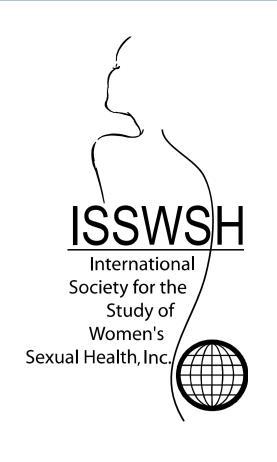
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The Safety and Efficacy of Ablative Fractional Er:YAG Laser Treatment for Vulvar Lichen Sclerosus

Michael Coyle, DO, FACOOG, FPMRS

Coyle Institute, Pensacola, FL, USA



Introduction

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Vulvar lichen sclerosus (VLS) is a chronic skin condition that predominantly affects the anogenital skin.(1) It is characterized by marked inflammation, epithelial thinning, and distinct dermal alteration; after an initial inflammatory phase, there is chronic scarring and skin atrophy.(2) Traditional treatment and

management has incorporated

topical steroid medications, immunomodulators, and supportive therapy.(3)

Studies have shown laser therapy as viable treatment modality for VLS.

The purpose of this study is to show the effectiveness of ablative fractional 2940nm Er:YAG laser treatment for VLS.

Δim

To evaluate the safety and efficacy of ablative fractional 2940nm Er:YAG laser

for treatment of Vulvar Lichen Sclerosus (VLS)

Method

- Female subjects with biopsy proven active VLS were included
- Symptoms of VLS dryness, itching, burning, bleeding, soreness, easily tears, ulcerated lesions
- Demographics, depth of disease, vulvovaginal symptoms (VSQ), 7-point global overall scores for vulvar atrophy symptoms were recorded at baseline, treatment and follow-up visits
- Biopsies were collected at baseline and 3 month follow-up
- Photographs taken at baseline, before treatment and follow-up

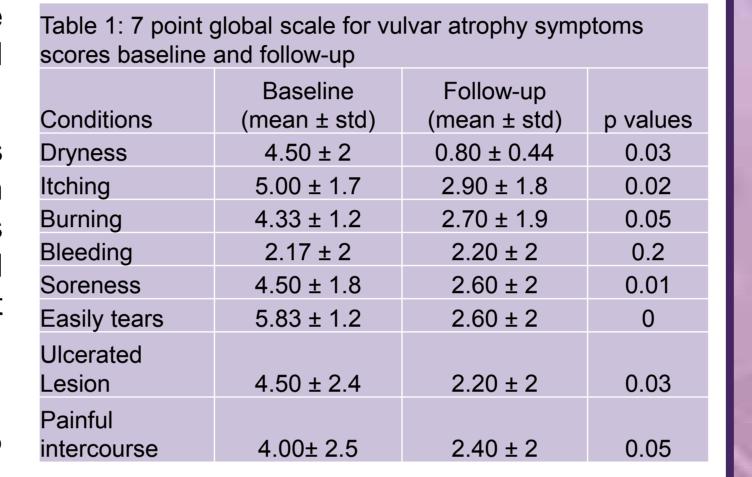
- Subjects received three treatments scheduled 4 weeks apart
- Treatment parameter based on depth of disease - depth 250-850um, 11% coverage, 1-2 passes
- Treatment numeric pain scores (0-10) and complications were recorded
- Follow-up was performed at one, three and six months post third treatment
- Paired t-test was used for data analysis

Results

Fifteen subjects ages 63±9.6 years were treated at a single center. All subjects had mild to moderate symptoms of VLS.

This poster reports finding of seven subjects at one and three months follow-up. Median numeric pain scores with the treatment was 5.7±2.3. No complications occurred. All conditions measured demonstrated significant improvement over baseline. Refer to figures 1 and 2, tables 1-4.

Three month biopsy data showed 50-100% improvement over baseline in VLS symptoms.



