Original Article

High effectiveness of peginterferon alfa-2a plus ribavirin therapy in Korean patients with chronic hepatitis C in clinical practice

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Background/Aims: Identifying the impact of a patient’s ethnicity on treatment responses in clinical practice may assist in providing individualized treatment regimens for chronic hepatitis C (CHC). The effectiveness of standard peginterferon plus ribavirin therapy and the need for triple combination therapy with protease inhibitors in Koreans remain matters of debate. These issues were investigated in the present study.

Methods: The clinical data of 272 treatment-naïve Korean CHC patients who were treated in a community-based clinical trial (Clinical Trial group; n=51) and in clinical practice (Cohort group; n=221), were analyzed and compared. All were treated with standard protocols of peginterferon alfa-2a plus ribavirin therapy.

Results: For patients with hepatitis C virus (HCV) genotype 1, the sustained virological response (SVR) rates in the Clinical Trial and Cohort groups were 81% (21/26) and 55% (58/106), respectively, by intention-to-treat (ITT) analysis (P=0.02), and 100% (13/13) and 80% (32/40), respectively, in treatment-adherent patients (P=0.18). For patients with HCV genotype 2, the SVR rates in these two groups were 96% (24/25) and 88% (101/115), respectively, by ITT analysis (P=0.31). Adherence and treatment duration were independent predictors of SVR for genotypes 1 and 2, respectively (P<0.01 for each). Korean patients with CHC achieved high SVR rates with peginterferon alfa-2a plus ribavirin in both the clinical trial and clinical practice settings.

Conclusions: Measures to raise adherence to standard therapy in clinical practice may improve the SVR rates in these patients as effectively as adding protease inhibitors, thus obviating the need for the latter. (Clin Mol Hepatol 2013;19:60-69)

Keywords: Medication adherence; Hepatitis C; Peginterferon alfa-2a; Ribavirin

Abbreviations:
ANC, absolute neutrophil count; CHC, chronic hepatitis C; cEVR, complete early virologic response; CI, 95% confidence interval; EVR, early virologic response; ETR, end-of-treatment response; HCV, hepatitis C virus; Hb, hemoglobin; IL28B, interleukin 28B; OR, odds ratio; PCR, polymerase chain reaction; PEG-IFN, peginterferon; RBV, ribavirin; RFMP, restriction fragment mass polymorphism; RNA, ribonucleic acid; SVR, sustained virological response

This study has been registered at ClinicalTrials.gov, number NCT01596517 (URL http://www.clinicaltrials.gov/ct2/show/NCT01596517).

Received : Sep. 21, 2012 / Revised : Dec. 27, 2012 / Accepted : Jan. 10, 2013

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INTRODUCTION

Chronic hepatitis C virus (HCV) infection is one of the major causes of chronic hepatitis, cirrhosis, and hepatocellular carcinoma. The prevalence of HCV infection varies geographically, with the majority of infected people (about 90 million) originating from Asian countries. Despite the high burden of HCV disease among Asians, very little is known about the treatment outcomes in these patients because most of the pivotal studies only included small numbers of Asian patients.

The standard-of-care treatment for patients with chronic hepatitis C (CHC) has been combination of peginterferon (PEG-IFN) and ribavirin (RBV), which induces sustained virological response (SVR) rates of 40-50% in cases with HCV genotype 1, and of 80% or more in cases with genotype 2 or 3 infections. The recent development of protease inhibitors has substantially improved the SVR to achieve SVR than Caucasians.

studies have demonstrated that Asian patients are more likely than the majority of infected people (about 90 million) originating from Asian countries. However, given the high cost and the more frequent occurrence of adverse events associated with triple combination therapy with PEG-IFN, RBV, and protease inhibitor, it is important to identify who would benefit the most from this therapy.

Although the HCV genotype is known to be the strongest predictor of SVR, several host genetic factors have also been found to affect the response to treatment. Recent studies have revealed that the likelihood of achieving an SVR with PEG-IFN and RBV depends on the nucleotide sequence near the interleukin (IL) 28B gene. Interestingly, the frequency of the favorable IL 28B allele is substantially higher in East Asians. Notably, several earlier studies have demonstrated that Asian patients are more likely to achieve SVR than Caucasians. However, contradicting the earlier reports described above, several recent studies have found that Asians have similar or even inferior SVR rates compared to Caucasian patients with the same HCV genotype.

When comparing data from different studies, the study design and methods of analysis should be considered. Real-world effectiveness data derived from ordinary clinical practice settings often differ markedly from the efficacy data obtained in the settings of randomized controlled registration trials.

We assessed the effectiveness of PEG-IFN α-2a and RBV therapy in treatment-naïve Korean patients with CHC and who had accurately diagnosed HCV genotype 1, 2, or 3.

PATIENTS AND METHODS

Patients

The study population was recruited from two groups of treatment-naïve patients with CHC who were treated with the PEG-IFN α-2a plus RBV combination. One group consisted of 100 patients in a prospective, industry-sponsored, open-label, uncontrolled, community-based clinical trial (Pegasys Expanded Access Program) conducted at six tertiary referral centers in Korea between 2003 and 2004 (Clinical Trial group). The second group consisted of 522 patients who were treated in a single tertiary referral hospital (Asan Medical Center, Seoul, Korea) between 2004 and 2008 (Cohort group).

