Immunocapture of dsRNA-bound proteins provides insight into Tobacco rattle virus replication complexes and reveals Arabidopsis DRB2 to be a wide-spectrum antiviral effector

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Short title: dsRNA-bound proteome from virus-infected Arabidopsis

One-sentence summary: Double-stranded RNA immunoprecipitation allows the identification of Arabidopsis host proteins associated with tobacco rattle virus replication complexes.

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ABSTRACT

Plant RNA viruses form organized membrane-bound replication complexes to replicate their genomes. This process requires virus- and host-encoded proteins and leads to the production of double-stranded RNA (dsRNA) replication intermediates. Here, we describe the use of Arabidopsis thaliana expressing GFP-tagged dsRNA-binding protein (B2:GFP) to pull down dsRNA and associated proteins in planta upon infection with Tobacco rattle virus (TRV). Mass spectrometry analysis of the dsRNA-B2:GFP-bound proteins from infected plants revealed the presence of viral proteins and numerous host proteins. Among a selection of nine host candidate proteins, eight showed re-localization upon infection, and seven of these co-localized with B2-labeled TRV replication complexes. Infection of A. thaliana T-DNA mutant lines for eight such factors revealed that genetic knock-out of DSRNA-BINDING PROTEIN 2 (DRB2) leads to increased TRV accumulation and DRB2 overexpression caused a decrease in the accumulation of four different plant RNA viruses, indicating that DRB2 has a potent and wide-ranging antiviral activity. We propose B2:GFP-mediated pull down of dsRNA to be a versatile method to explore virus replication complex proteomes and to discover key host virus replication factors. Given the universality of dsRNA, development of this tool holds great potential to investigate RNA viruses in other host organisms.

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INTRODUCTION

Viruses are obligate endocellular parasites that hijack their host’s molecular processes and machinery to multiply, a process that sometimes results in devastating diseases. Pivotal to a successful infection is the efficient replication of the viral genomic nucleic acid(s). In the majority of plant virus species, the genome consists of one or more molecules of single-stranded positive polarity RNA, or (+)ssRNA. Replication is carried out by the virus-encoded RNA-dependent RNA-polymerase (RdRp), often part of a larger protein known as the replicase. This enzyme first copies the viral (+) genome into (-)ssRNA that is then used as a template for the production of progeny (+)ssRNA. Intrinsic to the RNA replication process is the generation of long double-stranded RNA (dsRNA) intermediates by the viral RdRp.

The replication of all known (+) strand RNA virus takes place on host membranes whose origin, whether the endoplasmic reticulum, chloroplasts, mitochondria, peroxisomes, etc., depends on virus species (reviewed in (Ritzenhaler and Elamawi, 2006; Verchot, 2011; Grangeon et al., 2012; Jiang and Laliberté, 2016)). The progressive virus-induced recruitment of such membranes generally leads to dramatic reorganizations of the host endomembrane system into so called “viral factories”. These viral factories are the sites where all steps vital to the virus life are carried out including protein translation, RNA encapsidation and RNA replication sensu stricto.

The specialized molecular entities on which RNA replication occurs within the viral factories are known as the virus replication complexes (VRCs). While a minimal VRC arguably consists of single- and double-stranded viral RNA and replicase, their precise composition, which depends on the virus and host species, remains largely unexplored. A number of studies have shown that specific host proteins can be integral part of these VRC and exert positive (pro-viral) or negative (anti-viral) effects on replication. Our knowledge on these host proteins is summarized in several exhaustive reviews (Nagy and Pogany, 2011;
Wang, 2015; Nagy, 2016). These include among others RNA-binding proteins, RNA helicases, chaperones and proteins belonging to the RNA interference machinery (RNAi, or RNA silencing), which is the primary response to replicating viruses in plants and other eukaryotes (Wang, 2015; Barton et al., 2017).

