EDP-514, a Novel Pangenotypic Class II Hepatitis B Virus Core Inhibitor: Results of a 28-day Phase 1b Study in NUC-suppressed CHB Patients

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INTRODUCTION

Chronic hepatitis B (CHB) virus infection is a global public health challenge, with estimates of more than 296 million hepatitis B virus (HBV) carriers worldwide, of whom approximately 820,000 die annually from HBV-related liver disease. There is an unmet medical need for curative therapy, i.e., a finite treatment which yields a sustained post-treatment response.

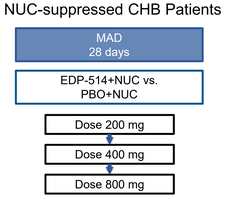
EDP-514 is a novel class II HBV core inhibitor. EDP-514 inhibits HBV replication with an *in vitro* EC_{50} of 18, 27 and 17 nM in HepAD38, HepDE19, and HepG2.2.15 cells, respectively, and a >4-log viral load reduction in HBV-infected chimeric mice with human liver cells. EDP-514 was shown to be generally safe and well tolerated over a broad range of single and multiple doses for up to 14 days in healthy adult subjects. Here, we present results of a Phase 1b, 28-day study in non-cirrhotic, HBeAg(+) or (-) CHB patients virologically suppressed on Nucleos(t)ide (NUC) therapy.

METHODS

Study Design

Eight subjects per cohort were randomized 3:1 to receive multiple once-daily oral doses of either EDP-514 at 200, 400, and 800 mg or placebo QD for 28 days.

The first cohort received 200 mg of EDP-514 and cohort progression was determined by a Study Adjudication Committee after review of blinded safety and available PK data.



Key Objectives

- Primary
  - To evaluate the safety and tolerability of multiple doses of EDP-514 administered to NUC-suppressed CHB patients
- Secondary
  - To evaluate the plasma PK of multiple doses of EDP-514 of NUC-suppressed CHB patients
  - To evaluate the antiviral activity of multiple doses of EDP-514 in NUC-suppressed CHB patients

Key Inclusion/Exclusion Criteria

- Inclusion Criteria:
  - Male and female subjects of any ethnic origin between the ages of 18 and 70 years
  - CHB subjects must have HBsAg detectable at screening and in most recent test at least 6 months prior
  - Screening HBV DNA < LLOQ and no HBV DNA ≥ LLOQ over previous 12 months
  - No change in HBV NUC treatment regimen for 12 months prior to screening
- Exclusion Criteria:
  - Documented prior diagnosis of cirrhosis
  - Documented extensive bridging fibrosis or cirrhosis
  - Evidence of hepatocellular carcinoma by imaging or screening AFP ≥ 50 ng/mL
  - Meeting defined safety laboratory parameters at screening
  - Coinfection with HIV, HAV, HCV, HDV, or HEV
  - Chronic liver disease of non-HBV etiology

Assessments

- Safety and tolerability assessments
  - Adverse events, clinical laboratory tests, physical examinations, vital signs, and electrocardiograms
- PK assessments
  - On Days 1 and 28, blood samples were collected at pre-dose and at 0.5, 1, 2, 3, 4, 5, 6 and 8 hours post-dose
  - On Days 3, 7, 14 and 21, blood samples were collected at pre-dose, 1-3 hrs post-dose and at least 1 hr after the first postd-ose sample and prior to next dose
  - Concentrations of EDP-514 and its metabolites were measured using a validated method
  - PK parameters were determined using non-compartmental methods in Phoenix WinNonlin (Pharsight Corporation)
- Antiviral activity assessments
  - HBV DNA levels
  - Incidences of virologic failure defined as HBV DNA level ≥LLOQ and which is confirmed to be ≥LLOQ on repeat testing

RESULTS

Subject Disposition and Demographics

- A total of 24 subjects were enrolled in the study
  - One subject from the EDP-514 200 mg arm discontinued from the study due to an adverse event of upper abdominal pain
  - Subjects were mostly male, Asian, HBeAg(-), with a mean age of 45 years, mean HBV DNA of 1.67 IU/mL, and mean HBV RNA of 1.13 log U/mL at baseline
- Subject demographics and disease characteristics are summarized in **Table 1**

