Inhibition of hepatitis C virus replication by chloroquine targeting virus-associated autophagy

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Abstract

**Background** Autophagy has been reported to play a pivotal role on the replication of various RNA viruses. In this study, we investigated the role of autophagy on hepatitis C virus (HCV) RNA replication and demonstrated anti-HCV effects of an autophagic proteolysis inhibitor, chloroquine.

**Methods** Induction of autophagy was evaluated following the transfection of HCV replicon to Huh-7 cells. Next, we investigated the replication of HCV subgenomic replicon in response to treatment with lysosomal protease inhibitors or pharmacological autophagy inhibitor. The effect on HCV replication was analyzed after transfection with siRNA of ATG5, ATG7 and light-chain (LC)-3 to replicon cells. The antiviral effect of chloroquine and/or interferon-α (IFNα) was evaluated.

**Results** The transfection of HCV replicon increased the number of autophagosomes to about twofold over untransfected cells. Pharmacological inhibition of autophagic proteolysis significantly suppressed expression level of HCV replicon. Silencing of autophagy-related genes by siRNA transfection significantly blunted the replication of HCV replicon. Treatment of replicon cells with chloroquine HCV replication was analyzed after transfection with siRNA of ATG5, ATG7 and light-chain (LC)-3 to replicon cells. The antiviral effect of chloroquine and/or interferon-α (IFNα) was evaluated.

**Conclusions** Chloroquine and/or IFNα inhibited HCV replication in replicon cells. These findings may be useful for developing new therapeutic strategies for HCV infection.
suppressed the replication of the HCV replicon in a dose-dependent manner. Furthermore, combination treatment of chloroquine to IFNα enhanced the antiviral effect of IFNα and prevented re-propagation of HCV replicon. Protein kinase R was activated in cells treated with IFNα but not with chloroquine. Incubation with chloroquine decreased degradation of long-lived protein leucine.

**Conclusion** The results of this study suggest that the replication of HCV replicon utilizes machinery involving cellular autophagic proteolysis. The therapy targeted to autophagic proteolysis by using chloroquine may provide a new therapeutic option against chronic hepatitis C.

**Keywords** Autophagy · Autophagosome · HCV replicon · Chloroquine

**Introduction**

The genome of HCV, a member of the family Flaviviridae, consists of a positive-sense single-stranded RNA. Peg-interferon/ribavirin combination therapy, which is the most effective therapy against HCV infection, is effective in around 50% for genotype 1 and 80% for genotypes 2 and 3 [1–3], however, many people cannot tolerate the serious side effects and are resistant to Peg-interferon/ribavirin combination therapy. Difficulties in eradicating HCV are attributable to the limited number of treatment options against HCV [4, 5]. Therefore, the search for novel therapeutic agents remains a strong aspiration.

Autophagy is an evolutionarily conserved cellular pathway in which the cytoplasm and organelles are engulfed within double-membraned vesicles, known as autophagosomes. While cellular autophagy is thought to be in preparation for the turnover and recycling of cellular constituents [6–8], this process has been proposed as a mechanism of virus replication complex formation in positive-stranded RNA viruses including poliovirus, equine arteritis virus and coronavirus [9–12]. In these viruses, the replication complexes consist of double membrane vesicles in the cytoplasm, suggestive of an autophagosome origin [9, 12]. Recently, it was reported that transfection of HCV replicon induced autophagy [11]. Additionally, Sir et al. [13] demonstrated that the suppression of autophagy inhibited the replication of HCV. These findings suggested that the autophagy plays a pivotal role in HCV replication.

Chloroquine, which is widely used for the treatment of malaria, is a well-established inhibitor of autophagic proteolysis which acts by inhibiting acidification of lysosomes and endosomes [14]. It has been reported that chloroquine exerts direct antiviral effects on several RNA viruses including coronaviruses, flaviviruses and human immuno-deficiency virus (HIV) [8, 15–17]. Moreover, clinical studies have demonstrated the safety, tolerability, and efficacy of chloroquine in the antiviral treatment of HIV infection [18, 19]. Here, we have demonstrated that autophagic proteolysis plays a pivotal role on HCV replication, moreover, the inhibition of autophagic proteolytic pathways can constitute an effective new therapeutic target against HCV.

