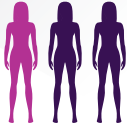


Testosterone for Women

Let's talk HSDD



1 in 3 women (aged 40-64) will experience HSDD which can severely impair relationships, mental health, social functioning and overall QOL.^{1,2,3}

12% Reported sexual problems during the first 5 minutes without being asked.

36% Reported sexual problems when asked about sexuality after 5 minutes.

Prevalence rose from 12% to 48% when they were asked about sexuality after 5 minutes.⁴

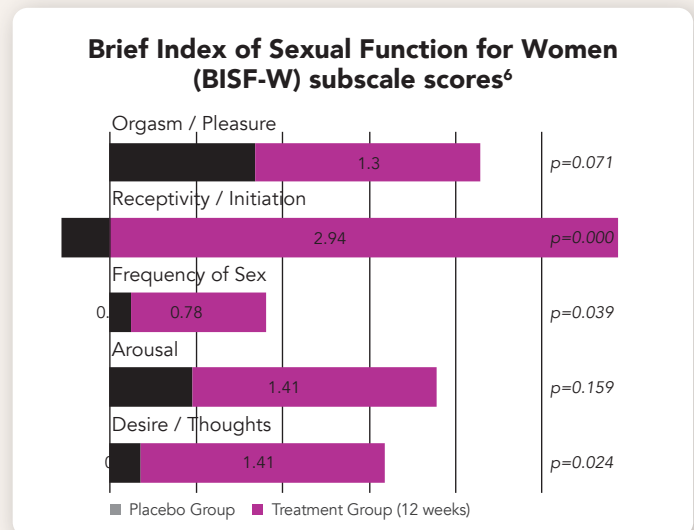
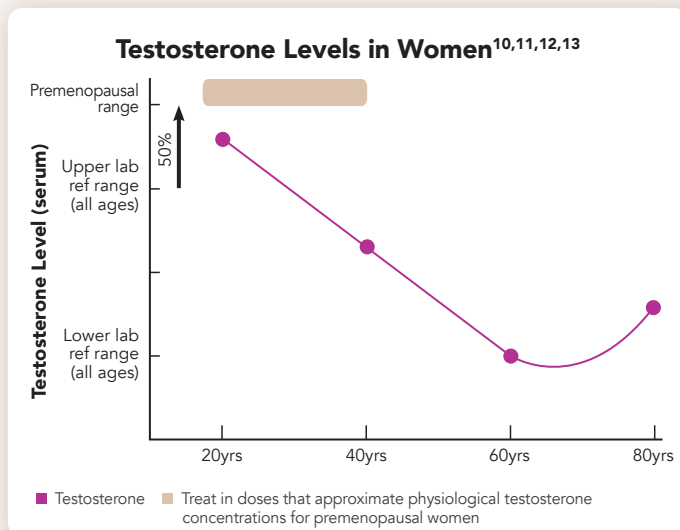
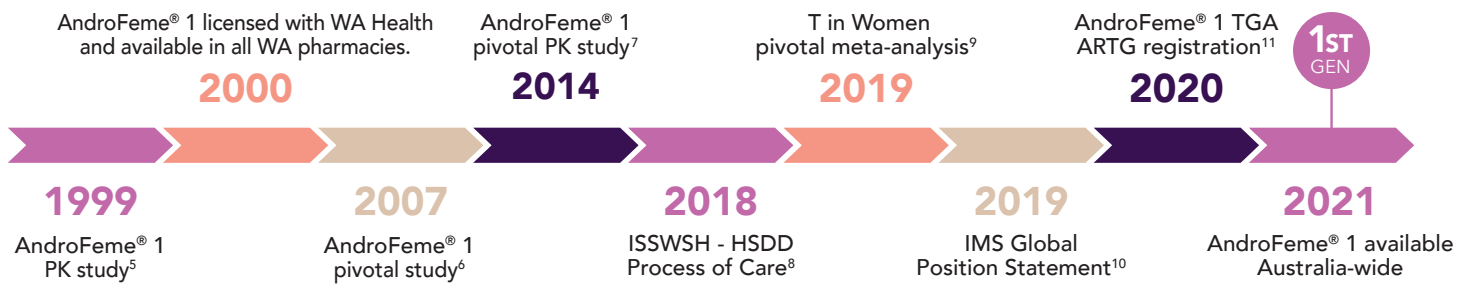
“Many women going through menopause have concerns with sexual function; is this a concern for you?”



