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Obeticholic acid for the treatment of non-alcoholic steatohepatitis: interim analysis from a multicentre, randomised, placebo-controlled phase 3 trial

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1 **Obeticholic Acid for the Treatment of Nonalcoholic Steatohepatitis—**
2 **Interim Analysis From a Randomised Phase 3 Study**
3

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SUMMARY

Background: Nonalcoholic steatohepatitis (NASH) is a common type of chronic liver disease that can lead to cirrhosis. Obeticholic acid (OCA), a farnesoid X receptor agonist, has been previously shown to improve the histologic features of NASH. Results of a planned interim analysis of an ongoing, randomised, double-blind, multicentre, placebo-controlled phase 3 global study of OCA for NASH are reported.

Methods: Patients with NASH, non-alcoholic fatty liver disease (NAFLD) activity score ≥ 4 and fibrosis stages F2–F3, and an exploratory cohort with fibrosis stage F1, were randomised to receive placebo, OCA 10-mg, or OCA 25-mg daily in a 1:1:1 ratio. The primary endpoints for the month 18 interim analysis were fibrosis improvement (≥ 1 stage) with no worsening of NASH, or NASH resolution with no worsening of fibrosis, with the study considered successful if either primary endpoint was met. The study also evaluated histologic and biochemical markers of NASH and fibrosis, and safety (NCT02548351; EudraCT 2015-002560-16).

Findings: The intent-to-treat population included 931 patients with stage F2–F3 fibrosis (placebo, n=311; OCA 10-mg, n=312; OCA 25-mg, n=308). Fibrosis improvement by ≥ 1 stage was met by 12% of placebo patients, 18% of OCA 10-mg patients, and 23% of OCA 25-mg patients ($p=0.0002$). Although NASH resolution was not met (placebo, 8%; OCA 10-mg, 11%; OCA 25-mg, 12%), more OCA-treated patients had absence of definite NASH at month 18 based on pathologist’s assessment. Improvements, including normalization, were noted in liver biochemistry. In the safety population (F1–F3, N=1968), the most common adverse event was pruritus (placebo, 19%; OCA 10-mg, 28%; OCA 25-mg, 51%). The overall safety profile was similar to that in previous studies, and incidence of serious adverse events was similar across treatment groups (11–14%).

1 **Interpretation:** Obeticholic acid 25-mg significantly improved fibrosis and NASH disease

2 activity among NASH patients with fibrosis.

3 **Funding:** Intercept Pharmaceuticals.

4

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1 The RandomizEd Global phase 3 Study to Evaluate the impact on NASH with fibRosis of
2 obeticholic Acid TreatmEnt (REGENERATE) study is an international, prospective, randomised,
3 double-blind, placebo-controlled phase 3 study of OCA in patients with NASH and fibrosis
4 (NCT02548351).¹¹ Here we report the final results of the prespecified month-18 interim analysis
5 on the safety and efficacy of OCA in improving fibrosis and underlying disease activity.

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METHODS

Study design and participants

This study is being conducted in 332 centres across 20 countries. Eligible patients were adults (aged ≥ 18 years) with histologic evidence (per central reading of a liver biopsy obtained within 6 months of randomisation) of steatohepatitis; a non-alcoholic fatty liver disease (NAFLD) activity score (NAS) ≥ 4 ; and fibrosis stage per the NASH CRN scoring criteria of F2 or F3, or F1 with ≥ 1 accompanying comorbidity (obesity, type 2 diabetes, or alanine amino transferase [ALT] > 1.5 times the upper limit of normal [ULN]). Patients were excluded if cirrhosis, other chronic liver disease, significant alcohol consumption (> 2 units/day for women or > 4 units/day for men for > 3 months within 1 year before screening), or confounding conditions were present. All patients provided written informed consent. Complete study design including inclusion and exclusion criteria were previously reported.¹¹

A planned interim analysis was to be performed after a minimum of 750 randomised patients with fibrosis stages F2 or F3 reached their actual/planned month-18 visit. The end-of-study analysis will evaluate the effect of OCA on liver-related clinical outcomes including progression to cirrhosis, all-cause mortality, and the long-term safety of OCA, and will be completed once approximately 291 adjudicated clinical outcome events occur in the OCA 25-mg and placebo groups combined in patients with fibrosis stage F2 or F3. Patients are expected to have a minimum follow-up time of approximately 4 years.

1 **Randomisation and blinding**

2 Eligible patients were randomised in a 1:1:1 ratio to receive daily placebo, OCA 10-mg, or OCA
3 25-mg orally. To determine eligibility at enrollment, two central pathologists were required to
4 confirm histological presence of NASH and fibrosis, and a NAS ≥ 4 with a score of at least 1 in
5 each component of NAS. Randomisation was performed using an Interactive Web Response
6 System (IWRS); for patients with fibrosis stage F2 or F3, randomization was stratified by both
7 the presence of type 2 diabetes at enrollment and the use of thiazolidinediones (TZD) or vitamin
8 E at baseline. Placebo and OCA were supplied as identical tablets in coded containers. All
9 patients, study investigators, and other site research staff were blinded to treatment assignment.

10

11 **Procedures and assessments**

12 Biopsies were obtained at screening, and month 18/end-of-treatment. Histologic assessments
13 followed standardised criteria to ensure consistency, and all biopsies were read centrally. The
14 month 18 or early termination biopsy slides were read with the screening biopsy slides by one of
15 the two pathologists who were blinded to the slide sequence and the patient's treatment.

16 Assessments of liver biochemistry were performed at each study visit. Safety and tolerability of
17 OCA were assessed by analysis of adverse events (AEs), vital signs, electrocardiograms, and
18 clinical laboratory assessments (including lipid profile changes). An independent data and safety
19 monitoring committee reviewed, and continues to review, safety and efficacy during the study.

20

21 **Endpoints**

22 REGENERATE was designed to assess liver histology at month 18 as a surrogate endpoint for
23 clinical outcomes.¹¹ The primary endpoints were defined as improvement in fibrosis (reduction

1 of ≥ 1 stage) with no worsening of NASH (defined as no increase of hepatocellular ballooning,
2 lobular inflammation, or steatosis), or NASH resolution (defined as the overall histopathologic
3 interpretation of “no fatty liver disease” or “fatty liver disease without steatohepatitis” and a
4 NAS of grade 0 for ballooning and 0–1 for inflammation) with no worsening of fibrosis. The key
5 secondary endpoint was improvement of fibrosis by ≥ 1 stage without worsening of NASH and/or
6 resolution of NASH, without worsening of fibrosis. Secondary endpoints also included
7 evaluation of the effect of OCA versus placebo on histologic improvement of features of NASH
8 as well as NAS, liver biochemistry, and markers of liver function.¹¹ A post hoc analysis
9 evaluated NASH resolution based on the pathologist diagnostic assessment of presence/absence
10 of definite steatohepatitis as determined by the overall pattern of injury rather than scoring of
11 individual NAS parameters.

12

13 **Statistical analyses**

14 For the month 18 primary efficacy endpoint of improvement in fibrosis with no worsening of
15 NASH, a sample size of 250 per group with an assumed 15% discontinuation rate will provide
16 98% power to demonstrate a statistically significant treatment difference between the OCA (10-
17 mg and 25-mg) and placebo groups based on the Cochran-Mantel-Haenszel test with a two-sided
18 type I error at the 0.01 level, assuming an adjusted response rate of 36.7% and 17.6% in the
19 OCA (10-mg and 25-mg) and placebo groups, respectively. Inferential testing was performed
20 sequentially in the dose level, adjusting for multiplicity using truncated Hochberg procedure, to
21 test the two primary endpoints within each dose level, starting by comparing the OCA 25-mg
22 group with placebo for the two primary endpoints, then comparing the OCA 10-mg group with
23 placebo in the intent-to-treat (ITT) population. All other testing and the associated p values

1 presented in this article are not controlled for type II error. Success of the study was defined as
2 meeting one of the two primary endpoints at the predetermined significance level. The analyses
3 of histologic endpoints at the month-18 interim analysis were performed with an alpha level of
4 0.02. The statistical analysis plan and primary endpoints were agreed with the FDA prior to
5 study initiation.

6 As shown in figure 1, all patients (fibrosis stages F1-F3) who received ≥ 1 dose of study
7 treatment by the pre-specified month-18 interim analysis cutoff date were included in the safety
8 population, which was used for all safety and tolerability analyses. The primary analysis
9 population for efficacy endpoints was the ITT population, comprised of patients with more
10 advanced disease (fibrosis stage F2-F3) who had received ≥ 1 dose of treatment and reached or
11 would have reached the month-18 visit by the pre-specified interim analysis cutoff date. Efficacy
12 endpoints were also analysed in the per-protocol population, defined as the ITT population who
13 completed ≥ 15 months of treatment, had a month-18/end of treatment biopsy, were on treatment
14 ≥ 30 days immediately preceding biopsy, and did not have any major protocol deviation

15

16 **Role of the funding source**

17 The REGENERATE study was designed by VR, AJS, and ZMY in collaboration with Intercept
18 Pharmaceuticals; operational and protocol-specific aspects of the study were supervised by a
19 steering committee comprising AJS, MR, PB, QMA, RL, SH, VR, ZG, and ZMY (chair). All
20 authors vouch for the fidelity of the study to the protocol, the accuracy and completeness of the
21 data, and approved publication of the manuscript. The first and corresponding authors had full
22 access to all the data in the study and had final responsibility for the decision to submit for

1 publication. Funding for the study was provided by Intercept Pharmaceuticals. No funding was
2 provided to any author for writing and editing of the manuscript.

