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ORIGINAL ARTICLE

Clinical Trials Study

Daclatasvir vs telaprevir plus peginterferon alfa/ribavirin for hepatitis C virus genotype 1

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Data sharing statement: Technical appendix, statistical code, and dataset are available from Ira Jacobson at ijacobson@chpnet.org. Patients gave informed consent regarding the relevant use and sharing of key-coded data.

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Abstract

AIM: To evaluate daclatasvir *vs* telaprevir, each

combined with peginterferon alfa-2a/ribavirin (pegIFN/RBV), in treatment-naïve hepatitis C virus (HCV) genotype (GT) 1-infected patients.

METHODS: In this phase 3, randomized, open-label, noninferiority study, 602 patients were randomly assigned (2:1) to daclatasvir *vs* telaprevir, stratified by *IL28B* rs12979860 host genotype (CC *vs* non-CC), cirrhosis status (compensated cirrhosis *vs* no cirrhosis), and HCV GT1 subtype (GT1a *vs* GT1b). Patients were selected by study inclusion criteria from a total of 793 enrolled patients. Patients received daclatasvir 60 mg once daily or telaprevir 750 mg 3 times daily plus pegIFN/RBV. Daclatasvir recipients received 24 wk of daclatasvir plus pegIFN/RBV; those without an extended rapid virologic response (eRVR; undetectable HCV-RNA at weeks 4 and 12) received an additional 24 wk of pegIFN/RBV. Telaprevir-treated patients received 12 wk of telaprevir plus pegIFN/RBV followed by 12 (with eRVR) or 36 (no eRVR) wk of pegIFN/RBV. The primary objective was to compare for noninferiority of sustained virologic response rates at posttreatment week 12 (SVR12) in GT1b-infected patients. Key secondary objectives were to demonstrate that the rates of anemia (hemoglobin < 10 g/dL) and rash-related events, through week 12, were lower with daclatasvir + pegIFN/RBV than with telaprevir + pegIFN/RBV among GT1b-infected patients. Resistance testing was performed using population-based sequencing of the NS5A region for all patients at baseline, and for patients with virologic failure or relapse and HCV-RNA \geq 1000 IU/mL, to investigate any link between NS5A polymorphisms associated with daclatasvir resistance and virologic outcome.

RESULTS: Patient demographics and disease characteristics were generally balanced across treatment arms; however, there was a higher proportion of black/African Americans in the daclatasvir groups (6.0% and 8.2% in the GT1b and GT1a groups, respectively) than in the telaprevir groups (2.2% and 3.0%). Among GT1b-infected patients, daclatasvir plus pegIFN/RBV was noninferior to telaprevir plus pegIFN/RBV for SVR12 [85% (228/268) *vs* 81% (109/134); difference, 4.3% (95%CI: -3.3% to 11.9%)]. Anemia (hemoglobin < 10 g/dL) was significantly less frequent with daclatasvir than with telaprevir [difference, -29.1% (95%CI: -38.8% to -19.4%)]. Rash-related events were also less common with daclatasvir than with telaprevir, but the difference was not statistically significant. In GT1a-infected patients, SVR12 was 64.9% with daclatasvir and 69.7% with telaprevir. Among both daclatasvir and telaprevir treatment groups, across GT1b- or GT1a-infected patients, lower response rates were observed in patients with *IL28B* non-CC and cirrhosis - factors known to affect response to pegIFN/RBV. Consistent with these observations, a multivariate logistic regression analysis in GT1b-infected patients demonstrated that SVR12 was associated with *IL28B* host genotype (CC *vs* non-CC, $P = 0.011$) and cirrhosis status (absent *vs* present, $P = 0.031$). NS5A polymorphisms associated with daclatasvir resistance

(at L28, R30, L31, or Y93) were observed in 17.3% of GT1b-infected patients at baseline; such variants did not appear to be absolute predictors of failure since 72.1% of these patients achieved SVR12 compared with 86.9% without these polymorphisms. Among GT1b-infected patients, treatment was completed by 85.4% (229/268) in the daclatasvir group, and by 85.1% (114/134) in the telaprevir group, and among GT1a-infected patients, by 67.2% (90/134) and 69.7% (46/66), respectively. Discontinuations (of all 3 agents) due to an AE were more frequent with telaprevir than with daclatasvir, whereas discontinuations due to lack of efficacy were more frequent with daclatasvir, due, in part, to differences in futility criteria.

CONCLUSION: Daclatasvir plus pegIFN/RBV demonstrated noninferiority to telaprevir plus pegIFN/RBV for SVR12 and was well-tolerated in treatment-naïve GT1b-infected patients, supporting the use of daclatasvir with other direct-acting antivirals.

Key words: Direct-acting antiviral; Chronic hepatitis C; Daclatasvir; Genotype 1b; NS5A inhibitor; Liver disease

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Core tip: This phase 3 study describes the first prospective comparison of an NS5A inhibitor and an NS3/4A protease inhibitor in peginterferon-based regimens. Combinations of peginterferon alfa-2a/ribavirin (pegIFN/RBV) with boceprevir or telaprevir were the standard-of-care for genotype (GT) 1-infected patients at the time of study design. In treatment-naïve GT1b-infected patients, daclatasvir (NS5A inhibitor) plus pegIFN/RBV achieved a sustained virologic response at posttreatment week 12 (SVR12) of 85% and demonstrated noninferiority to telaprevir plus pegIFN/RBV showing 81% SVR12. Daclatasvir plus pegIFN/RBV was well-tolerated, with a superior safety profile for anemia compared with telaprevir plus pegIFN/RBV. These results support the ongoing investigation of daclatasvir in all-oral combinations in multiple patient populations.

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INTRODUCTION

Chronic infection with hepatitis C virus (HCV) affects

130-150 million people globally and is a major cause of cirrhosis and hepatocellular carcinoma^[1]. Of the 7 HCV genotypes (GTs) identified^[2], GT1 is the most prevalent worldwide, and accounts for 75% of all infections in the United States (US)^[3]. GT1 can be classified into the two main subtypes GT1a and GT1b, of which GT1b is the most common worldwide, predominating in Europe, Japan, and China; in the US 36% of all GT1 infections are subtype 1b^[4].

Peginterferon alfa plus ribavirin (pegIFN/RBV) has traditionally been used to treat HCV. However, this regimen achieves only limited sustained virologic response (SVR) rates of 40%-50%^[5,6] and is associated with a high frequency of adverse events (AEs)^[7]. Although pegIFN/RBV-based therapies continue to be the standard of care in some countries^[8], HCV treatment has evolved toward direct-acting antiviral agents (DAAs) that target specific viral proteins^[5,6], with the first all-oral combinations for GT1 recently approved in Japan, Europe, the US, and Canada^[9-15].

Combinations of pegIFN/RBV with one of the NS3/4A protease inhibitors boceprevir or telaprevir were the first DAA-based regimens approved and at the time of study design, the standard-of-care for GT1-infected patients. With telaprevir plus pegIFN/RBV, SVR rates increased from < 50% with pegIFN/RBV alone to 72%-75% in GT1-infected, treatment-naïve patients^[16-18]. However, skin rash and anemia are frequent, and sometimes severe, adverse events (AEs) observed with telaprevir^[16-19]. In patients treated with telaprevir plus pegIFN/RBV, rash has been reported in 35%-37% of patients (compared with 24% with pegIFN/RBV), necessitating premature discontinuation of telaprevir in 7% of patients, and anemia in 37%-42% (compared with 19% with pegIFN/RBV)^[16-18]. Combinations of pegIFN/RBV with more recent DAAs, such as the NS5B inhibitor sofosbuvir or the NS3/4A protease inhibitor simeprevir, achieved SVR rates of 80%-90% and demonstrated a more favorable safety profile than telaprevir plus pegIFN/RBV^[20-22]. Telaprevir plus pegIFN/RBV has been compared with simeprevir plus pegIFN/RBV in treatment-experienced GT1-infected patients^[23]; however, for treatment-naïve patients, no direct comparison of telaprevir vs a non-protease inhibitor DAA has been performed to date.

Daclatasvir is a potent, once-daily, pangenotypic NS5A inhibitor^[24,25] that has been studied and shown to be well-tolerated in > 13000 patients. In phase 2 trials in treatment-naïve patients infected with GT1-4, daclatasvir + pegIFN/RBV demonstrated greater efficacy than pegIFN/RBV alone^[26,27]. In GT1-infected patients, daclatasvir plus pegIFN/RBV achieved SVR at posttreatment week 24 (SVR24) rates of 60% compared with 38% with pegIFN/RBV; response rates were consistently higher in patients with GT1b (77%) than in those with GT1a (55%)^[27], a finding that has also been observed with other DAA + pegIFN/RBV

combinations^[16,21,28]. Daclatasvir-containing pegIFN-free regimens are approved for treatment of chronic HCV infection in a number of countries: daclatasvir plus asunaprevir (ASV, NS3 inhibitor) was approved as the first all-oral treatment for GT1 in Japan^[9], and daclatasvir plus sofosbuvir (with or without ribavirin) is approved in Europe for GT1, 3, and 4^[10], and in Canada for GT1, 2, and 3^[29]. Daclatasvir is also approved in the US, indicated in combination with sofosbuvir for the treatment of chronic HCV GT3 infection^[30].

This phase 3 COMMAND-3 study compared the safety and efficacy of daclatasvir, an NS5A inhibitor, with that of telaprevir, a protease inhibitor, each in combination with pegIFN/RBV, in treatment-naïve patients with GT1 infection, with a focus on GT1b-infected patients.