Eligible patients were previously untreated adults aged 18-70 years who had polymerase chain reaction (PCR)-detectable HCV ribonucleic acid (RNA) and compensated liver disease. Patients were excluded if they had any of the following: a HCV genotype other than 1, 2, or 3; acute hepatitis C; decompensated cirrhosis; hepatocellular carcinoma; other forms of liver disease. Patients with human immunodeficiency virus, pre-existing severe depression or other psychiatric disease, previous organ transplantation, absolute neutrophil count (ANC) <1,000 cells/mm³, platelet count <75,000 cells/mm³, or hemoglobin (Hb) <13 g/dL for men or <12 g/dL for women were also excluded. Cirrhosis was based on the histological diagnosis in Clinical Trial group, and on histological or radiological diagnosis in Cohort group. All study patients were of Korean ethnicity. This study was approved by the Institutional Review Board at each participating center.

Treatment protocol

The HCV genotype was determined by using the restriction fragment mass polymorphism (RFMP) assay. Patients with genotype 1 were treated with PEG-IFN α-2a (Roche, Basel, Switzerland) 180 μg/week and a daily RBV (Roche for the Clinical Trial group; Shinpoong, Korea for the Cohort group) dose of 1,000 mg (for patients with body weight <75 kg) or 1,200 mg (for patients with body weight ≥75 kg) for 48 weeks. Patients with genotype 2 or 3 were treated with PEG-IFN α-2a 180 μg/week and a daily RBV dose of 800 mg for 24 weeks. All study medications for the patients in the Clinical Trial group were provided by Roche, whereas the Cohort group patients purchased their medications. HCV RNA was quantified (Roche AMPLICOR HCV Test v2.0) at pretreatment, weeks 12, 24 and 48 for both genotypes and at week 72 for genotype 1.
Dose modification and definition of adherence

In the Clinical Trial group, if ANC dropped below 750 cells/mm$^3$, PEG-IFN $\alpha$-2a was reduced to 135 $\mu$g. If ANC fell below 500 cells/mm$^3$, PEG-IFN $\alpha$-2a was stopped until ANC recovered to 1,000 cells/mm$^3$, and the restarting of medication was initiated from 90 $\mu$g. If platelet counts dropped below 50,000 cells/mm$^3$, PEG-IFN $\alpha$-2a was reduced to 90 $\mu$g. If platelet counts fell below 25,000 cells/mm$^3$, the discontinuation of administration was recommended. The RBV dose was reduced if Hb decreased to <10 g/dL and stopped if Hb decreased to <8.5 g/dL.

In the Cohort group, the dose modification depended on the decision of the physician, which was based on laboratory data and clinical information and referred to the guidelines used in the clinical trial. The information about the amount of each drug administered to a patient was obtained from the drug dispensing/return or prescription records.

None of the patients in either group were treated with growth factors for hematological side effects.

Adherence $\geq$80/80/80 was defined when a patient was treated for $\geq$80% of the assigned treatment duration and received $\geq$80% of both PEG-IFN $\alpha$-2a and RBV doses.

Main outcome measures

The primary outcome of interest in this study was the proportion of patients who achieved SVR, which was defined as documented PCR-undetectable serum HCV RNA levels at 24 weeks after cessation of treatment. Secondary outcomes included early virological response (EVR, reduction of HCV RNA levels by 2 log or more at 12 weeks of treatment), complete EVR (cEVR, PCR-undetectable HCV RNA at 12 weeks of treatment), and end-of-treatment response (ETR, HCV RNA undetectable at the end of treatment).

Statistical analysis

All response evaluations and adherence to treatment were assessed by modified intention-to-treat analysis unless specified (i.e., all patients who received at least one dose of medication were included in the analysis).

Between-group comparisons of continuous variables were determined by using independent t-tests, and categorized variables were compared by using the chi-square test or Fisher’s exact test, as appropriate. The predictive factors for SVR were analyzed by logistic regression. A $P$-value less than 0.05 was considered to be statistically significant in all analyses. Statistical analyses were performed by using SPSS version 17.0 (SPSS Inc., Chicago, IL, USA).

RESULTS

Baseline characteristics

For the Clinical Trial group, 100 patients with CHC were assessed: 5 refused to provide consent and 44 had previous treatment; thus, 51 treatment-naive patients were included in intention-to-treat analysis. Of these, 26 and 25 had genotype 1 and genotype 2 or 3, respectively (Fig. 1).

In the Cohort group, 522 consecutive patients with HCV were...
assessed: 301, including three patients with HCV genotype 6, did not meet the enrollment criteria and were excluded. The remaining 221 eligible patients were included in intention-to-treat analysis. Of these, 106 and 115 had genotype 1 and genotype 2, respectively (Fig. 1).

The Clinical Trial and the Cohort group were similar in most baseline characteristics (Table 1). However, the Clinical Trial group had a higher serum ALT levels than the Cohort group (P=0.03 and P<0.01 for genotypes 1 and 2, respectively). Of those with genotype 2, the Cohort patients were more likely to have cirrhosis than the Clinical Trial patients (0% vs. 21.7%, P<0.01).

