Productive Homologous and Non-homologous Recombination of Hepatitis C Virus in Cell Culture

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Abstract

Genetic recombination is an important mechanism for increasing diversity of RNA viruses, and constitutes a viral escape mechanism to host immune responses and to treatment with antiviral compounds. Although rare, epidemiologically important hepatitis C virus (HCV) recombinants have been reported. In addition, recombination is an important regulatory mechanism of cytopathogenicity for the related pestiviruses. Here we describe recombination of HCV RNA in cell culture leading to production of infectious virus. Initially, hepatoma cells were co-transfected with a replicating JFH1ΔE1E2 genome (genotype 2a) lacking functional envelope genes and strain J6 (2a), which has functional envelope genes but does not replicate in culture. After an initial decrease in the number of HCV positive cells, infection spread after 13–36 days. Sequencing of recovered viruses revealed non-homologous recombinants with J6 sequence from the 5′ end to the NS2–NS3 region followed by JFH1 sequence from Core to the 3′ end. These recombinants carried duplicated sequence of up to 2400 nucleotides. HCV replication was not required for recombination, as recombinants were observed in most experiments even when two replication incompetent genomes were co-transfected. Reverse genetic studies verified the viability of representative recombinants. After serial passage, subsequent recombination events reducing or eliminating the duplicated region were observed for some but not all recombinants. Furthermore, we found that inter-genotypic recombination could occur, but at a lower frequency than intra-genotypic recombination. Productive recombination of attenuated HCV genomes depended on expression of all HCV proteins and tolerated duplicated sequence. In general, no strong site specificity was observed. Non-homologous recombination was observed in most cases, while few homologous events were identified. A better understanding of HCV recombination could help identification of natural recombinants and thereby lead to improved therapy. Our findings suggest mechanisms for occurrence of recombinants observed in patients.

Citation: Scheel TKH, Galli A, Li Y-P, Mikkelsen LS, Gottwein JM, et al. (2013) Productive Homologous and Non-homologous Recombination of Hepatitis C Virus in Cell Culture. PLoS Pathog 9(3): e1003228. doi:10.1371/journal.ppat.1003228

Editor: Brett D. Lindenbach, Yale University, United States of America

Received May 23, 2012; Accepted January 21, 2013; Published March 28, 2013

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Funding: TKHS is supported by a Postdoctoral Fellowship and a Sapere Aude Research Talent award from The Danish Council for Independent Research. AG is the recipient of a Marie Curie International Reintegration Grant. The study was supported by research grants from Lundbeck Foundation (TKHS, AG, JM and JB), The Danish Cancer Society (YL, JM and JB), Novo Nordisk Foundation (YL, JM and JB), The Danish Medical Research Council (YL, JB), A. P. Møller and Chastine McKinney Møllers Medical Research Foundation (TKHS, JM and JB), Hvidovre Hospital Research Foundation (TKHS and JM), and Guangzhou Medical Research Foundation (AG). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

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Introduction

RNA viruses are rapidly adapting to their environment. The error-prone viral polymerases and the lack of proofreading mechanisms for most RNA viruses lead to high mutation rates. Genetic recombination between viral genomes is an additional mechanism increasing genetic diversity, which has proven to be epidemiologically relevant and allows RNA viruses to adapt to their surroundings [1]. Recombination could allow escape from natural or therapeutically induced immunity [2], or during antiviral treatment constitute an escape mechanism to antiviral compounds with an otherwise high barrier to resistance [3]. In addition, viral recombination has been associated with increased pathogenicity [4], and has caused the emergence of new human pathogens, such as Western equine encephalitis virus [5]. The use of live attenuated viral vaccines has led to re-emergence of disease due to recombination of vaccine strains with related viruses [6,7]; this remains a problem in poliovirus eradication. Thus, understanding the nature of viral recombination has general evolutionary implications, and might affect treatment and vaccination for important human pathogens.

Significant differences have been reported in recombination frequencies for different virus families, with high frequencies among Picornaviridae and lower frequencies among Flaviviridae and Alphaviridae [8]. Although hepatitis C virus (HCV) belongs to the Flaviviridae family, several epidemiologically important recombinant strains have been reported [9–11]. HCV constitutes a major public health burden with 130–170 million people chronically infected. Infection leads to increased risk of hepatitis, liver cirrhosis and hepatocellular carcinoma. The single positive-stranded HCV RNA genome of around 9600 nucleotides encodes one long open reading frame (ORF) flanked by 5′ and 3′ untranslated regions (UTRs). The HCV polyprotein is co- and post-translationally processed into structural (Core, E1 and E2), and nonstructural
Author Summary

Genetic recombination is the alternative joining of nucleic acids leading to novel combinations of genetic information. While DNA recombination in cells is of importance for evolution and adaptive immunity, RNA recombination often has only transient effects. However, RNA viruses are rapidly evolving and recombination can be an important evolutionary step in addition to mutations introduced by the viral polymerase. Recombination can allow escape from the host immune system and from antiviral treatment, and recombination of live attenuated viral vaccines has led to re-emergence of disease. Hepatitis C virus (HCV) is an important human pathogen that chronically infects more than 130 million worldwide and leads to serious liver disease. For HCV, naturally occurring recombinants are rare but clinically important. HCV recombination constitutes a challenge to antiviral treatment and can potentially provide an escape mechanism for the virus. In this study, we established an assay for HCV RNA recombination and characterized the emerging homologous and non-homologous recombinant viruses. Interestingly, recombination did not depend on viral replication, occurred most efficiently between isolates of the same genotype and did not occur with strong site-specificity. Better diagnosis of clinically important recombinants and an increased knowledge on viral recombination could strengthen antiviral and vaccine development.