MATERIALS AND METHODS

Study design

This was a phase 3, randomized, open-label, non-inferiority study in treatment-naïve patients with GT1 infection (Study AI444-052; ClinicalTrials.gov number NCT01492426). Overall, 602 patients were randomly assigned (2:1) to daclatasvir vs telaprevir, stratified by *IL28B* rs12979860 host genotype (CC vs non-CC), cirrhosis status (compensated cirrhosis vs no cirrhosis), and HCV GT1 subtype (GT1a vs GT1b). Patients with compensated cirrhosis were capped at 25%. GT1a-infected patients were capped at 200 (33%); this cap was introduced during enrollment (May 2012) based on data from a previous daclatasvir phase 2 study indicating that daclatasvir was more effective in GT1b-infected than in GT1a-infected patients^[27]. Patients were treated with daclatasvir 60 mg/d ($n = 402$) or telaprevir 750 mg 3 times/d ($n = 200$) in combination with pegIFN alfa-2a 180 µg once weekly and RBV [weight-based dosing of 1000 mg/d (< 75 kg) or 1200 mg/d (≥ 75 kg)]. Daclatasvir-treated patients with undetectable HCV-RNA at weeks 4 and 12 [extended rapid virologic response (eRVR)] had a planned treatment duration of 24 wk of daclatasvir plus pegIFN/RBV; those without eRVR received an additional 24 wk of pegIFN/RBV (total of 48 wk of therapy), provided they did not experience treatment futility. Telaprevir-treated patients received 12 wk of telaprevir plus pegIFN/RBV, followed by 12 (with eRVR) or 36 (without eRVR) weeks of pegIFN/RBV alone, provided they did not experience treatment futility. Patients in both groups were followed for 24 wk (without eRVR) or 48 wk (with eRVR) posttreatment.

In the daclatasvir group, treatment futility, which mandated discontinuation of all study drugs, was defined as: (1) virologic breakthrough [$> 1\text{-log}_{10}$ increase in HCV-RNA over nadir or confirmed HCV-RNA \geq lower limit of quantification (LLOQ) after confirmed undetectable HCV-RNA while on treatment beginning

at week 2 of therapy]; (2) week 12 HCV-RNA > 1000 IU/mL; or (3) week 24 HCV-RNA \geq LLOQ. In the telaprevir group, treatment futility was defined per prescribing information as week 4 or 12 HCV-RNA > 1000 IU/mL or week 24 HCV-RNA detectable confirmed; virologic breakthrough was not included in the telaprevir futility criteria, per the prescribing information^[19]. Relapse was defined as undetectable HCV-RNA at the end of treatment (EOT) followed by confirmed HCV-RNA \geq LLOQ at any follow-up visit.

Patients

The study included treatment-naïve patients aged 18 years or older, with GT1a or GT1b infection, and HCV-RNA ≥ 10000 IU/mL at screening. Patients with no cirrhosis or compensated cirrhosis [by liver biopsy at any time, or by FibroScanTM (≥ 14.6 kPa) within 1 year of screening] were eligible for inclusion. No previous treatment of HCV with interferon-based regimens or DAAs was allowed. Other exclusion criteria included evidence of decompensated liver disease (including a history or presence of ascites, bleeding varices, or hepatic encephalopathy), evidence of a medical condition contributing to chronic liver disease other than HCV, documented or suspected hepatocellular carcinoma or other malignancies, co-infection with HIV or hepatitis B virus, alanine aminotransferase $\geq 5 \times$ the upper limit of normal, hemoglobin < 12 g/dL (120 g/L) for women and < 13 g/dL (130 g/L) for men, platelet count $< 90 \times 10^9$ cells/L, international normalized ratio ≥ 1.7 , albumin < 3.5 g/dL (35 g/L), or any criterion that would exclude the patient from receiving pegIFN/RBV or telaprevir. Patients were randomized to a regimen within stratum *via* block randomization (block size of 6) and using an interactive voice response system prepared by the sponsor.

Objectives and assessments

Primary and secondary objectives were amended during enrollment to focus on GT1b-infected patients based on the results of previous phase 2 data^[27]. The primary objective was to demonstrate that daclatasvir plus pegIFN/RBV was noninferior to telaprevir plus pegIFN/RBV for SVR at posttreatment week 12 (HCV-RNA $< \text{LLOQ}$ at posttreatment week 12) in GT1b-infected patients. The first two secondary objectives were to demonstrate that the rates of anemia (hemoglobin < 10 g/dL) and of rash-related events, through week 12, were lower with daclatasvir plus pegIFN/RBV than with telaprevir plus pegIFN/RBV, among GT1b-infected patients (see Supplementary Material and Methods for the definition of rash-related events). Additional secondary objectives included noninferiority comparisons between arms of undetectable HCV-RNA at week 4 [rapid virologic response (RVR)], week 12 [complete early virologic response (cEVR)], and weeks 4 and 12 (eRVR), and HCV-RNA $< \text{LLOQ}$ at post-treatment week 24 (SVR24),

in GT1b-infected patients. The final secondary objective was a noninferiority comparison between arms of SVR12 in GT1a-infected patients.

HCV-RNA was assayed using the Roche HCV COBAS® TaqMan® test v2.0 (LLOQ = 25 IU/mL; limit of detection approximately 10 IU/mL). HCV GT and subtype were determined by Versant HCV GT 2.0 assay (LIPA) and were analyzed by ICON Central Laboratories, Inc. *IL28B* genotype was determined by polymerase chain reaction amplification coupled with allelic discrimination. Resistance testing was performed using population-based sequencing of the NS5A region for all patients at baseline, and for patients with virologic failure or relapse and with amplifiable (HCV-RNA \geq 1000 IU/mL) plasma samples. Safety monitoring was based on the incidences of AEs, serious AEs (SAEs), discontinuations due to AEs, laboratory abnormalities, vital signs, and physical examinations.

Statistical analysis

The statistical methods of this study were reviewed by the biometrics group at Bristol-Myers Squibb. A noninferiority margin of -12% was employed in this study. A 2-sided 95% confidence interval (CI) for the difference in rates, daclatasvir plus pegIFN/RBV minus telaprevir plus pegIFN/RBV, was used to test for noninferiority. To demonstrate noninferiority, the lower bound of the CI had to be > -0.12 . A sample size of 400 GT1b-infected patients, randomized 2:1 to daclatasvir plus pegIFN/RBV vs telaprevir plus pegIFN/RBV, provided 91% power to show that the SVR12 rate of daclatasvir plus pegIFN/RBV was noninferior to that of telaprevir plus pegIFN/RBV at the 5% significance level, assuming SVR12 rates of 85% for both regimens^[16].

Each secondary comparison was conducted at the 5% level and proceeded hierarchically according to the order of the objectives (see Objectives and assessments). Testing of an endpoint was performed only if the null hypothesis of the preceding endpoint was rejected. Safety comparisons, for anemia and rash-related events, were for superiority, *i.e.*, to show that daclatasvir plus pegIFN/RBV was less toxic than telaprevir plus pegIFN/RBV. Secondary efficacy comparisons were for noninferiority (noninferiority margin -12%).

Efficacy analyses were restricted to all treated patients and were performed using a modified intent-to-treat (mITT) analysis (patients with missing HCV-RNA measurements were considered failures). For the primary endpoint, an analysis based on SVR documented on or after (if follow-up week 12 HCV-RNA was missing) posttreatment week 12 was also conducted. A stratum-adjusted, 2-sided, asymptotic 95%CI was used to compute the difference in SVR12 rates between arms^[31]. The strata were those used in the randomization. Stratum-adjusted CIs were also used for testing differences between rates for

secondary efficacy endpoints.

All authors had access to the study data and have reviewed and approved the final manuscript.

RESULTS

Patient disposition

Overall, 793 patients were screened; of these, 402 GT1b-infected patients were randomized and treated with daclatasvir plus pegIFN/RBV ($n = 268$) or telaprevir plus pegIFN/RBV ($n = 134$), and 200 GT1a-infected patients were randomized and treated with daclatasvir plus pegIFN/RBV ($n = 134$) or telaprevir plus pegIFN/RBV ($n = 66$) (Figure 1). Among GT1b-infected patients, treatment was completed by 85.4% (229/268) in the daclatasvir group, and by 85.1% (114/134) in the telaprevir group, and among GT1a-infected patients, by 67.2% (90/134) and 69.7% (46/66), respectively. Discontinuations (of all three agents) due to an AE were more frequent with telaprevir than with daclatasvir, whereas discontinuations due to lack of efficacy were more frequent with daclatasvir. This was due, in part, to the different futility criteria in the two treatment groups. Posttreatment follow-up was initiated by 384 patients in the daclatasvir group and by 191 patients in the telaprevir group and was completed by 359 (93.5%) and 181 (94.8%), respectively.

Baseline characteristics

Patient demographics and disease characteristics were generally balanced across treatment arms (Table 1); however, there was a higher proportion of black/African Americans in the daclatasvir groups (6.0%-8.2%) than in the telaprevir groups (2.2%-3.0%). Mean HCV-RNA levels ranged from 6.23-6.31 log₁₀ IU/mL, 9.7%-13.6% of patients had cirrhosis, and there was a high proportion of patients with a non-CC *IL28B* genotype (68.7%-79.9%) across all groups.

