

has no inhibitory effect on CKI activity (62). Thus, it appears that two or more kinds of serine/threonine-specific protein kinases, including CKI- $\alpha$ , participate in NS5A phosphorylation that is important for the regulation of its subcellular distribution. It is tempting to speculate that host factors involved in membrane trafficking or lipogenesis possibly interact with NS5A in a phosphorylation-dependent manner and facilitate recruitment of NS5A to low-density membrane structures surrounding LDs. Although further study is needed to validate this speculation, NS5A-interacting factors, such as VAP-A and diacylglycerol acyltransferase-1 (19, 63, 64), might be candidates for involvement in the regulation of this process.

To identify potential phospho-acceptor sites for CKI- $\alpha$ , a phospho-proteome analysis was carried out with NS5A isolated from HCVcc-infected cell lines with and without CKI- $\alpha$  knockdown (Fig. 6). Three kinds of phosphopeptides were identified out of a total of 629 peptides after peptide selection, although fine mapping of the phosphorylation sites was not completely successful. Among them, only the relative frequency of phosphopeptide 1 (GSPPEASSSVSLSAPSLR) was decreased after CKI- $\alpha$  knockdown, suggesting the possibility that peptide 1 contains amino acids phosphorylated by CKI- $\alpha$ . Serine residues in peptide 1, which are well conserved among HCV genotypes, matched the consensus sequences for CKI- $\alpha$ -mediated phosphorylation (57). In contrast, a threonine residue in peptide 2 is not conserved and does not match the consensus sequences for phosphorylation by CKI- $\alpha$ . The frequency of phosphopeptide 3 was unchanged with and without CKI- $\alpha$  knockdown (0.1% versus 0.1%). Further mutagenesis analyses targeting the peptide 1 region suggested that S225 and S232 are possible CKI- $\alpha$  phosphorylation sites involved in NS5A hyperphosphorylation and infectious virus production, because alanine substitution for either of these serine residues reproduced the viral phenotype after CKI- $\alpha$  knockdown more accurately than the other mutants within the peptide 1 region (Fig. 7B to E). The S229A and S235A mutations severely impaired viral replication, suggesting that phosphorylation at S229 and S235 is essential for efficient viral replication (Fig. 7B). However, the HCV protein expression assay using vaccinia virus expressing the T7 RNA polymerase revealed that phosphorylation of these residues is not involved in NS5A hyperphosphorylation (Fig. 7E). In the case of genotype 1 isolates, NS5A mutations that reduce hyperphosphorylation enhanced viral RNA replication (17, 18). However, this was not the case for the S225A or S232A mutation in genotype 2a. In addition, S229A and S235A mutations in genotype 1b (Con1) markedly enhance viral replication (18), but the same mutations are lethal in genotype 2a. S232 has been shown to be a potential phosphorylation site for CKI- $\alpha$  by a peptide-based kinase assay *in vitro* (28). However, the present study is the first to demonstrate the significance of phosphorylation at S225 and S232 in infectious virus production. The sequence coverage of NS5A by our mass spectrometry analysis was less than 60% and was especially low in domain III of NS5A (Fig. 6B). We cannot exclude the possibility of the presence of additional CKI- $\alpha$  phosphorylation sites in NS5A.

Recently, two cellular kinases involved in the regulation of NS5A phosphorylation have been identified. PI4K-III $\alpha$  is essential for HCV replication (65–70) and catalyzes the synthesis of phosphatidylinositol 4-phosphate accumulating in HCV replicating cells through its enzymatic activation resulting from an interaction with NS5A (71, 72). PI4K-III $\alpha$  directly interacts with the

C-terminal end of NS5A domain I, and NS5A–PI4K-III $\alpha$  binding is essential for viral replication. Its depletion resulted in a relative increase of p58 abundance, while overexpression of enzymatically active PI4K-III $\alpha$  increased the relative abundance of p56 (73). Plk1 has been shown to play a role in viral replication through hyperphosphorylation of NS5A (25). Plk1 was coimmunoprecipitated with NS5A, and knockdown of Plk1 or treatment with a specific inhibitor decreased both NS5A hyperphosphorylation and HCV replication. Since the recognition sites for CKI- $\alpha$ , PI4K-III $\alpha$ , and Plk1 have been assumed to be spatially close to each other (25, 28, 73), it is interesting to analyze their interactive actions in regard to regulation of NS5A phosphorylation.

The exquisite balance between the two different phosphorylated forms of NS5A has been proposed to regulate the HCV life cycle; basally phosphorylated p56 abundance is hypothetically involved in viral RNA replication, and hyperphosphorylated p58 is required for virion assembly (9, 17, 18, 73, 74). However, this hypothesis has not yet been fully proven. Our results here provide strong evidence supporting the involvement of NS5A hyperphosphorylation in viral assembly and that of CKI- $\alpha$  in mediating this process. These results not only contribute to a better understanding of the regulatory details of the HCV life cycle but also illuminate targets for potential antiviral strategies.

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## Alternative endocytosis pathway for productive entry of hepatitis C virus

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Previous studies have shown that hepatitis C virus (HCV) enters human hepatic cells through interaction with a series of cellular receptors, followed by clathrin-mediated, pH-dependent endocytosis. Here, we investigated the mechanisms of HCV entry into multiple HCV-permissive human hepatocyte-derived cells using trans-complemented HCV particles (HCVtcp). Knockdown of CD81 and claudin-1, or treatment with bafilomycin A1, reduced infection in Huh-7 and Huh7.5.1 cells, suggesting that HCV entered both cell types via receptor-mediated, pH-dependent endocytosis. Interestingly, knockdown of the clathrin heavy chain or dynamin-2 (Dyn2), as well as expression of the dominant-negative form of Dyn2, reduced infection of Huh-7 cells with HCVtcp, whereas infectious entry of HCVtcp into Huh7.5.1 cells was not impaired. Infection of Huh7.5.1 cells with culture-derived HCV (HCVcc) via a clathrin-independent pathway was also observed. Knockdown of caveolin-1, ADP-ribosylation factor 6 (Arf6), flotillin, p21-activated kinase 1 (PAK1) and the PAK1 effector C-terminal binding protein 1 of E1A had no inhibitory effects on HCVtcp infection into Huh7.5.1 cells, thus suggesting that the infectious entry pathway of HCV into Huh7.5.1 cells was not caveolae-mediated, or Arf6- and flotillin-mediated endocytosis and macropinocytosis, but rather may have occurred via an undefined endocytic pathway. Further analysis revealed that HCV entry was clathrin- and dynamin-dependent in ORL8c and HepCD81/miR122 cells, but productive entry of HCV was clathrin- and dynamin-independent in Hep3B/miR122 cells. Collectively, these data indicated that HCV entered different target cells through different entry routes.

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### INTRODUCTION

Over 170 million people worldwide are chronically infected with hepatitis C virus (HCV), and are at risk of developing chronic hepatitis, cirrhosis and hepatocellular carcinoma (Hoofnagle, 2002). HCV is an enveloped virus belonging to the family *Flaviviridae*. Its genome is an uncapped 9.6 kb positive-stranded RNA consisting of the 5'-UTR, an ORF encoding viral proteins and the 3'-UTR (Suzuki *et al.*, 2007). A precursor polyprotein is further processed into structural proteins (core, E1, and E2), followed by p7 and non-structural (NS) proteins (NS2, NS3, NS4A, NS4B, NS5A and NS5B), by cellular and viral proteases.

Host–virus interactions are required during the initial steps of viral infection. Viruses enter the cells by various pathways, such as receptor-mediated endocytosis followed by pH-dependent or -independent fusion from endocytic compartments, or pH-independent fusion at the plasma membrane coupled with receptor-mediated signalling and coordinated disassembly of the actin cortex (Grove & Marsh, 2011). It was reported previously that CD81 (Bartosch *et al.*, 2003; McKeating *et al.*, 2004; Pileri *et al.*, 1998), scavenger receptor class B type I (SR-BI) (Bartosch *et al.*, 2003; Scarselli *et al.*, 2002), claudin-1 (Evans *et al.*, 2007; Liu *et al.*, 2009) and occludin (Benedicto *et al.*, 2009; Liu *et al.*, 2009; Ploss *et al.*, 2009) are critical molecules for HCV entry into cells. Recently, epidermal growth factor receptor and ephrin receptor type A2 were also identified as host cofactors for HCV entry, possibly by modulating interactions between CD81 and claudin-1 (Lupberger *et al.*,

Two supplementary figures are available with the online version of this paper.

2011). In addition, Niemann–Pick C1-like 1 (NPC1L1) cholesterol absorption receptor has been shown to play a role in HCV entry, probably at the fusion step (Sainz *et al.*, 2012).

Following receptor binding, HCV has been reported to enter cultured cells via clathrin-mediated endocytosis, the most common and best-characterized mode of endocytosis, following membrane fusion in early endosomes (Blanchard *et al.*, 2006; Codran *et al.*, 2006; Collier *et al.*, 2009; Meertens *et al.*, 2006; Trotard *et al.*, 2009) using retrovirus-based HCV pseudoparticles (HCVpp) and cell culture-produced HCV (HCVcc). Early steps in HCV infection, including the role of HCV glycoprotein heterodimers, receptor binding, internalization and pH-dependent endosomal fusion, have been at least in part mimicked by HCVpp. However, as HCVpp are generated in non-hepatic cells such as human embryo kidney 293T cells, it is likely that the cell-derived component(s) of HCVpp differ from those of HCVcc.

In the present study, we readdressed the HCV endocytosis pathway using trans-complemented HCV particles (HCVtcp) (Suzuki *et al.*, 2012), of which the packaged genome is a subgenomic replicon. HCVtcp, generated in Huh-7 or its derivative cell lines with two plasmids, are infectious, but support only single-round infection, thereby allowing us to examine infectious viral entry without the influence of reinfection. In addition, HCVtcp is useful for quantifying productive infection by measuring luciferase activity. Furthermore, it has been shown that the HCVtcp system is more relevant as a model of HCV infection than HCVpp (Suzuki *et al.*, 2012). Our results demonstrated conclusively that, in addition to the clathrin-mediated endocytosis pathway, HCV was capable of utilizing the clathrin- and dynamin-independent pathways for infectious entry of HCV into human liver-derived cells.

## RESULTS

### HCV entry depends on receptor-mediated, pH-dependent endocytosis

HCV has been shown to enter permissive cells through clathrin-mediated endocytosis and low pH-dependent fusion with endosomes mostly using HCVpp (Codran *et al.*, 2006; Meertens *et al.*, 2006; Trotard *et al.*, 2009), although some researchers have used HCVcc with limited cell lines (Blanchard *et al.*, 2006; Collier *et al.*, 2009). However, several distinct characteristics between HCVpp and HCVcc have recently been revealed with regard to morphogenesis and entry steps (Helle *et al.*, 2010; Sainz *et al.*, 2012; Suzuki *et al.*, 2012; Vieyres *et al.*, 2010). Therefore, in this study, we used HCVtcp, which exhibit similar characteristics to HCVcc when compared with HCVpp and support single-round infection (Suzuki *et al.*, 2012).

