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# Report

# WWDA Virtual Centre

# Co-Design Workshops

# 4 – 19 July 2019

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## Executive Summary

WWDA undertook a series of workshops around Australia with diverse groups of women, to find out what they needed and wanted from the Virtual Centre, a web-based information source to address five priority areas of need: rights, leadership, decision-making, safety from violence and sexual and reproductive health.

The workshops involved smart, wise, humble, proud, tenacious and energising women of all ages. The purpose of the workshops was to ensure that the Virtual Centre is both responsive and reflects the diversity of women with disability around Australia. The project brief stated:

The facilitation of the co-design workshops will be collaborative, adaptable to each workshop’s context and will harness the unique expertise, insight and skills of local women with disability.

Dr Julia Wolfson was hired to design and facilitate the workshops. The co-design format allowed the WWDA team and facilitator to be learners, and the participants to educate, inform and make visible what was on their minds and what they felt would be useful on the Virtual Centre. The workshops were iterative and adaptive in that each workshop benefitted from the experiences and insights of prior workshops in refinement of structure, team roles, facilitation style and content focus.

The WWDA project manager led the project and attended all workshops, accompanied in some locations by the WWDA Senior Content Officer and WWDA Communications Officer. A local co-facilitator – a women with a disability – was recruited in each location on the advice of the local host organisation to provide insight into the local context and to ensure that each workshop was truly for and by women with disability.

The website design agency [Handbuilt Creative](https://www.handbuiltcreative.com.au/) was contracted to undertake the design and development of the Virtual Centre. Handbuilt staff attended three of the six workshops to gain first-hand insight and experience.

The workshops were structured in two parts. Part one was a highly interactive and frank discussion across the five priority areas. Part two gave individuals the opportunity to take part in an initial prototype testing activity to inform the development of the Virtual Centre layout.

Workshops were held in six locations: Hobart, Canberra, Melbourne, Brisbane, Tennant Creek and Perth. In total 45 women between ages of 18 and 65 participated.

### Summary of key learnings from the workshops

1. Overwhelming support for the creation of the Virtual Centre and appreciation for non-tokenistic involvement of women with disability. The participants valued being part of the website development so early on, and the opportunity to contribute meaningfully through the workshops.
2. Stories, stories, stories. There was unanimous request for stories and videos to be a feature of the website. Women want to see themselves, their strengths and diversity reflected in the website. They want to see, hear and read stories from their peers as this can create a sense of “I am not alone”.
3. Practical guides, checklists and resources for navigating specific situations should be provided on the website. This could include situations such as going for a job interview, when and how to talk about your disability, how to tell your carer to stop being controlling.
4. The layout and design of the website should make it easy to use, including for those who require a support person to assist them.
5. The Virtual Centre has an important role as a self-advocacy tool because it is equipping and resourcing women to be self-advocates, and to advocate for others.
6. There was widespread appreciation that the Virtual Centre is offering information on sensitive and often taboo topics such as violence, gender and sex.
7. There was a widespread desire for the website to provide a virtual forum for mentoring, connectivity and interaction on topics that matter to women with disability.



Participants in discussion at the Canberra workshop

## Part 1: Context

### 1.1 Background

The Virtual Centre for women and girls with disability (‘the Virtual Centre’) is a new website that will provide accessible, inclusive and practical resources and information for women and girls with disability. The Virtual Centre is being created by Women with Disabilities Australia (WWDA) using funding provided by the National Disability Insurance Scheme (NDIS) Information Linkages and Capacity Building (ILC) program. Project Planning commenced in late 2018 and the Virtual Centre is due to launch in early 2020.

The goal of the Virtual Centre project is:

“To develop a centralised and sustainable Virtual Centre for Women and Girls with Disability that provides accessible and quality information on their rights to: decision-making and agency; leadership and participation; sexual and reproductive health; and safety from all forms of violence.”

The Virtual Centre will publish information and resources on the priority areas listed in the project goal. These priorities have been selected by women with disability in Australia through multiple structured consultation processes, including the [WWDA National Forum (2017](http://wwda.org.au/report-wwda-national-forum-for-women-and-girls-with-disability-proceedings-and-outcomes/)) and the [National Symposium on Violence Against Women](http://wwda.org.au/wp-content/uploads/2013/12/STV_Outcomes_Paper.pdf) (2013). WWDA has published position statements that provide more information on each of these priority areas, which are available here: <http://wwda.org.au/papers/toolkit/position-statements/>.

The primary target audience for the Virtual Centre is women and girls (aged 15 years+) with disability in Australia. WWDA recognises that the unique experiences of young women and girls with disability under the age of 15 years may require different considerations, which are beyond the scope of this project.

In early May 2019 a competitive tendering process was undertaken to recruit a website design agency. Handbuilt Creative, based in Richmond Tasmania,was the successful tenderer, largely due to their experience in designing user-driven, accessible websites.

#### 1.2 Participation and co-design

WWDA recognises that the meaningful participation of women and girls with disability in the design, implementation, monitoring and evaluation of targeted measures to advance their rights, is critical if they are to realise those rights.

The Virtual Centre project is utilising a co-design approach to ensure that the project is led and influenced by women and girls with disability at every stage.

Co-design is a collaborative, participatory approach to design which actively engages and places decision-making capacity in the hands of end users[[1]](#footnote-1). Co-design goes beyond simply involving stakeholders in the beginning stages of design. Participants engage in problem-solving, decision-making and feedback processes throughout the project cycle, allowing for genuine and meaningful contribution[[2]](#footnote-2),[[3]](#footnote-3). Co-design principles are widely recognised as best-practice in the disability sector and utilised by the National Disability Insurance Agency (NDIA) and National Disability Services (NDS) to ensure service-user inclusion and satisfaction[[4]](#footnote-4).

The Virtual Centre project’s co-design approach includes ensuring the meaningful participation of women with disability in governance and decision-making bodies. The Virtual Centre Project Steering Committee (PSC) oversees the governance and strategic direction of the Virtual Centre. It comprises seven women who are experts in their areas and have deep networks throughout the women’s and disability sectors. Four of these women identify as women with disability.

The Virtual Centre Expert Advisory Panel (EAP) is comprised of over 40 women with disability. This diverse group is consulted with regularly to review and provide feedback and input on Virtual Centre content and design. The EAP help to ensure the website is accessible, inclusive and relevant, and importantly, guarantee that the Virtual Centre is developed by women and girls with disability for women and girls with disability.

### 1.3 The workshops

As part of the co-design approach, participatory workshops were held around Australia in the mid stages of the Virtual Centre development. The purpose of these workshops was to seek feedback and input from a range of women with disability on the design, structure and content of the Virtual Centre. The workshops were to be engaging, inclusive and positive events that provided a safe and enjoyable space for women with disability to contribute to the design and development of the Virtual Centre. The workshops were intended also for the project team to capture input and feedback from women and girls with disability who may be harder to reach and/or who prefer to be engaged with in person, in order to have the opportunity to contribute meaningfully.

Collating the feedback from the co-design workshops in this report and acting on it will ensure that the Virtual Centre is relevant, accessible, collaborative and usable for a wide range of women and girls with disability.

WWDA’s commitment to genuine co-design led to hiring a workshop facilitator highly experienced in co-design in the disability space as well as across diverse sectors, in Australia and internationally[[5]](#footnote-5).

### 1.4 Report scope

Part 1 of this report presents the context of the co-design workshops. Part 2 covers the information and feedback gathered from participants at all six workshops. Part 3 offers some reflections and suggestions for the website, with possible directions for the Virtual Centre. To assist the reader to get an immediate overview of needs and wants that emerged, a summary of highlights is presented in the next section.

The report focuses on the content aspect of the Virtual Centre website, in five core focus areas, with some additional topic areas that emerged during the workshops. As the Virtual Centre is for and by women and girls with disability, decisions about prioritisation of the material will be made by the project team in consultation with the EAP and PSC and are beyond the brief of this report.

