Persistent Replication of a Chikungunya Virus Replicon in Human Cells Is Associated with Presence of Stable Cytoplasmic Granules Containing Nonstructural Protein 3

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ABSTRACT Chikungunya virus (CHIKV), a mosquito-borne human pathogen, causes a disabling disease characterized by severe joint pain that can persist for weeks, months, or even years in patients. The nonstructural protein 3 (nsP3) plays essential roles during acute infection, but little is known about the function of nsP3 during chronic disease. Here, we used subdiffraction multicolor microscopy for spatial and temporal analysis of CHIKV nsP3 within human cells that persistently replicate replicon RNA. Round cytoplasmic granules of various sizes (i) contained nsP3 and stress granule assembly factors 1 and 2 (G3BP1/2), (ii) were next to double-stranded RNA foci and nsP1-positive structures, and (iii) were close to the nuclear membrane and the nuclear pore complex protein Nup98. Analysis of protein turnover and mobility by live-cell microscopy revealed that the granules could persist for hours to days, accumulated newly synthesized protein, and moved through the cytoplasm at various speeds. The granules also had a static internal architecture and were stable in cell lysates. Refractory cells that had cleared the noncytotoxic replicon regained the ability to respond to arsenite-induced stress. In summary, nsP3 can form uniquely stable granular structures that persist long-term within the host cell. This continued presence of viral and cellular protein complexes has implications for the study of the pathogenic consequences of lingering CHIKV infection and the development of strategies to mitigate the burden of chronic musculoskeletal disease brought about by a medically important arthropod-borne virus (arbovirus).

IMPORTANCE Chikungunya virus (CHIKV) is a reemerging alphavirus transmitted by mosquitoes and causes transient sickness but also chronic disease affecting muscles and joints. No approved vaccines or antivirals are available. Thus, a better understanding of the viral life cycle and the role of viral proteins can aid in identifying new therapeutic targets. Advances in microscopy and development of noncytotoxic replicons (A. Utt, P. K. Das, M. Varjak, V. Lulla, A. Lulla, A. Merits, J Virol 89:3145-3162, 2015, https://doi.org/10.1128/JVI.03213-14) have allowed researchers to study viral proteins within controlled laboratory environments over extended durations. Here we established human cells that stably replicate replicon RNA and express tagged nonstructural protein 3 (nsP3). The ability to track nsP3 within the host cell and during persistent replication can benefit fundamental research efforts to better understand long-term consequences of the persistence of viral protein complexes and thereby provide the foundation for new therapeutic targets to control CHIKV infection and treat chronic disease symptoms.

KEYWORDS Airyscan, confocal microscopy, neglected tropical diseases, nonstructural
Chikungunya virus (CHIKV), a reemerging arbovirus of the Alphavirus genus, causes a transient illness with debilitating symptoms (fever, headache, rash, myalgia, and arthralgia). Chronic disease is common, and joint pain can persist for months to years (1–3). Half of the patients from the recent Latin American outbreak may develop chronic inflammatory rheumatism, raising the health burden of musculoskeletal disease in areas of endemcity (4, 5). During acute infection, this cytotoxic virus induces apoptosis, leading to direct tissue injury and local inflammation (6–8). Biopsies have also revealed the persistence of CHIKV antigens and RNA in synovial macrophages and muscle tissue (1, 9). CHIKV also persists in mice and nonhuman primate models (10–13). Chronic disease may be a consequence of persistent, replicating, and transcriptionally active CHIKV RNA (13), but an understanding of CHIKV’s long-term effect is still emerging.

The ~12-kb positive-sense RNA genome of CHIKV encodes four nonstructural proteins, nsP1 to nsP4, which make up the viral replication and transcription complex (Fig. 1A) (reviewed in reference 14). A subgenomic RNA expresses six structural proteins. Cellular responses to infection include apoptosis, interferon signaling, stress granule (SG) formation, unfolded protein response, host cell shutoff, and autophagy (reviewed in reference 15). Previous research on alphaviruses established the vital role that nsP3 plays in counteracting cellular responses (16–20) and identified essential protein-protein interactions between nsP3 and host proteins (16, 21–23). However, few studies have systematically investigated the long-term effect of persistently replicating CHIKV RNA and continued expression of proteins such as nsP3 on human cells. Although recent studies characterize the formation of organelles that contain nsP3 during acute infection and transient replication (16, 24–27), a corresponding characterization during persistent CHIKV replication is missing. To address these gaps, we sought to further develop CHIKV replicons capable of persistent replication in human cells and to harness this system for analysis by subdiffraction multicolor microscopy.

We previously characterized transient replication of CHIKV replicons in mammalian and invertebrate cell lines (27) and tagged nsP3 with the versatile SNAP tag for advanced fluorescence microscopy applications (26). The development of a noncytotoxic CHIKV replicon allowed the establishment of persistent replication in a human cell line (28). Here, we extended the SNAP-based labeling system to this noncytotoxic CHIKV replicon and generated a human cell line that persistently replicates replicon RNA and stably expresses SNAP-tagged nsP3. We then characterized nsP3-containing cytoplasmic granular organelles by subdiffraction multicolor microscopy. We used this technique to address questions relating to the subcellular localization of nsP3-containing granules and their stability, composition, and motility. This report is the first to shed light on the persistence of stable intracellular granules of nsP3 within human cells. In turn, understanding the link between the persistence of stable viral protein complexes and pathogenesis has relevance to future studies of chronic CHIKV disease.

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RESULTS

Development of a stable human-origin cell line carrying a SNAP-tagged CHIKV replicon and superresolution microscopy of nsP3-G3BP-containing granules. To determine the intracellular distribution of nsP3, we previously generated a SNAP-tagged replicon construct (26). Whereas this replicon is cytotoxic and replicates transiently, noncytotoxic replicons can establish persistent replication in the human cell line HuH-7 (28). To improve the HuH-7 CHIKV cell line, we added a SNAP-tagged nsP3 to a noncytotoxic replicon (Fig. 1A) and selected puromycin-resistant cells, which are called stable CHIKV cells throughout this paper. Silicon-rhodamine-conjugated Oβ-benzylguanine probes (BG-647-SiR) labeled SNAP-nsP3 and revealed nsP3-containing granules (Fig. 1B) comparable to those formed by a wild-type virus, CHIKVWT, and those
FIG 1 nsP3 has a granular distribution in stable CHIKV cells and infected Huh-7 cells. (A) Schematic representation of tagged reporter viruses and noncytotoxic replicon encoding SNAP-nsP3. SGP, subgenomic promoter; PAC, puromycin-N-acetyltransferase; 2A, foot-and-mouth disease virus (FMDV) 2A autoprotease. (B) Subdiffraction confocal microscopy of BG-647-SiR-labeled stable CHIKV cells, imaged in the far-red channel. Cells
formed by CHIKV\textsuperscript{SNAP-P3} and CHIKV\textsuperscript{ZsGreen-P3} viruses harboring SNAP- or ZsGreen-tagged nsP3 (Fig. 1C to E). Analysis of the sizes of nsP3-containing granules formed in stable CHIKV cells also showed size distributions and diameters comparable to those of granules formed during infection with tagged and untagged viruses (Fig. 1F). Further experiments focused on the characterization of these nsP3-containing granules. Whereas cells infected with CHIKV\textsuperscript{ZsGreen-P3} displayed only a granular nsP3-ZsGreen distribution pattern, cells infected with CHIKV\textsuperscript{SNAP-P3} also made rod-like structures (Fig. 1G), as described previously (26, 27). However, the presence of rods did not correlate with infectivity, as ZsGreen- and SNAP-tagged viruses replicated to similar titers (Table 1). Although the rest of this study used stable CHIKV cells derived from HuH-7 cells, the SNAP-tagged noncytotoxic replicon ChikvRepSNAP could also replicate in the C2C12 mouse myoblast cell line (Fig. 1H).

