

Long-Term Obeticholic Acid Treatment is Associated With Improvements in Collagen Morphometry in Patients With Primary Biliary Cholangitis

OCA

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Introduction

- Primary biliary cholangitis (PBC) is a rare autoimmune liver disease of the intrahepatic bile ducts, leading to progressive fibrosis and eventual cirrhosis¹
- In patients with PBC, cirrhosis-related events and clinical outcomes have been associated with the fibrosis stage²⁻⁴
- Measuring collagen content is emerging as a reliable method of quantifying liver fibrosis⁵ and has shown evidence of being an effective tool in patients with PBC⁶
- Second harmonic generation (SHG) microscopy is a new tissue imaging technology that allows the accurate quantification of several collagen parameters on unstained tissue sections⁷
- Obeticholic acid (OCA) is a selective, potent farnesoid X receptor (FXR) agonist approved as a second line therapy in patients with PBC and an inadequate response to or intolerance of ursodeoxycholic acid (UDCA)⁸
- Approval is based on a surrogate endpoint of biochemistry (alkaline) phosphatase [ALP] and bilirubin)

Objective

 This post-hoc analysis assessed the impact of 3 years of OCA treatment on collagen morphometry using biopsy samples from the PBC OCA International Study of Efficacy (POISE) study

Methods If on UDCA: Continue UDCA Placebo ± UDCA (n=73) All patients initiated OCA 5 mg for 3 months, after which patients had the OCA 5-10 mg ± UDCA (n=70) option to titrate up based on tolerability OCA 10 mg ± UDCA (n=73) **Open-Label Extension** † (Visits Every 3 Months, Up to 5 Years)

STUDY DESIGN:

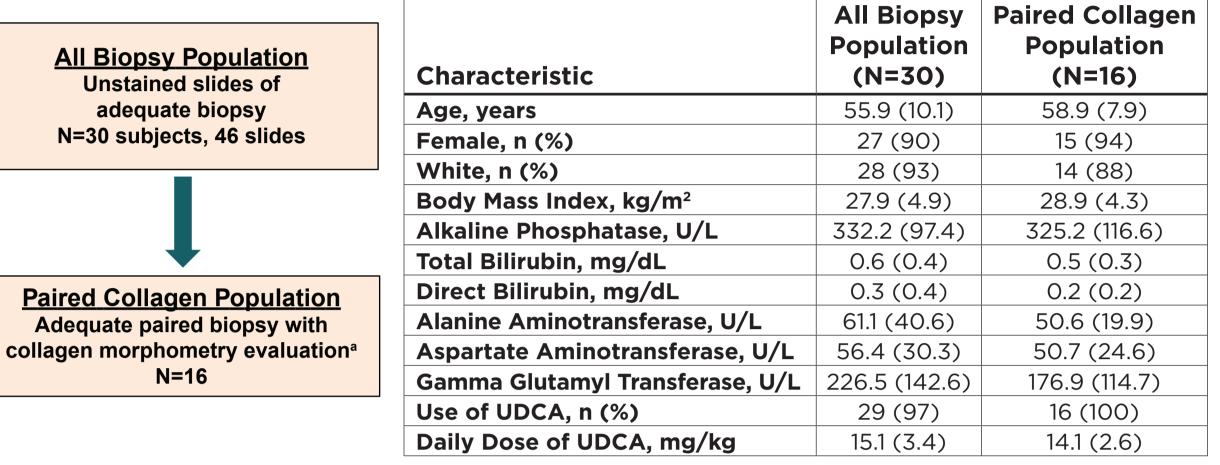
- POISE was a randomized, double-blind, placebo-controlled, pivotal Phase 3 study evaluating OCA treatment in 216 patients with PBC through a 12-month double-blind (DB) phase and 5-year open-label extension (OLE) phase⁸
- POISE included a prespecified substudy that evaluated biopsy samples at baseline (up to 1 year prior to the start of the DB phase) and after approximately 3 years of OCA treatment