### 1.5 Needs and wants that emerged: Summary

##### Overall learnings from the workshops

There was overwhelming support for the creation of the Virtual Centre, and non-tokenistic involvement. The women valued being part of its development so early on, and to contribute through the workshops.

Stories, stories, stories. There was unanimous request for stories, videos and showing things rather than just explaining them in text. Women want to see themselves, their strengths and diversity reflected in the website.

Practical guides, checklists and resources are needed for navigating specific situations such as: going for a job interview, when and how to talk about your disability and how to tell your carer to stop being controlling or patronising.

The Virtual Centre has an important role as a self-advocacy tool because it is equipping and resourcing women to be self-advocates, and advocate for others.

There is widespread appreciation that the Virtual Centre is offering information on sensitive and often taboo topics such as violence, gender relations and sex.

There is a widespread desire for the website to provide a virtual forum for mentoring, connectivity and interaction on topics that matter to women with disability.

##### Website design

It was clear from the feedback that there is a tension between accessibility and complex functionality built into design decisions. It became evident that the Virtual Centre website may need to trade some functionality aspects to extend the accessibility of the site to anyone who can use a computer (even with help).

This means cleaner design, clearer fonts, icons with thicker strokes, contrasting colours. It became clear that a well-designed, accessible website was far more preferred to a powerful online resource, and Handbuilt Creative is already taking steps to reflect this in their approach.

### 1.6 Workshop locations and participants

The local workshops offered a unique opportunity to engage with women who can most meaningfully contribute in a face-to-face environment. The workshops also welcomed and encouraged a diverse group of women to attend. The range of disabilities represented included physical, intellectual, learning and psychosocial. Workshop participant numbers ranged from four to 12 across the workshops. In total 45 women participated in the events, with ages ranging from 18 to 65. Further details on workshop participation is available in Appendix 1.

The workshops were held in partnership with a local advocacy or service organisation. The six locations and partners were:

1. Hobart, Tasmania (in partnership with [Speak Out Association of Tasmania](http://www.speakoutadvocacy.org/))
2. Canberra, ACT (in partnership with [Women with Disabilities ACT](https://www.wwdact.org.au/))
3. Melbourne, Victoria (in partnership with [Women with Disabilities Victoria](https://www.wdv.org.au/))
4. Brisbane, Queensland (in partnership with [WWILD](https://wwild.org.au/))
5. Tennant Creek, Northern Territory (in partnership with [First Peoples Disability Network](https://fpdn.org.au/))
6. Perth, Western Australia (in partnership with [Women with Disabilities WA](https://wwdwa.org.au/)).

### 1.7 Workshop approach

##### Guiding principles and outcomes

A key principle of the Virtual Centre is that it is for and by women and girls with disability. The principles that guide the Virtual Centre also guided the workshops:

1. Women and girls with disability actively contribute to leading, shaping and influencing their community
2. Women and girls with disability are connected and have the information they need to make decisions and choices

The workshops used a co-design approach(see section 1.2).

The purpose of the workshops was to augment the feedback and recommendations gathered thus far from the PSC and EAP members, with direct feedback in-person from women in diverse settings and with diverse requirements.

The focus of the co-design workshops was for the WWDA project team to learn how to make the Virtual Centre content and functionality most relevant, usable and accessible.

##### Facilitation team

Dr. Julia Wolfson of Turning Forward (see [www.turningforward.org](http://www.turningforward.org)) was hired to facilitate the workshops, report on the learnings and make recommendations. The WWDA Project Manager attended all workshops, and the WWDA Senior Content Officer and WWDA Communications and Marketing Officer attended most workshops.

Every workshop was co-facilitated by a woman with disability. Before the start of each workshop, the travelling WWDA team met with local facilitators to learn about the local context and to go through the workshop program.

The local facilitator welcomed participants, acknowledged country and elders, and took ownership for creating a welcoming ambience to encourage participation.

There was no lengthy lecture of content provided. The topic for discussion was given by the facilitators, the participants provided content and responded to open questions.

The flexible style of facilitation and the use of visual posters stuck up on the walls around the room enabled spontaneous contributions to come out across themes and be immediately captured. This also made it easy to pick up where the discussions left off if the discussion shifted to a deeper level of emerging feelings and concerns.

The WWDA facilitation team viewed themselves as listeners and the participants as teachers.

An example of a workshop agenda is provided in Appendix 4.

##### Feedback

Participants were invited to provide feedback at the end of the workshop about the extent to which they felt free to participate and express their views. They were also invited to give advice to the facilitators on how to make future workshop more inclusive and participative.

The WWDA Project Manager undertook to stay in touch with participants about ongoing progress of the website moving forward. This included future opportunities for user testing of the website in their own time, from their own devices.

All workshop participants were also given the opportunity to review and provide comments on this workshop report prior to its finalisation. An Easy Read version of the Executive Summary will also be made available.

### 1.8 Ensuring accessibility

All workshops were held in accessible venues. Local partner organisations in each location assisted the travelling WWDA team by providing information on each participant’s interests, communication style and accessibility requirements prior to each workshop. If relevant, the local team ensured that participants had a support person or assistant present to enable their full participation.

All workshop materials were provided in alternative formats where required – this included the development of Easy English summaries and consent forms (see Appendix 3). The flexible facilitation style and the use of both verbal and visual communication methods also helped to ensure inclusion of all participants.

### 1.9 Emotional well-being and safety

The workshops had an advocacy focus and were not primarily intended as a space for sharing personal experiences. Even so, difficult experiences from the past and present naturally surfaced. The facilitation approach was open to emotional experiences and gave them space, appreciating personal strengths and resilience that shone through.

Various visual cards were strewn around the tables that could be used by participants to indicate if they needed to stop, pause, take a break, or speak with someone privately.

The local teams and facilitators had a vital role in creating a welcoming, energised and culturally safe environment that encouraged participation. In some localities (such as ACT and QLD), local team members were certified in mental health first aid and/or violence response and were available to respond in the event of a disclosure or need for support.

Local teams were asked to prepare a referral list of counsellors, therapists and other specialist providers in their area, and have access to a local counsellor if not already available within their organisation. The WWDA project team brought along and distributed 1800RESPECT flyers.

When hot topics arose, the facilitator took time to pause and check whether the group wanted to keep going. Asking permission any time the atmosphere shifted and deepened, or when someone began to tell a painful story, helped to create safety in the group.

Another safety practice was to frame the more painful topics as an advocacy discussion. The facilitator began such conversations by thanking everyone for being present and appreciating the survival abilities and strengths in every person to go through the ups and downs of life, and to endure things that no human being should have to go through. Questions were posed: ‘What got you through?’ ‘What might the Virtual Centre show or do, to make this different for others coming after you?’ ‘What would you say to someone who went through what you went through, and didn't know what to do?’ ‘Has anyone else here experienced that? Ah! You are not alone!’. This made it easier to stay with the more generalised advocacy purpose.

All participants signed a consent form prior to the workshop commencing (see Appendix 2 and 3). This ensured that they were all comfortable with the workshops being recorded, allowed them to decide whether they were included in photos and explained how and where their feedback would be used.

### 1.10 Website technical design

The functionality, design and accessibility of the website was also discussed in the workshops. Handbuilt Creative CEO, David Shering was present at the co-design workshops in Hobart, Melbourne and Canberra. The workshops sought to learn how diverse users will most easily interact with the site, and understand accessibility needs and considerations. In particular, Handbuilt Creative wanted to learn who was going to use the website, and what functions the users might typically perform when using a website. Handbuilt Creative and the WWDA project team were working from the assumption that users are able to access the internet and navigate a website (either on their own or with assistance).

At each workshop participants were invited to try out three prototype wireframes in a personal interview with a WWDA/Handbuilt team member and provide feedback on their unique accessibility and user preferences. In total, 32 women completed the activity. The response form is attached at Appendix 5.