### Virological responses and adherence to treatment

In patients with genotype 1, the Clinical Trial and Cohort group did not differ in the proportion of patients who achieved EVR, cEVR, and ETR (EVR: 84.6% vs. 80.2%, P=0.78; cEVR: 84.6% vs. 73.6%, P=0.24; ETR: 92.3% vs. 80.2%, P=0.25; Table 2). However, the Clinical Trial group had a significantly higher SVR rates than the Cohort group (80.8% vs. 54.7%, P=0.02). The Clinical Trial group also had a significantly higher proportion of patients with treatment duration ≥80% than the Cohort group (96.2% vs. 69.8%, P<0.01). However, the difference in the proportion of patients with adherence ≥80/80/80 was not significant between the two groups (53.8% vs. 38.7%, P=0.16).

In patients with genotype 2, the Clinical Trial group had signifi-

| Table 1. Baseline characteristics of the study patients |
|--------------------------------------------------------|
| Genotype 1                                             |
|                                                        |
| Clinical trial | Cohort | P-value |
|--------------------------------------------------------|
| Number        | 26     | 106     |            |
| Age* (yr)     | 51.9±12.5 | 49.2±11.5 | 0.30 |
| Male          | 16 (61.5%) | 75 (70.8%) | 0.36 |
| Body weight† (kg) | 66.0±9.9  | 67.9±11.6  | 0.45 |
| Serum ALT† (IU/L) | 84 (61-155) | 68 (35-101) | 0.03 |
| Cirrhosis‡   | 2 (7.7%) | 23 (21.7%) | 0.16 |
| HCV RNA† (log_{10} IU/mL) | 5.91 (5.22-6.06) | 6.04 (5.62-6.51) | 0.22 |
| HCV RNA level >800,000 IU/mL | 13 (50.0%) | 55 (51.9%) | 0.86 |
| HCV Subtype  |            |            | 0.35 |
| 1b            | 21 (80.8%) | 93 (87.7%) |       |
| 1, 1a, 1a/b, 1c | 4, 1, 0, 0 (19.2%) | 1, 8, 2, 2 (12.3%) |       |
| Genotype 2                                             |
|                                                        |
| Clinical trial | Cohort | P-value |
|--------------------------------------------------------|
| Number        | 25     | 115     |            |
| Age* (yr)     | 49.6±9.1 | 52.9±11.5 | 0.17 |
| Male          | 16 (64.0%) | 59 (51.3%) | 0.25 |
| Body weight† (kg) | 68.4±11.5  | 64.6±11.3  | 0.13 |
| Serum ALT† (IU/L) | 99 (57-211) | 60 (40-93)  | <0.01 |
| Cirrhosis‡   | 0 (0.0%) | 25 (21.7%) | <0.01 |
| HCV RNA† (log_{10} IU/mL) | 5.53 (4.83-5.83) | 5.39 (4.32-5.85) | 0.27 |
| HCV RNA level >800,000 IU/mL | 4 (16.0%) | 28 (24.3%) | 0.37 |
| HCV subtype  |            |            | 0.86 |
| 2a/c          | 16 (64.0%) | 78 (67.8%) |       |
| 2             | 8 (32.0%) | 30 (26.1%) |       |
| 2a, 2b, 2e, 3a | 0, 0, 0, 1 (4.0%) | 3, 3, 1, 0 (6.1%) |       |

*Mean±SD; †Median (interquartile range); ‡Cirrhosis was based on the histological diagnosis in Clinical Trial group, and on histological or radiological diagnosis in Cohort group.

ALT, alanine aminotransferase; HCV, hepatitis C virus.
Table 2. Virological responses and adherence to treatment according to study group

|                    | Clinical trial | Cohort  | P-value |
|--------------------|----------------|---------|---------|
| Genotype 1         |                |         |         |
| Number             | 26             | 106     |         |
| Early virologic response | 22 (84.6%)   | 85 (80.2%) | 0.78   |
| Complete early virological response | 22 (84.6%)   | 78 (73.6%) | 0.24   |
| End-treatment response | 24 (92.3%)   | 85 (80.2%) | 0.25   |
| Sustained virological response | 21 (80.8%)   | 58 (54.7%) | 0.02   |
| Early drop-out*    | 0              | 16 (15.0%) | 0.04   |
| Treatment duration ≥80% | 25 (96.2%)   | 74 (69.8%) | 0.01   |
| Adherence ≥80/80/80† | 14 (53.8%)   | 41 (38.7%) | 0.16   |
| Genotype 2         |                |         |         |
| Number             | 25             | 115     |         |
| Early virologic response | 23 (92.0%)   | 83 (72.2%) | 0.04   |
| Complete early virological response | 23 (92.0%)   | 82 (71.3%) | 0.03   |
| End-treatment response | 24 (96.0%)   | 111 (96.5%) | 1.00   |
| Sustained virological response | 24 (96.0%)   | 101 (87.8%) | 0.31   |
| Early drop-out*    | 0              | 10 (8.7%) | 0.21   |
| Treatment duration ≥80% | 25 (100%)    | 92 (80.0%) | 0.01   |
| Adherence ≥80/80/80† | 18 (72.0%)   | 78 (67.8%) | 0.68   |

All endpoints were analyzed by modified intention-to-treat analysis, i.e. all patients who received at least one dose of medication were included in the analysis.