CHIK nsP3 sequesters G3BP1/2 when expressed alone (17), in the context of a replicon (16, 26), or during virus infection (24, 25), thereby interfering with SG responses. Recent subdiffraction microscopy revealed stable substructures of G3BP1 protein within SGs (30, 31). To determine whether nsP3-containing granules also sequestered G3BP1/2 proteins and contained similar substructures, we imaged stable CHIKV cells with Airyscan microscopy. Airyscan or image scanning microscopy (32, 33) relies on array detectors to reassign photon pixels and oversample the pattern from diffracted light, thereby improving image resolution (1.7-fold) and sensitivity (34). Airyscan outperformed standard confocal microscopy and was sensitive enough to detect small granular structures of nsP3 (Fig. 2A, region of interest [ROI] 1 to 3). Whereas nsP3 appeared to have a diffuse distribution in confocal images, the improved resolution of the Airyscan microscope uncovered an uneven distribution in a large (1.2-\textmu m-diameter) granule, consistent with the presence of substructure (Fig. 2A, ROI 4). Both G3BP1 (Fig. 2B) and G3BP2 (Fig. 2C) were colocalized with these granules, which also had a high fluorescence intensity. The higher sensitivity of the Airyscan method also made small clusters of nsP3 more visible; these clusters had full width at half maximum (FWHM) of 190 to 240 nm and were about 10 times less intense (Fig. 2D and E, see line profiles of fluorescence intensity) than larger granules (Fig. 2B and C, FWHM of 360 to 430 nm). Lastly, live-cell imaging also confirmed that fluorescent nsP3 signals in large granules were nonuniformly distributed within each granule (Fig. 2F).

**Juxtaposition of nsP3-containing granules, dsRNA foci, nsP1-positive structures, nuclear membrane, and Nup98.** During the viral life cycle, nsP3-containing granules sequester G3BP1, thereby blocking SG assembly (16, 17). The relationship between cytoplasmic nsP3-G3BP1 complexes and CHIKV RNA synthesis is less clear;
Characterization of nsP3-G3BP1/2 interaction by subdiffraction microscopy. (A) Comparison of confocal and Airyscan images. Airyscan provided improved resolution and signal-to-noise ratios (SNRs). Airyscan was able to image faint clusters that did not resolve well with standard confocal microscopy (ROI 1 to 3). Note that we use the term “granules” for these protein clusters. Airyscan also revealed differences in fluorescence intensity within large granules that were not apparent with confocal imaging (ROI 4). (B to E) SNAP-nsP3 (cyan) was stained with BG-647-SiR as described in the legend to Fig. 1. G3BP1 or G3BP2 (magenta) was immunostained with specific antibodies. Images were acquired with an Airyscan microscope operated in superresolution mode. For high-contrast display of nsP3-containing granules, contrast was optimized within each image by adjusting the view range in the histogram viewer window of Icy software. Overlaps between cyan and magenta layers appear in white (panels “nsP3/H11001 G3BP1/2”). Line profiles that plot the fluorescence intensity along the line of interest (green) are provided as well, along with the full-width at half maximum (FWHM) of granules. Fluorescence intensity was measured in arbitrary units (A.U.). Images represent single slices, which were extracted from Z-stacks. (F) Live-cell Airyscan microscopy of large granules. Stable CHIKV cells were stained with BG-647-SiR and imaged with an Airyscan microscope operated in the Fast Airyscan mode. Fluorescence intensity maps were created in Icy software and represent relative pixel intensity according to the Jet color map.
viral double-stranded RNA (dsRNA) foci overlap minimally with nsP3-G3BP1-positive clusters in replicons (16). Moreover, few dsRNA foci colocalized with G3BP2 puncta at 6 h postinfection, and even fewer overlapped after 8 h (25). Recently, it was reported that large cytoplasmic and small plasma-membrane-bound G3BP1-nsP3 complexes colocalize with viral genomic RNA during CHIKV infection, with dsRNA foci forming nearby (24). Structural and biochemical data also support a model in which a matrix of nsP3-G3BP1-containing complexes stabilizes replication complexes and shields viral replicative intermediates from the RNA degradation machinery or cytosolic dsRNA sensors (35). Thus, nsP3-G3BP1/2-containing granules appear to play roles in addition to sequestering SG-related proteins.

To further explore the spatial relationship between nsP3 and replication sites in stable CHIKV cells, we visualized SNAP tag-labeled nsP3, together with immunostaining for nsP1 and dsRNA. The antibody against dsRNA was previously used to identify alphavirus replication complexes (36). The fluorescence of ZsGreen in stable CHIKV cells served as an indirect readout of the viral subgenomic RNA (Fig. 1A, cartoon). Rather than completely overlapping with larger nsP3-containing granules, dsRNA foci were in a proximal location and often juxtaposed (Fig. 3, arrowheads). In another example, a dsRNA focus coincided with a smaller nsP3-containing cluster (Fig. 3, cell 2, ROI 1, arrowhead). Ring-like structures coated with nsP1 were also near these dsRNA foci (Fig. 3). The proximity of dsRNA foci, nsP1-coated structures, and nsP3-containing granules suggested that nsP3-containing granules not only sequestered G3BP1/2 protein but also played a role in viral replication.

As described above, nsP3-containing granules were part of a unique microenvironment that also housed dsRNA foci and nsP1. Moreover, a fraction of granules containing nsP3 and G3BP2 were located close to the nuclear membrane (Fig. 4A and B). The nuclear transport factor 2 (NTF2)-like domain of G3BP1 has previously been cocrystallized with FXFG (phenylalanine-glycine motif, where X is usually serine) nucleoporin (Nup) repeat peptides (37) and overexpressed G3BP1 interacts with some nucleoporins, whereas a mutant of G3BP lacking the binding site does not (38).

To further characterize the environment at the nuclear membrane of stable CHIKV cells, we probed for the nuclear pore complex protein Nup98. nsP3-containing granules were detected (i) at the nuclear membrane (Fig. 4C), flanked by Nup98-containing regions, and (ii) near cytoplasmic clusters of Nup98 (Fig. 4D). Thus, we were able to visualize a subset of nsP3-containing granules at the nuclear membrane and near Nup98-containing structures by Airyscan microscopy.

**Imaging the dynamics of nsP3-containing granules within stable CHIKV cells.** SNAP reagents can label live cells, allowing both the analysis of the movement of tagged proteins and pulse-chase studies to examine protein turnover. nsP3-containing granules labeled at the onset of a pulse-chase image experiment (Fig. 5A and B) could be tracked in live-cell imaging experiments over the entire length of a recording that lasted 16 h (Fig. 5B; see Video S1 in the supplemental material). Moreover, stable CHIKV cells still contained "aged" nsP3-containing granules after chase periods of 1 to 2 days (Fig. 5C). The addition of a nonfluorescent SNAP ligand (i.e., quench) in complementary quench-pulse-chase experiments (Fig. 5A) blocked all binding sites of the SNAP-tagged protein pool (Fig. 5D, field of view 1 [FOV1]). After a defined chase period of 3 and 6 h in unlabeled medium, pulsing with the fluorescent SNAP reagent uncovered an unblocked population of nsP3-containing granules, consistent with newly synthesized protein accumulating in granular structures (Fig. 5D, FOV2 and 3).