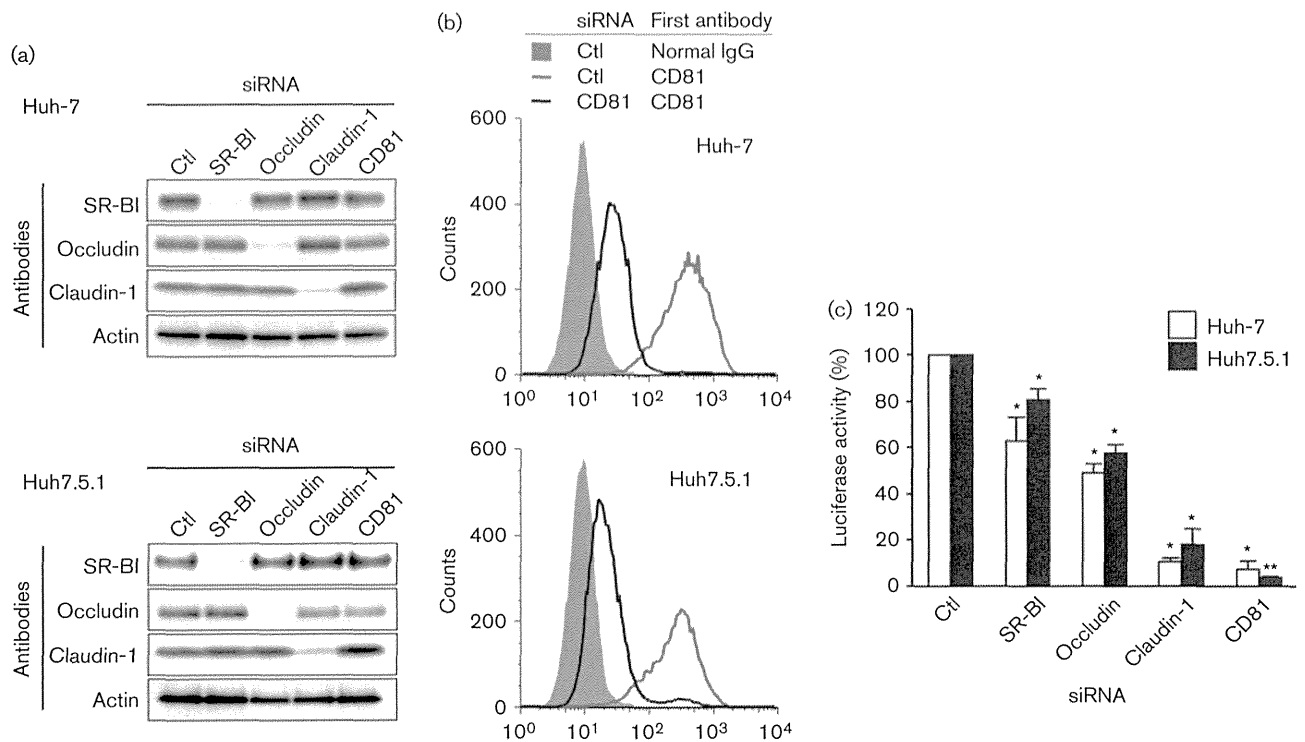
Initially, to determine whether receptor candidates such as CD81, claudin-1, occludin and SR-BI are essential for HCV

entry into Huh-7 and Huh7.5.1 cells, we examined the knockdown effect of these molecules on HCVtcp infection. Knockdown of these receptors was confirmed by immunoblotting (Fig. 1a) and FACS analysis (Fig. 1b). It should be noted that the luciferase activity in Huh7.5.1 was approximately four times higher than that in Huh-7 cells when the same amount of inoculum was used for infection (Fig. S1, available in the online Supplementary Material), and knockdown did not affect cell viability (data not shown). Knockdown of CD81 and claudin-1 significantly reduced the infection of Huh-7 and Huh7.5.1 cells with HCVtcp derived from genotype 2a (Fig. 1c). Knockdown of occludin led to a moderate reduction in infection; however, only a marginal effect was observed in SR-BI knockdown in both Huh-7 and Huh7.5.1 cells (Fig. 1c), possibly due to the reduced requirement for SR-BI during virus entry by adaptive mutation in E2 (Grove *et al.*, 2008).

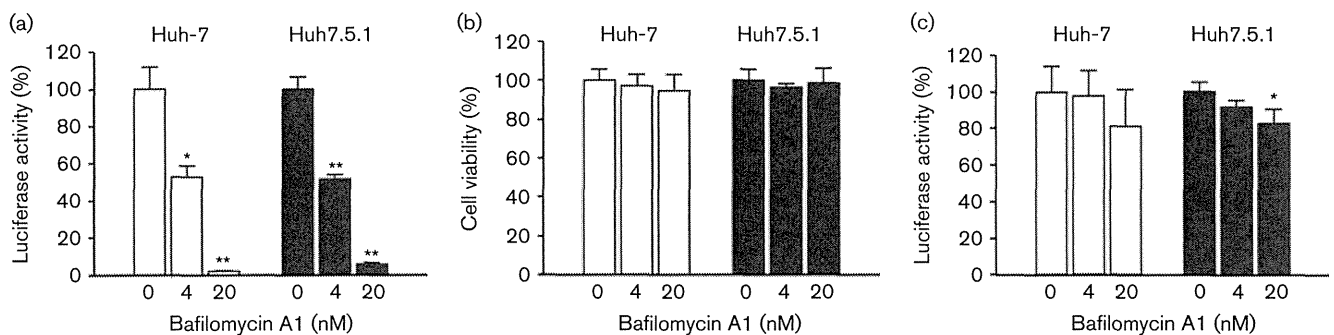
Next, to examine whether HCV entry was pH-dependent, Huh-7 and Huh7.5.1 cells were pretreated with bafilomycin A1, an inhibitor of vacuolar H<sup>+</sup>-ATPases that impairs vesicle acidification, and then infected with HCVtcp. At 72 h post-infection, luciferase activity and cell viability were determined. Bafilomycin A1 inhibited HCVtcp infection in a dose-dependent manner without affecting cell viability in both Huh-7 and Huh7.5.1 cells (Fig. 2a, b). We also confirmed that treatment with bafilomycin A1 after HCVtcp infection had a minor effect on luciferase activity (Fig. 2c). These results indicated that the infectious route of HCVtcp into Huh-7 and Huh7.5.1 cells is receptor-mediated and involves pH-dependent endocytosis.

### Knockdown of clathrin heavy chain (CHC) or dynamin-2 (Dyn2) reduces HCVtcp infection in Huh-7 cells, but not in Huh7.5.1 cells

Among the known pathways of pH-dependent viral endocytosis, clathrin-mediated dynamin-dependent endocytosis is a major endocytosis pathway. Chlorpromazine, an inhibitor of clathrin-dependent endocytosis, has been commonly used to study clathrin-mediated endocytosis; however, it exerts multiple side-effects on cell function as it targets numerous receptors and intracellular enzymes, and alters plasma membrane characteristics (Sieczkarski & Whittaker, 2002a). Therefore, we examined the HCV endocytosis pathway by knockdown of specific molecules required for the endocytosis pathway. CHC, a major structural protein in clathrin-coated vesicles, and Dyn2, a GTPase essential for clathrin-coated-pit scission from the plasma membrane, play important roles in the clathrin-mediated pathway. Another well-studied model of viral entry is caveolin-mediated endocytosis. The role of dynamin in both clathrin-mediated endocytosis and caveolae-dependent endocytosis has been established (Marsh & Helenius, 2006; Miaczynska & Stenmark, 2008). To examine the endocytosis pathways of HCV, small interfering RNAs (siRNAs) for CHC, Dyn2 and caveolin-1 (Cav1), or scrambled control siRNA, were transfected into Huh-7 or



**Fig. 1.** Knockdown effect of receptor candidate molecules on HCV infection. (a) Huh-7 or Huh7.5.1 cells were transfected with the indicated small interfering RNAs (siRNA), harvested at 48 h post-transfection and the specific knockdown of each protein was verified by immunoblotting. (b) Huh-7 or Huh7.5.1 cells were transfected with CD81 or control siRNAs, harvested at 48 h post-transfection and the cell surface expression of CD81 was verified by FACS analysis. (c) Cells transfected with siRNA were infected with the same amount of HCVtcp at 48 h post-transfection. Firefly luciferase activity in the cells was determined at 72 h post-infection and is expressed relative to the activity with control siRNA transfection. The value for control (Ctl) siRNA was set at 100%. Data represent the mean  $\pm$  sd. Statistical differences between controls and each siRNA were evaluated using Student's *t*-test. \**P* < 0.05, \*\**P* < 0.001 versus control.



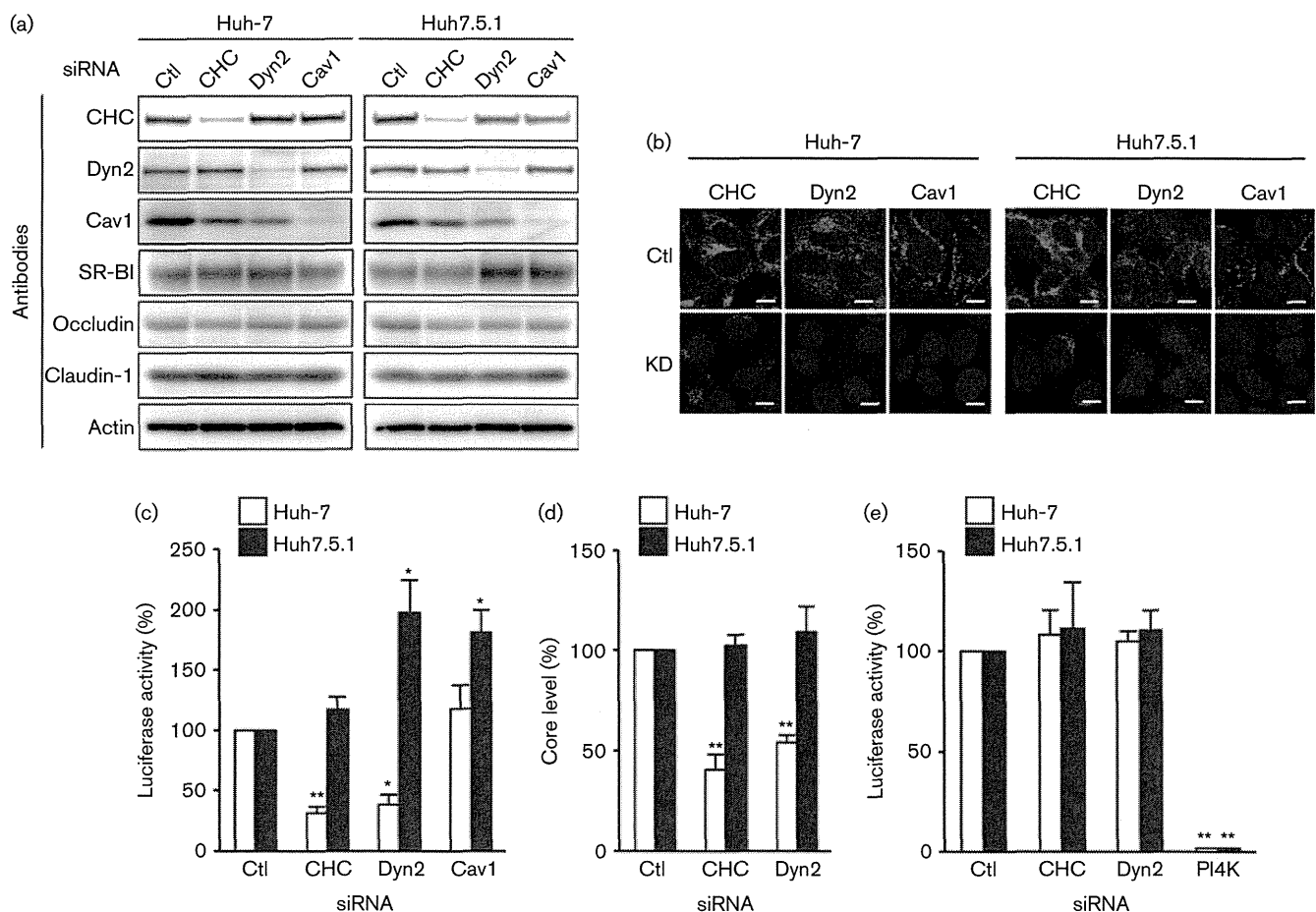
**Fig. 2.** Role of endosomal low pH in HCV infection. Cells were treated with bafilomycin A1 for 1 h at the indicated concentrations and infected with HCVtcp. (a, b) Luciferase activity (a) and cell viability (b) were determined at 72 h post-infection, and expressed relative to amounts observed in controls. (c) Cells were treated with bafilomycin A1 for 1 h at the indicated concentrations 48 h after HCVtcp infection. Luciferase activity was determined at 10 h post-treatment and expressed relative to amounts observed in controls. Data represent the mean  $\pm$  sd. Statistical differences between controls and indicated concentrations were evaluated using Student's *t*-test. \**P* < 0.05, \*\**P* < 0.001 versus control.

