Dengue Virus Uses a Non-Canonical Function of the Host GBF1-Arf-COPI System for Capsid Protein Accumulation on Lipid Droplets

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ABSTRACT

Dengue viruses cause the most important human viral disease transmitted by mosquitoes. In recent years, a great deal has been learned about molecular details of dengue virus genome replication; however, little is known about genome encapsidation and the functions of the viral capsid protein. During infection, dengue virus capsid progressively accumulates around lipid droplets by an unknown mechanism. Here, we examined the process by which the viral capsid is transported from the ER membrane, where the protein is synthesized, to lipid droplets. Using different methods of intervention, we found that the GBF1-Arf1/Arf4-COPI pathway is necessary for capsid transport to lipid droplets, while the process is independent of both COPII components and Golgi integrity. The transport was sensitive to brefeldin A, while a drug resistant form of GBF1 was sufficient to restore capsid subcellular distribution in infected cells. The mechanism by which lipid droplets gain or lose proteins is still an open question. Our results support a model in which the virus uses a non-canonical function of the COPI system for capsid accumulation on lipid droplets, providing new ideas for antiviral strategies.

INTRODUCTION

Dengue virus (DENV) is considered one of the most important arthropod-borne human pathogen worldwide. This virus is a member of the Flavivirus genus in the Flaviviridae family together with other important pathogens such as yellow fever virus (YFV), West Nile virus (WNV), Saint Luis encephalitis virus (SLEV), and Japanese encephalitis virus (JEV). DENV alone causes about 390 million infections per year. Despite this large public health problem and the urgent need to control DENV infections, vaccines and specific antivirals are still unavailable.

DENV, like other plus strand RNA viruses, induces a profound rearrangement of cellular membranes to provide platforms for viral replication (1-3). DENV RNA synthesis occurs in membranous structures called vesicle packets that function as viral genome factories, which are in close association to viral particle morphogenesis in the ER membrane (3). During this process, the viral capsid protein (C) is responsible for recruiting the viral genome during viral encapsidation, forming a nucleocapsid that buds into the ER lumen, acquiring membranes and the structural proteins E and prM (4, 5). The new viral particle travels through the secretory pathway to be released by exocytosis.

The viral C is a small, highly basic protein that forms homodimers in solution (6, 7). We have previously shown that C progressively accumulates around cellular lipid droplets (LDs) during DENV infection (8). Atomic force microscopy has been recently used to define biochemical properties of this C-LD interaction (9). LDs are dynamic organelles involved in lipid metabolism that regulate storage and turnover of neutral lipids. They contain mostly triacylglycerols and sterol esters surrounded by a phospholipid monolayer, decorated with proteins (10-12). Previous studies using leukocytes have demonstrated an increased amount of LDs per cell in samples from patients with dengue hemorrhagic fever when compared with that from healthy volunteers (13). Similar observations were reported in DENV infected human macrophages and in different infected cell lines, such as A549, Hep-G2 and BHK (8, 13-15). Interestingly, it has been shown that DENV infection triggers the redistribution of the rate-limiting enzyme in lipid biosynthesis, the fatty acid synthetase (FASN) to the ER. It has been demonstrated that binding of FASN to the viral protein NS3 increases cellular fatty acid synthesis (16); and that pharmacological inhibition of FASN activity blocks induction of LDs during DENV infection (8). A relationship between autophagy-mediated lipid droplet degradation and DENV infection has been also observed (17). It has been suggested that lipid degradation was necessary during DENV RNA synthesis. However, inhibition of autophagy appears to affect different stages of viral infection. In this regard, destabilization of beclin-Vps34-Agt14, which inhibits autophagy, was shown to have a modest effect on viral RNA synthesis but a profound effect on particle infectivity, due to defects on peptide pr release from prM during particle maturation (18). Thus, it is likely that lipid biosynthesis and degradation are both necessary during different stages of the viral life cycle. More recently, the Ras-related protein in brain (Rab), GTPase Rab18, which participates in lipid exchange between ER and LD compartments, was shown to be necessary for efficient DENV replication and to have a positive role in LD induction (19). Even though different lines of evidence link DENV infection to LD metabolism, the
functional significance and the mechanisms by which the virus manipulates LDs remain obscure. We have previously reported that C accumulation on LDs is necessary for efficient viral particle production; but the role of LDs in this process is still unknown. Budding viral particles in the ER have been observed next to the pores of vesicle packets, where the viral genome is replicated, suggesting that LDs are not the place of genome recruitment during encapsidation (3). It is possible that accumulation of C on LDs modulates the host lipid metabolism required for viral particle formation or serves as storage of the viral protein in early stages of viral replication (8).