Efficacy in GT1b-infected patients

In GT1b-infected patients, SVR12 rates were 85.1% (228/268) with daclatasvir plus pegIFN/RBV vs 81.3% (109/134) with telaprevir plus pegIFN/RBV (mITT; primary endpoint) (Table 2); the difference between treatment arms was 4.3% (95%CI: -3.3% to 11.9%), demonstrating noninferiority (lower bound of the 95%CI greater than -12%). Similar SVR12 rates in GT1b-infected patients were obtained when using the next available HCV-RNA value for patients with missing posttreatment week 12 measurements [daclatasvir plus pegIFN/RBV: 85.8% (230/268); telaprevir plus pegIFN/RBV: 82.1% (110/134)]. Response rates observed for the secondary efficacy endpoints (RVR, eRVR, cEVR, SVR24) in GT1b-infected patients appeared similar between treatment arms (Table 2); however, no formal comparisons could be made because the test for the difference in rash-

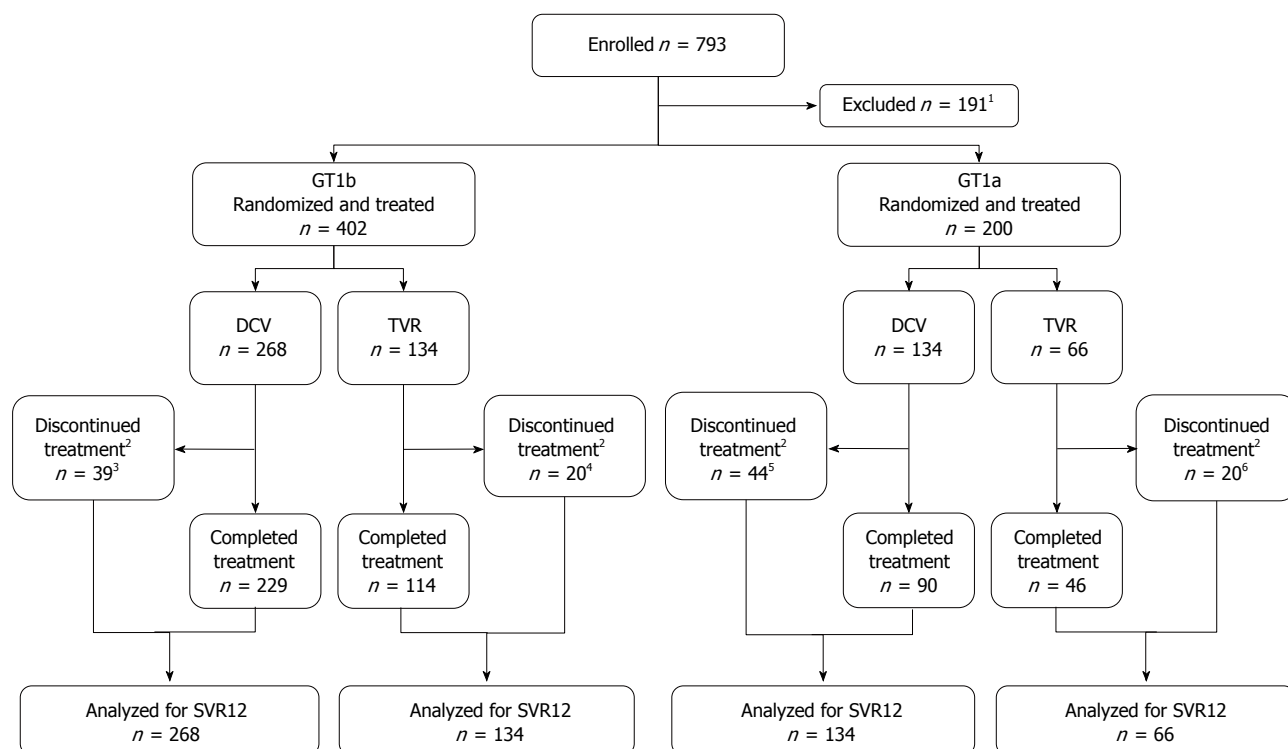


Figure 1 Patient disposition. Efficacy analyses were based on a modified intent-to-treat analysis (included all randomized and treated patients who had received ≥ 1 dose of study medication; patients with missing HCV RNA measurements were considered failures). ¹No longer met study entry criteria during the screening period ($n = 139$), withdrew consent ($n = 28$), administrative reason by sponsor ($n = 5$), lost to follow-up ($n = 3$), pretreatment AE ($n = 2$), or other reasons ($n = 11$); ²All 3 study drugs discontinued; ³Lack of efficacy ($n = 15$; 11 virologic breakthroughs, 3 futility, and 1 other), AE ($n = 14$), patient request ($n = 6$), lost to follow-up ($n = 3$), and withdrew consent ($n = 1$); ⁴AE ($n = 17$), withdrew consent ($n = 2$), and lost to follow-up ($n = 1$); ⁵Lack of efficacy ($n = 23$; 11 virologic breakthroughs, 4 futility, 8 other), adverse event ($n = 11$), lost to follow-up ($n = 6$), patient request ($n = 1$), and other ($n = 3$); ⁶Lack of efficacy ($n = 5$; 3 futility, 2 other), AE ($n = 8$), patient request ($n = 3$), lost to follow-up ($n = 3$), and death ($n = 1$; the death occurred in a patient who had discontinued treatment at week 16 due to bacteremia and died at posttreatment week 4 due to sepsis secondary to HCV-related cirrhosis). DCV: Daclatasvir; GT: Genotype; SVR12: Sustained virologic response (HCV-RNA < LLOQ) at posttreatment week 12; TVR: Telaprevir.

Table 1 Patient demographics and baseline disease characteristics

n (%)	GT1b		GT1a	
	DCV + pegIFN/RBV (n = 268)	TVR + pegIFN/RBV (n = 134)	DCV + pegIFN/RBV (n = 134)	TVR + pegIFN/RBV (n = 66)
Age (yr), median (range)	46.0 (18-71)	48.0 (19-69)	49.0 (19-67)	51.5 (28-69)
Male	159 (59.3)	72 (53.7)	98 (73.1)	47 (71.2)
Race				
White	243 (90.7)	129 (96.3)	120 (89.6)	63 (95.5)
Black/African American	16 (6.0)	3 (2.2)	11 (8.2)	2 (3.0)
Asian	6 (2.2)	2 (1.5)	1 (0.7)	0
Other	3 (1.1)	0	2 (1.5)	1 (1.5)
HCV-RNA log ₁₀ (IU/mL), mean (SD)	6.23 (0.701)	6.23 (0.577)	6.30 (0.637)	6.31 (0.636)
HCV-RNA ≥ 800000 IU/mL	196 (73.1)	97 (72.4)	104 (77.6)	51 (77.3)
IL28B genotype				
CC	53 (19.8)	27 (20.1)	42 (31.3)	20 (30.3)
CT	161 (60.1)	86 (64.2)	73 (54.5)	37 (56.1)
TT	53 (19.8)	21 (15.7)	19 (14.2)	9 (13.6)
Not reported	1 (0.4)	0	0	0
Cirrhosis				
Present	26 (9.7)	15 (11.2)	16 (11.9)	9 (13.6)

DCV: Daclatasvir; GT: Genotype; HCV: Hepatitis C virus; pegIFN: Peginterferon alfa-2a; RBV: Ribavirin; TVR: Telaprevir.

related events between arms, which preceded the comparisons of secondary efficacy endpoints in the testing hierarchy, was not statistically significant (see

Safety section). Among GT1b-infected patients with SVR12, three patients did not achieve SVR24 (one patient in each arm relapsed between posttreatment

Table 2 Efficacy endpoints and failures in GT1b-infected patients

Outcome, n/n (%)	DCV + pegIFN/RBV	TVR + pegIFN/RBV
Efficacy		
SVR12 (mITT) ^{1,2}	228/268 (85.1)	109/134 (81.3)
SVR12 on or after PT week 12 ³	230/268 (85.8)	110/134 (82.1)
RVR (HCV-RNA undetectable at week 4) ^{1,4}	207/268 (77.2)	106/134 (79.1)
cEVR (HCV-RNA undetectable at week 12) ^{1,4}	243/268 (90.7)	121/134 (90.3)
eRVR (HCV-RNA undetectable at weeks 4 and 12) ^{1,4}	201/268 (75.0)	98/134 (73.1)
EOTR (HCV-RNA undetectable at EOT)	244/268 (91.0)	131/134 (97.8)
SVR24 ^{1,4}	226/268 (84.3)	108/134 (80.6)
Failures		
Non-SVR12	40/268 (14.9)	25/134 (18.7)
On-treatment failures	21/268 (7.8)	3/134 (2.2)
Virologic breakthrough	11/268 (4.1)	NA ⁵
Treatment futility other than virologic breakthrough	3/268 (1.1)	0
HCV-RNA detectable at EOT	7/268 (2.6)	3/134 (2.2)
Posttreatment relapse ⁶	12/244 (4.9)	20/131 (15.3)
HCV-RNA undetectable at EOT but missing PT week 12 data	7/244 (2.9)	2/131 (1.5)

¹Patients with missing data at posttreatment week 12 were considered failures; ²Difference, 4.3%; 95%CI: -3.3% to 11.9%; noninferior at the 5% significance level (lower bound of the 95%CI was > -12%); ³Patients with missing data at posttreatment week 12 were considered responders if the next available HCV-RNA value was < LLOQ; ⁴Differences and 95%CI were as follows: RVR, -1.5% (-9.8% to 6.8%); cEVR, 0.6 (-5.5% to 6.6%); eRVR, 2.2% (-6.8% to 11.2%); SVR24, 4.4% (-3.5% to 12.2%); ⁵Not included in TVR futility criteria per prescribing information; ⁶Assessed in patients with undetectable HCV-RNA at EOT. cEVR: Complete early virologic response; DCV: Daclatasvir; EOT: End of treatment; EOTR: End of treatment response; eRVR: Early rapid virologic response; GT: Genotype; HCV: Hepatitis C virus; LLOQ: Lower limit of quantitation; mITT: Modified intent-to-treat; NA: Not applicable; pegIFN: Peginterferon alfa-2a; PT: Posttreatment; RBV: Ribavirin; SVR12: Sustained virologic response (HCV-RNA < LLOQ) at posttreatment week 12; RVR: Rapid virologic response; SVR24: Sustained virologic response at posttreatment week 24; TVR: Telaprevir.