Huh7.5.1 cells, followed by infection with HCVtcp. Expression of CHC, Dyn2 and Cav1 was downregulated by transfection of specific siRNAs (Fig. 3a, b), whereas expression of SR-BI, occludin, claudin-1 and CD81 was not reduced (Figs 3a and S2). As indicated in Fig. 3(c), luciferase activity from HCVtcp was significantly reduced by knockdown of CHC and Dyn2 in Huh-7 cells, but not in Huh7.5.1 cells. Knockdown of Cav1 showed no inhibitory effects on HCVtcp entry into either cell line. Dynamin-independent entry in Huh7.5.1 cells was also observed using HCVtcp derived from genotype 1b (data not shown). Knockdown of CHC or Dyn2 also reduced entry of HCVcc in Huh-7 cells, but had no inhibitory effects in Huh7.5.1 (Fig. 3d). To rule out the possibility of effects on CHC and Dyn2 knockdown on viral RNA replication, HCVtcp were also

inoculated before siRNA transfection. Luciferase activity was not affected by knockdown of CHC or Dyn2 in either cell line, whereas marked inhibition was observed for phosphatidylinositol 4-kinase (PI4K) (Fig. 3e). These data suggested that HCV entry was clathrin-mediated and dynamin-dependent in Huh-7 cells, but productive entry of HCV was clathrin- and dynamin-independent in Huh7.5.1 cells.

### Expression of the dominant-negative form of Dyn2 reduces HCV infection in Huh-7 cells, but not in Huh7.5.1 cells

We also examined the role of dynamin in infectious entry of HCV into Huh-7 and Huh7.5.1 cells by overexpression of the dominant-negative form of Dyn2 (Dyn-K44A), which



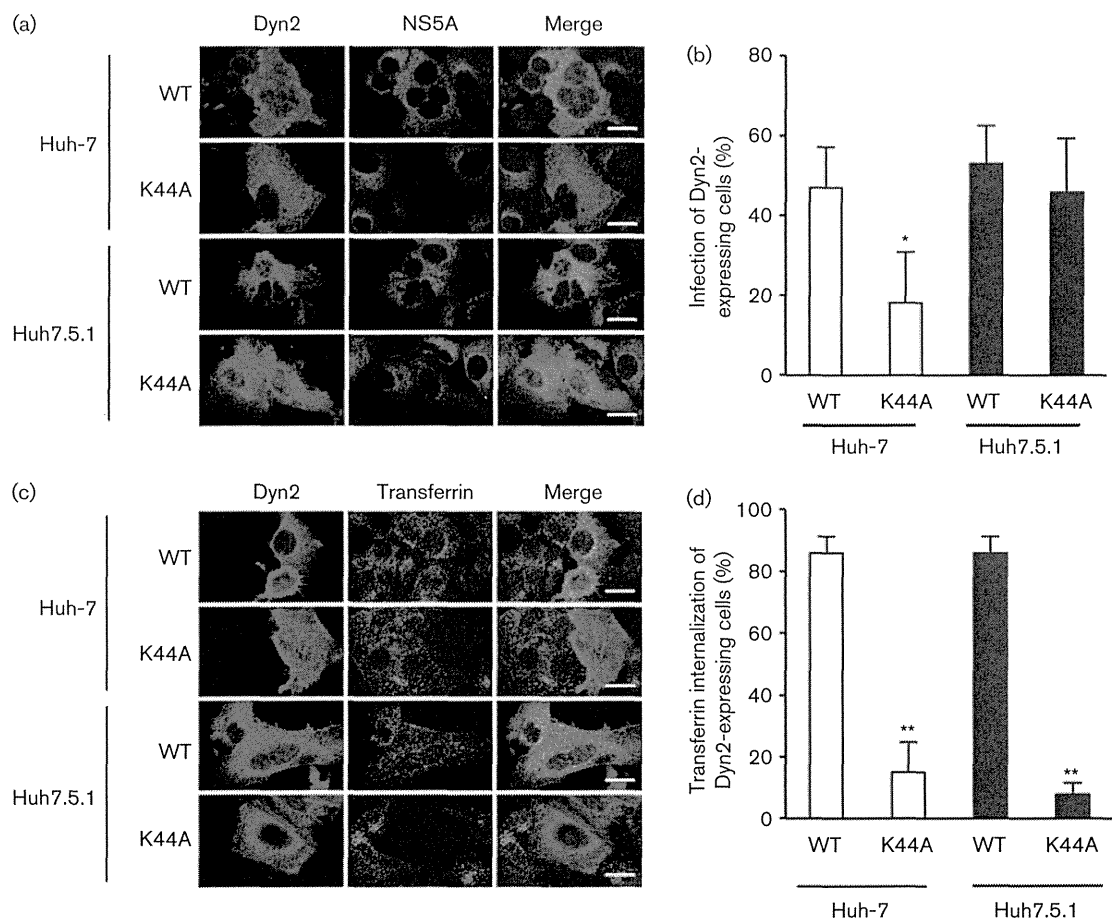
**Fig. 3.** Effects of CHC, Dyn2 and Cav1 knockdown on HCV infection. (a, b) Huh-7 cells or Huh7.5.1 cells were transfected with the indicated siRNAs and the specific knockdown (KD) of each protein was verified by immunoblotting (a) or immunostaining (b) at 48 h post-transfection. Bar, 50 μm. (c) Cells were transfected with the indicated siRNAs, followed by infection with HCVtcp at 48 h post-transfection. Firefly luciferase activity in the cells was subsequently determined at 3 days post-infection. The value for control (Ctl) siRNA was set at 100%. Data represent the mean ± SD. (d) Cells were transfected with siRNA, followed by infection with HCVcc at 48 h post-transfection. Intracellular core levels were quantified at 24 h post-infection. The value for control siRNA was set at 100%. Data represent the mean ± SD. (e) Cells were infected with HCVtcp, followed by transfection with the indicated siRNAs. Luciferase activity in the cells was subsequently determined at 2 days post-transfection. The value for control siRNA was set at 100%. Data represent the mean ± SD. Statistical differences between controls and each siRNA were evaluated using Student's *t*-test. \**P* < 0.05, \*\**P* < 0.001 versus control.

has been shown to effectively block clathrin-dependent and caveolar endocytosis (Damke *et al.*, 1995). Expression of haemagglutinin (HA)-tagged Dyn-K44A reduced the number of HCV-infected Huh-7 cells, but not Huh7.5.1 cells, as compared with WT HA-tagged Dyn2 (Dyn-WT), as shown in Fig. 4(a, b). Interestingly, internalization of transferrin, which is known to be mediated by clathrin-dependent endocytosis, was reduced in both Huh-7 and Huh7.5.1 cells expressing Dyn-K44A, whereas cells expressing Dyn-WT showed efficient endocytosis of transferrin (Fig. 4c, d). Collectively, these results suggested that dynamin participated in the internalization of HCV in Huh-7 cells, but was

not absolutely required in Huh7.5.1 cells, although transferrin was taken up via dynamin-dependent endocytosis in both Huh-7 and Huh7.5.1 cells.

#### Flotillin-1 or the GTPase regulator associated with focal adhesion kinase 1 (GRAF1) play no major role during HCV infection of Huh7.5.1 cells

In order to dissect the major endocytosis pathways of HCVtcp in Huh7.5.1 cells, we investigated the role of alternative routes of HCV entry by siRNA knockdown. We silenced essential factors for the clathrin- or dynamin-independent pathways



**Fig. 4.** Dynamin participates in the internalization of HCV in Huh-7 cells, but not in Huh7.5.1 cells. (a) Cells were transfected with HA-tagged WT Dyn2 (Dyn-WT) or dominant-negative Dyn2 (Dyn-K44A) expression plasmids. At 2 days post-transfection, cells were infected with HCVtcp, which possessed a subgenomic replicon without the luciferase gene. After 3 days, cells were fixed and HA-Dyn2 or HCV NS5A stained with anti-HA or anti-NS5A antibodies, respectively. Cell nuclei were counterstained with DAPI. Bar, 100 μm. (b) Data were quantified as the population of HCVtcp-infected cells among HA-positive cells. At least 20 HA-positive cells were evaluated in triplicate experiments. Data represent the mean ± SD. (c) Cells were transfected with HA-tagged Dyn-WT or Dyn-K44A expression plasmids. At 2 days post-transfection, cells were incubated with Alexa Fluor-488 labelled transferrin at 37 °C in a 5% CO<sub>2</sub> incubator. After 30 min of incubation, cells were washed, fixed and stained with anti-HA antibodies. Cell nuclei were counterstained with DAPI. Bar, 100 μm. (d) Data were quantified as the population of transferrin-internalized cells among HA-positive cells. At least 20 HA-positive cells were evaluated in triplicate experiments. Data represent the mean ± SD. Statistical differences between Dyn-WT and Dyn-K44A were evaluated using Student's *t*-test. \**P*<0.05, \*\**P*<0.001 versus Dyn-WT.