To further examine the function of C on LDs and explore new tools for antiviral intervention, we investigated the process by which C is transported from the ER membrane to LDs during DENV infection. How LDs receive or deliver proteins is still an open question. In recent years, more than 200 cellular proteins have been reported that associate to LDs, including lipid metabolism enzymes, Rab GTPases, SNARE proteins and coatamer components (12). The most prominent lipid droplet associated proteins are the PAT proteins (Perilipin, Adipophilin/ADRP, and the Tail interacting protein/TIP47) (19). Interestingly, a function of components of the vesicle trafficking systems, Coat Protein II and I (COPII and COPI), has been reported for the transport of different cellular proteins to LDs (20-22). Normally COPII vesicles mediate transport of proteins and lipids from the ER to Golgi, while COPI mainly participate in the retrieval of proteins from the Golgi back to the ER. How COP components mediate delivery of proteins to LDs is unclear, but might involve novel functions of this class of proteins. Here, we investigated the involvement of these host factors in the accumulation of C on LDs. Because DENV requires an active vesicle transport system for viral maturation and egress, we restricted our studies to evaluate the C transport in a single viral replication cycle. Using different methods of intervention, we defined that C association to LDs requires a functional GBF1 protein. The process was found to be COPI dependent, COPII independent and did not involve the Golgi complex, supporting a non-canonical function of this cellular transport system. We believe that dissection the host machinery used during DENV infection will provide novel ideas for employing host targets for antiviral approaches.

RESULTS

Delivery of capsid to LDs is BFA sensitive

It has been previously reported that the DENV C protein associates to different cellular compartments during viral infection (8, 23, 24). Using immunofluorescence assays and subcellular fractionation of DENV infected cells, we observed C in the nucleus and cytoplasm (Fig. 1A and 1B). During viral infection, C is translated as a polyprotein associated to ER membranes and it is proteolytically processed to release the mature protein. Mature C was detected by western blot at 12, 16 and 20 h post infection in membrane, cytoplasmic and nuclear fractions (Fig. 1B). Immunofluorescence studies showed that the cytoplasmic C partially co-localized with the host protein TIP-47 and the viral E protein, which associate to LDs and ER, respectively (Fig. 1A). In the nucleus, C co-localized with the nucleolar B23 protein (Fig. 1A). Visualization of proteins on LDs is limited when methanol is used as fixation method due to extraction of lipids (25). Because paraformaldehyde (PFA) preserves the integrity of LDs, this method was elected for detection of proteins on LDs (Fig. 1B). We conclude that in DENV infected cells, C accumulates in nucleolus, LDs and ER membranes.

The mechanism by which host proteins are transported to LDs is still not fully understood. It is possible that C directly associates to nascent droplets after proteolytic processing or, alternatively, uses a specific transport system. In this regard, recent reports have implicated components of COPII and COPI complexes in the transport of some cellular proteins to LDs (20, 22). It has been shown that COPI and COPII are necessary for ADRP and ATGL accumulation on LDs, and this process was sensitive to brefeldin A (BFA), a GTP exchange factor inhibitor that blocks COPI activity. In contrast, other proteins such as TIP47, were found to be independent of COPI and COPII systems (20, 22). Thus, we examined the requirement of host proteins for C accumulation on LDs in DENV infected cells. Because BFA was reported to inhibit ADRP and ATGL accumulation on LDs, we first evaluated C distribution in cells treated or not with this drug. It is important to clarify that DENV particle morphogenesis and release requires a functional secretory pathway (26); and it has been demonstrated that BFA inhibits viral particle secretion (27). Therefore, to avoid new rounds of infections that would be sensitive to BFA, we examined the impact of the drug on C subcellular distribution in a single round of infection. To this end, cells were treated with different concentrations of BFA at 8h post-infection and pre-treated with oleate to promote formation of LDs. The correlation coefficient of C with a LD marker was drastically reduced in the presence of BFA, while no effect on C accumulation in nucleus was observed (Fig. 2A and 2B). The total amount of mature C detected in cells treated or not with BFA was similar as determined by western blot (Fig. 2C), indicating that BFA did not affect viral translation or RNA amplification in the conditions used. The effect of different concentrations of drug indicated that concentrations as low as 0.05 μg/ml greatly reduced C on LDs (Fig. 2D). Moreover, the effect was found to be reversible. Localization of C on LDs was recovered 2h after incubation of cells in media without BFA (Fig. 2E). The results indicate that BFA did not significantly change the total amount of C neither the nucleolar accumulation of the viral protein in infected cells; however, it greatly reduced protein association to LDs.

