



Us Mob and HIV



www.usmobandhiv.org.au

5TH
EDITION
2024

Us Mob and HIV

Aboriginal and Torres Strait Islander people should be aware that this booklet may contain names of people who have passed away.

Us Mob and HIV (5th edition) is for Aboriginal and Torres Strait Islander people. It contains information that deals with both men's and women's business.

It was produced by Health Equity Matters in partnership with the Anwernekenhe National HIV Alliance (ANA). The first two editions of this booklet were known as 'HIV/AIDS and Us Mob'.

We recognise this resource may not represent all aspects of the diversity within Aboriginal and Torres Strait Islander communities and cultures. Text from this resource can be changed or adapted to better suit the needs of Indigenous communities.

We acknowledge and pay respects to Aboriginal and Torres Strait Islander people as the traditional custodians of the lands on which we work.

Artwork

Front cover: Arone Raymond Meeks

Back cover: Toby Cedar

Copyright remains with the artists.

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Us Mob and HIV

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**Find out the facts.
HIV in our community affects us all.**

Special thanks to the Aboriginal and Torres Strait Islander people who have given their yarns in the booklet. (Names may have been changed to protect their privacy.)



Cover Art

Title: **Dialogue**

Artist: **Arone Raymond Meeks**



This work represents communication between people on a verbal level and how they feel and respond to each other.

The figures are linked by their hands and crosshatching. Crosshatching is a symbol for fertility and the earth. The background depicts elements of the tropics, such as the reef, rainforest, and coral spawning. The kidney shapes within the figures represent the emptiness we may feel when responding to confronting situations.

Artist Acknowledgement

Arone Meeks was a Kuku Midigi man and an accomplished artist and leading advocate for Aboriginal and Torres Islanders with HIV.

Arone Meeks provided a unique contribution to the HIV sector and pioneered a new way of storytelling. Through art, he shared his story and that of his community by interconnecting Aboriginal culture, HIV and health promotion.

His art was ground-breaking, giving a voice to those previously voiceless. Arone Meeks' lasting legacy is a collection of artwork which will continue to have a profound impact on the way we understand the history of the HIV response in Australia.

Back Cover Art

Title: **Floral Nam Dari**

Artist: **Toby Cedar**



This art piece has the colours of the Torres Strait Islander flag throughout it – with green representing land, black representing our people, blue representing the sea, and white representing our Dari (Headdress). The coconut leaves, hibiscus and frangipani signify the tropical lifestyle as they are seen throughout the Torres Strait. The Turtle plays a special part in the island lifestyle as both a food source and many families' totem. Inlaid in the turtles' shells are the Dari which is the main symbol on our flag to represent all our Torres Strait Islands.

Us Mob
and HIV



Visit Us Mob and HIV Online

www.usmobandhiv.org.au



About HIV

What is HIV?

HIV is a type of germ called a virus.

HIV damages your immune system. This means your body cannot fight germs or other diseases, like a cold or flu.

HIV makes it hard for your body to keep itself well.

Unlike most other germs, there is no medicine to get rid of HIV. Once you get HIV, it stays in your body.

HIV stands for:

Human

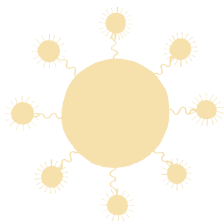
Only people get it.

Immunodeficiency

It damages your immune system which is what keeps your body well.

Virus

It is a virus (germ).





What is AIDS?

HIV is not AIDS. If people have HIV and do not get treatment, their immune system will become so weak that they will get unusual illnesses. When HIV has gone this far, doctors call it AIDS.

AIDS is very serious. In fact, people with AIDS are more likely to get diseases that can kill them.

Taking treatment can stop you getting AIDS.

AIDS stands for:

Acquired	You can only get HIV from someone who already has it.
Immune	It affects your immune system, which fights off germs.
Deficiency	Your immune system stops working properly.
Syndrome	People get sick from it in different ways.