Table 1. Baseline Demographics and Disease Characteristics

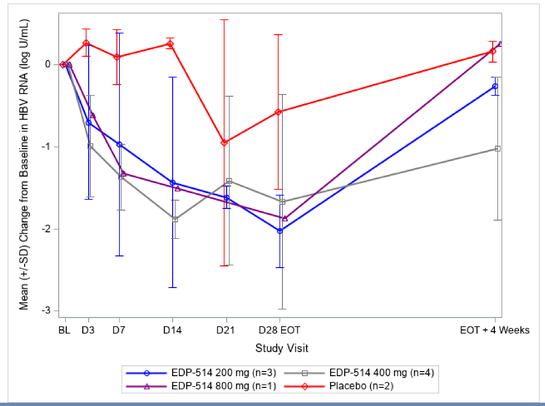
	200 mg QD (N=6)	400 mg QD (N=6)	800 mg QD (N=6)	Placebo (N=6)	Overall (N=24)
Sex [n (%)]					
Female	4 (66.7)	0	4 (66.7)	1 (16.7)	9 (37.5)
Male	2 (33.3)	6 (100)	2 (33.3)	5 (83.3)	15 (62.5)
Race [n (%)]					
White	1 (16.7)	1 (16.7)	1 (16.7)	3 (50.0)	6 (25.0)
Black or African American	0	2 (33.3)	0	0	2 (8.3)
Asian	5 (83.3)	3 (50.0)	5 (83.3)	3 (50.0)	16 (66.7)
Ethnicity [n (%)]					
Not Hispanic or Latino	6 (100)	6 (100)	6 (100)	6 (100)	24 (100)
Age (y)[a]	43.2 (25, 62)	41.8 (34, 52)	43.0 (31, 49)	52.7 (45, 61)	45.2 (25, 62)
BMI (kg/m^2)[a]	24.46 (18.3, 31.1)	27.22 (23.5, 34.6)	26.81 (24.8, 31.0)	28.09 (23.0, 34.4)	26.64 (18.3, 34.6)
HBeAg negative [n (%)]	5 (83.3)	5 (83.3)	5 (83.3)	6 (100)	21 (87.5)
HBV DNA (IU/mL)[a,b]	2.50 (0.0, 10.0)	2.50 (0.0, 5.0)	1.67 (0.0, 5.0)	0.00 (0.0, 0.0)	1.67 (0.0, 10.0)
HBV RNA (Log U/mL)[a,c]	1.18 (0.0, 2.5)	1.57 (0.0, 3.7)	0.83 (0.0, 4.2)	0.96 (0.0, 2.1)	1.13 (0.0, 4.2)
HBV RNA < LOD [n (%)][c]	3 (50.0)	2 (33.3)	5 (83.3)	4 (66.7)	14 (58.3)
Tenofovir (TAF/TDF) [n (%)]	5 (83.3)	5 (83.3)	6 (100)	6 (100)	22 (91.7)

a. Presented as Mean (Min, Max). b. HBV DNA Limit of Detection = 10 IU/mL, c. HBV RNA Limit of Detection = 1.65 Log U/mL
BMI = Body Mass Index; QD = Once Daily, y = Year, TAF = tenofovir alafenamide, TDF = tenofovir disoproxil fumarate

Antiviral Activity

- For subjects with detectable HBV RNA at baseline (**Table 1**), mean change from baseline to Day 28 for HBV RNA was -0.58, -2.03, -1.67, and -1.87 log U/mL in the placebo, 200 mg, 400 mg, and 800 mg groups, respectively. (**Figure 4**)
- EDP-514 led to a maximum HBV RNA reduction of 2.3 log in HBeAg(-) and 2.8 log in HBeAg(+) subjects in EDP-514 arms compared to 1.2 log in placebo
- As expected in this NUC-suppressed patient population, there were no discernible changes in HBV DNA, and also, no changes in HBeAg, HBcrAg, and HBsAg
- There were no instances of virologic failure were reported

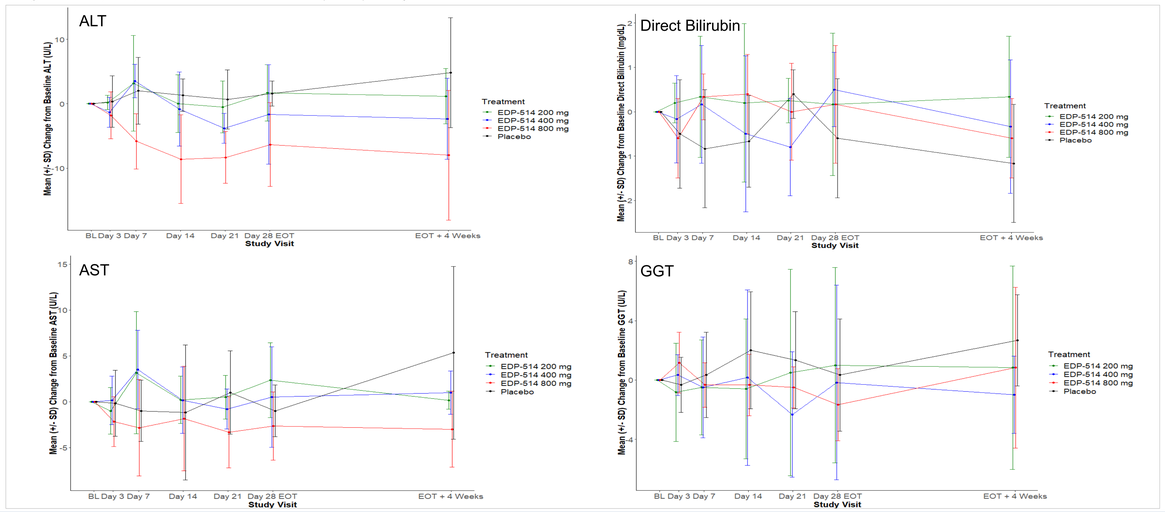
Figure 4. Antiviral Activity by HBV RNA Mean (±SD) Change from Baseline Over Time for Subjects with Detectable HBV RNA at Baseline