BFA inhibits several GTP exchange factors (GBF1, BIG1 and BIG2). Because golgicide A (GCA), has been shown to be more specific for GBF1 inhibition, which is the GEF involved in the Arf1-COPI pathway (28), the effect of GCA on C association to LDs was also examined. GCA greatly reduced C accumulation on LDs (Fig. 2F), suggesting that transport of C to LDs is sensitive to GBF1 inhibition.
In these studies we could not evaluate the impact of the lack of C association to LDs by BFA on viral particle production because the drug inhibits viral particle secretion independently of C (27). Accordingly, BFA greatly decreased viral titers in our experimental conditions (Fig. S1).

**Disruption of the Golgi complex does not alter capsid localization on LDs**

Inhibition of GBF1 and COPI functions produces alterations in the secretory pathway, including Golgi disassembly, phenotype observed in the presence of BFA (22, 29). To examine whether the BFA effect on C distribution was the result of a general alteration on the host vesicular transport system, a dominant negative form of Rab1b was used. This protein is a GTPase that plays a role in ER to Golgi transport (30). It has been previously demonstrated that a dominant negative form of Rab1b causes Golgi reabsorption into the ER, analogous to that described for BFA treatment (29). Thus, we expressed wild type or dominant negative Rab1b (N121I) protein fused to GFP in DENV infected cells and evaluated C association to LDs. As a control for function of the dominant negative protein, localization of giantin, a Golgi associated protein, was examined. N121I Rab1b was mainly cytosolic and induced rapid Golgi disruption as observed by giantin redistribution (Fig. 3A), but did not alter C association to LDs (Fig. 3B). Cells expressing Rab1b-GFP displayed the typical ring-like structures of C on LDs (Fig. 3B). The results suggest that components of the ER-Golgi machinery are involved in C localization on LDs, independently of Rab1b activity and Golgi integrity but sensitive to BFA.

**Capsid accumulation on LDs requires COPI components but is independent of COPII function**

COPI and COPII systems have been implicated in the transport of host proteins to LDs (20-22, 31); therefore, we analysed the requirement of components of these systems on C transport to LDs during DENV infection. We used siRNAs directed to different subunits of COPI and COPII in A549 cells. Cells were transfected with siRNAs specific for βCOP subunit (COPI component), Sec13 (COPII component) or nonrelated control siRNAs, and then infected with DENV (Fig. 4A). Depletion of βCOP was about 70% and showed a drastic effect on C association to LDs without altering TIP47 localization (used as control). In contrast, depletion of Sec13 did not alter the amount of C nor its association to LDs, which remained co-localized with TIP47 (Fig. 4A and 4B). The results suggest that C transport to LDs in infected cells depends on COPI but is independent of COPII components.

The possible involvement of COPI on C association to LDs lead us to analyze a function of Arf1 in this process. To study this possibility, we evaluated the depletion of Arf1 on C distribution in DENV infections. Depletion of Arf1 in infected A549 cells was almost complete, while C and TIP47 remained associated to LDs (Fig. 4D and 4E). This result was unexpected because Arf1 activity is proposed to be necessary for COPI recruitment. It is possible that COPI is directly recruited by GBF1, without Arf1 requirement (32).