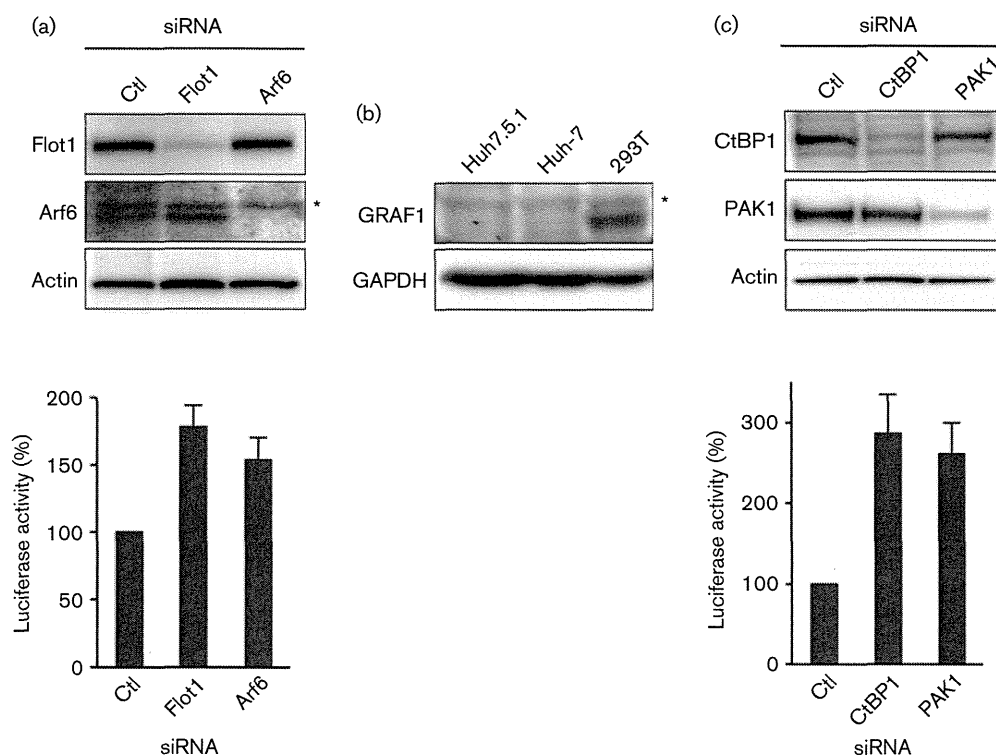


including flotillin-dependent endocytosis, ADP-ribosylation factor 6 (Arf6)-dependent endocytosis, clathrin-independent carrier/glycosylphosphatidylinositol-enriched early endosomal compartment (CLIC/GEEC) endocytic pathway and macropinocytosis in Huh7.5.1 cells. Flotillin-1 and Arf6 are indispensable components of the flotillin and Arf6 pathways, respectively. Knockdown of flotillin-1 or Arf6 had no inhibitory effects on HCVtcp infection in Huh7.5.1 cells (Fig. 5a). The CLIC/GEEC endocytic pathway has recently become better defined and is regulated by the GTPase regulator associated with focal adhesion kinase-1 (GRAF1). However, GRAF1 was not detected in Huh-7 or Huh7.5.1 cells (Fig. 5b); thus, it is unlikely that the CLIC/GEEC pathway was involved in HCV entry in Huh7.5.1 cells. In addition, knockdown of p21-activated kinase 1 (PAK1) and the PAK1 effector C-terminal binding protein 1 of E1A (CtBP1), which play important regulatory roles in the process of macropinocytosis, did not inhibit HCVtcp infection in Huh7.5.1 cells (Fig. 5c). Taken together, these results suggested that the entry of HCVtcp into Huh7.5.1 cells was not mediated mainly by flotillin-dependent endocytosis,

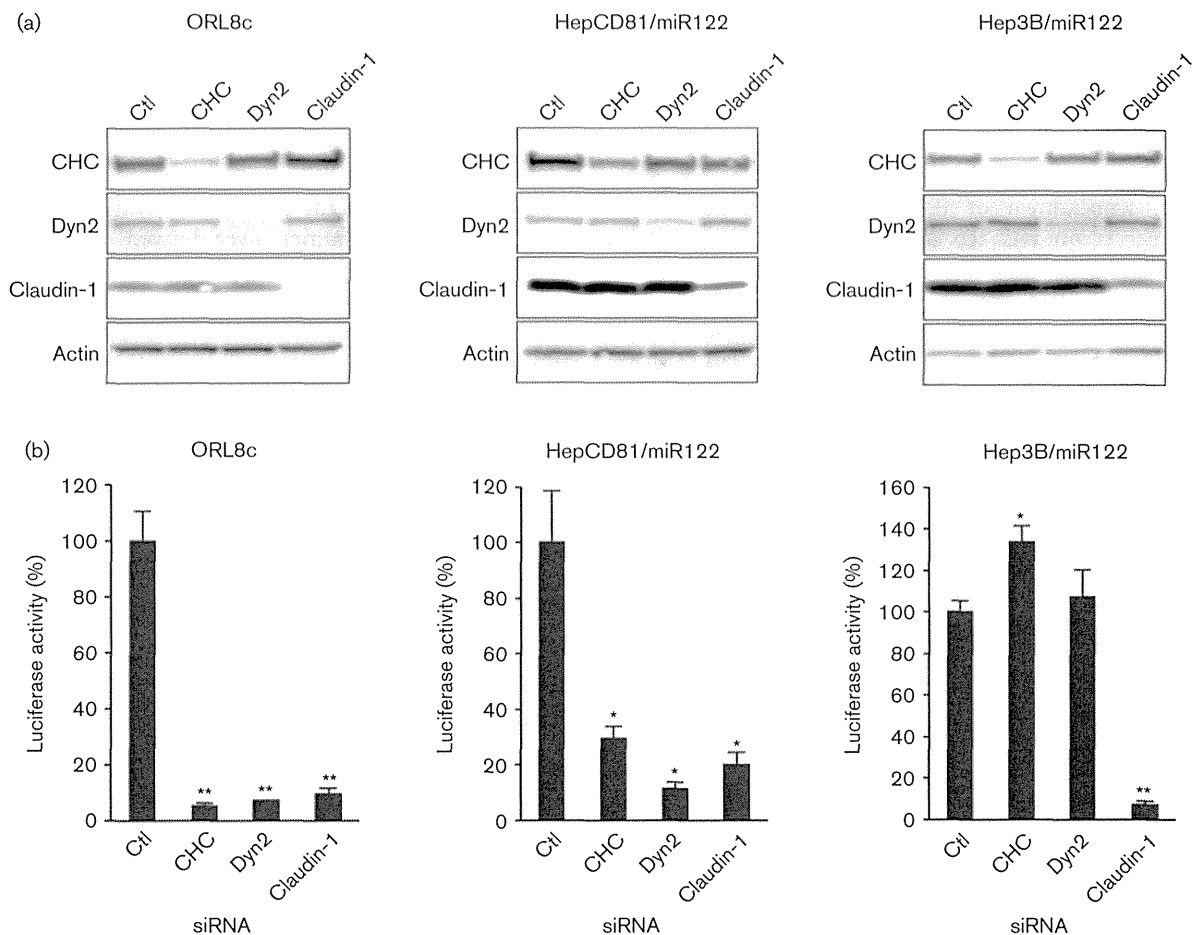
Arf6-dependent endocytosis, the CLIC/GEEC endocytic pathway and macropinocytosis.

### Clathrin-dependent and -independent pathways for HCV entry in other hepatic cells

We further examined the endocytosis pathways for HCV in non-Huh-7-related human liver-derived cell lines. Three HCVcc permissive hepatocellular carcinoma cell lines, Li23-derived ORL8c (Kato *et al.*, 2009), HepCD81/miR122 cells (HepG2/CD81 cells overexpressing miR122) and Hep3B/miR122 (Kambara *et al.*, 2012), were transfected with siRNA for CHC, Dyn2 or claudin-1, followed by infection with HCVtcp. Immunoblotting was performed in order to confirm knockdown of target proteins (Fig. 6a). Although knockdown of CHC or Dyn2 expression inhibited HCVtcp infection of ORL8c and HepCD81/miR122 cells, HCVtcp infection of Hep3B/miR122 cells was not affected (Fig. 6b), thus suggesting that productive entry of HCV is clathrin- and dynamin-independent in Hep3B/miR122 cells.



**Fig. 5.** Role of an alternative endocytosis pathway of HCV in Huh7.5.1 cells. (a) Huh7.5.1 cells were transfected with flotillin-1 (Flot1) or Arf6 siRNAs and specific knockdown of each protein was verified by immunoblotting (upper). Non-specific bands are marked with an asterisk. Cells transfected with siRNA were infected with HCVtcp. Luciferase activity (lower) was determined at 72 h post-infection and expressed relative to the amount observed in control (Ctl) siRNA transfection. Data represent the mean  $\pm$  SD. (b) Expression of GRAF1 and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) in Huh7.5.1, Huh-7 and 293T cells was analysed by immunoblotting. Non-specific bands are marked with an asterisk. (c) Huh7.5.1 cells were transfected with CtBP1 or PAK1 siRNA and specific knockdown of each protein was verified by immunoblotting (upper). Cells transfected with siRNA were infected with the HCVtcp. Luciferase activity (lower) was determined at 72 h post-infection and expressed relative to the amount observed in control (Ctl) siRNA transfection. Data represent the mean  $\pm$  SD.



**Fig. 6.** Clathrin-dependent and -independent pathway of HCV entry in other HCV-permissive cells. The indicated cells were transfected with the indicated siRNAs and then infected with HCVtpp at 48 h post-transfection. (a) Specific knockdown of each protein was verified by immunoblotting. (b) Luciferase activity was determined at 72 h post-infection and expressed relative to the amount observed in the control (Ctl) siRNA transfection. Data represent the mean  $\pm$  SD. Statistical differences between controls and each siRNA were evaluated using Student's *t*-test. \* $P < 0.05$ , \*\* $P < 0.001$  versus control.

In summary, we identified an alternative clathrin- and dynamin-independent entry pathway for HCV in at least two independent cell lines, Huh7.5.1 and Hep3B/miR122 cells, in addition to the previously reported clathrin- and dynamin-dependent pathway. These findings provided clues for understanding the molecular mechanisms of the endocytosis pathway for HCV infection.

## DISCUSSION

Many viruses have been shown to utilize a number of different endocytic pathways to productively infect their hosts. Clathrin-dependent endocytosis would appear to be the most commonly used, but it is increasingly clear that a number of clathrin-independent endocytosis pathways are also used by several different viruses (Mercer *et al.*, 2010). In the case of HCV, it has been reported that viral entry is mediated by clathrin-dependent endocytosis (Blanchard

*et al.*, 2006; Codran *et al.*, 2006; Coller *et al.*, 2009; Meertens *et al.*, 2006; Trotard *et al.*, 2009). In these papers, HCVpp was used at least in part for analysis of HCV entry pathway. However, recent reports have revealed several different characteristics between HCVpp and HCVcc.

Viral entry has been addressed primarily by pharmacologic inhibitor studies, immunofluorescence and electron microscopy, by transfection with dominant-negative constructs, and more recently by siRNA knockdown. Analysis of endocytosis pathways using pharmacological inhibitors has raised concerns about specificity. For example, chlorpromazine, an inhibitor of clathrin-mediated endocytosis, has been shown to exert multiple side-effects on cell function as it targets numerous receptors and intracellular enzymes, and alters plasma membrane characteristics (Sieczkarski & Whittaker, 2002a). Methods for elucidating the viral endocytosis pathway by co-localization of virus particles with host factor also have limitations. Electron and

fluorescence microscopy, which require a high particle number, do not allow the differentiation of infectious and non-infectious particles. Infectious particles of HCV in the supernatant of infected cells appeared to represent only a small portion of secreted virus particles (Akazawa *et al.*, 2008) and it is unclear whether the viral particles observed by microscopy could lead to productive infection. Therefore, we utilized HCVtcp, which is useful for determining productive entry of the virus without reinfection, and a combination of siRNA knockdown and dominant-negative mutants for analysis of the productive route of infection. Although HCVcc is also utilized in analysis of productive entry, it cannot completely exclude the effects of reinfection by virus produced by infected cells. Reduction of HCVcc infection by knockdown of CHC and Dyn2 was moderate when compared with that of HCVtcp (Fig. 3c, d), thus suggesting slight effects due to reinfection in HCVcc.

The data presented here demonstrate for the first time to our knowledge that HCV is able to enter cells via dynamin-independent endocytosis in addition to the previously described classical clathrin- and dynamin-dependent pathway. First, knockdown of CHC and Dyn2 had no inhibitory effects on HCVtcp and HCVcc entry into Huh7.5.1 cells. Second, overexpression of dominant-negative Dyn2 had no inhibitory effects on HCVtcp in Huh7.5.1 cells. Finally, in addition to Huh7.5.1 cells, Hep3B/miR122 cells were also shown to be infected with HCV via clathrin- and dynamin-independent pathways. We further investigated the role of alternative minor routes of HCV entry into Huh7.5.1 cells; however, the productive endocytosis pathway could not be defined. It should be noted that inhibition of alternative endocytosis routes by siRNA led to an increase of luciferase activity (Figs 3c and 5a, c). This could be explained by the inhibition of a particular endocytosis pathway resulting in a compensatory increase in alternative endocytosis pathways (Damke *et al.*, 1995).