20% and 2–10%, respectively, at the nucleotide and amino acid level. Genotypes, subtypes and isolates/strains differ at around 30%, grouped into seven major genotypes and many subtypes [12].

The epidemiologically most important HCV recombinant is the homologous recombinant of genotype 2k/1b that was first identified in St. Petersburg [13]. Since then, a number of naturally occurring inter- and intra-genotypic recombinants have been reported [9–11]; most inter-genotypic recombinants have junction in or close to the NS2 gene. Further, naturally occurring subgenomic deletion mutants have been described to persist in or close to the NS2 gene. Frequently, recombination between isolates of the same genotype and did not occur with strong site-specificity. Great HCV diversity is found among HCV isolates, which are grouped into seven major genotypes and many subtypes [12]. Genotypes, subtypes and isolates/stains differ at around 30%, 20% and 2–10%, respectively, at the nucleotide and amino acid levels.

Interestingly, recombination did not depend on viral replication, occurred most efficiently between isolates of the same genotype and did not occur with strong site-specificity. Better diagnosis of clinically important recombinants and an increased knowledge on viral recombination could strengthen antiviral and vaccine development.

PLOS Pathogens | www.plospathogens.org 2 March 2013 | Volume 9 | Issue 3 | e1003228

Results

Co-transfection of HCV genomes lacking viability in vitro led to productive non-homologous recombination

To study HCV recombination, an assay was established using the Huh7.5 hepatoma cell line. Since recombination of HCV is thought to be a relatively rare event, HCV genomes lacking viability in vitro were co-transfected to facilitate the identification of viable recombinants. RNA transcripts of the JFH1ΔE1E2 genome were transected alone or in combination with either the J6CF or J6/JFH1-GND genome (all genotype 2a, Figure 1). JFH1ΔE1E2 carries a partial deletion of the envelope genes, which allows replication but not viral particle production. The consensus full-length clone of the J6 isolate, J6CF, does not replicate in Huh7.5 cells [26] but has a functional 5’UTR-NS2 region in vitro [27], while the replication-deficient J6/JFH1-GND, carries an NS5B polymerase mutation in the viable J6/JFH1 background [28].

In all experiments, around 30% of cells were positive for HCV Core one day after transfection (Figure 2A); this percentage rapidly decreased due to lack of spread of infection and growth advantages of untransfected cells, as previously shown [29]. HCV RNA levels in the supernatant were comparable for all cultures during the first 8 days (Figure 2B) and no infectious particles were released from any of the cultures on day 3 and 6 (Figure 2C). An increase in percentage of HCV positive cells and HCV RNA levels was observed for the culture co-transfected with JFH1ΔE1E2 and J6CF from day 10 and infection spread to the almost entire culture on day 13. Similarly, infection spread to the majority of cells around day 36 in the culture co-transfected with JFH1ΔE1E2 and J6/JFH1-GND. After spread of infection in culture, infectivity titers of around 10^4 focus-forming units (FFU)/mL or 10^3 FFU/mL, respectively, were observed in supernatant from the two cultures (Figure 2C). After passage of supernatant from the J6CF co-transfected culture to naïve cells, HCV RNA titers above 10^3 IU/mL and infectivity titers around 10^5 FFU/mL were produced. Two additional co-transfections of JFH1ΔE1E2 and J6CF led to similar results, with spread of infection to the majority of the culture after 8 and 25 days, respectively.

To determine the nature of the infectious HCV genomes from the original co-transfection of JFH1ΔE1E2 with J6CF after passage to naïve cells, we performed direct sequencing of 12 overlapping PCR amplicons covering the entire ORF. While amplicons 1–2 (5’UTR-E2) had J6 sequence, amplicons 3–12 (E2-5’UTR) had JFH1 sequence; amplicons 2 and 3 contained overlapping sequence in E2 from both strains, which indicated the presence of a duplicated region. This was further analyzed for all
three cultures co-transfected with JFH1ΔE1E2 and J6CF by cloning of longer PCR amplicons and amplicons generated by inverted primer sets. The resulting sequences revealed non-homologous recombinant genomes with different genomic structures. The first recombinant had J6 sequence from the 5’UTR to nucleotide (nt) 2986 (NS2), recombined with JFH1ΔE1E2 from nt 872 (Core) to the 3’UTR (Rec#1; including the envelope deletion from nt 991 to 2040) (Figure 3). This recombination produced an in-frame non-homologous recombinant HCV ORF containing 1065 duplicated nts (355 amino acids) with a total predicted genome length of 10743 nts, compared to 9678 for JFH1 and 9711 for J6CF. A second recombinant had J6 sequence from the 5’UTR to nt 2070 (NS2), recombined with JFH1ΔE1E2 at nt 561 (Core) (Rec#2) (Figure 3). The third recombinant had breakpoint further downstream with J6 sequence from the 5’UTR to nt 4254 (NS3) joined to JFH1ΔE1E2 from nt 796 (Core) (Rec#3). The resulting genome had a predicted length of more than 12 kb, over 2400 nucleotides longer than natural HCV isolates. While this is longer than typical infectious HCV reporter constructs expressing fluorescent or luminescent markers [30], much longer BVDV recombinants (up to around 20 kb) were identified in similar cell culture recombination experiments [23].