3

4

RESULTS

1
2 Between November 2015 and October 2018, a total of 1968 patients were enrolled and randomly
3 assigned to one of the three treatment groups (figure 1). The ITT population included 931
4 patients randomised to receive placebo (n=311), OCA 10-mg (n=312), or OCA 25-mg (n=308).
5 At the time of the interim analysis, 23% of placebo patients, 23% of OCA 10-mg patients, and
6 25% of OCA 25-mg patients had discontinued treatment (figure 1); 81% of patients receiving
7 placebo or OCA 10-mg and 79% receiving OCA 25-mg completed the month 18 biopsy. An
8 additional 3% of patients in each treatment group completed any postbaseline biopsy (patients
9 who discontinued the treatment before month 18 and underwent an end-of-treatment biopsy).
10 The per-protocol population included 668 patients (placebo, n=224; OCA 10-mg, n=226; OCA
11 25-mg, n=218) and the safety population included 1968 patients (placebo, n=657; OCA 10-mg,
12 n=653; OCA 25-mg, n=658).
13 In the ITT population, baseline characteristics were balanced across treatment groups and
14 reflective of a noncirrhotic NASH population (table 1). A majority of patients had stage F3
15 fibrosis (54–58%) and NAS ≥ 6 (68–70%) indicative of advanced fibrosis and high disease
16 activity. Consistent with NASH epidemiology, more than half of the patients had type 2 diabetes
17 (55–56%), and 52-54% overall were receiving antidiabetic medication at baseline. Additionally,
18 41–46% of patients were receiving statin therapy and a minority were receiving NASH-
19 modifying agents, TZD (1-3%) and Vitamin E (10-14%). A similar pattern of baseline
20 characteristics was observed in the per-protocol population (table S1).
21 The primary endpoint of fibrosis improvement by ≥ 1 stage with no worsening of NASH was met
22 by 12% of placebo patients, 18% of OCA 10-mg patients (p=0.04 vs placebo) and 23% of OCA
23 25-mg patients (p=0.0002 vs placebo) with an OCA:placebo response ratio (95% confidence

1 interval [CI]) of 1.48(1.01, 2.18) and 1.94(1.35, 2.78) for OCA 10-mg and OCA 25-mg,
2 respectively (figure 2, table 2). OCA 25-mg was statistically significant per the inferential testing
3 method pre-specified in the statistical analysis plan. Similar results were observed in the per-
4 protocol population (placebo 13%, OCA 10-mg 21% [p=0.02], OCA 25-mg 28% [p<0.0001])
5 (figure 2, table 2). Across subgroups of interest in the ITT population, a ≥ 1 stage improvement in
6 fibrosis was consistently observed in the OCA 25-mg group (fibrosis stage F2 or F3 [p=0.006
7 and p=0.02]; NAS ≥ 6 [p=0.0003]; presence or absence of type 2 diabetes [p=0.02 and
8 p=0.005]). The NAS <6 subgroup did not include enough patients for a meaningful comparison
9 (figure S1).

10 In the per-protocol population, which includes patients with ≥ 15 months of treatment, three times
11 as many patients in the OCA 25-mg group achieved ≥ 1 stage improvement in fibrosis (38%) as
12 opposed to progression of fibrosis (13%) compared to the placebo group, which showed a similar
13 number of patients that improved (23%) or worsened (21%) (figure 3).

14 The primary endpoint of NASH resolution based on no hepatocellular ballooning and no or
15 residual lobular inflammation with no worsening of fibrosis did not meet statistical significance
16 in the ITT population (placebo 8%, OCA 10-mg 11% [p=0.18], OCA 25-mg 12% [p=0.13]) with
17 an OCA:placebo response ratio (95% CI) of 1.39(0.86, 2.25) and 1.45(0.90, 2.35) for OCA 10-
18 mg and OCA 25-mg, respectively (figure 2, table 2). Similar results were observed in the per-
19 protocol population (figure 2, table 2). Despite not meeting NASH resolution, a dose-dependent
20 response was observed in the ITT population with more OCA 25-mg patients compared to
21 placebo achieving ≥ 1 -point improvement in scores of lobular inflammation (44% vs 36%,
22 p=0.03) and hepatocellular ballooning (35% vs 23%, p=0.001), key features of NASH (figure
23 S2).

1 In a post hoc analysis, NASH resolution was evaluated by assessing a change in diagnosis from
2 presence of definite steatohepatitis at baseline to absence of definite steatohepatitis (without
3 worsening of fibrosis) at month 18. This pathologist diagnostic assessment of NASH, based on
4 the overall pattern of liver injury, showed that in the ITT population approximately twice as
5 many patients in the OCA 25-mg group achieved NASH resolution compared with the placebo
6 group (23% vs 12%, $p=0.0004$) (figure S3). A similar dose-dependent response was observed in
7 the per-protocol population (29% vs 16%, $p=0.0005$) (figure S3).

8 The key secondary endpoint of improvement of fibrosis by ≥ 1 stage and/or resolution of NASH,
9 without worsening of either fibrosis or NASH was achieved by 16% of placebo, 22% of OCA
10 10-mg ($p=0.07$) and 27% of OCA 25-mg patients ($p=0.0005$) (ITT population) (table 2, figure
11 S4). A significantly higher proportion of patients receiving OCA 25-mg compared to placebo
12 achieved improvement in NAS by ≥ 2 -points with no worsening of fibrosis (36% vs 24%,
13 $p=0.001$), had no disease progression as assessed by no worsening of fibrosis and no worsening
14 of NASH (48% vs 38%, $p=0.011$), and had improvement in fibrosis by ≥ 2 stages (10% vs 5%,
15 $p=0.018$) (table 2). Results of additional secondary NASH and fibrosis endpoints are provided in
16 table 2.

17 Favourable changes in key liver enzyme levels were observed in patients who received OCA.

18 Early dose-dependent decreases in ALT and aspartate aminotransferase (AST) were observed by
19 month 3 and continued through month 18 (mean [standard error (SE)] change at month-18 ALT:
20 placebo -15.6 [3.3] U/L, OCA 10-mg -23.8 [2.6] U/L, OCA 25-mg -36.0 [3.6] U/L; AST:
21 placebo -9.8 [2.4] U/L, OCA 10-mg -14.1 [2.1] U/L, OCA 25-mg -20.4 [2.3] U/L) (figure 4).

22 These changes correspond to a decrease in ALT of 6% for placebo, 26% for OCA 10-mg, and
23 33% for OCA 25-mg and in AST of 4%, 19%, and 24%, for placebo, OCA 10-mg, and OCA 25-

1 mg, respectively (figure 4). A post hoc analysis demonstrated that a higher proportion of patients
2 receiving OCA with elevated ALT and AST at baseline achieved levels below the ULN at month
3 18 compared with placebo (figure S5). Gamma-glutamyl transferase (GGT) levels dropped
4 rapidly and were generally stable after month 3 (change at month 18: placebo 1%, OCA 10-mg
5 -24%, OCA 25-mg -38%) (figure 4). Slight increases in alkaline phosphatase (ALP) were
6 observed with OCA treatment, but levels remained below ULN through month 18 (change at
7 month 18: placebo -1%, OCA 10-mg 9%, OCA 25-mg 20%) (figure 4).

8 Additionally, patients receiving OCA versus placebo had a higher, dose-dependent decrease in
9 body weight throughout the 18-month observation period (mean [SE] change: placebo, -0.7
10 [0.4] kg; OCA 10-mg, -1.8 [0.4] kg; OCA 25-mg -2.2 [0.3] kg).

11 A total of 1968 patients were included in the safety analysis, comprised of 15% with fibrosis
12 stage F1 (15%), stage F2 (35%), and stage F3 (50%). The duration of exposure was generally
13 similar across treatment groups. Overall, treatment-emergent AEs occurred in 83% of placebo,
14 89% of OCA 10-mg, and 91% of OCA 25-mg patients; most (69–74%) were mild to moderate in
15 severity (table S2). The frequency of serious AEs (SAEs) was similar across treatment groups
16 (11–14%) and no single SAE occurred in >1% of patients in any treatment group (table S2). The
17 most frequent AE was pruritus (placebo, 19%; OCA 10-mg, 28%; OCA 25-mg, 51%) (table 3).

18 The incidence of pruritus was highest during the first 3 months of treatment with OCA, and
19 generally mild to moderate in severity. Treatment discontinuation due to pruritus occurred in five
20 placebo patients (<1%), five OCA 10-mg patients (<1%), and 57 OCA 25-mg patients (9%). Of
21 those 57 patients in the OCA 25-mg group who discontinued due to pruritus, 36 discontinuations
22 were protocol-mandated based on the investigator-assessed grade of the event.

1 In patients receiving OCA, LDLc increased through month 1 (mean [SE]: placebo -3.0 [0.9]
2 mg/dL, OCA 10-mg, 17.8 [1.0] mg/dL, OCA 25-mg 23.8 [1.1] mg/dL) and decreased thereafter,
3 approaching baseline by month 18 (mean [SE]: placebo -7.1 [1.7] mg/dL, OCA 10-mg, 1.4 [2.0]
4 mg/dL, OCA 25-mg 2.7 [2.1] mg/dL) (figure 5). A total of 380 patients started statin therapy
5 during the study (placebo, n=66; OCA 10-mg, n=155; OCA 25-mg, n=159). Among OCA-
6 treated patients who initiated statins, increases from baseline in LDLc diminished with statin
7 treatment, and levels of LDLc were below baseline levels from month 6 through month 18
8 (figure S6). Levels of HDLc showed dose-dependent decreases through month 1 (mean [SE]:
9 placebo -0.7 [0.2] mg/dL, OCA 10-mg, -1.8 [0.2] mg/dL, OCA 25-mg -4.6 [0.3] mg/dL) and
10 were sustained through month 18; mean HDLc remained within the normal limit (<46 mg/dL) at
11 all timepoints. Changes in total cholesterol over time were similar to LDLc. A dose-dependent
12 decrease in triglycerides was observed by month 1 in the OCA groups and continued to gradually
13 decline over the study period with a maximum mean change from baseline of -37.4 mg/dL in the
14 OCA 25-mg group at month 18 (figure 5).