Antiviral RNAi against RNA viruses in the model plant *Arabidopsis thaliana* is initiated by RNAse-III Dicer-Like enzymes DCL4 and DCL2, which cleave dsRNA into 21- and 22-nt small-interfering RNA (siRNA), respectively. These siRNAs are then loaded into Argonaute (AGO) proteins, which use them as templates to recognize and cleave viral ssRNA in a sequence-specific manner. Viruses have evolved a vast array of strategies to evade or block RNAi, the best studied of which are viral suppressors of RNA silencing (VSRs). These proteins suppress silencing through a wide range of molecular strategies, from inhibition of dicing, to siRNA sequestration, to AGO degradation (Incarbone and Dunoyer, 2013). Of note, *A. thaliana* encodes several RNAse-III-like enzymes (RTLs) in addition to Dicers, but little is known regarding their function (Elvira-Matelot et al., 2016). The accumulation of viral dsRNA *in planta* varies widely among (+)ssRNA virus and host species, as we have recently shown (Monsion et al., 2018). While the precise molecular events occurring during virus RNA replication are often unclear, it can be argued that rapid and efficient separation of the (+) and (-) RNA strands could not only allow more replication cycles to take place, but also constitute a powerful mechanism of RNA silencing evasion/suppression through removal of dsRNA.

A great deal of our knowledge on plant virus VRCs emerged from a series of seminal studies conducted with *Tomato bushy stunt virus* (TBSV) on yeast (*Saccharomyces cerevisiae*), a surrogate host used as a powerful biological tool to conduct genetic screens and functional studies on host factors involved in TBSV VRC activity (reviewed in (Nagy et al., 2016)). While the authors of these studies, where possible, validated the results obtained in
yeast and in vitro in Nicotiana benthamiana, data obtained in planta on VRCs of other viruses remains sparse. Recent studies have reported methods to spatio-temporally visualize VRCs in vivo via fluorescently labeled dsRNA-binding proteins (Cheng et al., 2015; Barton et al., 2017; Monsion et al., 2018). Candidate-based, reverse genetic approaches have been used to probe the involvement of host factors in VRC formation and activity in plants (Wei et al., 2013; Li et al., 2016). These approaches, however, are necessarily based on prior discovery acquired by other experimental means. Another experimental strategy successfully used to characterize VRCs has been to pull down tagged viral proteins and analyze the resulting protein populations by mass spectrometry (Dufresne et al., 2008; Lohmus et al., 2016; Wang et al., 2018). While this last method has provided valid and compelling data, we decided to investigate VRCs from a viral RNA-centered, rather than viral protein-centered, perspective. We hypothesized that pull-down of dsRNA from virus-infected plants followed by mass spectrometry of dsRNA-associated proteins would provide insight into the molecular composition of VRCs.

In this study, we report that the dsRNA-binding domain of FHV protein B2, when fused to GFP (B2:GFP hereafter) can not only be used to visualize VRCs in planta as shown previously in N. benthamiana (Monsion et al., 2018), but additionally can be exploited as a means to pull-down dsRNA-associated nucleo-protein complexes from infected Arabidopsis plants. As a proof of concept, we used Tobacco rattle virus (TRV genus: Tobravirus, family: Virgaviridae), a well-studied (+)ssRNA virus, to perform B2:GFP immunoprecipitations (IPs). In this manner, a number of virus- and host-encoded proteins could be identified and validated for their localization in and around VRCs. Among these candidates, DRB2, a dsRNA-binding protein was found to display antiviral activity by using loss- and gain-of-function approaches. These results provide robust validation of dsRNA pull-down as an effective and high-throughput method for VRC characterization in planta. Furthermore, the
results offer detailed snapshots of TRV replication complexes and viral factories, with host factors showing unique and distinct localization patterns in and around these complexes.

RESULTS

B2:GFP-mediated isolation of tobacco rattle virus dsRNA from A. thaliana

The double-stranded RNA-binding B2:GFP protein, when ectopically expressed in transgenic N. benthamiana, has been previously shown to specifically associate with the VRCs of several positive-strand RNA viruses from plants and insects (Monsion et al., 2018). For the sake of clarity, VRCs refers hereafter to all the factors that directly and/or indirectly associate to the viral replicating dsRNA rather than to the replicase core complex sensu stricto.

Following these findings, we wished to further exploit B2:GFP as a biochemical bait to explore the composition and biology of RNA VRCs, the pivotal element of which is dsRNA. To do so, and given the versatility of A. thaliana as a model plant species, we first produced homozygous 35S:B2:GFP transgenic plants. Although in this work we focused essentially on the 35S:B2:GFP/Col-0 line, 35S:B2:GFP was also introduced into various genetic backgrounds including mutants of the core antiviral Dicer-Like genes, dcl2-1, dcl4-2 and triple dcl2-1/dcl3-1/dcl4-2 (Supplemental Figure S1A,B,C). The rationale behind this choice is that DCL proteins are arguably the best-known RNase III enzymes in plants, and the small-interfering RNA (siRNA) they generate from virus-derived dsRNA precursors are the effectors of RNA silencing, the main antiviral defense in plants (Incarbone and Dunoyer, 2013; Pumplin and Voinnet, 2013).