*Defined when the treatment was stopped within initial 12 weeks; †Defined when a patient received ≥80% of total peginterferon α-2a and ribavirin doses during ≥80% of the assigned duration of therapy.

Significantly higher EVR and cEVR rates than the Cohort group (EVR: 92.0% vs. 72.2%, P=0.04; cEVR: 92.0% vs. 71.3%, P=0.03; Table 2). However, the two groups did not differ in terms of ETR and SVR rates (ETR: 96.0% vs. 96.5%, P=1.00; SVR: 96.0% vs. 87.8%, P=0.31, respectively). The Clinical Trial group had a significantly higher proportion of patients with a treatment duration ≥80% than the Cohort group (100% vs. 80.0%, P=0.01). However, the two groups were similar in the proportion of patients with adherence ≥80/80/80 (72.0% vs. 67.8%, P=0.68).

Predictive factors for SVR

In patients with genotype 1, univariate analysis showed that age, cirrhosis, treatment in the Clinical Trial, achievement of EVR, treatment duration ≥80%, and adherence ≥80/80/80 were significantly associated with the achievement of SVR (Table 3). Of these significant factors, age (odds ratio [OR]=0.95, 95% confidence interval [CI]=0.92-0.99, P=0.02), treatment in the Clinical Trial (OR=4.94, CI=1.38-17.7, P=0.01), and adherence ≥80/80/80 (OR=4.67, CI=1.82-12.0, P<0.01) were independently associated with SVR by multivariate analysis.

In patients with genotype 2, univariate analysis showed that age, cirrhosis, achievement of EVR, treatment duration ≥80%, and adherence ≥80/80/80 were significantly associated with SVR. Multivariate analysis revealed that only cirrhosis (OR=0.17, CI=0.05-0.63, P<0.01) and treatment duration ≥80% (OR=8.16, CI=2.29-29.13, P<0.01) were independently associated with SVR.

SVR rate as a function of EVR and adherence

SVR rate as a function of the amount of treatment received was assessed at different adherence levels to determine any relationship between adherence and sustained response. As shown in Fig. 2, we found a stepwise, increasing relationship between adherence and SVR in either Clinical Trial or Cohort group regardless of HCV genotype. Thus, in genotype 1 patients with treatment duration ≥80%, at least 80% adherence to therapy increased SVR rates from 66.7% to 100% for those in Clinical Trial group and from 54.8% to 80.0% for those in Cohort group (Fig. 2A).
Table 3. Predictive factors of a sustained virological response

|                        | Univariate analysis | Multivariate analysis |
|------------------------|---------------------|-----------------------|
|                        | OR (95% CI)         | P-value               |
|                        |                     | OR (95% CI)           | P-value*                     |
| Genotype 1             |                     |                       |
| Age                    | 0.95 (0.91-0.98)    | <0.01                 |
| Gender (male)          | 1.25 (0.59-2.64)    | 0.56                  |
| Body weight (kg)       | 0.99 (0.96-1.02)    | 0.62                  |
| Serum ALT (IU/L)       | 1.00 (0.99-1.01)    | 0.35                  |
| HCV RNA <800,000 IU/mL | 1.60 (0.79-3.23)    | 0.19                  |
| Cirrhosis              | 0.37 (0.15-0.90)    | 0.03                  |
| Clinical trial†        | 3.48 (1.22-9.91)    | <0.01                 |
| EVR (+)                | 8.97 (3.10-25.95)   | <0.01                 |
| Treatment duration ≥80% | 7.92 (3.20-19.65)   | <0.01                 |
| Adherence ≥80/80/80    | 8.26 (3.44-19.83)   | <0.01                 |
| Genotype 2             |                     |                       |
| Age                    | 0.93 (0.88-0.99)    | 0.02                  |
| Gender (male)          | 1.85 (0.62-5.51)    | 0.27                  |
| Body weight (kg)       | 1.01 (0.96-1.06)    | 0.73                  |
| Serum ALT (IU/L)       | 1.00 (0.99-1.01)    | 0.88                  |
| HCV RNA <800,000 IU/mL | 1.82 (0.57-5.76)    | 0.31                  |
| Cirrhosis              | 0.19 (0.06-0.60)    | <0.01                 |
| Clinical trial†        | 3.33 (0.42-26.55)   | 0.26                  |
| EVR (+)                | 4.35 (1.45-13.10)   | <0.01                 |
| Treatment duration ≥80%| 8.38 (2.66-26.44)   | <0.01                 |
| Adherence ≥80/80/80    | 5.35 (1.71-16.80)   | <0.01                 |

*Generated by logistic regression analysis with backward elimination method; †With the Cohort group as a reference.