15 The incidence of cardiovascular AEs and SAEs was similar across the treatment groups (AEs:
16 5% placebo, 7% OCA 10-mg, and 6% OCA 25-mg; SAEs 2% placebo, 1% OCA 10-mg, 2%
17 OCA 25-mg). Treatment with OCA was associated with a generally modest and transient rise in
18 glycemic parameters (glucose, insulin, HOMA-IR, and HbA1c) that occurred early and returned
19 to levels similar to placebo after approximately 6 months of OCA treatment. In a subgroup
20 analysis of patients with type 2 diabetes, OCA treatment was associated with a transient increase
21 in glucose (month 1) and HbA1C (month 3) with return to levels similar to placebo by month 6.
22 In nondiabetic patients, small, sustained increases in glucose were seen with OCA 25-mg by
23 month 1 (mean [SE]: 8.2 [1.0] mg/dL); there were no changes in HbA1C in this group (figure

1 S7). Blood pressure was generally stable, but variable, with no significant difference between
2 treatment groups. Other vital signs were not affected by study treatments.
3 Gallstone-related AEs occurred at a rate of <1%, 1% and 3% in placebo, OCA 10-mg and OCA
4 25-mg patients respectively. Pancreatitis, a more serious and potentially cholelithiasis-related
5 event, was rare and evenly distributed across treatment groups (incidence <1%). Hepatic SAEs
6 were uncommon, and each case was reviewed by independent expert hepatologists. While more
7 events occurred in the OCA 25-mg group (6 [$<1\%$]) than the OCA 10-mg group (2 [$<1\%$]) or
8 placebo group (2 [$<1\%$]), expert reviewers did not identify any consistent pattern of liver injury
9 and all cases were associated with confounding severe intercurrent illness and/or concomitant
10 medications.
11 A total of three deaths occurred on study (two placebo [bone cancer and cardiac arrest], and one
12 OCA 25-mg [glioblastoma]); none were considered related to study treatment.

13

14

DISCUSSION

15

16 The REGENERATE study is the first positive phase 3 trial in NASH and represents a landmark
17 in the development of new therapies for an increasingly common chronic liver disease.¹²⁻¹⁵
18 Treatment with OCA 25-mg met the primary endpoint of improvement in fibrosis with no
19 worsening of NASH in patients with stage F2 or F3 fibrosis, at the month-18 interim analysis.
20 The robust antifibrotic effect of OCA was dose-dependent and consistent across different patient
21 populations, subgroups, and was further supported by fibrosis-related secondary endpoints
22 including a ≥ 2 -stage improvement in fibrosis. Per the draft guidance from the FDA on efficacy
23 endpoints for clinical trials in NASH, improvement in fibrosis by ≥ 1 stage with no worsening of

1 NASH is likely to predict clinical benefit.¹⁰ Patients with NASH have an almost 65 times greater
2 risk of liver-specific mortality and almost 3 times greater risk/rate of overall mortality compared
3 to healthy subjects¹⁴. Fibrosis has been shown to be the strongest histological predictor of liver-
4 related adverse outcomes, including liver-related death.¹⁶⁻¹⁹ Treatment with OCA 25-mg both
5 improved fibrosis and prevented progression of fibrotic disease, demonstrating a halting of
6 disease progression. To slow or reverse the progression of fibrosis is the ultimate goal of NASH
7 treatment as fibrosis is the most reliable predictor of liver-related mortality and once patients
8 progress to cirrhosis, preventing complications of cirrhosis may become even more difficult.^{16,18}
9 More OCA-treated patients relative to placebo achieved NASH resolution with no worsening of
10 fibrosis, the second primary endpoint; however, neither OCA dose achieved statistical
11 significance. More patients receiving OCA 25-mg showed improvements in hepatocellular
12 ballooning and lobular inflammation, the two key individual histologic features of the pre-
13 specified NASH resolution endpoint. These data are relevant given that features of
14 steatohepatitis, such as hepatocellular ballooning, are predictive of increased liver-related events
15 and reduced liver transplant-free survival.¹⁹ In addition, more patients receiving OCA 25-mg had
16 no worsening of fibrosis and ≥ 2 -point improvement in NAS, the primary endpoint traditionally
17 used in phase 2 studies such as FLINT⁷ and PIVENS,²⁰ indicating that OCA reduces NASH
18 disease activity.

19 A greater proportion of OCA 25-mg patients compared to placebo achieved NASH resolution as
20 defined in the pathologist diagnostic assessment of the absence of definite steatohepatitis at
21 month 18. This evaluation was based on an assessment of the overall pattern of histologic lesions
22 or injury, as opposed to the more rigid categorical scoring system of the pre-specified
23 methodology described above. This finding has clinical relevance given that this definition is

1 commonly used to diagnose NASH in clinical practice, as well as in natural history studies
2 evaluating any correlation between presence of NASH and mortality¹⁶. The apparent dichotomy
3 of substantial improvements in key individual components of NASH, while failing to meet the
4 pre-specified primary endpoint of NASH resolution demonstrates the challenges associated with
5 assessing histological response in complex, composite pathological patterns such as
6 steatohepatitis. The NAS, a tool designed to measure disease activity and severity in NASH, is
7 distinct from a clinical diagnosis of definite steatohepatitis. In an investigation into the
8 relationship between NAS and the diagnosis of steatohepatitis, threshold values of NAS did not
9 always correlate with pathologist overall assessment of presence of NASH.²¹ Therefore, as the
10 field continues to evolve it may be more appropriate to establish the presence/absence of NASH
11 using diagnostic criteria as an endpoint.

12 In addition to consistent improvements in multiple histologic parameters, improvement in liver
13 health was also evident based on clinically relevant, dose-dependent, improvements in markers
14 of liver injury (ALT and AST) and oxidative stress (GGT). The modest increases in ALP are
15 consistent with earlier observations and are associated with an on-target effect of FXR activation.
16 Lifestyle modifications including weight loss have been shown to be an effective
17 nonpharmacologic therapy for NAFLD. Weight loss >7% has been associated with improvement
18 in NAS, and weight loss $\geq 10\%$ has been associated with improvement in fibrosis.²² OCA-treated
19 patients in REGENERATE experienced weight loss of approximately 2%, an amount lower than
20 that expected to have an effect on histologic parameters of NASH. Although modest, the effect
21 of OCA on weight is important to note given the prevalence of obesity and metabolic
22 abnormalities in this population.

1 Based on a substantial database including almost 2000 patients, of whom approximately 900
2 were exposed for ≥ 18 months, OCA was generally well tolerated. The majority of AEs were
3 mild to moderate in severity and were generally consistent with the known safety profile of
4 OCA.⁷ As previously seen, mild to moderate pruritus was the most commonly reported AE, the
5 incidence of which was dose dependent. More subjects in the OCA 25-mg group experienced
6 pruritus that led to treatment discontinuation; however, the vast majority of randomized patients
7 were ongoing in the study through at least month 18. The impact of pruritus in this study on
8 patient-reported outcomes and its relationship to OCA is being investigated.²³ The incidence of
9 hepatic-related AEs was balanced across treatment groups, and serious hepatic complications
10 were rare; although numerically more occurred in the OCA 25-mg treated group, there was no
11 clear pathologic pattern seen consistently among these SAEs and all cases were confounded by
12 severe intercurrent illness. Treatment with OCA was associated with serum lipid changes that
13 were consistent with a class effect of FXR activation, as well as limited increases in glycemic
14 parameters. However, these increases were manageable by clinical practice measures. The
15 impact of lipid and glycemic laboratory changes on cardiovascular risk should be assessed in the
16 context of other OCA-related reductions in risk factors, including a decrease in weight, serum
17 triglyceride levels, and GGT, a promising marker for assessing cardiovascular risk.^{24,25} The
18 incidence of cardiovascular AEs and SAEs was low and similar across treatment groups.

19 The results of the interim analysis of REGENERATE reported here are clinically relevant in the
20 context of fibrosis due to NASH but may underestimate the long-term benefit of OCA on the
21 target illness. Improvement in fibrosis, a generally slow-changing feature, was observed at the
22 month-18 interim analysis of the ongoing study, and the effect size may increase with prolonged
23 therapy. This has been shown with other interventions that reported improvement in fibrosis at

1 early time points with a greater effect over the longer term. For example, tenofovir treatment
2 resulted in 10% fewer patients with hepatitis B virus-associated advanced fibrosis or cirrhosis
3 after the first year of treatment (28% vs 38% at baseline).²⁶ In the tenofovir study, patients
4 continued to improve on treatment, and the proportion of patients with advanced fibrosis or
5 cirrhosis declined to 12% at year 5.²⁶ In REGENERATE, the continuing improvement in liver
6 enzyme markers of fibrosis such as ALT and AST suggest the potential for further antifibrotic
7 response. Data from the ongoing long-term outcome portion of the study will inform whether
8 prolonged therapy will result in a higher rate of antifibrotic response.

