



Transgender Health  
A/Prof Katie Wynne



## Gender diversity: transgender, gender diverse and non-binary people

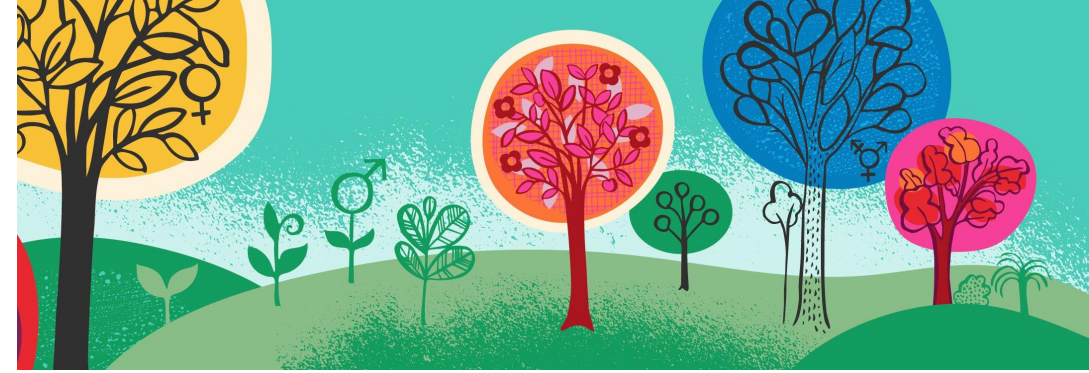
- Society has become more aware and accepting of gender diversity
- Estimates in young people have risen from <0.01% to 2.3%
- Increasingly recognized by healthcare as part of human diversity
- Gender diverse people are vulnerable due to stigma and minority stress
  - Risk of family rejection
  - Social exclusion
  - Reduced healthcare access
  - Bullying and assault
- Psychological distress, suicidal ideation/attempts and self-harm x3-5
- All-cause mortality x3



**3 x**

**all-cause mortality**





## Gender-affirming care

- Defined by the WHO as '*any single or combination of social, psychological, behavioural or medical interventions designed to support and affirm an individual's gender identity*'
- RACP (2020) states clinical care needs to be "*non-judgemental, supportive and welcoming for children, adolescents and their families*" "*withholding or limiting access to care and treatment would be unethical and would have serious impacts on the health and wellbeing of young people.*"

