ORIGINAL ARTICLE

Hepatitis C virus kinetics by administration of pegylated interferon-α in human and chimeric mice carrying human hepatocytes with variants of the IL28B gene

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ABSTRACT

Objective Recent studies have demonstrated that genetic polymorphisms near the IL28B gene are associated with the clinical outcome of pegylated interferon-α (peg-IFN-α) plus ribavirin therapy for patients with chronic hepatitis C virus (HCV). However, it is unclear whether genetic variations near the IL28B gene influence hepatic interferon (IFN)-stimulated gene (ISG) induction or cellular immune responses, lead to the viral reduction during IFN treatment.

Design Changes in HCV-RNA levels before therapy, at day 1 and weeks 1, 2, 4, 8 and 12 after administering peg-IFN-α plus ribavirin were measured in 54 patients infected with HCV genotype 1. Furthermore, we prepared four lines of chimeric mice having four different lots of human hepatocytes containing various single nucleotide polymorphisms (SNP) around the IL28B gene. HCV infecting chimeric mice were subcutaneously administered with peg-IFN-α for 2 weeks.

Results There were significant differences in the reduction of HCV-RNA levels after peg-IFN-α plus ribavirin therapy based on the IL28B SNP rs8099917 between TT (favourable) and TG/GG (unfavourable) genotypes in patients; the first-phase viral decline slope per day and second-phase slope per week in TT genotype were significantly higher than in TG/GG genotype. On peg-IFN-α administration to chimeric mice, however, no significant difference in the median reduction of HCV-RNA levels and the induction of antiviral ISG was observed between favourable and unfavourable human hepatocyte genotypes.

Conclusions As chimeric mice have the characteristic of immunodeficiency, the response to peg-IFN-α associated with the variation in IL28B alleles in chronic HCV patients would be composed of the intact immune system.

INTRODUCTION

Hepatitis C is a global health problem that affects a significant portion of the world’s population. The WHO estimated that, in 1999, 170 million hepatitis C virus (HCV)-infected patients were present worldwide, with 3–4 million new cases appearing per year.1

The standard therapy for hepatitis C still consists of pegylated interferon-α (peg-IFN-α), administered once weekly, plus daily oral ribavirin for 24–48 weeks in countries where protease inhibitors are not available.2 This combination therapy is quite successful in patients with HCV genotype 2 or 3.
immuno-destruction of viral-infected hepatocytes. However, it viral replication as well as the second phase associated with modulation affects the viral kinetics during peg-IFN-α treatment. Whether the immune response by the peg-IFN-α human hepatocytes in chimeric mice and the response to decline during peg-IFN-α IL28B homozygous favourable allele had a more rapid decline of IL28B gene region are also associated with spontaneous HCV clearance. Genetic polymorphism in the IL28B gene region are also associated with spontaneous HCV clearance.11–12

Interestingly, a recent report showed the effect of genetic polymorphisms near the IL28B gene on the dynamics of HCV during peg-IFN-α plus ribavirin therapy in Caucasian, African American and Hispanic individuals,13 HCV-infected patients with the IL28B homozygous favourable allele had a more rapid decline of HCV in the first phase, which is associated with the inhibition of viral replication as well as the second phase associated with immuno-destruction of viral-infected hepatocytes.14 However, it is unknown how a direct effect by the IL28B genetic variation, such as the induction of IFN-stimulated genes (ISG) or cellular immune responses, would influence the viral kinetics during IFN treatment. Over recent periods, engineered severe combined immunodeficient (SCID) mice transgenic for urokinase-type plasminogen activator (uPA) received human hepatocyte transplants (hereafter referred to as chimeric mice)15–17 and are suitable for experiments with hepatitis viruses in vivo.18, 19 We have also reported that these chimeric mice carrying human hepatocytes are a robust animal model to evaluate the efficacy of IFN and other anti-HCV agents.20–21

The purpose of this study was to reveal the association between genetic variations in the IL28B gene region and viral decline during peg-IFN-α treatment in patients with HCV, and to clarify the association between different IL28B alleles of human hepatocytes in chimeric mice and the response to peg-IFN-α without immune response. These studies will elucidate whether the immune response by the IL28B genetic variation affects the viral kinetics during peg-IFN-α treatment.

