Quantitative Analysis of Trans-Epithelial Corneal Riboflavin Loading

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Affiliations

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Dr. Talamo: Massachusetts Eye and Ear Infirmary and Harvard Medical School, Boston, MA
# Financial Interests

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<tr>
<th>R. Rubinfeld</th>
<th>J. Talamo</th>
<th>D. Stulting</th>
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<td>Aura Biosciences</td>
<td>Alcon Laboratories</td>
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<td>CXLUSA, LLC</td>
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Background

• Dresden protocol effective, epi removal risks
• Advantages of less invasive transepi CXL
  • Return to preop vision, function POD 1
  • Log scale shift in safety and risk/benefit
  • Treat upon diagnosis
Perforation after Epi-Off CXL
Transepithelial corneal collagen crosslinking for progressive keratoconus: 24-month clinical results

Aldo Caporossi, MD, FRCS, Cosimo Mazzotta, MD, PhD, Anna Lucia Paradiso, MD, Stefano Baiocchi, MD, PhD, Davide Marigliani, MD, Tomaso Caporossi, MD

PURPOSE: To assess the clinical results of transepithelial collagen crosslinking (CXL) in patients 26 years and younger with progressive keratoconus suitable for epithelium-off (epi-off) CXL.

SETTING: Department of Ophthalmology, Siena University Hospital, Siena, Italy.

DESIGN: Prospective case series.

METHODS: The study included 26 eyes (26 patients) treated by transepithelial (epithelium-on) CXL. The mean age was 22 years (range 11 to 26 years), 10 younger than 18 years, 16 between 19 years and 26 years. Preoperative and postoperative examinations included uncorrected (UDVA) and corrected (CDVA) distance visual acuities, simulated maximum keratometry (K), coma and spherical aberration, and corneal optical coherence tomography optical pachymetry. The solution for transepithelial CXL (Ricorin T) comprised riboflavin 0.1%, dextran 15.0%, trometamol (Tris), and ethylenediaminetetraacetic acid. Ultraviolet-A treatment was performed with the Caporossi Baiocchi Mazzotta X Linker Vega at 3 mW/cm².

RESULTS: After relative improvement in the first 3 to 6 months, the UDVA and CDVA gradually returned to baseline preoperative values. After 12 months of stability, the simulated maximum K value worsened at 24 months. Coma aberration showed no statistically significant change. Spherical aberration increased at 24 months. Pachymetry showed a progressive, statistically significant decrease at 24 months. Fifty percent of pediatric patients were retreated with epi-off CXL due to significant deterioration of all parameters after 12 months of follow-up.

CONCLUSIONS: Functional results after transepithelial CXL showed keratoconus instability, in particular in pediatric patients 18 years old and younger; there was also functional regression in patients between 19 years and 26 years old after 24 months of follow-up.

Financial Disclosure: No author has a financial or proprietary interest in any material or method mentioned.

CXL Photochemistry

UVA Light
Energy Source

Oxygen
Rate Limiting Reagent

Riboflavin
Energy Transfer

Amide

Aldehyde
Epi-On Success Requires

1. **Stroma**: Reliable, consistent riboflavin loading
2. **Epithelium**: Good UVA transmission, relatively clear of riboflavin, intact, non-edematous,
3. **Oxygen**: Rate limiting reagent if 1,2 present
Clear Epithelium, Loaded Stroma

No Visible Green Color in Epithelium

Well Loaded Green Stroma
Purpose

• Evaluate corneal penetration of riboflavin using a novel, proprietary, patent pending formulation and delivery system
• Compare results to those using a commercially available product
• Correlate slit lamp findings with quantitative measurements of corneal riboflavin concentration
Materials

- *Absorption Systems, Inc.*, San Diego California
- 3-4.4 kg New Zealand white rabbits
- **Group 1**: ParaCel™ q90 sec x 4 min., then VibeX Xtra™ q90 sec x 6 min. (per label)
- **Group 2**: Proprietary prep x 30-60 sec. then CXLUSA/CXLO riboflavin² via sponge with drops q1-3 min x 10 min.

¹Trans-epithelial Cross-Linking Kit, Avedro, Inc.
²Proprietary system from CXL Ophthalmics utilized by CXLUSA investigators under a physician-sponsored IRB approved study.
Methods

• Slit lamp photos at baseline
• Riboflavin application
• Masked grading and slit lamp photos at 10, 15, and 20 min. (2X labeling)
• Euthanasia, epithelial removal, and riboflavin assay by liquid chromatography/tandem mass spectrometry (LC-MS/MS) 22-25 min. after application
# Slit Lamp Grading System

<table>
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<tr>
<th>Grade</th>
<th>Findings</th>
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<tr>
<td>I</td>
<td>Mild green tint just visible</td>
</tr>
<tr>
<td>II</td>
<td>Substantial green visible</td>
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<tr>
<td>III</td>
<td>Obvious green color</td>
</tr>
<tr>
<td>IV</td>
<td>Bright green appearance</td>
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<tr>
<td>V</td>
<td>Strong, bright green color</td>
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Ex Vivo Rabbit Cornea SL Photos

Paracel™-VibeX Xtra™
At 20 min

CXLUSA Formulation
At 10 Min
Ex Vivo Rabbit Cornea SL Photos

Paracel™-VibeX Xtra™
At 20 min

CXLUSA Formulation
At 10 Min
Riboflavin Concentration
(μg/g at 20-25 min.)

Paracel™-VibeX Xtra™

CXLUSA

Rabbit

Riboflavin Concentration (μg/g)

1 2 3

0 5 10 15 20 25 30 35

Rabbit

Riboflavin Concentration (μg/g)

0 5 10 15 20 25 30 35

1 2 3 4
Correlation Between SLE and LC-MS/MS

Riboflavin Conc. μg/g vs. Slit Lamp Grade

The graph shows a positive correlation between riboflavin concentration (μg/g) and slit lamp grade. As the slit lamp grade increases, the riboflavin concentration also tends to increase.
15 Min Epi-On Human CXLUSA
Conclusions

• CXLUSA Riboflavin system produces a 4.0-fold greater corneal stromal concentration in the rabbit than commercially available system

• This correlates well with the 4.3-fold greater concentration of stromal riboflavin produced in the human by the epi-off technique [Gore et al., 2015; IOVS 56:5006] compared to the same commercially available system used in this study
Conclusions

- Failure of previous epi-on protocols to halt progression of ectatic corneal disease may, in part, be due to inadequate corneal riboflavin concentration.
- Slit lamp exam provides a valid estimate of stromal riboflavin concentration that can be used to assure adequate corneal loading prior to UVA exposure.
Conclusions

• The CXL technique used by CXLUSA may achieve corneal stiffening similar to that achieved by the classical Dresden epi-off technique without the risk of epithelial removal.

• Early 2-year clinical data suggest this is a valid hypothesis.