9 In conclusion, the totality of data from the month 18 interim analysis of this pivotal, phase 3
10 study provides strong evidence of an improvement in clinically significant histologic endpoints
11 deemed reasonably likely to predict clinical benefit with OCA treatment and affirms the positive
12 benefit-risk of OCA for the treatment of NASH with fibrosis. Beneficial effects of OCA on
13 fibrosis and components of NASH disease activity were robust, based on the observed
14 consistency of results across multiple histologic endpoints with reproducible response ratios, as
15 well as the evident dose-response and markedly consistent benefit across analysis populations.

16 Treatment with OCA had a beneficial effect on other markers of chronic liver disease
17 (hepatocellular ballooning and lobular inflammation), hepatocellular injury (ALT and AST), and
18 oxidative stress (GGT). OCA was generally well tolerated, with a profile that is generally
19 consistent with prior studies. Following the month-18 interim analysis, REGENERATE
20 continues in a blinded fashion, and patients will be followed over an extended period for clinical
21 outcomes, such as all-cause mortality, liver-related clinical outcomes, and long-term safety, to
22 confirm clinical benefit. In a chronic disease with no approved therapies and potential for serious

1 sequelae, these findings provide compelling evidence that patients with non-cirrhotic advanced
2 fibrosis due to NASH may benefit from OCA treatment.

3

4

1 **CONTRIBUTORS**

2
3 VR, AJS, and ZMY participated in initial study design in collaboration with the sponsor (DS,
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10
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12
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DATA SHARING STATEMENT

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21 The authors declare that all data supporting the findings of this interim analysis are available
22 within the article and its supplementary information files. The study is ongoing at the time of
23 publication and blinded at the individual level; patient-level data therefore are not available.

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TABLES

[[SUPPLEMENTARY TABLES ARE INCLUDED IN THIS SECTION FOR REVIEW, AND WILL BE MOVED TO A SUPPLEMENTARY APPENDIX PRIOR TO SUBMISSION; ADDITIONALLY, TABLES 1-3 WILL BE SUBMITTED AS A SEPARATE WORD DOCUMENT.]]

Table 1 Demographic and baseline clinical characteristics (ITT population, N=931)

	Placebo (n=311)	OCA 10 mg (n=312)	OCA 25 mg (n=308)
Age, years	55 (12)	55 (11)	55 (11)
Female, n (%)	187 (60)	177 (57)	175 (57)
White, n (%)*	264 (94)	263 (92)	249 (87)
Hispanic ethnicity, n (%) [†]	52 (18)	42 (15)	47 (17)
Fibrosis stage F3, n (%)	169 (54)	182 (58)	169 (55)
NAS ≥6, n (%)	215 (70)	211 (68)	208 (68)
Type 2 diabetes, [‡] n (%)	175 (56)	171 (55)	171 (56)
Dyslipidaemia, n (%)	211 (68)	217 (70)	205 (67)
Hypertension, n (%)	215 (69)	215 (69)	196 (64)
Lipids			
Total cholesterol, mg/dL	184.5 (42.7)	185.2 (53.0)	183.5 (44.7)
HDLc, mg/dL	45.6 (11.1)	44.9 (12.1)	44.3 (11.0)
LDLc, mg/dL	114.8 (38.2)	113.8 (38.4)	113.3 (38.8)
Triglycerides, mg/dL	178.7 (154.5)	184.6 (195.0)	181.7 (131.6)
Metabolic factors			
Fasting glucose, mg/dL	119.1 (38.3)	120.8 (43.6)	119.5 (40.3)
Body weight, kg	95 (19)	95 (19)	95.40 (19)
HOMA-IR	9.6 (11.8)	9.9 (16.9)	8.3 (10.2)
HbA1c, %	6.6 (1.22)	6.5 (1.2)	6.5 (1.3)
Laboratory parameters			
ALT, U/L	80 (57)	76 (47)	80 (56)
AST, U/L	59 (41)	57 (34)	57 (34)
Platelet count, x10 ⁹ /L	241.9 (67.0)	238.5 (68.0)	237.2 (69.0)
Total bilirubin, mg/dL	0.64 (0.3)	0.65 (0.3)	0.69 (0.3)
Concomitant medication use			
Lipid lowering, n (%) [§]	175 (56)	170 (54)	160 (52)
Statins, n (%)	144 (46)	142 (46)	127 (41)
Antidiabetic medication, n (%)	167 (54)	171 (55)	159 (52)

Thiazolidinediones, [‡] n (%)	5 (2)	9 (3)	4 (1)
Vitamin E, [*] n (%)	42 (14)	34 (11)	32 (10)

Data are mean (SD) unless otherwise noted.

^{*}Percentages calculated based on patients for whom race information was not missing.

[†]Percentages calculated based on patients for whom ethnicity information was not missing.

[§]In addition to statins, lipid lowering drugs included fibrates, cholesterol-absorbing resins, PCSK9 inhibitors, and omega-3 fatty acids

[‡]Randomisation was stratified based on presence of type 2 diabetes and treatment with thiazolidinediones or vitamin E.

ALT=alanine aminotransferase. AST=aspartate aminotransferase. NAS=NAFLD activity score. PCSK9=proprotein convertase subtilisin-kexin type 9. SD=standard deviation.

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2

	ITT population (N=931)			Per-protocol population (N=668)		
Primary Endpoint, RR (95% CI)	Placebo (n=311)	OCA 10 mg (n=312)	OCA 25 mg (n=308)	Placebo (n=224)	OCA 10 mg (n=226)	OCA 25 mg (n=218)
Improvement of fibrosis with no worsening of NASH	-	1.48 (1.01, 2.18) p=0.04	1.94 (1.35-2.78) p=0.0002	-	1.62 (1.06, 2.48) p=0.02	2.15 (1.44, 3.21) p<0.0001
Resolution of NASH with no worsening of fibrosis	-	1.39 (0.86, 2.25) p=0.18	1.45 (0.90, 2.35) p=0.13	-	1.49 (0.86, 2.33) p=0.11	1.41 (0.86, 2.33) p=0.18
Secondary Endpoint, n (%)	Placebo (n=311)	OCA 10 mg (n=312)	OCA 25 mg (n=308)	Placebo (n=224)	OCA 10 mg (n=226)	OCA 25 mg (n=218)
Improvement of fibrosis by ≥ 1 stage and/or resolution of NASH without worsening of either	49 (15.8)	67 (21.5) p=0.07	84 (27.3) p=0.0005	41 (18.3)	59 (26.1) p=0.04	71 (32.6) p=0.0004
No worsening of fibrosis and no worsening of NASH	117 (37.6)	127 (40.7) p=0.43	147 (47.7) p=0.011	100 (44.6)	109 (48.2) p=0.43	125 (57.3) p=0.006
Improvement of NAS by ≥ 2 with no worsening of fibrosis	76 (24.4)	94 (30.1) p=0.11	112 (36.4) p=0.001	69 (30.8)	82 (36.3) p=0.19	96 (44.0) p=0.004
Improvement of fibrosis and resolution of NASH as a composite endpoint	13 (4.2)	23 (7.4) p=0.090	23 (7.5) p=0.080	11 (4.9)	22 (9.7) p=0.045	20 (9.2) p=0.064
Improvement in fibrosis by ≥ 2 stages	15 (4.8)	19 (6.1) p=0.49	30 (9.7) p=0.018	10 (4.5)	16 (7.1) p=0.22	29 (13.3) p=0.0008
Resolution of fibrosis	4 (1.3)	8 (2.6) p=0.25	10 (3.2) p=0.10	4 (1.8)	8 (3.5) p=0.21	9 (4.1) p=0.14

CI=confidence interval, NAS=NAFLD activity score, NASH=nonalcoholic steatohepatitis, RR=response ratio.
P values compare OCA treatment with placebo.

1
2

Table 3 Adverse events occurring in ≥5% of patients in either OCA group (safety population, N=1968)			
System Organ Class Preferred Term, n (%)	Placebo (n=657)	OCA 10 mg (n=653)	OCA 25 mg (n=658)
Skin and Subcutaneous Tissue Disorders			
Pruritus	123 (19)	183 (28)	336 (51)
Grade 1 (mild or localised)	90 (14)	113 (17)	148 (22)
Grade 2 (intense or wide spread)	30 (5)	67 (10)	152 (23)
Grade 3 (intense or widespread and limit activities of daily living)	3 (<1)	3 (<1)	36 (5)
Gastrointestinal Disorders			
Nausea	77 (12)	72 (11)	83 (13)
Constipation	36 (5)	65 (10)	70 (11)
Abdominal pain	62 (9)	66 (10)	67 (10)
Diarrhoea	79 (12)	44 (7)	49 (7)
Abdominal pain upper	35 (5)	46 (7)	45 (7)
Vomiting	33 (5)	34 (5)	44 (7)
Abdominal distension	23 (4)	31 (5)	31 (5)
Infections and Infestations			
Urinary tract infection	49 (7)	54 (8)	62 (9)
Upper respiratory tract infection	44 (7)	47 (7)	54 (8)
Nasopharyngitis	41 (6)	34 (5)	45 (7)
Bronchitis	28 (4)	34 (5)	35 (5)
Sinusitis	35 (5)	36 (6)	30 (5)
Investigations			
Low density lipoprotein increased	47 (7)	109 (17)	115 (17)
Blood cholesterol increased	12 (2)	30 (5)	38 (6)
Musculoskeletal and Connective Tissue Disorders			
Arthralgia	55 (8)	50 (8)	50 (8)
Back pain	50 (8)	56 (9)	40 (6)
Metabolism and Nutrition Disorders			
Hyperlipidaemia	18 (3)	42 (6)	55 (8)
Diabetes mellitus	36 (5)	46 (7)	45 (7)
Hypercholesterolaemia	14 (2)	35 (5)	29 (4)
General Disorders and Administration Site Conditions			
Fatigue	88 (13)	78 (12)	71 (11)
Nervous System Disorders			
Headache	51 (8)	42 (6)	34 (5)
Dizziness	28 (4)	32 (5)	25 (4)
Respiratory, Thoracic and Mediastinal Disorders			
Cough	27 (4)	29 (4)	38 (6)
Vascular Disorders			
Hypertension	28 (4)	36 (6)	39 (6)
LDLc=low density lipoprotein cholesterol. Note: Arranged by descending order of incidence (system organ class and preferred term within system organ class) in the OCA 25-mg group, followed by descending order of incidence in the OCA 10-mg group.			