Figure 2. Proportion of the patients in the Clinical Trial and Cohort groups who had a sustained virological response (SVR) as a function of early virological response (EVR) and adherence to treatment for patients with HCV genotype 1 (A) or genotype 2 (B). All SVR rates were determined by intention-to-treat analysis. The total number of patient is shown at the base of each bar. Adherence <80/80 means reduced doses (i.e., <80% of one or both drug). Adherence ≥80/80/80 was defined when a patient was treated for ≥80% of the assigned treatment duration and received ≥80% of both PEG-IFN α-2a and RBV doses.
otype 1 patients who achieved EVR and adherence ≥80/80/80, the SVR rate was not significantly different between the Clinical Trial and Cohort groups (100% vs. 80.0%, P=0.18).

In patients with genotype 2, the stepwise, increasing relationship between adherence and SVR was observed only in Cohort group (Fig. 2B).

Causes of non-adherence

When the genotype 1 and 2 patients were combined, 19 of the 51 Clinical Trial patients (37.3%) and 102 of the 221 Cohort group patients (50.7%) did not show adherence ≥80/80/80 (Table 4). Overall, the most common cause of non-adherence in both groups was laboratory abnormalities, including anemia and neutropenia. However, the two groups differed significantly in terms of the distribution of the causes of non-adherence (P<0.01): compared to the Clinical Trial group, laboratory abnormalities were a significantly less frequent cause of non-adherence in the Cohort group (89.5% vs. 50.0%, P<0.01), while adverse symptoms were a more frequent cause (10.5% vs. 34.3%, P=0.04).

DISCUSSION

In the present study, we tried to adequately address all issues in assessing the treatment response in CHC patients including heterogeneity in ethnicity of patients, prior treatment-experience, errors in HCV genotyping, study design, and dataset for endpoint analysis. Thus, we clearly demonstrate that treatment-naïve Korean patients with CHC achieve high SVR rates with PEG-IFN α-2a plus RBV therapy in both the community-based clinical trial and the clinical practice settings. For patients with HCV genotype 1, the overall SVR rates of the Clinical Trial and the clinical practice Cohort group were 80.8% and 54.7%, respectively, by an intention-to-treat analysis. The SVR rates of the Clinical Trial and the Cohort group increased to as high as 100% and 80.0%, respectively, when a per-protocol analysis was applied (i.e., in patients with EVR and adherence ≥80/80/80). For patients with genotype 2, the overall SVR rates of the both groups were 96.0% and 87.8%, respectively.

The SVR rates of this real-world effectiveness study in East Asian patients with CHC are higher than the SVR rates of published randomized controlled registration trials that employed a similar regimen and duration of treatment and mainly included Caucasian patients.4,5 In the latter trials, the overall SVR rates

| Table 4. Causes of nonadherence according to treatment group†‡ |
|---------------------------------------------------------------|
| **Frequency, N (%)** | **Clinical trial (n=19)** | **Cohort (n=102)** | **P-value** |
| --- | --- | --- | --- |
| Laboratory abnormalities | 17 (89.5%) | 51 (50.0%) | <0.01 |
| Anemia | 8 | 40 |  |
| Neutropenia | 13 | 11 |  |
| Thrombocytopenia | 4 |  |  |
| Pancytopenia | 3 |  |  |
| Elevated ALT | 4 |  |  |
| Adverse symptoms | 2 (10.5%) | 35 (34.3%) | 0.04 |
| Flu-like symptoms | 8 |  |  |
| Gastrointestinal symptoms | 1 | 6 |  |
| Pruritus | 6 |  |  |
| Dizziness | 4 |  |  |
| General weakness | 4 |  |  |
| Alopecia | 4 |  |  |
| Dyspnea | 3 |  |  |
| Skin rash | 1 | 2 |  |
| Insomnia | 2 |  |  |
| Tinnitus | 2 |  |  |
| Depression | 2 |  |  |
| Nocturia | 2 |  |  |
| Anxiety | 1 |  |  |
| Hematochezia | 1 |  |  |
| Generalized edema | 1 |  |  |
| Anorexia | 1 |  |  |
| Hyperthyroidism | 1 |  |  |
| Hypothyroidism | 1 |  |  |
| Blurred vision | 1 |  |  |
| Muscle cramping | 1 |  |  |
| Ascites | 1 |  |  |
| Others | 2 (10.5%) | 24 (23.5%) | 0.36 |
| Comorbid illnesses | 1 | 7 |  |
| Lack of EVR | 1 | 1 |  |
| Economic issues | 2 |  |  |
| Non-compliance | 2 |  |  |
| Loss of follow up | 12 |  |  |

*Non-adherence was defined when the patient did not fulfill the criteria of adherence ≥80/80/80; †Multiple causes of non-adherence in a patient were expressed separately; ‡Hepatocellular carcinoma (n=3), ileus (n=1), Gastrointestinal stromal tumor (n=1), appendicitis (n=1), thyroid cancer (n=1), tuberculosis (n=1).

ALT, alanine aminotransferase.
ranged between 46% and 52% for genotype 1 patients and between 76% and 80% for genotype 2 or 3 patients. The SVR rate of 81% in our Clinical Trial group and 80% of treatment-adherent patients in our clinical practice Cohort group with HCV genotype 1 were even higher or comparable to the results from recent registration trials on triple combination therapy with boceprevir or telaprevir, PEG-IFN, and RBV.2,8

The high effectiveness of PEG-IFN and RBV therapy in our patients with CHC are interesting because patients treated in community practices generally have lower, or similar at best, response rates compared to those in registration trials; this is due to the inclusion of a broader spectrum of patients, less frequent monitoring, lack of financial support, low insurance coverage, and lower adherence to treatment.21-23-26 Thus, the data of this study strongly suggest that East Asian patients with CHC are more responsive to PEG-IFN plus RBV therapy than Caucasian patients.