Taz's Yarn

As a Brotherboy I see there is a huge amount of stigma and misunderstanding of the needs of Brotherboys and a need for culturally safe and affirming testing sites and other service centres. It is important that assumptions are not made about our different sexualities and the services that we require, and that clinicians and health workers are appropriately trained to better meet the needs of our communities. As Brotherboys it is important not only that we remain educated about safe practices in all aspects whether it is sex, taking substances or even tattooing or piercing, but also to reach out to services where we feel safe for HIV testing and our other needs so that we have better health outcomes for the community.

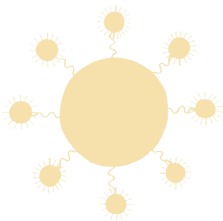


What happens when HIV gets into your body?

There is often no sign (symptoms) that you have HIV, and many people with HIV feel well for years.

But HIV slowly hurts the immune system so it cannot protect you from germs. You can also pass on HIV to others more easily during this time because there is a lot of HIV in your body.

If you do not take treatments, after a while you will begin to get very sick. You may also get sick from illnesses your body would normally fight off. Treatments can help you stay strong and stop HIV making you sick. Taking HIV treatments properly also means you can reduce the amount of HIV in your body so you cannot pass it on to other people through sex.



How do you get HIV?

Anyone can get HIV. HIV can be passed on through contact with:

- blood
- vaginal fluids
- semen (cum)
- rectal (anal) fluids
- pre-cum
- breast milk

HIV can be passed on from:

- Vaginal and anal sex without condoms
- Sharing needles and other injecting equipment
- Carrying parent to baby, through pregnancy, birth, and breastfeeding.

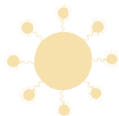
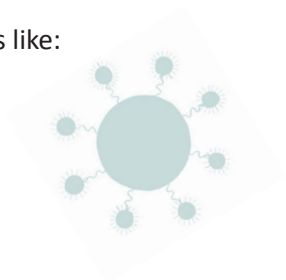
It is not likely but occasionally it can also get passed on by:

- Sharing razors and other cutting, piercing or tattooing equipment
- Getting body fluids that can pass on HIV in cuts or in your mouth. This includes a person bleeding from accidents or during fights.



You **cannot** get HIV from things like:

- saliva
- kissing and hugging
- sharing cups or forks
- toilet seats or towels
- mosquitoes
- sweat



How can I make sure I do not get HIV or pass it on?

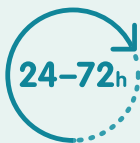


Use condoms and lube for sex.

Using lube can stop condoms from breaking and can feel good.



Take PEP. If you forget to use a condom or it breaks, or if you share injecting equipment, have no shame, and get PEP (Post-Exposure Prophylaxis).



You take PEP medications for 1 month to prevent HIV after a risk. PEP needs to be started within 72 hours after a risk, but it is better if you can start within 24 hours. You can get PEP from Emergency Departments of most public hospitals or at sexual health clinics and some other doctors.

For a fact sheet and video on PEP go to usmobandhiv.org.au

For more information visit www.getpep.info



Take treatment for HIV. If you have HIV, taking treatment keeps you well. It can also reduce the amount of HIV in your body to an undetectable level (so low that tests cannot find it). That means you **cannot** pass HIV on to someone else. This is called 'Treatment as Prevention'. This is also known as 'Undetectable = Untransmittable' or 'U=U'.



Take PrEP. PrEP stands for Pre- Exposure Prophylaxis. It is a medication that HIV-negative people can take to prevent HIV.

Taking PrEP before being exposed to HIV means there is enough medicine in your system to stop HIV if it gets into your body.

PrEP protects you from HIV, but it does not prevent other STIs, or pregnancy.

For a fact sheet and video on PrEP go to www.usmobandhiv.org.au

Aboriginal and Torres Strait Islander people can get PrEP at a reduced cost or for free under the 'Closing the Gap' scheme.

Talk to your doctor.





Use new gear (needle, syringe, and other injecting equipment) every time if you inject drugs. Used syringes contain small amounts of blood. Even a small amount of blood can carry germs or disease.

Make sure you return your used needles and syringes to a Needle and Syringe Program (NSP) or throw them away safely inside something hard, like a plastic bottle with a lid, so other people do not get hurt by them.

To find NSPs in your area:
www.nspdirectory.aivl.org.au



Cover up cuts, and if you have to clean up blood, wear gloves (or use plastic bags if you do not have gloves).