POST-HOC ANALYSIS:

- For patients that had paired evaluations (both baseline and on-treatment) biopsies underwent:
- Nakanuma scoring⁹ by 2 liver pathologists in a consensual reading, blinded to randomization and timing of biopsies
- Collagen quantification by second harmonic generation (SHG) and 2-photon excitation (2PE) microscopy on unstained slides
- Collagen area ratio (CAR): area of collagen (collagen pixel count) / total area of region of interest (total pixel count) • Collagen fiber density (CFD): total "brightness" of collagen (intensity) / collagen surface area
- Collagen reticulation index (CRI): measure of complexity of collagen network (collagen branch points / collagen length)
- Fibrosis composite score (FCS): composed of 15 unique morphometric parameters

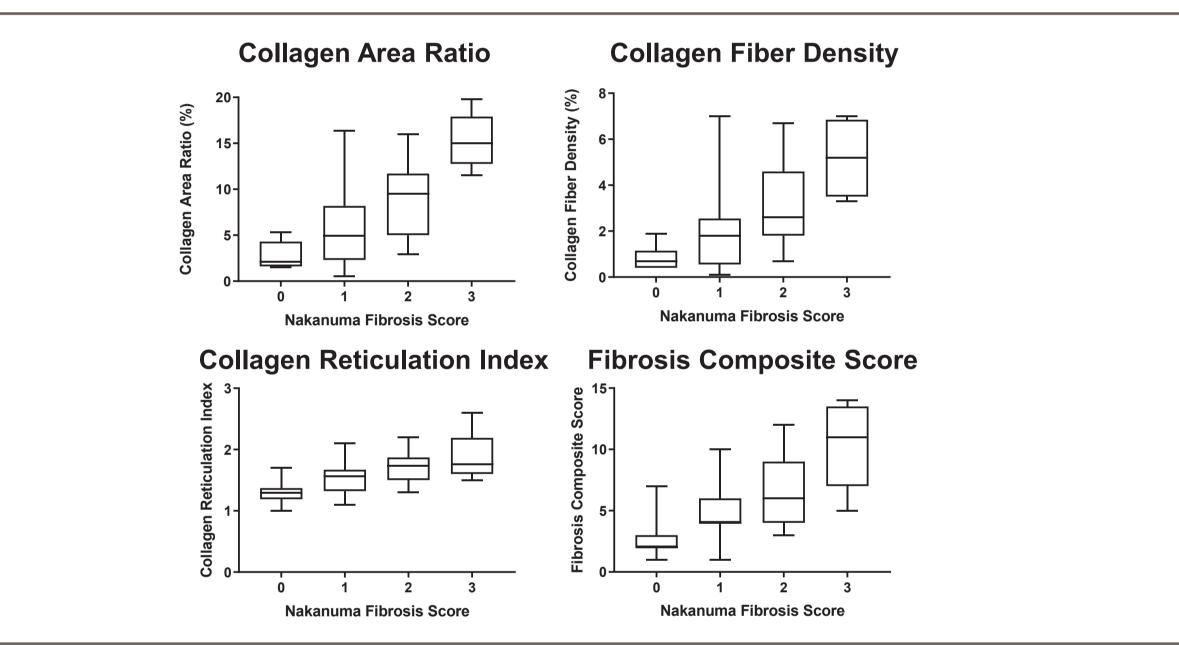
Results

Table 1. Baseline characteristics



Data are mean (standard deviation) unless otherwise indicated ^a17 patients had adequate paired biopsies used for evaluation of Nakanuma score: 16 of these patients had adequate paired biopsies and unstained slides for evaluation of collagen by second harmonic generation/2-photon excitation microscopy. OCA, obeticholic acid; UDCA, ursodeoxycholic acid.