To further study the intracellular transport of nsP3-containing granules, we used instant structured illumination microscopy (iSIM) for live-cell recordings at high frame rates (39). iSIM increases spatial resolution by a factor of \( \sqrt{2} \) compared with wide-field microscopy and by a further factor of \( \sqrt{2} \) with postprocessing, while rapid image capture provides the temporal resolution needed for dynamic events within cells. A variety of fast-moving nsP3-containing objects with linear displacements and intermittent bursts of speed were tracked in live-cell recordings (Fig. 6A; Videos S2 to S5). As can be seen in the videos, the movement of these nsP3-containing objects differed from...
FIG 3: Four-color microscopy of nsP3, dsRNA, nsP1, and ZsGreen. Stable CHIKV cells were fixed and probed for nsP3 (cyan), dsRNA (magenta), nsP1 (nsP1), and ZsGreen (green) by a combination of SNAP tag labeling and indirect immunofluorescence assays. Images were taken with an Airyscan microscope operated in the superresolution mode. Overlay images are a combination of the nsP3, nsP1, and dsRNA layers as indicated. The zoomed-out ZsGreen channel is shown as a separate reference, with the corresponding ROI marked by a white box. Arrowheads indicate regions of proximity between nsP3, dsRNA, and nsP1.
FIG 4 Association of nsP3-containing granules with nuclear membrane and Nup98. (A) Fixed, stable CHIKV cells were examined for the presence of nsP3 (cyan) and G3BP2 (magenta) at the nucleus (stained with DAPI; yellow). As shown in Fig. 2, both G3BP1 and G3BP2 colocalize with nsP3. Here we probed for G3BP2, since immunostaining had a higher SNR than G3BP1 staining. ROIs 1 to 4 are high-magnification views of granules associating with nuclei. (B) Quantification (Continued on next page)
Static objects containing nsP3, which were imaged in the same recording. Fast-moving granules containing nsP3 could reach speeds of 0.8 to 5.9 μm/s (Table 2) with net displacements of 9.77 and 10.58 μm measured in the longest tracks (Table 2, tracks 1 and 4). Next, to complement our iSIM live-cell imaging analysis, we took advantage of total internal reflection fluorescence (TIRF) microscopy, which images only fluorescent molecules located close to the glass/specimen interface. TIRF microscopy produced images with good signal-to-noise ratio and reduced the background fluorescence from out-of-focus planes, enabling the observation of nsP3-containing granules located in or close to the plasma membrane (Fig. 6B; Video S6). Tracks analyzed in iSIM and TIRF images revealed similar object displacements and peak velocities (Table 2). Prior to TIRF imaging, we also labeled the plasma membrane with the CellMask orange plasma membrane stain and captured the images at least 1 h after staining to ensure that some of the dye was internalized in membrane-containing vesicles. Multicolor TIRF microscopy then allowed us to visualize the cotrafficking of nsP3- and CellMask-containing structures (Fig. 6B, Zoom, ROI 1 and 2). In summary, the dynamic analysis of nsP3-containing granules showed that they (i) could persist in cells for days, (ii) accumulated newly synthesized protein, (iii) could be classified into static and motile subclasses with characteristic displacements and speeds, and (iv) cotrafficked with membrane-containing structures.

**Static internal architecture of nsP3-containing granules during persistent replication.** To determine the dynamic behavior of nsP3 in granules, we performed fluorescence recovery after photobleaching (FRAP) experiments. Stable CHIKV cells were labeled with BG–6-carboxytetramethylrhodamine (TMR)–Star, defined regions with diameters of about 0.8 μm were photobleached, and fluorescent recovery was measured within each region of interest (Fig. 7A). No fluorescence recovery occurred over the duration of the experiment (Fig. 7A and B), suggesting that nsP3 remained fixed within the granular architecture and did not undergo dynamic exchange within each granule or with the surrounding cytoplasm. In contrast, SGs formed during overexpression of an enhanced green fluorescent protein (EGFP)-G3BP1 fusion in uninfected HuH-7 cells showed a characteristic rapid recovery of fluorescence after the photobleaching (Fig. 7B, inset), consistent with G3BP1 rapidly shuttling into and out of SGs.

Previously, G3BP1-containing SGs were shown to be stable in cell lysates, suggesting that stable core structures make up these membrane-less organelles (30). To further test the stability of nsP3-containing granules, we microscopically examined lysates of stable CHIKV cells (Fig. 8A). Bright-field images of cell lysates indicated the presence of refractive granules, while fluorescence microscopy identified granules that had incorporated the BG-TMR-Star label (Fig. 8B). We also confirmed that the stability of nsP3-containing granules was not unique to stable CHIKV cells, as we could also detect ZsGreen-positive granules following lysis of cells infected with CHIKVZsGreen-P3 (Fig. 8C).

In stable CHIKV cells, ZsGreen serves as a fluorescent marker of subgenomic replicon-RNA synthesis (Fig. 1A, cartoon). The green fluorescence also allowed us to use fluorescence-activated cell sorting (FACS) to eliminate any ZsGreen-negative cells. Although stable CHIKV cells maintained high levels of SNAP-nsP3 and ZsGreen for up to 2 months, cells with reduced or undetectable ZsGreen fluorescence accumulated in the absence of puromycin selection after only 1 week of culturing (Fig. 8D and E). These cells were sensitive to puromycin (Fig. 8D and F), suggesting that they no longer harbored the replicon. To test whether ZsGreen-negative cells that had emerged during...
FIG 5 Long-term imaging of SNAP-nsP3 using pulse-chase and quench-pulse-chase approaches. (A) Schematic drawing of pulse-chase image and quench-pulse-chase approaches. “Pulse” refers to labeling with a SNAP-specific dye; “chase” refers to incubation in standard culture medium free of SNAP-specific dyes. (B) Stable CHIKV cells were plated in glass-bottom dishes. Labeling with BG-647-SiR was carried out the following day. Cells were imaged with a Nikon Ti2-E inverted microscope equipped with a light-emitting diode (LED) light source, a 60× oil 1.4-NA objective, and a
culturing in puromycin-free medium regained some of the characteristics of uninfected cells, we subjected a mixed population (with both ZsGreen-positive and -negative cells) to a restress experiment with sodium arsenite. Consistent with previous experiments that examined the effects of “restressing” alphavirus-infected cells (17, 40), ZsGreen-positive cells sequestered G3BP2 into nsP3-containing granules even in the absence of sodium arsenite (Fig. 8G, FOV1), and new G3BP2-containing granules were absent after arsenite-induced stress (Fig. 8G, FOV3). In contrast, cells lacking ZsGreen did not have any G3BP2-containing granules in the absence of arsenite stress (Fig. 8G, FOV2) but were able to form G3BP2-positive clusters after arsenite treatment (Fig. 8G, FOV4). Therefore, the renewed ability to respond to arsenite-induced stress was associated with a loss of viral replication and nsP3-containing granules.

DISCUSSION

The objectives of this study were, first, to characterize the interaction between CHIKV nsP3 and cellular components during persistent replication and, second, to evaluate the persistence of cytoplasmic granules composed of viral and cellular proteins. To achieve these objectives, we expanded the utility of a noncytotoxic replicon by combining it with SNAP tag-based fluorescent labeling and subdiffraction multicolor microscopy to provide unprecedented insights into the substructure of persistent nsP3-G3BP-containing granules. These studies revealed their relationship with dsrRNA, nsP1-positive structures, and the nuclear membrane. Examining the dynamics of nsP3-containing granules uncovered a stable population of nsP3-containing granules along with a subclass of nsP3-positive structures trafficking through the cell cytoplasm. Importantly, we observed that nsP3-containing granules lacked a dynamic internal architecture and remained stable in cell lysates. Lastly, we showed that the ability to respond to oxidative stress was associated with the loss of CHIKV replication and nsP3-containing granules.

Stable CHIKV cells as a versatile tool for studying cytoplasmic nsP3-containing granules. Previous reports on noncytotoxic Old World alphaviruses elucidated the relationship between cytotoxicity, nsP2, and viral genome replication (28, 41–43). The typical cytotoxicity of CHIKV replicons precluded long-term studies of a previously described SNAP-tagged replicon. We have now overcome this limitation with a new HuH-7 cell line that harbors replicating CHIKV replicon RNA and encodes both SNAP-tagged nsP3 and ZsGreen as a genetic reporter for subgenomic replicon RNA. Whether this replicon establishes persistent replication only in specific cell types, as has been observed for other noncytotoxic replicons (28, 41), remains to be determined. We found that the SNAP-tagged replicon also persisted in C2C12 mouse myoblasts, albeit less efficiently.