Alternatively, other Arf isoforms could participate in this process. Kahn and colleagues have previously reported a redundancy of Arf functions in the cell (33). Therefore, we analyzed depletion of different Arfs and found that simultaneous knock down of Arf1 and Arf4 largely reduced C transport to LDs, while cells lacking only Arf4 did not affect C subcellular distribution (Fig. 4D and 4E). These studies support the idea that localization of C on LDs requires recruitment of COPI by either Arf1 or Arf4.

**BFA resistant GBF1 restores capsid delivery to LDs in DENV infected cells**

To further examine the role of GBF1 on C association to LDs different methods were used. First, the effect of expressing a WT or a dominant negative form of GBF1 (E794K) was evaluated. A549 cells were infected with DENV and then transfected with expression constructs encoding GBF1 fused to YFP (Fig. 5A). While expression of the WT protein did not affect C association to LD, the E794K inactive protein reduced about 70% the amount of cells with C on LDs (Fig. 5A), suggesting the requirement of an active GBF1 protein. To confirm this observation, the effect of RNAi mediated depletion of GBF1 on C localization was also tested. The efficiency of GBF1 depletion was observed by western blots (Fig. 5B). Immunofluorescence analysis using antibodies to detect both GBF1 and C indicated that infected cells lacking GBF1 (without green signal in Fig. 5B) also lacked C localization on LDs, while cells expressing detectable levels of endogenous GBF1 showed a clear association of C to LDs.

Taken together, it is likely that BFA inhibition of GBF1 is responsible for the phenotype observed. Thus, we examined whether a GBF1 mutant that is resistant to BFA was able to reverse the effect of the drug. Cells were transfected with plasmids expressing YFP alone, GBF1-WT or BFA-resistant GBF1 (M832L) fused to YFP, and then infected with DENV in the presence or absence of BFA (Fig. 6A). In the absence of BFA, between 90 to 95% of the cells that expressed YFP showed C on LDs, whether they were transfected with the control plasmid or either GBF1 constructs, suggesting that GBF1 overexpression had no effect on C localization (Fig. 6B). In the presence of BFA, the amount of cells expressing YFP control with C on LDs was reduced to 18%, whereas it was about 40% and 80% in WT or M832L GBF1 expressing cells (Fig. 6B). It was evident that cells expressing or not the BFA resistant form of GBF1 showed a remarkable difference on C distribution in the presence of drug (Fig. 6A). The results indicate that a BFA resistant form of GBF1 was sufficient to reverse the effect of the drug, confirming the requirement of an active GBF1 for C subcellular distribution during DENV infection.

**DISCUSSION**

Different lines of evidence support a link between DENV replication and LD metabolism. Thus, it is important to further understand why and how the virus manipulates LDs. Here, we examined the requirement of host factors for C accumulation around LDs during infection with the final goal of identifying possible antiviral targets. We defined that an active GBF1-Arf-
COPI pathway is necessary for C transport from the ER to LDs in human infected cells. The process was found to be dependent of COPI but independent of Golgi integrity, supporting a non-canonical function of components of this vesicle transport system.

During DENV infection C distributes between different cellular compartments: nucleolus, LDs and ER membranes. This distribution is observed early after infection, suggesting that it is not a consequence of cell damage during viral replication (Fig. 1). In the viral polyprotein, C is followed by an anchor peptide that directs prM-E to the ER lumen. It has been shown that the anchor peptide is not necessary for C accumulation on LDs (8). Release of C by the viral NS2B3 cleaves the anchor peptide rendering a mature protein that remains attached to the cytoplasmic side of the ER membrane (34), and then it is transported to LDs and nucleus. Why the protein accumulates in these cellular compartments is still unknown. We have previously shown that C could be toxic for viral RNA synthesis due to its high affinity for nucleic acids. In this regard, it is possible that C is sequestered in different organelles early during infection to avoid premature interaction with the viral RNA. However, more studies are necessary to define mechanistic aspects of C functions in the nucleus and LDs.