**Materials and methods**

Cell culture and treatment

Huh-7 cells were stably transfected with HCV replicon expressing chimeric protein of firefly luciferase and neomycin phosphotransferase [20, 21]. They were cultured in Dulbecco’s modified essential medium (DMEM) (Sigma, St. Louis, MO) supplemented with 10% foetal bovine serum (FBS) at 37°C under 5% CO2. To maintain cell lines carrying the HCV replicon, G418 (Wako, Osaka, Japan) was added to the medium at a final concentration 500 μg/ml.

**Luciferase assay**

Luciferase activities were quantified to evaluate the replication of HCV replicon by a luminometer (Lumat LB9507; Berthold, Germany) using a Bright-Glo Luciferase Assay System (Promega, Madison, WI). Assays were performed in triplicate, and the results were expressed as mean ± SD as percents of controls.

**Cell viability assay**

The viability of cells was assessed by WST-1 assay. Cells were cultured in 96-well plates at 5 × 10³/well for 24 h, and then treated with 3-methyladenine [22] (10 mM), mixture of E64d (1 μg/ml) and pepstatin A [23] (1 μg/ml), and chloroquine (10⁻⁶–10⁻³ M) for 18 h. Cell proliferation reagent WST-1 (Roche, Swiss) was added to each well, and the cells were incubated for another 1 h at 37°C. The absorbance was measured against a background control by microplates reader (SPECTRA max 340PC, Molecular Devices, Sunnyvale, CA) at 450 nm. The reference wavelength was 650 nm.

**Inhibition of autophagy and replication of HCV replicon**

Cells were treated with 3-methyladenine (10 mM) or mixture of E64d (1 μg/ml) and pepstatin A (1 μg/ml), chloroquine (10⁻⁶–10⁻³ M), interferon (IFN)α (100 U/ml) for 18 h, the levels of replication of HCV replicon were assessed by luciferase assay. Moreover, cells were cultured...
with chloroquine (10^{-5} \text{ M}) and/or IFN\(_\alpha\) (100 U/ml) for 7 days, then continued to incubate without drugs for another 21 days. Replication levels of HCV replicon were determined by luciferase assay at 7th and 21st days from cessation of drugs.

Identification of autophagosomes

Naïve Huh7 cells, Huh7/Rep-Feo cells, and Huh7/Rep-Feo treated with IFN\(_\alpha\) for 14 days were seeded on 30 mm dishes and incubated for 48 h. In addition, Huh7/Rep-Feo cells were treated with chloroquine (10^{-5} \text{ M}) for 18 h. Cells were prefixed with 2\% glutaraldehyde, post-fixed with 1\% osmic acid, dehydrated in graded ethanol, embedded in resin, and cut into sections on an ultramicrotome. The cells were analyzed by a transmission electron microscope (Hitachi H7100, Japan). The number of autolysosomes in 100 \mu m^2 of cytoplasm was counted by using transmission electron microscopy.

Small interfering RNA knockdown of ATG5, 7, LC-3

A combination of four chemically synthesized siRNA duplex molecules targeted to the human ATG5, 7, LC-3\_\alpha, LC-3\_\beta mRNA sequence (Dharmacon, Lafayette, CO) was transiently transfected (final concentration 50 nM) into Huh7/Rep-Feo cells using a transfection reagent (Dharmacon, Lafayette, CO). siRNA targeted to enhanced green fluorescence protein was used as a control. Forty-eight hours after transfection, levels of HCV replication were analyzed by luciferase assay.

Western blot analysis

Twenty-five micrograms of total cell lysates were subjected to SDS/PAGE on a 10\% gradient gel and electro-photoretically transferred onto polyvinylidene fluoride membranes. After blocking with 5\% non-fat dry milk in Tris-buffered saline, membranes were incubated with primary rabbit monoclonal antibody against Phospho-protein kinase R (P-PKR) (Cell Signaling Technology, Danvers, MA) or light-chain 3 (LC3), followed by a secondary horseradish peroxidase (HRP)-conjugated anti-rabbit IgG antibody (Cell Signaling Technology, Danvers, MA). Subsequently, specific bands were visualized using ECL detection kit (Amersham Pharmacia Biotech, Midland, ON, Canada).