To our knowledge, the system presented here is the first to allow intracellular tracking of nsP3 during persistent replication of CHIKV RNA in a replicon system. A similar accumulation of nsP3 in cytoplasmic granules occurs in transient replicons (16, 26, 27) and during late stages of infection (24, 25, 27). Strikingly, SNAP-nsP3 in stable CHIKV cells did not form rod-like structures, which were observed in cells infected with CHIKVNSAP-P3. Rod-like structures appear not only during transient replication in HuH-7 cells (26) but also in other cell types (mouse myoblasts, glial cells, dermal fibroblasts) during transient replication and late stages of infection (25, 27). Moreover, mutagenesis

FIG 5 Legend (Continued)

heated stage insert (set to 37°C with 5% CO2). Z-stacks of the same position were taken every 30 min for a total of 16 h. The entire time-lapse recording is also provided in Video S1 in the supplemental material. (C) Cells were prepared as described for panel B but imaged with an Airyscan microscope operated in the Fast Airyscan mode after the final wash (0 h), after 24 h (day 1), and after 48 h (day 2). Cell dishes were returned to a heated incubator after separate FOVs were imaged at each time point. (D) Quench-pulse-chase experiment. Stable CHIKV cells were plated in 24-well plates containing glass coverslips. The next day, nonfluorescent bromothenylpteridine was used to block the reactivity of intracellular SNAP-nsP3. Blocked cells were fixed with 4% formaldehyde at the indicated times postblock (0 h, 3 h, 6 h), and newly synthesized SNAP-nsP3 was stained with BG-647-SiR postfixation. Total nsP3 was stained with an indirect immunofluorescence assay using nsP3-specific antibodies. Stained samples were imaged with an LSM880 system operated in the Fast Airyscan mode. One representative field of view (FOV) is shown from each sample. The same laser power and detector settings were used to image each FOV. Z-stacks were acquired to capture all the granules present within cells. Images are maximum-intensity projections. The SNAP-nsP3 channel was pseudocolored with the Fire lookup table.
FIG 6 Live imaging of SNAP-nsP3 in stable CHIKV cells showing movement patterns of nsP3-containing granules. (A) Live imaging of SNAP-nsP3 in stable CHIKV cells by instant structured illumination microscopy (iSIM). Entire recordings are also included in Video S2 to S5 in the supplemental material. The two-dimensional (2-D) time-lapse series consisted of 100 to 200 (Continued on next page)
of nsP3’s C-terminal domain results in the formation of long rod-like structures (16, 44).
In contrast, CHIKV nsP3 preferentially forms granules in specific cell types, such as human muscle and epithelial cell lines (27). Surprisingly, infection with CHIKVZsGreen-P3 was not associated with the presence of rod-like structures. However, we cannot rule out the possibility that rod-like structures only form transiently and are no longer present at the observed time point.

Nonetheless, the lack of rods was not accompanied by a reduction in infectious titers. Thus, our results suggest that the ability to form rod-like structures can be affected by the sequence of the inserted tag in the C-terminal domain (SNAP versus ZsGreen) but also by whether nsP3 is expressed during persistent replication or infection. Interestingly, the noncytotoxic replicon also encodes a leucine residue instead of isoleucine at position 175, in a presumed unstructured region between predicted domains of nsP3 (28). Although this mutation may primarily stabilize replication complexes in conjunction with other noncytotoxic mutations (28), we do not know yet whether it affects the formation of rod-like structures. Taken together, SNAP-nsP3 can form a cytoplasmic mixture of rod-like and granular structures during CHIKV infection, but only granules persist in cells that persistently replicate CHIKV replicon RNA.

Persistence of nsP3-G3BP-containing granules within a microenvironment containing dsRNA, nsP1, and cellular markers. Subdiffraction multicolor microscopy of

### TABLE 2 Quantification of tracks

| Microscopy type | Start time (s) | End time (s) | Duration (s) | Total displacement (μm) | Net displacement (μm) | Peak velocity (μm/s) |
|-----------------|---------------|-------------|--------------|-------------------------|----------------------|---------------------|
| ISIM            |               |             |             |                         |                      |                     |
| 1               | 0             | 8           | 8           | 14.15                   | 9.77                 | 4.7                 |
| 2               | 0             | 3           | 3           | 3.96                    | 1.67                 | 3.1                 |
| 3               | 0             | 13          | 13          | 7.82                    | 2.71                 | 1.4                 |
| 4               | 0             | 8           | 8           | 12.92                   | 10.58                | 5.9                 |
| 5               | 0             | 9           | 9           | 9.23                    | 2.23                 | 3.8                 |
| 6               | 0             | 9           | 9           | 8.80                    | 0.69                 | 3.3                 |
| 7               | 0             | 18          | 18          | 3.09                    | 2.1                  | 0.8                 |
| 8               | 0             | 14          | 14          | 9.96                    | 5.71                 | 2.8                 |
| 9               | 0             | 13          | 13          | 10.45                   | 1.37                 | 2.8                 |
| 10              | 0             | 7           | 7           | 5.9                     | 4.65                 | 2.3                 |
| TIRF            |               |             |             |                         |                      |                     |
| 0               | 0             | 84          | 84          | 23.76                   | 3.54                 | 1.2                 |
| 1               | 85            | 120         | 35          | 11.58                   | 10.23                | 1.5                 |
| 2               | 83            | 119         | 36          | 4.58                    | 3.62                 | 0.6                 |
| 3               | 81            | 110         | 30          | 14.07                   | 12.77                | 2.3                 |
| 4               | 77            | 88          | 12          | 11.05                   | 9.09                 | 1.8                 |
| 5               | 23            | 47          | 24          | 11.41                   | 2.33                 | 2.4                 |
| 6               | 0             | 120         | 120         | 42.83                   | 6.25                 | 2.5                 |
| 7               | 14            | 26          | 12          | 9.76                    | 6.78                 | 1.6                 |
| 8               | 103           | 120         | 17          | 4.23                    | 3.55                 | 0.8                 |
| 9               | 1             | 24          | 23          | 5.06                    | 3.22                 | 1.1                 |
| 10              | 5             | 120         | 115         | 27.12                   | 2.42                 | 0.8                 |

aISIM, instant structured illumination microscopy; TIRF, total internal reflection fluorescence microscopy.
bTotal displacement, the sum of all consecutive displacements in each track, which corresponds to the total distance traveled by the object.
cNet displacement, the distance between the starting and ending positions of each track.
dPeak velocity, highest velocity of the object over the duration of each track.

of nsP3’s C-terminal domain results in the formation of long rod-like structures (16, 44). In contrast, CHIKV nsP3 preferentially forms granules in specific cell types, such as human muscle and epithelial cell lines (27). Surprisingly, infection with CHIKVZsGreen-P3 was not associated with the presence of rod-like structures. However, we cannot rule out the possibility that rod-like structures only form transiently and are no longer present at the observed time point.

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Persistence of nsP3-G3BP-containing granules within a microenvironment containing dsRNA, nsP1, and cellular markers. Subdiffraction multicolor microscopy of

**FIG 6 Legend (Continued)**
frames. The original 2-D time-lapse series consisted of 100 to 200 frames but were cropped to the relevant frames in zoomed-in views. Images were acquired at intervals of 88 ms. The relative axial position of each ZsGreen-positive cell is indicated as top, bottom, or middle of the cell. The statistics for tracks 1 to 10 are listed in Table 2. (B) Live imaging of SNAP-nsP3 in stable CHIKV cells by total internal reflection fluorescence (TIRF) microscopy. After labeling SNAP-nsP3 with BG-647-SiR (cyan), the plasma membrane was stained with CellMask dye (magenta). Live-cell images were acquired after at least 1 h to allow for internalization of the CellMask dye via membrane-containing vesicles. Entire recordings are also included in Video S6 in the supplemental material. The statistics for tracks 0 to 10 are listed in Table 2. Arrows serve as digital fiducial markers and point toward nsP3- and CellMask-containing structures showing cotrafficking.
stable cells revealed that nsP3-containing granules were (i) G3BP1 and G3BP2 positive, (ii) juxtaposed to dsRNA foci and nsP1-positive structures, (iii) associated with the nuclear membrane, and (iv) proximal to Nup98-positive organelles. Alphavirus nsP3 forms cytoplasmic granules with vertebrate G3BP1/2 and the mosquito homolog Rasputin (16, 17, 21, 24, 25, 45–47). The noncytotoxic replicon preserved this interaction in cytoplasmic granules whose diameters and protein contents varied. Moreover, Airyscan microscopy allowed us to address the internal substructure of larger granules (>1 μm), which had detectable differences in fluorescence intensity within the granule, suggesting intragranular variations in the density of nsP3. In the future, stochastic optical reconstruction microscopy (STORM), which can provide an even higher resolution to Airyscan microscopy, may be necessary to reveal the detailed substructure of smaller granules (<500 nm). For example, STORM revealed that G3BP-containing SGs had stable core structures with diameters of ~200 nm (30).