Table 3 Proportion of GT1b-infected patients with SVR12 by subgroups

SVR12 ¹ , n/n (%)	DCV + pegIFN/RBV (n = 268)	TVR + pegIFN/RBV (n = 134)
Age (yr)		
< 65	218/256 (85.2)	103/126 (81.7)
≥ 65	10/12 (83.3)	6/8 (75.0)
Sex		
Male	134/159 (84.3)	61/72 (84.7)
Female	94/109 (86.2)	48/62 (77.4)
Race		
White	208/243 (85.6)	105/129 (81.4)
Black/African American	11/16 (68.8)	2/3 (66.7)
Asian	6/6 (100.0)	2/2 (100.0)
Other	3/3 (100.0)	0
Baseline HCV-RNA		
< 800000 IU/mL	66/72 (91.7)	33/37 (89.2)
≥ 800000 IU/mL	162/196 (82.7)	76/97 (78.4)
Cirrhosis		
Absent	208/242 (86.0)	99/119 (83.2)
Present	20/26 (76.9)	10/15 (66.7)
IL28B genotype		
CC	51/53 (96.2)	23/27 (85.2)
CT	132/161 (82.0)	69/86 (80.2)
TT	44/53 (83.0)	17/21 (81.0)

¹mITT analysis (patients with missing data at posttreatment week 12 were considered failures). DCV: Daclatasvir; GT: Genotype; HCV: Hepatitis C virus; pegIFN: Peginterferon alfa-2a; RBV: Ribavirin; SVR12: Sustained virologic response (HCV-RNA < LLOQ) at posttreatment week 12; TVR: Telaprevir.

weeks 12 and 24, and one patient in the daclatasvir group had a missing posttreatment week 24 HCV-RNA measurement).

In GT1b-infected patients, observed SVR12 rates

tended to be higher in the daclatasvir arm than in the telaprevir arm in subgroups based on demographic and disease status, such as age, sex, *IL28B*, cirrhosis, and baseline HCV-RNA level (Table 3). Of note, among cirrhotics, a higher proportion of patients receiving daclatasvir vs telaprevir achieved SVR12 (76.9% vs 66.7%). In the daclatasvir group, SVR12 appeared to be independent of age and sex. In both treatment groups, lower response rates were observed in patients with *IL28B* non-CC, baseline HCV-RNA ≥ 800000 IU/mL, or cirrhosis, baseline factors known to affect response to pegIFN/RBV^[32]. Consistent with these observations, a multivariate logistic regression analysis in GT1b-infected patients demonstrated that SVR12 was associated with *IL28B* host genotype (CC vs non-CC, *P* = 0.011), baseline HCV-RNA (< 800000 IU/mL vs ≥ 800000 IU/mL, *P* = 0.016), and cirrhosis status (absent vs present, *P* = 0.031). Virologic response was not associated with age, sex, race, or type of treatment (Supplementary Table 3).

Virologic failure in GT1b-infected patients

Forty (14.9%) patients in the daclatasvir group and 25 (18.7%) patients in the telaprevir group did not achieve SVR12 (Table 2). Virologic breakthrough occurred in 11 (4.1%) patients in the daclatasvir group. Virologic breakthrough was not assessed as a futility criteria for the telaprevir group, per the telaprevir prescribing information^[19]. Relapse (among patients with undetectable HCV-RNA at EOT) was reported in 12/244 (4.9%) patients in the daclatasvir group and 20/131 (15.3%) patients in the telaprevir group. Other posttreatment failures, occurring in

Table 4 Overall on-treatment safety in GT1-infected patients

Event, <i>n</i> (%)	DCV + pegIFN/RBV (<i>n</i> = 402)	TVR + pegIFN/RBV (<i>n</i> = 200) ¹
Death	1 (0.2) ²	1 (0.5) ²
SAEs	26 (6.5) ²	20 (10.0) ²
AEs leading to discontinuation of any study drug	28 (7.0) ²	37 (18.5) ²
AEs leading to discontinuation of all 3 study drugs	25 (6.2)	25 (12.5)
AEs (grade 1-4) ≥ 20%		
Fatigue	140 (34.8)	81 (40.5)
Headache	137 (34.1)	57 (28.5)
Asthenia	109 (27.1)	53 (26.5)
Pruritus	107 (26.6)	75 (37.5)
Anemia	96 (23.9)	99 (49.5)
Rash	93 (23.1)	69 (34.5)
Nausea	88 (21.9)	74 (37.0)
Neutropenia	87 (21.6)	27 (13.5)
Alopecia	86 (21.4)	32 (16.0)
Influenza-like illness	85 (21.1)	38 (19.0)
Dry skin	84 (20.9)	34 (17.0)
Pyrexia	80 (19.9)	42 (21.0)
Grade 3 or 4 emergent laboratory abnormalities		
Hemoglobin	26 (6.5)	41 (20.5)
Absolute neutrophil count	104 (25.9)	41 (20.5)
Lymphocytes	67 (16.7)	44 (22.0)
Platelet count	15 (3.7)	8 (4.0)
ALT	3 (0.7)	4 (2.0)
AST	7 (1.7)	1 (0.5)
Total bilirubin	4 (1.0)	6 (3.0)
Serum creatinine increased	1 (0.2)	0

¹*n* = 198 for grade 3/4 laboratory abnormalities; ²Further information provided in supplementary material. AE: Adverse event; ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; DCV: Daclatasvir; GT: Genotype; pegIFN: Peginterferon alfa-2a; RBV: Ribavirin; SAE: Serious adverse event; TVR: Telaprevir.

7/244 (2.9%) patients in the daclatasvir group and 2/131 (1.5%) patients in the telaprevir group, were due to undetectable HCV-RNA at EOT but missing posttreatment week 12 HCV-RNA.

At baseline, 249/268 GT1b-infected patients treated with daclatasvir plus pegIFN/RBV had available NS5A population-based sequencing data. In 43/249 (17.3%) of these patients, one or more of the NS5A polymorphisms L28M/V, R30H/Q, L31M, or Y93H were detected at baseline; of these, 72.1% (31 patients) achieved SVR12, of which 61.3% (19 patients) had a non-CC *IL28B* genotype. Of the remaining 12 patients who did not achieve SVR12, 11 had a non-CC *IL28B* genotype. Among patients without NS5A polymorphisms at baseline, 87% (179/206) achieved SVR12.

Among the 40 GT-1b patients in the daclatasvir group who did not achieve SVR12, 32 had evaluable samples at baseline and at the time of failure. In two patients, the same NS5A resistance-associated variants (RAVs) were detected at both baseline and failure: L31M-Y93H in one patient, and L28V-R30Q-L31M-Q62D in a second patient. Among the remaining 30 patients, the most common treatment-emergent NS5A RAVs were L31F/I/M/V (22 patients) and Y93H (21 patients); RAVs at L31 and Y93 emerged together in 18 patients. NS5A L31 and Y93 RAVs also emerged together in one daclatasvir-treated patient who achieved SVR12 but relapsed at posttreatment

week 24. Of the eight non-SVR12 patients who were not tested at failure [due to undetectable HCV-RNA at last available visit (*n* = 5), missing HCV-RNA measurement (*n* = 2), or lost to follow-up while HCV-RNA was < 1000 IU/mL (*n* = 1)], seven had no NS5A polymorphisms at baseline, and one had no baseline NS5A sequence available.

Efficacy outcomes in GT1a-infected patients

In GT1a-infected patients, SVR12 rates were 64.9% in the daclatasvir group and 69.7% in the telaprevir group. Other outcomes and treatment failures in GT1a-infected patients are shown in Supplementary Table 1. Baseline NS5A sequences were available for 123 GT1a-infected patients treated with daclatasvir; the polymorphisms Q30R, L31M, and/or Y93N were detected in 6 (5%) patients, of whom 5 achieved SVR12. Among 34 GT1a-infected virologic failures with evaluable samples, the most common emergent NS5A RAVs at failure were Q30E/H/R (32 patients). In patients with emergent Q30 RAVs, L31 variants were also frequently detected, either as on-treatment emergent RAVs (19 patients) or pre-existing at baseline (one patient). In a daclatasvir-treated patient who relapsed after achieving SVR12, the NS5A RAV Q30E was detected at posttreatment week 24.