Protein degradation assay

Long lived protein is mainly degraded by autophagy [24]. Cells were incubated with Williams’ E/10\% FBS containing 0.5 \mu Ci/ml \[^{14}\text{C}]\text{leucine for 24 h to label long-lived proteins. Cells were washed with Williams’ E/10\% FBS containing 10^{-5} M of unlabeled leucine and incubated with the medium for 2 h to allow degradation of short-lived proteins and minimize the incorporation of labeled leucine. The cells were then washed with phosphate-buffered saline (PBS) and incubated at 37^\circ C with Williams’ E/10\% FBS in the presence or absence of chloroquine (10^{-5} M). After 4 h, aliquots of the medium were taken and a one-tenth volume of 100\% trichloroacetic acid was added to each aliquot. The mixtures were centrifuged at 12,000 \text{g} for 5 min, and the acid-soluble radioactivity was determined using a liquid scintillation counter. At the end of the experiment, the cultures were washed twice with PBS, and 1 ml of cold trichloroacetic acid was added to fix the cell proteins. The fixed cell monolayers were washed with trichloroacetic acid and dissolved in 1 ml of 1 N NaOH at 37^\circ C. Radioactivity in an aliquot of 1 N NaOH was determined by liquid scintillation counting. The percentage of protein degradation was calculated according to published procedures [25].

Fig. 1 Expression of autophagy is changed by presence or absence of HCV replicon. a Naïve Huh7 cells, Huh7/Rep-Feo cells, and Huh7/Rep-Feo treated with IFN\(_\alpha\) for 14 days were seeded on 30 mm dishes and incubated for 48 h. The cells were analyzed by a transmission electron microscopy. Autophagosomes (arrow heads) were detected by transmission electron microscopy. b The number of autolysosomes in 100 \mu m^2 of cytoplasm was counted by using transmission electron microscopy.
Statistical analysis

Differences were compared using ANOVA. Basically $P$ values less than 0.05 were considered as statistically significant.

Results

The inhibition of autophagy suppressed replication of HCV replicon

We counted numbers of autophagosome and autolysosome in cells transduced with HCV replicon Rep-Feo by using electron microscopy. Double membrane vesicles with the morphology of autophagosomes were identified at 2.3 vacuoles/cells in naïve Huh-7 cells, while transfection of HCV replicon increased the number of vacuoles to about fourfold over untransfected Huh-7 cells (Fig. 1a, b).

Subsequent treatment of the cells with IFNα (100 U/ml) for 14 days to eliminate HCV replicon substantially decreased the autophagolysosome in cytoplasm of Huh7/Rep-Feo cells (Fig. 1a, b). These observations suggested that HCV replicon induces formation of autophagosomes. To clarify the role of autophagy on the replication of HCV, Huh7/Rep-Feo cells were treated with 3-methyladenine (10 mM) or a mixture of E64d (1 µg/ml) and pepstatin A (1 µg/ml) for 18 h. Cell proliferation reagent WST-1 was added to each well, and the cells were incubated for 1 more hour at 37°C. The absorbance was measured against a background control by microplates reader at 450 nm. The reference wavelength was 650 nm. A combination of four chemically synthesized siRNA duplex molecules targeted to the human ATG5, 7, LC-3α, LC-3β mRNA sequence was transiently transfected into Huh7/Rep-Feo cells using a transfection reagent. siRNA targeted to enhanced green fluorescence protein was used as the control. Forty-eight hours after transfection, levels of HCV replication were analyzed by luciferase assay.
E64d and pepstatin A did not affect cell viability (Fig. 2b). To clarify the role of autophagy induction in the replication of HCV, we suppressed the induction of autophagy by silencing autophagy-related genes (ATG5, ATG7, LC-3α and LC-3β) by siRNA transfection. Silencing of autophagy-related genes reduced the replication of HCV replicon to about 70% of control (Fig. 2c). Transfection with siRNA of autophagy related genes decreased the number of autophagosomes to about 30% of control. These results indicated that autophagy plays a pivotal role in replication of HCV.