3

Supplementary Table S1 Demographic and baseline clinical characteristics (PP population, N=668)			
Characteristics	Placebo (n=224)	OCA 10 mg (n=226)	OCA 25 mg (n=218)
Age, years, mean (SD)	55 (12)	55 (11)	54 (12)
Female, n (%)	133 (59)	126 (56)	122 (56)
White, n (%) [*]	189 (94)	192 (94)	185 (89)
Hispanic ethnicity, n (%) [†]	29 (14)	29 (14)	33 (16)
Fibrosis stage F3, n (%)	122 (54)	135 (60)	117 (54)
NAS ≥6, n (%)	159 (72)	152 (67)	144 (66)
Type 2 diabetes, [‡] n (%)	120 (54)	121 (54)	119 (55)
Weight, mean kg (SD)	96 (19)	94 (19)	97 (20)
Laboratory parameters, mean (SD)			
ALT, U/L	77 (53)	73 (44)	82 (61)
AST, U/L	57 (38)	55 (31)	56 (34)
Concomitant medication use			
Lipid lowering, n (%) [§]	131 (58)	128 (57)	113 (52)
Statins, n (%)	108 (48)	108 (48)	86 (39)
Antidiabetic medication, n (%)	117 (52)	120 (53)	111 (51)
Thiazolidinediones, [‡] n (%)	3 (1)	8 (4)	2 (<1)
Vitamin E, [*] n (%)	29 (13)	25 (11)	23 (11)
[*] Percentages calculated based on patients for whom race information was not missing [†] Percentages calculated based on patients for whom ethnicity information was not missing [§] In addition to statins, lipid lowering drugs included fibrates, cholesterol-absorbing resins, PCSK9 inhibitors, and omega-3 fatty acids [‡] Randomisation was stratified based on presence of type 2 diabetes and treatment with thiazolidinediones or vitamin E. ALT=alanine aminotransferase. AST=aspartate aminotransferase. NAS=NAFLD activity score. PCSK9=proprotein convertase subtilisin-kexin type 9. SD=standard deviation.			

Supplementary Table S2 Summary of treatment-emergent adverse events (safety population, N=1968)			
n (%)	Placebo (n=657)	OCA 10 mg (n=653)	OCA 25 mg (n=658)
≥1 Treatment-emergent adverse event (TEAE)	548 (83)	579 (89)	601 (91)
TEAEs by severity ^a			
Mild	160 (24)	163 (25)	130 (20)
Moderate	294 (45)	323 (49)	338 (51)
Severe	87 (13)	89 (14)	130 (20)
Life-threatening	5 (<1)	4 (<1)	2 (<1)
Death	2 (<1)	0	1 (<1)
TEAEs leading to treatment discontinuation	41 (6)	39 (6)	83 (13)
Serious adverse events (SAEs)	75 (11)	72 (11)	93 (14)

^a Subjects reporting more than one adverse event are counted only once using the highest severity. Adverse events are graded for severity using Common Terminology Criteria for Adverse Events (CTCAE) Version 4.03.

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1 **FIGURE LEGENDS**

2
3 **Figure 1.** Patient flow diagram.

4
5 **Figure 2.** Primary endpoints. The proportion of patients with improvement in fibrosis ≥ 1 stage
6 and no worsening of NASH in the ITT (Panel A) and per protocol (Panel B) populations, and the
7 proportion of patients with resolution of NASH and no worsening of fibrosis in the ITT (Panel
8 C) and per protocol (Panel D) populations. Fibrosis improvement was evaluated per NASH CRN
9 criteria; no worsening of NASH defined as defined as no worsening of hepatocellular ballooning,
10 lobular inflammation or steatosis. NASH resolution defined as: (i) overall pathologist assessment
11 of “no steatohepatitis,” and (ii) hepatocellular ballooning = 0 and lobular inflammation = 0 or 1.

12 *Statistically significant in accordance with the statistical analysis plan as agreed with the FDA.

13
14 **Figure 3.** Regression or progression of fibrosis by ≥ 1 stage. The proportion of patients with
15 improved or worsened fibrosis by ≥ 1 stage is shown for patients in the per-protocol population
16 with available fibrosis stage data at month-18/end of treatment (n=656).

17
18
19 **Figure 4.** Changes in liver biochemistry over time. Mean (SE) values of change from baseline up
20 to month 18 are shown for patients from each treatment group in the ITT population (\circ placebo,
21 \blacktriangle OCA 10-mg, \blacktriangledown OCA 25-mg).

22

1 **Figure 5.** Changes in serum lipids over time. Mean (SE) values of change from baseline up to
2 month 18 are shown for patients from each treatment group in the safety population (○ placebo,
3 ▲ OCA 10-mg, ▼ OCA 25-mg).

4
5 **Supplementary Figure S1.** Subgroup analysis of fibrosis improvement by ≥ 1 stage with no
6 worsening of NASH. Odds ratios and 95% confidence intervals of obeticholic acid versus
7 placebo for patients in the ITT population grouped by fibrosis stage, NAFLD Activity Score
8 (NAS), and type 2 diabetes. An odds ratio greater than 1 favours obeticholic acid.

9
10 **Supplementary Figure S2.** The proportion of patients with improvements in histologic features
11 of NASH (steatosis, lobular inflammation, and hepatocellular ballooning) in the ITT population.

12
13 **Supplementary Figure S3.** The proportion of patients with resolution of NASH with no
14 worsening of fibrosis, based on the pathologist diagnostic assessment of the absence of definite
15 steatohepatitis in the ITT (Panel A) and per protocol (Panel B) populations.

16
17 **Supplementary Figure S4.** The proportion of patients with improvement of fibrosis and/or
18 resolution of NASH with no worsening of either in the ITT population. NASH resolution defined
19 as hepatocellular ballooning = 0 and lobular inflammation = 0 or 1.

20
21 **Supplementary Figure S5.** The proportion of patients in the ITT population with elevated ALT
22 (n=546) (Panel A) or AST (n=665) (Panel B) at baseline who achieved transaminase levels
23 \leq ULN. For ALT, ULN = 55U/L; for AST, ULN = 34 U/L.

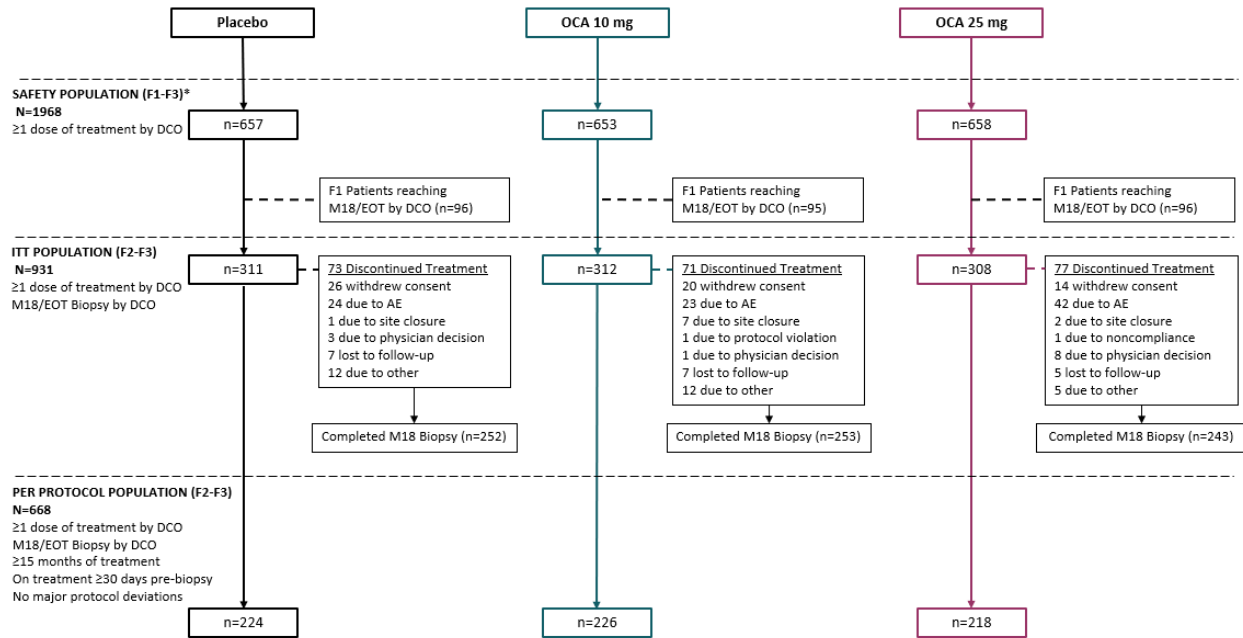
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2 **Supplementary Figure S6.** Changes in LDLc over time by statin use. Mean (SE) values of
3 change in LDLc from baseline up to month 18 are shown patients who never used statins,
4 patients who were using statins at baseline, and patients who started using statins during the
5 study for each treatment group in the safety population (○ placebo, ▲ OCA 10-mg, ▼ OCA 25-
6 mg).