## Goals

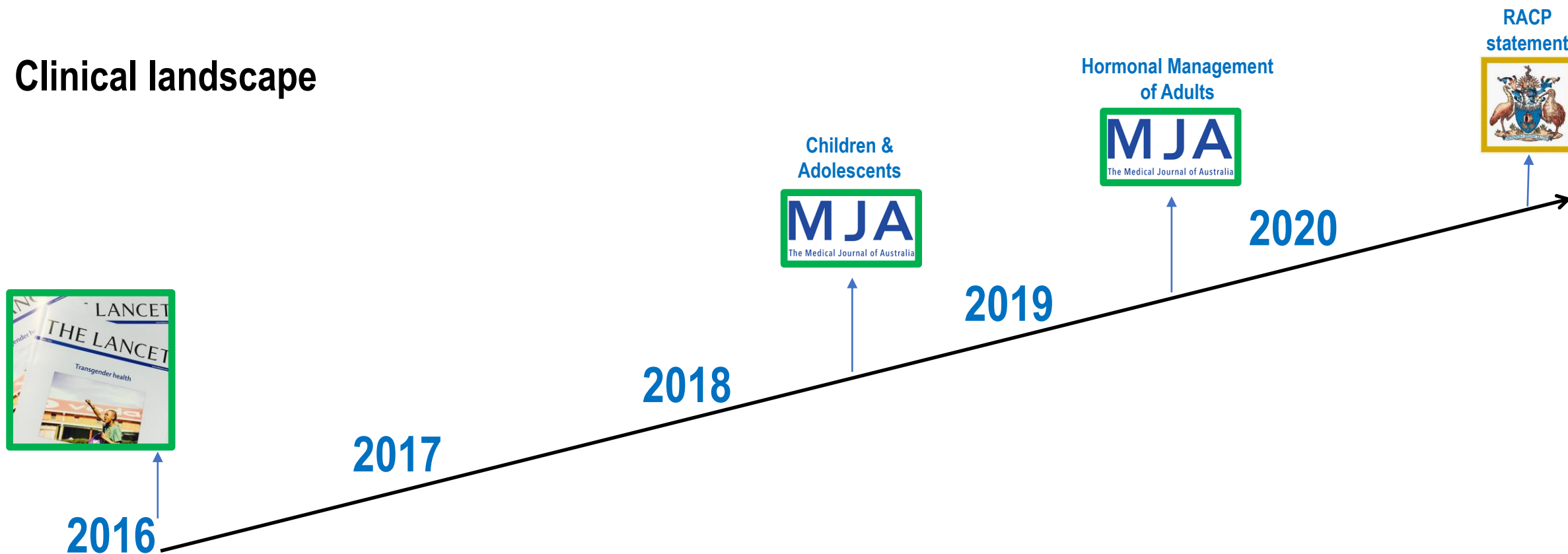
**Improve health outcomes**

**Improve social outcomes**

**Alleviate gender incongruence**

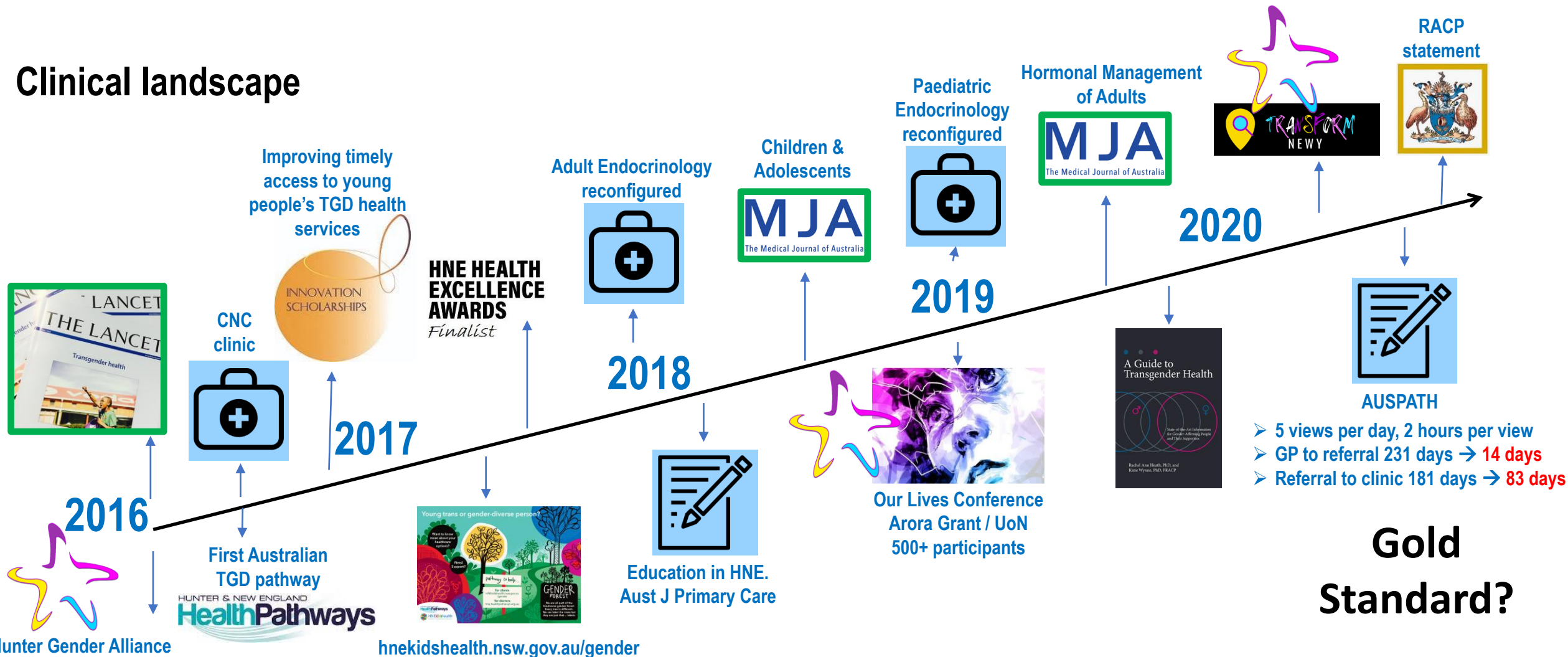


## Clinical landscape





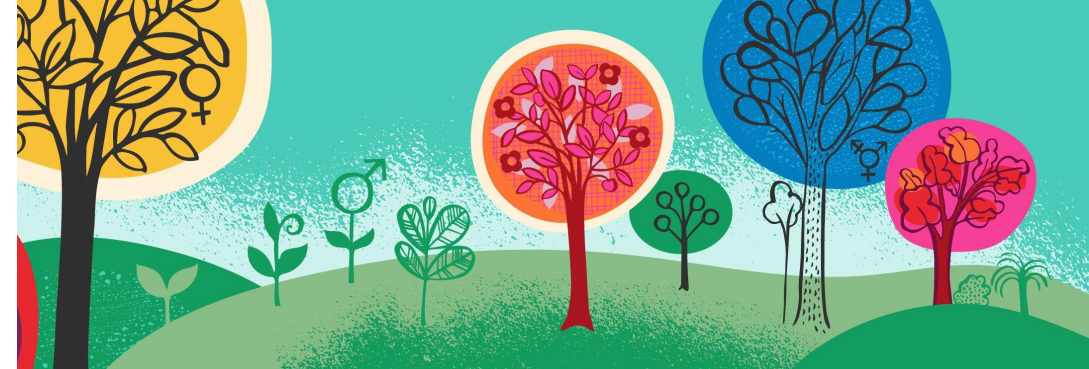
## Clinical landscape





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Children, Young People & Families



## Legal complexities: <18 years

***Re Jamie***

**2013**

- Overturned existing law that required Family Court authority for puberty blockers <18 years
- Providing the parents and young person's medical practitioner in agreement

***Re Kelvin***

**2017**

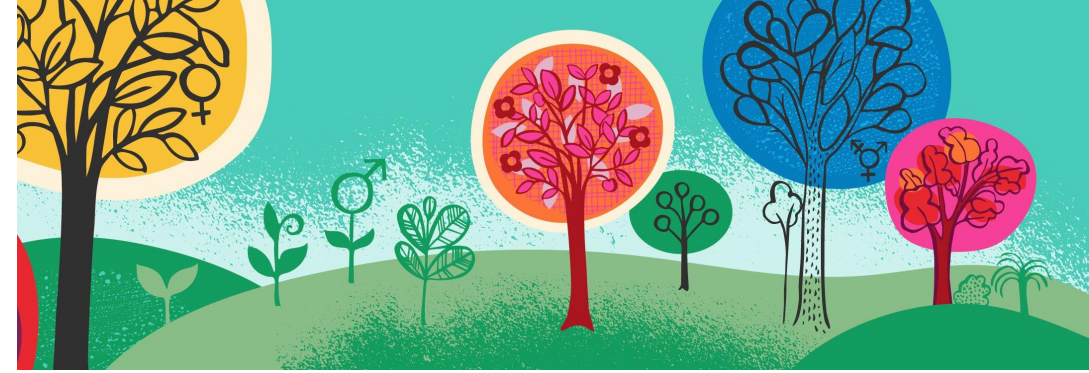
- Enabled young people to commence gender-affirming hormones <18 years without Family court authorisation
- No comment on what to do if parents were in dispute. The effect was to bring hormonal therapy within parental consent, and allow the individual to consent if deemed '*Gillick*-competent'

***Re Imogen***

**2020**

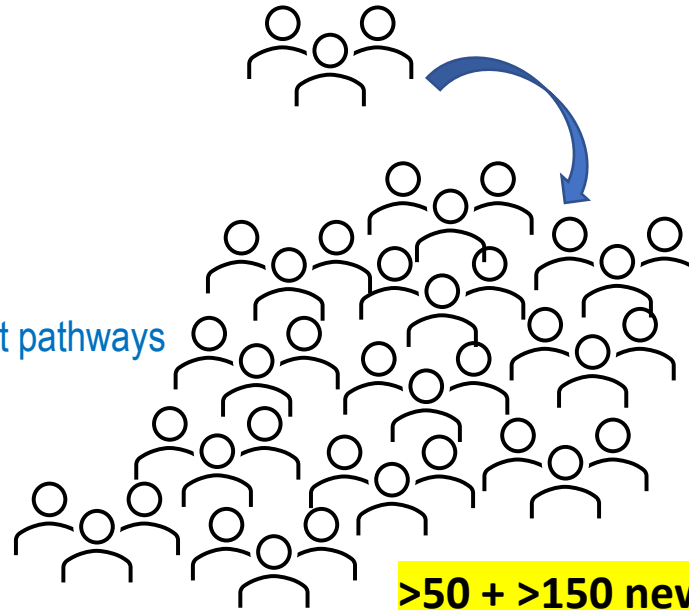
- Young people can only receive hormone treatment when there is no dispute between parents, the physician and young person regarding the diagnosis, treatment and *Gillick*-competence.
- Therefore, physicians must ascertain if both parents consent, even if the individual has capacity, and if there is not consensus the Family Court must provide authority.