**MATERIALS AND METHODS**

**Patients**

Fifty-four Japanese patients with chronic HCV genotype 1 infection at Nagasaki Medical Center and Nagoya City University were enrolled in this study (table 1). Patients received peg-IFN-α2a (180 μg) or 2b (1.5 μg/kg) subcutaneously every week and were administered a weight-adjusted dose of ribavirin (600 mg for <60 kg, 800 mg for 60–80 kg, and 1000 mg for >80 kg daily), which is the recommended dosage in Japan. Patients with other hepatitis virus infection or HIV coinfection were not included in the study. The study protocol conformed to the ethics guidelines of the 1975 Declaration of Helsinki as reflected by earlier approval by the institutions’ human research committees.

**Laboratory tests**

Blood samples were obtained before therapy, as well as on day 1 and at weeks 1, 2, 4, 8 and 12 after the start of therapy and were analysed for the HCV-RNA level by the commercial Abbott Real-Time HCV test with a lower limit of detection of 12 IU/ml (Abbott Molecular Inc., Des Plains, Illinois, USA). Genetic polymorphism in the IL28B gene (rs8099917), a single nucleotide polymorphism (SNP) recently identified to be associated with treatment response,6–8 was tested by the TaqMan SNP genotyping assay (Applied Biosystems, Foster City, California, USA).

**HCV infection of chimeric mice with the liver repopulated for human hepatocytes**

SCID mice carrying the uPA transgene controlled by an albumin promoter were injected with 5.0–7.5×10^7 viable hepatocytes through a small left-flank incision into the inferior splenic pole, thereafter chimeric mice were generated. The chimeric mice were purchased from PhoenixBio Co, Ltd (Hiroshima, Japan). Human hepatocytes with the IL28B homozygous favourable allele, heterozygous allele or homozygous unfavourable allele were imported from BD Biosciences (San Jose, California, USA) (table 2). Murine serum levels of human albumin and the body weight were not significantly different among four chimeric mice groups, providing a reliable comparison for anti-HCV agents.22 Three different serum samples were obtained from three chronic HCV patients (genotype 1b),21 22 Each mouse was intravenously infected with serum sample containing 10^7 copies of HCV genotype 1b. Administration of peg-IFN-α2a (Pegasys; Chugai Pharmaceutical Co., Ltd., Tokyo, Japan) at the dose formulation (30 μg/kg) was consecutively applied to each mouse on days 0, 3, 7 and 10 (table 3).

**HCV-RNA quantification**

HCV-RNA in mice sera (days 0, 1, 3, 7 and 14) was quantified by an in-house real-time detection PCR assay with a lower quantitative limit of detection of 10 copies/assay, as previously reported.21

**Quantification of IFN-stimulated gene-expression levels**

For analysis of endogenous ISG levels, total RNA was isolated from the liver using the RNeasy RNA extraction kit (Qiagen, Valencia, California, USA) and complementary DNA synthesis was performed using 2.0 μg of total RNA (High Capacity RNA-to-cDNA kit; Applied Biosystems). Fluorescence real-time PCR analysis was performed using an ABI 7500 instrument (Applied Biosystems) and TaqMan Fast Advanced gene expression assay (Applied Biosystems). TaqMan Gene Expression Assay primer and probe sets (Applied Biosystems) are shown in the supplementary information (available online only). Relative amounts of messenger RNA, determined using a FAM-Labeled TaqMan probe, were normalised to the endogenous RNA levels of the housekeeping reference gene, glyceraldehyde-3-phosphate dehydrogenase. The delta Ct method (2^−ΔΔCt) was used for quantification of relative mRNA levels and fold induction.23–24

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**Table 1 Characteristics of 54 patients infected HCV genotype 1**