The results of this study are consistent with those of several previous clinical trials and cohort studies, which showed that Asian patients with CHC have higher SVR rates when directly compared to Caucasian patients.9,10,27-29 As we and others have found, the vast majority (82-100%) of Asians who had EVR also achieved cEVR, whereas only about half of Caucasians with EVR achieved cEVR.17 These results suggest that a higher proportion of the Asian subjects in the present study were rapid virological responders, and as a result, they had lower rates of breakthrough or relapse and a higher SVR rate.

Recent understanding for the effect of IL-28B gene-related single nucleotide polymorphisms on response to PEG-IFN and RBV treatment may provide the biological basis for the racial differences in viral kinetics during treatment.9,11 Patients with the C/C genotype showed a greater decline in HCV RNA during first 4 weeks of treatment when compared to patients with non-C/C genotypes, irrespective of race,27 and that the C/C genotype is also associated with a three times higher probability of SVR than the C/T and T/T genotypes combined.9,27 Geographical variation in the IL28B gene supports the notion that East Asian patients should have higher response rates as they have a higher frequency of the C/C genotype than their white European counterparts.9,10,28,29

In contrast to our results, others have recently demonstrated that Asians have similar or even inferior SVR rates compared to Caucasian patients with the same HCV genotype.17,20 They suggested that the SVR rates in Asians in the earlier studies were erroneously inflated by the possible inclusion of easier-to-treat genotype 6 patients in the genotype 1 group. Supporting this is that the earlier studies used the INNO-LiPA probe assay for HCV genotyping which may mistype genotype 6 as genotype 1. This is a plausible possibility because up to one-third of HCV-infected patients from Southern China and Southeast Asia are estimated to have HCV genotype 6.30 However, the mistyping error was eliminated from our study because we used the RFMP genotyping assay, which accurately differentiates HCV genotype 6 from genotype 1.31 A bias might also have arisen by including diverse ethnic groups as a single Asian race in previous studies. Each of Asian ethnic groups encompasses diverse genetic backgrounds, and racial classifications do not convey genetic homogeneity.19,25 For example, grouping subjects from Indian sub-continent and East Asia as Asians may introduce bias, as genetic heterogeneity may exist between these groups.19 By including only a single East Asian ethnic group, this kind of selection bias could be avoided in this study.

Adherence to treatment was an independent determinant of the achievement of SVR in this study. The importance of adherence on SVR rates has been well described previously25,32-35 We also found that adherence to treatment was more important for patients with genotype 1 than those with genotype 2, and adherence tended to be lower in the Cohort group than in the Clinical Trial group. Accordingly, the overall SVR rate for genotype 1 patients in the Cohort group was lower than patients in the Clinical Trial group. However, this difference disappeared when the outcome of only the patients with a high level of adherence was evaluated. These results suggest that measures to raise adherence to standard treatment in East Asian patients with CHC would be more cost-effective way to improve the overall SVR rate than adding protease inhibitors, especially in clinical practice settings.

The most common cause of non-adherence was dose reduction due to laboratory abnormalities. Asians seem to be much more likely to undergo RBV dose reductions and discontinuations of treatment because of anemia.16,17 Hence, it is possible that the use of growth factors may be particularly beneficial for Asian patients in terms of improving adherence to treatment and SVR in clinical practice.24

Treatment-related adverse symptoms were more frequent cause of non-adherence in the Cohort group compared with Clinical Trial group. Although it is unclear whether the dose reduction or treatment discontinuation was patient-initiated or physician-instructed, the importance of the role of health care providers should not be overlooked. Previous studies have reported that decreased drug exposure due to physician-directed dosage reductions of interferon and RBV can reduce virologic response rates.33,36 In addition to the physician experience, the role of a dedicated physician as-

http://www.e-cmh.org http://dx.doi.org/10.3350/cmh.2013.19.1.60
sistant who provides education and support to patients undergoing hepatitis C therapy has been suggested to be important. Unfortunately, we did not have such a physician assistant during the period of this study, which might have contributed to the low adherence in our clinical practice cohort patients.

The main limitation of our study is its retrospective design. However, we used the standard definition of SVR as our primary endpoint which is an objective and well-defined outcome variable. Moreover, all results were analyzed according to the intention-to-treat method. Second, there is a possibility of type 2 error, that is, the small number of Clinical Trial group may fail to show the significant difference in the SVR rate between the Clinical Trial group and the Cohort group in per-protocol analysis (100% vs. 80%) in genotype 1 and in intention-to-treat analysis and per protocol analysis in genotype 2 (96% vs. 88%, and 94% vs. 95%, respectively). However, the SVR rate was already near the maximum level of 100% in the Clinical Trial group, thus, there is low possibility to discriminate the significant difference by increasing the sample size of the Clinical Trial group. Third, since our study only included patients of a single Asian ethnicity, we were not able to compare our results directly to those of Caucasian patients. Lastly, since few patients had HCV RNA quantitation data at treatment week 4, we could not analyze the impact of a rapid virological response on the achievement of SVR.