Chloroquine inhibits the replication of HCV replicon

Next, we evaluated the anti-HCV effect of chloroquine, which is a lysosomotropic agent that raises intralysosomal pH and impairs autophagic protein degradation. To assess the effects of chloroquine on the intracellular replication of the HCV replicon, Huh7/Rep-Feo cells were cultured with various concentrations of chloroquine in the medium. The replication of the HCV replicon was increased to about twofold within 18 h in the control media, however, which was suppressed by chloroquine in a dose-dependent manner (Fig. 3a). Next, cytotoxicity of chloroquine was analysed by WST-1 assays. Huh7/Rep-Feo cells treated with chloroquine showed no significant effect on cell viability in doses of lower than 10⁻⁵ M (Fig. 3b). However, incubation with 10⁻⁴ M of chloroquine reduced the cell viability to 25% of control. On the basis of the toxicity curve, the IC₅₀ of the drug was calculated to be 3.6 × 10⁻⁵ M (Fig. 3c). The average EC₅₀ of chloroquine was calculated as 2.2 × 10⁻⁷ M (Fig. 3c). The replication of HCV replicon was suppressed to nearly 40% of control at 10⁻⁵ M of chloroquine, which did not affect cell viability. These data indicated that chloroquine efficiently inhibited the replication of HCV replicon in the absence of toxic effect to cells at the concentration of 10⁻⁵ M. Accordingly, we used 10⁻⁵ M of chloroquine for the following study.

Next, we conducted the following assay to determine the synergistic inhibitory effect of chloroquine to IFNα on HCV replication. Treatment with chloroquine for 18 h
resulted in a significant decrease of HCV replicon to about 40% of control. On the other hand, incubation with IFNα for 18 h inhibited the replication of HCV replicon to the levels about 15% of controls as expected. However, co-incubation with 100U/ml of IFNα and 10⁻⁵ M of chloroquine further decreased HCV replication significantly (Fig. 4a).

To determine whether long-term chloroquine treatment inhibits post-treatment re-propagation of HCV replicon, we followed up luciferase activity of the cells at the 7th and 21st days after 7 days of treatment with chloroquine and/or IFNα (Fig. 4b). In HCV replicon cells treated by chloroquine, luciferase activities recovered to 53 and 88% on 7 and 21 days after cessation of treatment. In cells that were treated by IFNα, luciferase activity maintained background level for 7 days post-treatment. However, it reappeared in 21 days. In sharp contrast, co-incubation with IFNα and chloroquine for 7 days suppressed HCV replication for the extensive period up to 21 days, even in the absence of these drugs (Fig. 4b).

Anti-HCV effect of chloroquine independent of IFN signaling pathway

IFN-inducible double-stranded RNA-activated protein kinase R (PKR) plays a key antiviral role against hepatitis C virus [26, 27]. To elucidate the mechanisms of the inhibitory effect of chloroquine on HCV replication, phosphorylated PKR (P-PKR) was evaluated by western blotting analysis. P-PKR was detectable in cells treated with IFNα after 24 h; this increase in P-PKR expression peaked at 24 h after IFNα treatment and was reduced at 48 h (Fig. 5a). In contrast, P-PKR was not observed in cells treated with chloroquine at any time point.

Chloroquine blunts autophagic proteolysis in cells transfected with HCV replicon

It is reported that chloroquine disrupts lysosomal function, preventing effective autophagic protein degradation, leading to the accumulation of ineffective autophagosomes [28]. Therefore, we investigated if chloroquine led to the accumulation of autolysosomes as a result of suppression of proteolysis. We performed electron microscopic investigation to evaluate quantities of autophagosomes and autolysosomes. Ultrastructural analysis identified 0.94 ± 0.1 vacuoles/100 μm² of autolysosomes in control cells; however, treatment with chloroquine increased the number of autolysosomes dramatically to about 13-fold over control (Fig. 5b). Furthermore, the molecular form of LC3 protein of the cells, which is a component of autophagosomes, was examined by western blot analysis to ensure that chloroquine treatment leads to the accumulation of autophagosomes and autolysosomes. As shown in Fig. 5c, immunopositive protein bands for LC3-I and LC3-II forms were clearly evident in control cells [⁺⁺P < 0.05 vs. IFNα treatment group (day 28) by ANOVA].
Discussion