It was clear [from workshop discussions] that a well-designed, accessible website was far more preferred to a powerful online resource, and we have taken steps to reflect this in our approach. This means cleaner design, clearer fonts, icons with thicker strokes, more contrasting colours and a range of other techniques.

David Shering, Handbuilt Creative

A ‘user profile’ spreadsheet was completed after each workshop which detailed each participant’s responses to the prototype wireframe testing activity and captured any additional feedback and insight they provided.



A participant at the Brisbane workshop holds up her design of the Virtual Centre logo

## Part 2: Needs and wants: what we learned

This section provides a synthesis of the content learning from the workshops under the five topic headings. The WWDA project team and Expert Advisory Panel can use this information to prioritise feedback and requests.

The order of presentation here is not intended to convey a hierarchy of importance. Direct quotes from the workshops are provided in italics. Where relevant to the content, the specific location is identified. In some instances, a participant asked for their name to be attached to the quote. [Words in brackets] indicates they were added by the authors.

### 2.1 Distinct themes across the six workshops

There were themes common to all six workshops (as outlined in the executive summary and section 1.3). Each workshop also had its own unique flavour.

A variety of hot topics popped up and were given space to be processed in each session. When someone had something to say, however seemingly off topic, the group listened. Invariably the person was carrying something urgent and meaningful to the whole group that was relevant.

In Hobart, women living in residential settings spoke of the everyday restrictions on their environment, freedom of movement, and choices and decisions made for them by staff.

In Canberra, the women emphasised accessibility and adaptive communication to enable women with a myriad of communication styles to express themselves. Hidden disabilities folded under mental health stigma was also a topic – women expressed that they are not seen or believed, and their situation is dismissed. Women expressed a lot of anger at unwanted touch by people trying to be helpful but actually “*clutching your arm, holding your cane, or touching your wheelchair”*.

In Melbourne many women expressed immense frustrations with NDIS planning and responsiveness. Some expressed the painful experience of being seen as less-than within the disability community. One’s worth ­­– we were told – varies depending on whether you were born with your situation or acquired it later in life.

In Brisbane, women living in the aftermath of violence spoke of the difficulties getting support to get back on their feet.

In Tennant Creek the women strongly advocated for the content and presentation of the Virtual Centre to be culturally relevant and mirror the lives of indigenous women and girls. In addition, the women spoke of the cycle of violence: who and what creates it, and what might help to break it.

In Perth the women emphasised the vexed topic of power and misuse of power by support workers. We learned of the agony of put downs, basic disrespect and restrictions on basic life choices. As one participant said: “*It starts out all right, then they take over. I’m not your client, I’m your boss*”. Another strong theme was the power of individual agency in influencing practical improvements to a more inclusive and accessible public environment.

### 2.2 Feedback on overall structure

This section covers the feedback in each topic area from all six workshops, comments on design and accessibility, as well as additional topic areas that emerged across the workshops, such as the National Disability Insurance Scheme (NDIS).

#### 2.2.1 Sections to add:

What is disability? “Your disability may be invisible on the outside, but its real!”

United Nations (UN) definitions

Relevant statistics: e.g. % of population have a disability, what kinds of disabilities.

#### 2.2.2 Ideas for each of the five focus area sections:

Provide a brief summary of the topic, a definition of the topic area, Frequently Asked Questions (FAQ) and reliable and verified links to more information.

On each page include contact information to make it easy to find without going back.

Provide legal frameworks – national and state-by-state pulled together in one spot

Statistics: Include summary statistics in every section relevant to that section

#### 2.2.3 General requests:

Definitions of all key terms used

Examples and case studies

A suggestion box and ability for users to submit content.

Ensure the site is constantly updated.

### 2.3 Your Rights: Overview page

While every topic on the Virtual Centre covers rights, the Your Rights sections will give an overview. Some suggestions from participants:

#### 2.3.1 Summary – key messages

The Australian rights framework

UN rights framework

Legal rights – being a decision maker

Information on consent

Information on discrimination

Rights are equitable and universal

Intersectionality – rights cut across various areas of life.

#### 2.3.2 Statistics

Incarceration rates of people with disabilities

Gender inequities.

#### 2.3.3 Frameworks

The Australian rights framework

UN Conventions (include Easy English/Read versions)

Your legal rights as a decision maker

About consent

Advocacy information.

#### 2.3.4 About rights: what are your rights?

Clarify in practical ways what exercising rights might mean to different people. This could be in a story, talking book or some other lively way.

Respect, equity and opportunity: The right to be happy. The right to enjoy an everyday life with the same opportunities that people in your community and in Australia expect and enjoy. The right to non-discrimination. The right to be seen, heard and respected.

“The right to respect: to be listened to, not to be told ‘you are stupid because you have a disability’. The right to get help when you don't understand”.

Safety: The right to be and feel safe.

Home: The right to move house if you want to. The right to have access to country and be on country for end of life.

“Government housing has no insulation. It’s freezing. Great rate of deaths from the flu here. People are freezing in their homes” and “People over 50 are facing homelessness and women with disabilities are in violent situations”.

Fair treatment: The right to complain about rights violations, restrictive practices and unfair treatment, and get redress.

Relationships: The right to an intimate relationship, where intimacy is defined by you. The right to sexual relationships and respectful relationships.

Care: The right to good care and support that respects your rights. The right to be supported by a person of your preferred gender.

“I want to say to my support worker, I am not your client, I’m your boss! One lady berated me for being a control freak. I’m allowed to be! It’s my life! They infantilise you. It starts out alright, after a while they take over”.

Family: The right to have a child and raise your child and show what a great parent you can be – not have your child taken away.

Self-advocacy: The right to make your own decisions. Your right to be included in decisions about things that affect you and are important to you. Right to use bush tucker.

Participation: Your right to be included and participate in a group you are part of: *“Don't just assume I can’t do it”.* The right to a case-worker to help you get out of the house and find low-cost or free opportunities in your local community. To participate, learn, develop interests such as find an exercise group or an art group, contribute, meet people.

Information: Your right to information about things that are important to you, in an accessible format.

Education: Your right to be informed about how society works. Your right to education. The right of students in special schools to be provided with equitable education.

Privacy: Your right to privacy regarding your records and personal information. When there is a third party involved, how to assert yourself to get accurate information and assert your right to give permission to have your information passed on. Your right to non-disclosure and privacy about your disability.

Consent to touch: Your rights with your body. Your right to not be touched without consent. “*Ask me if you can touch me, even if you are helping me get upstairs”.*

Employment: Your rights at work. Your right to equity in employment. Your right to gender equity in the workplace, and an equitable wage in the workplace. The right to progress on a career path. Your right to choose your work.

“My right to job support to learn the job and not get fired for making a mistake”.

Advocate: Your right to an external advocate/supporter.

#### 2.3.5 Videos, stories moving books

Stories of survival. Role models living ordinary lives. Messages of encouragement to keep going through the tough times. “How to” stories

Stories to encourage you to reinforce what you are good at and keep moving forward

Stories that show what to do if people are mean to you

Message to parents: “Let me do my own thing”.

#### 2.3.6 Tips, tools and checklists

How to make a complaint

Hiring your own staff to get the support person you like and the support you want and need

What to do if…. with examples: “*My support worker texts while driving. It’s dangerous!” “The tap in my shower is broken, I have asked and asked to get it fixed I’m scared to get in the shower in case I get burned*”   
Go to someone you trust

What accommodations do you have the right to ask for in different areas of life (e.g. work), and how to go about it?

Some tips as to how to enter a space with any professional – and have adequate time and space to tell what is going on

“*A checklist to help me understand my own access requirements”*

What you can expect when you go to a service for assistance?

How to strengthen your self-esteem.

#### 2.3.7 Links to quality information

Your rights in the criminal justice system – for instance, if you are arrested.