Multicolor Airyscan microscopy provided a convenient workflow to examine ZsGreen-expressing stable cells for interactions between nsP3, dsRNA, and nsP1. Alphavirus nsP1 can bind membranes (48, 49) and may use its membrane-binding domain to tether replication complexes to cellular membranes (50). During infection of the related Semliki Forest virus (SFV), nsP1 colocalizes with G3BPs in putative replication complexes (17). However, the nsP1-nsP3 and nsP1-G3BP associations could not be clearly detected during transient CHIKV replication and CHIKV infection (16, 25, 26). We were able to image a partial overlap of nsP1-positive structures with nsP3 granules in stable cells. Occasionally, nsP1 coated ring-like structures, which may represent virus-induced membranous organelles. Furthermore, we could detect dsRNA-positive foci in contact with nsP3-containing granules. Large cytoplasmic G3BP-nsP3 structures contain viral genomic RNA but not dsRNA, and these complexes grow over the course of an infection (24). During SFV infection, replication complexes initially form at the plasma membrane within so-called spherules, which have a characteristic bulb shape and a diameter of about 50 nm. Later, spherules are internalized and incorporated in large intracellular cytopathic vacuoles, which are derived from endolysosomal membranes and about 0.6 to 2 μm in diameter (51–54). Replication complexes are comprised not only of the complementary negative strand but also the full-length positive strand and the subgenomic mRNA. However, negative-strand synthesis occurs only during the first few hours of SFV infection (55, 56). During Sindbis virus (SINV) infection, dsRNA intermediates are packed into membrane spherules at the plasma membrane and also
FIG 8 (A) Cell lysates from stable CHIKV cells. Live cells were stained with BG-TMR-Star and lysed with Glasgow lysis buffer. The lysate was then bound to plastic chamber slides overnight and imaged the following day. Images were acquired with an LSM880 microscope operated in Fast Airyscan mode. Cell lysates from uninfected, naive HuH-7 cells are shown as a control. (B) Zoomed-in views of the sample shown in panel A, providing higher-magnification views of nsP3-containing granules and corresponding bright-field images (transmission). (C) Cell lysates from HuH-7 cells infected with CHIKV-ZsGreen-P3. Samples were prepared as described for panel A and imaged in the green channel. (D) Schematic overview of different populations (P1 to P4) obtained during culture of stable CHIKV cells. (E) Wide-field microscopy of stable CHIKV cells passaged for 1 week in the absence of puromycin (P2). Naive HuH-7 cells and cells treated for 1 week with puromycin (P1) are shown as controls. Images were obtained with an IncuCyte Zoom live-cell imaging system. (F) Effect of puromycin treatment on mixed populations containing both ZsGreen-positive and ZsGreen-negative cells. Confluence of the two populations was determined from images taken with an IncuCyte Zoom live-cell imaging system. (G) Effect of sodium arsenite treatment on mixed populations containing both ZsGreen-positive and ZsGreen-negative cells. To induce cellular stress granules, sodium arsenite was added for at least 30 min. Cells were fixed and then stained for SNAP-nsP3 (cyan) and G3BP2. (Continued on next page)
contain nsP1 early during infection (2 h), whereas nsP1-nsP3-dsRNA cytoplasmic complexes appear later in infection (36). Thus, nsP3 structures that are associated with dsRNA and ring-like structures of nsP1 in this study may be related to cytopathic vacuoles. The fraction of nsP3 associated with active RNA replication complexes varies between alphaviruses (reviewed in reference 57), and a smaller fraction of nsP3-containing structures colocalizes with dsRNA in CHIKV-infected cells than in SFV-infected cells, where most of the staining colocalizes. Moreover, replication complexes are efficiently internalized from the cell periphery for SFV but not CHIKV, and this reduction in CHIKV replication complex internalization correlates with a reduced stimulation of the prosurvival PI3K-Akt-mTOR pathway in comparison to that in SFV infection. Hence, persistent replication of the noncytotoxic replicon may also be associated with an increased internalization of membrane-bound replication complexes and increased colocalization of large, cytoplasmic nsP3-containing granules with dsRNA. Ultimately, correlative light and electron microscopy (CLEM) of stable CHIKV cells can elucidate the ultrastructure of nsP3-containing granules and their relationship with membranous organelles, as was done for SFV (58). Stable CHIKV cells offer particular advantages during CLEM sample preparation: (i) tetramethylrhodamine-coupled SNAP ligands are compatible with CLEM approaches (59), (ii) every puromycin-selected cell is guaranteed to harbor the replicon, and (iii) ZsGreen fluorescence marks the cytoplasm of imaged cells.

We also captured high-resolution images of an association between nsP3-containing granules and the nuclear membrane. Moreover, we investigated the previously unexplored relationship between nsP3 and the nucleoporin Nup98. Little is known about the nuclear transport of nsP3, while the localization of nsP2 to the nucleus is well documented (16, 46, 60, 61). Intriguingly, a role for G3BP1 as a nuclear transport factor has been proposed, and SINV nsP3 has been identified at the nuclear membrane (21). Our results imply that nsP3-containing granules are associated with a nucleoporin during persistent replication and may connect to RNA transport pathways at the nuclear membrane. Viral proteins that bind to Nups or RNA transport factors have been shown to stimulate remodeling of the nuclear membrane and affect the nuclear transport of cellular mRNA and proteins (62, 63). During SFV infection, many nuclear proteins relocate to the cytoplasm, where they play both proviral and antiviral roles (64). We also observed an association of nsP3 granules with cytoplasmic Nup98. During hepatitis C virus (HCV) infection, cytoplasmic nucleoporins accumulate at sites rich in viral proteins, including virus-induced membranous organelles and cytosolic lipid droplets (65, 66). In summary, Nups may play a role in persistent replication of CHIKV, which could hijack the physiological functions of nucleoporins to transport CHIKV nonstructural protein components, mRNA, viral RNA, or cellular proteins. Our data warrant a further investigation of this hypothesis.

**Stable CHIKV cells contain a mixture of static and dynamic nsP3-containing granules, which lack a dynamic internal architecture and are stable in cell lysates.** Self-labeling enzyme tags such as the SNAP tag provide experimental control over the time of labeling, thereby allowing us to study protein turnover. nsP3-containing granules were stable for hours and persisted for days. Granules were also the site where newly synthesized nsP3 accumulated. Thus, old and new populations of nsP3 may continuously mix within cytoplasmic granules, as was seen during transient replication (26). Live-cell microscopy also provided the first real-time tracking of CHIKV nsP3-containing granules and in-depth view of granule dynamics in mammalian cells. Previous live-cell microscopy revealed three subclasses of nsP3 structures during SFV infection: (i) small, nonacidic, nsP3-positive vesicles undergoing multidirectional and short-distance (2–μm) movement reminiscent of actin-based movement, (ii) large,
acidic vesicles displaying less-frequent jumps over distances of >10 μm, and (iii) large, acidic vesicles that were immobile and concentrated in the perinuclear area (51). Blebbistatin, an inhibitor of the actin motor protein myosin II, inhibited the dynamic movements of small vesicles, while nocodazole, a tubulin-disrupting agent, inhibited saltatory movements (51). We report similar movement patterns, including (i) the presence of immobile granules within perinuclear regions and (ii) granules moving over short (1- to 3-μm) and long (>4-μm) distances at maximum speeds between 0.8 and 5.9 μm/s. We also visualized cotrafficking of nsP3- and membrane-containing structures, which suggests that nsP3 moves through the cell by hijacking components of the cellular secretory machinery. In the future, stable CHIKV cells can provide invaluable real-time insight into interactions between CHIKV and the host through multicolor imaging of ZsGreen, far-red-fluorescent SNAP-nsP3 labeling, and a third, blue or red, fluorescent marker.