Do NOT share anything that could have had blood on it including knives, razor blades or needles.

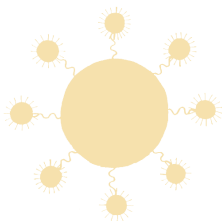


If you are pregnant, talk to your doctor or health worker about what you need to do. People with HIV can have healthy babies without HIV.

HIV can have a big impact on our community and culture.

Sometimes people take more risks when they are away from home.

Whether you are at home or away for some fun, keep yourself safe!



Getting tested



How do I know if I have HIV?

You may have HIV but not know it.



Some people do not get sick at all when they first get HIV.



A lot of people do feel a bit sick when they first get HIV but do not realise it is because of HIV.



It feels a bit like having a bad cold or flu, so that is what people can think they have.



After the first couple of weeks, many people do not have any illness from HIV for months or years.



You may have HIV even if you currently feel well. It is important to get tested before your immune system is too damaged.



The only way to know whether you have HIV is to have an HIV test.

If you are worried because you have been having unprotected sex or sharing needles, talk to your doctor or health worker about having a HIV test. Let them know if you have had a recent illness that felt like a bad cold or flu.

You can find out where your local sexual health clinic and Aboriginal Medical Service is on the Better to Know website at **www.bettertoknow.org.au**



The 715 Health Check is an annual health check for Aboriginal and Torres Strait Islander people.

It is free at Aboriginal Medical Services and bulk billing GP clinics. Yarn to your doctor about including tests for HIV and other STIs in your 715 Health Check.



Jayden's Yarn

As a gay man HIV has always been something that scared me and sometimes cause anxiety when having sex. I was always sure I used a condom which eased my anxiety during sex. Although one time a condom broke and I did not know at the time but thankfully I was able to get PEP (Post-Exposure Prophylaxis). I was lucky that I was able to get PEP within the time after the potential risk to be protected. After my treatment I spoke to my doctor about PrEP (Pre-Exposure Prophylaxis) and we both decided it was a good idea I start taking PrEP so I can now feel less anxiety and more confidence when having sex knowing I am protected.



Your HIV test is your business

Before you get tested you should think carefully about who you want to tell about your plans. It can be good to tell someone you trust so they can give you support. But if you want to keep your business private, think about who you can trust.

Getting tested for HIV is your own private business. Only your doctor, nurse or Aboriginal Health Worker needs to know. They are not allowed to tell anyone about your HIV test results unless you say they can. They are not even allowed to tell anyone whether you have had a HIV test.

If you are worried about seeing someone at your local health centre, you could think about seeing a different clinic, or someone in another town or area.



Get tested for HIV!



And while you're there, ask for tests for other illnesses that can be passed through blood and sex.





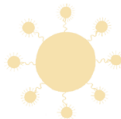
Everything about your HIV test is private:

-  Your doctor will have a yarn with you about your test. Those conversations are private
-  Your results are private business.

Getting the results

Only your doctor, nurse or Indigenous health worker can tell you your HIV test results.

-  If your HIV test result is positive, it means you have HIV.
-  If your test result is negative, it means HIV has not shown up in your blood. That could be because you do not have HIV, but it could mean you are in 'the window period'.





The **window period** means HIV might be in your blood, but it is too early to show up in a test. It usually shows up in a few weeks after a risk for HIV but can take up to 6 to 12 weeks, depending on the type of test. If you have had a test in the window period, you will need to have another test after the window period to double check the results. These results will be accurate if you do not have unprotected sex or share needles in the window period.

Your doctor or health care worker can explain a negative test result and help work out exactly what it means.

Getting tested for other illnesses

STIs Having sex means you could also have other STIs (sexually transmissible infections). Condoms can help protect you from most STIs, but you can still get some STIs even when you use condoms. They can make you really sick and damage your body if you do not get treatment.

If you have an STI, it is easier to get or pass on HIV. You should ask for an STI test at the same time as your HIV test.



There are very high rates of some STIs in many Aboriginal and Torres Strait Islander communities. Getting an STI test is a step to improving the health of our communities.