In conclusion, our study highlights the importance of ethnicity as an integral component of the tailored treatment approach to CHC. Our results also emphasize the need to enhance adherence to HCV therapies. With adequate adherence to PEG-IFN α-2a plus RBV treatment, Korean patients with HCV genotype 1 could achieve similar or even higher SVR rate than that can be achieved by Caucasian patients with triple combination therapy including protease inhibitors. Thus, measures to raise adherence to the standard therapy in clinical practice may improve the SVR rates in East Asian CHC patients as effectively as adding protease inhibitors.

Acknowledgements

This study was funded by Roche in part. Roche had no role in study design, data collection, analysis, or decision to submit the manuscript.

The authors thank Drs. Jung Hyun Kwon and Si Hyun Bae for their help in data collection.

Conflicts of Interest

The authors have no conflicts to disclose.

REFERENCES

1. Lavanchy D. The global burden of hepatitis C. Liver Int 2009;29(Suppl 1):74-81.
2. Shepard CW, Finelli L, Alter MJ. Global epidemiology of hepatitis C virus infection. Lancet Infect Dis 2005;5:558-567.
3. WHO. Hepatitis C: global prevalence. Wkly Epidemiol Rec 1997;72:341-344.
4. Fried MW, Shiffman ML, Reddy KR, Smith C, Marinos G, Gonçalves FL Jr, et al. Peginterferon alfa-2a plus ribavirin for chronic hepatitis C virus infection. N Engl J Med 2002;347:975-982.
5. Hadziyannis SJ, Sette H Jr, Morgan TR, Balan V, Diago M, Marcellin P, et al. Peginterferon-alpha2a and ribavirin combination therapy in chronic hepatitis C: a randomized study of treatment duration and ribavirin dose. Ann Intern Med 2004;140:346-355.
6. Manns MP, McHutchison JG, Gordon SC, Rustgi VK, Shiffman M, Reindollar R, et al. Peginterferon alfa-2b plus ribavirin compared with interferon alfa-2b plus ribavirin for initial treatment of chronic hepatitis C: a randomised trial. Lancet 2001;358:958-965.
7. Jacobson IM, McHutchison JG, Dusheiko G, Di Bisceglie AM, Reddy KR, Bzowej NH, et al. Telaprevir for previously untreated chronic hepatitis C virus infection. N Engl J Med 2011;364:2405-2416.
8. Poordad F, McCone J Jr, Bacon BR, Bruno S, Manns MP, Sulkowski MS, et al. Boceprevir for untreated chronic HCV genotype 1 infection. N Engl J Med 2011;364:1195-1206.
9. Ge D, Fellay J, Thompson AJ, Simon JS, Shinaa KV, Urban TJ, et al. Genetic variation in IL28B predicts hepatitis C treatment-induced viral clearance. Nature 2009;461:399-401.
10. Thomas DL, Thio CL, Martin MP, Qi Y, Ge D, O’Huigin C, et al. Genetic variation in IL28B and spontaneous clearance of hepatitis C virus. Nature 2009;461:798-801.
11. Tanaka Y, Nishida N, Sugiyama M, Kurosaki M, Matsuura K, Sakanoto N, et al. Genome-wide association of IL28B with response to pegylated interferon-alpha and ribavirin therapy for chronic hepatitis C. Nat Genet 2009;41:1105-1109.
12. Dev AT, McCaw R, Sundararajan V, Bowden S, Sievert W. Southeast Asian patients with chronic hepatitis C: the impact of novel genotypes and race on treatment outcome. Hepatology 2002;36:1259-1265.
13. Hepburn MJ, Hepburn LM, Cantu NS, Lapeer MG, Lawitz EJ. Differences in treatment outcome for hepatitis C among ethnic groups. Am J Med 2004;117:163-168.
14. Missiha S, Heathcote J, Arenovich T, Khan K; Canadian Pegasys
Expanded Access Group. Impact of asian race on response to combination therapy with peginterferon alfa-2a and ribavirin in chronic hepatitis C. Am J Gastroenterol 2007;102:2181-2188.

15. Yan KK, Guirgis M, Dinh T, George J, Dev A, Lee A, et al. Treatment responses in Asians and Caucasians with chronic hepatitis C infection. World J Gastroenterol 2008;14:3416-3420.

16. Hu KQ, Freilich B, Brown RS, Brass C, Jacobson IM. Impact of Hispanic or Asian ethnicity on the treatment outcomes of chronic hepatitis C: results from the WIN-R trial. J Clin Gastroenterol 2011;45:720-726.

17. Vutien P, Nguyen NH, Trinh HN, Li J, Garcia RT, Garcia G, et al. Similar treatment response to peginterferon and ribavirin in Asian and Caucasian patients with chronic hepatitis C. Am J Gastroenterol 2010;105:1110-1115.