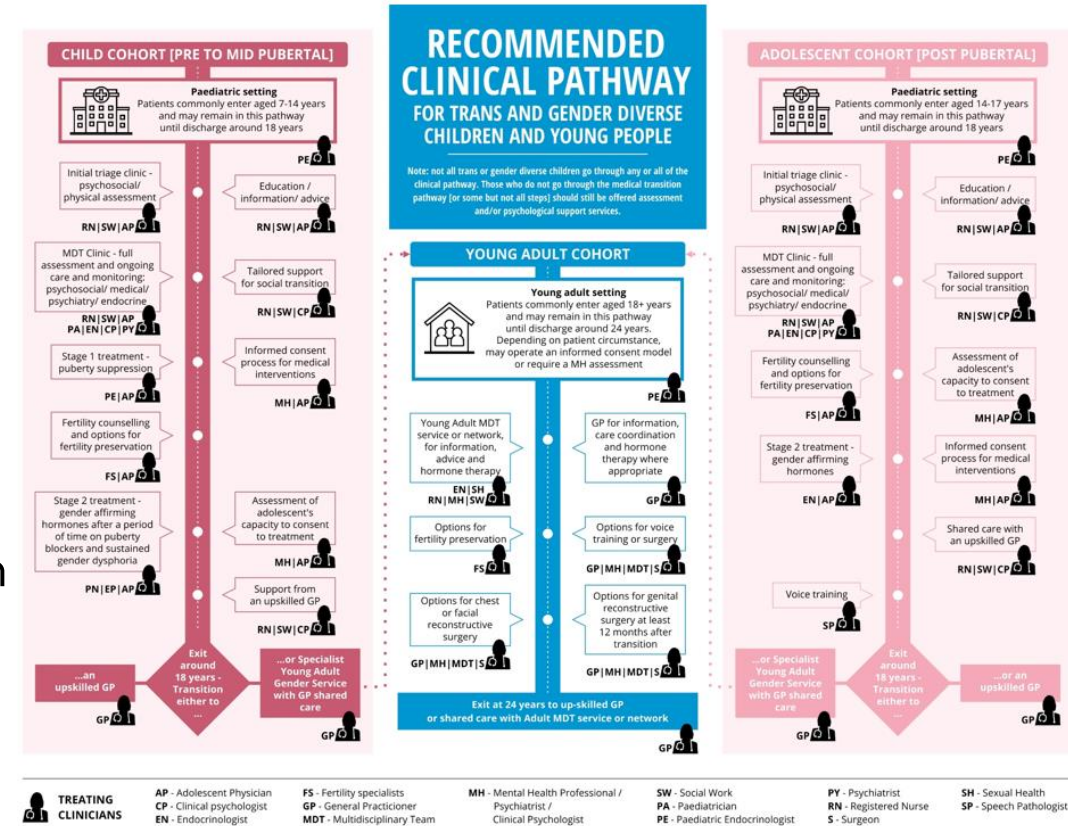


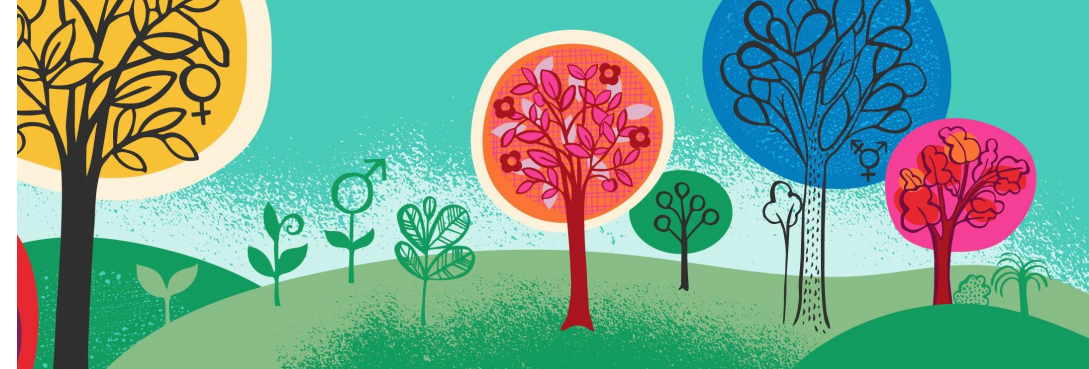
## NSW Ministry of Health

- National and International guidelines endorsed by recognised professional associations
- In NSW has inequity of access, a poorly integrated system and multiple service gaps
- Increasing referrals to services
- Funded core services in the public system
- A service model endorsed by NSW Health
- Recommended child, adolescent and adult pathways
- Multidisciplinary team as gold standard