It was previously demonstrated that the NS3 helicase contributes to the unique replication abilities of the JFH1 isolate [31]. Since this might have restricted the region of recombination in co-transfections of JFH1ΔE1E2 and J6CF, we investigated whether a different type of recombination event had occurred in the culture co-transfected with JFH1ΔE1E2 and J6/JFH1-GND, where both genomes carried an NS3 protein of JFH1 origin. After passage of viral supernatant to naïve cells, sequencing of the entire ORF from recovered viruses again showed J6 sequence for amplicons 1–2 and JFH1 sequence for amplicons 3–12. In further analysis, PCR amplicon clones covering the junction revealed a recombinant genome with J6/JFH1-GND sequence from the 5’UTR to nt 2971 (NS2), followed by JFH1ΔE1E2 from nt 860 (Core) to 3’UTR (Rec#4) (Figure 3), similar in structure to those already identified.

Recombination does not depend on a functional HCV polymerase

In the initial recombination assay, a replicating genome (JFH1ΔE1E2) was co-transfected with a non-replicating genome

![Figure 1. HCV genomes of strains J6 and JFH1 used for co-transfection experiments in the recombination assay. Genomes from the top panel were co-transfected with genomes from the bottom panel. Genomes are color coded according to isolate (J6: red, JFH1: blue). The black oval indicates replacement of 3’UTR sequence by an irrelevant cellular RNA sequence. Triangle denotes cleavage of pJ6CF by restriction enzyme; where no triangle is indicated plasmids were constructed with the HCV sequence shown. Details of individual genomes are given in Materials & Methods. doi:10.1371/journal.ppat.1003228.g001](http://www.plospathogens.org/)

![Figure 2. Co-transfection of JFH1ΔE1E2 and replication deficient genomes into Huh7.5 cells. HCV genomic RNA transcripts of JFH1ΔE1E2 were transfected alone or in combination with J6CF or J6/JFH1-GND. In addition, J6/JFH1-GND was transfected alone as a replication negative control. Cultures were followed until day 23, at which time the JFH1ΔE1E2 control had become negative; co-transfection of JFH1ΔE1E2 and J6/JFH1-GND was followed until day 41 and never became negative. (A) Percentage of HCV Core positive cells as determined by immunostainings. No positive cells were observed when J6/JFH1-GND was transfected alone. (B) HCV RNA titers (IU/mL) in supernatant after transfection. (C) Infectivity titers (FFU/mL) in supernatant after transfection. *Titrations were negative for all cultures on day 3 and 6. Other time points were not measured. doi:10.1371/journal.ppat.1003228.g002](http://www.plospathogens.org/)
Figure 3. Characteristics of recombined HCV genomes. For each observed recombination event (Rec#), the 30 nt sequence around the recombination breakpoint is shown for the parental 5' and 3' genomes. Grey shading indicates the sequence of the recombined genome. Conserved nucleotides around the junction site are shown as dots. In cases where breakpoints were located at stretches of conserved nucleotides in the two parental sequences, numbering is consistently done to include most of the 5' fragment and is indicated by space separation of the sequence.
Homologous (homol.) recombination events are indicated. The predicted total genome length is given, assuming that the recombination breakpoint was the only recombination event present. Schematic drawings of the genome structure of individual recombinants are shown. Junction identified by direct sequencing of PCR products. A Junction identified by sequencing of cloned fragments. One of seven clones contained an in-frame deletion of JFH1(5′-RNA)-EI. The same junction was subsequently also found for co-transfections of JFH1-A5’-p7 and JFH1-A5’-NS4A.

doi:10.1371/journal.ppat.1003228.g003

Viability of non-homologous recombinants confirmed by reverse genetic studies

To confirm that the identified non-homologous recombinants were viable, two representative clones, J6/JFH1-AE1E2(Rec#1) and J6/JFH1(Rec#10) were generated based on the original J6CF, JFH1-AE1E2 and JFH1 consensus clones. After transfection into Huh7.5 cells, J6/JFH1-AE1E2(Rec#1) and J6/JFH1(Rec#10) immediately spread in culture and produced infectivity titers greater than 10^6 FFU/mL (Figure 5). Similar infectivity titers were produced after passage of J6/JFH1-AE1E2(Rec#1) and J6/JFH1(Rec#10) supernatant to naïve cells. Sequencing of the entire ORF confirmed the identity of the replicating recombinants. J6/JFH1(Rec#10) did not acquire mutations, while J6/JFH1-AE1E2(Rec#1) had acquired A2071S and C2574R (A1712S and C2215R according to the H77 reference polyprotein, AF009606). These changes were not observed from the original co-transfected culture. Thus, the recombinated genomes were fully viable in cell culture and the initially identified genomic structures were confirmed.

Sequential recombination events observed after serial passage in culture

To determine whether sequential recombination events could occur on the same genome, we performed long term passaging of

![Figure 4. Measurement of translation from input RNA. To evaluate translation from input JFH1-A5’ RNA using luciferase reporter genomes, Huh7.5 cells were transfected with JFH1-M14444, J6/JFH1-RLucΔ40 (positive control for translation and replication), J6/JFH1-GND-RLucΔ40 (positive control for translation, negative control for replication) and J6/JFH1 (replicating, negative control for luciferase expression). Relative light units (RLU) of Renilla luminescence were measured at indicated time points and the mean and standard error of the mean of five replicates are shown. Differences in signal intensities at the individual time points were evaluated statistically using ANOVA with Bonferroni correction. Highly significant (p<0.0001) differences to JFH1-A5’-RLucΔ40 levels are indicated (**), other differences to JFH1-A5’-RLucΔ40 were not significant. doi:10.1371/journal.ppat.1003228.g004](image)
Recombination of HCV in Cell Culture