FRAP experiments revealed the static internal architecture of nsP3-containing granules, whereas arsenite-induced G3BP-containing granules had a fluorescence recovery similar to that seen in human osteosarcoma cells (67). The absence of a rapid exchange in CHIKV-induced granules implies that nsP3 may play a role that differs biochemically from the dynamic role of G3BP1 in SGs (30, 31, 68). For example, nsP3 may create a scaffold similar to the one formed by Fas-activated serine/threonine kinase (FASTK) in SGs (68). Although we cannot rule out that G3BP1 or G3BP2 shuttles in and out of nsP3-containing granules, we predict that G3BP1/2 would be similarly fixed in granules: nsP3 completely overlapped G3BP, and nsP3-containing granules were stable enough to be preserved in cell lysates. Moreover, previous studies demonstrated that alphavirus nsP3-G3BP-containing granules lack canonical SG markers (16, 17) and remain stable during cycloheximide treatment (16), which dissolves SGs (69). FRAP experiments of membrane-associated foci containing nonstructural proteins of another RNA virus, HCV, also found a limited exchange between clusters of nonstructural proteins and the periphery (70–72). Thus, some of the nsP3 structures may represent cytopathic vacuoles, in which nsP3 has a limited exchange with the surrounding cytoplasm.

Unlike cytopathic vacuoles, which would be sensitive to detergents, a population of nsP3-containing granules was detergent resistant and stable in cell lysates. This persistence in lysates mimics that of mammalian SG cores, where a dynamic shell around core structures gives SGs biochemical qualities akin to liquid-liquid phase separations (30). We propose that nsP3-G3BP-containing granules lack canonical SG markers (16, 17) and remain stable during cycloheximide treatment (16), which dissolves SGs (69). Recent studies show that environmental conditions can cause proteins bearing intrinsically disordered protein regions to undergo liquid-liquid phase separation and assemble droplets, hydrogels, and aggregates; this concentration of proteins into discrete subcellular domains appears to be essential for cellular metabolism and stress responses (for a recent review, see reference 73). In turn, defects in the regulation of such membrane-less organelles could impair cellular functions, alter stress responses, and form the basis of pathogenic inclusions linked with neurodegenerative disease (74). Intriguingly, the C terminus of alphavirus nsP3 itself is unstructured, which is a prerequisite not only for proteins to undergo phase separation but also to form more solid gel-like granules. Moreover, viral genomic RNA colocalizes with nsP3-G3BP (24), providing evidence that nsP3-G3BP-containing granules are made up not only of protein but also of RNA, a key component of cellular ribonucleoprotein granules (for a review on ribonucleoprotein granules, see reference 75). The link between liquid-liquid phase separation, membrane-less organelles, stress responses, and toxic protein clusters forms the basis of a new hypothesis that nsP3-containing granules can perturb cellular responses to environmental conditions. However, more experiments are needed to (i) further characterize persistent nsP3-containing granules biochemically, (ii) identify other cellular or viral proteins within granules, and (iii) induce granular disassembly. Clearing cells of these stable cytoplasmic complexes could be essential for preventing any toxicity that emerges during prolonged exposure to CHIKV proteins. Moreover, directly targeting persistent nsP3-containing granules could lead to new approaches to combat chikungunya virus infections. Cells that had turned ZsGreen

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natural during culturing in puromycin-free medium were not irreversibly perturbed but rather had regained the ability to form SGs in response to arsenite treatment.

In summary, our results present the first evidence that granules containing the viral protein nsP3 and cellular protein G3BP persist in human cells with autonomously replicating CHIKV replicon RNA. Generation of a cell line harboring a persistently replicating SNAP-tagged replicon and advances in microscopy technology allowed us to reveal interactions between SNAP/nsP3, viral components (nsP1, dsRNA), and the nuclear membrane. Overall, nsP3-containing granules were stable, differed in their mobility, lacked a dynamic internal architecture, and were stable in cell lysates. These findings may also have clinical relevance, as CHIKV can cause chronic infection and persist in various cell types, such as macrophages, muscle, and liver cells. However, whether prolonged exposure to nsP3-containing granules causes pathogenic changes within the cell and can contribute to chronic chikungunya disease remains to be determined. Lastly, the reagent presented in this study adds a new dimension for future explorations of host-pathogen interactions, in particular as they relate to nsP3, and for the search for inhibitors that specifically target nsP3.

MATERIALS AND METHODS

CHIKV constructs. The replicon CHIKVRepRLuc-FL-SA-PG-IL was described previously and allows for stable, noncytotoxic growth in HuH-7 cells (28). It contains a cassette encoding a puromycin-N-acetyltransferase (Pac-FMDV 2A autoprotease-ZsGreen fusion under the control of the subgenomic promoter. In the CHIKVRepLuc-FL-Sg-A-PG-IL replicon, a Renilla luciferase (Rluc) flanked by SpeI restriction sites was inserted into nsP3. The SNAP-tagged replicon, which has a SNAP sequence (also flanked by SpeI restriction sites) inserted into nsP3, has also been described previously (26). The parental replicon used in the generation of the SNAP-tagged replicon was originally assembled from DNA constructs containing the CHIKV replicon cDNA from the LR2006 OPY1 strain, which was isolated from the serum of a febrile patient traveling from La Réunion (76); cDNA fragments (Geneart) were synthesized based on the published sequence of the LR2006 OPY1 strain and assembled in vitro to generate fully synthetic replicons. To generate a noncytotoxic SNAP-tagged replicon (CHIKVRepSnap), we ligated a DNA fragment corresponding to the region encoding the SNAP tag (excised by SpeI digestion of SNAP-tagged nsP3) to SpeI-digested CHIKVRepLuc-FL-SA-PG-IL vector.

Restriction site cloning via SpeI was used to replace a gene encoding the green fluorescent ZsGreen protein (originally derived from an Anthozoa species of reef corals (77)) with the SNAP sequence, in the context of an infectious CHIKV virus (CHIKVZsGreen-P3). This infectious clone was synthesized previously based on the sequence from CHIKV LR2006 OPY1 (43).

CHIKV constructs were verified by DNA sequencing of nsP3 regions (to confirm correct orientation of SNAP tag after ligation at SpeI sites) and the subgenomic region, as well as analysis of EcoRI/BamHI restriction digest patterns to test for the overall integrity of CHIKV replicons and infectious constructs.

Cells, media, transfection, and infection. HuH-7 cells were maintained in complete medium (Dulbecco’s modified Eagle’s medium supplemented with fetal calf serum, penicillin, streptomycin, nonessential amino acids, and HEPES buffer) as described previously (28). HuH-7 is a well-differentiated hepatocyte-derived cellular carcinoma cell line taken from the liver tumor of a male Japanese patient in 1982 (78); these cells were from John McLauchlan (Centre for Virus Research, Glasgow). Growth medium supplemented with puromycin (final concentration, 5 μg/ml) was used for antibiotic selection.