Safety

Safety, pooled across GT1 subtypes, is shown in

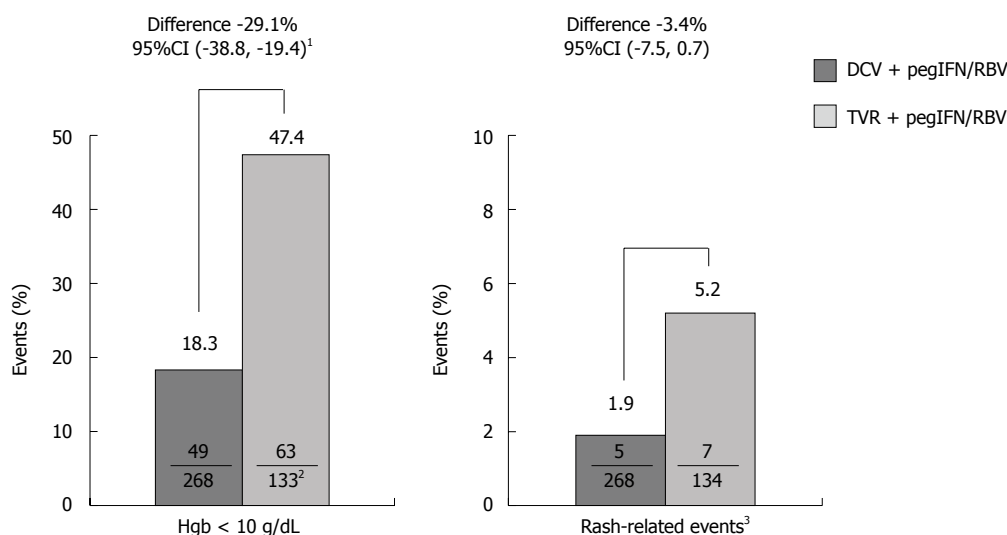


Figure 2 Safety secondary endpoints in GT1b-infected patients (first 12 wk). mITT analysis (patients with missing data at posttreatment week 12 were considered failures). ¹Superior at the 5% significance level; upper bound of the 95%CI was 0%; ²One patient in the TVR arm had no measurement due to discontinuation prior to on-treatment laboratory assessment; ³Includes rash-related SAEs, AEs leading to discontinuation, and grade 3 or 4 AEs. AE: Adverse event; CI: Confidence interval; DCV: Daclatasvir; GT: Genotype; Hgb: Hemoglobin; pegIFN: Peginterferon alfa-2a; RBV: Ribavirin; SAE: Serious adverse event; TVR: Telaprevir.

Table 4. AEs leading to discontinuation of any study drug were more frequent with telaprevir (18.5%) vs daclatasvir (7.0%). In the daclatasvir group, the most common ($\geq 1\%$) AEs leading to discontinuation of any study drug were psychiatric (1.5%), skin (1.5%), or hemolytic events (1.2%); in the telaprevir group the most common AEs were skin events (9.0%), hemolytic events (4.5%), general disorders (3.0%), gastrointestinal events (2.0%), psychiatric events (1.5%), infections and infestations (1.5%), nervous system disorders (1.5%), or renal events (1.0%). Serious AEs (SAEs) were reported in 6.5% of daclatasvir-treated patients and in 10.0% of telaprevir-treated patients. Drug-related SAEs were reported in 3.5% ($n = 14$) and 8.0% ($n = 16$) of treated patients in the daclatasvir and telaprevir arms, respectively, including one case of drug reaction with eosinophilia and systemic symptoms (DRESS) syndrome in each treatment group and five cases of anemia in the telaprevir group. The daclatasvir-treated patient with DRESS syndrome, a 62-year-old male with GT1b-infection, developed grade 3 rash after approximately 2.5 mo of therapy, followed by hyperthermia and hypereosinophilia (eosinophils 1.12×10^9 cells/L) leading to hospitalization. Topical corticosteroid (betamethasone) and trimeprazine resulted in rapid improvement and normalization of the eosinophil count (0.41×10^9 cells/L) within 10 d of discontinuing study therapy. The patient achieved SVR12. The five telaprevir-treated patients with a related SAE of anemia all had grade 4 anemia with hemoglobin levels of 54-72 g/L; all 5 patients received blood transfusions and 2 discontinued study therapy. The most common (> 25%) AEs were fatigue, headache, asthenia, and pruritus in the daclatasvir group, and anemia, fatigue, nausea, rash, pruritus, headache, and asthenia in the

telaprevir group. Grade 3 or 4 laboratory abnormalities were comparable between treatment groups, except for grade 3 or 4 hemoglobin levels, which were more frequent with telaprevir (20.5%) than with daclatasvir (6.5%). Grade 3 or 4 bilirubin elevations occurred in 3% of patients in the telaprevir group compared with 1% in the daclatasvir group.

One death was reported in each treatment group, both of which occurred during off-treatment follow-up and were considered unrelated to therapy. In the daclatasvir group, the patient died during posttreatment week 4 from multiple fractures and a subdural hematoma related to a fall. In the telaprevir group, the patient, who had discontinued treatment at week 16 (due to bacteremia), died during posttreatment week 4 from sepsis secondary to cirrhosis due to hepatitis C.

Grade 1-4 anemia was experienced by 23.9% of GT1-infected patients in the daclatasvir group, and 49.5% in the telaprevir group, including 3.2% ($n = 13$) and 13.5% ($n = 27$), respectively, with grade 3 or 4 anemia. Incidences for the secondary endpoint of hemoglobin < 10 g/dL in GT1b-infected patients through week 12 were significantly lower in the daclatasvir group than in the telaprevir group (daclatasvir: 18.3% vs telaprevir: 47.4%; difference, -29.1; 95%CI: -38.8% to -19.4%; Figure 2). Grade 1-4 rash [single Medical Dictionary for Regulatory Activities (MedDRA) term] was reported in 23.1% of GT1-infected patients in the daclatasvir group and in 34.5% in the telaprevir group, including 1.0% ($n = 4$) and 3.5% ($n = 7$), respectively, with grade 3 or 4 rash. Incidences for the secondary endpoint of rash-related events (composite MedDRA term) in GT1b-infected patients through week 12 were also lower with daclatasvir than with telaprevir, however the difference was not statistically significant given the

low event rates in both arms (daclatasvir: 1.9%; telaprevir: 5.2%; difference, -3.4; 95%CI: -7.5% to 0.7%; Figure 2).

Safety was also assessed in subgroups of patients with and without cirrhosis at baseline (Supplementary Table 2). In both treatment arms, anemia, reduced platelet count, and elevated total bilirubin were more frequent in patients with than in those without cirrhosis. Among cirrhotic patients, those treated with telaprevir experienced a higher frequency of the AEs anemia (58.3% vs 33.3%), rash (41.7% vs 21.4%), fatigue (50.0% vs 38.1%), and nausea (37.5% vs 11.9%), and of SAEs (16.7% vs 7.1%); in contrast, neutropenia was more common with daclatasvir (26.2%) than with telaprevir (16.7%) but was not associated with an increase in infections (daclatasvir: 21.4%; telaprevir: 41.7%).

DISCUSSION

This study evaluated the efficacy and safety of daclatasvir plus pegIFN/RBV compared with telaprevir plus pegIFN/RBV in HCV GT1-infected patients and is the first head-to-head comparison of two different DAA classes - an NS5A inhibitor vs an NS3/4A protease inhibitor - in combination with pegIFN/RBV in treatment-naive patients. In treatment-experienced GT1-infected patients simeprevir (NS3/4A protease inhibitor) plus pegIFN/RBV has been compared with telaprevir plus pegIFN/RBV, with simeprevir plus pegIFN/RBV demonstrating noninferiority to telaprevir plus pegIFN/RBV for SVR12 (54% vs 55%) and a more favourable safety profile^[23].

In the present study in treatment-naive, GT1b-infected patients, SVR12 rates achieved with daclatasvir plus pegIFN/RBV were noninferior to those observed with telaprevir plus pegIFN/RBV, and anemia was significantly less frequent among daclatasvir-treated patients than in telaprevir-treated patients. Rash-related events were also less common among patients receiving daclatasvir vs telaprevir. Daclatasvir plus pegIFN/RBV demonstrated high SVR12 rates in GT1b-infected patients across all subgroups of baseline factors known to affect response rates to pegIFN/RBV (cirrhosis, *IL28B* genotype, age, sex, baseline viral load). Importantly, in difficult-to-cure patients with cirrhosis, SVR12 rates were higher with daclatasvir plus pegIFN/RBV than with telaprevir plus pegIFN/RBV (76.9% vs 66.7%). On-treatment treatment futility was more common with daclatasvir than with telaprevir; however this may have been related to the different futility criteria between the two treatment groups. However, posttreatment relapse was more frequent with telaprevir than with daclatasvir (15% vs 5%).

Using population-based sequencing of the NS5A region, NS5A polymorphisms associated with daclatasvir resistance (L28, R30, L31, and/or Y93) were observed

in 17% of GT1b-infected patients at baseline. Although such variants may be associated with virologic outcome to daclatasvir plus pegIFN/RBV, they do not appear to be absolute predictors of failure because 72% of GT1b-infected patients with these polymorphisms at baseline achieved SVR12 in this study. Furthermore, since most (11/12) patients with baseline NS5A polymorphisms who did not achieve SVR12 also had a non-CC *IL28B* genotype, it may be that *IL28B* genotype has a stronger association with virologic failure. A similar observation was reported in a previous daclatasvir plus pegIFN/RBV phase 2 study, in which only 2/10 GT1b-infected patients with L31 or Y93 variants failed to achieve a response, but 30/32 GT1b virologic failures were non-CC *IL28B*^[27].

A relevant consideration is that the persistence of RAVs may influence further treatment options in an IFN-free context. In this study, patients were not monitored beyond post-treatment week 24, thus analysis of the persistence of NS5A RAVs was not undertaken. However, persistence beyond one year has been described previously in GT1b- and GT1a-infected patients^[33,34]. Persistence of emergent NS5A RAVs has also been described from an interim analysis of a three-year follow-up study^[35]. Detection of viral variants by direct sequencing limits the ability to detect low-frequency variants within a viral population; however, despite this limitation, the results of these analyses have been useful for describing relationships between RAVs and clinical outcomes.