Figure 5. Transfection of the cloned recombinants J6/JFH1ΔE1E2(Rec#1) (A), or J6/JFH1(Rec#10) (B) in Huh7.5 cells. HCV genomic RNA transcripts were transfected and compared to J6/JFH1. The J6/JFH1-GND control remained negative throughout the experiment shown in (A). Percentage of HCV Core positive cells as determined by immunostainings (lines) and viral infectivity titers measured in supernatant (bars) are shown. doi:10.1371/journal.ppat.1003228.g005

Evaluation of recombination frequency between isolates of the same genotype

All 14 co-transfection experiments with J6 and JFH1-based genomes performed so far led to emergence of viable recombinants. To get a more quantitative understanding of recombination frequencies we re-plated cells co-transfected with JFH1AΔ5’ and J6176G into 96-well format before virus production was expected to occur. This would allow an estimation of recombination frequency between the genotype 2a isolates J6 and JFH1 over the Core-NS5A region. Through 22 days of follow up, 8 controls transfected with J6/JFH1 were positive, while recombination occurred in 4/72 (5.6%) co-transfected wells (Figure 7). Taking into account that 7000 cells were plated per well and that the transfection efficiency was 50% (assuming that co-transfection had the same efficiency as observed when evaluating NS5A positive cells one day post transfection of J6/JFH1) this equals to one productive recombination event for every 63,000 co-transfected cells, or recombination in 0.0016% of the cells.

Infrequent recombination between isolates of different genotypes

Intergenotypic recombinants were previously identified in vivo [9–11], and synthetic intergenotypic recombinants could establish infection in cell culture [32,33]. Thus, we next investigated whether recombination in vitro could also occur between isolates of different genotypes. Since efficient replication in the infectious cell culture system at the outset of this study relied on the JFH1 isolate, we co-transfected JFH1AE1E2 with consensus clones of genotype 1a (H77C and HC-TN), 1b (JHLS8), 3a (S52) or 4a (ED43) or with 3’ truncated versions (truncation in NS5B) of the same genomes. Similarly to J6CF, these clones are infectious in chimpanzees but cannot replicate in HuH7.5 cells [26]. RNA transcripts of JFH1AE1E2 were co-transfected with H77C, HC-TN (3 replicates each), H77CA3’, HC-TN3’, JHLS8, JHLS6A3’, S52, S52A3’, ED43 or ED43A3’ (1 replicate each). The percentage of HCV positive cells in most cultures was similar to transfection of JFH1AE1E2 alone, with a rapid decrease leading to no positive cells from around day 20. However, few HCV positive cells remained in the culture co-transfected with S52A3’ and infection eventually spread to the almost entire culture after 82 days (Figure 8). After passage of supernatant to naive cells, cloning of PCR amplicons identified intergenotypic non-homologous recombination events. Of 13 clones, 6 contained S52 sequence until nt 2823 (NS2) and JFH1 sequence from nt 2291 (E2) (Rec#1.1, Figure 3). The peak supernatant infectivity titer increased in passage 8, when the shorter Rec#1.1 genome dominated from passage 3 and dominated the virus population from passage 6 (Figure 6A and B). The peak supernatant infectivity titer increased in passage 8, where the shorter Rec#1.1 genome dominated (Figure 6A and C). In contrast, no changes occurred to the comparably short duplicated junction region of J6/JFH1(Rec#10) during 8 serial passages. Infectivity titers of almost 10^5 FFU/mL were observed in most passages for this apparently genomically stable recombinant (Figure 6C). Thus, sequential recombination events could take place in culture to eliminate long duplicated and presumably non-functional genome regions, apparently leading to increase of viral fitness.

the J6/JFH1ΔE1E2(Rec#1) and J6/JFH1(Rec#10) recombinants by serial inoculation of naive cells with supernatant from fully infected cultures. Interestingly, after three passages to naive cells a novel recombinant was detected in the J6/JFH1ΔE1E2(Rec#1) culture. The genetic structure of the new genome showed that an additional non-homologous recombination event had taken place and removed most of the duplicated region, resulting in a junction from nt 2823 (NS2) of J6 to nt 2638 (p7) of JFH1 (Rec#1.1, Figure 3). This second-generation recombinant was detectable from passage 3 and dominated the virus population from passage 6 (Figure 6A and B). The peak supernatant infectivity titer increased in passage 8, where the shorter Rec#1.1 genome dominated (Figure 6A and C). In contrast, no changes occurred to the comparably short duplicated junction region of J6/JFH1(Rec#10) during 8 serial passages. Infectivity titers of almost 10^5 FFU/mL were observed in most passages for this apparently genomically stable recombinant (Figure 6C). Thus, sequential recombination events could take place in culture to eliminate long duplicated and presumably non-functional genome regions, apparently leading to increase of viral fitness.
Figure 6. Characterization of sequential recombination events. After long-term passage in Huh7.5 cell culture a second sequential recombination event occurred for J6/JFH1ΔE1E2(Rec#1) but not for J6/JFH1(Rec#10). (A) PCR validation of the recombination region of Rec#1. A PCR was designed to cover the primary and secondary recombination events (see Materials & Methods). A Rec#1 type junction yielded an amplicon of 2321 nts (evident until passage 6), while a Rec#1.1 type junction yielded an amplicon of 1442 nts (evident from passage 6 onwards, and as early as passage 3 on long exposure images). Exact recombination sites are given in Figure 3. M, size marker. No size change was observed for amplicons covering the Rec#10 junction. (B) Schematic overview of recombinant types found in the original co-transfection experiment (J6/JFH1ΔE1E2(Rec#1)) and in passage 2–8 of the cloned Rec#1 to naïve cells. Regions within the PCR amplicon shown in (A) that were sequenced to reveal the recombinant junction are shown with blue bars; gaps (deletions) are shown with black lines. The genome structure included NS2/Core and E1/E2 fusion proteins for the original Rec#1 and an NS2/p7 fusion protein after the second recombination event. (C) Peak infectivity titers in serial passage of J6/JFH1ΔE1E2(Rec#1) and J6/JFH1(Rec#10) in culture. A representative titer after infection of naïve cells (passage 1) with J6/JFH1 is shown for comparison.