In vitro transcription and electroporation of CHIKV RNA. Plasmids containing cDNA of SNAP-tagged noncytotoxic CHIKV replicon were linearized by NotI digestion. Purified DNA was used as the template for an in vitro transcription reaction using the mMESSAGE mMACHINE SP6 transcription kit (Ambion). RNA was purified with the PureLink RNA minikit (Thermo Fisher Scientific) and stored in aliquots of distilled water at −80°C until the day of electroporation. RNA was transfected into cells via electroporation as described before (26). Electroporated cells were seeded in 10-cm dishes. Cells were incubated in puromycin-free medium for a minimum of 2 days before starting puromycin selection. During puromycin selection, cells were monitored with a wide-field fluorescence microscope and a fluorescein isothiocyanate (FITC) filter setup for ZsGreen fluorescence. After ZsGreen-positive cells reached a high proportion (2 to 5 days), cells were expanded in puromycin-free medium. Heterogeneous populations of ZsGreen-positive cells, which we call stable CHIKV cells, were collected from confluent T75 flasks to make frozen cell stocks in fetal calf serum supplemented with 10% dimethyl sulfoxide (DMSO) (about 2 weeks after electroporation). At the same time, stable CHIKV cells were passaged under standard cell culture conditions and used in microscopy experiments. To study the appearance of a subpopulation of ZsGreen-negative cells, a pure population of ZsGreen-positive cells was obtained with fluorescence-activated cell sorting (FACS) of live cells. Cell populations were sorted with a FACS Melody instrument (BD Biosciences) based on green fluorescence (488-nm laser and 527/32 filter). Only singlets were picked to avoid artificially high fluorescence. Following a cell sort of 1 million cells, cells were plated into a T25 flask and allowed to expand in puromycin-free medium to allow for the appearance of ZsGreen-negative cells.

For infection experiments, plasmids containing cDNA of CHIKV SNAP-P3 and CHIKVZsGreen-P3 were linearized by NotI digestion. In vitro transcription was carried out as described above. HuH-7 cells were harvested from T175 flasks, electroporated at 0.5 × 10⁷ cells/ml using a square-wave protocol at 260 V...
for 25 ms, seeded into T175 flasks, and allowed to incubate for multiple days. Supernatants were frozen and used as virus stocks. The working stock of CHIKV was plaque titrated in BHK-21 cells (ATCC CCL10). For microscopy analysis of viral infection, this working stock was added to naive HuH-7 cells at a multiplicity of infection (MOI) of 10 and fixed 24 h later. To compare the multiplication of ZsGreen- and SNAP-tagged virus, HuH-7 cells were infected with viral stocks at the same MOI; supernatants were collected at 24, 48, and 72 h postinfection and plaque titrated in BHK-21 cells.

**Primary and secondary antibodies.** Polyclonal anti-G3BP2 was obtained from Bethyl Laboratories, mouse anti-G3BP1 antibody was from BD Biosciences, and antibodies detecting Nup98 were from Cell Signaling. For immunofluorescence labeling of dsRNA, mouse monoclonal anti-dsRNA (J2; Scicons) was used. J2 specifically recognizes dsRNA of more than 40 bp in length (79). Polyclonal rabbit antibodies against CHIKV nsP3 and nsP1 were produced in-house (Merits laboratory). Whole species-specific IgG secondary antibodies were either anti-rabbit Alexa Fluor 594-conjugated IgG (Thermo Fisher Scientific) (Fig. 2 and 4), anti-rabbit DyLight 405 IgG (to detect nsP1 in Fig. 3), or anti-mouse Alexa Fluor 594-conjugated IgG (to detect J2 in Fig. 3).

**Intracellular SNAP tag staining.** To stain intracellular SNAP-tagged proteins with the standard protocol, benzylguanine (BG), conjugated to fluorophores (silicon rhodamine [SiR], or TMR-Star, commercially available as SNAP-Cell 647-SiR and SNAP-Cell TMR-Star [NEB]), was added to live cells and incubated for at least 15 min at 37°C, 5% CO2. This was followed by three washes in complete medium and an extended incubation in complete medium for at least 30 min to remove background fluorescence. For fixed-cell microscopy analysis shown in Fig. 1, cells were fixed overnight at room temperature with 4% formaldehyde for 30 min. Cells were then counterstained with 4′,6-diamidino-2-phenylindole (DAPI) and mounted onto glass slides by the addition of ProLong diamond antifade mountant (Thermo Fisher Scientific).

**IFAs.** For indirect immunofluorescence assay (IFA) and staining with G3BP1, G3BP2, or J2 antibodies, formaldehyde-fixed cells were permeabilized with 100% methanol for 10 min at −20°C. For all other antibodies, cells were permeabilized with a buffer containing 5% fetal calf serum and 0.3% Triton X-100. Cells were incubated with primary antibody solution containing 1% bovine serum albumin (BSA) overnight at 4°C, except the mouse J2 antibody, which was incubated for 2 h at room temperature in diethylpyrocarbonate-treated phosphate-buffered saline (PBS). After three washes in PBS, secondary antibody (anti-rabbit Alexa Fluor 594-conjugated IgG or anti-mouse Alexa Fluor 594-conjugated IgG; Molecular Probes) was added. For nsP3/J2/nsP1 triple staining, rabbit nsP1 antibody was added overnight at 4°C to cells already stained with BG-647-SiR (benzylguanine-silicon-rhodamine) and mouse J2. The following day, cells were washed three times in PBS, and secondary antibody (anti-rabbit Alexa Fluor DyLight 405) was added. These cells were not counterstained with DAPI. However, where indicated (Fig. 1, 2, and 4), DAPI was added to visualize nuclei. Coverslips were mounted onto glass slides by the addition of ProLong diamond antifade mountant (Molecular Probes).

**Pulse-chase and quench-pulse-chase experiments.** For long-term pulse-chase experiments (Fig. 5B and C), stable CHIKV cells were plated in 35-mm glass-bottom dishes with a no. 1.5 grided coverslip (Nunc); labeling with BG-SiR was carried out the following day using the live-cell protocol described above. Live-cell imaging solution (supplemented with HEPES, 10% FBS, nonessential amino acids, and ProLong live antifade reagent) was added after the final wash. A Nikon Ti2-E inverted microscope was used to image the turnover of SNAP-nsP3 over 16 h. To image live cells after 24 h and 48 h chase periods, an LSM880 imaging system was operated in Fast Airyscan mode. Different fields of view were taken with the same imaging settings as those for the 0-h time point.

For quench-pulse-chase experiments, stable CHIKV cells were plated in 24-well plates containing 13-mm glass coverslips. The next day, 10 μM nonfluorescent bromothymylxylene (SNAP-Cell block; NEB) was used to block the reactivity of intracellular SNAP-nsP3 in stable CHIKV cells. After a 45-min incubation, blocked cells were washed three times with complete medium, followed by a 30-min incubation in complete medium. Cells were fixed with 4% formaldehyde at the indicated times (Fig. 5) postblock (0 h, 3 h, 6 h), and newly synthesized SNAP-nsP3 was stained with BG-SiR. Total nsP3 was stained with a rabbit antiserum against nsP3 and dye-conjugated secondary antibodies (anti-rabbit Alexa Fluor DyLight 405). Stained coverslips were mounted onto glass slides in ProLong diamond (Molecular Probes), and Z-stacks were acquired with an LSM880 system (Zeiss) operated in the Fast Airyscan mode. A Plan-Apochromat 63×/1.4 oil pH3 M27 objective was used for these experiments.

**Subdiffraction light microscopy.** An LSM880 upright confocal microscope with Airyscan (Zeiss) was used to acquire subdiffraction microscopy images as described previously (26, 80). This microscope provides a maximum lateral resolution of 140 nm and an axial resolution of 400 nm for a fluorophore emitting at 480 nm. Z-stacks were acquired with a 63×/1.4NA Plan-Apochromat oil objective at a step size of 0.16 μm. Pixel size was 40 nm by 40 nm by 160 nm. Sequential scans (scan zoom = 4 in frame mode, 1-s frame time, averaging set to 1 or 2) were acquired in four channels, as follows: channel 1 = 633 nm laser, channel 2 = 561 nm laser, channel 3 = 488 nm, channel 4 = 405 laser. Z-stacks in Fig. 2, SC, SD, 8A, 8B, and 8G and single-slice images of live cells (Fig. 2F) were acquired with the Fast Airyscan mode. To increase signal-to-noise ratio and resolution, image stacks were processed by Airyscan processing within Zen Black. Single-slice images were extracted to produce panels in Fig. 2, 3, and 4.