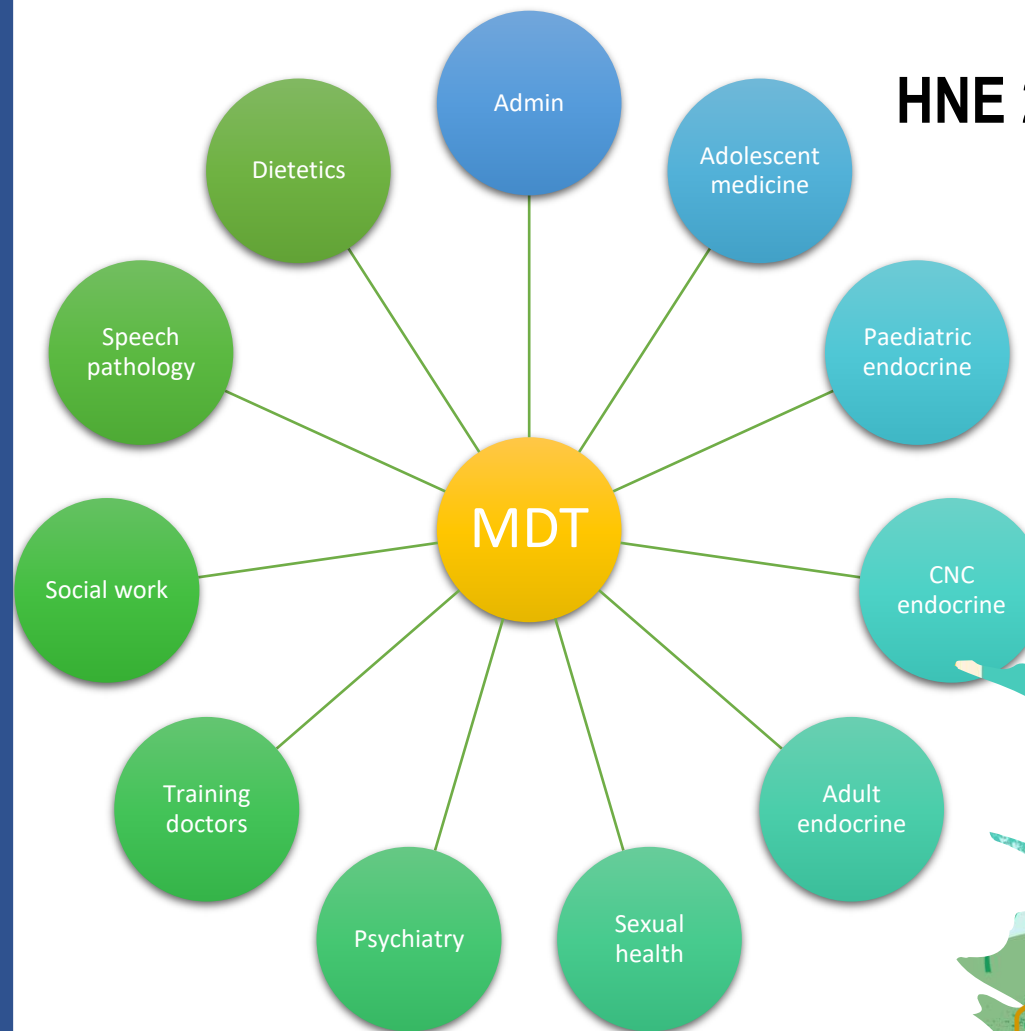


>50 + >150 new referrals/year

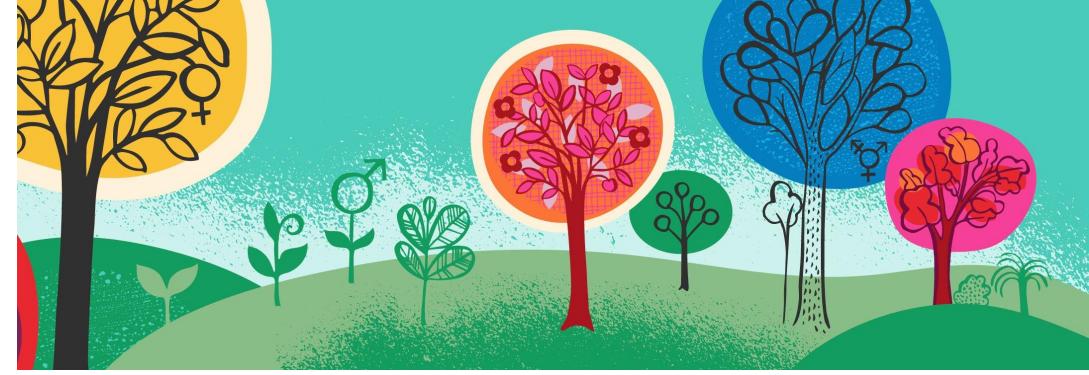




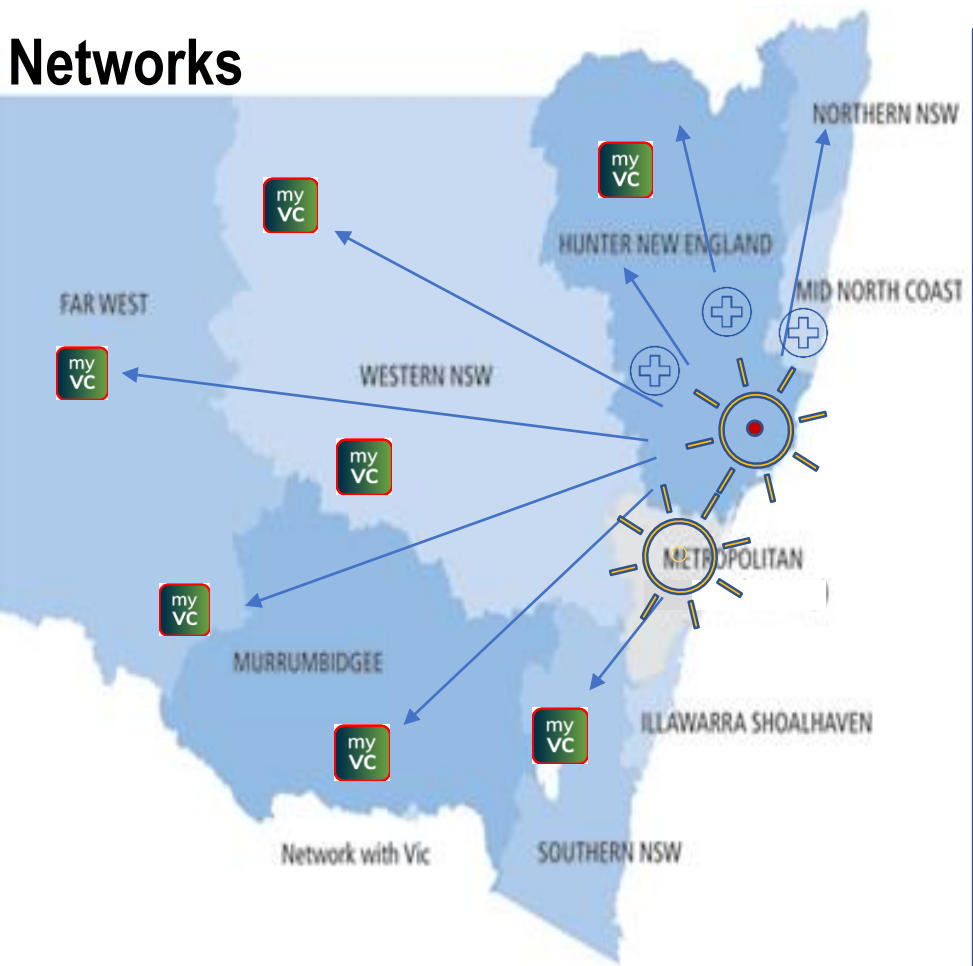
## HNE 2021: from clinic to service Maple Leaf House























