

Enhanced Uptake of 10% Ascorbic Acid After 1440-nm or 1927-nm Non-ablative Fractional Diode Laser Treatment

OBJECTIVE

- To quantify uptake of 10% ascorbic acid following pretreatment with low-power 1440-nm or 1927-nm non-ablative fractional diode lasers (Clear + Brilliant® laser system; Solta Medical, Bothell, WA) with varying treatment densities

CONCLUSIONS

- In this ex vivo analysis, the greatest enhancement of 10% ascorbic acid uptake was seen with 1927-nm pretreatment at 320 MTZ/cm² and 1.0 W, compared to 1440-nm wavelengths at varying wattage and treatment densities
- This provides a foundation for clinical studies on laser-enhanced uptake of ascorbic acid and other topicals, which can allow clinicians to better understand the relationship between quantifiable uptake enhancement and patient-centered outcomes

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SYNOPSIS

- The stratum corneum forms a vital protective barrier along the outer layer of the skin, but also prevents optimal uptake of topical formulations¹
- Lasers can facilitate better penetration and absorption of topicals by disrupting the stratum corneum and tight junctions in the epidermis²
- Non-ablative lasers generally target dermal tissue and largely spare the stratum corneum, which minimizes overall thermal side effects and postprocedural recovery time, while fractionation further reduces postprocedural downtime^{3,4}
- The relationship between topical uptake and laser device settings, such as wavelength, peak power, and spot density, must be quantified to optimize treatment benefits

METHODS

- Excised human abdominal skin samples of 500-µm thickness were pretreated with a 1440-nm laser with 80 microscopic treatment zones (MTZ)/cm² (1.2 W), 1440-nm laser with 320 MTZ/cm² (3 W), 1927-nm laser with 320 MTZ/cm² (1 W), or received no pretreatment (Table 1)

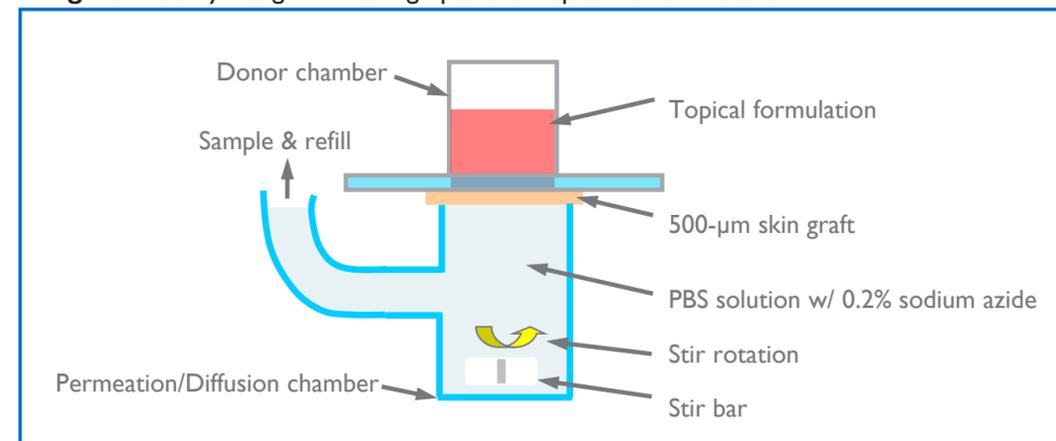
Table 1. Experimental Parameters

Parameter	Setting		
Device wavelength, nm	1440	1440	1927
Spot density, MTZ/cm ²	80	320	320
Peak power, W	1.2	3	1
Spot size, µm	130	130	130
Pulse energy, mJ	9	9	9

MTZ, microscopic treatment zones.