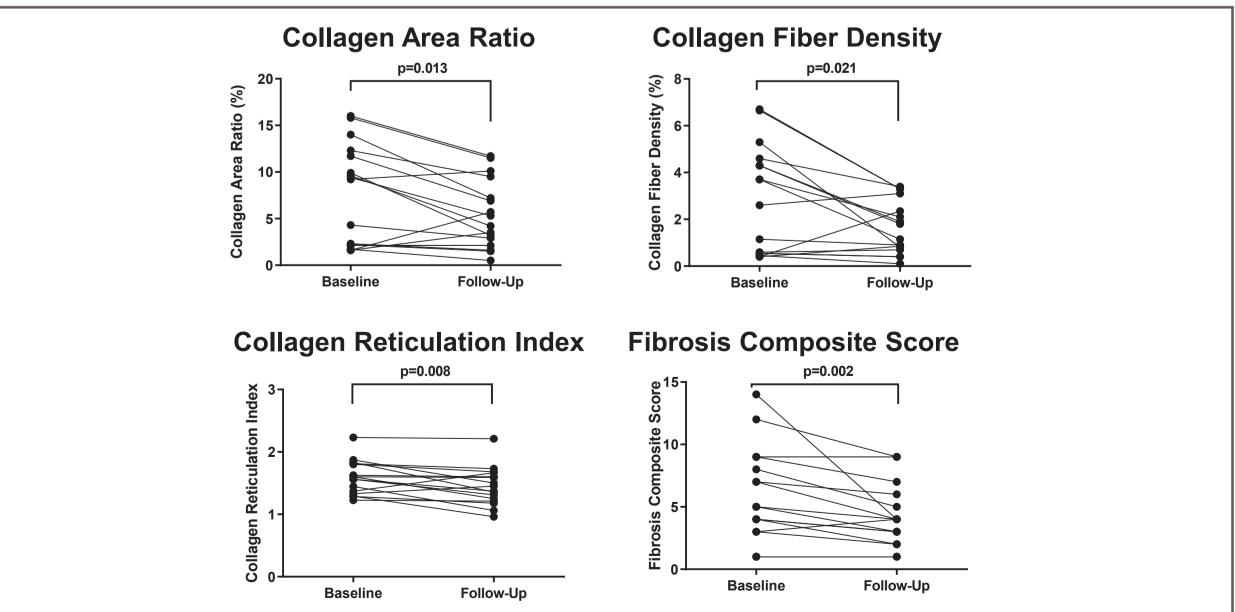
Figure 1. Collagen Morphometry vs Nakanuma Fibrosis Score (N=30 subjects, 46 slides)



Line within box represents median: top and bottom of box represent IQR: bars represent min and max.

 In the all biopsy population (N=30), the CAR, CFD, CRI, and FCS all increased in parallel with the Nakanuma fibrosis score

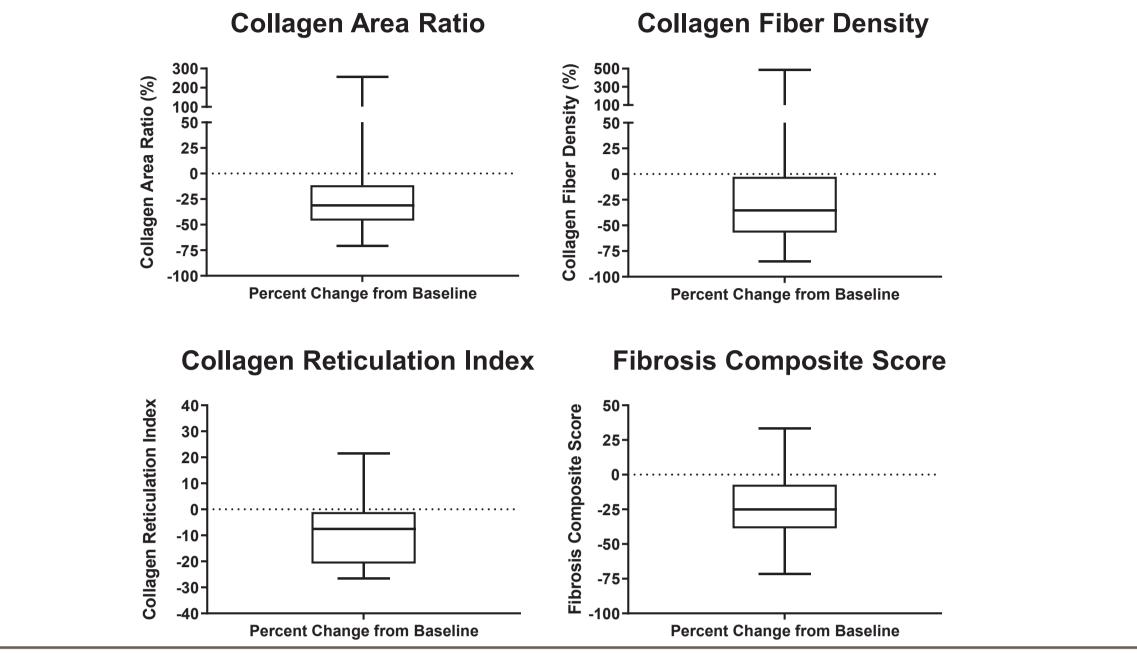
Figure 2. Individual Patient Collagen Morphometry From Baseline to Follow-Up (N=16)