Hepatitis Sharing needles and syringes can spread HIV but also infections like hepatitis B or hepatitis C. If you have been sharing needles, syringes, or other injecting equipment tell your doctor and get tested for hepatitis B and C at the same time as your HIV test.

Lots of STIs have no symptoms. Have regular sexual health checks. And make sure you use condoms and lube.

You can talk to your doctor if you are worried about anything.








I have HIV – What now?



Having HIV is serious, but it is not the end of the world.

If you do nothing, HIV will lead to illnesses that can kill you. But there are treatments you can take to control HIV so you can be well. You will also need to look after yourself and make sure you have people around to support you.

If you have just found out that you are HIV positive, you will need to think about:

-  Who you want or need to tell
-  Getting support from family and friends
-  Getting help from local health services
-  Choosing an HIV specialist doctor who is right for you
-  Starting HIV treatment (medicines)

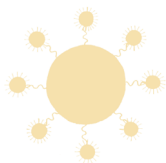


Working out who else might need an HIV test

Your doctor or health care worker might ask you for the names of people you have had sex with or who you have shared needles with. If they think those people might also have HIV, they will contact them and ask them to come in for an HIV test. This is not about blaming you.

It is to make sure everyone who might have HIV gets a test. The doctor can also talk to them about prevention and if needed treatment to stay healthy.

If the health care worker talks to your previous partners, they will **NOT** tell them your name.





Telling sexual partners

It can be really hard to tell someone you have HIV. Your health care worker, counsellor or peer worker can help you work out who you want to tell, what you might want to say, and where and how to say it. There are also services in the back of the booklet that can help.

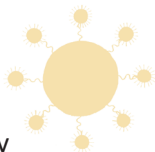
In some parts of Australia, the law says you must tell your partner you have HIV before you have sex or share injecting equipment.

You and your community

You need to look after yourself - your body and spirit.

HIV is an important issue facing Aboriginal and Torres Strait Islander communities. Our communities have high rates of STIs (sexually transmissible infections) and HIV rates have been increasing.

There is no shame having HIV, we all belong and are a part of our community. We all have a big role to play looking after family, community, and culture.



Michael's Yarn



I did not know that I was at “risk of HIV” until after a Positive HIV test, and I did not know about PrEP, (HIV prevention). HIV and treatment get easier over time. I started treatment straight away and was undetectable within a few months. Keeping an undetectable viral load is good for your health because it stops the virus doing damage to your body and Undetectable = Untransmittable (U=U) means you cannot pass HIV on.

At first it was hard to remember to take my medication, so I would set an alarm on my phone. If people asked, what is the alarm for? I would tell them, it is time to take my vitamins! Why do you take vitamins? I would tell them, to stay healthy. So, because I am undetectable, I do not have to worry about passing HIV on to my sexual partners. I see my doctor for regular check-ups and sexual health screening. Once my HIV is under control, I can focus on living my life.



Treatments

Taking treatments (medicines)

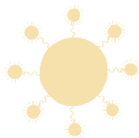


Treatments for HIV are called **'antiretroviral therapy'**.

It cannot get rid of HIV, but it reduces the amount of HIV in your body and keeps your immune system strong. Treatments are easier to take than before. They keep people with HIV healthier, so they live as long as everyone else.

If you have Medicare, you can get HIV treatments from a local chemist at a set cost and is cheaper if you have a concession card. In some states in Australia, these costs are waived, and treatment is free.

If you do not have a Medicare card, there are other options for you to access treatments, so speak to your doctor.



If you are worried about cost, talk to your doctor, Aboriginal Health Worker or the organisations listed in the back of this booklet.

Some people need to take a few different treatments, but many people take just one pill each day. Your doctor will talk to you about which treatment might be best for you.



It is important to take your treatment the way the doctor tells you or it may not work! This means remembering to take it every day and at the same time every day.

When to start treatment

The earlier someone starts treatment after they are diagnosed, the better this will be for their health. It stops HIV from doing more damage. It is often recommended to start treatment straight after being diagnosed.








Once you start treatment, you will need to take them for the rest of your life. Talk to your doctor about when you are ready to start.





There are a few things to think about.