7
8 **Supplementary Figure S7.** Changes in glucose and HbA1c over time. Mean (SE) values of
9 change in glucose (Panels A and C) and HbA1c (Panels B and D) from baseline up to month 18
10 are shown based on diabetes status for patients from each treatment group in the safety
11 population (○ placebo, ▲ OCA 10-mg, ▼ OCA 25-mg).

FIGURES

[[SUPPLEMENTARY FIGURES ARE INCLUDED IN THIS SECTION FOR REVIEW, AND WILL BE MOVED TO A SUPPLEMENTARY APPENDIX PRIOR TO SUBMISSION]]

Figure 1 Patient flow diagram



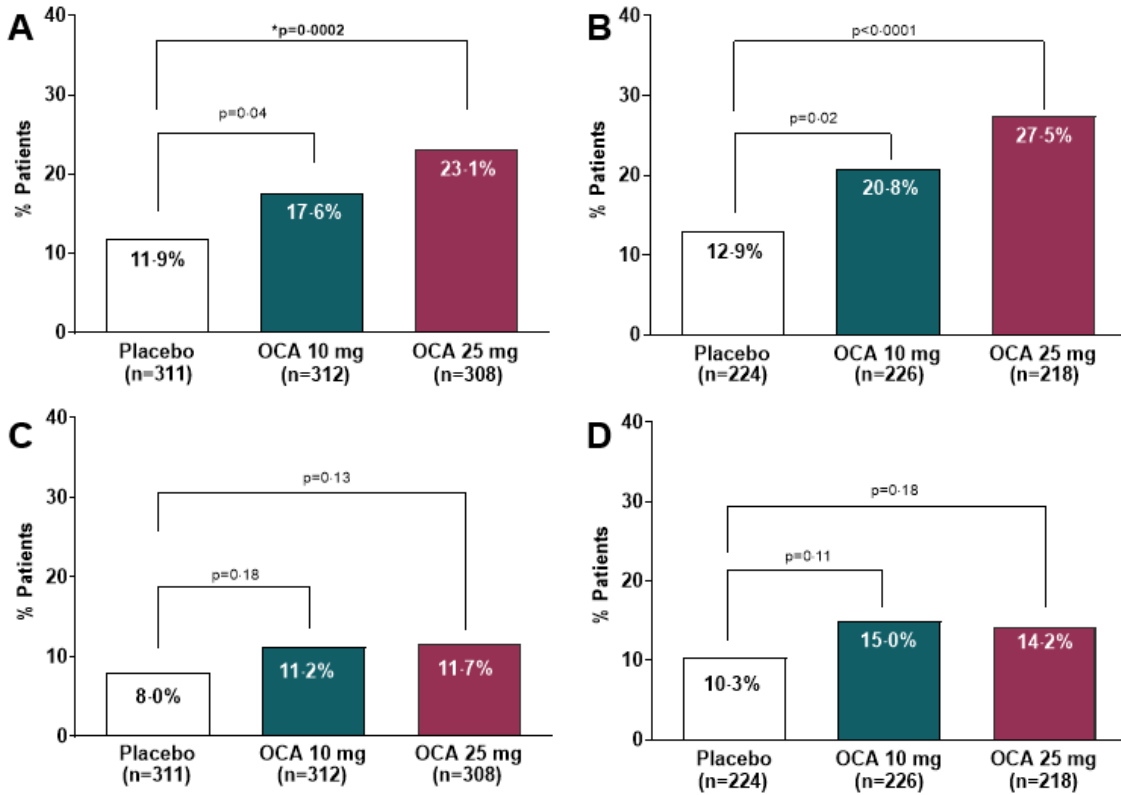
*750 patients included in the safety population had not reached their M18/EOT visit by DCO and were therefore not included in the ITT or per protocol populations.
AE: Adverse Event, DCO: Data Cutoff, ITT: intent-to-treat, OCA: Obeticholic Acid

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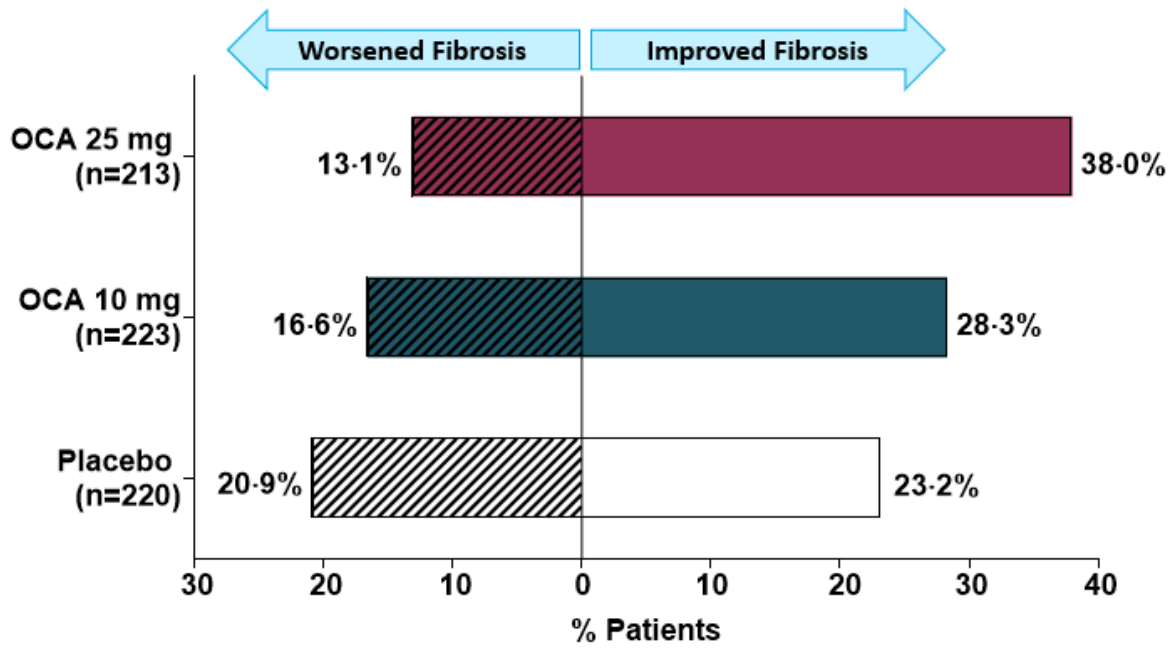
1 **Figure 2 Primary endpoint**