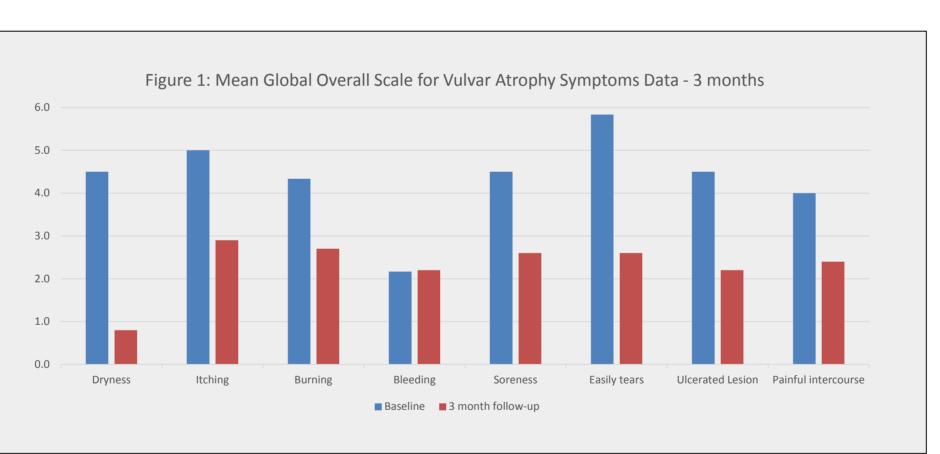
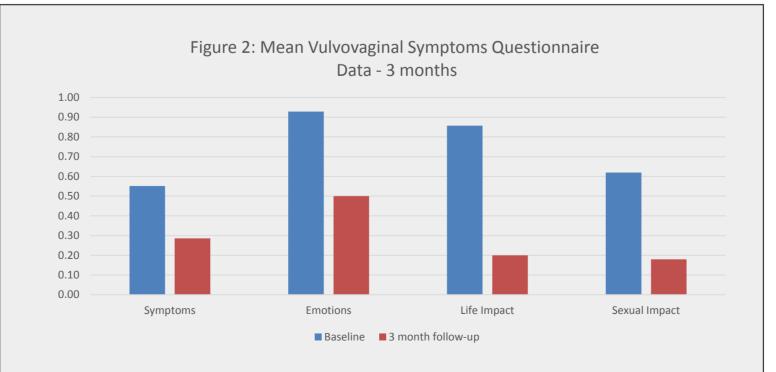