doi:10.1371/journal.ppat.1003228.g006

Figure 7. Emergence of positive recombinants in frequency experiment. Cells were transfected and 18 hours later distributed into 96-well format (7000 cells plated per well) to study recombination frequency. The number of HCV positive cells per well of replica staining plates plated ever 2–3 days (as indicated in Materials & Methods) was followed over time and is shown for the 8 J6/JFH1 positive controls and the 4/72 wells co-transfected with J6lı–7666 and JFH1Δ5’, where recombinants emerged. Contamination of these four cultures by J6/JFH1 was excluded by passaging of virus to naïve cells and sequencing the NS2/NS3 junction, except for one recombinant (*) that was too attenuated to efficiently re-infect naïve cells. Cell numbers below 10, corresponding to background, are not plotted. Decline in number of infected cells correlated with massive virus induced cell death.

doi:10.1371/journal.ppat.1003228.g007

Figure 8. Co-transfection of JFH1ΔE1E2 and replication deficient genomes of other HCV genotypes into Huh7.5 cells. HCV genomic RNA transcripts of JFH1ΔE1E2 were transfected alone or in combination with S52Δ3’ or J4L6ΔG540. Percentage of HCV Core positive cells as determined by immunostaining is shown. The JFH1ΔE1E2 culture was followed until day 35; no positive cells were observed after day 19 in this culture. For 16 other intergenotypic co-transfections, no infectious virus emerged and data similar to JFH1ΔE1E2 transfection alone were observed.

doi:10.1371/journal.ppat.1003228.g008
recombinants, J4/JFH1 and ED43/JFH1 [32,35]. While no recombination occurred in triplicate co-transfections with ED43-
TR77A;9TR77C, co-transfection with JFH1NS5BUTR resulted in spread of infection to the majority of cells 93 days post-transfection (Figure 6). After passage to naïve cells, sequencing identified intergenotypic non-homologous recombination. The replicating genome contained JFH1UTR, sequence from the 5’UTR to NS3 and JFH1 from NS2 to the 3’UTR, and carried the introduced mutation F886L (Rec#16, Figure 3). Thus, introduction of mutations conferring adaptation to synthetic intergenotypic JFH1-based Core-NS2 recombinants had only limited effect, on recombination frequency.

While all intragenotypic co-transfections performed with high input RNA led to emergence of viable recombinants, only two intergenotypic recombination events were identified from a total of 18 co-transfection experiments. Considering all co-transfection experiments with JFH1AE1E2 and different clones of other genotypes, an estimated generalized recombination frequency would be one productive recombination event per million co-transfected cells, or recombination in 0.0001% of the cells, taking into account two productive recombination events, the starting number of 400,000 cells in each of 18 experiments and an estimated average transfection efficiency of 30%.

Recombination sites were not restricted to specific regions of the HCV genome

The recombination events identified so far all had breakpoints in the p7-NS3 region of the 5’ fragment and the Core-NS2 region of the 3’ fragment. Due to the lack of functional envelope genes in the JFH1AE1E2 construct, many recombination breakpoints were however restricted from occurring further upstream. Likewise, due to the importance of the NS3 helicase for the unique replication abilities of the JFH1 isolate [31], breakpoints could be restricted from occurring further downstream of non-JFH1 genomes. To investigate whether recombination events could occur in other regions, we co-transfected JFH1Δ5’ with versions of J6CF truncated at nt 708, 1344, 2407, 2564, 2972 or 3479 (Figure 1). While no spread of infection was identified in two cultures (J6i-708 and J6i-2564), the majority of cells in the other cultures became infected after 13–22 days. Recombined genomes were identified after passage to naïve cells. Another case of homologous recombination was identified in the culture co-transfected with J6i-1344, occurring in the nt 858–883 region (Core) (Rec#17). The three other recombination events were non-homologous with junctions from E1 to Core (Rec#18; J6i-2407), a mixed population of 2878 (NS2)/2261 (E2) and 2901 (NS2)/2521 (E2) (Rec#19a/b; J6i-2972), and from NS2 to E2 (Rec#20; J6i-3479) (Figure 3). Thus, recombination of J6 and JFH1 occurred outside the NS2 region, even in the most upstream gene, Core.

Next, we wanted to determine whether recombination could occur downstream of NS3. Since JFH1 exhibits efficient function of the NS3-NS5B region in Huh7.5 cells, we transfected 5’ truncated transcripts of J6/JFH1 together with J6/JFH1/3’X, which carried the 5’UTR-NS2 from J6CF, NS3-3’UTR(polyU) from JFH1 and an irrelevant human mRNA sequence replacing the 3’X region (Figure 1). No HCV positive cells were observed when any of these genomes were transfected alone. Thus, J6/JFH1/3’X was co-transfected with J6/JFH1Δ5’ lacking the 5’UTR, J6/JFH1A(5’-p7), J6/JFH1A(5’-NS4A), J6/JFH1A(5’-NS4B), or J6/JFH1A(5’-NS5A). While no productive recombination occurred in the J6/JFH1A(5’-NS4B) co-transfected culture, infection spread in all other cultures after 8–17 days. Interestingly, identical recombinants were identified after passage of virus from all four positive cultures to naïve cells. The breakpoint was in NS5B from nt 9338 to nt 8517 (Rec#21) (Figure 3); this recombination took place in a region where 11 of 12 consecutive bases were conserved. Depending on the primers used, wild-type NS5B sequence could also be amplified from these cultures. Independent confirmation of the junction site by RT-PCR excluded cross contamination between the samples with identical breakpoint.