## Networks



Professional bodies	Community organisations
     	   
	
<b>Clinical services</b>   	 



**John Hunter**  
**Children's Hospital**  
 Children, Young People & Families



**Hunter**  
**Healthcare**  
**Experience**

n=76  
 7-77 years  
 90.6% GP

Satisfaction  
 index **36/45**

My GP helps me

My GP has enough time for me

I trust my GP

My GP understands me

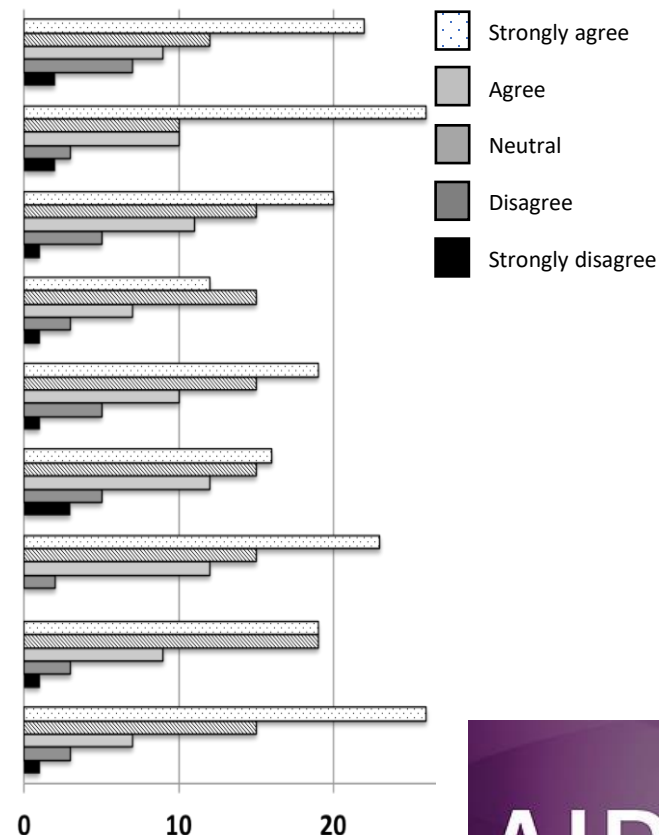
My GP is dedicated to helping me

My GP and I agree on my problems

I can talk to my GP

I feel content with my GP's treatment

My GP is easily accessible



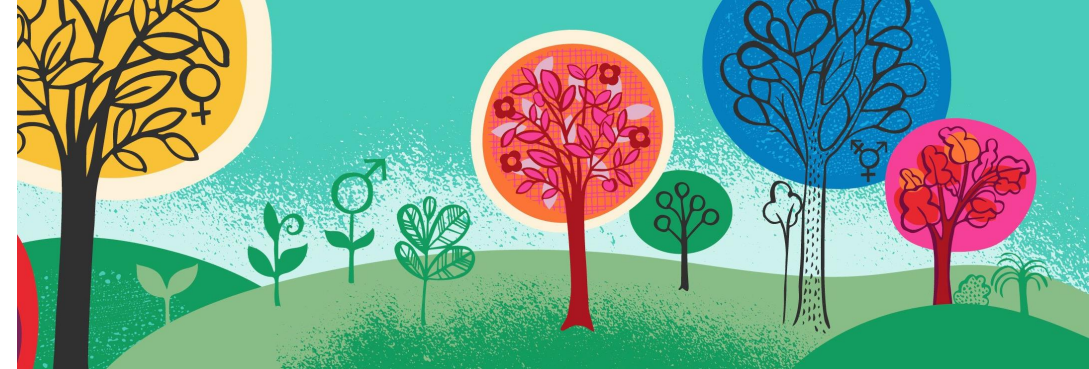
A good patient-doctor relationship is associated with better treatment adherence and improved outcomes. The **constructive GP relationship** provides a **key platform to deliver care**





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01

## Name, pronoun and gender identity

Screen MH red flags  
Self-medicating

02

## Referral to MLH

Prue Lopez or Rowan Seckold  
Adolescent physician  
Katie Wynne or Judy Luu

03

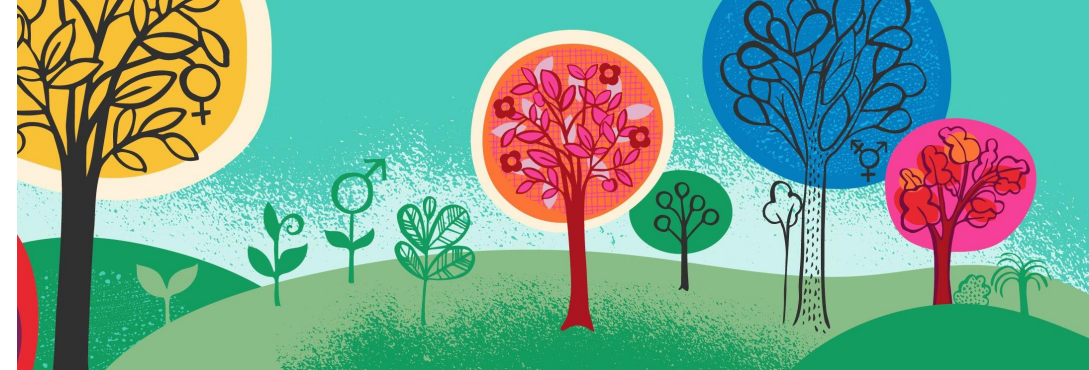
## Management

Norethisterone for puberty suppression > Tanner 3  
Bridging hormones using informed consent 18-24 years



- Clinical decision-making for young people **<18 years** requires expert specialist assessment and support.
- Family Court is not needed if the child, parents/guardians, and treating clinicians agree.
- The treating medical practitioner must seek and document written consent of all parents/guardians and child (if Gillick competent).
- **Family Court is needed** in the case of dispute about Gillick competence, diagnosis or proposed treatment
- For a child **<16 years** clinicians must also consider if the intervention is a 'special medical treatment': *a intended or reasonably likely to render the child permanently infertile, unless the treatment is administered to treat a life-threatening condition*.
- If so, consent of the NSW Civil and Administrative Tribunal - **NCAT is required**.