Similar to the B2:FP N. benthamiana lines (Monsion et al., 2018) and despite the clear expression of B2:GFP (Supplemental Figure S1A), the transgenic lines in the different dcl mutant backgrounds showed little to moderate developmental phenotypes (Supplemental Figure S1C). These were reminiscent of (but distinct from) those caused by ectopic
expression of other RNA silencing suppressors such as P19 or HC-Pro (Kasschau et al., 2003; Incarbone et al., 2017). In addition to leaf serration, the elongated and slightly downward-curled leaves of the B2:GFP line are reminiscent of mutants in the miR390/TAS3 pathway such as rdr6, dcl4 and ago7 (Adenot et al., 2006). Such phenotypes may be determined (i) by inhibition of long dsRNA processing into siRNA and/or (ii) by disruption of miRNA function through their sequestration.

The full-length FHV B2 protein has been shown to be a suppressor of RNA silencing (Li et al., 2002; Seo et al., 2012), and proposed to act through both inhibition of dicing and sequestration of siRNA duplexes (Chao et al., 2005). To investigate whether the GFP-tagged 73 amino acid dsRNA binding domain of B2 that lacks the residues involved in the interaction with PAZ domains of Dicer proteins (Singh et al., 2009; Liu et al., 2012) also acts as a suppressor of RNA silencing, we performed a standard GFP silencing patch test on N. benthamiana leaves (Supplemental Figure S1D,E). B2, as a C-terminal fusion to tRFP (B2:tRFP) was able to suppress silencing of the GFP transgene, as was turnip crinkle virus suppressor P38. By contrast, a C44S, K47A double-mutated version of B2:tRFP impaired in dsRNA binding (Monsion et al., 2018) was unable to suppress silencing, suggesting that suppression activity is dsRNA-binding-dependent and likely DCL-binding-independent.

Next, we investigated the effects of stably expressed B2:GFP on endogenous small RNA pathways: biogenesis of microRNAs 159 and 160 was not perturbed in 35S:B2:GFP/Col-0 plants, while biogenesis of siRNA such as endo-siRNA (IR71) and trans-acting siRNA (TAS1) was completely abolished (Supplemental Figure S1B). Whether these defects in endo-siRNA and trans-acting siRNA biogenesis are responsible of the observed developmental phenotypes remains to be determined.

We then infected 35S:B2:GFP/Col-0 plants with a recombinant TRV carrying part of the PHYTOENE DESATURASE (PDS) gene (Liu et al., 2002). As expected, the control Col-0
plants showed minor viral symptoms and the typical bleaching phenotype linked to PDS gene silencing. In contrast, the B2:GFP-expressing plants showed no significant leaf discoloration but severe viral symptoms (Figure 1A) and death of the plants occurred before flowering (not shown), well in agreement with the efficient RNA silencing suppression activity of the B2 dsRNA-binding domain. Observation of systemically infected leaves by confocal microscopy showed TRV-induced re-localization of B2:GFP to distinct cytosolic mesh-like structures (Figure 1B) reminiscent to those observed in 35S:B2:GFP/N. benthamiana and shown to correspond to TRV-induced VRCs (Monsion et al., 2018). RNA gel blot analysis of RNA from TRV-PDS systemically infected 35S:B2:GFP/N. benthamiana and 35S:B2:GFP/Col-0 revealed that B2:GFP caused a striking over-accumulation of viral (+)ssRNA in both plant species (Figure 1C). This is also in agreement with B2:GFP activity as a suppressor of RNA silencing, and could be recapitulated in a dcl2-1 dcl4-2 double mutant (dcl24), which lacks the two main antiviral Dicers (Supplemental Figure S1F)(Deleris et al., 2006). B2:GFP also caused a tremendous increase in long double-stranded RNA content, likely corresponding to replication intermediates, as determined by dsRNA–protein gel blot (northwestern) blotting in both B2:GFP-expressing Col-0 and N. benthamiana plants (Figure 1D).