Although we confirmed an alternative endocytosis pathway for the productive entry of HCV, it is not clear why and how the two independent endocytosis pathways operate in different cell lines. SV40 can enter cells via caveolae-dependent (Norkin *et al.*, 2002; Pelkmans *et al.*, 2001) and -independent (Damm *et al.*, 2005) pathways. Influenza virus enters cells via clathrin-mediated endocytosis (Matlin *et al.*, 1981) in addition to non-clathrin-mediated, non-caveola-mediated internalization pathways (Sieczkarski & Whittaker, 2002b). Entry of dengue virus type 2 is clathrin-dependent in HeLa and C6/36 cells (Acosta *et al.*, 2008; Mosso *et al.*, 2008; van der Schaar *et al.*, 2008), and is clathrin-independent in Vero cells (Acosta *et al.*, 2009). Different receptor usage may determine the consequential route of entry. However, we did not observe any differences between Huh-7 and Huh7.5.1 cells in terms of knockdown effects of receptor candidate molecules on HCV infection, as shown in Fig. 1(c), although we cannot exclude the possibility that other undefined receptors are associated with viral entry. Huh7.5.1 cells were established by

elimination of the HCV genome from replicon cells derived from Huh-7 cells (Blight *et al.*, 2002; Zhong *et al.*, 2005) and they exhibit more potent replication of HCV than the original Huh-7 cells. Further study showed that the increased permissiveness of cured cells results from a mutation in the retinoic acid-inducible gene I (Sumpter *et al.*, 2005), which impairs IFN signalling. In addition, it has been shown that cured cell lines express higher levels of miR122 than parental cells participating in the efficient propagation of HCVcc (Kambara *et al.*, 2012). As it is unclear whether these changes are the reason for a distinct endocytosis pathway, it will be of interest to explore these associations in further studies.

In conclusion, we confirmed an alternative clathrin-independent endocytosis pathway in HCV-permissive human hepatic-derived cells, in addition to the previously reported clathrin-dependent endocytosis pathway. This paper highlights the fact that clathrin- and dynamin-mediated endocytosis is the main route of HCV entry for Huh-7, HepCD81/miR122 and ORL8c cells, whilst clathrin and dynamin do not play a major role during the productive route of HCV infection in Huh7.5.1 and Hep3B/miR122 cells. Taken together, these studies suggest that different cell entry pathways for HCV infection may be utilized in different cell types, although further studies are necessary in order to understand this phenomenon.

## METHODS

**Cells.** The human hepatocellular carcinoma cell lines Huh-7, Huh7.5.1, Hep3B/miR122 and HepG2/CD81, which overexpressed miR122 (Kambara *et al.*, 2012), were maintained in Dulbecco's modified Eagle's medium (DMEM; Wako Pure Chemical Industries) containing non-essential amino acids, penicillin (100 U ml<sup>-1</sup>), streptomycin (100 µg ml<sup>-1</sup>) and 10% FBS. Li23-derived ORL8c cells (Kato *et al.*, 2009) were maintained in F12 medium and DMEM (1:1, v/v) supplemented with 1% FBS, epidermal growth factor (50 ng ml<sup>-1</sup>), insulin (10 µg ml<sup>-1</sup>), hydrocortisone (0.36 µg ml<sup>-1</sup>), transferrin (5 µg ml<sup>-1</sup>), linoleic acid (5 µg ml<sup>-1</sup>), selenium (20 ng ml<sup>-1</sup>), prolactin (10 ng ml<sup>-1</sup>), gentamicin (10 µg ml<sup>-1</sup>), kanamycin monosulfate (0.2 mg ml<sup>-1</sup>) and fungizone (0.5 µg ml<sup>-1</sup>). All cell lines were cultured at 37 °C in a 5% CO<sub>2</sub> incubator.

**Preparation of viruses.** HCVtcp and HCVcc derived from JFH-1 with adaptive mutations in E2 (N417S), p7 (N765D) and NS2 (Q1012R) were generated as described previously (Suzuki *et al.*, 2012). For HepCD81/miR122 and ORL8c cells, HCVtcp containing the *Gaussia* luciferase (GLuc) reporter gene were used. To do this, plasmid pHH/SGR-JFH1/GLuc/NS3m carrying the bicistronic sub-genomic HCV replicon containing the GLuc reporter gene and the NS3 adaptive mutation was constructed by replacement of the firefly luciferase (FLuc) gene of pHH/SGR-Luc containing the NS3 mutation (N1586D) (Suzuki *et al.*, 2012) with the GLuc gene of pCMV-GLuc (NEB).

**Plasmids.** HA-tagged Dyn2, a dominant-negative Dyn2 (K44A) in which Lys44 was replaced with Ala, was cloned into pcDNA3.1 as described previously (Kataoka *et al.*, 2012).

**Gene silencing by siRNA.** siRNAs were purchased from Sigma-Aldrich and were introduced into the cells at a final concentration of

30 nM using Lipofectamine RNAiMAX (Invitrogen) in accordance with the manufacturer's instructions. Target sequences of the siRNAs were: occludin (5'-GCAAGAUCAUAUGAGACA-3'), SR-BI (5'-GAGCUUUGCCUUGGUCUA-3'), CD81 (5'-CUGUGAUCAUGAUUCUGA-3'), CHC (5'-CUAGCUUUGCACAGUUUAA-3'), Dyn2 (5'-CCCUCAGGAGGCGCUCAA-3'), Cav1 (5'-CCCUAAACACCUCACGAU-3'), flotillin-1 (5'-CCUAUGACAUCGAGGUCAA-3'), Arf6 (5'-CAGUUCUUGGUAAGUCCU-3'), CtBP1 (5'-GACUCGACGCUGUGCCACA-3') and PAK1 (5'-GCAUCAAUUCCUGAAGAUU-3'). Target sequences of the siRNAs for claudin-1, PI4K and scrambled negative control were as described previously (Suzuki *et al.*, 2013).

**Immunoblotting.** Cells were washed with PBS and incubated with passive lysis buffer (Promega). Lysates were sonicated for 10 min and added to the same volume of 2 × SDS-PAGE sample buffer. Protein samples were boiled for 10 min, separated by SDS-PAGE and then transferred to PVDF membranes (Merck Millipore). After blocking, membranes were probed with primary antibodies, followed by incubation with peroxidase-conjugated secondary antibody. Antigen-antibody complexes were visualized using an enhanced chemiluminescence detection system (SuperSignal West Pico Chemiluminescent Substrate; Thermo Scientific) in accordance with the manufacturer's protocols.

**Flow cytometry.** Cultured cells detached by treatment with trypsin were incubated with anti-CD81 antibody or anti-mouse IgG antibody for 1 h at 4 °C. After being washed with PBS containing 0.1 % BSA, cells were incubated with an Alexa Fluor 488-conjugated anti-mouse secondary antibody (Invitrogen) for 1 h at 4 °C, washed repeatedly and resuspended in PBS. Analyses were performed using a FACSCalibur system (Becton Dickinson).

**Reagents and antibodies.** Bafilomycin A1 was obtained from Wako Pure Chemical Industries. Alexa Fluor 488-conjugated transferrin was obtained from Invitrogen. For immunoblotting, anti-SR-BI (NB400-104; Novus Biologicals), anti-occludin (71-1500; Invitrogen), anti-claudin-1 (51-9000; Invitrogen), anti-Dyn2 (ab3457; Abcam), anti-Cav1 (N-20; Santa Cruz Biotechnology), anti-flotillin (H-104; Santa Cruz Biotechnology), anti-Arf6 (ab77581; Abcam) and anti-PAK1 (2602; Cell Signaling Technology) rabbit polyclonal antibodies; anti-CD81 (JS-81; BD Biosciences), anti-β-actin (AC-15; Sigma-Aldrich), anti-CHC (23; BD Biosciences), anti-GRAF1 (SAB1400439; Sigma-Aldrich) and anti-glyceraldehyde 3-phosphate dehydrogenase (6C5; Merck Millipore) mouse mAb; and anti-CtBP1 goat polyclonal antibody (C-17; Santa Cruz Biotechnology) were used. For immunofluorescence staining, anti-CHC mAb (X22) and anti-HA rat polyclonal antibody (3F10) were obtained from Thermo Scientific and Roche Applied Science, respectively. Anti-NS5A antibody was a rabbit polyclonal antibody against synthetic peptides. Alexa Fluor 488- or 555-labelled secondary antibodies were obtained from Invitrogen.

**DNA transfection.** Cell monolayers were transfected with plasmid DNA using TransIT-LT1 transfection reagent (Mirus) in accordance with the manufacturer's instructions.

**Treatment of cells with bafilomycin A1 and cell viability.** Cells were preincubated with various concentrations of bafilomycin A1 for 60 min at 37 °C. Preincubated cells were then infected with HCVtcp. Cells treated with 0.1 % DMSO were used as controls. Cell viability was analysed by the Cell Titre-Glo Luminescent Cell Viability Assay (Promega).

**Uptake of transferrin.** Cells were grown on glass coverslips. After cells were transfected with HA-tagged Dyn2 expression plasmids, Alexa Fluor 488-conjugated transferrin at 20 µg ml<sup>-1</sup> was added and incubated for 30 min. Cells were washed with PBS and fixed in 4 % paraformaldehyde.

**Immunofluorescence analysis.** Huh7.5.1 and Huh-7 cells were fixed with 4 % paraformaldehyde in PBS for 30 min, and were then blocked and permeabilized with 0.3 % Triton X-100 in a non-fat milk solution (Block Ace; Snow Brand Milk Products) for 60 min at room temperature. Samples were then incubated with anti-CHC, anti-Dyn2, anti-Cav1, anti-NS5A or anti-HA for 60 min at room temperature, washed three times with PBS, and then incubated with secondary antibodies for 60 min at room temperature. Finally, samples were washed three times with PBS, rinsed briefly in double-distilled H<sub>2</sub>O and mounted with DAPI mounting medium. The signal was analysed using a Leica TCS SPE confocal microscope.

**Luciferase assay.** For quantification of FLuc activity in HCVtcp-infected cells, cells were lysed with passive lysis buffer (Promega) at 72 h post-infection. FLuc activity of the cells was determined using a luciferase assay system (Promega). For quantification of GLuc activity in supernatants of HCVtcp-infected cells, the *Renilla* Luciferase Assay System (Promega) was used. All luciferase assays were performed at least in triplicate.

**Quantification of HCV core protein.** HCV core protein was quantified using a highly sensitive enzyme immunoassay (Lumipulse G1200; Fujirebio) in accordance with the manufacturer's instructions.

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## Characterization of human bocavirus-like particles generated by recombinant baculoviruses



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Human bocavirus (HBoV) is a nonenveloped, single-stranded DNA virus, classified recently into the genus *Bocavirus* in the family *Parvoviridae*. A recombinant baculovirus expression system was used to express the major capsid protein VP2 of HBoV1, HBoV2, HBoV3 and HBoV4 in insect cells. A large amount of the 61-kDa VP2 capsid protein (p61) of HBoVs was generated and efficiently released into the supernatant. The capsid protein was self-assembled into 22-nm-dia. virus-like particles (VLPs) with a buoyant density of 1.30 g/cm<sup>3</sup>. The morphology of HBoVs-LPs was similar to that of the native HBoV particles, and immunogenic studies demonstrated the cross-reactivity among HBoV1, HBoV2, HBoV3 and HBoV4. When VP1 and VP2 protein of HBoV1 were co-expressed in insect cells, both proteins were detected in the same fraction after CsCl gradient centrifugation, suggesting that the VP1 protein is a minor structural protein of HBoVs. We developed an ELISA using purified VLPs as the antigen and used it to detect antibodies against HBoV1, HBoV2, HBoV3 and HBoV4. A high prevalence of antibodies against HBoVs was found in a general population of healthy Japanese, indicating that HBoVs have spread throughout Japan.

