessment sessions, with each lasting for $\bf 1$ - $\bf 2\frac{1}{2}$ hours.





2. Treatment

A **computer** will decide which **treatment** you will **receive.** This is done **randomly** and the research team have **no control over** which treatment you might receive.



You may receive:

CIAT to work on your **talking**

or

M-MAT to work on talking, writing, drawing and gestures

<u>or</u>

Usual Care – your **normal** activities.





3. Assessment



You will complete tests after the treatment period.

This will happen at a **clinic** or at your **home**.

There will be $\mathbf{2} - \mathbf{3}$ assessment sessions, lasting for $\mathbf{1} - \mathbf{2}$ hours.

Between this assessment and the last assessment (12 weeks), you will fill in a diary. We will text or call you regularly to remind you.





4. Follow-up assessment



You will complete **tests twelve weeks** after treatment.

This will happen at a **clinic** or at your **home**.

There will be 2 - 3 assessment sessions, lasting for $1 - 2\frac{1}{2}$ hours.





5. Extra Therapy for Group C

After the follow up assessment if you are in **Group C** we will **check** that you are **still able to participate** in therapy. If **yes**, you may receive **CIAT** or **M-MAT**.



You will receive:

CIAT to work on your **talking**

<u>or</u>

M-MAT to work on talking, writing, drawing and gestures.





6. Extra Therapy Assessment



You will complete tests after the treatment period.

This will happen at a **clinic** or at your **home**.

There will be 2 - 3 assessment sessions, lasting for $1 - 2\frac{1}{2}$ hours.

Between this assessment and the last assessment (12 weeks), you will fill in a diary. We will text or call you regularly to remind you.





7. Extra Therapy Follow-up assessment



You will complete **tests twelve weeks** after treatment.

This will happen at a **clinic** or at your **home**.

There will be 2 - 3 assessment sessions, lasting for $1 - 2\frac{1}{2}$ hours.





VIDEO RECORDING

During assessment and treatment, we will use a voice recorder and

camcorder to record what is happening.





Why?

- to collect information about the treatments
- make sure the **treatment** is being given **correctly.**





AM I THE RIGHT PERSON FOR THE RESEARCH?

You have to:

- be over 18 (eighteen) years of age



- have had your stroke/brain injury more than 6 months s

ago



- have **Aphasia**



- have used English well before your stroke/brain injury
- attend all study visits





have a carer/close other who can attend assessment visits







AM I THE RIGHT PERSON FOR THE RESEARCH?

You **cannot** participate if you:

- have a diagnosed cognitive deficit (problem with thinking or memory)
- have severe **apraxia** of speech



Cannot attend all study visits



If you have other **problems** such as:

- current depression, anxiety, or other mental illness
- taking illegal drugs
- vision or hearing problems

they need to be **managed** for you to **participate**.

For example: by wearing glasses or taking medication.





WHO WILL BE IN THE TREATMENT GROUPS?



There will be 2-3 (two-three)



people with aphasia in each group.



There will also be a **speech pathologist** and a **member of the research team** may also **be present**.



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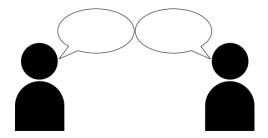
WHAT WILL THE



ACTIVITIES BE?

Constraint-Induced Aphasia Therapy (CIAT)

Practising words and sentences



2. Multi-Modality Aphasia Treatment (M-MAT)

- Practising words and sentences
- Practising drawing
- Practising writing
- Practising using **gestures**









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WHERE WILL THE GROUP HAPPEN?

| Room: | | | |
|-------------|--|------|------|
| Building: _ | | | |
| Address: _ | | | |

Family members are **not** included in the group.





We will collect the following types of information:

- your answers to questions and tests
- video recordings of you having a conversation with your carer or close other
- interviews with you

This information will be **used** and **stored**:

- on paper and
- on the **computer**

The information will be **kept** for at least **15 years**.





All information about you will be:

• labelled with a code (no names) of numbers and letters.

M.A. 123

M.R. 456

L.L. 059

The information will be stored safely in:

• a locked filing cabinet at La Trobe University



a computer with a password







Information about you will be seen by

- the research team
- and **staff** working on the **project**

After 15 years, the information may be destroyed.







We will only **share** your information **if**:

- we are asked under law



you or someone else is at risk of harm











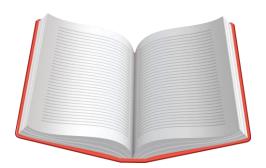
WHAT WILL HAPPEN TO THE RESULTS OF THE STUDY?

The **results** of this study will be:

- presented at conferences



- published in **journals**



Your identity will stay private.







HOW MIGHT THE RESEARCH BENEFIT ME?

You might benefit from:

Meeting new people and having conversations



- Practising your communication skills
- Trying **new things**
- Building confidence in yourself
- Taking part in something that will help other people
 with aphasia

We cannot guarantee that you will benefit.

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WHAT ARE THE COSTS OF TAKING PART?

You will need to **travel** to the treatment sessions. This will take time.



We will pay your travel expenses.







WHAT ARE THE RISKS OF TAKING PART?

You may feel **tired** after the assessment and treatment sessions.



You may feel **sad** or **depressed** after the assessment and treatment sessions.







WILL MY DECISION HAVE ANY CONSEQUENCES?

Your decision to:

- participate,
- not participate or
- stop participating



will not affect your relationship with:

- La Trobe University,
- your relationship with any of the researchers, or
- your **opportunity** to **receive** other **services** from them.





WHAT HAPPENS IF I AGREE TO PARTICIPATE?

We will ask you to sign a consent form



This means you:

- understand what we have explained to you,
- have had the chance to **ask** us **questions** and
- agree to all of this information







WHAT IF I WANT TO CHANGE MY MIND?

You have the **right** at **any time** to:



- stop the questions and testing,
- stop going to the treatment sessions, and
- receive free counselling if you want it.

If you decide to withdraw it may not be possible to withdraw your data from the study, however all data will be coded and will not identify you by name.





HOW DO I CONTACT THE RESEARCHERS?

Miranda Rose, Associate Professor, La Trobe University

Phone: (03) 9479 2776

Email: compareaphasia@latrobe.edu.au



WHAT CAN I CONTACT THE RESEARCHERS ABOUT?

- any questions or concerns



WHO DO I CONTACT ABOUT A COMPLAINT?

If you have any complaints or concerns about your participation in the study that the researcher has not been able to answer to your satisfaction, you may contact the Senior Human Ethics Officer, Ethics and Integrity, Research Office, La Trobe University, Victoria, 3086 (P: 03 9479 1443, E: humanethics@latrobe.edu.au). Please quote the application reference number 15-043.

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