<table>
<thead>
<tr>
<th>IL28B SNP rs8099917</th>
<th>TT (n=34)</th>
<th>TG (n=19) + GG (n=1)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>55.6±10.1</td>
<td>54.7±11.3</td>
<td>0.746</td>
</tr>
<tr>
<td>Gender (male %)</td>
<td>70</td>
<td>50</td>
<td>0.199</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>24.6±3.1</td>
<td>24.7±3.3</td>
<td>0.870</td>
</tr>
<tr>
<td>Viral load at therapy (log IU/ml)</td>
<td>6.0±0.7</td>
<td>5.8±0.8</td>
<td>0.357</td>
</tr>
<tr>
<td>SVR rate (%)</td>
<td>50</td>
<td>11</td>
<td>0.012</td>
</tr>
<tr>
<td>Serum ALT level (IU/l)</td>
<td>100.3±80.8</td>
<td>79.3±45.0</td>
<td>0.226</td>
</tr>
<tr>
<td>Platelet count (&lt;10^12/µl)</td>
<td>17.1±9.0</td>
<td>16.5±8.5</td>
<td>0.771</td>
</tr>
<tr>
<td>Fibrosis (F3-F4 %)</td>
<td>42</td>
<td>40</td>
<td>0.877</td>
</tr>
</tbody>
</table>

HCV, hepatitis C virus; SNP, single nucleotide polymorphism; SVR, sustained virological response.

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plus ribavirin therapy in patients infected with HCV genotype 1.

**RESULTS**

**Characteristics of the study patients**

Genotypes (rs8099917) TT, TG and GG were detected in 34, 19 and one patient infected with HCV genotype 1, respectively. SVR rates were significantly higher in HCV patients with genotype TT than in those with genotype TG/GG (50% vs 11%, p=0.012). The initial HCV serum load was comparable between genotypes TT and TG/GG (6.0±0.7 vs 5.8±0.8 log IU/ml). There were no significant differences in sex (male 6%, 70% vs 50%), age (55.6±10.1 vs 54.7±11.3 years), serum alanine aminotransferase level (100.3±80.8 vs 79.3±45.0 IU/L), platelet count (17.1±9.0 vs 16.5±5.8 ×10⁴/μl) and fibrosis stages (3.9±4%, 42% vs 40%) between HCV patients with the favourable (rs8099917 TT) and unfavourable (rs8099917 TG/GG) IL28B genotypes (table 1).

**Changes in serum HCV-RNA levels in patients treated by peg-IFN-α plus ribavirin**

Figure 1 shows the initial change in the serum HCV-RNA level for 14 days after peg-IFN-α plus ribavirin therapy in patients infected with HCV genotype 1 based on the genetic polymorphism near the IL28B gene. The immediate antiviral response (viral drop 24 h after the first IFN injection) was significantly higher in HCV patients with genotype TT than genotype TG/GG (−1.08 vs −0.39 log IU/ml, p<0.001). Figure 2 also shows the subsequent change in the serum HCV-RNA reduction after peg-IFN-α plus ribavirin therapy in patients infected with HCV genotype 1.

Similarly, during peg-IFN-α plus ribavirin therapy, a statistically significant difference in the median reduction in serum HCV-RNA levels was noted according to the genotype (TT vs TG/GG). The median reduction in the serum HCV-RNA levels (log IU/ml) at 1, 2, 4, 8 and 12 weeks between genotypes TT and TG/GG was as follows: −1.58 vs −0.62, p<0.001; −2.35 vs −0.91, p<0.001; −3.48 vs −1.56, p<0.001; −4.53 vs −2.37, p<0.01; −4.93 vs −2.86, p<0.001. Furthermore, the initial first-phase viral decline slope per day (Ph1/day) and subsequent second-phase viral decline slope per week (Ph2/week) in TT genotype were significantly higher than in genotype TG/GG (Ph1/day 0.94±0.83 vs 0.38±0.40 log IU/ml, p<0.001; Ph2/week 0.08±0.06 vs 0.04±0.03 log IU/ml, p<0.001) (figure 3).