- Support of children and adolescent's gender identity by family and community is associated with better mental health<sup>1,2</sup>
- GnRH treatment has been offered since the 1990's, is supported by a moderate level of evidence and is recommended in international evidence-based and consensus-based guidelines<sup>3</sup>
- Young people who have gender dysphoria as children, continue to identify as other gender in adolescence, and who show increasing distress about unwanted pubertal body changes are **likely** to continue to experience gender incongruence through adolescence and adulthood<sup>3,4</sup>
- Puberty suppression prescribed according to guidelines is associated with good or improved mental health and reduced suicidality in transgender adolescents<sup>5-10</sup>

**"There is now moderate evidence supporting the use of puberty suppression in earlier adolescence and the use of estrogen or testosterone in later adolescence,** for adolescents who have gender dysphoria which is worsening in early adolescence who strongly wish for and request these treatments. The evidence is strong and unequivocal for the physical effects of pausing puberty. The evidence is moderate for short and medium term improvements in gender dysphoria, mental health and quality of life. The evidence is limited beyond a 7 year follow-up period into young adulthood. **This is acceptable, because the treatment is used when distress and risks (on no treatment) are so grave..."**







> 18  
years

## Informed consent

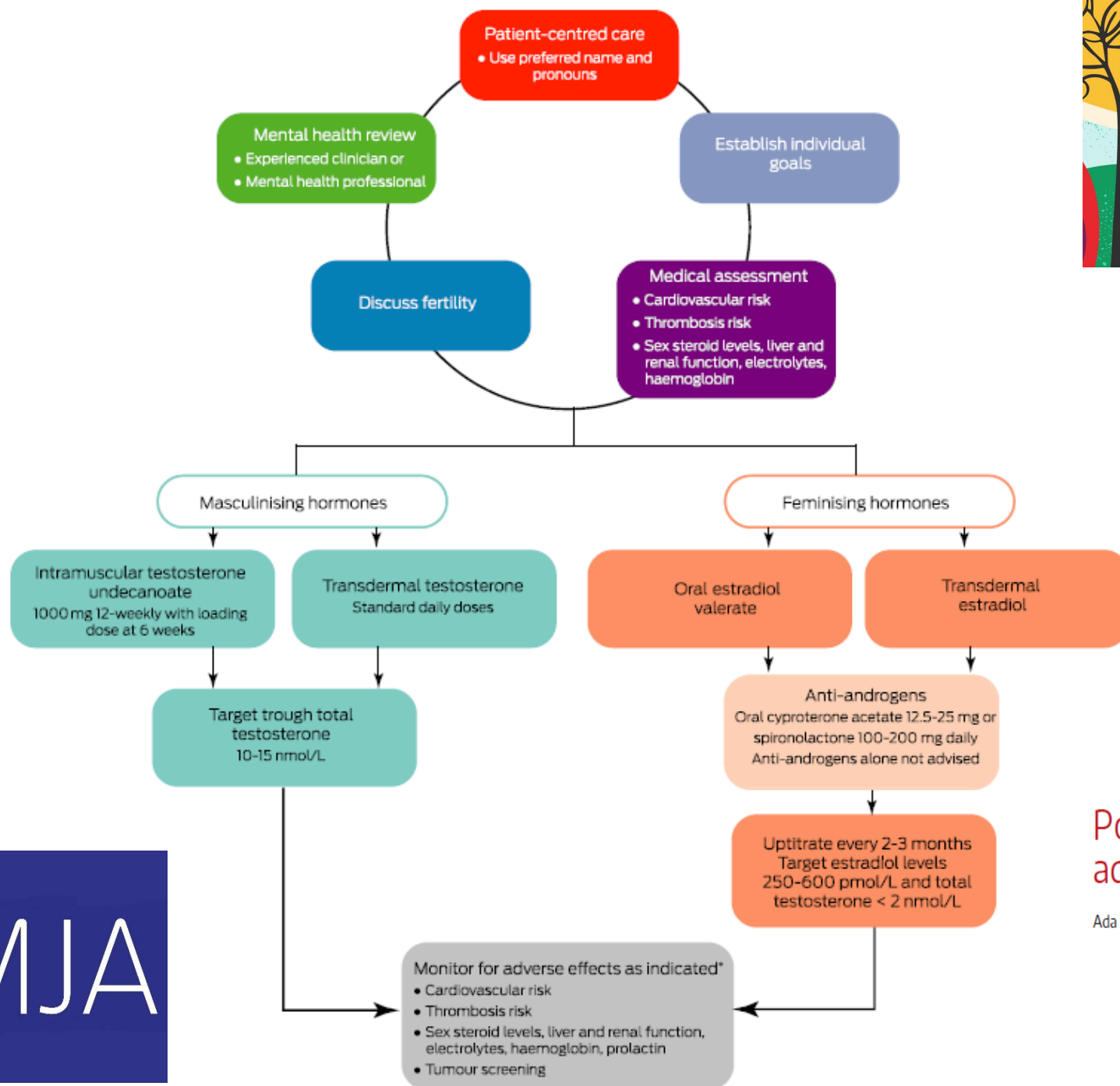
Physical changes  
Time course  
Potential adverse effects  
Irreversible nature of some changes

Soc 7 WPATH 2011

## Position statement on the hormonal management of adult transgender and gender diverse individuals

Ada S Cheung<sup>1</sup> , Katie Wynne<sup>2</sup> , Jaco Erasmus<sup>3</sup>, Sally Murray<sup>4</sup>, Jeffrey D Zajac<sup>1</sup>

Med J Aust 2019; 211 (3):127-133







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## Androgen Blockers

- **Spironolactone** 100mg od (50-100mg): aldosterone antagonist that also acts at the testosterone receptor, and weak estrogen-like activity. Risks: hyperkalaemia, rarely GI bleed.<sup>1</sup>
- **Cyproterone Acetate** 12.5mg od (12.5-25mg): progestin → gonadotrophins reduces testosterone synthesis, testosterone receptor antagonist. Risks: mood, LFTs, rarely meningioma<sup>2</sup>
- **GnRH analogues** (Zoladex 10.8mg 12-weekly, 3.6mg monthly)