HIV treatment:

-  reduces the amount of HIV in your body. This will make you feel better and prevent you from getting sick.
 -  can reduce the amount of HIV in your body to an undetectable level. That means you cannot pass HIV on to someone else.
 -  does not cause side effects (feeling sick) for most people. Some people still get side effects, but they are usually not so bad and do not last for long.
 -  is something you take for the rest of your life. Once you start treatment, you should keep taking it. If you miss tablets regularly HIV can become 'resistant' to treatment – meaning the HIV treatments will not work properly. You will need to change to another treatment.
 -  Your doctor or health worker should have some suggestions about how to take your treatment every day.
 -  Talk with your doctor. Do not be afraid to ask questions. It is recommended for people with HIV to start treatment as soon as possible after they find out they have HIV to make sure they stay healthy.
- 



HIV positive people who are pregnant should start treatment early.

This will help keep the baby safe from HIV. If you have HIV and are planning to get pregnant, see your doctor. There are a number of things you can do to protect your baby from HIV, including taking treatments. Pregnant people who take HIV treatment regularly almost always have babies that do not have HIV. Your actions will make a big difference to whether or not your baby has HIV.

If you find out you are pregnant, see your doctor as soon as you can!



Michelle's Yarn

As a woman being diagnosed with HIV in the early 90's then having a baby, and not knowing if I passed on the virus was scary, luckily she was ok. Many things have changed since the 90's, such as new medication, taking PrEP which can prevent HIV or using condoms. If you get tested for HIV and the result is that you are now HIV Positive, you can start medication straight away so that you can now live a long life. Taking your HIV medication every day stops you from passing it on. Life does not stop if your test comes back HIV positive, you can still work, you can still play sport, Women can still have babies and we can drink from the same cup. It is important to talk to your doctor if you have other health conditions that might cause problems with your HIV medication. Yarning with Mob who are living with HIV can help you get through the tough times, so you are not alone.



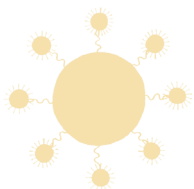
Regular check-ups

If you have HIV, it is important to keep checking your health. You should visit your doctor every few months to check how much HIV is in your blood and whether it is damaging your immune system.

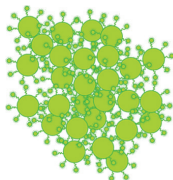
Your doctor will do blood tests to check how much HIV you have in your blood and whether it is damaging your immune system. Everyone is different, so it is important to talk to your doctor about exactly what these tests mean for you.

Checking the virus – The viral load test shows how much HIV is in your blood. If your viral load is high, there is a big chance of getting sick and HIV can pass to someone else quite easily.

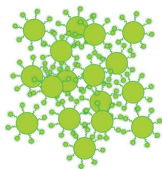
If you go on treatments, your viral load should go way down. Most people will get down to an ‘undetectable’ level. That means you still have HIV but there is so little it does not show up in the tests. This will keep you well and means you cannot pass on HIV.



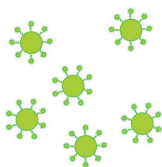
What your viral load means



High viral load – There is a lot of HIV in your blood. This can do a lot of damage, make you sick and makes it easier to pass HIV on to someone else.



Low viral load – HIV is being controlled and does not have much power to make you sick. It is not as easy to pass on HIV.



Undetectable viral load – You still have HIV, but it does not show in tests. This keeps you well and means you **cannot** pass on HIV to your partners.

‘Undetectable = Untransmittable’ or ‘U=U’

Checking your immune system

CD4 cell count tests show how strong your immune system is. CD4 cells fight germs, so HIV tries to get rid of them. If your CD4 count is low, you are more likely to get sick. When you go on treatments, your CD4 count should go way up, which will stop you from getting sick.

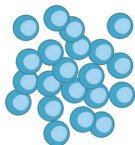
What your CD4 cell count test can tell you



Less than 350 – Your immune system is very weak.



350 to 500 – Your immune system is a little bit weak.



More than 500 – Your immune system is strong.



Kirra's Yarn

I first became more conscious of sexual health when I first started taking PrEP (Pre-Exposure Prophylaxis) before I started my transition as a Sistergirl (Aboriginal Transgender woman) three years ago. Once I became more sexually active, I also learnt more about sexual health and particularly the importance for Mob to also be conscious of sexual health as a high-risk community. STIs and HIV is still a leading concern in rural and remote communities, and we must stand together against the spread of HIV.