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### 1. Introduction

Human bocavirus type 1 (HBoV1), HBoV2, HBoV3 and HBoV4 are members of the genus *Bocavirus* in the subfamily *Parvovirinae* of the family *Parvoviridae* (Jartti et al., 2012). Following the first identification of HBoV1 in the nasopharyngeal secretion of a child with respiratory manifestations in 2005 (Allander et al., 2005), HBoV2, HBoV3 and HBoV4 were identified in fecal samples from children with non-polio acute flaccid paralysis and diarrhea in 2009 and 2010 (Arthur et al., 2009; Kapoor et al., 2009, 2010). HBoV is a small, nonenveloped icosahedral virus with a linear, single-stranded DNA genome of approx. 5.3 kilobases (kb). Sequence analyses of the HBoV genome revealed that it contains three open reading frames (ORFs) encoding four proteins in the following order: two nonstructural proteins (NS1 and NP1) and two viral capsid proteins (VP1

and VP2) (Allander et al., 2005). As in other members of the family *Parvoviridae*, the two capsid proteins, VP1 and VP2, have identical nucleotide and amino acid sequences except for an extra unique phospholipase-A motif (VP1u) at the amino-terminal end of the VP1 protein (Dijkman et al., 2009; Chen et al., 2010). The capsid protein exhibits *T*=1 symmetry with 60 copies of the coat protein (Gurda et al., 2010). The functions of NS1 and NP1 are unknown.

HBoV infection is associated mainly with pediatric respiratory diseases and gastrointestinal diseases (Jartti et al., 2012). HBoV DNAs have been detected in 5% to 15% of patients with respiratory illness or gastrointestinal symptoms in North America, South America, Europe, Asia, Australia and Africa, indicating that HBoV infection is a global health concern (Longtin et al., 2008; Arthur et al., 2009; Fabbiani et al., 2009; Moreno et al., 2009; Soderlund-Venermo et al., 2009; Tozer et al., 2009; Vallet et al., 2009; Chow et al., 2010; Huang et al., 2010; Kapoor et al., 2009, 2010; Santos et al., 2010). The lack of an efficient cell culture system for HBoV has hampered the preparation of antigens for the serology assay. It is generally accepted that noninfectious virus-like particles (VLPs)

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assembled from the viral structural proteins are similar to native virions in size, shape and antigenicity, and thus that VLPs are useful as antigens for immunoassays.

To understand the antigenic properties of HBoVs, we established an efficient production system for HBoV1–4 VLPs using a recombinant baculovirus expression system, and we characterized the antigenicity of the purified VLPs. We performed seroepidemiological studies on HBoVs with sera from healthy populations in Japan using an enzyme-linked immunosorbent assay (ELISA) that was developed based on VLPs as the antigen.

## 2. Materials and methods

### 2.1. Construction of recombinant baculoviruses and expression of HBoV capsid proteins

We synthesized the viral genes encoding VP1 of HBoV1, VP2 of HBoV1, VP2 of HBoV2, VP2 of HBoV3 and VP2 of HBoV4, each of which contains a *Bam*HI site before the start codon and an *Xba*I site after the stop codon, based on published sequences (GenBank accession nos.: AB481080, FJ170279, FJ948861, and NC\_012729), and cloned them into vector pUC57 (GeneScript, Piscataway, NJ) to generate plasmids pUC57-HBoV1-VP1, pUC57-HBoV1-VP2, pUC57-HBoV2-VP2, pUC57-HBoV3-VP2, and pUC57-HBoV4-VP2. These plasmids were digested with *Bam*HI and *Xba*I, and the purified VP1 or VP2 gene fragment was ligated to the transfer vector pVL1393 (Pharmingen, San Diego, CA), yielding plasmids pVL1393-HBoV1-VP1, pVL1393-HBoV1-VP2, pVL1393-HBoV2-VP2, pVL1393-HBoV3-VP2, and pVL1393-HBoV4-VP2.

An insect cell line, Sf9, derived from the armyworm *Spodoptera frugiperda* (Riken Cell Bank, Tsukuba, Japan), was co-transfected with a linearized wild-type *Autographa californica* nuclear polyhedrosis virus DNA (BaculoGold 21100D; Pharmingen) and the transfer plasmid by a lipofectin-mediated method as specified by the manufacturer (GIBCO-BRL, Gaithersburg, MD). The cells were incubated at 26.5 °C in TC-100 medium (GIBCO-BRL) supplemented with 8% fetal bovine serum (FBS) and 0.26% bacto tryptose phosphate broth (Difco Laboratories, Detroit, MI). Each recombinant virus was plaque-purified three times in Sf9 cells. The baculovirus recombinants thus obtained were designated as AchBoV1-VP1, AchBoV1-VP2, AchBoV2-VP2, AchBoV3-VP2 and AchBoV4-VP2, respectively. For large-scale expression, we used an insect cell line from the cabbage looper *Trichoplusia ni*, BTL-Tn 5B1-4 (Tn5) (Invitrogen, San Diego, CA). Tn5 cells were infected with the recombinant baculoviruses at a multiplicity of infection (m.o.i.) of 10 and cultured in EX-CELL 405 medium (JRH Biosciences, Lenexa, KS) at 26.5 °C as described (Li et al., 1997, 2011).

### 2.2. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and Western blot analysis

The proteins in the cell lysate and culture medium were separated by 5–20% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and stained with Coomassie blue. For Western blotting, the proteins in the SDS-PAGE gel were electrophoretically transferred onto a nitrocellulose membrane. The membrane was then blocked with 50 mM Tris-HCl (pH 7.4) containing 5% skim milk and 150 mM NaCl, and incubated with a human serum positive for anti-HBoV IgG. Detection of human IgG antibody was achieved using alkaline phosphatase conjugate goat anti-human IgG (1:1000 dilution; Dako, Copenhagen, Denmark). Nitroblue tetrazolium chloride and 5-bromo-4-chloro-3-indolyl phosphate P-toluidine were used as coloring agents (Bio-Rad Laboratories, Hercules, CA).

### 2.3. Purification of VLPs

The culture medium of the recombinant baculovirus-infected Tn5 cells was harvested on day 7 post-infection (p.i.). The intact cells, cell debris, and progeny baculoviruses were removed by centrifugation at 10,000 × g for 60 min. The supernatant was then spun at 126,000 × g for 3 h in a rotor (SW32Ti, Beckman Coulter, Indianapolis, IN), and the resulting pellet was resuspended in EX-CELL™ 405 medium at 4 °C overnight. For the CsCl gradient centrifugation, 4.5 mL of the samples were mixed with 2.1 g of CsCl, and then centrifuged at 116,000 × g for 24 h at 10 °C in the SW55Ti rotor. The gradient was fractionated into 250-μL aliquots, and each fraction was weighed in order to estimate the buoyant density and isopycnic point. Each fraction was diluted with EX-CELL™ 405 medium and centrifuged for 2 h at 237,000 × g in a Beckman TLA55 rotor to sediment the VLPs.

### 2.4. Electron microscopy (EM)

The purified VLPs were placed on a carbon-coated grid for 45 s, rinsed with distilled water, stained with a 2% uranyl acetate, and examined with an electron microscope (TEM-1400, JEOL, Tokyo) at 80 kV.

### 2.5. N-terminal amino acid sequence analysis

VLPs were purified by CsCl gradient centrifugation. N-terminal amino acid (aa) microsequencing was carried out using 100 pmol of the protein by Edman automated degradation on a protein sequencer (model 477, Applied Biosystems, Foster City, CA).

### 2.6. Healthy human serum samples

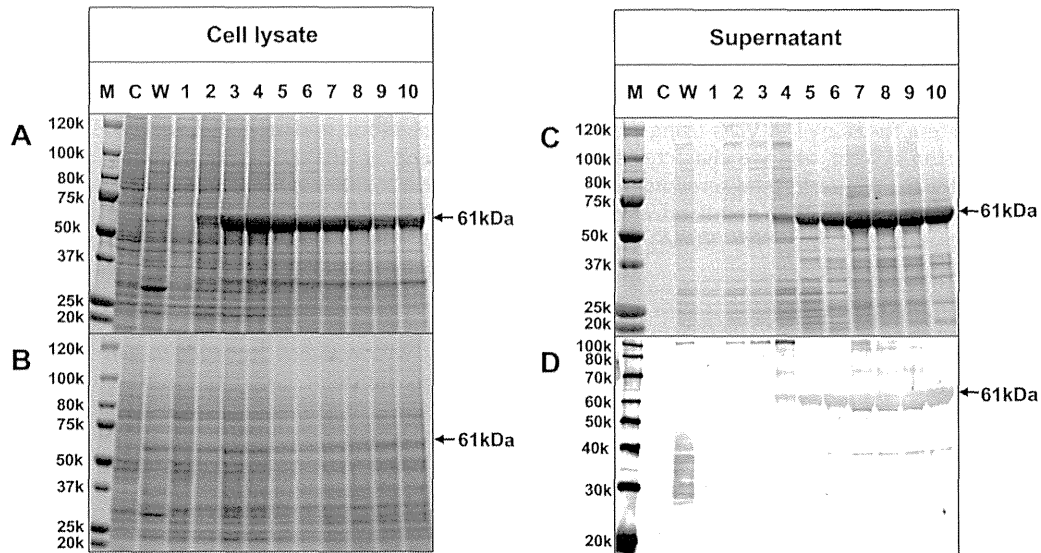
The specimens tested for HBoV serology consisted of 372 serum samples from healthy Japanese subjects collected in 1993. Serum samples were obtained from the Serum Bank of the National Institute of Infectious Diseases, Japan. The ages of the subjects ranged from 1 to 80 years old. Sixty-four percent (238/372) of the subjects were female, and 36% (134/372) were male.

### 2.7. Hyperimmune sera

Rabbits (Japanese White, 8 weeks old, female) were immunized with VLPs. The immunization was performed by a thigh muscle injection of purified VLPs at a dose of 500 μg per animal, and the booster injections were carried out at 4 and 6 weeks after the first injection with half doses of VLPs. All of the injections including the booster injections were carried out without any adjuvant. Immunized animals were bled 1 week after the last injection. The rabbit experiments were reviewed by the National Institute of Infectious Diseases (NIID) Ethics Committee and carried out according to the “Guides for Animal Experiments Performed at NIID” under code 111054.