## Estrogen at 6-12 weeks

- Estradiol **oral** tablets start 1-2mg, up to 1-8mg (**Progy Nova**)
- Estradiol **gel** start 1mg, up to 1-5mg (**Sandrena** / Estragel)
- Estradiol **patch** start 25-50mcg, up to 50-100mcg/d (**Estradot** / **Estraderm**)



17b estradiol

1. Gulmez BJCP 2008 (RRx13, >55 years on 100mg+)  
2. Gill BJCP 2011 (RRx11, 60/100,000 person-years)



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<b><i>Preferred first-line feminising estradiol preparations</i></b>	
Estradiol valerate (oral)	25 (71.4%)
Estradiol patch (transdermal)	3 (8.6%)
Other (depends on age of patient; transdermal in older, oral in younger)	1 (2.9%)
No response	6 (17.1%)
<b><i>Do you use anti-androgen treatments in addition to estradiol therapy?</i></b>	
Almost always	20 (57.1%)
Often	6 (17.1%)
Sometimes	2 (5.7%)
Only if I can't suppress the testosterone on estradiol alone	1 (2.9%)
No response	6 (17.1%)
<b><i>Anti-androgen medications used (more than one option could be selected)</i></b>	
Spironolactone	27 (93.1%)
Cyproterone acetate	28 (96.6%)
5-alpha reductase inhibitors (finasteride, dutasteride)	10 (34.5%)
Bicalutamide	1 (3.5%)
No response	6 (17.1%)

Anonymous survey of prescribing given to ANZPATH members 2017.<sup>1</sup>

N=43 (GP (62%), Endo (20%), Sex Health (17%))

## Targets

- No data on gradual vs. rapid titration
- Induction of puberty
- Gradual up-titration every 2-3 months<sup>2</sup>
- Estradiol 250-600pmol/l<sup>2,3</sup>
- Testosterone <2nmol/l and/or symptoms<sup>3</sup>
- Timing: 4 hours post-dose or 48 post-dose for patches
- Adjust according to the patient's biochemical response
- Menopause estradiol 100-250pmol/l?



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## Testosterone

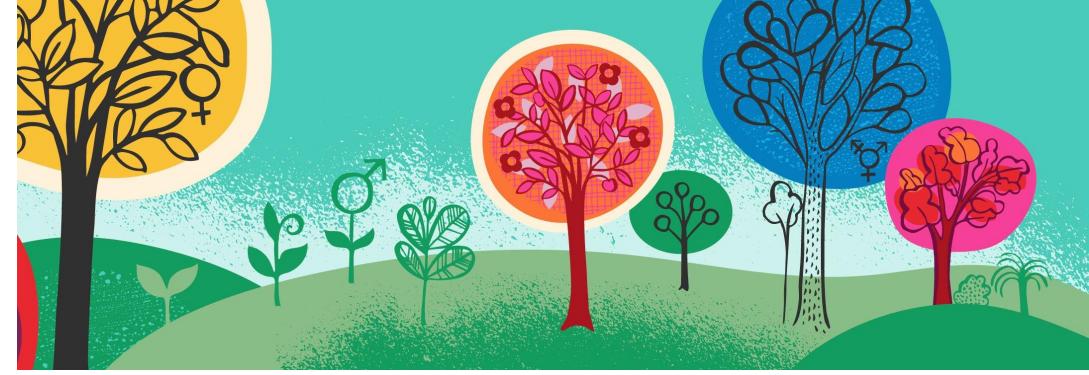
'Androgen deficiency due to an established testicular disorder'. In consultation with a paediatrician, endocrinologist, urologist or sex health physician. Any gender marker.

- Testosterone undecanoate 1000mg im, 12-(9-15) weekly, first two doses 6 weeks apart ([Reandron](#))
- Testosterone 1% (50mg/5g) gel ([Testogel](#))
- Testosterone 1% (12.5mg/actuation) gel pump, 1-4 actuations ([Testagel](#))
- Testosterone 2% (23mg/actuation) gel pump 1-2 actuations, ([Testavan](#))
- Testosterone 5% (50mcg/ml) cream, 0.5-2mg ([Androforte 5](#))

## Non-PBS

- Testosterone enantate ([Primoteston](#)) 250mg im 2-4 weeks or 75-80mg (50-100mg) sc weekly<sup>1</sup> Testosterone esters ([Sustanon](#)) - 250mg im 3-6 weeks





<i>Preferred first-line masculinising testosterone preparations</i>	
Testosterone undecanoate (intramuscular)	17 (48.6%)
Testosterone enantate (intramuscular)*	11 (31.4%)
Testosterone gel/cream (transdermal)	5 (14.3%)
No response	1 (3.0%)