Historically PrEP was targeted at gay men or men who have sex with men, however as a Sistergirl on PrEP I know I have taken the right steps to protect myself from HIV regardless of my gender and sexual life. I encourage my Trans, Sistergirl and Brotherboy communities to be active in protecting themselves against HIV and STIs and spreading the awareness of sexual health within their communities. Sexual health does not have to be shame, we all do it and it is better that we are protecting ourselves and protecting others by spreading the word.

Care and support




Having HIV can feel like riding a roller coaster, with lots of 'ups' and 'downs'. The good news is that people with HIV are getting on well in their lives - working, studying, in relationships, having children, looking after their families, and doing all kinds of things they enjoy.

You will probably need support, particularly in the early days. It can be good to have a yarn, but it is important to find the right balance - to get support but make sure you do not have regrets about who you have told. You need to feel safe.

It is up to you who you tell. Care and support can come from different places including family, friends, and health care workers.


Sometimes it is useful to talk to other people with HIV because they may have had similar experiences and they will have some idea what you are going through.



You might also want to talk to a counsellor. Remember, counsellors are not just for people who are desperate. They can help you work out everyday stuff, help you sort out your feelings, and help you make plans to improve things.

There are services in the back of the booklet that can tell you about support services near you.

You are not alone!



There are thousands of people in Australia living with HIV, and many, many more who care about them.

If you have HIV, it is important to keep checking your health. You should visit your doctor every few months to check how much HIV is in your blood and whether it is damaging your immune system.

You will also need to keep checking your health in other ways, like looking after your teeth, eyes, heart, bones, and liver. Your doctor or health worker will help you check these things and anything else that is important.



What can I do if someone I know has HIV?



It is important not to listen to rumours or gossip.

Be careful what you say and do and think how you would like people to be if you had HIV.

People with HIV need to make their own decisions and live their own lives but there are some things you can do to help:



Find out about HIV so you can help them and also stay safe.



Ask the person with HIV how they would like you to help (support) them.



Be there, hug them and show that you care.



Help them eat the right foods and take their treatment when and how the doctor tells them.



Respect their business (privacy). It is up to each person with HIV who they tell.



Talk to other people in your community about HIV so they can be supportive, stay safe and not worry about getting HIV.



We all have a job to stop the shame around HIV and STIs.



Services that can help



This section tells you about the types of services that are available and how to contact them.

1800 telephone numbers are free when dialed from any Australian phone line including mobile.

Some HIV and LGBTIQ+ organisations have Aboriginal and Torres Strait Islander project officers who can help.

You can find out where your local sexual health clinic and Aboriginal Medical Service is on the Better to Know website at www.bettertoknow.org.au

HIV and LGBTIQ+ Organisations



Most states and territories have an HIV and LGBTIQ+ organisations, and some have offices in regional towns. These organisations can tell you more about HIV as well as the best places to get help. Some of the bigger organisations offer their own support services.



PWHIV organisations and Positive Living Centres

PWHIV stands for People with HIV. There are PWHIV organisations in most states and territories.

They provide a safe place for people with HIV to get together, as well as confidential (private) support, advice, and referral to services.

In some states, PWHIV services also have Positive Living Centres, which offer support services like free/cheap meals, vitamin supplements and counselling.



Needle and Syringe Program (NSPs)

If you inject drugs, you can get new gear (needles, syringes, other injecting equipment) and helpful information from Needle and Syringe Programs (NSPs). You can ask the HIV and LGBTIQ+ organisation or your health worker where to find one near you.

To find NSPs in your area:
www.nspdirectory.aivl.org.au





Aboriginal Community Controlled Health Services

Aboriginal Medical Services (AMSs) provide confidential testing for HIV and other STI's (sexually transmissible infections), as well as general health care, counselling, and support. Some can provide HIV treatments.