Since four identical recombinants were observed, we also cloned this recombinant type, J6/JFH1(Rec#21), and analyzed it in reverse genetic studies. Surprisingly, the input recombinant with the duplicated region could only be detected one day after transfection, while wild-type virus was detected thereafter. A silent mutation introduced in NS4B was amplified together with the duplicated region to exclude contamination. Thus, Rec#21 apparently resulted from one recombination event leading to a transient state, which was rapidly followed by a second recombination event leading to wild-type J6/JFH1 sequence. The presence of wild-type NS5B sequence also in the original cultures was in accordance with Rec#21 representing a transient state.

Thus, efficient recombination was demonstrated also in the 3’-end of the HCV genome. In co-transfections unbiased by the selection of HCV isolates [both J6/JFH1Δ5’ and J6/JFH1/3’X carried the complete J6/JFH1 ORF], a longer stretch of conserved nucleotides seemed to be preferred over the NS2 region for the recombination breakpoint.

Discussion

In this study, efficient HCV RNA recombination leading to robust virus production was demonstrated in cell culture. Most recombination events were non-homologous with large in-frame insertions of up to 2400 nucleotides, while fewer homologous events were identified. Almost all recombinants identified from replication defective genomes were of different nature, and we thus found no strong site specificity. Further, recombination occurred most efficiently between isolates of the same genotype. Most identified recombinants maintained at least one complete copy of each HCV protein and many recombinants carried two copies of one or more genes. It remains to be determined whether such duplications could produce two different functional protein copies, e.g. leading to viral particles carrying envelope proteins of different isolates or give any advantage to the virus. Only one recombinant type did not have at least one intact copy of all HCV genes (Rec#21). Though this recombinant type had an internal junction in NS5B, it carried an intact globular finger-palm-thumb structure followed by duplicated sequence and finally the C-terminal membrane anchor [36].

Interestingly, HCV RNA recombination did not depend on HCV replication as co-transfection of two replication incompetent genomes led to productive recombination (Rec#6-14 and #17-21). Further, the frequencies of recombination and the time until spread of infection in culture did not seem to differ between co-transfections with and without replication competent genomes. A non-replicative mechanism for HCV recombination is in agreement with findings in cell culture for the related BVDV [23] and for poliovirus [22]. This type of recombination was shown primarily to take place at single-stranded RNA structures [37], and it is hypothesized to occur through endoribonucleolytic cleavage and subsequent ligation of 3’-phosphate and 5’-hydroxyl partners. It remains to be determined by which mechanism(s) HCV recombination occurs in patients. The replicative copy-choice mechanism has previously been favored, since it is straightforward to envision how this strategy could produce the homologous recombinants observed in vivo. Accordingly, a model
that could explain the generation of the 2k/1b recombinant from St. Petersburg by template switching was previously suggested [30]. Here we demonstrated that homologous recombinants could be produced through a non-replicative mechanism (Rec#9 and Rec#17), which could represent an alternative or parallel pathway to replicative recombination in vivo.

After long-term passage in culture of the non-homologous recombinant J6/JFH1ÆE1E2(Rec#1) a more fit variant emerged, replacing the original replicating genome and leading to higher viral titers. This new recombinant resulted from a second recombination event and carried a duplication of only 106 nts compared to the original 1065 nts. Since the original recombinant was cloned and the second event occurred after a new transfection and subsequent cell-free passages, recombination must have occurred from the same genome or among genomes with identical structures and sequentially led to a more fit variant with a smaller insertion. Similar deletions of heterologous sequences have been observed in cytopathogenic BVDV genomes with heterologous sequences [39], and HCV genomes with inserted reporter genes in cell culture and in vivo [30], reflecting the virus ability to evolve and increase its fitness. Non-homologous recombinants have not been observed in patients [9–11], potentially due to strong fitness selection for homologous recombinants. However, non-homologous recombinants could represent precursors to more fit homologous recombinants through sequential recombination events, as we observed in reverse genetic experiments with J6/JFH1(Rec#21).

Co-transfections with two genomes of the same genotype led to productive recombination events in 22 of 25 experiments (86%) or 0.0016% of cells, whereas only 2 out of 18 (11%) or 0.0001% of cells in intergenotypic experiments led to productive recombination. Except for two cases of homologous recombination, all identified events were non-homologous. Reiter et al. previously described homologous recombination in the HCV replicon system [25]. However, since duplicated regions generated by non-homologous recombination between fragments of the same isolate could be obscured in direct sequencing from PCR products, non-homologous recombinants could possibly also have been occurring in that study. The recombination frequency in the replicon-based study ranged from one event per 3,000 to 30,000 cells, depending on the length of the genomic region available for recombination, or 0.003 to 0.03% of cells replicating wild-type replicons in parallel experiments [25]. This was slightly higher than frequencies observed in the present study, however in the replicon system, selection could allow less fit recombinants to survive and some recombination events might be compatible with replication but not with the complete viral life-cycle. In a study of cells infected with a non-cytopathogenic BVDV strain, which were subsequently transfected with a defective cytopathogenic genome, recombination events were observed in 33–58% of cultures when electroporated cells were plated in 24-well format, or roughly equivalent to one event per 0.001% of cells (assuming around 105 cells per culture) [23]. This was in the range of what was observed in the present study on HCV. A notable difference, however, is that for BVDV this occurred for a viable genome, while observation of similar recombination frequencies for HCV depended on two non-viable genomes. A direct comparison of frequencies is complicated, since recombination is thought to be affected by the length of the genomic region available for recombination [25], replication capacity and constraints on genome organization of productive recombinants. Studies with poliovirus and BVDV previously showed that the frequency of homologous recombination decreased with decreasing sequence homology between the RNA molecules [21,39], and that non-homologous recombination was the most frequent for recombination between different BVDV strains.