In contrast, virus-derived siRNAs (vsiRNA) accumulated differentially between Arabidopsis and N. benthamiana. Thus, while B2:GFP expression led to an overall reduction in 21 and 22 nt vsiRNA species in Col-0 plants, the opposite effect was observed in N. benthamiana (Figure 1E). Conversely, miR159 and U6-derived small nucleolar RNA (snRNA) accumulation was unaffected by the presence of B2:GFP in both plant species (Figure 1E, Supplemental Figure S1B). These results suggest that B2:GFP interferes with TRV RNA processing by DCL enzymes either by promoting (N. benthamiana) or preventing (Arabidopsis) siRNA production, in a manner that is associated with enhanced viral
replication in both plant species. Whether increased TRV replication is a cause or a consequence in siRNA levels is unknown.

As a first experiment establishing B2:GFP as a tool to study VRC composition, we performed anti-GFP IPs from TRV-PDS-infected 35S:B2:GFP/Col-0 plants and analyzed their composition in (+)strand viral RNA (Figure 1C), long dsRNA (Figure 1D), siRNA (Figure 1E) and proteins (Figure 1F). As a negative control, we included TRV-PDS-infected 35S:GFP/Col-0 plants. RNA gel blot and dsRNA–protein gel blot analyses performed on IPed RNA revealed that immune complexes contained (+)ssRNA, long dsRNA and 22nt vsiRNA, but no U6 and miR159 (Figure 1C-E). Interestingly, and in contrast with our previous report in vitro (Monsion et al., 2018) but well in agreement with the capacity of B2 to bind dsRNAs longer than 18 bp (Chao et al., 2005), antiviral siRNA were immunoprecipitated (Figure 1E). Immunoblot analysis of proteins from the same experiment revealed efficient IP of both GFP and B2:GFP (Figure 1F). Altogether we concluded that immunoprecipitation allowed the isolation of TRV double-stranded replication intermediates, the core element of VRCs.

**Immunoprecipitation and identification of B2:GFP-associated viral and host proteins by mass spectrometry**

Once established that immunoprecipitation of B2:GFP from plants allowed efficient isolation of virus replication dsRNA intermediates, we wondered whether these complexes contain specific virus- and host-encoded proteins. To address this question, we performed anti-GFP IP on TRV-PDS-infected 35S:GFP/Col-0 vs. 35S:B2:GFP/Col-0 in triplicate, and analyzed the immunoprecipitated proteins by mass spectrometry (MS). The complete list of identified viral and host proteins from this analysis is shown in Supplemental Data Set S1. We also
performed the same IP and MS analysis on non-infected 35S:GFP/Col-0 vs. 35S:B2:GFP/Col-0 plants (Supplemental Data Set S2).

A preliminary analysis by immunoblot confirmed efficient and reproducible B2:GFP and GFP immunoprecipitation (Supplemental Figure S2), which could be confirmed by MS, reads from B2:GFP and GFP being the most abundant (Figure 2A,B, Supplemental Data Set S1). We next searched and ranked accessions that were identified only - or enriched - in TRV-infected 35S:B2:GFP/Col-0 samples (Figure 2A,B, Supplemental Data Set S1). As expected, the TRV replicase (Uniprot accession Q9J942) was among the most abundant proteins detected in IPs from B2:GFP samples (Figure 2). This result, along with the previously described detection of viral dsRNA in analogous IPs (Figure 1), suggests that B2:GFP immunoprecipitation allows the isolation of TRV VRCs. Detection of TRV coat protein (CP, Uniprot Q88897) and 16k suppressor of silencing (Uniprot Q77JX3) in B2:GFP IPs (Figure 2) also suggests that these viral proteins, not known to participate in replication, associate directly or indirectly to dsRNA. Although unlikely, we can’t at this point rule out that one or more of these TRV proteins bind B2:GFP and not dsRNA.

In addition to TRV-encoded proteins, MS analysis allowed also the identification of 110 host proteins exclusively present in IPs from B2:GFP-expressing plants (Figure 2 and Supplemental Data Set S1), which we considered as replication complex-associated host protein candidates. 29 of these proteins were significantly enriched in the IPs with an adjusted p-value < 0.05 (Figure 2B). Volcano plots generated by the comparison of the TRV-infected 35S:B2:GFP/Col-0 samples with combinations of all controls (non-infected 35S:B2:GFP/Col-0, non-infected 35S:GFP/Col-0 and TRV-infected 35S: GFP/Col-0) are displayed in Supplemental Figure S3.