Finding an advocate. Some girls and women are good at advocating for themselves. For some the barriers are too great to advocate and negotiate on their own. Find an advocate if you need it. Sometimes this is someone beyond your paid support and natural support network whose sole function is to support you to advocate for your rights

Your rights to have the carer you want, and what to do about a carer who overpowers you

Link to UN conventions, including Easy Read

Links to UN Declaration on the Rights of Indigenous Peoples

Links to Facebook groups for peer exchange, information and support to know and exercise your rights

Link to information about carer boundaries and limits – what you can expect

Link to Human Right Commission

Link to health rights – a document on People with Disabilities (PWD) website

Links to advocacy information and organisations (e.g. Persons with Disabilities Australia)

Link to heath consumers councils

Link to AdvoCare – for women on the aged pension

Link to culturally and linguistically diverse (CALD) websites with information about rights

Links to Aboriginal and Torres Strait Islander (ATSI) websites with information about rights

Links to vision impaired and deaf community websites

Where to get practical assistance to navigate services, including how to fill out forms etc (e.g. Anglicare’s Alina in Brisbane).

#### 2.3.8 Myths and facts

To educate and raise awareness among carers, family members, service providers, the broader community, civic institutions and civil society.

“Don't compare me with others. I am an individual with unique gifts”

Women with disability are often seen as inadequate mothers because of their disability. Include a mother’s story.



A participant contributes to the discussion at the Tennant Creek workshop

### 2.4 Leadership and Opportunities

#### 2.4.1 About leadership

“Be a leader by leading others to become leaders”

##### What is leadership?

What it is and isn’t - what does good leadership look like?

What is leadership mentoring? The value of finding a leadership mentor and being a mentor. The boundaries of mentoring

Tokenism – and what to do about it

Getting a job for some is an act of leadership, and provides leadership opportunities

What is advocacy? The right to advocacy

Encourage younger women to progress on a career path

What is involved in being a leader and what are some benefits?

Write letters to government

Support others to speak up

Join an executive committee

Be a role model for other self-advocates

Find a self-advocate mentor

Learn how to take responsibility and get organised

Be a representative – think broader than just your own needs

Experience diversity.

“Role models and mentors are very important to help you build confidence by showing others how. I went to the United Nations in the USA. Presenting on stage is scary, but you can just do it, and get courage by seeing others do it”.

“Mentors are like a best friend – there for you no matter what”

“Carer pressure can be limiting, for example, the carer’s mindset and values can limit your sexuality. If you come up against barriers, NO, keep at it, don't give up. Seek support/mentoring/assistance to pursue your path”.

#### 2.4.2 Stories, talking books, videos, resources

“There are opportunities everywhere. Be open. They present themselves when you least expect it” Liz, Perth

“You can grow up to be anything you want to be” Jackie, Perth

“Don't dream your life away, believe in yourself” Elizabeth, Perth

[What helps you find your personal power when the situation is agony? Finding a role model. Easing the path for others by sharing your story and what got you through. You have important lessons to teach about surviving, bouncing back from hard situations and using your experience to help others].

Get courage to overcome shyness: “*People used to walk all over me*”.

Videos and quotes in the words of indigenous elders

Leadership books – inspiring autobiographies and picture books are uplifting

Music – songwriters who create their stories in song

Guidance for starting a project (e.g. a business, a group etc)

How to talk about your disability at an initial job interview

Include inspirational quotes – see Brain Injury Matters

Examples of people who have gone through various leadership courses

Crystal – Sistergirl Australian of the Year Awards NT

Purple house – dialysis on country – Sarah Brown

Elizabeth’s story: Elizabeth went through the public toilets and recorded exactly what was needed to make them accessible, wrote to the council, and got action.

“Snap, send, solve” App – you can use this to take a photo of something that isn’t accessible (e.g. footpath) and it will be sent to the council to action.

#### 2.4.3 Opportunities: Things you can do to get involved and make a difference

Get asked to join a group

Join a youth group

Join health, fitness and good eating groups

Join a special interest group: *“I went on the cat walk!”*

Link up with international groups

Start a self-advocacy group, join a self-advocacy group and learn self-advocacy skills

Go to leadership camps – where you can make new friends, meet people and learn to trust

Meet people

Sign up for newsletters which often have information about events and groups

Join a mother’s group

Join activity groups and interest groups in your region/state to meet people, pursue interests, and develop a sense of belonging and expand your network

Join leadership trainings, camps and workshops. Learn to develop yourself and help others

Volunteer in your local community

Make a difference in your local council’s Disability and Inclusion Plan

Join advisory groups at your local hospital/council

#### 2.4.4 Checklists, tips and tools

How to do publicity and media

How to plan for accessibility in getting around safely when travelling

How to prepare and dress for an interview. How to talk about disability in a job interview. Where to access interview coaching

Writing a CV and including your leadership role. Provide an example of a good CV

How to fill in an expression of Interest form – provide an example

Things you can do to facilitate end of life on country

Where to go and how to go about finding a mentor

Links

#### 2.4.5 Information about governance opportunities and development

##### Leadership events and links

Redefining Leadership Exhibition in ACT

Stories on social media

Self-advocacy groups – national and state by state. Who you can contact in advocacy organisations?

Organisations offering mentoring opportunities

Organisations willing to be contacted if you want to offer yourself as a mentor

A.N.D employment link

Leadership events and training opportunities, where to learn self-advocacy skills (state-by-state links)

Applying for leadership scholarships

Disability Leadership Institute Australia resources

ATchat Facebook page – Peer support for getting a mentoring program up and running

Links to forums such as PWD – where you can learn, build up networks and ask questions.

##### Forums

Have an online forum for exchange of information. A space to grow naturally into mentorship roles for younger and older women.

Link to Disability Leadership Institute’s closed Facebook page. This page is reported to be well moderated and an opportunity to exchange, learn, network and mentor.



Participants in discussion at the Hobart Workshop

### 2.5 Choices and Decisions

“You are the only one who knows you! You know!”

“I have the right to make choices”

#### 2.5.1 Definitions

Informed decision making

Supported decision making

Assisted decision making

Assumed capacity

Consent

Autonomy

Supported decision making

Guardianship and the role of guardianship boards. [The purpose of guardianship is to expand your exercise of rights, not lessen it].

#### 2.5.2 Information

To make informed decisions, you need to have information on the scope of what is available

Gendered impact on access to choice and decision making for women

Decision making can take time

The importance of being believed and listened to

An advocate is someone who can help you speak up and speak for you

Who can step in and when.

#### 2.5.3 Everyday life choices and decisions

Work you want, where you want

Where you live and who you live with

What activities you do and get involved in

Learn things, and attend courses

Choose what to cook, eat and when to eat

The right to self-expression – access to communication devices and communication support

The right to respect – to be treated like an adult and to be asked – not talked over

The right to direct communication with health professionals

The right to privacy – for a person with a psycho-social disability to refuse unwanted risk assessment that you feel corrodes your right to privacy

The right to assert your right to say no and state your needs

The right to choose your support

The right to go to your country, to eat bush tucker. Access to traditional ways. Access to going bush.

Residential settings – the right to have a friend over to visit, to stay the night. The right to have a pet.

“I made a decision to go to Hobart [for this workshop]”

“It’s your choice to make, not anyone to make it”

#### 2.5.3 Stories, videos, talking books

Share your story with other women and girls and show: “*You are not alone” “I look after myself”*

Show women with disability leading ordinary everyday lives in pictures. Washing, cooking, looking after each other

“*We don't leave people with disabilities on their own. Caring and sharing. Family help me around the house. That's our culture*” Marjorie (Tennant Creek).

#### 2.5.4 Tips and tools

Using your inner power: Taking control of your lifestyle choices. What to do about people who stop you from doing things? How to respond when people take over? How to be assertive, speak up and stand up for your rights? What language you can use when you feel shut down, infantilised. Some easy ways to say “No, *this* is what I need”.