p-value obtained using Wilcoxon Signed Rank test.

 OCA treatment resulted in significant reductions from baseline in the median (Q1, Q3) CAR (-2.1 [-4.6, -0.3], p=0.013), CFD (-0.8 [-2.5, 0.0], p=0.021), CRI (-0.1 [-0.3, 0.0], p=0.008), and FCS (-1.0 [-2.5, -0.5], p=0.002)

Figure 3. Percent Change From Baseline in Collagen Morphometry and the Fibrosis Composite Score (N=16)



- Reductions represent -31%, -35%, -7%, and -25% percent change from baseline in CAR, CFD, CRI, and FCS, respectively

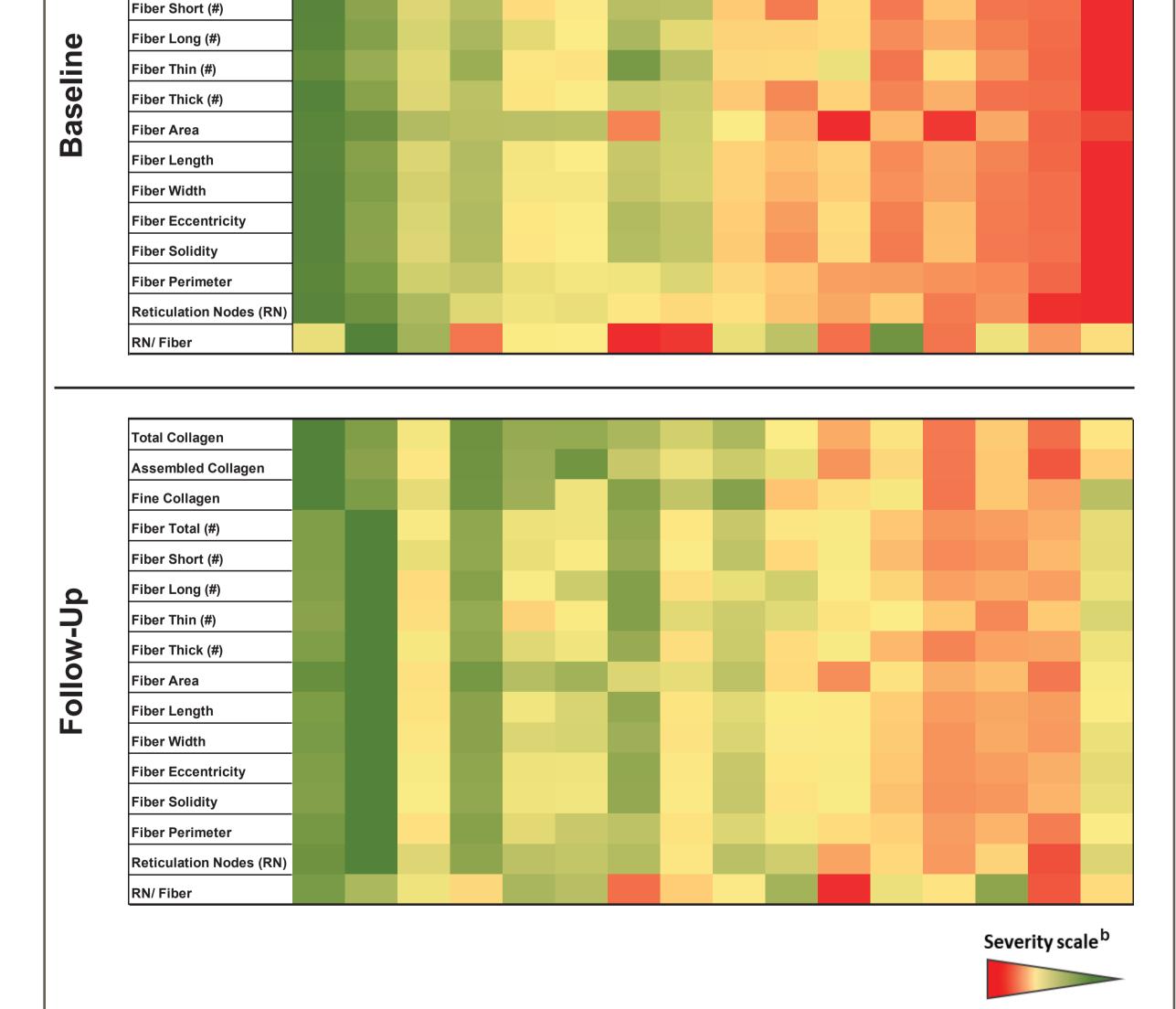
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16