Previous reports have disclosed that autophagy plays a pivotal role on the replication of several RNA viruses [10–12]. Our present results demonstrate that autophagy is induced by transfection of HCV replicon and is reduced by deletion of replicon due to IFNα (Fig. 1a, b). These results suggest that autophagy is induced in the presence of HCV replication in its host cells. However, the role of autophagy in the pathogenesis of HCV is largely unclear. We found that the inhibition of autophagosome formation and autophagic proteolysis blunt the replication of genotype 1b subgenomic HCV replicon (Fig. 2a, c). Sir et al. [13] reported that inhibition of autophagy also reduced the replication of the JFH1-based full length genotype 2a genome. Therefore, the utilization of autophagy on viral replication is shown by HCV strains across different genotypes.

On the other hand, not only a silencing of autophagic gene but also pharmacological inhibition of autophagic proteolysis possesses anti-HCV effects (Fig. 2a, c). However, treatment with both chloroquine and the mixture of E64d and pepstatin induced the accumulation of...
autophagosomes in cytoplasm. Therefore, it is likely that HCV does not utilize the double membrane structure as the localization of the viral replication formation. These results support the hypothesis that protein degradation due to autophagy is important for HCV replication.

Chloroquine is a well-known inhibitor of autophagic protein degradation and is often used as an anti-malarial agent. Moreover, the anti-viral effect of chloroquine on other RNA viruses has been already reported in clinical trials [15, 16]. In our results, chloroquine inhibits the intracellular replication of an HCV replicon in a dose-dependent manner (Fig. 3a). This antiviral effect of chloroquine was clearly not due to cytotoxic effects (Fig. 3b). Moreover, chloroquine possesses a synergistic effect with IFNz on HCV replication (Fig. 4a). Although IFNz possesses strong anti-HCV effects, re-propagation of HCV replicon was observed after 3 weeks following 7 days of treatment with IFNz. Interestingly, co-incubation with IFNz and chloroquine for 7 days prevented re-propagation of HCV replicon (Fig. 4b). Chloroquine is a lysosomal weak base that is known to affect acid vesicles leading to dysfunction of several proteins [29]. It was demonstrated that disruption of lysosomal function impairs maturation of viruses through inhibiting the low-pH dependent proteases in trans-Golgi vesicles in HIV and the SARS coronavirus infection in vitro [15, 29]. However, little is understood about the mechanism of its antiviral effect. In previous reports, various drugs which possess inhibitory effects on the replication of HCV and have a synergistic action with IFNz have been proposed as new therapeutic agents to treat HCV. Some of them have proved to exhibit their anti-HCV effects through augmentation of IFNz-induced antiviral gene responses [30, 31]. However, the anti-HCV effect of chloroquine was not associated with activation of one of IFN receptors signaling molecule PKR (Fig. 5a). Our results showed chloroquine induced the accumulation of ineffective autophagosomes in cytoplasm of Huh7/Rep-Feo cells (Fig. 5b) and inhibited the degradation of long-lived protein leucine (Fig. 6). These findings imply that chloroquine effectively impairs the function of autophagy in our experiment. These results indicated that chloroquine is a new anti-HCV agent that targets the autophagic proteolysis.