In this study, productive recombination more often took place between isolates of the same HCV genotype. The identification of several recombination events at a conserved nucleotide sequence in NS5B supports the importance of similar sequences for recombination to occur. Another explanation could be the higher functional compatibility between proteins of the same genotype expressed by the recombined RNA. The lack of sequence conservation at a number of recombination sites (Figure 3) indicates that sequence similarity is not a prerequisite for non-homologous recombination to occur. On the other hand, the high frequency of ambiguous nucleotides in recombination sites in this study (residues around the recombination site that are identical in the two parental sequences; Figure 3), indicates a role for primary sequences in dictating junction sites. Random joining would leave one ambiguous nucleotide in one of four recombination events, two ambiguous nucleotides in one of 16 events etc. Thus, the frequency of ambiguous nucleotides in cross over sites in this study is higher than expected. The low frequency of intergenotypic recombination events identified in this study is in some contradiction to the ratio of inter- and intragenotypic recombinants identified in patients [10]. However, since intragenotypic recombinants by nature are harder to define, their existence could be underestimated in the literature.

The recombination frequency calculations from the replicon study indicated that no recombination hotspots are present in the HCV genome [25]. This is in agreement with our findings that productive HCV recombination in the infectious cell culture system is not restricted to certain regions of the genome. However, several cases of recombination between two nearly identical 12 nt stretches in NS5B indicated some preference for conserved sequences. Interestingly, the experimental setup in the replicon study did not allow recombination to occur at this potential hotspot [25]. Recombination site specificity remains to be fully investigated in the absence of constraints using identical HCV isolates covering the entire genome. It could be speculated that some restrictions on recombination sites could apply at least to non-homologous recombination. Interestingly, all recombination sites identified in this study fall in regions where recombination of natural strains was also described [10].

All recombination events identified in the present study led to joining of viral RNA fragments. While insertion of cellular sequences has been reported for several other viruses [4,40–42], and is an important regulatory process for cytopathogenicity of the related BVDV [40,43], this has not been reported for HCV in vivo. However, by cell culture transfection of deletion mutants of stem loop I of the HCV 5’UTR, we previously recovered viable genomes that acquired RNA stem loop structures derived from viral or host sequences compensating for the deletion [27]. Now knowing that replication independent recombination is possible for HCV, these variants could have arisen by such a mechanism.

Non-homologous recombination could initiate important evolutionary steps in generation of novel types of viral genomes or cause diversity in genome regions tolerating insertions and deletions. Such productive non-homologous recombination events might potentially be followed by another recombination event to get rid of duplicate fitness-lowering sequences. The importance of RNA recombination for the evolution of RNA viruses is well documented [19], and many recently emerged human diseases are caused by viruses that display active recombination or reassortment [1,5]. The presence of reverse transcriptase could even fix such sequences in the cellular genome [44]. Thus, RNA
recombination could have played an important role in cellular and viral genetic evolution.

The prevalence of HCV recombinants in patients is relatively low, which could in part be caused by the super-infection exclusion principle [45,46], which would reduce the chance of having two different HCV strains replicating in the same cell. In vitro, the amount of replicating RNA is further expected to be much lower than the amounts of RNA present after co-transfection in vivo. Thus, the recombination frequency reported here could well be overrepresented compared to the in vivo setting. Further, fitness of novel recombinants in vivo should be high for the recombinant to eventually dominate over the parental strains. In a treatment setting this might however be accomplished, e.g. if parental genomes each carried resistance to one of two antiviral compounds in a combination therapy, with recombination leading to a double-resistant recombinant genome. Subgenomic deletion mutants [14,15] are naturally occurring in patients and are similar in structure to the JFH1AE1E2 construct used in this study. These could therefore constitute a reservoir of independent genomes that could potentially recombine with the wild-type to generate treatment-resistant or otherwise high-fitness genomes. With an increased knowledge on HCV recombination, better diagnosis of clinically important recombinants could become available, thereby facilitating selection of optimal therapeutic regimens for the patients. Our findings shed new light on how HCV recombination could occur in patients. Further, viral recombination might be an important escape mechanism to specific antiviral therapy in general, which could be important to consider in design of treatment regimens for certain viruses.

Materials and Methods

Plasmids

The HCV plasmids pJFH1AE1E2 [29], pJ6/JFH1 [28], pJ6/JFH1-GND [28], pJ6CF [47], pH7C [48], pHC-TN [19], pJH4L6S [50], pS52 [26] and pED43 [26] were previously described. Introduction of single mutations and construction of JFH1 [51]. For J4L6S and S52 intergenotypic recombination a scanning approach was used, [34,51]. Supernatants collected during experiments were sterile-filtered and stored at −80°C. HCV RNA titers were determined as previously described [51]. Infectivity titers were determined by adding 100 μL of triplicate sample dilutions (diluted 1:2 or more) to 6 × 10^5 Huh7.5 cells/well plated 24 hours before infection on poly-D-lysine-coated 96-well plates (Nunc). Cells were fixed 48 hours post-infection and immunostained for HCV following a previously established protocol [51] using anti-NS5A 9E10 as primary antibody [28]. FFU were determined as previously described [51]. Infectivity titers were determined by manual counting or by using an automated counter (ImmunoSpot Series 5 UV Analyzer, CTL Europe GmbH) with customized software, as previously described [26,52].

