Citrus psorosis virus movement protein contains an aspartic protease required for autocleavage and the formation of tubule-like structures at plasmodesmata

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Running title: MP\textsuperscript{CpV} forms tubule structures at PD upon autocleavage

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Abstract

Plant virus cell-to-cell movement is an essential step in viral infections. This process is facilitated by specific virus-encoded movement proteins (MPs), which manipulate the cell wall channels between neighboring cells known as plasmodesmata (PD). Citrus psorosis virus (CPsV) infection in sweet orange involves the formation of tubule-like structures within PD, suggesting that CPsV belongs to “tubule-forming” viruses that encode MPs able to assemble a hollow tubule extending between cells to allow virus movement. Consistent with this hypothesis, we show that the MP of CPsV (MP\textsubscript{CPsV}) indeed forms tubule-like structures at PD upon transient expression in Nicotiana benthamiana leaves. Tubule formation by MP\textsubscript{CPsV} depends on its cleavage capacity, mediated by a specific aspartic protease motif present in its primary sequence. A single aminoacid mutation in this motif abolishes MP\textsubscript{CPsV} cleavage, alters the subcellular localization of the protein and negatively affects its activity in facilitating virus movement. The amino terminal 34 kDa cleavage product (34K\textsubscript{CPsV}), but not the 20 kDa fragment (20K\textsubscript{CPsV}), supports virus movement. Moreover, similar to tubule-forming MPs of other viruses, the MP\textsubscript{CPsV} (and also the 34K\textsubscript{CPsV} cleavage product) can homo-oligomerize, interact with PD-Located Protein 1 (PDLP1) and assemble tubule-like structures at PD by a mechanism dependent on the secretory pathway. 20K\textsubscript{CPsV} retains the protease activity and is able to cleave a cleavage-deficient MP\textsubscript{CPsV} \textit{in trans}. Altogether, these results demonstrate that CPsV movement depends on the autolytic cleavage of MP\textsubscript{CPsV} by an aspartic protease activity, which removes the 20K\textsubscript{CPsV} protease and thereby releases the 34K\textsubscript{CPsV} protein for PDLP1-dependent tubule formation at PD.

Importance

Infection by citrus psorosis virus (CPsV) involves a self-cleaving aspartic protease activity within the viral movement protein (MP), which results in the production of two peptides termed 34K\textsubscript{CPsV} and 20K\textsubscript{CPsV} that carry the MP and viral protease activities, respectively. The underlying protease motif within the MP is also found in the MPs of other members of the \textit{Aspiriviridae} family suggesting that protease-mediated protein processing represents a conserved mechanism of protein expression in this virus family. The results also demonstrate that CPsV and potentially other ophioviruses move by a tubule-guided
mechanism. Although several viruses from different genera were shown to use this mechanism for cell-to-cell movement, our results also demonstrate that this mechanism is controlled by post-translational protein cleavage. Moreover, given that tubule formation and virus movement could be inhibited by a mutation in the protease motif, targeting the protease activity for inactivation could represent an important approach for ophiovirus control.
Introduction

The cell-to-cell spread of virus infection generally depends on the activity of one or more virus-encoded movement proteins (MPs), which target intercellular communication channels within the plant cell walls known as plasmodesmata (PD) (1, 2). These channels provide both membrane and cytoplasmic continuity between cells and thus function as intercellular conduits for both soluble and membrane-associated compounds, ranging from small molecules such as salts, hormones and metabolites, to macromolecules, such as proteins, short and long RNAs and protein:RNA complexes (3). Structurally, PD represent unique structures, in which the plasma membrane (PM) and the endoplasmic reticulum (ER) form two concentric membrane tubules extending through the cell wall and are in close opposition to each other (ca. 10 nm) (4). The ER tubule (the ‘desmotubule’) is linked from all its surface to the PM and, potentially, to the cell wall by proteinaceous spoke-like extensions, which may represent specialized protein tethers that regulate the distance between the two membranes and, thereby, the size of the cytoplasmic annulus available for cytoplasmic transport (5). However, while this model for the regulation of the PD size exclusion limit (SEL) remains speculative, numerous studies correlated the regulation of PD function in controlling intercellular communication with the degree of callose deposition in the cell wall surrounding the PD neck regions (6-8). Whereas the synthesis and accumulation of callose in the cell wall causes the closure of the cytoplasmic compartment by forcing the plasma membrane against the desmotubule, the degradation of callose by beta-glucanases opens this compartment for intercellular transport (5).

Plant viruses exploit PD for their cell-to-cell movement by co-opting the cellular machinery of PD-mediated transport. The cell-to-cell movement strategies used by viruses have been deeply reviewed (9, 10) and classified into two general mechanisms based on the type of alteration of PD structure. Whereas the tubule-guided mechanism involves the displacement of the desmotubule by the formation of a tubule-like transport structure assembled by multimerization of viral MP and the transport whole virions through the tubule(9, 11-16), the non-tubule-guided mechanism usually occurs in the absence of virions and does not involve structural alterations within PD but rather a viral interference with the normal regulation of PD. Tobacco mosaic virus (TMV), the type virus using a non-tubule guided mechanism, moves cell-to-cell independently of the coat protein (CP) (9, 17), by
targeting viral replication complexes (VRCs) to PD with the help of myosin motor proteins (18). Moreover, this virus interferes with callose depositions at PD, thus allowing the passage of the VRCs into the adjacent cell (19). The MP of this virus interacts with the ER, microtubules and microtubule-associated proteins proposed to play a role in the formation of movement-competent VRCs (20-22). There is evidence that the viral movement process also involves the severing of actin filaments (23) and interactions of the MP with synaptotagmin A (SYTA), which is proposed to act in endosomal recycling (24) as well as a membrane tethering protein (5, 10, 25, 26).

Unlike the MP of TMV and presumably the MPs of other viruses moving by the non-tubule-mediated mechanism, the MPs of tubule-forming viruses interact with members of the PD-Located Protein (PDLP) family and this interaction is required for tubule assembly and the spread of infection (27, 28). PDLPs require the ER-Golgi pathway for their targeting to PD (27, 29), which may explain the observed sensitivity of tubule-formation and tubule-mediated virus movement to secretory pathway inhibitors (27, 30, 31). DNA and RNA viruses that move cell-to-cell by a tubule-guided mechanism have been found in the Caulimo- (11, 27, 32, 33), Seco- (15, 27, 34-37), Bunya- (13, 38), and Bromoviridae (39) families.

Citrus psorosis virus (CPsV), the type member of the Aspiviridae family (formerly Ophioviridae), genus Ophiovirus (40, 41), is a three-partite, non-enveloped, negative-sense, single-stranded (ss) RNA virus. RNA 1 encodes a 280 kDa replicase (42) as well as a 24 kDa protein that affects miRNA maturation (43) and has RNA silencing-suppressing activity (44). RNA 2 encodes a protein of 54 kDa (named hereafter MP<sub>CPsV</sub>), which displays several features of a MP (45, 46) and has RNA silencing-suppressing activity as well (44). RNA 3 encodes a CP of 48 kDa (47). In addition to CPsV, the Ophiovirus genus contains six more members, Miraflori lettuce big-vein virus (MiLBVV), blueberry mosaic associated virus (BIMaV), lettuce ring necrosis virus (LRNV), freesia sneak virus (FSV), ranunculus white mottle virus (RWMV) and tulip mild mottle mosaic virus (TMMMV), of which the last two have not been completely sequenced and are less characterized.