**Changes in serum HCV-RNA levels in chimeric mice treated by peg-IFN-α**

In order to clarify the association between IL28B alleles of human hepatocytes and the response to peg-IFN-α, we prepared four lines of uPA/SCID mice and four different lots of human hepatocytes containing various rs8099917, rs8103142 and rs12979860 SNPs around the IL28B gene (table 2). The chimeric mice were inoculated with serum samples from each HCV-1b patient, and then HCV-RNA levels had increased and reached more than 10⁶ copies/ml in all chimeric mice sera at 2 weeks after inoculation. After confirming the peak of HCV-RNA in all chimeric mice, they were subcutaneously administered with four times injections of the bolus dose of peg-IFN-α2a for 2 weeks (table 3). Figure 4 shows the change in the serum HCV-RNA levels for 14 days during IFN injection into chimeric mice transplanted with IL28B favourable or unfavourable human hepatocyte genotypes. On peg-IFN-α administration, no significant difference in the median reduction in HCV-RNA levels in the serum A-infected chimeric mice sera was observed between favourable (n=7) and unfavourable genotypes (rs8099917 TT) and unfavourable (rs8099917 TG/GG) IL28B genotypes (table 1).

### Table 3: Dosage and time schedule of pegIFN-α2a+ treatment for HCV genotype 1b infected chimeric mice

<table>
<thead>
<tr>
<th>Donor hepatocytes†</th>
<th>No of chimeric mice</th>
<th>Inoculum</th>
<th>Test compound</th>
<th>Level (μg/kg)</th>
<th>Concentration (μg/ml)</th>
<th>Volume (ml/kg)</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>3</td>
<td>Serum A</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>B</td>
<td>4</td>
<td>Serum A</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>C</td>
<td>3</td>
<td>Serum A</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>D</td>
<td>3</td>
<td>Serum A</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>A</td>
<td>2</td>
<td>Serum B</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>C</td>
<td>2</td>
<td>Serum B</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
<tr>
<td>A</td>
<td>2</td>
<td>Serum C</td>
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<td>Serum C</td>
<td>Peg-IFN-α2a</td>
<td>30</td>
<td>3</td>
<td>10</td>
<td>Day 0, 3, 7, 10</td>
</tr>
</tbody>
</table>

†Pegasys; Chugai Pharmaceutical Co., Ltd., Tokyo, Japan.
†The IL28B genetic variation of the donor hepatocytes was indicated in table 2.
HCV, hepatitis C virus; peg-IFN-α; pegylated interferon α.
of representative antiviral ISG (eg, myxovirus resistance protein A, oligoadenylate synthetase 1, RNA-dependent protein kinase) and other ISG for promoting antiviral signalling (eg, Toll-like receptor 3, retinoic acid-inducible gene 1) were significantly induced at least 8 h after treatment, and prolonged at 24 h. No significant difference in ISG expression levels in HCV-infected livers was observed between favourable and unfavourable IL28B genotypes. The other inoculum for persistent infection of HCV-1b also demonstrated no significant difference in ISG expression levels between favourable and unfavourable IL28B genotypes (figure 5B,C). Interestingly, IFN-α expression levels by treatment of peg-IFN-α were significantly induced in HCV-infected human hepatocytes harbouring the favourable IL28B genotype (figure 5 A–C).

DISCUSSION

Several recent studies have demonstrated a marked association between the chronic hepatitis C treatment response and SNP (rs8099917, rs8099917, rs1031142 and rs12979860) near or within the region of the IL28B gene, which affected the viral dynamics during peg-IFN-α plus ribavirin therapy in Caucasian, African American and Hispanic individuals.

It has been reported that when patients with chronic hepatitis C are treated by IFN-α or peg-IFN-α plus ribavirin, HCV-RNA generally declines after a 7–10 h delay. The typical decline is biphasic and consists of a rapid first phase lasting for approximately 1–2 days during which HCV-RNA may fall 1–2 logs in patients infected with genotype 1, and subsequently a slower second phase of HCV-RNA decline. The viral kinetics had a predictive value in evaluating antiviral efficacy. In this study, biphasic decline of the HCV-RNA level during peg-IFN-α treatment was observed in both patients and chimeric mice infected with HCV genotype 1; however, in the first and second phases of viral kinetics, a difference between IL28B genotypes was observed only in HCV-infected patients; a more rapid decline in serum HCV-RNA levels after administering peg-IFN-α plus ribavirin was confirmed in patients with the TT genotype of rs8099917 compared to those with the TG/GG genotype.