Safety

- There were no clinically significant laboratory abnormalities, including no clinically significant ALT, AST, direct bilirubin and GGT elevations or notable differences in mean (+/- SD) change from baseline of all arms (**Figure 3**)

Figure 3: ALT, AST, Direct Bilirubin and GGT Mean (+/- SD) Change from Baseline Over Time



Safety Summary

- Overall, EDP-514 was generally safe and well-tolerated in 200, 400 and 800 mg doses (**Table 3**)
- Eight patients reported treatment emergent adverse events (TEAEs); all were mild except for 1 moderate event (upper abdominal pain) in the EDP-514 200 mg arm that led to study drug discontinuation, and 1 severe event (drug hypersensitivity [allergic reaction to aloe cream]) in the EDP-514 800 mg arm that was unrelated to study drug
- There were no Grade 4 or serious TEAEs
- There were no clinically meaningful changes in ECG or vital signs

Table 3: Summary of TEAEs Following Administration of EDP-514 in the MAD Phase

System Organ Class Preferred Term [n (%)]	200 mg QD (N=6)	400 mg QD (N=6)	800 mg QD (N=6)	Placebo (N=6)	Overall (N=24)
Total Subjects with at Least One TEAE	5 (83.3)	1 (16.7)	2 (33.3)	0	8 (33.3)
Gastrointestinal disorders					
Nausea	1 (16.7)ˇ	1 (16.7)	0	0	2 (8.3)
Abdominal pain upper	1 (16.7)ˇ	0	0	0	1 (4.2)
Diarrhoea	1 (16.7)ˇ	0	0	0	1 (4.2)
Investigations					
Blood creatine phosphokinase increased	0	0	2 (33.3)^	0	2 (8.3)
Neutrophil count decreased	0	0	1 (16.7)^	0	1 (4.2)
White blood cell count decreased	0	0	1 (16.7)^	0	1 (4.2)
Nervous system disorders					
Dizziness	1 (16.7)	0	0	0	1 (4.2)
Headache	0	1 (16.7)	0	0	1 (4.2)
Respiratory, thoracic and mediastinal disorders					
Hyperventilation	1 (16.7)	0	0	0	1 (4.2)
Rhinorrhoea	1 (16.7)	0	0	0	1 (4.2)
General disorders and administration site conditions					
Fatigue	1 (16.7)	0	0	0	1 (4.2)
Immune system disorders					
Drug hypersensitivity	0	0	1 (16.7)	0	1 (4.2)
Infections and infestations					
Urinary tract infection	1 (16.7)	0	0	0	1 (4.2)
Skin and subcutaneous tissue disorders					
Pruritus	1 (16.7)	0	0	0	1 (4.2)
Vascular disorders					
Flushing	1 (16.7)ˇ	0	0	0	1 (4.2)

n = Number of Subjects
ˇOccurred in same subject, ^Occurred in same subject

CONCLUSIONS

- EDP-514 was generally safe and well-tolerated at 200, 400 and 800 mg QD for 28 days in NUC-suppressed CHB patients
- At Day 28, EDP-514 demonstrated reductions in circulating HBV RNA levels, consistent with its mechanism of action as an HBV core inhibitor

REFERENCES

- EDP-514, a novel HBV core inhibitor with potent antiviral activity both *in vitro* and *in vivo*. M Vaine, et al. J Hepatology, VOLUME 70, ISSUE 1, SUPPLEMENT, E474-E475
- EDP-514, a novel pangenotypic class II hepatitis B virus core inhibitor: preliminary results of a phase 1 study in healthy adult subjects. K Larson, et al. J Hepatology, VOLUME 73, SUPPLEMENT 1, S871

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DISCLOSURES

- GDLR, AA, EL, ALC and NA are employees and stockholders of Enanta Pharmaceuticals, Inc.