Here we show that CPsV induces the formation of tubular structures at PD during infection in citrus. Similar structures protruding from PD are observed by confocal laser scanning microscopy (CLSM) upon ectopic expression of MP<sub>CPsV</sub> fused to fluorescent markers. The
formation of the tubules is dependent on the proper localization of PDLP at PD. We also show that the MP<sup>CPsV</sup> is autocatalytically processed during both infection and transient expression and that a conserved aspartic protease motif within MP<sup>CPsV</sup> is responsible for this activity. By further functional characterization, we demonstrate that the autocatalytic cleavage releases an N-terminal peptide fundamental for tubule formation and a C-terminal protein fragment with viral protease activity.

**Results**

**Cell walls of CPsV-infected citrus plants exhibit tubule-like structures**

To get insight into the cell-to-cell movement mechanism used by CPsV, we analyzed the PD structure in CPsV-infected sweet orange plants. Ultrathin sections of healthy and infected leaves harvested at similar developmental stages were sectioned and observed by Transmission electron microscopy (TEM). In contrast to healthy samples (FIG. 1Ai) the PD in the CPsV-infected sample contained a double line of electron-dense and well-organized proteinaceous material, compatible with a longitudinal sectioning through a hollow tubular structure that extended from the cell wall into the cytoplasm (FIG. 1Aii, arrow).

Given the capacity of ophiovirus MPs to target PD upon transient expression in *N. benthamiana* leaves (46), we wanted to know whether these proteins form tubule-like structures under these conditions. Expression of MP<sup>CPsV</sup> N-terminally (FIG. 1Bi, arrowhead) and C-terminally (FIG. 1Bii, arrows) fused to fluorescent proteins (*e.g.* enhanced green fluorescent protein, GFP) revealed that MP<sup>CPsV</sup> indeed can form tubules; however, only the C-terminal fusion (MP<sup>CPsV</sup>:GFP) showed tubule-like structures at PD (FIG. 1Bii, arrows), whereas N-terminal fusions (GFP:MP<sup>CPsV</sup>) labeled PD without forming such structures. For cells expressing C-terminal fusion protein, the number of tubules per cell was variable and each cell showed both tubular and non-tubular MP<sup>CPsV</sup>:GFP at PD (FIG. 1Bii). These observations indicate although both orientations target PD, only C-terminal fusions to GFP can form tubular structures.

**The MP<sup>CPsV</sup> interacts with PD-localized PDLP1 for tubule assembly**

Consistent with the tubule assembly from MP, the monomers of the tubule-forming MPs of caulimo- and nepoviruses showed MP-MP interaction *in vivo* (27). Moreover, these
proteins were shown to interact with members of the PDLP family at PD (27, 28). Amari et al. (27) showed that both the MP of grapevine fanleaf virus (GFLV) and the MP of cauliflower mosaic virus (CaMV) interact with PDLPs at PDs, and that the systemic movement of both viruses was inhibited in pdlp1 pdlp2 pdlp3 triple knock-out Arabidopsis thaliana mutants (27). The interaction with the PDLP family of proteins might be a common feature of the tubule-forming viruses. Consistently, also the tubule-forming MP of cowpea mosaic virus (CPMV) was shown to interact with PDLP1 in vivo (28). To determine whether the MP<sup>CPsV</sup> has the capacity to interact with PDLP1 we used fluorescence lifetime imaging microscopy (FLIM) to measure the degree of fluorescence resonance emission transfer (FRET) between the GFP and monomeric red fluorescent protein (RFP) moieties of MP<sup>CPsV</sup>:GFP and PDLP1:RFP expressed in N. benthamiana epidermal cells. When MP<sup>CPsV</sup>:GFP was expressed alone, the average fluorescence lifetime (τ) of GFP was 2.1 ± 0.1 ns (n = 65) (FIG. 2Ai and v). A similar τ value of 2.1 ± 0.08 ns (n = 59) was measured when this protein was co-expressed with PD callose-binding protein 1 fused to the red fluorescent protein Cherry (PDCB1:Cherry) (48) (FIG. 2Aii and v). When MP<sup>CPsV</sup>:GFP was co-expressed with an RFP-tagged version of the MP<sup>CPsV</sup> (MP<sup>CPsV</sup>:RFP), the τ value was 1.9 ± 0.1 ns (n = 30). This value is significantly different (P<0.01) from the τ value observed when MP<sup>CPsV</sup>:GFP was expressed alone and represents a FRET efficiency of 10% (FIG. 2Aiii and v), thus indicating the capacity of MP<sup>CPsV</sup> to oligomerize. Co-expression of MP<sup>CPsV</sup>:GFP with PDLP1:RFP resulted in a τ-value of 2.0 ± 0.1 ns (n = 78) (5% FRET), which also differed significantly (P<0.01) from the τ-value obtained when MP<sup>CPsV</sup>:GFP was expressed alone (FIG. 2Aiv and v). Given that the efficiency of FRET depends on intermolecular vicinity between GFP and RFP moieties, we wondered whether the FRET efficiency could be altered upon changing the position of the GFP fusion to MP<sup>CPsV</sup> from the C-terminus to the N-terminus. The average fluorescence lifetime (τ) of GFP was 2.47 ± 0.04 ns (n = 65) when the GFP:MP<sup>CPsV</sup> fusion protein was expressed alone (FIG. 2Bi and v). A similar τ value was measured when this protein was co-expressed with PD callose-binding protein 1 PDCB1:Cherry (48) (FIG. 2Bii and v). When GFP:MP<sup>CPsV</sup> was co-expressed with MP<sup>CPsV</sup>:RFP, the τ value was 2.1 ± 0.1 ns (n = 76). This value is significantly different (P<0.01) from the τ value observed when GFP:MP<sup>CPsV</sup> was expressed alone and represents a FRET efficiency of 15% (FIG. 2Biii and v), thus showing
again MP-MP oligomerization irrespective of the orientation of the GFP fusion to the protein. Co-expression of GFP:MP\textsuperscript{CPsV} with PDLP1:RFP resulted in a $\tau$-value of 1.9 ± 0.1 ns (n = 42) (23% FRET), which also differed significantly ($P<0.01$) from the $\tau$-value obtained when GFP:MP\textsuperscript{CPsV} was expressed alone (FIG. 2Biv and v). These results confirm that the MP\textsuperscript{CPsV} has the capacity to oligomerize and to interact with PDLP1.

PDLPs are targeted to PD via the ER-Golgi secretory pathway (29) through COPII vesicles which are formed at ER-exit sites (49). The vesicle formation is dependent on Sar 1, a Ras-like small GTPase. Consistently, the expression of a dominant-negative mutant of Sar1[H74L], inhibited COPII vesicle formation and thus PDLP targeting to PD (49, 50). To test if the inhibition of PDLP targeting to PD affects the tubule formation by MP\textsuperscript{CPsV} at PD, we transiently expressed PDLP1:RFP or MP\textsuperscript{CPsV}:RFP together with either Sar1[H74L]:GFP or with the wild type Sar1:GFP as control. Co-expression with the wild type Sar1:GFP showed no effect on the targeting of PDLP1:RFP to PD (FIG. 2Ci and ii) and the ability of MP\textsuperscript{CPsV} to assemble tubule-like structures at PD (FIG. 2Di, ii and Table 1). However, co-expression with Sar1[H74L]:GFP inhibited PDLP1:RFP accumulation of at PD (FIG. 2Ciii and iv) and as previously shown, it remains located at the ER (27, 29). In addition, although the MP\textsuperscript{CPsV}:RFP was detected at PD, as seen by callose co-staining with aniline blue, the ability of MP\textsuperscript{CPsV}:RFP to form tubule-like structures was inhibited (FIG. 2Diii, iv and Table 1). Thus, the ability of MP\textsuperscript{CPsV} to assemble tubules requires an intact ER-Golgi pathway.