Managing your support worker: How to get action when there is non-action by support workers e.g. getting something fixed. How to dismiss/change a support worker when it’s not a good match.

Safe environment: How to be safe in your home – appliances work, things in good order, the environment is physically safe.

#### 2.5.5 Links

UN framework

Legal framework for consent and guardianship state by state

Advocacy organisations

Peak bodies

Public Advocate state by state

Guardianship frameworks, boards, tribunals state by state

Multi-Cap – whole of life disability support

Links to learn who can help you make decisions, and how to make them

Links to independent living centre

Create a noticeboard for individuals and advocates to learn about funding sources

Environmental design and adaptations for people with disabilities.

#### 2.5.6 Resources

A user guide for guardians, supporters and various stakeholders in how to support decision making in order to enable exercising of rights

Decision making tools

How to support another person to make a decision

Where to find out about local activities? Join a choir, a theatre group, an art group, participate in community events etc.

### 2.6 Sexuality, women’s health and family

“You should always get consent and don't let them force you to do it or anything you don't want to do. Use a condom.”

#### 2.6.1 Definitions

Sexuality as distinct from sex. What is sexuality?

Sexual orientation, gender diversity, LGBTQIA+ etc. also in Easy English

Consent: Enthusiastic, informed, participatory consent. “She didn't say no” is not consent. Often you might freeze of be unable to verbalise on the spot.

Sexual abuse

Information on basic terms in sex education in Easy English. Safe sex. Terms and body parts

Contraception

Sterilisation

Spectrum of normal

Terminology – e.g. Sistergirl is an indigenous term for transgender woman.

#### 2.6.2 Myths and facts

Myth: *“Women with disability are inadequate mothers because of their disability”*. Show a video of a mother’s story. The agony of having your baby removed.

Myth: “*Don't breastfeed, your boobs will go saggy”.*

#### 2.6.3 Provide accurate, relevant and accessible information on:

Consent and health providers: Your rights when seeing a heath/medical practitioner. The right to say no to getting undressed, to request and get personal space. Asking for more time in an appointment. Dealing with having to pay double because you needed more time.

Sex: You have the right to have and to enjoy sex, and not be hurt. The right to consent to sex – in Plain English. Positive messages about sex and sexuality. Accessibility barriers in being sexually active – advice and tips. Show safe sex visually with objects. No means No!

Family: The right to access fertility treatments (e.g. IVF). The right to have a baby and to be informed about what is involved in taking care of a baby, such as hygiene, food, bottles. Information about how to be healthy during pregnancy– links to good videos. Information about miscarriage and grief. Advice and information about family health state by state.

Guardianship: Guardianship and marriage.

*“I’m 50 and my parents have guardianship over me and won’t allow me to get married”*

Sexuality, sex, sexual health and education: Your right to accurate, accessible, relevant, situational sex education that is easy to understand. The right to information about women’s health. The right to have sex with a consenting person/partner. The right to education to understand consent and safe sex. The right to wear what you like as a way to express your unique self.

* + Your rights regarding your sexual health and freedoms and how to protect them
  + Consent – resources and awareness
  + Setting boundaries
  + Safe sex, sexually transmitted infections (STIs)
  + What are your options if you are pregnant?
  + Pregnancy stigma and how to prevent it
  + State by state options regarding abortion
  + How and where to get help
  + Contraception, sterilisation
  + Assistive technology – to explore your own sexuality.

Health: “*If you know/feel there something wrong, keep at it, don't take NO – keep doing your research, seek second and third opinions. It can be life threatening*”. Jo in Brisbane.

* + Importance of breast screening and pap smears
  + Range of complementary heath care options available
  + Consent around youth and sterilisation
  + Menopause
  + Include a list of some of the common challenges, obstacles, experiences, structural barriers and inequalities women with disability have encountered when trying to navigate/access healthcare (e.g. substitute decision making, ‘carers’ talking on behalf to health practitioners, medical procedures or conditions not being properly explained to women, reproductive health, mother with disability).

#### 2.6.4 Links

Getting information/help with STIs

Getting information/help with unplanned pregnancy

Where to get the morning after pill

Consentability Facebook page

Contraception and consent (WWDA report)

Gender identity (WWDA report)

Intimately Disabled Facebook page

Links to websites on how to enjoy sex and intimacy and not be hurt

State by state links to health services

Accessing genetic counselling regarding having children

Queer community websites

Living safe and sexual lives – for people with intellectual disability

Parenting podcast on ABC Life Matters: “We’ve got this”

Breast screening and pap smears

Links to health professional recommended allies

iTalk studios for animated education and stories

Kulila! Indigenous language app (by NPY)

Mental health – who to talk to

Where to get help on the journey to passing – end of life conversations

Link to home heating subsidies for women with thermo regulatory disfunction.

#### 2.6.5 Stories and quotes – graffiti wall

Experience of using sex to feel self-worth

Ageism about being sexually active

Expectations of health practitioners.

#### 2.6.6 Forums

Forum for parents with an intellectual disability. Parenting boys and adolescents e.g. refer to closed Facebook group for parents of children with ADHD

Parenting support links state-by-state (e.g. Anglicare’s Alina Services in Brisbane).

#### 2.6.7 Tips and tools, checklists

How to get healthy, how to stay healthy. Exercise, aqua aerobics, lifting bars, dancing, go for a walk along the river, join a Zumba class, Tai Chi …

Respect, safety and consent in relationships

Tips for making appointments with health providers: How to be confident in a phone call to ask, open up and get what you need. Getting the best out of a service when you call them – know what the service is for, so you are going to the right place for what you want.

Dispel tension and get your needs understood. “I’m your problem patient today” in a friendly tone, can sometimes allow the person on the phone to want to help you!

Check the physical access in detail. How close is the clinic/hospital to public transport?

#### 2.6.8 Content

##### Messages – maybe in stories

Everyone is a sexual being and has a right to sex and to explore sexuality – that's inclusion

Love yourself, think for yourself

Do things you love

What is a good relationship? Loving each other, holding hands, sharing interests and ideas

Eat healthy, be heathy, care for your body

Help your children be healthy

Talk to someone you trust

Build strong networks – talk about taboo topics.

### 2.7 Safety from violence

#### 2.7.1 Summary

Some core messages:   
Family violence affects everyone in the family. Violence is nuanced. It may be invisible, but its real!! Find someone you trust and respect. Talk about sexual violence. Important that every person has a network and feels they are valued and they belong.

#### 2.7.2 Definitions

Provide pictures, words, moving pictures to help women to understand what violence is, in all its forms.

Provide UN definitions

Define all forms of violence and abuse [mistreatment and neglect]:

* + What is violence? Types of violence and what they look like. Includes hitting, being pushed off the bus…. What is safety from violence?
  + What is sexual assault versus sexual abuse, and what it is not
  + What is abuse? Include financial abuse, social abuse and how they may specifically impact people with a disability.

Institutionalised and systemic violence. Exclusionary policies, restrictive practices

Intersectionality and violence against ATSI women and LGBTQI+ women with disability.

“*Financial abuse is when people make you buy things for them*”

Mandatory reporting.

#### 2.7.2 Statistics

Statistics on violence, gender and disabilities, women with disability

Prevalence, who it happens to, ages, who perpetrates.

#### 2.7.3 Information

What is the role of police?

Your rights regarding records

The fear of not being believed

Being safe in all environments – work, home, leisure

Knowledge is power – get yourself informed.

#### 2.7.4 What to do and getting help

“Go to someone you trust and get help. Don’t stay with the person. Go to the cops and get help. Stay: STOP! I don't like it!”

This section should provide a clear pathway if this happens to you and advice on what to do. You should not have to wade through information.

You can call 1800 RESPECT, Download Sunny App, go to someone you trust

Discreet instructions for incognito browsing and quick ‘get out’ link

What actions to take if you are in an unsafe situation and need immediate and urgent help

Who you can talk to for support

Mandatory reporting rules.