There are more than 190 AMSs in Australia. You can search for your nearest AMSs on the following state and territory organisations' websites:

Australian Capital Territory



Winnunga Nimmityjah
Aboriginal Health and
Community Services
(WNAHCS)

www.winnunga.org.au

New South Wales



Aboriginal Health & Medical
Research Council of NSW
(AH&MRC)

www.ahmrc.org.au

Northern Territory

AMSANT

Aboriginal Medical Services
Alliance Northern Territory
(AMSANT)

www.amsant.org.au

Queensland



Queensland Aboriginal &
Islander Health Council (QAIHC)
www.qaihc.com.au

South Australia



Aboriginal Health Council of
South Australia (AHCSA)
www.ahcsa.org.au

Tasmania



Tasmanian Aboriginal Centre
www.tacinc.com.au

Victoria



Winnunga Nimmityjah
Aboriginal Health and
Community Services
(WNAHCS)
www.winnunga.org.au

Western Australia



Aboriginal Health Council
of WA (AHCWA)
www.ahcwa.org.au



HIV Services

Australian Capital Territory



Meridian
02 6257 2855
www.meridianact.org.au

New South Wales



ACON
Sydney – 02 9206 2000
Hunter – 02 4962 7700
Northern Rivers – 02 6622 1555
1800 063 060
www.acon.org.au



Positive Life
NSW 02 8357 8386
1800 245 677
www.positivelife.org.au

Bobby Goldsmith
Foundation

Bobby Goldsmith Foundation
(BGF)
02 9283 8666
www.bgf.org.au

Northern Territory



NTAHC
08 8944 7777
Alice Springs
08 8953 3172
www.ntahc.org.au

Queensland



Queensland Council for LGBTI
Health
Brisbane – 07 3017 1777
1800 177 434 (freecall outside
Brisbane only)
Cairns – 07 4041 5451
1800 884 401
www.quac.org.au



2Spirits
(07) 3017 1777
www.qc.org.au



Queensland Positive People
(QPP)
07 3013 5555
1800 636 21
www.qpp.org.au

Tasmania

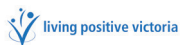


Tasmanian Council on AIDS,
Hepatitis & Related Diseases
(TasCAHRD)
03 6234 1242
1800 005 900
www.redthread.org.au

Victoria



Thorne Harbour Health
03 9865 6700
www.thorneharbour.org



Living Positive Victoria
03 9863 8733
www.livingpositivevictoria.org.au

Western Australia



WA AIDS Council (WAAC)
08 9482 0000
www.waac.com.au

Drug User Organisations

To find NSPs in your area: www.nspdirectory.aivl.org.au

Australian Capital Territory



Canberra Alliance for Harm
Minimisation and Advocacy
(CAHMA)
02 6253 3643
www.cahma.org.au

New South Wales



NSW Users and AIDS Association
(NUAA)
(02) 9171 6650
www.nuaa.org.au

Northern Territory



NTAHC
08 8944 7777
www.ntahc.org.au
Alice Springs
08 8953 3172



Queensland



QuIHN
07 3620 8111
1800 172 076 (free call)
www.quihn.org

South Australia



Hepatitis SA
08 8362 8443
hepatitissa.asn.au

Tasmania



Tasmanian Users and Health
Support League (TUHSL)
www.aivl.org.au/aivl-member-organisations

Victoria



Harm Reduction Victoria
03 9329 1500
www.hrvic.org.au

Western Australia



Peer Based Harm Reduction WA
08 9325 8387
www.harmreductionwa.org
Bunbury
08 9791 6699

National Organisations



Health Equity Matters
02 9557 9399
www.healthequitymatters.org.au



Anwernekenhe National HIV Alliance
(ANA)
02 9557 9399
ana@healthequitymatters.org.au



National Association of People
with HIV Australia (NAPWHA)
02 8568 0300
1800 259 666
www.napwha.org.au



Positive Aboriginal and Torres
Strait Islander Network (PAT SIN)
1800 259 666
[www.napwha.org.au/about-us/
patsin](http://www.napwha.org.au/about-us/patsin)



Australian Injecting & Illicit Drug
Users League (AIVL)
02 5110 3018
www.aivl.org.au



Scarlet Alliance, Australian Sex
Workers Association
02 9517 2577
www.scarletalliance.org

Glossary



AIDS

Acquired Immune Deficiency Syndrome – HIV will become AIDS if it is not treated. AIDS can kill.