**MP\textsuperscript{CPsV} GFP-fusion orientation and protein processing determines cell-to-cell movement activity**

Given that MP\textsuperscript{CPsV}:GFP, but not GFP:MP\textsuperscript{CPsV}, is capable of forming tubules at PD, we wondered whether this has functional relevance in virus movement. Since an infectious cDNA clone of CPsV to test this hypothesis by a reverse genetic approach is not available, we used an alternative assay to evaluate this activity (45, 46). In this assay we analyzed the capacity of MP\textsuperscript{CPsV}:GFP and GFP:MP\textsuperscript{CPsV} to trans-complement TMV\textDeltaMP\textDeltaCP-GFP, a previously reported movement-deficient TMV derivative (45). Highly diluted Agrobacterium cultures harboring the TMV\textDeltaMP\textDeltaCP-GFP-encoding plasmid were infiltrated together with cultures for the expression of either MP\textsuperscript{CPsV}:GFP, GFP:MP\textsuperscript{CPsV} or GFP as a negative control. The sizes of the highly GFP-fluorescent infection foci grown by
viral cell-to-cell movement from initially TMVΔMPΔCP-GFP inoculated cells were measured at 5 days post agroinfiltration (dpai). MpCPv-GFP- and GFP:MPCPv-expressing tissues exhibited larger foci than the GFP-expressing control tissues, indicating that MPCPv complements TMVΔMPΔCP-GFP cell-to-cell movement irrespective whether fused to GFP at the N- or C-terminus (FIG. 3A). Nevertheless, the infection foci complemented by GFP:MPCPv were significantly smaller (P<0.01) than those complemented by MPCPv:GFP (FIG. 3A).

Immunoblot analysis using antibodies against the fused fluorescent protein tags showed that both fusion proteins were expressed at a similar level (FIG. 3B), suggesting that the difference in complementation efficiency between GFP:MPCPv and MPCPv:GFP is not caused by a dose-dependent effect on movement activity. Surprisingly, expression of the C-terminal fusion protein (MPCPv-GFP) led to the production of a 79 kDa protein and a smaller GFP-containing protein of ≈48 kDa. Interestingly, expression of the N-terminal fusion protein (GFP:MPCPv) led to production of a GFP containing protein of ≈69 kDa in addition to the expected 82 kDa protein (FIG. 3B). In silico analysis with TargetP software (51) showed that MPCPv encodes a N-terminal chloroplast transit peptide (cTP) (FIG 3.C). Because these signals are cleaved upon chloroplast import, the 3 kDa mass difference observed between MBPCpV.GFP and GFP:MPCPv, could be a consequence of protein import at the chloroplast. In agreement with this hypothesis, we found that only the C-terminal fusion protein accumulates at chloroplasts (FIG. 3C).

Importantly, the expression profiles of the additional GFP-containing proteins are compatible with a post-translational cleavage event, in which two MPCPv peptides of 34kDa and 20 kDa (N- and C-terminal fragments, respectively) are produced.

The MPCPv contains a functional aspartic protease motif

The occurrence of MPCPv protein fragments was also observed in protein extracts of CPsV-infected Chenopodium quinoa plants using an antiserum against MPCPv. In these extracts a band of approximately 54 kDa corresponding to full length MPCPv, and two other bands of 34kDa and 20 kDa were seen (52). To further test these previous observations, we repeated the immunoblot analysis using protein extracts from the systemic host Nicotiana occidentalis P1 infected with CPsV. As shown in FIG. 4A, the MPCPv-
specific antibody indeed detected again the three distinct bands of 54 kDa, 34 kDa and 20 kDa, thus confirming the expression of three RNA 2-specific proteins during infection.

To address the origin of the observed MP\textsuperscript{CPsV} fragments, we analyzed the aminoacid sequence of the protein using the HHpred software package (53). We identified a region between aminoacids 331 and 413 of MP\textsuperscript{CPsV} showing aminoacid sequence similarity with the catalytic aspartic site of cathepsin D as well as retroviral proteases, such as the protease of HIV-2 (FIG. 4Bi). Consistently, also the MP of the ophioviruses MLBVV (Mp\textsubscript{MLBVV}) (protein id: AAN60448.1, region: 360-438) and BlMaV (Mp\textsubscript{BlMaV}) (protein id: AIF28243.1, region: 368-450) showed sequence similarity with retroviral proteases and cathepsin D (FIG. 4Bii and iii). The identified region in MP\textsuperscript{CPsV} contains an aspartic residue (D), which is also the first aminoacid of an aminoacid triad that is strictly conserved among ophioviruses MPs (45). This finding suggests that MP\textsuperscript{CPsV} and other ophiovirus MPs have an autocalytic protein cleavage activity.

To determine the location of the proteolytic cleavage site within the MP\textsuperscript{CPsV} aminoacid sequence, we aligned the MP\textsuperscript{CPsV} sequence with HIV-1 protease substrate peptides (54). We found that all these peptides aligned to the aminoacid sequence \textsuperscript{305}NLSNFLADQR\textsuperscript{314} of MP\textsuperscript{CPsV} (FIG. 4B), which is compatible with the location of a cleavage site expected to result in the formation of 34kDa and 20 kDa cleavage products. To identify the cleavage site, we expressed MP\textsuperscript{CPsV}:GFP and GFP (negative control) in \textit{N. benthamiana} and immunopurified these proteins with anti-GFP agarose beads followed by on-bead trypptic digestion and identification of the peptide by LC MS/MS. MS/MS spectra corresponding to peptides with only one end compatible with trypsin digestion and found in three independent MP\textsuperscript{CPsV}:GFP expressing samples and immunopurification experiments were analyzed. MS/MS spectra indicate the existence of peptides derived from the C-terminal end of the 34 kDa protein (FIG. 4 Ci) and another from the N-terminal end of the 20 kDa protein (FIG. 4Cii) compatible with a cleavage between the aminoacids \textsuperscript{310}LA\textsuperscript{311}.

To further prove that MP\textsuperscript{CPsV} contains an active aspartic protease motif responsible for the observed cleavage products, we mutated the respective sequence motifs within MP\textsuperscript{CPsV} and studied the effect of the mutations on the proteolytic processing of the protein. Site-directed mutagenesis was used to replace the predicted catalytic Asp residue (D340) by Ala (A) or Asn (N), and the mutant derivatives of MP\textsuperscript{CPsV} were named hereafter MP\textsuperscript{CPsV}D340A and
MPCpV:D340N, respectively (FIG. 4D). Additionally, we constructed GFP and RFP fusions to the 34KcPcV (N-terminal cleavage product) and 20KcPcV (C-terminal cleavage product) proteins (FIG. 4D). The same approach was used to construct protein with mutations at the cleavage site. Thus amino acids at 310LA311 were replaced by an Asp residue, thereby leading to two protein mutants named MPCpV:L310D and MPCpV:A311D. We also created MP mutants named MPCpV:A311R and MPCpV:A311H in which the Ala at the position 311 was replaced by Arg or His respectively. Upon transient expression in N. benthamiana, both MPCpV:D340A and MPCpV:D340N fused to RFP occurred with their expected sizes of 79 kDa (FIG. 4E). However, the 48 kDa cleavage product observed for MPCpV:RFP was not detected for these mutants, which indicates that the amino acid D340 is indeed critical for protein cleavage (FIG. 4E). As expected, expression of the fluorescent protein-fused 20KcPcV and 34KcPcV proteins were detected as immuno-reactive bands of 48 kDa and 62 kDa, respectively (FIG. 4E). The four mutant proteins MPCpV:L310D, MPCpV:A311D, MPCpV:A311R and MPCpV:A311H that carry mutations at the cleavage site showed a cleavage product as observed for the MPCpV upon expression in C-terminal fusion to RFP. Thus, none of the mutations introduced at the protein cleavage location prevented the aspartic protease motif to recognize and cleave the protein at this location (FIG. 4E).