On the other hand, in-vivo data using the chimeric mouse model showed no significant difference in the reduction of HCV-RNA titers in mouse serum among four different lots of human hepatocytes containing IL28B favourable (rs8099917 TT) or unfavourable (rs8099917 TG) genotypes, which was confirmed by the inoculation of two additional HCV strains. These results indicated that variants of the IL28B gene in donor hepatocytes had no influence on the response to peg-IFN-α under immunosuppressive conditions, suggesting that the immune response according to IL28B genetic variants could contribute to the first and second phases of HCV-RNA decline and might be critical for HCV clearance by peg-IFN-α-based therapy.

Two recent studies indeed revealed an association between the IL28B genotype and the expression level of hepatic ISG in human studies. Quiescent hepatic ISG before treatment among patients with the IL28B favourable genotype have been associated with sensitivity to exogenous IFN treatment and viral eradication; however, it is difficult to establish whether the hepatic ISG expression level contributes to viral clearance independently or appears as a direct consequence of the IL28B genotype. Another recent study addressed this question and the results suggested that there is no absolute correlation with the IL28B genotype and hepatic expression of ISG. Our results on the hepatic ISG expression level in immunodeficient chimeric mice also suggested that no significant difference in ISG expression levels was observed between favourable and unfavourable IL28B genotypes.
**IL28B** genotypes. However, these results were not consistent with a previous report using chimeric mice that the favourable **IL28B** genotype was associated with an early reduction in HCV-RNA by ISG induction. The reasons for the discrepancy might depend on the dose and type of IFN treatment, as well as the time point when ISG expression was examined in the liver. In addition, although IFN-λ transcript levels measured in peripheral blood mononuclear cells or liver revealed inconsistent results in the context of an association with the **IL28B** genotype, our preliminary assay on the **IL28A**, **IL28B** and **IL29** transcripts in the liver first indicated that the induction of IFN-λ on peg-IFN-α administration could be associated with the **IL28B** genotype. Therefore, the induction of IFN-λ followed by immune response might contribute to different viral kinetics and treatment outcomes in HCV-infected patients, because no difference was found in chimeric mice without immune response.

It has also been reported that the mechanism of the association of genetic variations in the **IL28B** gene and spontaneous clearance of HCV may be related to the host innate immune response. Interestingly, participants with seroconversion illness with jaundice were more frequently rs8099917 homozygous favourable allele (TT) than other genotypes (32% vs 5%, p=0.047). This suggests that a stronger immune response during the acute phase of HCV infection among patients with the **IL28B** favourable genotype would induce more frequent spontaneous clearance of HCV.

**Figure 3** (A) The first-phase viral decline slope per day (Ph1/day) and (B) second-phase viral decline slope per week (Ph2/week) in hepatitis C virus (HCV) genotype 1-infected patients treated with pegylated interferon α plus ribavirin. The lines across the boxes indicate the median values. The hash marks above and below the boxes indicate the 90th and 10th percentiles for each group, respectively.

**Figure 4** Median reduction of hepatitis C virus (HCV)-RNA levels (log copies/ml) after administering pegylated interferon α to chimeric mice having human hepatocytes containing various single nucleotide polymorphisms around the **IL28B** gene as favourable (rs8099917 TT) and unfavourable (rs8099917 TG) genotypes. Data are represented as mean+SD. Chimeric mice infected with a) serum A (n=7; favourable genotype, n=6; unfavourable genotype), b) serum B (n=2, each genotype), and c) serum C (n=2, each genotype). All serum samples were obtained from HCV-1b patients.
Taking into account both the above results in acute HCV infection and our results conducted on chimeric mice that have the characteristic of immunodeficiency, it is suggested that the response to peg-IFN-α associated with the variation in IL28B alleles in chronic hepatitis C patients would be composed of the intact immune system.

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Contributors YT and MM conceived the study. TW and FS and YT conducted the study equally. TW and FS coordinated the analysis and manuscript preparation. All the authors had input into the study design, patient recruitment and management or mouse management and critical revision of the manuscript for intellectual content. TW, FS and YT contributed equally.

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Competing interests None.

Patient consent Obtained.

Ethics approval This study was conducted with the approval of each ethics committee at the Nagoya City University and Nagasaki Medical Center (see supplementary information, available online only).

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