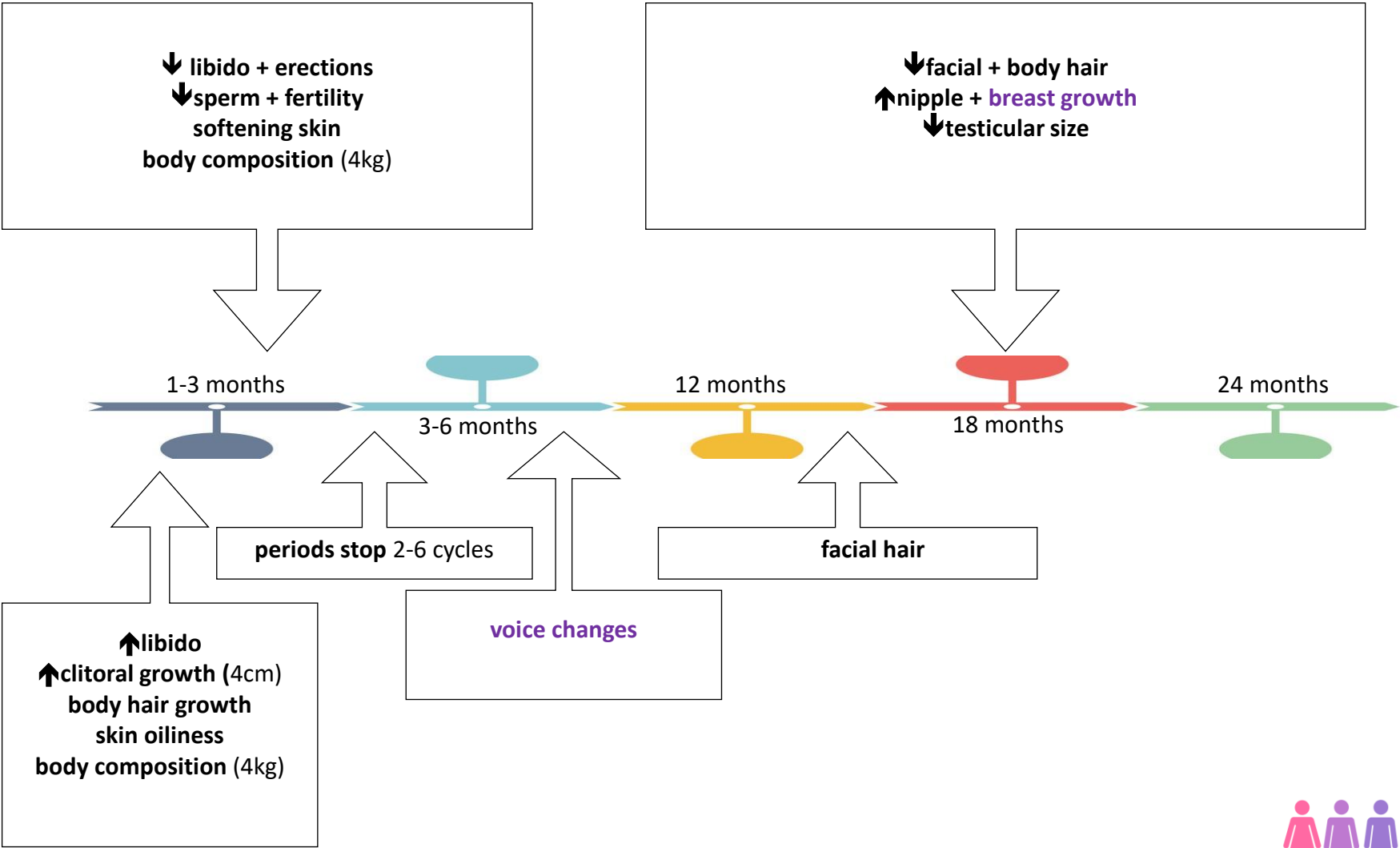
Anonymous health provider survey<sup>1</sup>



## Biochemical target

- Booster initiation (first two 6 weeks apart) or start with 12-weekly if patient choice.
- Trough testosterone 10-15nmol/l
- Peak testosterone 15-20nmol/l

1. Bretherton IMJ 2018
2. Wittert Andrology 2016





## Gender diversity: transgender, gender diverse and non-binary people

**In summary.....** Hormonal transition is safe and effective.

A systematic review of all studies to 2014. showed no increased risk of cancer or early mortality.<sup>1</sup>

For >18 years, can be delivered in **primary care** with specialist referral for difficult cases<sup>2</sup>

Further large cohorts:

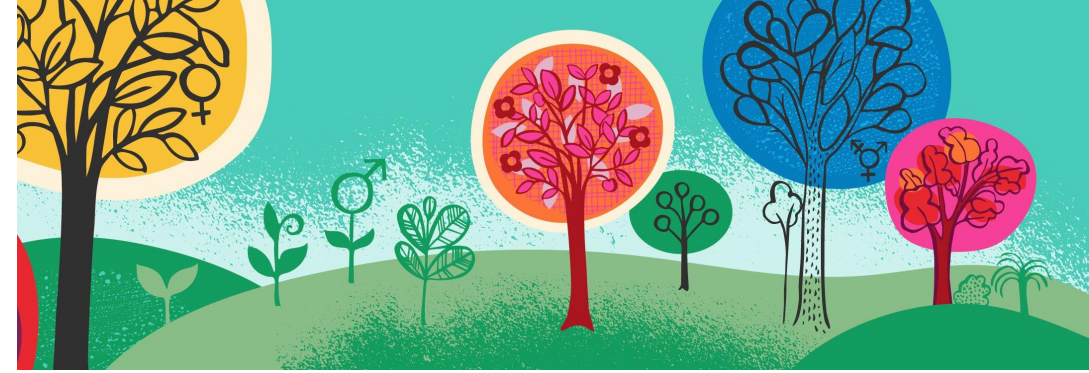
- ENIGI, Europe
- STRONG, US
- Veteran's Health, US
- **Trans 20, Australia<sup>3</sup>**

**Australian Position Statements** for children, adolescents<sup>4</sup> and adults<sup>5</sup>

A National Research Consortium for TGD young people has been established.

1. Weinand JTE 2015
2. Arora AJPH *in press* 2019
3. Tollit BMJ Open 2019
4. Telfer Med J Aust 2018
5. Cheung Med J Aust 2019



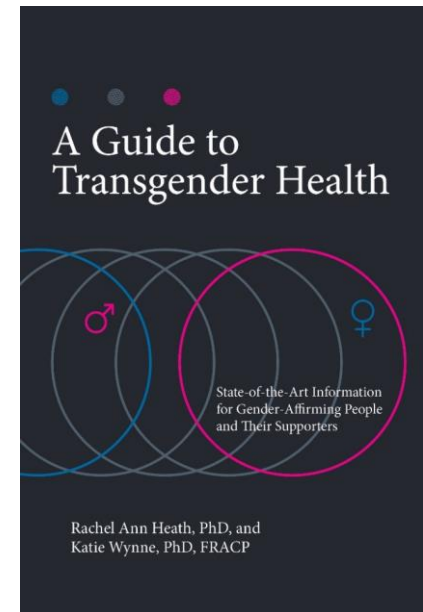


<https://www.gires.org.uk/e-learning/>

- for GP's
- caring for gender non-conforming young people
- transgender awareness for employers and service providers
- promoting trans equality in further education

[www.lgbthealtheducation.org/resources/in/transgender-health/](http://www.lgbthealtheducation.org/resources/in/transgender-health/)

- Gender Affirmative Health Care
- Cross Sex Hormone Therapy
- Surgical Gender Affirmation
- Primary and Preventative Care
- Caring for Gender Non-conforming Youth in Pediatrics and Primary Care



Heath and Wynne (2019) available at [amazon.com.au](https://www.amazon.com.au)