**MPCpV self-cleavage determines subcellular localization**

Next, we used confocal fluorescence microscopy to determine whether the introduced mutations in the sites involved in catalytic cleavage of MPCpV affect the subcellular localization of the protein. MPCpV:GFP localized to PD, tubule-like structures at PD, nucleus, chloroplast (FIG. 3C), cytoplasm and microtubules, as previously described (52) (FIG. 5A). In comparison, expression of MPCpV:D340A:GFP led to a strongly reduced nuclear localization of the GFP signal suggesting that the nucleus may accumulate the MP cleavage products rather than the full-length protein. Instead, GFP signal was more strongly associated with chloroplasts. Importantly, MPCpV:D340A:GFP accumulated at PD but failed to form tubule-like structures (FIG. 5B). The same subcellular localization pattern was also obtained for MPCpV:D340N:GFP (FIG. 5C). Expression of 34KcPcV:GFP led to the localization of GFP fluorescence exclusively at chloroplasts and also at PD, where it formed tubular-like structures (FIG. 5D). In contrast, expression of 20KcPcV:GFP led to
diffuse GFP fluorescence in the cytoplasm and nucleus (FIG. 5E) and showed no other specific subcellular localization. The presence of tubule-like structures at PD in cells expressing either MP<sup>CPsV</sup>:GFP or 34K<sup>CPsV</sup>:GFP, and the absence of the tubule-like structures in cells expressing MP<sup>CPsV</sup>D340A:GFP or MP<sup>CPsV</sup>D340N:GFP indicates that the cleavage of MP<sup>CPsV</sup> by the aspartic protease motif is a prerequisite for tubule formation. Since the full-length MP<sup>CPsV</sup> is not able to form tubules, the fluorescent tubules observed in MP<sup>CPsV</sup>:GFP samples are likely formed by the unfused 34K<sup>CPsV</sup> cleavage product, which allow the incorporation of the full-length MP<sup>CPsV</sup>:GFP protein into these structures.

The 20K<sup>CPsV</sup> cleavage product is an aspartic viral protease

Since the protein cleavage mechanism used by aspartic proteases involves two catalytic triads (55), the 20K<sup>CPsV</sup> cleavage product that retains the aspartic protease activity should also retain a capacity to dimerize. To test this hypothesis, we measured the fluorescence lifetime of 20K<sup>CPsV</sup>:GFP upon co-expression with 20K<sup>CPsV</sup>:RFP. Under these conditions, the τ of 20K<sup>CPsV</sup>:GFP localized at the nucleus was 2.37 ±0.05 ns (n = 23) and a very similar fluorescence lifetime of 20K<sup>CPsV</sup>:GFP of 2.32 ±0.05 ns (n = 14) was measured when expressed together with RFP as a negative control (P<0.01). A significantly lower τ value was determined when 20K<sup>CPsV</sup>:GFP was co-expressed with 20K<sup>CPsV</sup>:RFP (2.16 ±0.08 ns; n = 19;P<0.01), representing a FRET efficiency of 9%. Similar fluorescence lifetime measurements were repeated by focusing on the proteins localized in the cytoplasm. Here, the 20K<sup>CPsV</sup>:GFP and 20K<sup>CPsV</sup>:RFP underwent FRET with an efficiency of 14% (FIG. 6A iii and v). These observations indicate that the cleaved20K<sup>CPsV</sup> peptide retains the ability to dimerize and thereby to form the dimeric catalytic triad proposed to be involved in its protease activity.

Next, we wondered whether the aspartic protease activity of MP<sup>CPsV</sup> can recognize and process its target in trans. Immunoblot analysis of extracts derived from agro-infiltrated <i>N. benthamiana</i> leaves co-expressing MP<sup>CPsV</sup>:RFP together with the proteolytic activity-deficient mutant MP<sup>CPsV</sup>D340A:GFP revealed that full length protein, and/or the 20K<sup>CPsV</sup>:RFP autocleavage product efficiently cleaves the MP<sup>CPsV</sup>D340A:GFP in trans (FIG. 6B; immunoblots on the left). The trans-acting proteolytic activity indeed resides in the cleaved 20K<sup>CPsV</sup> peptide, as was confirmed upon coexpression of MP<sup>CPsV</sup>D340A:GFP
with 20KCPVsRFP (Fig. 6B; immunoblots on the right). Consistent with MP340A:GFP cleavage by MP340A::RFP or 20KCPVsRFP, green fluorescent tubule-like structures were seen at PD (Fig. 6C). Thus, MP340A::RFP has the capacity to execute cleavage in trans as well as in cis and the protease activity segregates with the aspartic protease within the cleaved 20KCPVsRFP protein fragment upon cleavage.

**Cell–to-cell movement activity depends on the 34K protein**

The effect of the MP340A::RFP protease domain mutations on supporting viral movement was tested in our functional complementation assay using movement-deficient TMV (45, 52).

As shown in Fig. 7A, TMVΔCPΔMP-GFP movement was complemented in leaves transiently expressing MP340A::RFP as well as in leaves transiently expressing the MP of TMV (MP340A::RFP) as demonstrated by the occurrence of infection foci at 5 dpai under these conditions. Such efficient functional complementation TMVΔCPΔMP-GFP also occurred in the presence of 34KCPVsRFP but not by 20KCPVsRFP. MP340A::RFP and MP340N::RFP were also able to complement TMVΔCPΔMP-GFP; however, the average sizes of infection foci were smaller than those formed in the presence of MP340A::RFP or MP340A::RFP (Fig. 7B). Interestingly, foci formed in the presence of 34KCPVsRFP, which contains the 30K superfamily domain (45), were larger than the foci formed in the presence MP340A::RFP, although they were still smaller than those formed in the presence of MP340A::RFP (Fig. 7B and 7C). The observation that the expression of the 34KCPVs cleavage fragment is sufficient and even more efficient in complementing viral movement than the expression of the full length MP340A::RFP underscores the importance of efficient MP340A::RFP cleavage with MP340A::RFP function in supporting virus movement.

**34KCPVs behaves as a tubule forming MP**

The above-described results indicate that MP340A::RFP processing into 34 kDa and 20 kDa cleavage products is essential to enable the formation of tubule-like structures at PD and that this feature of MP340A::RFP contributes to the function as a MP. Given that the expression of the N-terminal 34KCPVs fragment, but not the expression of the C-terminal 20KCPVs fragment, leads to tubule formation at PD and is sufficient for complementing the cell-to-cell movement of a MP-deficient TMV construct, we wondered whether 34KCPVs has the
ability to interact with other 34K<sub>CpSv</sub> molecules as well as with PDLP1, as observed for
MP<sub>CpSv</sub>. FLIM measurements on transiently expressed 34K<sub>CpSv</sub>:GFP revealed an average
GFP fluorescence lifetime of 2.2 ns when this protein was expressed alone (FIG. 8Ai), and
a similar value was measured when this protein was co-expressed with PDCB1:Cherry
(Fig. 8Aii). However, when 34K<sub>CpSv</sub>:GFP was co-expressed with 34K<sub>CpSv</sub>:RFP, the
average fluorescence lifetime of GFP was reduced to 1.7 ns, thus revealing a FRET
efficiency of 23% (P<0.01). When 34K<sub>CpSv</sub>:GFP was co-expressed with PDLP1:RFP, the
average fluorescence lifetime of GFP was 2.0 ns (FIG. 8Ciii), which represents a FRET
efficiency of 9% (P<0.01). These data indicate that 34K<sub>CpSv</sub> has the capacity to
oligomerize and to interact with PDLP1 similar like the full-length MP<sub>CpSv</sub>.