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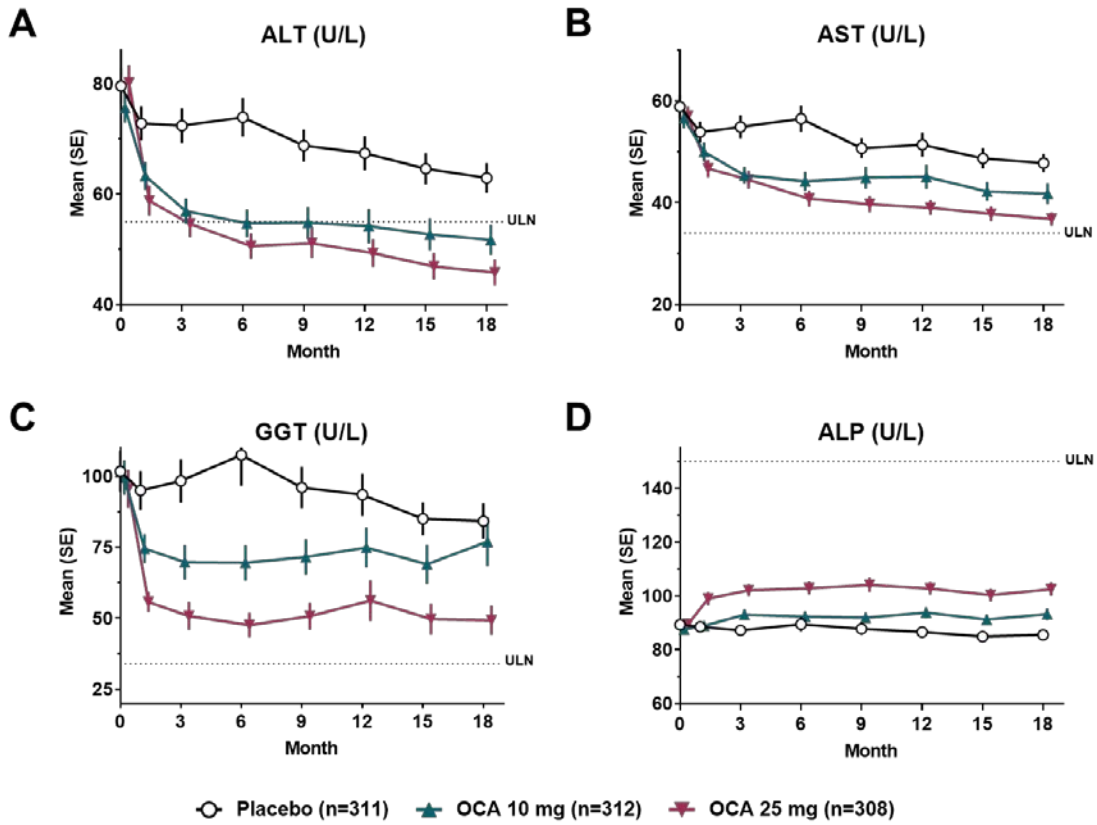
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1 **Figure 3. Improvement versus progression of fibrosis by ≥ 1 stage**
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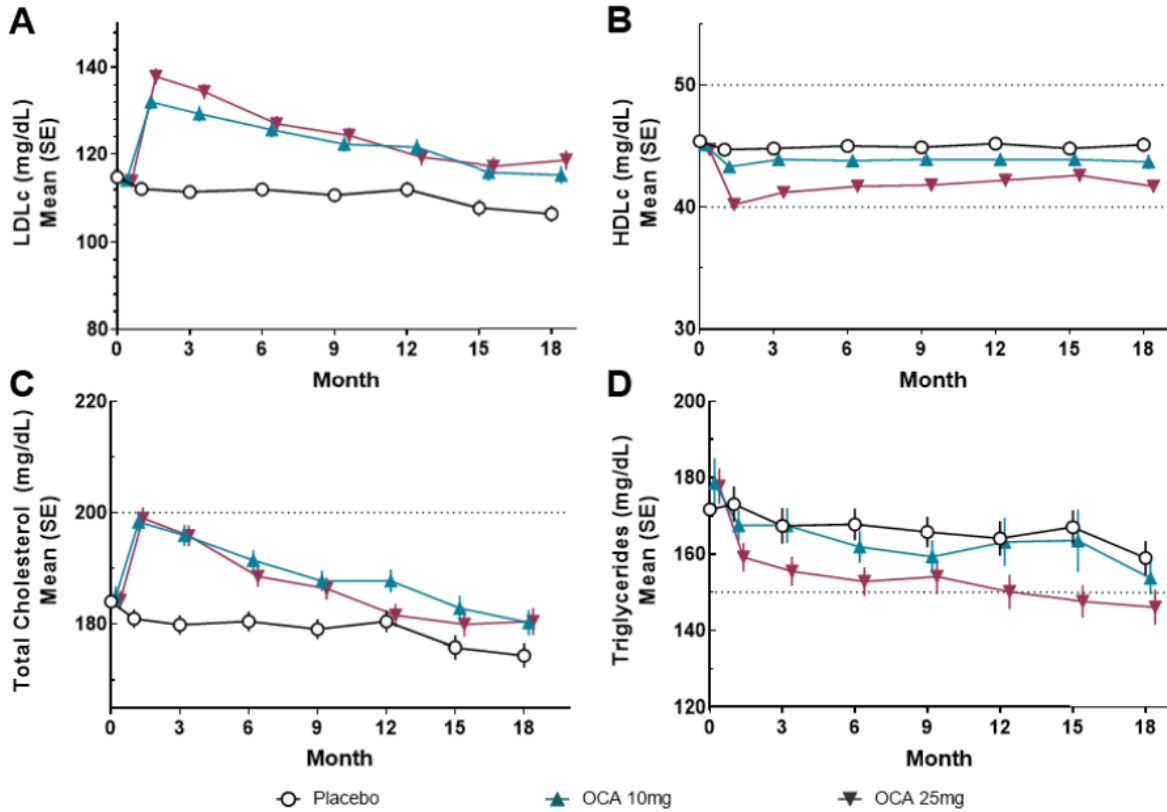
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1 Figure 4 Changes in liver biochemistry over time



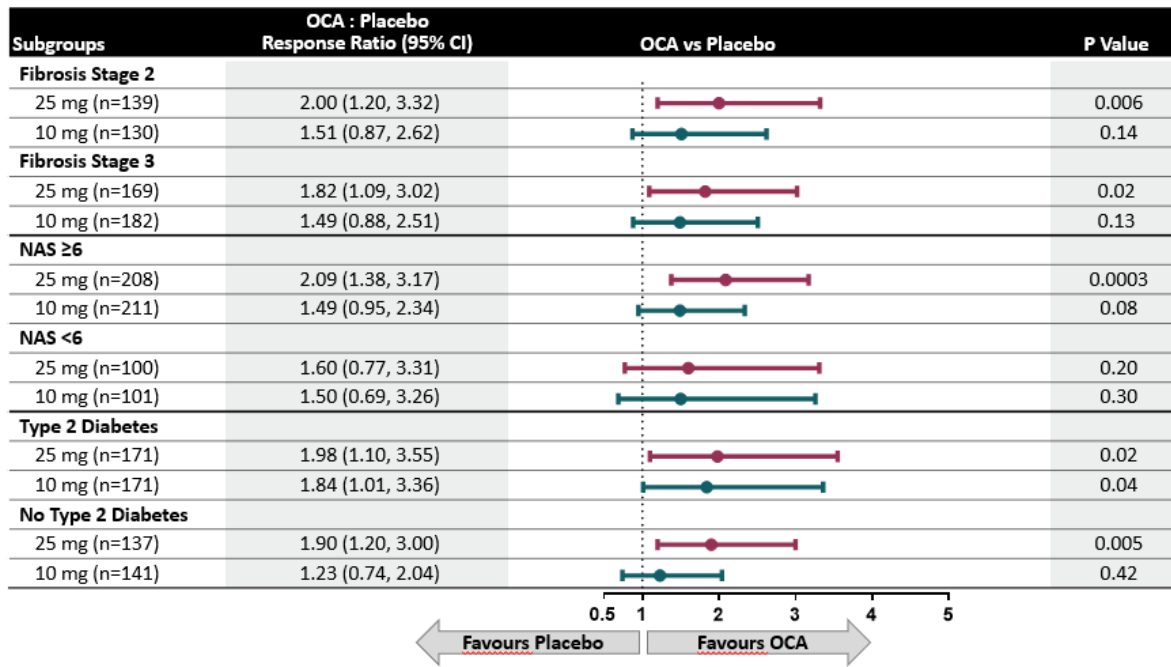
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1 Figure 5 Changes in serum lipids over time
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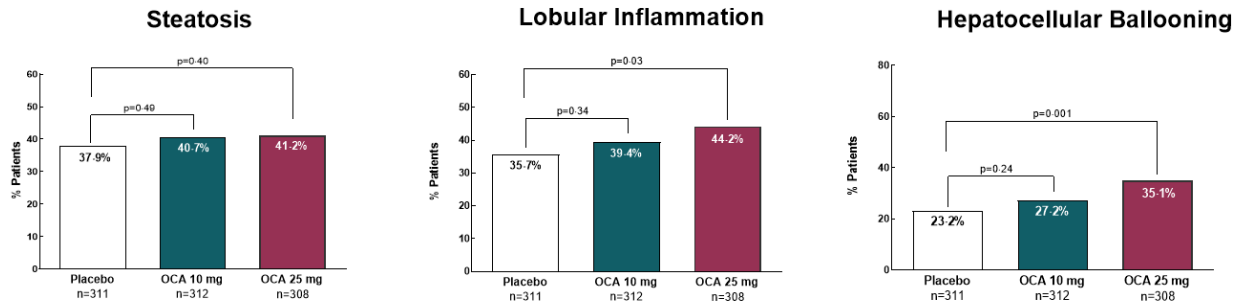
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1 **Supplementary Figure S1 Subgroup analysis of fibrosis improvement by ≥1 stage with no worsening of NASH**
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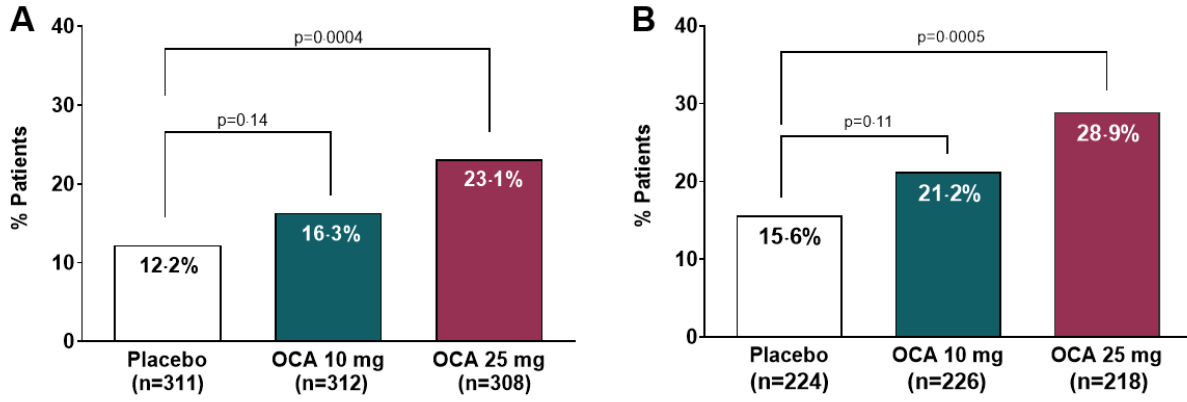
1 **Supplementary Figure S2 Improvements in histologic features of NASH (steatosis, lobular inflammation, and**
2 **hepatocellular ballooning)**
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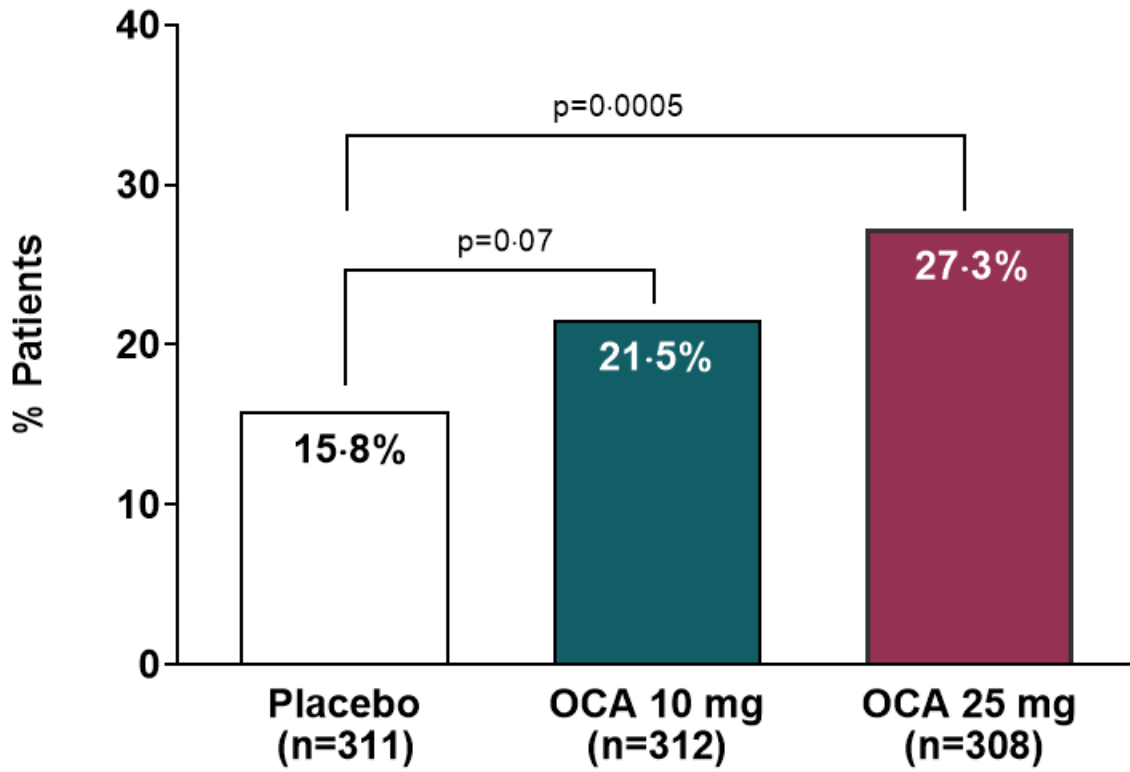
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Supplementary Figure S3 Pathologist diagnostic assessment of NASH: Resolution of NASH with no worsening of fibrosis based on the absence of definite steatohepatitis