#### 2.7.5 Tips, tools and checklists

Provide different scenarios and checklists for teenagers, parents and carers

“I want to say to my carer: Stop telling me what to do!”

Downloadable fact sheets

Guidelines to support women with disability to become free from violence – messages of hope (WWD Victoria)

Make a poster for your wall at home with things that help you feel safe and strong

Make reminders for yourself and put them around you

Staying calm – tips

Try journaling. Sometimes you can’t talk about it, but you can write or draw it and show people that way

Join a support group. Check it out first to be sure it’s right for you and that you like the people. Does it feel safe and good to be there?

Things you can do to be aware of your safety

Self-respect checklist – things you can say/do about unwanted touch of any kind. Say No! if you don't like it or want it

Taxis: once you find a good one, stick with it! Book direct, not through central switchboard

The importance of valuing your personal space – be aware of it in different situations and how to hold on to it. Trust your instincts.

#### 2.7.6 Reclaiming power – information, stories

Protective behaviours to strengthen women’s awareness to take back power

Bullying, and verbal abuse – learn to get in touch with your personal power to push back/get help

How to stay in control through the process of getting help. For instance: You can disclose to police and request it not be taken further at this stage

Finding and using your personal power to push back and get help

Saying no to unwanted touch.

#### 2.7.7 Stories, videos, case studies

Confidentiality and support for those sharing stories. For instance, someone else can tell your story on your behalf so you are not identified

Content alert: Give a warning at what point in the recording or video there is content some may find disturbing

It’s OK to say NO! Stop! and be assertive

Messages of hope. We can stop it from happening

Stories of people who have gone through it and come out strong

What causes violence? e.g. jealousy

What does a non-violent relationship look like? What does a good relationship look like?

Stories of success.

#### 2.7.8 Links

Resources for all ages. Online, national, state and local services

Response services

Prevention peak bodies

1800 RESPECT

SUNNY App

Domestic Violence Resource Centre Victoria (DVRCV)

Salvation Army – DV and financial support (Brisbane)

DV services – (Leona, Brisbane)

Self-care cards for people with low energy – full of ideas that don't require a lot of physical activity or movement

Positive action cards

Counselling services

Where to find professionals to talk to

Clutterbugs WA – will take your clutter/household goods to a women’s refuge

CrisCare – beds in refuges

Self-defence training

Links to good information

SECCA – including people with intellectual disability

### 2.8 Navigating the National Disability Insurance Scheme (NDIS)

“The NDIS will help [the problem of overpowering support workers] because it’s about self-managing” Elizabeth, Perth.

The NDIS came up repeatedly within so many of the discussions, therefore it has its own section here.

Right to apply for support though the NDIS to pursue your goals. Having access to ordinary everyday life opportunities that are available to people without a disability. Such as: work I love, being paid, a home I am happy in, pursuing my interests and goals, access to a good therapist

How to apply for the NDIS and develop a Plan. How to get your NDIS person to help you get what you need and help you to be able to pursue your goals

How to get NDIS funding for sexual rights

Need support to navigate the NDIS and all the admin involved. Things you can say to get more information in order to make better decisions.

[One participant wrote an entire submission within the workshop about the NDIS and it is presented here:]

“Choice and decisions – NDIS marketed giving disabled people more choices but there are many instances (often hard to detect) where choice is being manufactured.

Private NDIS providers to keep funding going into their own pockets will often limit clients to the services they provide, even when a service offered outside their business is more appropriate. Clients have to ask specifically for their NDIS worker to research services of other providers.

Private NDIS providers can be very insistent and pushy when it comes to clients signing new service agreements. Simple English, braille or audio versions of service agreements are often not provided.

A guide, or set of questions, criteria for people with disabilities to draw on to better assist them or a person close to them to choose an NDIS support worker that will best meet the individual needs. I.e. how many years of experience have you had working with people with disabilities (or more specifically what is your experience/knowledge of my disability, what have you studied? How long have you been in the disability sector?

Often a private NDIS provider will assign a random support worker to a client without much consideration as to whether they’re an adequate fit. Often people with disability have been waiting for services or for someone to contact them from the NDIS for such a long time that the moment a support worker is assigned to them they immediately ‘just go with it’ because of the urgency of needing to get support.

It needs to be more explicitly stated to people with disability that the worker assigned to them is not the one they have to work with.

I have mentioned the phrase ‘private NDIS provider’ a few times now and it occurred to me that perhaps some disabled people who are contacted by a ‘private NDIS provider’ might think they’re being contacted by the NDIA directly – and as a consequence of not comprehending this distinction they aren’t able to recognise how ultimately these providers are businesses (many businesses started under the NDIS have been started by people with no previous background in the disability sector”.

The Tennant Creek Barkly region was a first site for the NDIS. Equipment needs were the first priority. But wheel chairs were often not accessible in sand on country. We learned that there is no word for disability in the local language/s. The term ‘mad’ is used for both mental health and intellectual disabilities. Many don't think of themselves as living with a disability.

### 2.9 Website design and usability

##### Needs, suggestions and requests

“*Easy English and Easy Read. Should have it on everything so people with disabilities can read it and understand it and read it properly. Loud and big!”*

Easy read and plain English. No big words

Conversion button for easy read

Large font, well-spaced out

Not too much stuff!

Colour contrast between background and print e.g.: white and pink don't work!

Make it interesting and colourful

Use boxes, bold, clear, colour

Videos. Moving stories. Short. Clear message. Music. Acting. Colour

Stories in language. Use subtitles

Visual information. Buttons with icons.

*“We want viewers to see diversity, see country, see social life. Indigenous women want to see our own age, our own situation in the videos*” Tennant Creek.

Suggestion:

Summary section at the top (Easy English)

Plain English version under that with more detail

Links for more in-depth information for those who seek that

Screen reader compatible.

From a woman with autism:

Have a uniform pattern throughout so you don't get anxiety about what’s coming next

A pattern to the colours

Make it simple, “if it’s too overwhelming, I shut down”.

Have a section: What is this website and how do I use it?

Have audio format for those that don't read.

##### Layout

Under each category, after a ‘look out for these things’ sort of list, perhaps a recommendation of further resources/supports? Such as YouTube videos about the issue.

##### Networking

Create a forum to talk with other women and girls with disabilities

Facebook link to website moderated by WWDA for exchange about topics on the Virtual Centre.

##### Commonly used devices

Phone and tablet. Many without traditional computer literacy have workaround shortcuts to use computer – keyboard shortcuts by counting keys – shortcuts to various websites.

##### Access to internet in Tennant Creek

The local park will soon have a space to sit with free Wi-Fi. Michelle, hub manager, hopes to acquire computers for the Paterson St Hub.



Workshop facilitator, Julia, listens to participants feedback at the Perth Workshop

### 2.10 Workshop Evaluation

In most workshops an evaluation was conducted. In one setting the evaluation was not logistically possible because of time constraints, comings and goings (Tennant Creek, Northern Territory). The evaluation happened in two parts:

Participants were asked to place dots on a large poster with a line from left to right that represented a continuum - starting at ‘I was not able to contribute as much as I would like. I have more to say’ to ‘I got say what I wanted’. A green dot indicated ‘I got to say what I wanted. Easy to have my voice heard’. A red dot indicated ‘I had more to say that didn't come out’. For those who could not perform the task physically, a local co-ordinator assisted. In five of the six workshops one red dot was used, the rest were green.

In addition, team members and local co-ordinators (not the facilitators) discreetly went to each person privately to ask what they felt had worked well, and what suggestions they may have had for improvement. The responses were as follows:

##### Suggestions for improvement:

More time

More meetings like this

More of these workshops

The last few topics were a little rushed

It was a little long, I got tired

I got a bit distracted with personal things

More pictures to explain what we were talking about

More money on the gift cards

I would like to talk about more subjects

More groups like this.