In this study, response rates with daclatasvir plus pegIFN/RBV were lower and virologic failure was more frequent in patients infected with HCV GT1a than in patients infected with GT1b. Most likely, this is due to a lower resistance barrier of daclatasvir in GT1a-infected patients rather than to a higher potency in GT1b-infected patients, since the EC₅₀ of daclatasvir was low in both GT1a-infected patients (20 pM) and GT1b-infected patients (4 pM)^[24]. These lower response rates observed among GT1a-infected patients are consistent with previous studies of daclatasvir^[27] and have also been observed with other DAAs, including telaprevir^[16], boceprevir^[28], and simeprevir^[21].

Daclatasvir plus pegIFN/RBV was generally well-tolerated, with an overall safety profile similar to that of pegIFN/RBV alone, with no new safety or tolerability concerns attributable to daclatasvir. In the overall GT1-infected population, anemia events (grade 1-4 and grade 3 or 4) were less common in the daclatasvir group (24% and 3%, respectively) than in the telaprevir group (50% and 14%, respectively), and were also less frequently a reason for treatment discontinuation (daclatasvir: 0.5%; telaprevir: 4.0%). Among GT1b-infected patients, daclatasvir plus pegIFN/RBV demonstrated superiority over telaprevir plus pegIFN/RBV for the secondary endpoint of hemoglobin < 10 g/dL (18% vs 47%).

Grade 1-4 and grade 3 or 4 rash (single MedDRA

term) were also less frequent in the overall GT1-infected population in patients receiving daclatasvir (23% and 1%, respectively) than in those receiving telaprevir (35% and 4%, respectively), as were discontinuations due to rash (daclatasvir: 0.2%; telaprevir: 4.5%). The grade 1-4 events of rash observed in the daclatasvir group are most likely related to pegIFN/RBV, because the observed rate (23%) is comparable to that historically reported with pegIFN/RBV alone^[16,27], and no cases of rash have been reported with daclatasvir all-oral regimens, including daclatasvir plus sofosbuvir^[36]. Rash-related events (composite MedDRA term) were evaluated as a secondary endpoint in this study in GT1b-infected patients, and a lower incidence was observed in those receiving daclatasvir than in those receiving telaprevir, but this difference was not statistically significant due to low event rates in both arms. The single case of DRESS syndrome observed with daclatasvir is the only case reported to date in the daclatasvir development program and real-world experience.

At the time this study was designed, telaprevir plus pegIFN/RBV was the standard of care for the treatment of HCV GT1 infection; however, other DAAs have now been approved, both in combination with pegIFN/RBV and as part of IFN-free regimens. Currently approved all-oral treatment options for GT1-infected patients include daclatasvir plus sofosbuvir ± RBV (approved for GT1, 3, and 4 in the EU^[10]), which has been shown to achieve SVR12 rates of up to 98% in treatment-naïve GT1-infected patients^[36], and the dual combination daclatasvir plus asunaprevir (approved for GT1b in Japan^[9]), which has been shown to achieve SVR12 rates of 90% in GT1b-infected treatment-naïve patients^[37]. Other approved all-oral regimens include the combinations of sofosbuvir + ledipasvir ± RBV^[11,12] and simeprevir plus sofosbuvir ± RBV^[13,14,38], which achieved SVR rates of up to 98% and 94%, respectively, in this patient population. Ombitasvir/paritaprevir/ritonavir plus dasabuvir, which was recently approved, provided SVR rates of 90% (without RBV) and 95%-97% (with RBV) in treatment-naïve noncirrhotic patients infected with GT1a, and of 99% (without RBV) and 98%-100% (with RBV) in those infected with GT1b^[36,39,40]; in GT1-infected patients with cirrhosis SVR12 was 96%^[41]. Daclatasvir is also being evaluated as part of an all-oral, fixed-dose combination with asunaprevir and beclabuvir (formerly BMS-791325). In phase 3 studies, this regimen without RBV has provided SVR12 rates of 92% in GT1a- and 1b-infected treatment-naïve patients without cirrhosis^[42], and SVR12 rates of 93%-98% in GT1a- and 1b-infected patients with compensated cirrhosis, with or without RBV^[43]; in these studies, SVR12 rates were lower in patients infected with GT1a than in those infected with GT1b when RBV was excluded from the regimen.

Although all-oral regimens have become the new

standard of care in chronic hepatitis C management, pegIFN/RBV-based therapies will remain a potentially important treatment option in such settings as low-income countries or possibly for patients who have failed DAA-only therapies.

In conclusion, this first and only head-to-head comparison of two classes of DAAs in treatment-naïve patients demonstrates that daclatasvir plus pegIFN/RBV is noninferior to telaprevir plus pegIFN/RBV for SVR12 for treatment of HCV GT1b infection. Daclatasvir plus pegIFN/RBV was generally well-tolerated, with a significantly lower rate of anemia, and an observed lower rate of rash-related events compared with telaprevir plus pegIFN/RBV. The results of this study and other studies support the role of daclatasvir as an effective and well-tolerated component of interferon-free, all-oral HCV regimens across multiple genotypes.

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COMMENTS

Background

Chronic infection with hepatitis C virus (HCV) affects 130-150 million people globally and is a major cause of cirrhosis and hepatocellular carcinoma. Peginterferon alfa plus ribavirin (pegIFN/RBV) has historically been used to treat HCV, although this regimen achieves only limited sustained virologic response (SVR) rates of 40%-50% in genotype 1 (GT1) infection. Whilst pegIFN/RBV-based therapies remain the standard of care in some countries, HCV treatment has evolved toward direct-acting antiviral agents (DAAs) that target specific viral proteins.

Research frontiers

The combination of pegIFN/RBV with the NS3/4A protease inhibitor telaprevir was one of the first approved DAA-based regimens, and replaced pegIFN/RBV as the standard-of-care for GT1-infected patients. Skin rash and anemia are frequent, and at times severe, adverse events (AEs) observed with telaprevir. No direct comparison of telaprevir vs a non-protease inhibitor DAA has been performed to date. Daclatasvir is a potent, once-daily, pangenotypic NS5A inhibitor; this phase 3 study compared the safety and efficacy of daclatasvir with that of telaprevir, each in combination with pegIFN/RBV, in treatment-naïve patients with GT1 infection, with a focus on GT1b-infected patients.

Innovations and breakthroughs

Among GT1b-infected patients, daclatasvir plus pegIFN/RBV was noninferior to telaprevir plus pegIFN/RBV for SVR12 [85% (228/268) vs 81% (109/134); difference, 4.3% (95%CI: -3.3% to 11.9%)]. Daclatasvir was generally well-tolerated: anemia (hemoglobin < 10 g/dL) was significantly less frequent with daclatasvir than with telaprevir [difference, -29.1% (95%CI: -38.8% to -19.4%)], and rash-related events were less frequent with daclatasvir plus pegIFN/RBV compared with telaprevir plus pegIFN/RBV. Discontinuations of all 3 agents due to an adverse event were more frequent with telaprevir than with daclatasvir, whereas discontinuations due to lack of efficacy were more frequent with daclatasvir, due, in part, to differences in futility criteria.

Applications

Although all-oral regimens have become the new standard of care in chronic HCV management - for which daclatasvir is currently approved for the treatment of HCV GT1, GT3, and GT4 infections (approvals vary by country) - pegIFN/

RBV-based therapies will remain a potentially important treatment option in such settings as low-income countries or possible for patients who have failed DAA-only therapies. The results of this study and other studies support the role of daclatasvir as an effective and well-tolerated component of interferon-free, all-oral HCV regimens across multiple genotypes.

Terminology

SVR12/24, sustained virologic response [HCV-RNA < lower limit of quantitation (LLOQ)] at posttreatment week 12/24; Relapse, undetectable HCV-RNA at the end of treatment followed by confirmed HCV-RNA \geq LLOQ at any follow-up visit. In the daclatasvir group, treatment futility, which mandated discontinuation of all study drugs, was defined as (1) virologic breakthrough ($> 1\text{-log}_{10}$ increase in HCV-RNA over nadir or confirmed HCV-RNA \geq LLOQ after confirmed undetectable HCV-RNA while on treatment beginning at week 2 of therapy); (2) week 12 HCV-RNA > 1000 IU/mL; or (3) week 24 HCV-RNA \geq LLOQ. In the telaprevir group, treatment futility was defined per prescribing information as HCV-RNA > 1000 IU/mL (at weeks 4 or 12) or HCV-RNA detectable confirmed (at week 24).

Peer-review

The arena within which HCV is treated has moved at a very rapid pace over the past 5 years. The range of DAA options has expanded since telaprevir and boceprevir were the first protease inhibitors to become available. While globally, the world is focused on DAAs as the treatments of choice for HCV, these drugs are expensive and not all countries will be in a position to treat infection vs disease. In fact, some first-world countries are currently only treating patients with advanced disease due to the cost of these drugs. The authors make this point nicely and rightly state that there may be a place for IFN based anti-viral strategies for the management and treatment of HCV.

