

Clinical evidence for StrataMGT®

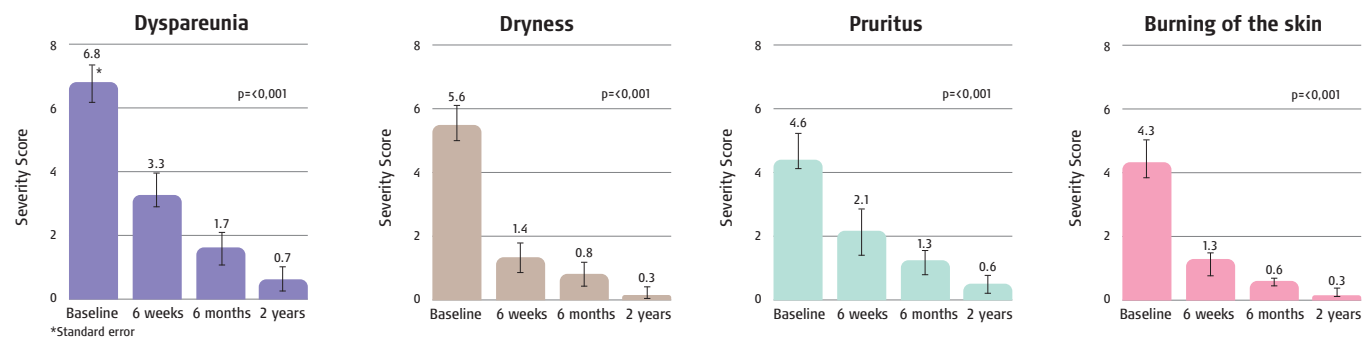
A 52-women open-label, single-arm intervention study was conducted in St. Andrews War Memorial Hospital in Brisbane, Australia to evaluate the efficacy and safety of StrataMGT for the management of genital skin conditions.²¹

Patients were diagnosed with Vaginal atrophy (VA), mean age of 66.3 (range 50 - 74 years); Lichen Sclerosus (LS) and Lichen Simplex Chronicus (LSC), mean age of 54.7 (range 19 - 81 years).

Patients applied StrataMGT daily internally and externally to the genital area for approximately 6 months (short-term analysis, n=52) and up to 2 years (long-term analysis, n=30).

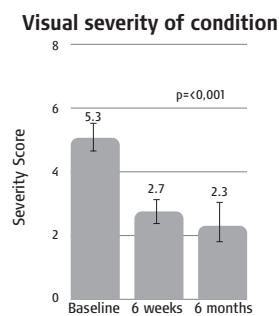
- **Severity of the symptoms:** Patients rated severity of the symptoms using a 10-point scale ranging from 0=normal to 10=worst possible.

The most significant **improvements were seen during the first 6 weeks**, where **symptoms improved up to a 100% resolution** for some patients. **Favorable results** were maintained over **2 years post start of the treatment**.



- **Severity of the condition:** The investigator rated the visual severity of the condition using a 10-point scale ranging from 0=normal to 10=worst possible.

The visual **severity of pathology and clinical signs improved significantly compared to baseline**. The visual pathology of the clitoral hood, urethral area, labia majora and minora, fourchette and perineum showed statistical improvement.



StrataMGT reduces Trans-Epidermal Water Loss (TEWL) by preventing water loss from epidermal and dermal layers, and gas permeability enables **better environment for epidermal migration and healing**.

Treatment **compliance** was very high throughout the study, between 86% and 97% in each visit interval. **No adverse events** were observed.

- StrataMGT is a **suitable alternative** treatment for women suffering from various highly symptomatic vulvovaginal conditions.
- StrataMGT is recommended as a **long-term treatment option without the side effects** of topical corticosteroids and vaginally administered hormonal therapy.

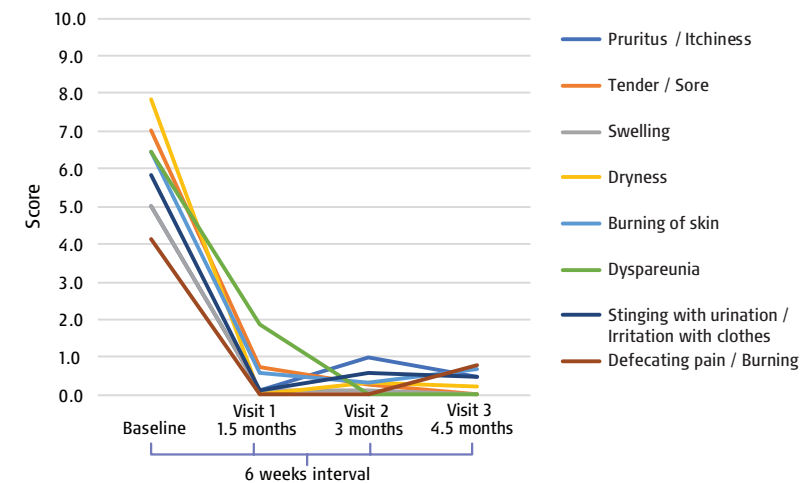
Clinical evidence for StrataMGT®

Case series with 19 patients to test the efficacy of StrataMGT on patients presenting for treatment of chronic uncontrolled vulvovaginal conditions, such as Vaginal atrophy (VA) and Lichen Sclerosus (LS).²²

Dr. Philip Hall, Gynaecologist, MBBS MRMed, FRANZCOG, FRCOG, FACRRM. Pelvic Medicine Centre, St Andrew's War Memorial Hospital Spring Hill Brisbane Australia (2018).



Severity of the symptoms: Patients rated severity of the symptoms using a 10-point scale ranging from 0=normal to 10=worst possible.



Patients were followed up over 3 visits (V1-3), averaging 6 week intervals. Patients mean age of 53.5 (range 30 - 77 years).

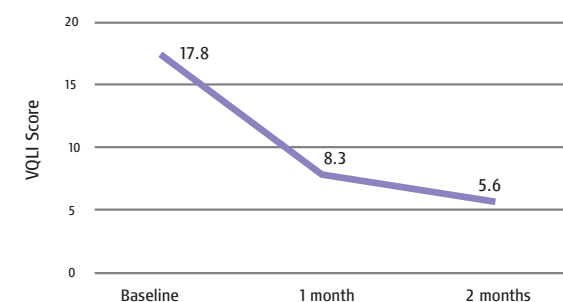
StrataMGT provides **hydration whilst protecting the epithelium and mucosa from irritants** such as urine, infection and friction.

The **most significant improvement** was seen for all patients within the first 6 weeks, where **symptoms improved from 33% through to 100% resolution** in some patients.

The clinical control of symptoms was maintained over Visit 2 (3 months) and Visit 3 (4.5 months) in the majority of patients.

A 9-women non-uncontrolled, open-label intervention study was conducted in Orange Coast Women's Medical Group in Laguna Hills, California, USA to evaluate the efficacy and safety of StrataMGT and the improvement of patient's Quality-of-Life (QoL). All patients were postmenopausal, diagnosed with VA, and treated with StrataMGT for 2 months.²³

Patients rated the impact of vulvar conditions on QoL using Vulvar Quality of Life Index (VQLI) ranging from 0=not impacted, 1-15=mildly impacted, 16-30=moderately impacted, 31-45=severely impacted.



- The **most significant improvement** was seen during the **first month** of treatment - the VQLI score **was reduced by 54%**.
- StrataMGT is **highly efficient** in reducing clinical symptoms associated with VA and improving QoL.