Figure 4. Collagen Morphometry Heat Map (N=16)

ssembled Collagen

Fine Collagen

Fiber Total (#)



^aEach row corresponds to individual collagen morphometric parameters; each column corresponds to individual patient reads bColorimetric scale displays shades of colors ranging from red (most severe fibrosis) to yellow to green (least severe fibrosis).

 OCA treatment resulted in an improvement in most collagen parameters in most patients as observed qualitatively by a reduction in red and increase in green within the heat map at the follow-up biopsy relative to baseline

Table 2. Cumulative Safety Across 3 Years of OCA Treatment

Adverse Events	Total OCA N=16
Pruritus	11 (69)
Fatigue	8 (50)
Arthralgia	5 (31)
Upper respiratory tract infection	4 (25)
Diarrhea	4 (25)
Nasopharyngitis	4 (25)
Urinary tract infection	4 (25)
Pain in extremity	4 (25)
Influenza	4 (25)

Data are n (%). OCA, obeticholic acid.

 All serious adverse events were considered unlikely or not related to OCA

Conclusions

- The majority of patients with PBC receiving 3 years of OCA treatment in this study showed improvements or stabilization in collagen morphometry
- Significant reductions were observed in collagen area ratio, collagen fiber density, and collagen reticulation index as assessed by SHG/2PE microscopy
- Morphometric measures of fibrosis increased with increasing histologic disease severity as assessed by the Nakanuma fibrosis score, supporting the potential validity of collagen measurements by SHG/2PE
- The data from this POISE subgroup analysis support that in patients with an inadequate response to UDCA, 3 years of OCA treatment results in an improvement or stabilization in fibrosis progression

References

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Disclosure

AK - Personal fees from AbbVie, Beiersdorf, BMS, CymaBay, Gilead, GSK, Intercept Pharmaceuticals, MSD. Grants from Intercept Pharmaceuticals

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Dr. Andreas E. Kremer Clinical Trial Information: CT.gov: NCT01473524; Eudra CT: 2011-004728-36