Finally, we wanted to know whether the formation of tubules by 34K<sub>CpSv</sub> depends on the
ER-Golgi pathway and the targeting of PDLP1 to PD, as in the case of MP<sub>CpSv</sub>. Co-
expression of 34K<sub>CpSv</sub>:RFP with Sar1:GFP wt (FIG 8Ci, ii and v) allowed the formation of
34K<sub>CpSv</sub>:RFP tubules in 57% of the PD within the observed leaf area (Table 2). On the
contrary, when 34K<sub>CpSv</sub>:RFP was co-expressed with Sar1[H74L]:GFP, 34K<sub>CpSv</sub>:RFP
localized to PD but the percentage of PD with 34K<sub>CpSv</sub>:RFP tubules was significantly lower
(10%, P<0.01; Table 2). The PDLP1:RFP targeting to PD was again inhibited by co-
expression of Sar1[H74L]:GFP but not by co-expression of Sar1:GFP, as already shown.

Thus, inhibition of ER-Golgi pathway reduces PDLP1 targeting to PD and tubule formation
by 34K<sub>CpSv</sub> and MP<sub>CpSv</sub>. Both the MP<sub>CpSv</sub> and its cleavage fragment 34K<sub>CpSv</sub> interact with
PDLP1, which shows reduced targeting to PD upon inhibition of the ER-Golgi pathway.
Thus, the 34K<sub>CpSv</sub> cleavage product carries the PDLP1-interacting and tubule-forming
functions required for virus movement and activated upon MP<sub>CpSv</sub> cleavage.

**Discussion**

We have shown that MP<sub>CpSv</sub> has the capacity to target PD, to interact at PD with PDLP1 for
oligomerization and tubule assembly, and to function in virus movement. Moreover, we
found that MP<sub>CpSv</sub> is cleaved into an N-terminal 34 kDa (34K<sub>CpSv</sub>) fragment carrying the
30K super-family domain and into a C-terminal 20 kDa (20K<sub>CpSv</sub>) fragment carrying an
aspartic protease motif responsible for this cleavage. Consistent with the presence of the
30K super-family domain, the 34 kDa fragment carries the ability to target PD, to interact
with PDLP, and to function in virus movement. Mutations within the catalytic domain of
the protease motif abolish tubule formation, alter subcellular localization and decrease the
efficiency of the protein in complementing spread of a MP-deficient TMV, thus indicating
that the formation of the 34 kDa fragment is a prerequisite for tubule-guided movement.
However, the autocleavage-deficient MP\textsuperscript{CPsV}\textsubscript{D340A}:RFP and MP\textsuperscript{CPsV}\textsubscript{D340N}:RFP proteins
retained some of their movement function and complemented the movement of a MP-
deficient TMV construct to some extent without being able to form tubules. This, together
with the fact that TMV is the prototype for non-tubule-guided movement, suggests that the
tubule-guided mechanism may not be the only mechanism by which MP\textsuperscript{CPsV} supports for
virus movement. Nevertheless, tubule-like structures were found at the PD of CPsV-
infected sweet orange plants, suggesting that the virus indeed uses a tubule-guided
movement as a mechanism for movement in its natural host. Ophiovirus particles have been
described as open circular CP decorated flexuous filaments of 3-4 nm thickness (56). Since
tubule-forming viruses usually have an icosahedral particle morphology, such as CPMV
(15, 37), CaMV (11, 27, 32, 33), GFLV (27, 34-36), alfalfa mosaic virus and brome mosaic
virus (39) and tomato spotted wild virus (13), the use of a tubule-guided mechanism by
CPsV may be unexpected. Tubule-guided movement by icosahedral viruses depends on
CP-MP interactions that may allow the viral particles to be guided along the inner tubule
wall (57-60). A tubule-guided mechanism also used by CPsV may therefore be supported
by our previous observation that the MP and CP of CPsV are able to interact (46). Based on
the importance of the 34K fragment of MP\textsuperscript{CPsV} for virus movement, it may be expected that
this fragment interacts with CP within the tubules.
Plant virus-encoded proteases belong to three classes (61). Whereas serine and serine-like
proteases have been described in Poty- (62, 63) and Secoviridae (64-66) families, cysteine
proteases are present in Beny- (67), Marafi- (68), Tymo- (69) and Closterovirus (70), while
aspartic proteases occur in the Caulimoviridae family (55, 71). The protease of CaMV
carries sequence similarity with retrovirus protease (72), thus similar to MP\textsuperscript{CPsV} as shown
here. To our knowledge this is the first report of an aspartic protease encoded by a plant
virus with a negative-strand RNA genome. Virus-encoded aspartic proteases are small
proteins of 10-15 kDa carrying an invariant catalytic aminoacid motif D (T or S) G. The
catalytic form of the enzyme is usually composed of two monomers each providing one D
(T or S) G motif. Cellular aspartic proteases contain two motifs in a single polypeptide chain (55). The 20KCPsV fragment contains only one DTG motif, which suggested that this fragment should dimerize. As shown here 20K-20K interaction takes place at both the nucleus and cytoplasm. At this latter location 20KCPsV co-localized with the MPCPsVD340A self-cleaving deficient mutant, which is cleaved in trans by the 20KCPsV but not when expressed alone. Therefore, the 20KCPsV acts as a viral protease involved in the maturation of the MPCPsV full-length protein. In both transient expression experiments as well as during infection we see that the processing of the full-length MPCPsV is not exhaustive and a high proportion of the MPCPsV remains full length. This is also observed for the trans-cleavage of MPCPsV-D340A by 20KCPsV, suggesting that the protease activity is under regulation (61). Mutants at the cleavage site evaluated in this work showed the capacity to undergo autocleavage, even though when the aminoacids 310LA311 were replaced for residues with different physicochemical properties, suggesting that these positions are not critical for recognition of the cleavage site. Further studies are required to determine the mechanism by which the cleavage site is recognized and cleaved by the protease. Moreover, it would also be important to know whether the 20KCPsV protease targets also other CPsV proteins, or even host proteins, during infection.

We recently showed that MPCPsV can suppress RNA silencing (44). It would be interesting to know whether the RNA silencing suppressing activity resides in the N-terminal 20 kDa or 34 kDa cleavage fragment of the protein.

In conclusion, we propose here a model where the 54K protein encoded in RNA 2 of CPsV, is a poly-protein containing MP and protease activity and should be renamed hereafter as MP-PRO. The MP-PRO polyprotein is cleaved by its aspartic protease activity. This processing event generates two fragments, an N-terminal fragment (34KCPsV) capable of PD targeting and tubule formation renamed as MPCPsV, and a C-terminal fragment (20KCPsV) carrying the aspartic protease activity, renamed as PRO. The MPCPsV fragment is sufficient and even more efficient than the full length MP-PRO in complementing a movement-deficient TMV, although the mechanism can be more specific in the natural host. The observation that the cleavage-deficient mutant MP-PRO D340A supports TMV movement, albeit to a lower efficiency, although it does not form tubules, indicates that the full-length protein complements movement by a different mechanism.
Material and Methods

Virus isolates and plant inoculation
Young leaves of sweet orange plants [Citrus sinensis (L.) Osb] that were either healthy or systemically infected with the CPV-4 isolate of CPsV (73) were used for transmission electron microscopy analysis. The CPsV isolate 90-1-1 (74) was used for mechanical inoculation of Nicotiana occidentalis P1 leaves to obtain protein extracts from infected plants for Immunoblot assays.