**Live-cell microscopy of stable CHIKV cells.** Live-cell wide-field imaging in Fig. 5B was done with a Nikon Ti2-E inverted microscope equipped with a Lumencor Spectra X LED light source, CFI Plan Apo Lambda 60× oil/1.4NA objective, a photometric Prime 95B sCMOS monochrome camera, and a heated stage insert (set to 37°C with 5% CO2). Z-stacks were taken every 30 min for a total of 16 h. Cells were grown in 35-mm glass (no. 1.5)-bottom dishes with a 27-mm viewing area (Nunc). Stable CHIKV cells were stained with BG-647-SiR and then maintained at 37°C in an optically clear, physiological, and CO2-
independent imaging buffer (Molecular Probes; live-cell imaging solution supplemented with 10% fetal calf serum, nonessential amino acids, and buffered with 10 mM HEPES). To suppress photobleaching, ProLong live antifade reagent was added according to the manufacturer’s instructions (Molecular Probes).

A home-built instant structured illumination microscope (iSIM) was used to acquire additional subdiffraction time-lapse series at high frame rates (one image every 88 ms) (Fig. 6A and B). This instrument is fitted with an Olympus water immersion objective 1.2-numerical aperture (NA) UPLSAPO 60xW and 488-nm and 561-nm lasers (81). Stable CHIKV cells were stained with red-fluorescent BG-TMR-Star before image acquisition. The heated stage was set to 37°C. The same live-cell imaging medium described above was used, supplemented with ProLong live antifade reagent. Regions of interest were found using the live iSIM display in the green channel (ZsGreen) to avoid bleaching of the red channel (nsP3). A single-slice two-color image of the green and red channels was taken as a reference. Two-color reference images were processed to remove striped scanning artifacts (39) with the stripes filter in ImageJ plugin Xlib (82). The Z-position corresponded to either the bottom, middle, or top of the cell. Time-lapse series were acquired by taking images of the red channel at intervals of 88 ms for 100 to 200 cycles. Cropped ROIs from these time-lapse series were processed using the Richardson-Lucy algorithm in the ImageJ plugin DeconvolutionLab (six iterations) (83) and a Gaussian filter (σ = 1 pixel).

Image contrast was adjusted for each cropped time-lapse series within the iCy (http://icy.bioimageanalysis.org) platform (84) by dragging the adjustable bounds of the histogram viewer, which enhances the contrast in the selected channel without altering the data (84). A viewing range that provided the best contrast for the moving objects within the time lapse was selected.

For TIRF (total internal reflection fluorescence) microscopy, a Ti2-E inverted microscope equipped with an LU-NA laser bed (405, 488, 561, 647), a CF-APo.NTars SR TIRF 100x: oil 1.5-NA objective, and a photometric Prime 95B sCMOS monochrome camera was used. Stable CHIKV cells were stained with BG-SiR. The plasma membrane was stained with red-fluorescent CellMask orange (Molecular Probes). Three-color images (green, red, far-red) of the ZsGreen, CellMask orange, and BG-SiR signals were taken. A single-slice two-color image of the green and red channels was taken as a reference. Two-color reference images were processed to remove striped scanning artifacts (39) with the stripes filter in ImageJ plugin Xlib (82). The Z-position corresponded to either the bottom, middle, or top of the cell. Time-lapse series were acquired by taking images of the red channel at intervals of 88 ms for 100 to 200 cycles. Cropped ROIs from these time-lapse series were processed using the Richardson-Lucy algorithm in the ImageJ plugin DeconvolutionLab (six iterations) (83) and a Gaussian filter (σ = 1 pixel).

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For FRAP analysis. Stable CHIKV cells, stained with BG-TMR-Star with the live-cell protocol, were used for experiments imaging SNAP-nsP3. An LSM700 imaging system (Zeiss) was used for FRAP experiments. Circular bleach areas were drawn within the Zen Black software (diameter of about 0.8 μm). Analyzed ROIs were pooled from recordings of 10 FOVs. One reference region of identical size was drawn over a granule and left unbleached. Another reference region was drawn within the cytoplasm to measure fluorescence background. Bleaching was set with the 405-nm, 488-nm, and 555-nm laser lines at 100% output. Bleaching was started after 3 frames, and another 97 frames were taken every 320 ms during the recovery period. Values of mean ROI intensities were extracted with Zen Black software, exported to Microsoft Excel, and graphed with GraphPad Prism. To induce genuine stress granules in HuH-7 cells, the plasmid pEgfp-G3BP (kindly provided by Richard Lloyd, Baylor University), encoding an EGFP-G3BP1 fusion protein (85), was transfected with Lipofectamine 2000 reagent (Thermo Fischer Scientific) in cells plated in a 35-mm glass (no. 1.5)-bottom dish with a 27-mm viewing area (Nunc). After 24 h, cells containing G3BP1 granules were identified by live-cell microscopy on an LSM700 confocal system set to 37°C.

Isolation of SNAP-nsP3 from cell lysates. Stable CHIKV cells grown in six-well plates were labeled with BG-TMR-Star according to the live-cell staining protocol outlined above. Cells were collected by scraping them into PBS using plastic cell scrapers, followed by centrifugation in 1.5-ml microcentrifuge tubes. Cell pellets were lysed with 300 μl ice-cold Glasgow lysis buffer (1% Triton X-100, 120 mM KC1, 30 mM NaCl, 5 mM MgCl2, 10% glycerol, and 10 mM piperazine-N,N’-bis(2-ethanesulfonic acid) (PIPS)-NaOH, pH 7.2) containing protease inhibitors. Lysates were vortexed for 30 s for four cycles and returned to ice between cycles. A final spin at 850 × g was included to remove the remaining cellular debris. The final supernatant was added to a two-well ibidi plastic slide with an ibitreat surface for optimal cell adhesion (Ibidi). After an overnight incubation at 4°C, 1 ml of 4% formaldehyde was added to each well for 1 h at room temperature. Wells were washed with PBS, and images were captured with an LSM880 system operated in Fast Aireyscan mode. The same protocol was used to analyze lysates from HuH-7 cells infected with CHIKVnsP3 (83).

Bioimage analysis. Data sets from Fig. 5B and 6B and from Video S1 in the supplemental material were processed with NIS-Elements AR imaging software (Nikon). Wide-field images were deconvolved within the Elements software according to the Richardson-Lucy algorithm (set to 10 iterations). All other
microscopy images and videos were processed on the Icy (http://icy.bioimageanalysis.org) platform (84). Contrast was optimized in individual images by dragging the adjustable bounds of the histogram viewer, which enhances the contrast in each channel without altering the data (84). Color maps (cyan, magenta, green, gray, yellow, fire, or jet) were applied with the lookup table manager to each channel in combination with the corresponding histogram bounds. The HK-Means and Active Contours plugins (A. Dufour, V. Meas-Yedid, A. Grassart, and J. C. Olivo-Marin, presented at the 19th International Conference on Pattern Recognition, 8 to 11 December 2008) were used for segmentation of nsP3-containing granules, and the ROI viewer provided information about the maximum Feret diameter of the granules. Tracking of nsP3-containing granules was done with the Manual Tracking (written by Alexandre Dufour) plugin, which allows clicking on the center of individual granules for every time point and integration with the software’s Track Manager (written by Fabrice de Chaumont) to visualize each track. Motion statistics (total displacement and relative/net displacement) for each track and peak velocity detected within an individual track were extracted from the Track Manager plugin (written by Fabrice de Chaumont) by adding the Motion Profiler track processor (written by Alexandre Dufour).

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at https://doi.org/10.1128/JVI.00477-18.

SUPPLEMENTAL FILE 1, MP4 file, 14.8 MB.
SUPPLEMENTAL FILE 2, MP4 file, 18.5 MB.
SUPPLEMENTAL FILE 3, MP4 file, 16.8 MB.
SUPPLEMENTAL FILE 4, MP4 file, 19.4 MB.
SUPPLEMENTAL FILE 5, MP4 file, 19.4 MB.
SUPPLEMENTAL FILE 6, MP4 file, 19.4 MB.
SUPPLEMENTAL FILE 7, PDF file, 0.1 MB.

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