Why use AndroFeme® 1?



Indication: The management of hypoactive sexual desire dysfunction (HSDD) in postmenopausal women.



Diagnosis



Screening

1. Menopause Checklist¹⁴
2. DSDS: Decreased Sexual Desire Screener¹⁵

Decreased Sexual Desire Screener (DSDS)

Patient MAY qualify for acquired, generalized HSDD if YES to Q 1-4 and NO to all factors in Q 5.

Patient MAY qualify for acquired, generalized HSDD if YES to Q 1-4 and YES to any factors in Q 5 if determined by clinical judgment*

*see reference 15

Modifiable Factors

Contributing Factors

- Menopausal symptoms
- Physical examination
- Psychological evaluation
- Medical conditions
- Medications

Gynaecology Factors

- Vaginal atrophy
- Dryness and/or pain

Sexual / Relationship Factors

- Relationship problems
- Partner's health
- Sexual assault / abuse

Serum – what to measure and why?

- It is recommended that serum testosterone monitoring be used as an aid to treatment rather than as the primary measure of efficacy.¹¹
- No cut-off blood level can be used for any measured circulating testosterone to differentiate women with and without sexual dysfunction.¹⁰
- Serum testosterone concentrations **must not** be used as a treatment target.
- Attend the same laboratory for each blood test.¹¹
- Baseline testosterone and sex hormone-binding globulin (SHBG) levels should be obtained prior to initiation of testosterone therapy and 3-6 weeks after therapy initiation.
- Measuring testosterone should be used for possibly overuse, but **must not** be used as the primary guide for patient management.

Total Testosterone Measuring total testosterone as the main biomarker rather than 'free' testosterone, as evidence that 'free' testosterone is the biologically active testosterone fraction is lacking.

SHBG

Testosterone binds to SHBG and bioavailability needs to be considered.

LOW

Be mindful of concurrent therapies that reduce SHBG.
Tibolone
Glucocorticoids

HIGH

Review concurrent therapies to reduce SHBG to mid-range, e.g. oral oestrogen, thyroxine dose.
Change to transdermal oestrogen and re-test SHBG in 12 weeks.

Treatment

The aim of treatment is the resolution of symptoms by administering testosterone in doses that approximate physiological testosterone concentrations in premenopausal women.¹⁰



AndroFeme® 1 1% testosterone cream

Application site	Upper outer thigh or buttock
Starting dose	0.5mL (5mg) once daily
Maximum dose	1mL (10mg)
Dose adjustment	Titrate up or down by 0.25mL increments depending upon symptom response. See monitoring below.
Pack size	50mL tube 100 days using 0.5mL once daily
Private script	\$100 per tube ≈ \$30 per month
PBS status	Not PBS listed

Monitoring / Follow up

The primary indicator of efficacy is symptom improvement in sexual function as reported by each woman. Improvement is not immediate and generally onset takes 4-8 weeks; peaking at 12 weeks.¹¹

Timeline	SHBG/Serum Testosterone	Efficacy/Safety review	Dose Modification (if required)
3-6 weeks	✓		✓
12 weeks	✓	✓	✓
6 months	✓	✓	

Morning blood sample to be taken prior to application of daily dose.

Women with total testosterone concentrations greater than 50% above the upper limit of the **premenopausal reference range** for the assay being used should be advised to reduce the dose of the applied cream.

Monitor dose and efficacy at 3 months and review 6 monthly.
If no efficacy reported at 6 months, cease therapy.

Therapy beyond 24 months should be an informed decision by physician and patient.

Safety

- It is recommended that if the serum testosterone concentration exceeds the upper limit of the **premenopausal range** of the assay being used that clinical evaluation is needed to screen for evidence of hyperandrogenism and a dose reduction considered. Typically serum T levels return to baseline 2-5 days post cessation of therapy.
- Patients should be made aware that long-term skin-to-skin contact particularly with children, can lead to adverse events including signs of virilisation.
- There is a lack of clinical trial safety data beyond 24 months.

PBS Information: Non PBS listed. Available nationally as a private prescription.

Please review full AndroFeme® 1 Product Information before prescribing at www.lawleypharm.com.au/products

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