Plasmid constructs, bacterial strains and agroinfiltration assays
Protein fusions MpCPsV:GFP and MpCPsV:RFP have been described earlier (52). Mutant MpCPsV derivatives D340A, D340N, 20K and 34K were obtained by site-directed mutagenesis with specific primers and Pfu DNA polymerase (Inbio, Argentina), and using the GATEWAY system-based plasmid pTOPO-MpCPsV (46) as template. Primer sequences are available upon request. The PCR products were digested with DpnI (NEB, USA) to remove the methylated DNA template before transformation into Escherichia coli DH5α competent cells. The introduced mutations were verified by DNA sequencing. The resulting pTOPO-MpCPsV derivatives carrying the desired mutations were subjected to LR recombination (Thermofisher, USA) with destination vectors pB7RWG2 and pB7FWG2 (75), and the resulting plasmids that now encode the mutant MpCPsV proteins fused to either RFP or GFP were transferred into Agrobacterium tumefaciens GV3101.

For transient expression of the fluorescent fusion proteins, A. tumefaciens cultures were harvested by centrifugation, resuspended in water to a final OD_{600nm} of 0.3 (unless stated differently) and infiltrated into the abaxial side of the leaf using a syringe without needle. Leaves were observed at 2-3 days post agroinfiltration (dpai). The expression, size and integrity of the fusion proteins were confirmed by Immunoblot assays.
Protein analysis

Four leaf discs (1.0 cm in diameter) were excised from *N. benthamiana* leaves expressing the viral proteins, ground in liquid nitrogen to fine powder and resuspended in 200 µl of protein extraction buffer (Tris-HCl 75 mM pH = 6.8, 30% glycerol, 5% β-mercaptoethanol, 2% SDS, protease complete inhibitor cocktail (Roche, Germany). This extract was centrifuged at 16000 g for 2 min, and the supernatant was used for immunoblot analysis by adding 200 µl of 4X Laemmli buffer. Samples were boiled for 5 min and centrifuged for 2 min at 16000 g. MP<sup>CP</sup>V (54 K protein) was detected with anti-54K serum as previously described (52). GFP and RFP fusion proteins were detected with anti-GFP (JL-8) monoclonal antibody (BD Biosciences, Clontech, USA) and anti-RFP (6G6) monoclonal antibody (Chromotek, Germany), respectively. Horseradish peroxidase-conjugated anti-mouse (BioRad, USA) was used as secondary antibody. Chemiluminescent reagent was used for detection of peroxidase activity according to the manufacturer’s instructions (GE, ECL Plus Western Blotting Detection Reagents, UK). Densitometry of the protein bands was applied to quantify signal strength using ImageJ(76).

Immunopurification and peptide identification by LC MS/MS

Four grams of tissue powder were resuspended in 8 ml of ice-cold extraction buffer (10 mM Tris/HCl pH 7.5; 150 mM NaCl; 5 mM EDTA; 0.5% NP-40, 1mM PMSF) supplemented with one tablet of complete protease inhibitor cocktail (Roche, country) per 10 ml of buffer and incubated for 30 min, with occasionally inversion of the tube. This extract was centrifuged at 11000 x g for 30 min, filtered through miracloth paper, and centrifuged again for 30 min at 11000 x g at 4 °C. The cleared extract was incubated with 50 µl of GFP-Trap agarose beads (Chromotek, Germany) for 1 hr 30 min at 4 °C. The beads were collected by centrifugation at 2500 x g for 2 min at 4 °C. The supernatant was discarded and the beads were washed four times with washing buffer 1 (WB1: 10 mM Tris/HCl pH 7.5; 150 mM NaCl; 5 mM EDTA, 1 mM PMSF). Bound proteins were reduced with 50 mM dithiothreitol in 50 mM ammonium bicarbonate and alkylated with 50 mM iodoacetamide in ammonium bicarbonate buffer before they were on-bead digested with Trypsin Gold over-night according to manufacture instructions (Promega, USA). Samples were centrifugated at 4 °C and 16000 x g in a microcentrifuge, the supernatants
containing the tryptic peptides were desalted with Zip-Tip C18 (Millipore, USA), and samples were lyophilized and finally resuspended in 10 μl of 0,1% Formic acid solution.

The resulting peptides were then separated by reverse phase nanoHPLC (Thermo Scientific, EASY-Spray Accucore (P/N ES801)) with a continuous gradient of two solutions, i.e. 0,1% formic acid in water and 0,1% formic acid in acetonitrile. The nanoHPLC column was coupled to an Electro Spray ionization source (EASY-SPRAY, Thermo Scientific) at a spray voltage of 3,5 kV. Following ionization, the ions were further separated and analyzed by a mass spectrometer Q-Exactive (Thermo Scientific) equipped with a High Collision Dissociation and an Orbitrap analyzer. Protein identification was carried out with MaxQuant 4.0 software, using an *N. benthamiana* database (Boyce Thompson Institute) where the 54K:GFP protein sequence was added. The digestion mode was semispecific to allow identification of the peptides derived from the aspartic protease. All the parameters left were set with the default values. The peptides were analyzed based on MS/MS count, which is the number of sequencing events for this sequence and the Posterior Error Probability (PEP) of the identification. The PEP value essentially operates as a p-value. Peptides with the highest MS/MS count and lowest PEP value were selected.

**Microscopy**

Confocal laser scanning microscopy (CLSM) and Fluorescence lifetime imaging microscopy (FLIM) were performed as described previously (77). Briefly, for CLSM a Leica TCS SP5 II microscope equipped with a HCX PL APO CS 63.0x 1.40 OIL UV objective was used. Excitation/emission wavelengths were 488/524-550 nm for GFP and 543/566-634 nm for RFP and 405/473-579nm for Aniline blue. Chloroplast autofluorescence was detected by excitation at 488 nm and emission filtering at 654-730nm. Images were acquired with LAS AF version 2.2.1 4842 software and processed with ImageJ software. Callose staining was achieved by infiltrating leaf disks with 0,01% Methy blue (Fluka, UK) solution in PBS buffer prior to observation. FLIM analysis of GFP fluorescence was done with a Lambert Instruments Fluorescence Lifetime Attachment (LIFA) mounted on a Nikon TE2000 inverted microscope. The microscope was equipped with a 63x NA 1.4 oil objective and specific filters for excitation/emission wavelengths of
460–500/510–560 nm for detection of GFP and of 550–600/615–665 nm for detection of RFP. FLIM images were acquired and processed with LI-FLIM software version 1.2.9.117 (Lambert Instruments). ANOVA analysis was performed, followed by a Turkey test (α=0.01). FRET efficiency was calculated as %FRET = 1 − τ_{DA}/τ_D × 100, whereby τ_D is the lifetime of the donor in the absence of the acceptor and τ_{DA} is the lifetime of the donor in the presence of the acceptor.

For transmission electron microscopy (TEM), symptomatic leaves from systemically infected sweet orange plants and healthy leaves from non-infected plants were harvested at a similar developmental stage and fixed at 4 °C with 2% glutaraldehyde in phosphate buffer under smooth vacuum during 2 hrs. Secondary fixation was carried out at 4 °C with 1% osmium tetroxide under vacuum for 1 hr. Samples were dehydrated and embedded in epoxy resin followed by ultrathin sectioning (70 nm). Uranyl acetate and lead citrate were used to contrast the samples prior to observation with a JEM 1200 EX II transmission electron microscope (JEOL Ltd., Tokio, Japan). Images were acquired with an Erlangshen ES1000W (Model 785) CCD camera (Gatan Inc., Pleasanton, California, USA).