REFERENCES

- World Health Organization.** Hepatitis C key facts. WHO factsheet No 164. (Updated April 2014, accessed June 23, 2015). Available from: URL: <http://www.who.int/mediacentre/factsheets/fs164/en/>
- Smith DB,** Bukh J, Kuiken C, Muerhoff AS, Rice CM, Stapleton JT, Simmonds P. Expanded classification of hepatitis C virus into 7 genotypes and 67 subtypes: updated criteria and genotype assignment web resource. *Hepatology* 2014; **59**: 318-327 [PMID: 24115039 DOI: 10.1002/hep.26744]
- Messina JP,** Humphreys I, Flaxman A, Brown A, Cooke GS, Pybus OG, Barnes E. Global distribution and prevalence of hepatitis C virus genotypes. *Hepatology* 2015; **61**: 77-87 [PMID: 25069599 DOI: 10.1002/hep.27259]
- Gower E,** Estes C, Blach S, Razavi-Shearer K, Razavi H. Global epidemiology and genotype distribution of the hepatitis C virus infection. *J Hepatol* 2014; **61**: S45-S57 [PMID: 25086286 DOI: S0168-8278(14)00526-1]
- European Association for Study of Liver.** EASL Clinical Practice Guidelines: management of hepatitis C virus infection. *J Hepatol* 2014; **60**: 392-420 [PMID: 24331294 DOI: 10.1016/j.jhep.2013.11.003]
- American Association for the Study of Liver Diseases and the Infectious Diseases Society of America.** Recommendations for testing, managing, and treating hepatitis C. (Updated July 7, 2014, accessed June 23, 2015). Available from: URL: <http://www.hcvguidelines.org/fullreport7>
- Ghany MG,** Strader DB, Thomas DL, Seeff LB. Diagnosis, management, and treatment of hepatitis C: an update. *Hepatology* 2009; **49**: 1335-1374 [PMID: 19330875 DOI: 10.1002/hep.22759]
- Kretzer IF,** do Livramento A, da Cunha J, Gonçalves S, Tosin I, Spada C, Treitinger A. Hepatitis C worldwide and in Brazil: silent epidemic—data on disease including incidence, transmission, prevention, and treatment. *ScientificWorldJournal* 2014; **2014**: 827849 [PMID: 25013871 DOI: 10.1155/2014/827849]
- Bristol-Myers Squibb.** Japan approves first all-oral, interferon- and ribavirin-free hepatitis C treatment, Daklinza® (daclatasvir) and Sunvepra® (asunaprevir) dual regimen. (Updated July 7, 2014, accessed June 23, 2015). Available from: URL: <http://news.bms.com/press-release/japan-approves-first-all-oral-interferon-and-ribavirin-free-hepatitis-c-treatment-dakl>
- Bristol-Myers Squibb.** European Commission approves Bristol-Myers Squibb's Daklinza (daclatasvir) across multiple genotypes for the treatment of chronic hepatitis C infection. (Updated August 27, 2014, accessed June 23, 2015). Available from: URL: <http://news.bms.com/press-release/rd-news/european-commission-approves-bristol-myers-squibbs-daklinza-daclatasvir-across>
- Gilead Sciences, Inc.** Harvoni (ledipasvir and sofosbuvir) prescribing information. (Updated March 2015, accessed June 23 2015). Available from: URL: http://www.gilead.com/~media/Files/pdfs/medicines/liverdisease/harvoni/harvoni_pi.pdf
- Gilead Sciences, Inc.** Harvoni (ledipasvir and sofosbuvir) [summary of product characteristics]. Cambridge, United Kingdom: Gilead Sciences International Ltd, 2014
- Janssen Therapeutics.** OLYSIO™ (simeprevir) prescribing information. (Updated April 2015, accessed June 23, 2015). Available from: URL: <https://www.oly시오.com/shared/product/olysio/prescribing-information.pdf>
- Janssen Therapeutics.** Olysio (simeprevir) summary of product characteristics. (Updated April 7, 2015, accessed June 23, 2015). Available from: URL: http://ec.europa.eu/health/documents/community-register/2014/20140514128513/anx_128513_en.pdf
- AbbVie, Inc.** Viekira Pak (ombitasvir, paritaprevir, and ritonavir tablets; dasabuvir tablets) prescribing information. (Updated March 2015, accessed June 23, 2015). Available from: URL: http://www.rxabbvie.com/pdf/viekirapak_pi.pdf
- Jacobson IM,** McHutchison JG, Dusheiko G, Di Bisceglie AM, Reddy KR, Bzowej NH, Marcellin P, Muir AJ, Ferenci P, Flisiak R, George J, Rizzetto M, Shouval D, Sola R, Terg RA, Yoshida EM, Adda N, Bengtsson L, Sankoh AJ, Kieffer TL, George S, Kauffman RS, Zeuzem S. Telaprevir for previously untreated chronic hepatitis C virus infection. *N Engl J Med* 2011; **364**: 2405-2416 [PMID: 21696307 DOI: 10.1056/NEJMoa1012912]
- Sherman KE,** Flamm SL, Afdhal NH, Nelson DR, Sulkowski MS, Everson GT, Fried MW, Adler M, Reesink HW, Martin M, Sankoh AJ, Adda N, Kauffman RS, George S, Wright CI, Poordad F. Response-guided telaprevir combination treatment for hepatitis C virus infection. *N Engl J Med* 2011; **365**: 1014-1024 [PMID: 21916639 DOI: 10.1056/NEJMoa1014463]
- Buti M,** Agarwal K, Horsmans Y, Sievert W, Janczewska E, Zeuzem S, Nyberg L, Brown RS, Hézode C, Rizzetto M, Paraná R, De Meyer S, De Masi R, Luo D, Bertelsen K, Witek J. Telaprevir twice daily is noninferior to telaprevir every 8 hours for patients with chronic hepatitis C. *Gastroenterology* 2014; **146**: 744-753.e3 [PMID: 24316262 DOI: 10.1053/j.gastro.2013.11.047]
- Vertex Pharmaceuticals.** Incivek (telaprevir) prescribing information. (Updated October 2013, accessed June 23, 2015). Available from: URL: http://pi.vrtx.com/files/uspi_telaprevir.pdf
- Lawitz E,** Mangia A, Wyles D, Rodriguez-Torres M, Hassanein T, Gordon SC, Schultz M, Davis MN, Kayali Z, Reddy KR, Jacobson IM, Kowdley KV, Nyberg L, Subramanian GM, Hyland RH, Arterburn S, Jiang D, McNally J, Brainard D, Symonds WT, McHutchison JG, Sheikh AM, Younossi Z, Gane EJ. Sofosbuvir for previously untreated chronic hepatitis C infection. *N Engl J Med* 2013; **368**: 1878-1887 [PMID: 23607594 DOI: 10.1056/NEJMoa1214853]
- Jacobson IM,** Dore GJ, Foster GR, Fried MW, Radu M, Rafalsky VV, Moroz L, Craxi A, Peeters M, Lenz O, Ouwerkerk-Mahadevan S, De La Rosa G, Kalmeijer R, Scott J, Sinha R, Beumont-Mauviel M. Simeprevir with pegylated interferon alfa 2a plus ribavirin in treatment-naïve patients with chronic hepatitis C virus genotype 1 infection (QUEST-1): a phase 3, randomised, double-blind, placebo-controlled trial. *Lancet* 2014; **384**: 403-413 [PMID: 24907225 DOI: S0140-6736(14)60494-3]
- Manns M,** Marcellin P, Poordad F, de Araujo ES, Buti M, Horsmans Y, Janczewska E, Villamil F, Scott J, Peeters M, Lenz O, Ouwerkerk-Mahadevan S, De La Rosa G, Kalmeijer R, Sinha R, Beumont-Mauviel M. Simeprevir with pegylated interferon alfa