Previous studies using genome wide RNAi screens in drosophila, have linked COPI components with LD metabolism and composition (21, 22). It has been shown that COPI regulate lipid homeostasis, likely by promoting transport of enzymes, like ATGL, to the surface of LDs (22). Here, we used different methods to evaluate the involvement of components of this trafficking system on C transport. Because depletion or inhibition of these components could alter the composition and abundance of LDs, the experiments were performed in cells pretreated with oleate. Our results support the idea that an active GBF1 participates in the transport of C to LDs (Fig. 5 and 6). It is possible that the cellular protein accumulates on viral induced membranes and activates Arf1 and/or Arf4 to recruit COPI (Fig. 4). Depletion of COPI subunits reduced transport of C to LDs, while depletion of Sec13 of COPII did not alter C distribution in DENV infected cells (Fig. 4). Therefore, it is unlikely that C is first transported to a cis-Golgi compartment before it is directed to LDs. In this regard, a dominant negative form of Rab1b that induces Golgi disruption did not affect C subcellular distribution. We also observed that BFA greatly reduced C accumulation on LDs while it did not alter its transport to the nucleolus (Fig. 2). The effect of BFA was reversible by drug removal and its specificity on GBF1 was demonstrated by the recovery of C on LDs in the presence of a BFA resistant form of GBF1 (Fig 6). We propose that COPI/Arf/GBF1 system mediate transport of C from the place where it is synthesized on the ER membranes to LDs, independently of Golgi integrity.

The best described function of COPI vesicles is to mediate retrograde vesicle transport between Golgi cisternae and from the Golgi to the ER, while COPII mediates anterograde transport mainly from the ER; however, it is evident that the canonical vesicular transport of cargo, as occurs in the secretory pathway, is not possible for protein transport to LDs. The monolayer of phospholipids that constitute the LD surface cannot fuse to a typical bilayer of a coated vesicle. Different possibilities that explain the transport of proteins have been suggested. For instance, lateral movement of proteins from ERES/ERGIC sites to LDs, or loading of nascent droplets of specific proteins (20, 35). A very recent report showed that cells lacking Arf1 and COPII have an increased amount of phospholipids on LDs, resulting in decreased LD surface tension and impairment to form bridges with the ER (36). Based on these recent findings, modulation of ER-LD membrane bridges, without involving vesicle formation, was proposed as a new mechanism for controlling protein transport from the ER to LDs (36). Our results are in agreement with this model, supporting the idea that C is transported from the ER to LDs using new functions of COPI.

An emerging theme in pathogen biology is the interaction of bacteria, viruses, and protozoan with LDs (37-39). The mechanism by which these pathogens co-opt LDs remains to be elucidated. In the case of viruses, a link between hepatitis C virus (HCV) replication and LDs has been extensively documented (40-42). During HCV infection, the core protein, as well as other viral proteins, accumulates on the surface of LDs (43-45). This localization of core has been proposed to play an important role in viral encapsidation (40, 42, 44-49). Interestingly, a recent study analyzing the trafficking of core during HCV infection suggested that BFA treatment increase the amount of the protein on LDs (50). Thus, DENV C and HCV core appear to use distinct mechanisms for LD association, one sensitive and the other one resistant to BFA, respectively. Taking into account previous studies, there are at least two different pathways involved in delivering proteins to LDs, one dependent on COPI (and COPII in some cases), sensitive to BFA (such as ADRP, ATGL and DENV C); and the other one independent on these trafficking systems and resistant to BFA (such as TIP47 and possibly HCV core).

Different viruses subvert the GBF1/Arf/COPI system during infection. In several cases, components of this pathway are involved in non-canonical functions such as the case of GBF1 on picornavirus replication (51-54) and GBF1-Arf1 in Ebola virion formation (55). Here, we found that DENV uses this host trafficking system for C accumulation on LDs. Because the role of GBF1/Arf/COPI on LD protein transport is just emerging as a novel function of these proteins, further studies using different viruses that co-opt LDs will help to uncover details of this cellular process and hopefully provide new ideas for antiviral approaches.