**TMV trans-complementation assays**

Trans-complementation assays were performed as previously described (45). Briefly, agrobacteria cultures carrying a TMVΔMPΔCP-GFP-expressing binary plasmid were resuspended in water to an OD_{600nm} of 1 x 10^{-5} and infiltrated into *N. benthamiana* leaves together with agrobacteria resuspended in water to an OD_{600nm} of 0.3 and carrying plasmids encoding either RFP, MP^{TMV}, MP^{CPV}, MP^{CPV}{D340A}, MP^{CPV}{D340N}, 34K^{CPV}, or 20K^{CPV}.

The development of infection foci was observed at 3, 4, and 5 dpai. At these time points, scaled images were acquired and the size of the infection foci was measured with ImageJ software (76).

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France. We would like to thank Dr. Susana Jurado and Dr. Roxana Peralta for their technical assistance with TEM sample preparations conducted at the Servicio Central de Microscopia Electrónica, Fac. Ciencias Veterinarias-UNLP and to Dra. Pía Valacco from the Centro de Estudios Químicos y Biológicos por Espectrometría de Masa (CEQUIBIEM, UBA) for help with the proteomic analysis. We are also grateful for support by Jérôme Mutterer (IBMP, Strasbourg) in performing FLIM measurements. Author Contributions: Conceived and designed the experiments: GRL, EJP, MLG, MH. Performed the experiments: GRL, MBB, EJP. Analyzed the data: GRL, EJP, MBB. Contributed reagents/materials/analysis tools: MLG, MH. Wrote the paper: GRL, EJP, MLG, MH.

Figure legends

FIG. 1. Tubule-like structures in the cell walls of CPsV-infected C. sinensis plants. (A) Transmission electron microscopy images showing PD in healthy (i) and infected (ii) C. sinensis plants. PD, plasmodesmata; CW, cell wall. The arrow in (ii) points to a tubule-like structure protruding from a modified PD. Scale bar, 100 nm. (B) CLSM of N. benthamiana epidermal cells expressing GFP:MP<sub>CPsV</sub> (i) or MP<sub>CPsV</sub>:GFP (ii). Arrowheads in (i) and (ii) indicate the presence of GFP:MP<sub>CPsV</sub> and MP<sub>CPsV</sub>:GFP at PD; arrows in (ii) indicate the presence of tubules containing MP<sub>CPsV</sub>:GFP at PD. Scale bar, 30µm.

FIG. 2. MP<sub>CPsV</sub>-PDLP interaction is necessary for MP-tubule formation at PD. (A) FRET-FLIM measurements of N. benthamiana epidermal cells expressing (i) MP<sub>CPsV</sub>:GFP alone, or together with either (ii) PDCB1:Cherry, (iii) MP<sub>CPsV</sub>:RFP, or (iv) PDLP1:RFP. Fluorescent intensity images (top) are combined with fluorescence lifetime images (bottom) showing lifetime in false color code according to the color scale on the left. Scale bar, 10 µm. A representative fluorescence lifetime analysis based on three independent replicate experiments is shown (v). τ, fluorescent lifetime (ns); SD, standard deviation; N, number of cells analyzed. Asterisks represent significant differences compared to MP<sub>CPsV</sub>:GFP expressed alone (P<0.01). (B) FRET-FLIM measurements of N. benthamiana expressing (i) GFP:MP<sub>CPsV</sub> alone or together with (ii) PDCB1:Cherry, (iii) MP<sub>CPsV</sub>:RFP, or (iv) PDLP1:RFP. Fluorescent intensity images (top) are combined with fluorescence lifetime images (bottom) showing lifetime in false color code according to the color scale on the left. Scale bar, 10 µm. A representative fluorescence lifetime analysis based on three independent replicate experiments is shown (v). τ, fluorescent lifetime (ns); SD, standard deviation.
deviation; N, number of cells analyzed. Asterisks represent significant differences respect to GFP:MP<sub>CPsV</sub> expressed alone in GFP fluorescence lifetime (P<0.01). (C) Expression of PDLP1:RFP together with either Sar1:GFP (i and ii) or Sar1[H74L]:GFP (iii and iv) (only RFP channel is shown in magenta). The images (ii) and (iv) (scale bar, 10 µm) show enlargements of the framed tissue regions indicated in (i) and (ii) (scale bar, 50 µm); arrowheads indicate PD. Immunoblots (v) show protein expression levels in each experiment. (D) Expression of MP<sub>CPsV</sub>:RFP together with either Sar1:GFP (i and ii) or Sar1[H74L]:GFP (iii and iv). RFP channel is shown in magenta and callose staining is showed in yellow. The images (ii) and (iv) (scale bar, 5 µm) show enlargements of the framed tissue regions indicated in (i) and (ii) (scale bar, 10 µm); arrowheads indicate PD; arrows indicate tubule-like structures at PD. Immunoblots (v) show protein expression levels in each experiment.

**FIG.3.** Fluorescence protein orientation affects MP<sup>CPsV</sup> movement activity and cleavage. (A) (Left) Representative images of *N. benthamiana* leaves showing the development of TMVΔCPΔMP-GFP infection foci at 5 dpai in the presence of either GFP:MP<sub>CPsV</sub>, MP<sub>CPsV</sub>:GFP or GFP (negative control). Scale bar, 10 mm. (Right) Analysis of the sizes of infection foci shown in (A). Letters above columns indicate statistical differences (P<0.01). (B) Immunoblot with anti-GFP antibody. The upper band corresponds to the full-length fusion protein; asterisks indicate the presence of additional GFP-containg proteins. Molecular masses are indicated on the right of the immunoblot. (C) (Top) Schematic representation of the predicted chloroplast transit peptide (cTP) at the N-terminal end of MP<sub>CPsV</sub>; (bottom) subcellular localization of MP<sub>CPsV</sub>:GFP (upper panel yellow) and GFP:MP<sub>CPsV</sub> (lower panel yellow) in relation to the chloroplasts (autofluorescence, in red). Scale bar, 10 µm.

**FIG. 4.** MP<sub>CPsV</sub> encodes an aspartic protease motif. (A) Immunoblot analysis of healthy and CPsV infected *N. occidentalis* P1 plants using anti-MP<sub>CPsV</sub> (anti-54) serum. Molecular masses of virus-specific bands are indicated on the right of the immunoblot. (B) (i) (Top) Representation of part of the aminoacids sequence of the MP<sub>CPsV</sub> aligning the aspartic protease domain with the respective domains in Cathepsin D (PDB structure ID: 4Od9_A) and HIV-2 protease (PDB structure ID: 3ec0_A), the catalytic D residue is underlined in...
each case. (Bottom) Alignment of the aminoacid sequences of HIV protease substrate peptides against the aminoacid sequence of MP\textsuperscript{CPsV}. The specific HIV protease cleavage releases specific proteins; MA, matrix; CA, capsid; NC, nucleocapsid; TF, trans-frame peptide; PR, protease; AutoP, auto proteolysis site; RT, reverse transcriptase; RH, RNAse H; IN, integrase. Cleavage site is indicated with an “|”. (ii and iii) Representation of part of the aminoacids sequence of the MP\textsuperscript{MELBVV} and MP\textsuperscript{BLMaV} respectively, aligning the aspartic protease domain with the respective domains in Cathepsin D (PDB structure ID: 4Od9\textunderscore A) and aspartic protease (PDB structure ID: 3LIY\textunderscore D). The catalytic D residue is underlined.