Antibodies

Antibodies are made by the immune system to fight germs

Antiretroviral therapy

Medicine that attacks HIV

CD4 (T4) cell

Part of your blood that protects you from illness and that is damaged by HIV.

HIV

Human Immunodeficiency Virus - a germ you can get from sex, sharing needles, and can be passed on from mothers to baby.

Immune System

Your body's way of protecting itself against illness

Immunodeficiency

Your body is not able to protect itself

Hep B

Hepatitis B – a type of liver sickness caused by a germ mainly spread by sex, sharing needles, and mothers to baby.

Hep C

Hepatitis C – a type of liver sickness mainly spread by sharing injecting equipment.



NSP

Needle and Syringe Program – a place where people who inject drugs can get injecting equipment, information and support

PEP

Post-Exposure Prophylaxis – a medication you take everyday for a month very soon after a possible exposure to HIV to prevent HIV.

PLHIV

People Living with HIV

PrEP

Pre-Exposure Prophylaxis – a medication that HIV negative people take to prevent HIV.

STIs

Sexually Transmissible Infections – diseases that can be passed on when having sex

Treatment as prevention

When a person with HIV takes treatment and has an ‘undetectable viral load’, they cannot pass on HIV.

Undetectable viral load

When a person with HIV takes treatment and the amount of HIV in their blood is so small, it doesn’t show up in the tests.

Viral load

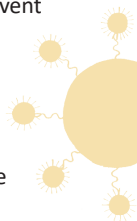
How much HIV is in your blood

Virus

A small organism (germ) often causing illness

Window period

How long it takes for HIV to show up in a test – usually 6 to 12 weeks after it enters your body

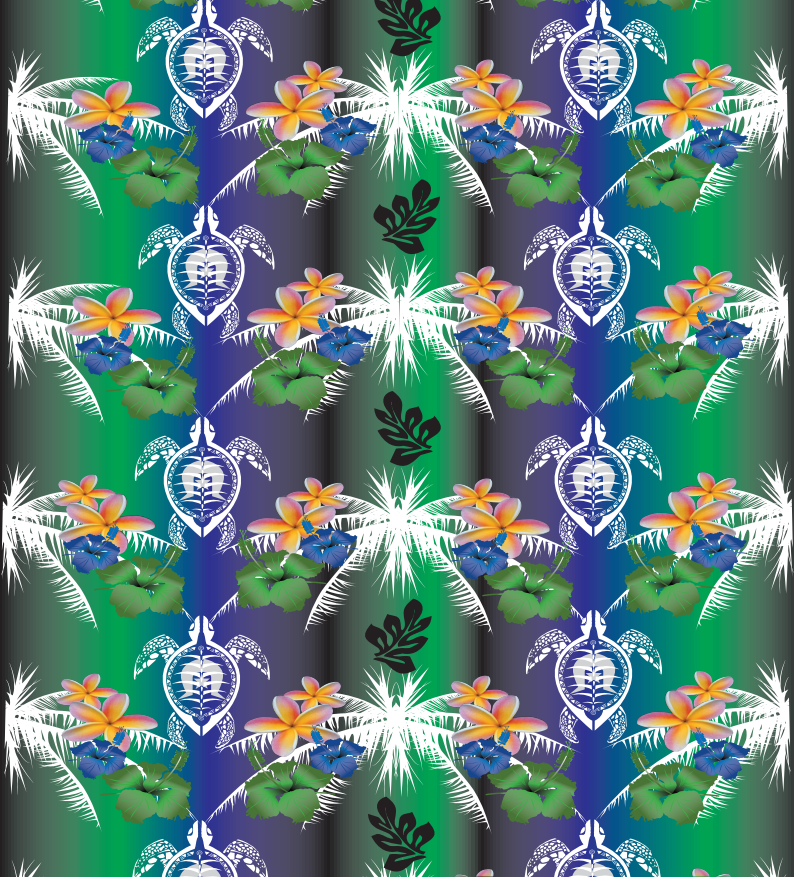


Us Mob and HIV



Visit Us Mob and HIV Online

www.usmobandhiv.org.au



www.usmobandhiv.org.au