MATERIALS AND METHODS

Cells and viruses
A549 human epithelial lung cells were cultured in Dulbecco’s modified Eagle’s medium F-12 (Ham) supplemented with 10% fetal bovine serum, 100 U/ml penicillin, and 100 µg/ml streptomycin. Stocks of DENV-2 16681 were prepared in mosquito cells

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Antibodies and immunofluorescence assays

For immunofluorescence and Western blot assays the following antibodies were used: mouse monoclonal anti-B23 (Abcam), Guinea pig polyclonal anti-TIP47 (Fitzgerald), rabbit polyclonal anti-Sec 13 (Abcam), mouse monoclonal anti-Tubulin (Santa Cruz), mouse monoclonal anti-Arf1 (Abcam), rabbit polyclonal anti-Arf4 (Proteintech), rabbit polyclonal anti-Giantin (Abcam), and mouse monoclonal anti-GBF1 (BD Transduction Laboratories). Rabbit polyclonal and mouse polyclonal anti-DENV capsid were obtained in our laboratory. For DENV envelope detection, mouse monoclonal anti-E antibody E18 was used (57). For lipid droplets staining, cells were incubated with BODIPY493/503 (Molecular Probes).

Immunofluorescence assays were performed as previously described (6). Briefly, A549 cells were seeded into 24-well plates containing glass coverslip. Twenty four hours later, the cells were infected with DENV-2 using a multiplicity of infection of 10. At 24 hours post-infection coverslip were collected and the cells were fixed with paraformaldehyde 4%, sucrose 4%, in PBS pH 7.4 at room temperature for 20 minutes. Alternatively, cells were fixed with methanol for 20 minutes at -20°C. PFA fixed cells were then permeated with 0.1% Triton X-100 for 4 minutes at room temperature. Images were obtained with a Carl Zeiss LSM 5 Pascal confocal microscope. For quantification of LD associated capsid, about 200 cells were scored for each experiment. Mean and standard deviation were indicated in each case. Three independent experiments were performed in each condition.

Subcellular fractionation

For subcellular fractionation Qproteome Cell Compartment kit (QIAGEN) was used according manufacturer’s protocol. The kit allows sequential isolation of proteins associated with the cytosol, membranes and nucleus from cell lysates. Different extraction buffers were added sequentially to an infected cell pellet, and the respective fractions were isolated by centrifugation. Each fraction was evaluated by Western blot with specific markers: anti-TIP47, anti-calnexin, and anti-Histone H3 antibodies. Additionally, each fraction was evaluated with anti-capsid antibodies at different times post-infection.

Brefeldin A and Golgicide A treatments

Confluent A549 cells were infected with DENV-2 16681 using a multiplicity of infection of 10. After 1 hour, cells were washed with PBS and DMEM-F12 medium supplemented with 10% serum was added. At 8 hours post-infection fresh medium with 0, 0.05, 0.1 or 0.5 µg/ml of Brefeldin A (Sigma), or 50µM of Golgicide A (Sigma) was added. The medium was supplemented with 100 µM of oleic acid (Sigma) complexed with BSA (58). For immunofluorescence the cells were fixed at 24 hours post-infection.

Plasmids and transfection

YFP-tagged GBF1 constructs were kindly provided by J. Bonifacino and C. Jackson (59). GFP-Rab1b N121I mutant has been previously described (29). For expression of heterologous proteins, confluent A549 cells were infected with DENV-2 and at 4 hours post-infection transfected with different plasmids using Lipofectamine 2000 (Invitrogen), according to manufacturer’s protocol. At 24 hours post-infection the cells were fixed with paraformaldehyde and immunofluorescences were performed.

RNA interference

RNA interference experiments were carried out using siGENOME ON-TARGET plus SMART pool siRNA oligonucleotides (Dharmacon RNA Technologies, Lafayette, CO, USA). RNAi were directed to GBF1 (8729), Sec13 (6396), β COP (1314 and 1315), Arf1 (375) or Arf4 (378). The control no-related siRNA used was directed to Renilla luciferase. At 24 hours after seeded in 24-well plates, A549 cells were transfected with the corresponding siRNA using Oligofectamine (Invitrogen). Briefly, 25 pmol of siRNA in 50 µl of Opti-MEM (Invitrogen) were mixed with 2 µl of Oligofectamine supplemented with 100 µM of oleic acid (Sigma) complexed with BSA. The mix was added to a 50% confluent A549 cells monolayer and incubated overnight. Then, the medium was replaced with complete DMEM F12 medium. After 48 hours of transfection, except in cases that indicate otherwise, cells were infected with DENV-2 and incubated for 24 hours.