Previous reports have shown that chloroquine possesses anti-viral effects on various RNA viruses. Its best-studied effects are those against HIV replication, which are being tested in clinical trials [17, 18]. HCV co-infection is common in HIV-positive patients in USA and Europe [32, 33]. Since HIV infection accelerates the progression of HCV-related liver disease, treatment of HCV is generally recommended. However, co-infected patients have a greater risk of antiretroviral therapy-associated hepatotoxicity than patients with HIV only [34]. Moreover, treatment with ribavirin is believed to increase the risk of anemia in patients taking the HIV drug zidovudine [35]. A clinical study designed for HIV patients showed the safety and efficacy of chloroquine used for long terms up to 48 weeks [36]. Therefore, the combination therapy of interferon and chloroquine is, possibly, a hopeful therapy for HCV–HIV co-infected patients. Since chloroquine is known as one of the inexpensive drugs, therefore, chloroquine might provide a new effective, safe and economical therapeutic option for patients with HCV. In conclusion, autophagic proteolysis might be a new therapeutic target on the replication of HCV.

References

1. Yotsuyanagi H, Koike K. Drug resistance in antiviral treatment for infections with hepatitis B and C viruses. J Gastroenterol. 2007;42:329–35.
2. Bruno S, Stroffolini T, Colombo M, Bollani S, Benvegnú L, Mazzella G, et al. Sustained virological response to interferon-alpha is associated with improved outcome in HCV-related cirrhosis: a retrospective study. Hepatology. 2007;45:579–87.
3. Shiffman ML, Cooksley WG, Dusheiko GM, Lee SS, Balart L, Reindollar R, et al. Peginterferon alfa-2a in patients with chronic hepatitis C and cirrhosis. N Engl J Med. 2000;343:1673–80.
4. Sezaki H, Suzuki F, Kawamura Y, Yatsuji H, Hosaka T, Akuta N, et al. Poor response to pegylated interferon and ribavirin in older women infected with hepatitis C virus of genotype 1b in high viral load. Dig Dis Sci. 2009;54:1317–24.
5. Honda T, Katano Y, Urano F, Murayama M, Hayashi K, Ishigami M, et al. Efficacy of ribavirin plus interferon-alpha is associated with improved outcome in HCV-related cirrhosis: a retrospective study. Hepatology. 2000;29:1717–21.
6. Reggiori F, Klionsky DJ. Autophagy in the eukaryotic cell. Eukaryot Cell. 2002;1:11–21.
7. Prentice E, Jerome WG, Yoshimori T, Mizushima N, Denison MR. Coronavirus replication complex formation utilizes components of cellular autophagy. J Biol Chem. 2004;279:10136–41.
8. Randolph VB, Winkler G, Stollar V. Acidotropic amines inhibit proteolytic processing of flavivirus prM protein. Virology. 1990;174:450–8.
9. Posthuma CC, Pedersen KW, Lu Z, Joosten RG, Roos N, Zevenhoven-Dobbe JC, et al. Formation of the arterivirus replication/transcription complex: a key role for nonstructural protein 3 in the remodeling of intracellular membranes. J Virol. 2008;82:4480–91.
10. Sir D, Chen WL, Choi J, Wakita T, Yen TS, Ou JH. Induction of incomplete autophagic response by hepatitis C virus via the unfolded protein response. Hepatology. 2008;48:1054–61.
14. Poole B, Ohkuma S. Effect of weak bases on the intralysosomal pH in mouse peritoneal macrophages. J Cell Biol. 1981;90:665–9.
15. Vincent MJ, Bergeron E, Benjannet S, Erickson BR, Rollin PE, Ksiazek TG, et al. Chloroquine is a potent inhibitor of SARS coronavirus infection and spread. Virol J. 2005;2:69.
16. Savarino A, Gennero L, Chen HC, Serrano D, Malavasi F, Boelaert JR, et al. Anti-HIV effects of chloroquine: mechanisms of inhibition and spectrum of activity. AIDS. 2001;15:2221–9.
17. Savarino A, Boelaert JR, Cassone A, Majori G, Cauda R. Effects of chloroquine on viral infections: an old drug against today’s diseases? Lancet Infect Dis. 2003;3:722–7.