##### What worked well:

Talking, sharing, doing pictures, drawing, writing

I like the people moving [around] where you search [for information on the posters]

All!

Good to talk about complaints, to make sure we don't have complaints

I loved the leadership from people, and people sharing their stories. Feel privileged to be learning here

Get to see changes before they will happen, to make a difference for those who don't have a voice

Life experiences

Very interactive. Excited to see what comes out!

Grateful for the experience

Liked the sense of safety in discussion around sex and violence

Deep work

Time flies!

Feeling comfortable enough to speak about my/our experience

Atmosphere was calm and laid back

Well prepared and executed. Great work.



Facilitator, Julia, hands a participant a card at the Tennant Creek Workshop

## Part 3: Reflections, outcomes and suggestions

### Women want the website

The fact of doing these workshops in itself was important. Participants felt engaged. They appreciated being so involved, having their voices heard and influencing the outcome early on. Women across the board overwhelmingly expressed that the Virtual Centre is needed.

### Stories, stories, stories

Across the board women requested stories and in particular, video stories. Women expressed over and over the need for role models that mirror their own lives.

Participants wanted to see and hear women with disability of all ages speak of their experience, how to live with pride, how to navigate the reality of everyday lives, to dream, and to show what is possible.

In response to the significance placed on this need, the project manager has now hired a professional film maker.

### Practical guides for specific situations

It is not so easy to do things differently if you have never seen it done. Across the board, women indicated it would be a great help to have practical checklists and resources for specific situations, such as going for a job interview, stopping your carer from taking over and limiting you, talking about your disability at work, and many others.

### Navigating services

The hunger for information about local services and clear and easy referral pathways is clearly a need for women with disability. In every location, the facilitators needed to explain that that the Virtual Centre is not a service directory. The Virtual Centre can, however, provide information that will help women to navigate services and referral pathways and help them to work out who to talk to that can assist them with this locally.

### Self-advocacy tool

It became evident through the workshop experience that the Virtual Centre is providing a self-advocacy tool. It is equipping and resourcing women with information, role models, examples and the confidence to be able to advocate for themselves, as well as for and with other women with disability. Many women commented, “*I know my rights, but how do I speak up about them? Particularly in situations where I feel less powerful – such as with my support worker”*.

Some of the women who attended the workshops are already well aware of their rights, and many are already in advocacy roles. They were able to speak about what women need more broadly, and advocate for women in similar situations who many not know their rights and the potential impact the Virtual Centre could have in their lives.

### Taboo topics

Many participants expressed appreciation that the Virtual Centre is providing information on more sensitive and often taboo topics in some settings –such as violence, gender relations and sex. It was stated multiple times that an online resource is a great way for women to research these areas themselves and learn about things they may not feel comfortable asking others about for information.

### Connectivity and interaction

It was recognised that the online environment opens up all kinds of opportunities for women with disability to connect with other women all around the country and the world. Women wanted the website to provide some form of connectivity to be able to interact with each other on topics that matter to them. For example: a face book page, a virtual forum, on-line mentoring and networking opportunities. This request should be included in the overall Victual Centre strategy.

### The Alice Springs experience

In Alice Springs it was not possible to organise a workshop with women with disability for local logistical reasons. The team did however, meet with various agencies providing services to women with disabilities to seek their direct input and feedback on the Virtual Centre plan. These meetings have the potential to strengthen the WWDA network. These organisations included:

The Purple House – an indigenous led and managed project for mobile dialysis on country. The team visited Purple House’s clinic, community centre and social enterprise hub in the centre of town.

National Disability Services (NDS), Alice Springs – the team met with the leadership team of NDS providing accommodation, employment and services to people with disability in the area.

### The Tennant Creek experience

In Tennant creek the team witnessed the strengths of community networks, of resilient women with inner power, and the power of influence. They met women with the savvy and humour to use the material resources they have access to, for practical solutions.

The workshop team also experienced first-hand the impact of economic and social barriers of systemic racism that are often invisible to many Australians. It was clear that material resources for simple solutions are scarce. Tennant Creek was a trial site for the NDIS, but women reported how difficult it has been to source adaptable equipment suited to local terrain. Money for service providers was allocated, but in the region there is a lack of therapists and suitable service providers to respond to the demand and need.

Like many ‘helpers’, the team flew in and out. It is recognized that for future projects there should be more effort made to make time to just be with people, and to get to know their lives and situation on their terms.

## A participant makes notes at the Brisbane workshop

A participant makes notes at the Brisbane workshop

## Conclusion: Unity in diversity

A guiding principle for the Virtual Centre is that women with disability see themselves reflected in it. A website that equips and resources women with disability, empowers them to realise their human rights and reflects and celebrates their diverse stories is an important and achievable goal.

The Virtual Centre is an opportunity to contribute to the vision of an equitable and inclusive Australia, and to reflect the diversity of communication styles, language, faces and perspectives of women with disability across Australia.

## Acknowledgement and thank you

The WWDA facilitation team owes our gratitude to all the women who participated in the workshops. We met incredible, smart, wise, humble, tenacious women of all ages, cultures and backgrounds. We learned so much from each and every one of you.

We thank our partner organsiations in every location who welcomed us, hosted the event, and provided invaluable assistance and local knowledge. Thank you for making us feel at home with you!



The co-facilitator Monique at the Hobart workshop

## Appendix 1: Workshop attendance summary

#### Table . Breakdown of the workshop sessions and attendee details

|  |  |  |
| --- | --- | --- |
| **Location** | **Participants** | **Age range** |
| Hobart | 7 | 18 – 60 |
| Canberra | 6 | 20 – 60 |
| Melbourne | 10 | 21 – 60 |
| Brisbane | 8 | 18 – 55 |
| Tennant Creek | 10 | 22 – 55 |
| Perth | 4 | 40 – 65 |

## Appendix 2: Consent form (Plain English)



## Workshop Information Sheet

Thank you for coming to the Women With Disabilities Australia (WWDA) Virtual Centre Co-Design Workshop in …… (place) at ….. (venue) On ….. (date, time)

WWDA is making a Virtual Centre website for women and girls with disability around Australia.

We will use the feedback and ideas that you give us in this workshop to help to make sure that the Virtual Centre website has information that is useful and easy for you to use.

We will be taking notes at the workshop. We will also take photos and will record what you say using an audio recording device. The notes and audio recording will only be used by us to help us remember what you say and to make sure we record all of your feedback and ideas.

We may also take photos and short videos at the workshop. These may be used for promotional purposes – including things like social media and WWDA newsletters. You can tell us if you do not want us to use your photo or video.

We will write a report about this workshop. This report will summarise all of the feedback and ideas that you and other workshop participants give us about the Virtual Centre. This will be used by WWDA and the website design organisation (Handbuilt Creative) to help us create the Virtual Centre. We will provide this report to the government department that has given us funds to create the Virtual Centre.

The report will not include your personal name, except in the list of participants (if you agree). The report might have ideas for a particular state or location. The report might include photos taken at the workshops.

The report will be kept on a password-protected computer at WWDA for at least 5 years. We will send you a copy of the report so you can read it.

The report may also be made public on the WWDA website.

WWDA may use parts of the report for things in the future like:

Statements to the media and the public

WWDA Newsletters.

We really appreciate your participation and the workshop and will provide you with a $50 gift voucher to thank you for your time at the end.

If you have any questions you contact the Project Manager, Naomi Thomson at [pm@wwda.org.au](mailto:pm@wwda.org.au)**.**

Thank you for participating.

The WWDA project team

Naomi Thomson (Project Manager)

Julia Wolfson (Workshop Facilitator)

Heidi La Paglia (Senior Content Officer)

Dana Endelmanis (Communications and Media)

# Women with Disabilities Australia Virtual Centre for Women and Girls with Disability. (logo)

## Workshop Consent Form

I agree to take part in the Virtual Centre workshop.

I agree to allow WWDA to audio record the workshop.