To evaluate the association of candidate host proteins to replication complexes and considering their high number (Supplemental Data Set S1), we arbitrarily restricted our
analysis to a set of nine *A. thaliana* gene products that were either detected with high (>60) spectral counts (AT5G02500, AT3G09440, AT3G12580) or confirmed/potential RNA-binding/interacting proteins from literature or NCBI annotation (AT1G24450, AT5G04430 (Fujisaki and Ishikawa, 2008), AT3G62800 and AT2G28380 (Clavel et al., 2015)), or both (AT1G23410, AT3G45570)(Figure 2A,B). The distribution of peptide reads along these selected proteins, along with the TRV and bait proteins, is shown in Figure 2C. It should be noted that four of the candidates (AT1G23410, AT3G09440, AT5G02500 and AT3G12580) were also present in B2:GFP IPs from non-infected plants (Supplemental Data Set S2) which may reflect their dsRNA binding activity in both healthy and virus-infected plants. However, the spectral count of peptides from these proteins was a fraction of that detected in IPs from TRV-infected plants, despite the spectral counts of the bait proteins being comparable. All other candidates were not detected in B2 IPs from non-infected plants. Finally, we excluded from our priority list a number of proteins that were significantly enriched in B2:GFP vs. GFP plants due to the number of candidates to analyze and their apparent lack of significance in viral replication process based on literature. This includes for instance the most enriched protein, a myrosinase with anti-microbial activity that is present in Brassica crops and involved in defense against herbivores (Bhat and Vyas, 2019).

**tRFP does not label TRV replication complexes in planta**

In a second step, we tested the subcellular localization of the selected candidates in relation to B2:GFP in healthy and TRV-infected plants. To do so, we opted for the 35S-driven transient expression of the Arabidopsis candidates as N- or C-terminal fusions to tRFP in healthy or TRV-infected 35S:B2:GFP/N. benthamiana. In all cases, confocal imaging was performed 3-4 days post agro-infiltration, a time that was found to be optimal for TRV-infection and transient expression of protein candidates.
As an absolute prerequisite to our validation pipeline of candidate proteins and considering tRFP was used as reporter tag, we first carefully analyzed the intracellular distribution of tRFP with respect to TRV replication complexes in 35S:B2:GFP/N. benthamiana. As expected, tRFP as well as B2:GFP showed a typical nucleo-cytoplastic localization in cells from healthy plants (Figure 3A), well in agreement with our previous report using the same experimental system (Monsion et al., 2018). Crucially, upon infection the intracellular distribution of tRFP remained unchanged, while B2:GFP concentrated into bright cytoplasmic cotton-ball-like structures often adjacent to the nucleus (Figure 3B). These large structures were previously shown to correspond to TRV viral factories enriched in mitochondria-derived membranes (Monsion et al., 2018) on which replication of TRV is thought to occur (Otulak et al., 2015). Importantly, our data clearly show that while B2:GFP is highly enriched in TRV replication factories, tRFP alone is significantly depleted from these structures (Figure 3B), in agreement with the behavior of tRFP as cytoplasmic and validating tRFP as a reporter protein with which to tag the candidates of interest.

Double-stranded RNA-binding proteins (DRBs) perfectly colocalize with B2-labeled viral replication complexes

DRBs are proteins with dual dsRNA-binding motifs with five representatives in the Arabidopsis genome (Schauer et al., 2002; Clavel et al., 2015). Despite showing low spectral counts in our IPs, two DRBs were identified in our analysis: DRB2 (AT2G28380, total counts: 4, Figure 2A) and DRB4 (AT3G62800, total counts: 5, Figure 2A) that were obvious candidates to test (Figure 4).