To analyze recombination frequency, cells were split 18 hours after transfection and 7000 cells were plated per well in 96-well format. Starting day 5, cells were split every 2–3 days using split ratios adjusted to ensure ≥50% confluence in all wells throughout the experiment. At each split a replica 96-well plate was plated and incubated for 2–3 days before staining as described above for infectivity titration. In this experiment, single infected cells were counted using the ImmunoSpot Series 5 UV Analyzer.

Luciferase translation assay

For luciferase assays, RNA was transfected into 10^5 Huh7.5 cells/well of 24-well plates. At indicated time points, cells were lysed for 15 min according to the Renilla Luciferase Assay System (Promega) protocol, and luciferase signals were measured in 5 replicates using optical bottom 96-well plates on a FluoStarOptima (BMG) plate reader.

Sequence determination of culture-derived HCV

HCV RNA was extracted from culture supernatant using High pure viral nucleic acid kit (Roche). For direct sequencing of the complete HCV ORF, reverse transcription, 1st round PCR covering the entire ORF and 12 overlapping 2nd round PCR amplifications were performed as previously described for J6/JFH1 [51]. For JH4L6S and S52 intergenotypic recombination events, primers designed for the corresponding JFH1-based recombinants were used [34,51]. Non-homologous recombination events that resulted in duplicated primer binding sites for the 2nd round PCR could not be identified using the direct sequencing approach. In these cases, additional 2nd round PCRs were set up with forward primers downstream of reverse primers (inverted primer sets), to specifically amplify the region containing a non-homologous recombination breakpoint with duplicated sequence. For supernatants originating from co-transfection of different HCV isolates, the initial 12 amplicon ORF direct sequencing was used to determine in which region such primers should be designed. For supernatants originating from co-transfection of RNA from the same HCV isolate, a scanning approach was used, in which inverted primer pairs placed for each ~500 nts were tested positive or negative by PCR. Amplified PCR bands were
sequenced to identify breakpoints of non-homologous recombination. In selected cases, and in cases where the recombination site could not be uniquely identified by direct sequencing, PCR products were TOPO-cloned (Invitrogen) and sequenced. The occurrence of sequential recombination events for Rec#1 over time was monitored by PCR using primers JF1848 (CTGTGTGTGGCCCAGTGTAC) and 2763R_J6 (AGCGTGAGCCCTGACGAACTGACG) on cDNA. The amplified product sizes varied according to the recombinant and thereby allowed differentiation. Sequence analysis was performed with Sequencher (Gene Codes Corp.).

Control experiments

As a control for correct identification of recombinant junctions, independent RNA extraction and RT-PCR were done on supernatant of selected recombinants (Rec#1, Rec#17 and all four cultures leading to Rec#21). This confirmed the identified breakpoints. It was further confirmed that recombinant-specific PCR products could be amplified from supernatant-derived cDNA and from cloned recombinant plasmids but not from pJ6/JFH1 using inverted primer sets JF2845 (ACACC CGGGTTATAAAGACC)/2111R_JFH1 (TGTACGTCCACGATGTTCTGGTG) (Rec#1) or JF1848 and the junction-specific reverse primer Rec10_R (GTGCGACAGTGCGTCTAGGCTCC-TATCGCCAGCATGACAG) (Rec#10). To exclude in vitro introduced recombination during T7-driven transcription or during reverse transcription after RNA extraction from supernatant, RNA produced by T7 transcription was subjected to 3 sequential rounds of DNaseI (Ferments) digestion using the RNaseq kit (Qiagen), mixed to yield combinations of 5’ and 3’ partners that previously led to successful recombination in cell culture, diluted to 50 pg (equivalent to around 10⁷ copies) and subjected to RT-PCR. PCR amplification using inverted primer sets JF2845/2111R_JFH1 on J6CF and JFH1AE1E2 RNA (Rec#1), JF1848 and the junction-specific reverse primer Rec10_R on J6CF and JFH1AE1E2 RNA (Rec#10), or inverted primer sets JR9606 (CAGATACACTGACCAGAGC)/JR9606 (TCCGTGAAAGCTCAGGTTC) on J6/JFH1/3’X and J6/JFH1(A5’-NS5A) RNA (Rec#21), did not lead to the specific ampiclons that were observed for the cloned plasmids of the respective recombinants. Further, the viable phenotypes of all cloned recombinants and the fact that all recombination events identified led to in-frame recombinant ORFs supported that RT-PCR-induced artifacts were not misleading our conclusions.

Acknowledgments

We thank A.-L. Sorensen and Labha Ghanem (CO-HEP, for general laboratory assistance and J. O. Nielsen and O. Andersen (Copenhagen University Hospital, Hvidovre) for valuable support. R. Purcell (NIH), T. Wakitani (National Institute of Infectious Diseases, Tokyo) and C. Rice (Rockefeller University) provided reagents.

Author Contributions

Conceived and designed the experiments: TKHS AG YL JMG JB. Performed the experiments: TKHS AGYL LSM JMG. Analyzed the data: TKHS AGYL JMG JB. Wrote the paper: TKHS AG JB.
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