### 2.8. Enzyme-linked immunosorbent assay (ELISA)

We developed an ELISA to detect anti-HBoV antibodies in human sera. Flat-bottom 96-well polystyrene microplates (Immunolon 2; Dynex Technologies, Chantilly, VA) were coated with the purified VLPs (1 μg/mL, 100 μL/well) and incubated overnight at 4 °C. Unbound VLPs were removed, and the plates were washed twice with 10-mM phosphate-buffered saline (PBS) containing 0.05% Tween 20 (PBS-T) and then blocked with 200 μL of 5% skim milk (Difco Laboratories) dissolved in PBS-T for 1 h at 37 °C. After three washes with PBS-T, diluted rabbit or human serum (100 μL/well) was added in duplicate. The plates were incubated



**Fig. 1.** Time course of HBoV1-VP2 expression. Insect cells, Tn5 ((A), (C) and (D)) and Sf9 (B), were infected with the recombinant baculovirus AchBoV1-VP2 and incubated at 26.5 °C. The cells were harvested on the indicated days (days 1 to 10), and 5  $\mu$ L of the sample from 25-times concentrated culture medium or the equivalent of the cell lysate from  $10^5$  cells was analyzed by SDS-PAGE. Protein bands were visualized by Coomassie blue staining or Western blot assay with an anti-HBoV1 IgG-positive human serum. (A) Coomassie blue staining of the Tn5 cell lysate. (B) Coomassie blue staining of the Sf9 cell lysate. (C) Coomassie blue staining of the Tn5 cell supernatant. (D) Western blot assay of the Tn5 cell supernatant. Arrows, p61; M, molecular weight marker; C, mock-infected cells; W, wild-type baculovirus-infected cells; lanes 1 to 10, AchBoV1-VP2-infected Sf9 or Tn5 cells were harvested on 1 to 10 days p.i.

at 37 °C for 1 h and washed three times as described above. The wells were incubated with 100  $\mu$ L of horseradish peroxidase-conjugated goat anti-human IgG (H+L) (Cappel, West Chester, PA) (1:10,000 dilution), IgM (Cappel) (1:2000 dilution) or horseradish peroxidase-conjugated goat anti-rabbit IgG (Cappel, Durham, NC) (1:2000 dilution) diluted with PBS-T containing 1% skim milk.

The plates were incubated at 37 °C for 1 h and washed four times with PBS-T. The substrate orthophenylenediamine (100  $\mu$ L) (Sigma Chemical, St. Louis, MO) and H<sub>2</sub>O<sub>2</sub> were added to each well. The plates were incubated in a dark room at room temperature for 30 min, then 50  $\mu$ L of 4N H<sub>2</sub>SO<sub>4</sub> was added into each well. Absorbance was measured at 492 nm. Test samples were considered positive when the absorbance was  $\geq 0.200$ .

### 2.9. Statistics method

Proportional analysis between groups and correlation analysis among anti-HBoV antibody levels were performed with the chi-squared test and Spearman test, respectively, using SPSS software.

## 3. Results

### 3.1. Efficient expression of HBoV VP2 proteins in Tn5 cells

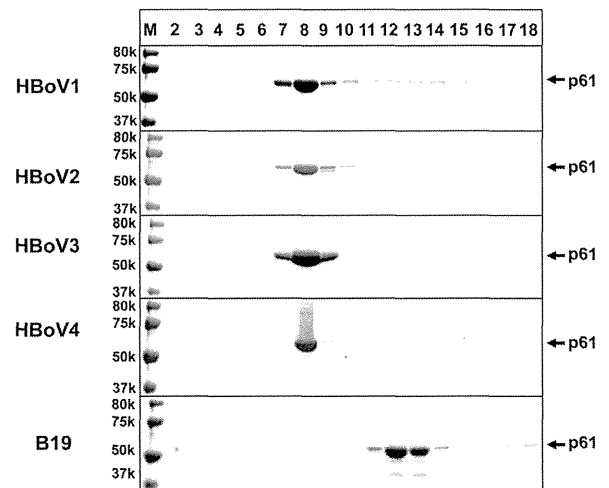
To examine which insect cell line is suitable for the expression of HBoV capsid proteins, we used the recombinant baculovirus AchBoV1-VP2 to infect Sf9 or Tn5 cells. The infected cells were harvested daily until 10 days p.i., and the protein expression was analyzed by SDS-PAGE followed by Coomassie blue staining. In the Tn5 cell lysates, one strong protein band with molecular mass of 61 kDa (p61) was first detected on day 2 and reached a peak on day 4 p.i. (Fig. 1A) In contrast, the expression level of VP2 in Sf9 cells was quite low, and we were unable to specify the VP2 protein by Coomassie blue staining (Fig. 1B); hence, we used Tn5 cells exclusively thereafter.

A large amount of p61 was also detected in the supernatant in infected Tn5 cells on day 4 and peaked on day 7 p.i. (Fig. 1C), suggesting that the HBoV-VP2 protein was efficiently released into

the supernatant. The p61 in the supernatant reacted with human serum that was positive for anti-HBoV IgG in Western blotting (Fig. 1D). The expression level and pattern of the VP2 capsid protein of HBoV2, HBoV3 and HBoV4 were similar to those of HBoV1-VP2 (data not shown).

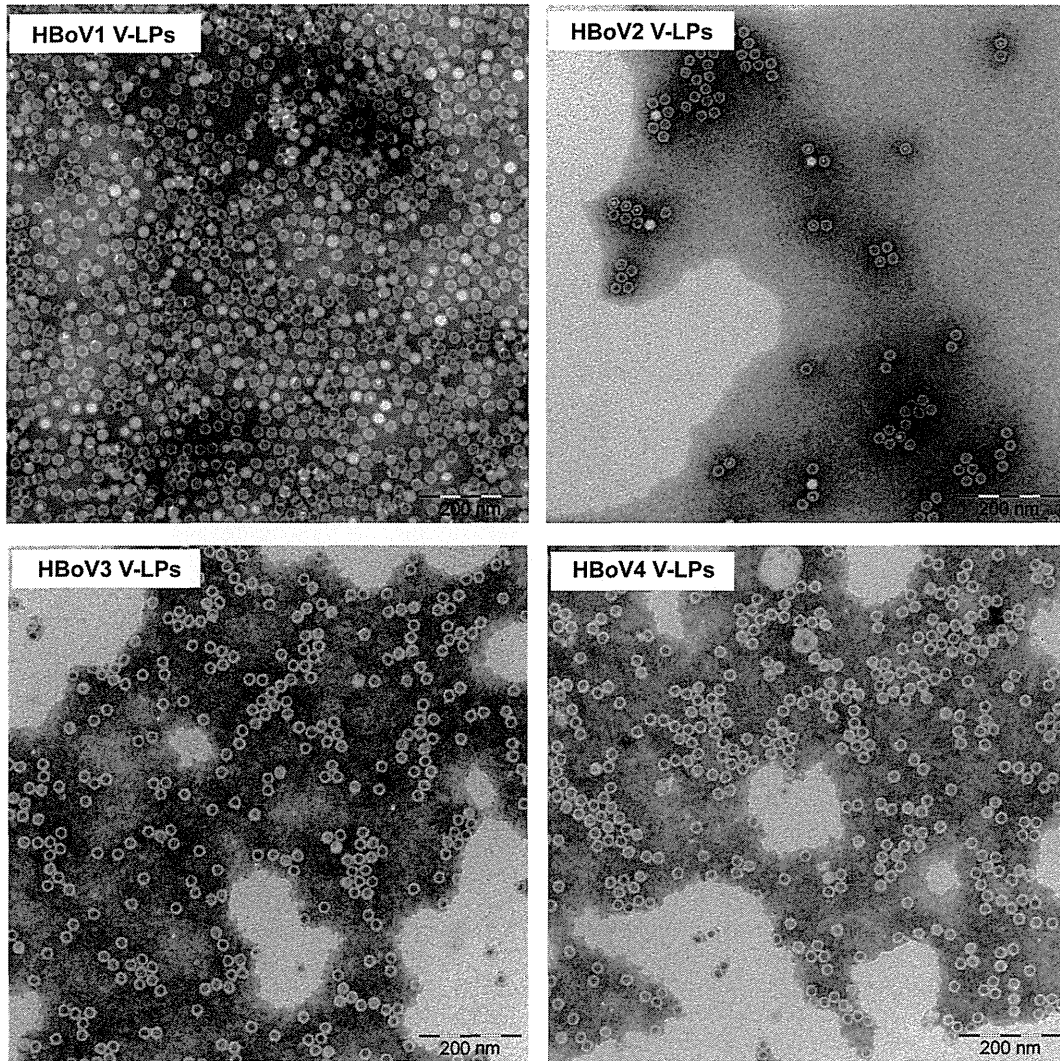
### 3.2. Recombinant VP2 of HBoV1–4 self-assembled into virus-like particles

The supernatant of recombinant baculovirus-infected Tn5 cells was harvested at 7 days p.i. and subjected to a CsCl density gradient centrifugation. The gradient was fractionated into 20 aliquots, and analyzed by SDS-PAGE followed by Coomassie blue staining. As shown in Fig. 2, p61 appeared in fraction 8 at a density of 1.30 g/cm<sup>3</sup>



**Fig. 2.** Purification of HBoVLPs. The recombinant baculovirus-infected Tn5 cells were harvested on 7 day p.i., and the supernatant was centrifuged for 3 h at 32,000 rpm in a Beckman SW32Ti rotor. The pellet was resuspended in 500  $\mu$ L EX-CELL™ 405, and then purified by CsCl equilibrium density gradient centrifugation. Aliquots from each fraction were analyzed by electrophoresis on 5% to 20% polyacrylamide gel, and stained with Coomassie blue.





**Fig. 3.** Electron micrographs of HBoVLPs. The fraction containing p61 protein was stained with 2% uranyl acetate and observed by EM. Bar, 200 nm.

in all HBoVs1–4. The control VLPs derived from B19 VP2 appeared in fractions 12 and 13 with the density of 1.28 g/cm<sup>3</sup>. The p61-containing fraction no. 8 was diluted and VLPs were sedimented by centrifugation and observed by electron microscopy. As shown in Fig. 3, empty spherical particles with an approx. 22-nm dia. were observed. The size of these particles was similar to that of the authentic HBoV particles (Brieu et al., 2007; Christensen et al., 2010; Santos et al., 2010), indicating that the recombinant VP2 self-assembled into virus-like particles (HBoV1 VLPs, HBoV2 VLPs, HBoV3 VLPs and HBoV4 VLPs). The yields of the purified HBoV VLPs obtained from 10<sup>7</sup> infected Tn5 cells were 1.2 mg (HBoV1 VLPs), 0.5 mg (HBoV2 VLPs), 1.0 mg (HBoV3 VLPs), and 0.8 mg (HBoV4 VLPs), respectively.

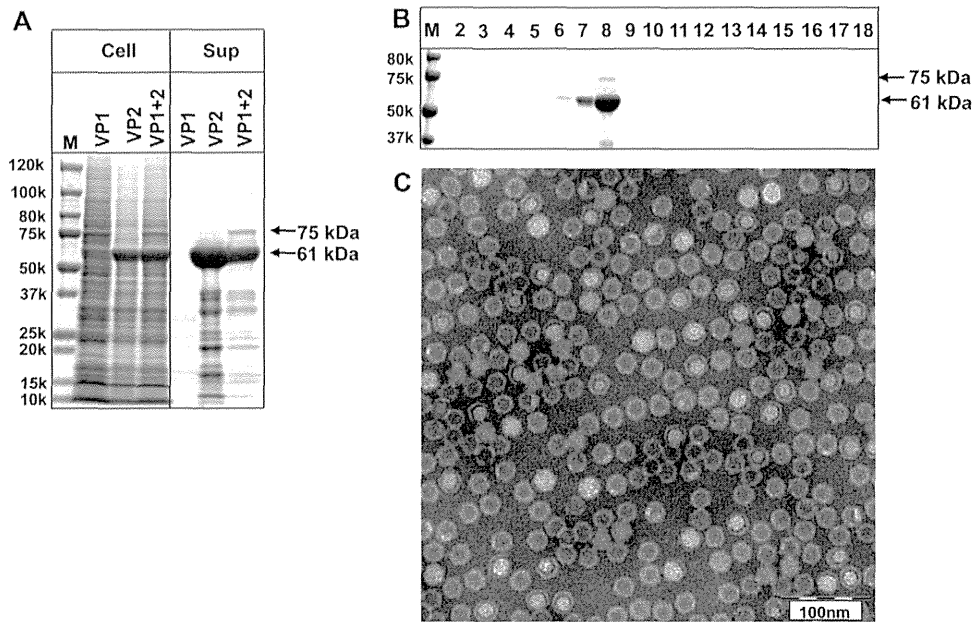
### 3.3. Co-expression of VP1 and VP2 proteins of HBoV1

Although VP1 of HBoV is known as a minor component of the viral particles, its role in the morphogenesis of HBoV has not yet been well determined. When HBoV1 VP1 was expressed by the recombinant baculovirus AchBoV1-VP1 in Tn5 cells, the VP1 protein with a molecular mass of 75 kDa (p75) was detected in infected Tn5 cells. However, the expression level of the VP1 protein was quite low and not released into the culture supernatant (Fig. 4A).