I agree that WWDA can include my comments and suggestions in the workshop report.

I understand my name will not be next to my comments in the workshop report.

I agree that WWDA can write my name in the list of participants in the workshop report.

I agree that WWDA can use the information in the report for public statements in the future.

I agree that WWDA can take photos of me at the workshop

I agree that WWDA can take videos of me at the workshop

I agree that WWDA can use photos and video footage of me for promotional purposes. I know I can change my mind about this at any time by contacting WWDA.

I would like to get information and continue to be involved in the development of the Virtual Centre after the workshop.

I understand that WWDA will not give my personal information to anyone else.

**Name**

**Address**

**Telephone**

**Email**

**Contact person (name and telephone) (optional)**

**Signature Date**

## Appendix 3: Consent form (Easy English)

## Women with Disabilities Australia (WWDA)

# 

### Consent Form

****



**Need a translator**

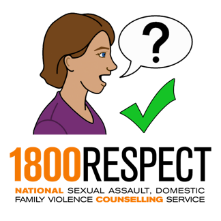
Call 131 450

Ask for 1800 737 732



**In danger now**

Call 000



To talk to some one about **violence**.

Call 1800 RESPECT on 1800 737 732.

or

Chat online at [www.1800respect.org.au](http://www.1800respect.org.au)

****

**WWDA** means

**Women with Disabilities Australia.**



WWDA is developing a Virtual Centre **website for women and girls with disability**



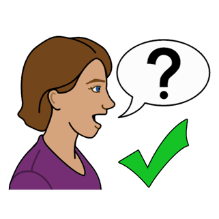
The Virtual Centre website is being funded by the NDIS



We are doing **this workshop** to find out what **you want** on the website



We will **tell you about the website** in the workshop



We will **ask you questions** about the website in the workshop



We will take notes in the workshop about what you say



We will **audio record** the workshop



* We will use the Information you give us to write reports for people. Like

****

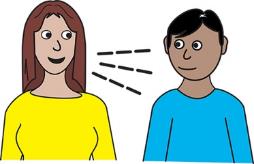
WWDA

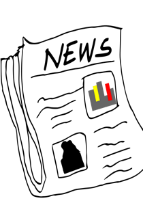


The people who make the website



The NDIS

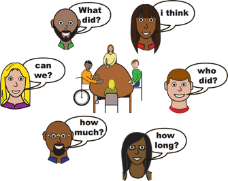
We may also use the things you say in other things like



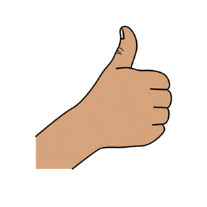
* News articles

****

* Reports about WWDA.



* Other workshops

****

We would like to know if you are happy for us to use the things you tell us in these ways

We would also like to know if we can take your



* **Photo**
* 

**You can tell us on this form**



1. Read the form



1. Tick the boxes next to the things you would like your photo to be used for



1. Sign this form

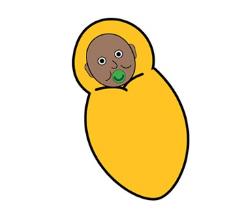


**You can ask someone to help you fill out this form.**

**About you**



**My name is**



**I was born on**

Write the day / month / year

****

**I live at**



**My phone number is**

****

**My email is**

****I am happy to take part in the

**workshop about the**

**Virtual Centre website**



I am happy for WWDA to use

the things I tell them to **make**

**sure the website is easy to use**



I am happy for WWDA to use

the things I tell them in **reports**



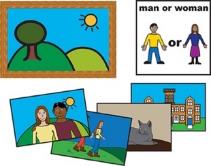
I am happy for WWDA to **record**

**the Workshop by audio**

****

I am happy for WWDA to

take **photos** of me at the workshop



**WWDA can use my photo**

****

* On the WWDA website
* wwda.org.au

****

* On the Virtual Centre

**WWDA can also use my photo**

On Social Media sites like



* Facebook



* Twitter



* Instagram



* In an email newsletter

A close up of a logo

Description automatically generated

**My signature:**

**Thank you for your time!**



We will give you a **$50 Coles Voucher** at the end of the workshop as a **payment.**



If you have **any questions** about the workshop you can contact our Project Manager Naomi:

* email [**pm@wwda.org.au**](mailto:pm@wwda.org.au)
* call **0448 417 875**

**Acknowledgements**

**Authorship**

Women with Disabilities Australia (WWDA) wrote the content for this booklet.

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## Appendix 4: Workshop Agenda – example

### WWDA Virtual Centre Co-design Workshop

**Location: Perth, WA**

**Date: Friday 16 July 2019**

**Time: 11am-2:30pm**

|  |  |  |
| --- | --- | --- |
| **Time** | **Topic** | **Speaker** |
| **11:00am** | **Welcome and Introductions**  **Consent and wellbeing**  **Agenda – what will happen today?** | **Local facilitator and**  **Julia Wolfson** |
| **11:20am** | **Introduction to the Virtual Centre**  **What is it? Why are we meeting today?** | **Naomi Thomson** |
| **11:30am** | **Questions and Discussion**  **What do you want to see on the Virtual Centre?** | **All** |
| **12:30pm** | **Break** |  |
| **1:00pm** | **Questions and Discussion**  **What do you want to see on the Virtual Centre?** | **All** |
| **1.45pm** | **Break** |  |
| **2:00pm** | **The Virtual Centre website layout**  **Trying it out on computers** | **All** |
| **2:15pm** | **What did you think of the workshop?**  **Summary of what we talked about** | **Local Facilitator & Julia** |
| **2:25pm** | **Thank you for coming and what we will do next** | **Julia & Naomi** |
| **2:30pm** | **Finish** |  |

## Appendix 5: Wireframe Prototype Activity Response Form

### Wireframe Prototype Activity Response Form

Form to be completed by interviewer

**Location:**

**Name:**

**Support requirements:**

**Requires a support person to view websites? (Yes/No)**

**Preferred device for using websites? (eg ipad, smartphone, laptop):**

**Uses assistive technology to view websites? (if yes, please detail):**

**Remind them – this is not a test!**

We will show you 3 different possible homepage designs (which are very simple, barebone ‘shells’ of website layout). They don’t contain images or correct content, and they are not functioning websites. We will give you a simple task to do for each design. Watching what you do will give us very helpful information and help us to choose which layout we use.

**Please talk out loud and tell us what you are thinking and doing as you do it.**

**Wireframe 1** – <https://wwdavc-beta01.webflow.io/>

1. Ask: Can you find information about Health?

**Comments/reactions**:

1. Ask: Did you know you could scroll?

**Wireframe 2** – <https://wwdavc-beta02.webflow.io/>

1. Ask: Can you find information about being Safe?

**Comments/reactions**:

1. Ask: Did you see the ‘Tips’? Were they helpful?

**Wireframe 3** – <https://wwdavc-beta03.webflow.io/>

1. Ask: Can you find information about Events?

**Comments/reactions**:

1. Ask: Did you notice the video? Do you like it?

**Any other comments/ notes**

1. NSW Council of Social Service (2017). Principles of Co-design. [online] NSW: NCOSS, pp.1-3. Available at:1<https://www.ncoss.org.au/sites/default/files/public/resources/Codesign%20principles.pdf> [Accessed 5 Aug. 2019]. [↑](#footnote-ref-1)
2. [Vichealth.vic.gov.au](http://Vichealth.vic.gov.au) (2019). How co-design delivers agency, advocacy and real-world impact. [online] Available at: <https://www.vichealth.vic.gov.au/letter/articles/vh-letter-45-co-design> [Accessed 13 Aug. 2019]. [↑](#footnote-ref-2)
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5. See Julia Wolfson’s [website Turning Forward](http://www.turningforward.org/). Julia is also a faculty member of the global [Deep Democracy Institute](http://www.deepdemocracyintensive.com/). [↑](#footnote-ref-5)