







June is Aphasia Awareness Month! And 2025 is the Australian Aphasia Association's 25th birthday! 🮏

Aphasia is very common. More than 140,000 people in Australia have aphasia. Hundreds of thousands of other people love and care for someone with aphasia. However, most people have not heard of aphasia, and do not understand what aphasia is.

So, for Aphasia Awareness Month in 2025, we are sharing 25 ways to help raise awareness of aphasia!

- Some of these ideas are for people with aphasia.
- Some ideas are for family members.
- Some ideas are for health professionals.
- Some ideas are for aphasia researchers.
- Many of these ideas can be used by anyone.

We are sharing **ALL 25 ideas** at the start of the month. This way, you can have a look and start planning! Each week in June, we will also talk about some of these ideas in more detail - so keep an eye out.

If you use these ideas, please share them with us on social media!



Tell 5 people about aphasia. Then, ask them to tell 5 more people about **aphasia** as well.

- You could tell them in person, OR
- You could send a video, Facebook post, or flyer.

Look below for lots of ideas!



Follow the AAA Facebook page! Then, share our Facebook posts with your friends and family.



Click here to visit the AAA Facebook Page.

Make an aphasia ID card to keep in your wallet. Show the card to other people when you are out. You could show it to café staff, librarians, bus drivers or taxi drivers.



Click here to make a FREE printable Aphasia ID card.

Click here to buy "I have aphasia" cards from the AAA shop.

Show a video about aphasia to a friend, neighbour, or co-worker.

This can help them understand what aphasia is like.



Click here for a list of videos about aphasia.

Click here for aphasia-friendly videos about aphasia research.

Download our "communication tips" flyers.

• Show the flyers to friends and family members, OR



Print some flyers, and take them to a **local café**, gym or library.

This can help other people learn how to support communication.

Click here to download the AAA "communication tips" flyers. Click here to visit our aphasia resources webpage.



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Send a letter to your local politician to tell them about aphasia. This can help them understand aphasia, and support aphasia in government decisions.



Click here for a template that can help you write a letter.

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Print some "Join the AAA" brochures.

Take them to your GP clinic or speech pathology department. This will help more people with aphasia to know about the AAA.



Click here to download the "Join the AAA" brochure.

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Join (or start) a community aphasia group. You can meet other people with aphasia, and share your story.

Click here to learn more about aphasia groups.

Click here for a list of local aphasia groups.



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Come to the AAA National Conference! AAA

Conferences are for people with aphasia, by people
with aphasia. In 2025, the AAA Conference will be



in **Sydney** on **8**th **and 9**th **November.** Please **spread the word!**

Click here to read more and buy your tickets

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Make some art to show what aphasia means to you.

Draw or paint a picture, make a collage, or take a

photo. Share it on social media, with the hashtag #Aphasia.

Click here to see the aphasia artwork from our 2023 conference!





Join in the #WordsForAphasia challenge on social media!

- Think of a word that shows what aphasia means to you.
- Write the word on a piece of paper in large letters.
- Take a photo of yourself holding up your word.
 (Or, just take a photo of the piece of paper by itself).
- Post your photo on social media with the hashtags #WordsForAphasia and #Aphasia

Click here for some examples and templates



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Visit the Stroke Foundation webpage on aphasia and young stroke. Share the page on Facebook, or show it to friends and family. This web page has lots of information about aphasia for younger people.

Click here to visit the Young Stroke "Aphasia" webpage.

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"Wear" your aphasia loud and proud! This could be:

- a t-shirt or jumper about aphasia
- a **hat** with the word "aphasia"
- some people even have an aphasia tattoo!

These are **great ways** to **start conversations** about **aphasia** and **raise awareness** about aphasia.

Click here to see some examples of people "wearing" aphasia.

Click here to buy a shirt from Aphasia Recovery Connection.



Talk with others about mental health and wellbeing with aphasia. This can help them to understand how aphasia can affect people. It can also remind them to check in on their loved ones with aphasia.



Click here to read about mental health and aphasia.





Run a **fund-raising event**. This could be a **bake sale**, a **morning tea**, a **dinner**, or you could **sell tickets** to a **trivia or movie night**.



Click here for more ideas about aphasia fundraising.

Click here to read about our Long Lunches for Aphasia.



Ask local businesses to use aphasia-friendly information. This means using shorter sentences, larger font and pictures. This can be helpful on menus, signs, flyers, and forms. Aphasia-friendly documents are easier for everyone to understand!



Click here to see guidelines on aphasia-friendly information.



Download the **Aphasia Handbook** and **show it to a friend.** The **AAA** and the **Stroke Foundation**made this handbook **together**. It has **lots of information** about **aphasia**.



Click here to download the Aphasia Handbook for FREE.

Click here to BUY hard copies of the Aphasia Handbook.

Click here to visit our Aphasia Resources webpage.



Download the Primary Progressive Aphasia (PPA) Guide and show it to a friend. PPA is a rare type of dementia that affects language. The PPA Guide is a booklet with lots of information about PPA.



Click here to download the PPA Guide for FREE.

Click here to visit our Aphasia Resources webpage.





Join in with #RockingAphasia on social media.

- Find a smooth rock or pebble.
- Write **#RockingAphasia** on the rock.





- Hide the rock somewhere (park, bus shelter, anywhere)
- Take a photo of your rock in its hiding place.
- Post the photo with the hashtag #RockingAphasia

Curious people will find the rocks, and look up "aphasia".

Click here to learn more about #RockingAphasia.



Be a part of aphasia research! You can be a research participant, or help to design aphasia research. This is a great way to help people, and



raise aphasia awareness. We share research opportunities with our members on Facebook and our member mailing list.

Join the AAA to hear about future research opportunities.

Visit the Aphasia Research Library to read aphasia research.



Run an **information session** about **aphasia** in your **local community**. You could do this at a **library**, a **community centre**, a **Rotary club**, or a **U3A meeting**. These organisations are often **happy to help!** Your session could be about:

- Understanding aphasia
- Supporting communication with aphasia
- Making businesses more aphasia-friendly



<u>Click here for a template of PowerPoint slides you can modify.</u>

(They were made by the Aphasia CRE at La Trobe University)





If you are a **family member – share** your **own experience** of **aphasia**. You could **contact** your **local radio station**, or **write** to a **newspaper**. This can **help**

to raise awareness of aphasia and how it affects families.

<u>Click here for an example radio interview (Bridget Noble).</u>
<u>Click here for an example video interview (Kim Beesley).</u>



about the AAA! The AAA is here to help and support people with aphasia and their loved ones. We want everyone with aphasia to know about the AAA.

Click here to download the "Join the AAA" brochure.



If you are an **aphasia researcher**, make **aphasia-friendly video abstracts**. These are **short videos** that **explain research** in **aphasia-friendly**



ways. This can raise awareness of important aphasia research.

Click here to learn about making aphasia-friendly abstracts.

Click here for examples of aphasia-friendly abstracts.



Join the AAA!

This helps us to keep working for the aphasia community. We organise events and share resources to support people with aphasia. We work hard to raise awareness about aphasia.

Click here to become a member of the AAA!