- 2a or 2b plus ribavirin in treatment-naive patients with chronic hepatitis C virus genotype 1 infection (QUEST-2): a randomised, double-blind, placebo-controlled phase 3 trial. *Lancet* 2014; **384**: 414-426 [PMID: 24907224 DOI: S0140-6736(14)60538-9]
- 23 **Reddy KR**, Zeuzem S, Zoulim F, Weiland O, Horban A, Stanciu C, Villamil FG, Andreone P, George J, Dammers E, Fu M, Kurland D, Lenz O, Ouwerkerk-Mahadevan S, Verbinen T, Scott J, Jessner W. Simeprevir versus telaprevir with peginterferon and ribavirin in previous null or partial responders with chronic hepatitis C virus genotype 1 infection (ATTAIN): a randomised, double-blind, non-inferiority phase 3 trial. *Lancet Infect Dis* 2015; **15**: 27-35 [PMID: 25482330 DOI: 10.1016/S1473-3099(14)71002-3]
- 24 **Gao M**, Nettles RE, Belema M, Snyder LB, Nguyen VN, Fridell RA, Serrano-Wu MH, Langley DR, Sun JH, O'Boyle DR, Lemm JA, Wang C, Knipe JO, Chien C, Colonno RJ, Grasela DM, Meanwell NA, Hamann LG. Chemical genetics strategy identifies an HCV NS5A inhibitor with a potent clinical effect. *Nature* 2010; **465**: 96-100 [PMID: 20410884 DOI: 10.1038/nature08960]
- 25 **Nettles RE**, Gao M, Bifano M, Chung E, Persson A, Marbury TC, Goldwater R, DeMicco MP, Rodriguez-Torres M, Vutikullird A, Fuentes E, Lawitz E, Lopez-Talavera JC, Grasela DM. Multiple ascending dose study of BMS-790052, a nonstructural protein 5A replication complex inhibitor, in patients infected with hepatitis C virus genotype 1. *Hepatology* 2011; **54**: 1956-1965 [PMID: 21837752 DOI: 10.1002/hep.24609]
- 26 **Dore GJ**, Lawitz E, Hézode C, Shafraan SD, Ramji A, Tatum HA, Taliani G, Tran A, Brunetto MR, Zaltron S, Strasser SI, Weis N, Ghesquiere W, Lee SS, Larrey D, Pol S, Harley H, George J, Fung SK, de Lédighen V, Hagens P, McPhee F, Hernandez D, Cohen D, Cooney E, Noviello S, Hughes EA. Daclatasvir plus peginterferon and ribavirin is noninferior to peginterferon and ribavirin alone, and reduces the duration of treatment for HCV genotype 2 or 3 infection. *Gastroenterology* 2015; **148**: 355-366.e1 [PMID: 25311593 DOI: 10.1053/j.gastro.2014.10.007]
- 27 **Hézode C**, Hirschfield GM, Ghesquiere W, Sievert W, Rodriguez-Torres M, Shafraan SD, Thuluvath PJ, Tatum HA, Waked I, Esmat G, Lawitz EJ, Rustgi VK, Pol S, Weis N, Pockros PJ, Bourlière M, Serfaty L, Vierling JM, Fried MW, Weiland O, Brunetto MR, Everson GT, Zeuzem S, Kwo PY, Sulkowski M, Bräu N, Hernandez D, McPhee F, Wind-Rotolo M, Liu Z, Noviello S, Hughes EA, Yin PD, Schnittman S. Daclatasvir plus peginterferon alfa and ribavirin for treatment-naive chronic hepatitis C genotype 1 or 4 infection: a randomised study. *Gut* 2015; **64**: 948-956 [PMID: 25080450 DOI: 10.1136/gutjnl-2014-307498]
- 28 **Kwo PY**, Lawitz EJ, McCone J, Schiff ER, Vierling JM, Pound D, Davis MN, Galati JS, Gordon SC, Ravendhran N, Rossaro L, Anderson FH, Jacobson IM, Rubin R, Koury K, Pedicone LD, Brass CA, Chaudhri E, Albrecht JK. Efficacy of boceprevir, an NS3 protease inhibitor, in combination with peginterferon alfa-2b and ribavirin in treatment-naive patients with genotype 1 hepatitis C infection (SPRINT-1): an open-label, randomised, multicentre phase 2 trial. *Lancet* 2010; **376**: 705-716 [PMID: 20692693 DOI: 10.1016/S0140-6736(10)60934-8]
- 29 **Bristol-Myers Squibb**. Health Canada approves Daklinza™ (daclatasvir) for the treatment of chronic hepatitis C infection across multiple genotypes including genotype 3. (Accessed October 14, 2015). Available from: URL: <http://www.bmscanada.ca/en/news/release/health-canada-approves-daklinza-daclatasvir-for-the-treatment-of-chronic-hepatitis-c-infection-across>
- 30 **Bristol-Myers Squibb**. Daklinza™ (daclatasvir) Prescribing Information 2015. (Accessed October 14, 2015). Available from URL: http://www.accessdata.fda.gov/drugsatfda_docs/label/2015/206843Orig1s000lbl.pdf
- 31 **Fleiss J**, Levin B, Paik M. Statistical Methods for Rates and Proportions. 3rd ed. Hoboken: John Wiley & Sons, 2013
- 32 **Asselah T**, Estrabaud E, Bieche I, Lapalus M, De Muynck S, Vidaud M, Saadoun D, Soumelis V, Marcellin P. Hepatitis C: viral and host factors associated with non-response to pegylated interferon plus ribavirin. *Liver Int* 2010; **30**: 1259-1269 [PMID: 20633102 DOI: 10.1111/j.1478-3231.2010.02283.x]
- 33 **Karino Y**, Toyota J, Ikeda K, Suzuki F, Chayama K, Kawakami Y, Ishikawa H, Watanabe H, Hernandez D, Yu F, McPhee F, Kumada H. Characterization of virologic escape in hepatitis C virus genotype 1b patients treated with the direct-acting antivirals daclatasvir and asunaprevir. *J Hepatol* 2013; **58**: 646-654 [PMID: 23178977 DOI: 10.1016/j.jhep.2012.11.012]
- 34 **McPhee F**, Hernandez D, Yu F, Ueland J, Monikowski A, Carifa A, Falk P, Wang C, Fridell R, Eley T, Zhou N, Gardiner D. Resistance analysis of hepatitis C virus genotype 1 prior treatment null responders receiving daclatasvir and asunaprevir. *Hepatology* 2013; **58**: 902-911 [PMID: 23504694 DOI: 10.1002/hep.26388.]
- 35 **Reddy KR**, Thuluvath PJ, Kumada Y, Toyota J, Chayama K, Levin J, Lawitz E, Gadano A, Ghesquiere W, Gerken G, Brunetto M, Peng CY, Silva M, Strasser S, Heo J, McPhee F, Liu Z, Linaberry M, Hughes EA, Noviello S. Long-term follow-up of patients treated with daclatasvir-based regimens in phase 2 and 3 studies. Hoboken: Wiley-Blackwell, 2014: 1154A-1155A
- 36 **Sulkowski MS**, Gardiner DF, Rodriguez-Torres M, Reddy KR, Hassanein T, Jacobson I, Lawitz E, Lok AS, Hinesstrova F, Thuluvath PJ, Schwartz H, Nelson DR, Everson GT, Eley T, Wind-Rotolo M, Huang SP, Gao M, Hernandez D, McPhee F, Sherman D, Hindes R, Symonds W, Pasquinelli C, Grasela DM. Daclatasvir plus sofosbuvir for previously treated or untreated chronic HCV infection. *N Engl J Med* 2014; **370**: 211-221 [PMID: 24428467 DOI: 10.1056/NEJMoa1306218]
- 37 **Manns M**, Pol S, Jacobson IM, Marcellin P, Gordon SC, Peng CY, Chang TT, Everson GT, Heo J, Gerken G, Yoffe B, Towner WJ, Bourliere M, Metivier S, Chu CJ, Sievert W, Bronowicki JP, Thabut D, Lee YJ, Kao JH, McPhee F, Kopit J, Mendez P, Linaberry M, Hughes E, Noviello S. All-oral daclatasvir plus asunaprevir for hepatitis C virus genotype 1b: a multinational, phase 3, multicohort study. *Lancet* 2014; **384**: 1597-1605 [PMID: 25078304 DOI: 10.1016/S0140-6736(14)61059-X]
- 38 **Lawitz E**, Sulkowski MS, Ghalib R, Rodriguez-Torres M, Younossi ZM, Corregidor A, DeJesus E, Pearlman B, Rabinovitz M, Gitlin N, Lim JK, Pockros PJ, Scott JD, Fevery B, Lambrecht T, Ouwerkerk-Mahadevan S, Callewaert K, Symonds WT, Picchio G, Lindsay KL, Beumont M, Jacobson IM. Simeprevir plus sofosbuvir, with or without ribavirin, to treat chronic infection with hepatitis C virus genotype 1 in non-responders to pegylated interferon and ribavirin and treatment-naive patients: the COSMOS randomised study. *Lancet* 2014; **384**: 1756-1765 [PMID: 25078309 DOI: 10.1016/S0140-6736(14)61036-9]
- 39 **Ferenci P**, Bernstein D, Lalezari J, Cohen D, Luo Y, Cooper C, Tam E, Marinho RT, Tsai N, Nyberg A, Box TD, Younes Z, Enayati P, Green S, Baruch Y, Bhandari BR, Caruntu FA, Sepe T, Chulanov V, Janczewska E, Rizzardini G, Gervain J, Planas R, Moreno C, Hassanein T, Xie W, King M, Podsadecki T, Reddy KR. ABT-450/r-ombitasvir and dasabuvir with or without ribavirin for HCV. *N Engl J Med* 2014; **370**: 1983-1992 [PMID: 24795200 DOI: 10.1056/NEJMoa1402338]
- 40 **Feld JJ**, Kowdley KV, Coakley E, Sigal S, Nelson DR, Crawford D, Weiland O, Aguilar H, Xiong J, Pilot-Matias T, DaSilva-Tillmann B, Larsen L, Podsadecki T, Bernstein B. Treatment of HCV with ABT-450/r-ombitasvir and dasabuvir with ribavirin. *N Engl J Med* 2014; **370**: 1594-1603 [PMID: 24720703 DOI: 10.1056/NEJMoa1315722]
- 41 **Poordad F**, Hezode C, Trinh R, Kowdley KV, Zeuzem S, Agarwal K, Shiffman ML, Wedemeyer H, Berg T, Yoshida EM, Forns X, Lovell SS, Da Silva-Tillmann B, Collins CA, Campbell AL, Podsadecki T, Bernstein B. ABT-450/r-ombitasvir and dasabuvir with ribavirin for hepatitis C with cirrhosis. *N Engl J Med* 2014; **370**: 1973-1982 [PMID: 24725237 DOI: 10.1056/NEJMoa1402869]
- 42 **Poordad F**, Sievert W, Mollison L, Bennett M, Tse E, Bräu N, Levin J, Sepe T, Lee SS, Angus P, Conway B, Pol S, Boyer N, Bronowicki JP, Jacobson I, Muir AJ, Reddy KR, Tam E, Ortiz-Lasanta G, de Lédighen V, Sulkowski M, Boparai N, McPhee F, Hughes E, Swenson ES, Yin PD. Fixed-dose combination therapy with daclatasvir, asunaprevir, and beclabuvir for noncirrhotic

patients with HCV genotype 1 infection. *JAMA* 2015; **313**: 1728-1735 [PMID: 25942723 DOI: 10.1001/jama.2015.3860]

- 43 **Muir AJ**, Poordad F, Lalezari J, Everson G, Dore GJ, Herring R, Sheikh A, Kwo P, Hézode C, Pockros PJ, Tran A, Yozviak J, Reau N, Ramji A, Stuart K, Thompson AJ, Vierling J, Freilich B, Cooper J,

Ghesquiere W, Yang R, McPhee F, Hughes EA, Swenson ES, Yin PD. Daclatasvir in combination with asunaprevir and beclabuvir for hepatitis C virus genotype 1 infection with compensated cirrhosis. *JAMA* 2015; **313**: 1736-1744 [PMID: 25942724 DOI: 10.1001/jama.2015.3868]

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