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FIGURE LEGENDS
Figure 1. Subcellular localization of DENV C in A549 infected cells. A. Immunofluorescence of C in infected cells using methanol as fixation method. A549 cells were infected with DENV and localization of C was analyzed in respect to different markers using antibodies anti-B23 (nucleolus), anti-DENV envelope E protein and anti-TIP47 (lipid droplets). Arrows indicate co-localization of C with markers for each case. B. Western blots with lysates of infected cells fractionated into cytoplasmic (Cyt), membranes (Me), and nuclear (Nu) fractions. The viral protein was detected with anti-C antibodies at 12, 16, and 20 hours post-infection, as indicated. The samples were immunoblotted with anti-histone H3, anti-calnexina and anti-TIP47, as markers for each fraction. C. Immunofluorescence of C in infected cells using paraformaldehyde as fixation method allows lipid droplet visualization. Lipid droplets were stained with BODIPY and localization of viral C and the host TIP47 was determined with specific antibodies as indicated in each case.

Figure 2. Brefeldin A blocks C localization on LDs in DENV infected cells. A. Immunofluorescence showing the effect of BFA on C associated to LDs. Specific antibodies against C and TIP47 were used in infected cells as indicated on the top. LDs were stained with BODIPY. Representative confocal images are shown for each case in the absence or presence of 0.05 µg/ml of BFA. Quantification of the percentage of C that co-localizes with TIP47 (mean +/- standard deviation) was performed using ImajeJ (JACoP plug-in), Manders’ coefficient is shown. B. Immunofluorescence showing the effect of BFA on C accumulation in nucleolus using methanol as fixation method. Representative confocal images using antibodies against C and B23 (marker of nucleolus) are shown in the absence or presence of 0.05 µg/ml of BFA. Quantification of the percentage of C that co-localizes with B23 (mean +/- standard deviation) was performed using ImajeJ (JACoP plug-in), Manders’ coefficient is shown. C. Total amount of C in cells treated or not with BFA. Cell extracts were used for immunoblots with specific anti-C antibodies. D. Effect of different concentrations of BFA on C association to LDs. Confluent A549 infected cells were treated with 0.1 µg/ml of BFA; then, the BFA was removed and cells incubated with fresh media. Immunofluorescence at 60, 90 and 120 min post-infection, as indicated. The samples were immunoblotted with anti-histone H3, anti-calnexina and anti-TIP47, as markers for each fraction. E. Immunofluorescence of C on LDs 8 h post-infection with 0, 0.05, 0.1 or 0.5 µg/ml of BFA. At 24 h post-infection cells were immunostained with anti-TIP47 and anti-C antibodies and the percentage of cells with C on LDs was determined. About 200 cells were scored; bars indicate the standard error of the mean. F. The effect of BFA on C association to LDs is reversible. A549 infected cells were treated with 0.1 µg/ml of BFA; then, the BFA was removed and cells incubated with fresh media. Immunofluorescence at 60, 90 and 120 min were used to evaluate C and TIP47 on LDs. The percentage of cells with C on LDs is shown for each time. About 200 cells were scored in duplicates. Bars indicate the standard error of the mean. G. Golgicide A impairs C localization on LDs. Immunofluorescence showing the effect of 50 µM of Golgicide A on C association to LDs in A549 cells. Representative confocal images of cells treated or not with the drug (Control or GCA) are shown stained with BODIPY for LDs, or labeled with anti-TIP47 and anti-C antibodies as indicated in each case. Right panel, quantification of the percentage of capsid that co-localizes with TIP47 (mean +/- standard deviation), Manders’ coefficient is shown.