(C) MS/MS spectra showing the ions matching with 54K derived peptides indicated at the bottom of each spectra. (i) Spectra of the peptide KSVSINLSNFL corresponding to the C-terminal end of the 34K fragment and (ii), spectra of the peptide ADQRRAPPQPLEKR corresponding to the N-terminal end of the 20K protein. MS/MS counts and PEP values are indicated. (D) MP\textsuperscript{CPsV} mutants used in this work. Aminoacid replacements are indicated underlined. (E) Immunoblot of protein extracts of \textit{N. benthamiana} tissues transiently expressing either RFP, MP\textsuperscript{CPsV}:RFP, MP\textsuperscript{CPsV}\textsuperscript{D340A}:RFP, MP\textsuperscript{CPsV}\textsuperscript{D340N}:RFP, 34K\textsuperscript{CPsV}:RFP, 20K\textsuperscript{CPsV}:RFP, MP\textsuperscript{CPsV}\textsuperscript{L310D}:RFP, MP\textsuperscript{CPsV}\textsuperscript{A311D}:RFP, MP\textsuperscript{CPsV}\textsuperscript{A311R}:RFP or MP\textsuperscript{CPsV}\textsuperscript{A311H}:RFP with anti-RFP monoclonal antibody. Molecular masses are indicated on the left of the immunoblot.

**FIG. 5.** Subcellular localizations of MP\textsuperscript{CPsV} mutants. Co-expression of PDCB1:Cherry with together with (A) MP\textsuperscript{CPsV}:GFP, (B) MP\textsuperscript{CPsV}\textsuperscript{D340A}:GFP, (C) MP\textsuperscript{CPsV}\textsuperscript{D340N}:GFP (D) 34K\textsuperscript{CPsV}:GFP or (E) 20K\textsuperscript{CPsV}:GFP in \textit{N. benthamiana} epidermal cells at 3 dpai. The patterns of GFP fluorescence (yellow), Cherry fluorescence (magenta), and chloroplast autofluorescence (red) are shown. Arrows indicate locations of tubule-like structures at PD; arrowheads GFP-fluorescent PD without tubules; white rectangles, chloroplasts; Nu, nucleus. Scale bar, 20 µm.

**FIG. 6.** 20K\textsuperscript{CPsV} cleavage product retains the aspartic protease activity. (A) FRET-FLIM measurements of \textit{N. benthamiana} epidermal cells expressing (i) 20K\textsuperscript{CPsV}:GFP either alone or together with (ii) RFP or (iii) 20K\textsuperscript{CPsV}:RFP. Fluorescent intensity images (top) are combined with fluorescence lifetime images (bottom) showing lifetime in false color code according to the color scale on the left. Scale bar, 10 µm. A representative fluorescence
lifetime analysis based on two independent replicate experiments is shown (iv and v), in
which the lifetime was measured in the nucleus (iv) and cytoplasm (v). $\tau$, fluorescent
lifetime (ns); SD, standard deviation; N, number of nucleus or cytoplasm analyzed.
Asterisks represent significant differences compared to 20K$^{CP\alpha V}$:GFP expressed alone
(P<0.01). (B) Immunoblot analysis of N. benthamiana plants expressing
MP$^{CP\alpha V}$D340A:GFP either alone or in combination with MP$^{CP\alpha V}$:RFP or 20K$^{CP\alpha V}$:RFP. Top
panel, anti-RFP antibody; bottom panel, anti-GFP antibody. Arrows indicate the fragment
produced by in trans proteolytic processing of MP$^{CP\alpha V}$D340A:GFP by either MP$^{CP\alpha V}$:RFP
(left) or 20K$^{CP\alpha V}$:RFP (right). (C) Expression of MP$^{CP\alpha V}$D340A:GFP (yellow) together with
(i) MP$^{CP\alpha V}$:RFP (magenta) or (ii) with 20K$^{CP\alpha V}$:RFP (magenta) in N. benthamiana leaves at
3 dpai. Arrowheads indicate tubule-like structures at PD. Scale bar, 10 µm.

FIG. 7. Virus movement activity of MP$^{CP\alpha V}$ mutants. (A) Representative images
showing TMVΔCPΔMP-GFP infection foci at 5 dpai in N. benthamiana leaves expressing
the indicated protein. Scale bar, 10 mm. (B) Size of infection foci in the presence of
indicated proteins measured at 5 dpai. Each letter indicates protein treatments with
statistical differences between them (P<0.01). (C) Size distribution of the infection foci
according to specific treatment at 5 dpai.

FIG. 8. 34K$^{CP\alpha V}$-PDLP interaction is necessary for tubule formation at PD. A) FRET-
FLIM measurements of N. benthamiana epidermal cells expressing (i) 34K$^{CP\alpha V}$:GFP either
alone, or together with (ii) PDCB1:Cherry, (iii) 34K$^{CP\alpha V}$:RFP, or (iv) PDLP1:RFP.
Fluorescent intensity images (top) are combined with GFP fluorescence lifetime images
(bottom) indicating lifetime in false color code according to the scale on the left. Scale bar,
10 µm. Lifetime analysis (v): $\tau$, fluorescent lifetime (ns); SD, standard deviation; n, number
of cells analyzed. Asterisks represent a significant reduction in GFP fluorescence lifetime
(P<0.01) compared to 34K$^{CP\alpha V}$:GFP. (B) Expression of 34K$^{CP\alpha V}$:RFP together with
Sar1:GFP (i and ii) or with Sar1[H74L]:GFP (iii and iv). RFP channel is shown in magenta
and callose staining is showed in yellow. The images (ii) and (iv) (scale bar, 5 µm) are
enlargements of the framed leaf regions shown in (i) and (ii) (scale bar, 10 µm);
arrowsheads, PD; arrows, tubule-like structures at PD. Immunoblot analysis showing protein
expression levels (v).


74. García ML, Derrick KS, Grau O. Citrus psorosis associated virus and citrus ringspot virus belong to a new virus group, p 430-431. In In P. Moreno JVdGaLWT (ed), IOCV,


<table>
<thead>
<tr>
<th>Assay</th>
<th>N° fields with tubule-like structures at PD</th>
<th>N° fields without tubule-like structures at PD</th>
<th>N° fields observed</th>
<th>% of fields with tubule-like structures at PD</th>
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*represent statistical difference compared to Sar [H74L] unpaired t-test with p<0.01.
TABLE 2. 34K<sup>CPnV</sup> tubule formation upon inhibition of the secretory pathway

<table>
<thead>
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<th></th>
<th>34K&lt;sup&gt;CPnV&lt;/sup&gt;:RFP + Sar1:GFP</th>
<th>34K&lt;sup&gt;CPnV&lt;/sup&gt;:RFP + Sar1[H74L]:GFP</th>
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<tr>
<td></td>
<td>N° of fields with tubule-like structures at PD</td>
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<td>29</td>
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*represent statistical difference compared to Sar [H74L] unpaired t-test with p<0.05.
Figure A: Imaging of FRET (Förster resonance energy transfer) at different percentages (10%, 5%, 15%, 23%) and lifetimes (ns).

Figure B: Life span analysis at various stages.

Figure C: Imaging of PDL1:RFP and Sarl1:GFP.

Figure D: Imaging of MP11:RFP, Callose, and merged images.

Figure V: Western blot analysis showing protein levels and Coomassie blue staining.