18. Sperber K, Chiang G, Chen H, Ross W, Chusid E, Gonchar M, et al. Comparison of hydroxychloroquine with zidovudine in asymptomatic patients infected with human immunodeficiency virus type 1. Clin Ther. 1997;19:913–23.
19. Paton NI, Aboulhab J. Hydroxychloroquine, hydroxyurea and didanosine as initial therapy for HIV-infected patients with low viral load: safety, efficacy and resistance profile after 144 weeks. HIV Med. 2005;6:13–20.
20. Tanabe Y, Sakamoto N, Enomoto N, Kurosaki M, Ueda E, Maekawa S, et al. Synergistic inhibition of intracellular hepatitis C virus replication by combination of ribavirin and interferon-alpha. J Infect Dis. 2004;189:1129–39.
21. Yokota T, Sakamoto N, Enomoto N, Tanabe Y, Miyagishi M, Maekawa S, et al. Inhibition of intracellular hepatitis C virus replication by synthetic and vector-derived small interfering RNAs. EMBO Rep. 2003;4:602–8.
22. Seglen PO, Gordon PB. 3-Methyladenine: specific inhibitor of autophagic/lysosomal protein degradation in isolated rat hepatocytes. Proc Natl Acad Sci USA. 1982;79:1889–92.
23. Ueno T, Ishidoh K, Mineki R, Tanida I, Murayama K, Kadowaki M, et al. Autolysosomal membrane-associated betaine homocysteine methyltransferase. Limited degradation fragment of a sequestered cystolic enzyme monitoring autophagy. J Biol Chem. 1999;274:15222–9.
24. Klionsky DJ, Emr SD. Autophagy as a regulated pathway of cellular degradation. Science. 2000;290:1717–21.
25. Gronostajski RM, Pardee AB. Protein degradation in 3T3 cells and tumorigenic transformed 3T3 cells. J Cell Physiol. 1984;119:127–32.
26. Clemens MJ. PKR—a protein kinase regulated by double-stranded RNA. Int J Biochem Cell Biol. 1997;29:945–9.
27. Gale MJ Jr, Korth MJ, Tang NM, Tan SL, Hopkins DA, Dever TE, et al. Evidence that hepatitis C virus resistance to interferon is mediated through repression of the PKR protein kinase by the nonstructural 5A protein. Virology. 1997;230:217–27.
28. Glaumann H, Ahlberg J. Comparison of different autophagic vacuoles with regard to ultrastructure, enzymatic composition, and degradation capacity—formation of crinosomes. Exp Mol Pathol. 1987;47:346–62.
29. Thorens B, Vassalli P. Chloroquine and ammonium chloride prevent terminal glycosylation of immunoglobulins in plasma cells without affecting secretion. Nature. 1986;321:618–20.
30. Dev A, Patel K, McHutchison JG. New therapies for chronic hepatitis C virus infection. Curr Gastroenterol Rep. 2004;6:77–86.
31. Lin K, Kwong AD, Lin C. Combination of a hepatitis C virus NS3-NS4A protease inhibitor and alpha interferon synergistically inhibits viral RNA replication and facilitates viral RNA clearance in replicon cells. Antimicrob Agents Chemother. 2004;48:4784–92.
32. Staples CT Jr, Rimland D, Dudas D. Hepatitis C in the HIV (human immunodeficiency virus) Atlanta V.A. (Veterans Affairs Medical Center) Cohort Study (HAVACS): the effect of coinfection on survival. Clin Infect Dis. 1999;29:150–4.
33. Denis F, Adjide CC, Rogez S, Delpeyrroux C, Rogez JP, Weinbreck P. Seroprevalence of HBV, HCV and HDV hepatitis markers in 500 patients infected with the human immunodeficiency virus. Pathol Biol (Paris). 1997;45:701–8.
34. Sulkowski MS, Benhamou Y. Therapeutic issues in HIV/HCV-coinfected patients. J Viral Hepat. 2007;14:371–86.
35. Alvarez D, Dieterich DT, Brau N, Moorehead L, Ball L, Sulkowski MS. Zidovudine use but not weight-based ribavirin dosing impacts anaemia during HCV treatment in HIV-infected persons. J Viral Hepat. 2006;13:683–9.
36. Paton NI, Aboulhab J, Karim F. Hydroxychloroquine, hydroxyurea and didanosine as economic treatment for HIV-1. Lancet. 2002;359:1667–8.