In contrast, when two recombinant baculoviruses, AchBoV1-VP1 and AchBoV1-VP2, were co-infected to Tn5 cells, both VP1 and VP2 proteins were observed in both the cell lysate and culture supernatant (Fig. 4A). As shown in Fig. 4B, p75 and p61 appeared in the same fraction no. 8 after CsCl gradient centrifugation. Amino acids analysis showed that the N-terminal sequences of p75 was Pro-Pro-Ile-Lys-Arg, which is identical to that of the VP1 N-terminus amino acid sequence. When fraction no. 8 was observed by electron microscopy, spherical empty particles with an approx. 22-nm dia. were observed, and its morphology observed by electron microscopy was similar to that of VP2 VLPs (Fig. 4C). These results indicate that the ratio between VP1 and VP2 in a VLP is quite different if VP1 protein was incorporated into the VLP evenly. These results also suggested that the VP1 protein is a minor structural protein of HBoVs.

### 3.4. Antigenic cross-reactivity among HBoV VLPs

To examine the antigenicity of HBoV VLPs, rabbits were immunized with HBoV1 VLPs, HBoV2 VLPs, HBoV3 VLPs, and HBoV4 VLPs, respectively, and we determined anti-VLPs IgG titers by ELISA (Fig. 5). After three injections, all of those rabbits elicited high levels of IgG antibody against each homologous antigen at titers of 1:409,600 (HBoV1 VLPs), 1:819,200 (HBoV2 VLPs), 1:409,600



**Fig. 4.** Co-expression of HBoV1-VP1 and HBoV1-VP2. Insect cells (Tn5) were infected with either AchBoV1-VP1 or AchBoV1-VP2, or co-infected with AchBoV1-VP1 and AchBoV1-VP2. Expression of VP1 and VP2 proteins in cell lysates was detected at 3 days p.i. The supernatant of the infected Tn5 cells was harvested on 7 days p.i., and concentrated by ultracentrifugation. (A) The proteins were separated by SDS-PAGE and stained with Coomassie blue. (B) The VLPs recovered from the supernatant of the co-infected cells were purified by CsCl gradient centrifugation. Aliquots from each fraction were analyzed by SDS-PAGE, and stained with Coomassie blue. (C) EM of fraction no. 8. Bar, 100 nm.

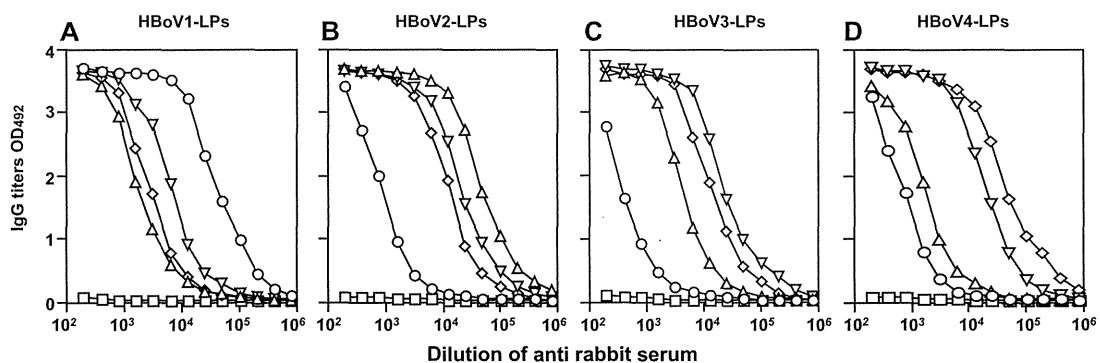
(HBoV3 VLPs), and 1:819,200 (HBoV4 VLPs), respectively. None of the pre-immune sera showed any reaction against heterologous or homologous VLPs. Each serum was cross-reacted with heterologous VLPs, but the titers were lower than those with the homologous antigen. The anti-HBoV1 VLPs IgG reacted with HBoV2 VLPs, HBoV3 VLPs and HBoV4 VLPs at titers of 6400. In contrast, anti-HBoV2 VLPs IgG reacted with HBoV3 VLPs and HBoV4 VLPs at titers of 25,600, and anti-HBoV3 VLPs IgG reacted with HBoV2 VLPs and HBoV4 VLPs at titers of 204,800. Anti-HBoV4 VLPs IgG reacted with HBoV2 VLPs and HBoV3 VLPs with titers of 102,400 to 204,800, respectively. The results indicated that HBoVs are serologically cross-reactive with each other, suggesting that HBoVs possess the same antigenic epitope(s).

### 3.5. Seroprevalence of HBoVs in a general population in Japan

We measured the seroprevalences of anti-HBoV1, HBoV2, HBoV3 and HBoV4 IgG and IgM using 372 sera collected from healthy individuals who lived in a southern prefecture of Japan.

The average positive rates of IgG were 93.0% (346/372), 70.1% (261/372), 67.7% (252/372) and 76.6% (285/372) for HBoV1, HBoV2, HBoV3 and HBoV4, respectively, whereas no significant difference was found in those of IgG among different age groups ( $p < 0.05$ ). A high prevalence of antibodies against HBoVs was already found in the population of healthy 1- to 9-year-old children in Japan, indicating that HBoVs infection is common and the infection occurs during childhood.

The average positive rates for HBoV1, HBoV2, HBoV3 and HBoV4 in serum IgG from males and females were 93.3% and 92.9% for HBoV1, 73.9% and 68.3% for HBoV2, 68.7.6% and 65.8% for HBoV3, and 78.2% and 75.8% for HBoV4, respectively; no significant difference was seen between males and females in the IgG-positive rate for all genotypes ( $p < 0.05$ ). In contrast to the relatively high positive rate in serum IgG, only one individual each out of the 372 (0.54%) was found to be IgM-positive for HBoV1 and HBoV3, with low OD values of 0.35 and 0.43, respectively. No serum IgM antibody against HBoV2 and HBoV4 was detectable in those serum samples.



**Fig. 5.** Cross-reactivity among HBoVs. We prepared hyperimmune sera against HBoV1-LPs (○), HBoV2-LPs (△), HBoV3-LPs (▽) and HBoV4-LPs (◇) by inoculating rabbits with HBoVs-LPs. Cross-reactivities of HBoVs-LPs found by ELISA are shown. The ELISA was performed using HBoV1-LPs (A), HBoV2-LPs (B), HBoV3-LPs (C), and HBoV4-LPs (D) as the antigen, respectively.

#### 4. Discussion

To date, four species of human bocavirus, HBoV1, HBoV2, HBoV3 and HBoV4, have been identified. Human bocaviruses are associated with respiratory illness and gastroenteric diseases. Because no culture system is available for HBoV growth, the serological diagnosis and vaccine development are difficult, and studies focusing on how to express the capsid protein of HBoVs are necessary. In the present study, we expressed the major capsid protein VP2 of four HBoVs in insect Tn5 cells which self-assembled into VLPs.

Recombinant baculovirus expression systems have long been used to express proteins, especially to generate VLPs of various DNA and RNA viruses. The advantages of this system are efficient expression and the self-assembly of the capsid protein into VLPs. VLPs produced by this system usually release into the cell culture supernatant and retain immunogenicity as well as physicochemical properties of their native virions (Li et al., 1997; Li et al., 2004; Murakami et al., 1999; Baumert et al., 1999; Matsuo et al., 2006). Sf9 and Tn5 cells are commonly used cell lines in baculovirus expression systems (Stewart and Possee, 1993; Wickham and Nemerow, 1993).

Previous studies have shown that the HBoV1 capsid protein VP2 expressed in Sf9 cells self-assembled into VLPs in the cell lysates, but were not released into the culture medium (Kahn et al., 2008; Soderlund-Venermo et al., 2009). Attempts were made to express the HBoV1 capsid protein VP2 in Sf9 cells, but the level of expression was significantly lower than that in Tn5, and it was difficult to obtain a larger amount of the protein (Fig. 1). In contrast, in the case of Tn5, a large amount of HBoV1 VLPs was released into the culture medium, making the purification of the VLPs much easier.

The morphologies of the VLPs derived from four genetically different HBoVs were similar. All of these VLPs had a dia. of 22 nm, similar to those of individual authentic virus particles. The density of HBoVs-LPs was approx. 1.30 g/cm<sup>3</sup>, slightly less than that of the native virus particles, perhaps because the VLPs were empty and did not contain the virus genome. Although VLPs were obtained by the co-expression of HBoV1 VP1 and VP2, the density and morphology of the VLPs did not significantly differ from those of the VP2 VLPs. Further structural studies are needed to clarify the role of VP1 in the VLP formation.

Our immunogenic and antigenic analyses showed that HBoVs-LPs are capable of inducing IgG antibodies as high as 1:409,600 to 1:819,200 in rabbits even without any adjuvant, and we observed different degrees of cross-reactivity among the four HBoVs. The high immunogenicity of the VLPs suggests that HBoVs-LPs are promising candidates for a vaccine against HBoV infection. Phylogenetic studies have shown that the amino acid identity of VP2 among HBoVs was as high as 76.9% to 90.8%, and the antigenic cross-reactivity was confirmed in this study. Notably, based on phylogenetic analyses of the amino acid sequences deduced from the VP1/2 gene, HBoV1 shared amino acid identities of 79.4% to 79.9% with HBoV2, HBoV3, and HBoV4; however, those among HBoV2, HBoV3 and HBoV4 were 90.1% to 91.8%, suggesting that HBoV1 is slightly different from HBoV2, HBoV3, and HBoV4. In fact, the ELISA titers of anti-HBoV1 IgG against VLPs from HBoV2, HBoV3, or HBoV4 were 1:6400, weaker than other heterogeneous anti-HBoVs-LPs IgG.

Consistent with the previous findings obtained from a general population, the IgG-positive rate to HBoV1 reached 93.0% in the healthy Japanese population in the present study, indicating widespread infection in Japan. In addition, the seroprevalences of HBoV2, HBoV3, and HBoV4 were examined for the first time in a general population in Japan and were also found to be relatively high, at 70.1%, 67.7%, and 76%, respectively. Considering the cross-reactivity among HBoVs, diseases associated with HBoVs infection should be further investigated, and other immunological

diagnostic methods such as the detection of HBoV-specific antigens with monoclonal antibodies are needed.

In conclusion, VLPs of four HBoVs – HBoV1, HBoV2, HBoV3 and HBoV4 – were generated by recombinant baculovirus, and their immunogenicity and antigenicity were examined. The VLPs-based ELISA results showed that HBoV infection is common in Japan. The production of HBoV VLPs will contribute to a better understanding of the epidemiology and biological characteristics of HBoVs. HBoV VLPs are also a promising resource for vaccine development.

#### Acknowledgements

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