18. Nguyen NH, VuTien P, Garcia RT, Trinh H, Nguyen H, Nguyen K, et al. Response to pegylated interferon and ribavirin in Asian American patients with chronic hepatitis C genotypes 1 vs 2/3 vs 6. J Viral Hepat 2010;17:691-697.

19. Lawson A; Trent Hepatitis C Study Group. A comparison of the natural history and outcome of treatment for Asian and non-Asian hepatitis C-infected patients. J Viral Hepat 2011;18:e270-e277.

20. Freshwater DA, O’Donnell K, Mutimer DI. Inferior response of Asian vs non-Asian hepatitis C genotype 3 infection to combination antiviral therapy. J Viral Hepat 2008;15:115-119.

21. Feuerstadt P, Bunim AL, Garcia H, Karlitz JJ, Massoumi H, Thosani AJ, et al. Effectiveness of hepatitis C treatment with pegylated interferon and ribavirin in urban minority patients. Hepatology 2010;51:1137-1143.

22. Lindor RA, Lindor KD. The value of observational research in liver diseases. Hepatology 2011;53:1-3.

23. Pariente A, Lahmek P, Duprat C, Denis J, Faroux R, Renou C, et al. Treatment of chronic hepatitis C with pegylated interferon and ribavirin in treatment-naive patients in 'true life': a plea in favor of independent postmarketing evaluations. Eur J Gastroenterol Hepatol 2010;22:1297-1302.

24. Shehab TM, Fontana RJ, Oberhelman K, Marrero JA, Su GL, Lok AS. Effectiveness of interferon alpha-2b and ribavirin combination therapy in the treatment of naive chronic hepatitis C patients in clinical practice. Clin Gastroenterol Hepatol 2004;2:425-431.

25. Lo Re V 3rd, Teal V, Localio AR, Amorosa VK, Kaplan DE, Gross R. Relationship between adherence to hepatitis C virus therapy and virologic outcomes: a cohort study. Ann Intern Med 2011;155:353-360.

26. Marotta P, Hueppe D, Zehnter E, Kwo P, Jacobson I. Efficacy of chronic hepatitis C therapy in community-based trials. Clin Gastroenterol Hepatol 2009;7:1028-1036; quiz 1022.

27. Howell CD, Gorden A, Ryan KA, Thompson AJ, Ibrahim C, Fried M, et al. Single nucleotide polymorphism upstream of interleukin 28B associated with phase 1 and phase 2 of early viral kinetics in patients infected with HCV genotype 1. J Hepatol 2012;56:557-563.

28. Sinn DH, Kim YI, Lee ST, Gwak GY, Choi MS, Lee JH, et al. Association of a single nucleotide polymorphism near the interleukin-28B gene with response to hepatitis C therapy in Asian patients. J Gastroenterol Hepatol 2011;26:1374-1379.

29. Lyoo K, Song MJ, Hur W, Choi JE, Hong SW, Kim CW, et al. Polymorphism near the IL28B gene in Korean hepatitis C virus-infected patients treated with peg-interferon plus ribavirin. J Clin Virol 2011;52:363-366.

30. Sievert W, Attraif I, Razavi HA, Abdo A, Ahmed EA, Alomair A, et al. A systematic review of hepatitis C virus epidemiology in Asia, Australia and Egypt. Liver Int 2011;31(Suppl 2):61-80.

31. Oh HB, Kim SO, Cha CH, Hong SP, Folk WR, Kim KM, et al. Identification of hepatitis C virus genotype 6 in Korean patients by analysis of 5’ untranslated region using a matrix assisted laser desorption/ionization time of flight-based assay, restriction fragment length polymorphism. J Med Virol 2008;80:1712-1719.

32. McHutchison JG, Manns M, Patel K, Poynard T, Lindsay KL, Trepo C, et al. Adherence to combination therapy enhances sustained response in genotype-1-infected patients with chronic hepatitis C. Gastroenterology 2002;123:1061-1069.

33. Reddy KR, Shiffman ML, Morgan TR, Zeuzem S, Hadziyannis S, Hamzeh FM, et al. Impact of ribavirin dose reductions in hepatitis C virus genotype 1 patients completing peginterferon alfa-2a/ribavirin treatment. Clin Gastroenterol Hepatol 2007;5:124-129.

34. Shiffman ML, Ghany MG, Morgan TR, Wright EC, Everson GT, Lindsay KL, et al. Impact of reducing peginterferon alfa-2a and ribavirin dose during retreatment in patients with chronic hepatitis C. Gastroenterology 2007;132:103-112.

35. Weiss JJ, Bräu N, Strivala A, Swan T, Fishbein D. Review article: adherence to medication for chronic hepatitis C - building on the model of human immunodeficiency virus antiretroviral adherence research. Aliment Pharmacol Ther 2009;30:14-27.

36. Raptopoulou M, Tsantoulas D, Vafiadi I, Ketikoglou I, Paraskevas E, Vassiliadis T, et al. The effect of adherence to therapy on sustained response in daily or three times a week interferon alpha-2b plus ribavirin treatment of naive and nonresponder chronic hepatitis C patients. J Viral Hepat 2005;12:91-95.

37. Clark CH, Ghalib RH. Hepatitis C: role of the advanced practice nurse. AACN Clin Issues 1999;10:455-463.