- Following laser pretreatment, 10% ascorbic acid (Obagi®, Long Beach, CA; 2010 formulation) was applied, and permeation was measured up to 24 hours after application (Figure 1)
- Samples were filtered and analyzed using high-performance liquid chromatography to measure topical permeation and retention for laser-treated samples and untreated controls
- Total uptake was calculated as the sum of the normalized cumulative permeation and retention in each sample
- Average total uptake was compared between laser-treated samples and untreated controls to determine the uptake enhancement ratio

Figure 1. Study design for testing uptake of topicals on skin tissue.



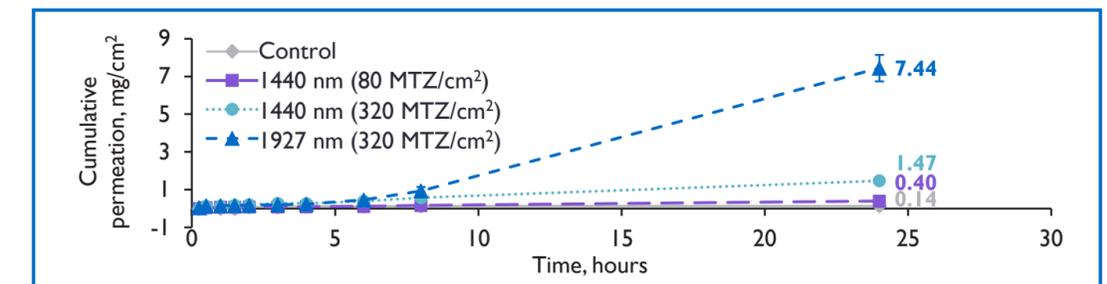
PBS, phosphate-buffered saline.

RESULTS

Permeation

- Pretreatment with the 1927-nm laser with 320 MTZ/cm² enhanced permeation of 10% ascorbic acid at 24 hours posttreatment relative to other pretreatments and untreated control (Figure 2)

Figure 2. Cumulative permeation of 10% ascorbic acid after laser pretreatment.



Values are mean ± standard deviation. MTZ, microscopic treatment zones.

Uptake

- Pretreatment with the 1927-nm laser with 320 MTZ/cm² enhanced uptake by >4 times compared to the 1440-nm laser with 320 MTZ/cm² (7.8 vs 1.8 mg/cm²; Table 2)
 - Compared to the 1440-nm laser with 80 MTZ/cm² (0.5 mg/cm²), uptake was enhanced by >15 times
 - Compared to untreated control (0.2 mg/cm²), uptake was enhanced by >33 times
- Pretreatment with the 1440-nm laser with 320 MTZ/cm² was associated with
 - >3-times greater uptake compared to the 1440-nm laser with 80 MTZ/cm² (1.8 vs 0.5 mg/cm²)
 - >7-times greater uptake compared to untreated control (1.8 vs 0.2 mg/cm²)

Table 2. Uptake Ratios of 10% Ascorbic Acid

	1440 nm (80 MTZ/cm ²)	1440 nm (320 MTZ/cm ²)	1927 nm (320 MTZ/cm ²)
Control	2.18 ± 0.55 x	7.75 ± 2.02 x	33.61 ± 0.02 x
1440 nm (80 MTZ/cm ²)	—	3.56 ± 0.70 x	15.45 ± 0.02 x
1440 nm (320 MTZ/cm ²)	—	—	4.34 ± 0.07 x

Values are mean ± root mean square. MTZ, microscopic treatment zones.

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Disclosures: JWV is an investigator for Solta Medical. PMF serves on the advisory board and speaker bureau for Solta Medical. AK and CP are employees of and may hold stock or stock options in Solta Medical. RGG is an investigator and advisory board member for Solta Medical.

References: 1. Lee et al. *Eur J Pharm Sci.* 2016;92:1-10. 2. Machado et al. *Aesthetic Plast Surg.* 2021;45:1020-1032. 3. Friedman et al. *J Drugs Dermatol.* 2020;19:s3-s11. 4. Farkas et al. *Aesthet Surg J.* 2013;33:1059-1064.