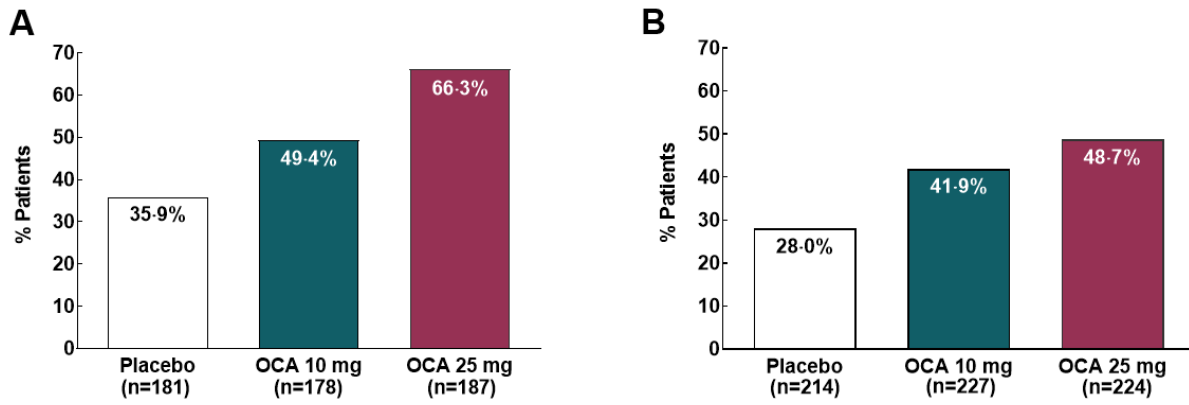
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1 **Supplementary Figure S4 Improvement of fibrosis and/or resolution of NASH with no worsening of either**
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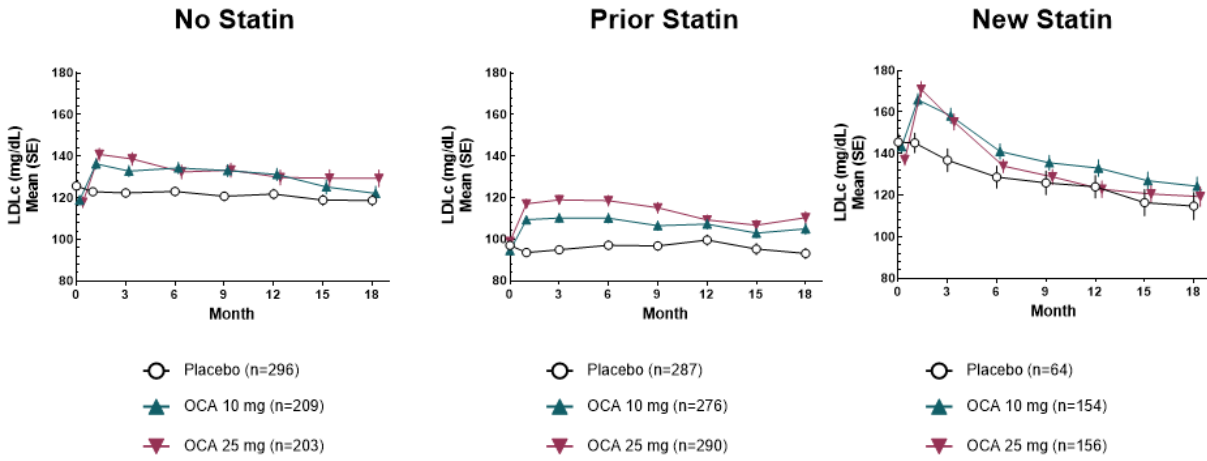
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1 **Supplementary Figure S5 Normalisation of elevated transaminase levels**
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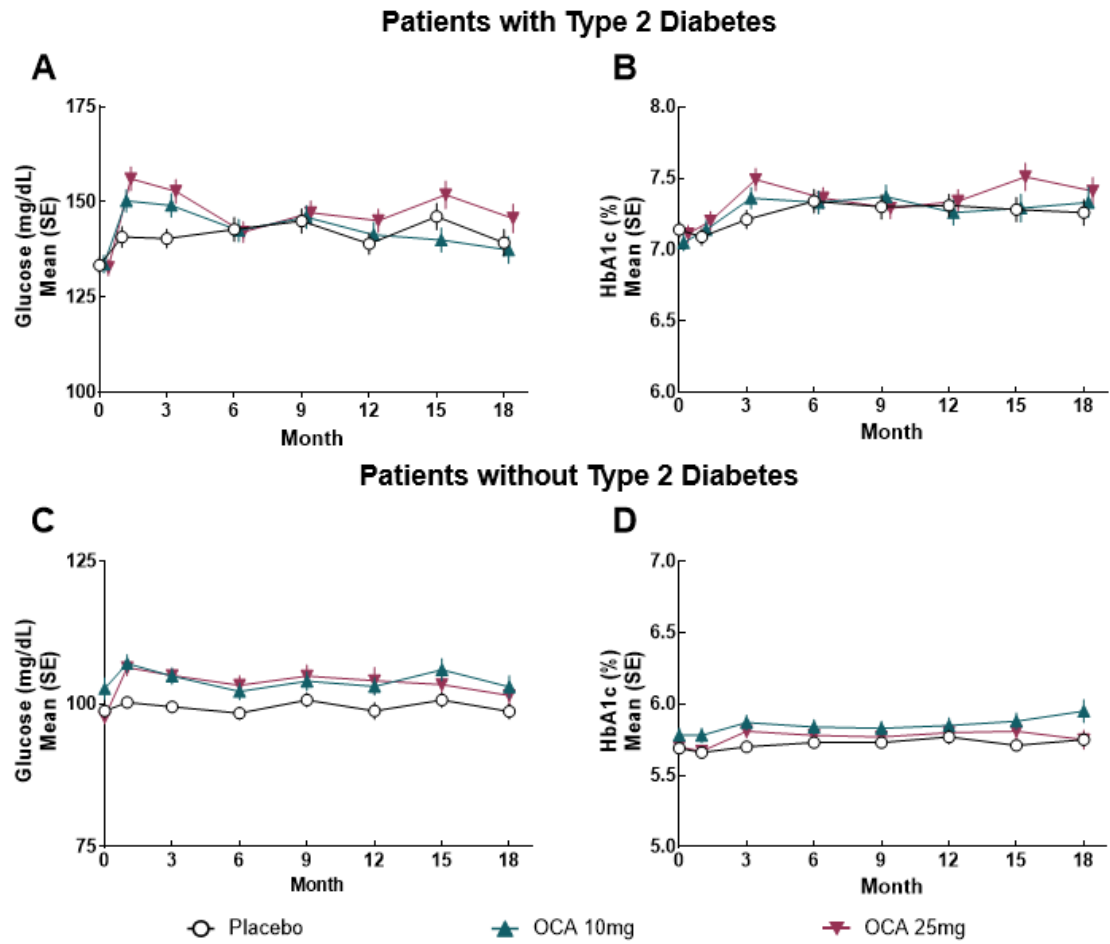
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Supplementary Figure S6 Changes in LDLc over time by statin use



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1 Supplementary Figure S7 Changes in glucose and HbA1C over time by diabetes status.
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