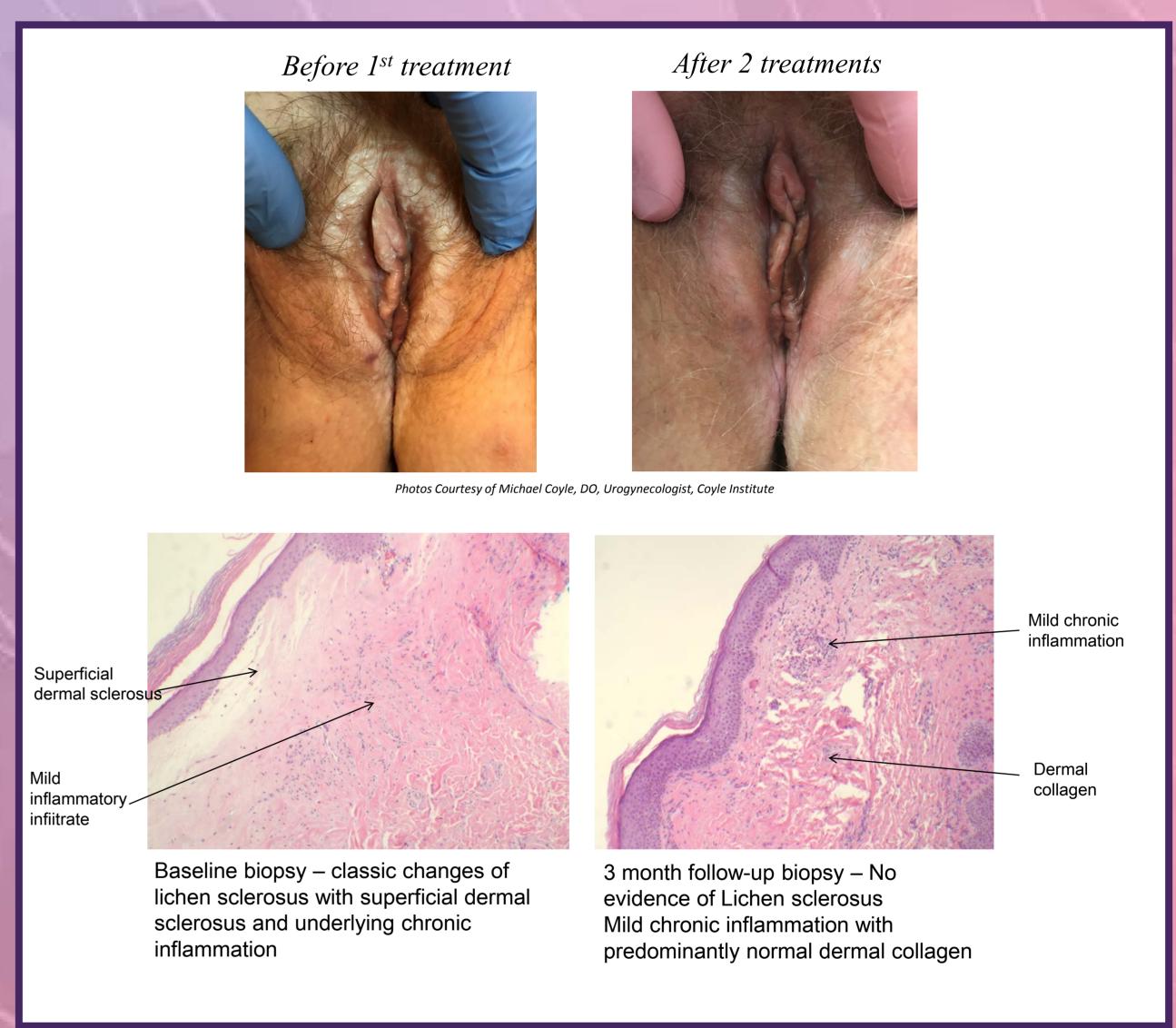
Table 2: 7 poir	•				
vulvar atrophy percent improvement					
	1 month	3 months			
Conditions	follow-up	follow-up			
Dryness	48%	82%			
Itching	46%	42%			
Burning	53%	37%			
Bleeding	47%	39%			
Soreness	57%	43%			
Easily tears	60%	54%			
Ulcerated					
Lesion	50%	54%			
Painful					
intercourse	53%	47%			



VSQ percent improvement		
1 month follow-up	3 months follow-up	
56%	48%	
61%	42%	
60%	72%	
74%	71%	
	1 month follow-up 56% 61% 60%	

Table 4: VSQ scores baseline and follow-up

	Condition	Baseline (mean ± std)	Follow-up (mean ± std)	p value
	Symptoms	0.55 ± 0.45	0.29 ± 0.25	0.06
	Emotions	0.93 ± 0.1	0.51 ± 0.26	0.008
	Life Impact	0.86 ± 0.14	0.2 ± 0.08	0.007
	Sexual Impact	0.62 ± 0.22	0.18 ± 0.1	0.07



Conclusion

The data indicates that the ablative fractional 2940nm Er:YAG laser appears to be a safe and effective treatment method for VLS. Additional follow-up data and comparative study with larger sample size are needed.

Acknowledgements

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- 3. Shahriari M, Makkar H, Finch J. Laser therapy in dermatology. *Clin Dermatol* 2015;33(6):681-6