Figure 3. Golgi integrity is not required for C association to LDs. A. Effect of Rab1b dominant negative on giantin distribution. On the left, quantification of cells with giantin in Golgi, expressing either the WT or the dominant negative (DN) form of Rab1b (N121I), as indicated. On the right, Representative confocal images showing the effect of expressing a DN form of GFP-Rab1b (N121I) on the cellular distribution of giantin. Arrows indicate cells expressing or not GFP-Rab1b (N121I), labeled with anti-giantin antibodies. B. On the left, quantification of cells with C on LDs, expressing either the WT or the DN form of GFP-Rab1b, as indicated. On the right, representative confocal images showing the effect of expressing the DN form of GFP-Rab1b (N121I) on the cellular distribution of the viral C protein.

Figure 4. COPI but not COPII components are required for C localization on LDs. A. Immunofluorescence showing the effect of COPI depletion on C distribution in DENV infected cells. A549 cells were transfected with non-related siRNA directed to Renilla luciferase (siRNA NR), siRNAs against the β subunit of COPI (siRNA βCOP), or siRNAs against the COPII component Sec13 (siRNA Sec13), as indicated on the top of each panel. The cells were infected with DENV-2 and immune-stained with anti-TIP47 as control and anti-C. Representative confocal images are shown. B. Immunoblot analysis showing βCOP and Sec13 depletion. Extracts of A549 cells transfected with siRNA βCOP (on the left) and siRNA Sec13 (on the right), were immunoblotted with specific antibodies. The same membranes were immunoblotted with anti-tubulin as a loading control, and antibodies against the viral C protein, as indicated on the left of each panel. C. Quantification of the percentage of cells with C on LDs after interference with siRNA against βCOP, Sec13 or a nonrelated siRNA control (NR). In each case, 200 cells were scored for the presence of C on LDs in three independent experiments. The bars indicate the standard error of the mean. D. Redundancy of Arf1 and Arf4 for C association to LDs in DENV infected cells. Immunoblot showing cells treated with siRNA directed to Arf1, Arf4, Arf1+Arf4, or control (NR). Antibodies anti-Arf1, anti-Arf4 or anti-tubulin were used as indicated on each case. E. Effect of Arfs
depletion on C accumulation on LDs. Quantification of the percentage of cells with C on LDs in cells interfered with a NR siRNA or siRNAs against Arf1, Arf4 or both. In each case, 200 cells were scored for the presence of C on LDs in three independent experiments. The bars indicate standard error of the mean.

**Figure 5.** GBF1 function is necessary for C accumulation on LDs. **A.** On the left, representative confocal images showing the effect of expressing YFP-GBF1 WT or E794K (dominant negative, DN) on the cellular distribution of C. A549 cells were infected with DENV-2 and 4h later transfected with the GBF1 expressing plasmids. Immunofluorescence was performed 24 h post-infection using anti-capsid antibodies. On the right, the amount of YFP positive cells with C on LDs were scored and shown as percentage of cells with C on LDs. About 100 cells were scored in each case in duplicates. Errors bars indicate the standard error of the mean. **B.** Effect of GBF1 depletion on C accumulation on LDs. On the left, immunoblot showing GBF1 depletion. Extracts of A549 cells transfected with siRNAs for GBF1 or NR siRNAs were immunoblotted with anti-GBF1 antibodies. The same membrane was immunoblotted with anti-tubulin as a loading control, as indicated. On the left, representative confocal image showing A549 cells depleted or not of GBF1 using anti-GBF1 antibodies (green) and indicating the distribution of the viral C in infected cells by anti-C antibodies (red).

**Figure 6.** BFA-resistant GBF1 reverts the effect of BFA on C localization. **A.** Representative confocal images showing the effect of expressing YFP-GBF1- M832L (BFA resistant) on the subcellular distribution of C in the presence and absence of BFA. A549 cells were infected with DENV-2 and 4h later transfected with the GBF1 expressing plasmids. Immunofluorescence was performed 24 h post-infection using anti-C antibodies. **B.** Percentage of C on LDs of cells expressing YFP control, YFP-GBF1 WT, or YFP-GBF1- M832L, in BFA-treated or non-treated cells, as indicated for each case. About 100 cells expressing YFP were scored for the presence of C on LDs in two independent experiments. Bars indicate standard error of the mean.