DRB2 localizes to the replication complexes of different RNA viruses (Barton et al., 2017), binds dsRNA (Tschopp et al., 2017) and plays a role in endogenous small RNA biogenesis (Péllissier et al., 2011; Eamens et al., 2012; Clavel et al., 2015). In non-infected
plants DRB2:tRFP and B2:GFP localized to partially overlapping cytoplasmic and nuclear structures. Interestingly, over-expression of DRB2 changed the localization pattern of B2 from a predominantly nuclear localization (Figure 3A and (Monsion et al., 2018)) to DRB2-labeled cytoplasmic structures as if B2:GFP was recruited to DRB2 localization sites (Figure 4A). Remarkably, such redistribution of B2 was not observed upon overexpression of DRB4 (Figure 4C). Crucially, near-perfect colocalization of DRB2:tRFP and B2:GFP was observed in the VRCs upon TRV-PDS infection (Figure 4A), which was particularly evident at high magnification (Figure 4B). Moreover, while DRB2:tRFP was almost exclusively found in the cytoplasmic VRCs upon infection, a substantial fraction of B2:GFP remained associated to nuclear structures likely containing dsRNA (Figure 4B). This suggests that although both proteins bind dsRNA, their intracellular targeting is likely not exclusively dsRNA-dependent.

DRB4 has been shown to be both a co-factor of DCL4 in small RNA biogenesis and an inhibitor of DCL3 in endogenous inverted-repeat RNA processing in A. thaliana (Fukudome et al., 2011; Montavon et al., 2017). More relevant here, DRB4 is involved in the defense against RNA viruses (Qu et al., 2008; Jakubiec et al., 2012). When we expressed DRB4:tRFP in non-infected tissue, this protein accumulated predominantly to the nucleus where it colocalized with B2:GFP (Figure 4C), in agreement with previous reports (Barton et al., 2017; Monsion et al., 2018). Upon TRV infection DRB4:tRFP was clearly redistributed to VRCs, where it perfectly colocalized with B2:GFP (Figure 4C,D). In contrast to DRB2 that was barely detectable in the nucleus (Figure 4B), a significant fraction of DRB4:tRFP remained nuclear upon infection (Figure 4C,D).

Altogether, the robust colocalization of both double-stranded RNA-binding proteins DRB2 and DRB4 with B2:GFP during infection provide (i) further evidence that the TRV-viral factories are indeed cytoplasmic dsRNA hotspots and, more importantly, (ii) a validation of the immunoprecipitation procedure.
Proteins previously linked to viral infection localize at/near VRCs

A family of proteins that emerged with high spectral counts were those belonging to the HSP70 family: HSP70 (AT3G12580, 63 counts), HSP70-1 (or HSC70-1, AT5G02500, 81 counts) and HSP70-3 (or HSC70-3, AT3G09440, 71 counts) (Figure 2A). Members of this family of chaperones have been shown in several studies to play key roles in virus infection cycles (reviewed in (Verchot, 2012; Nagy, 2016)). They can regulate viral life cycles both positively and negatively, and depending on the virus, they affect VRC formation, virus movement and coat protein homeostasis, among other processes. Three recent studies showed that unrelated plant viruses hijack HSP70 to greatly enhance virus replication (Pogany and Nagy, 2015; Lohmus et al., 2017; Yang et al., 2017).

All three HSP70 members were tested in TRV-infected and non-infected 35S:B2:GFP/N. benthamiana (Figure 5 and Supplemental Figure S4). When overexpressed in healthy plants HSP70:tRFP, HSP70-1:tRFP, HSP70-3:tRFP located essentially to distinct cytoplasmic foci whose number, size and distribution were specific for each of the three HSP70 observed (Supplemental Figure S4). Remarkably, upon infection, HSP70-1:tRFP (Figure 5B) and HSP70-3:tRFP (Figure 5C) were clearly redistributed to TRV viral factories enriched in B2:GFP. In contrast, the localization pattern of HSP70:tRFP remained essentially unaffected upon infection, with no obvious colocalization of B2:GFP with HSP70:tRFP-labeled foci (Figure 5A, Supplemental Figure S4A).

It should be noted that despite the clear redistribution of HSP70-1:tRFP and HSP70-3:tRFP upon infection, only partial colocalization was detected between these proteins and B2:GFP (Figure 5B,C). The latter appeared engulfed in large HSP70-1 or HSP70-3-containing bodies, likely corresponding to larger viral factories. This sub-localization is in sharp contrast to the near perfect colocalization of B2:GFP with DRB2 and DRB4 upon
infection (Figure 4). Altogether our results suggest that HSP70-1 and HSP70-3, contrarily to HSP70, are components of the TRV viral factories. However, in contrast to B2, DRB2 and DRB4, which directly interact with dsRNA, HSP70-1 and HSP70-3 are likely involved in indirect interactions with TRV replication complexes, perhaps via the TRV replicase or other viral or host components. Interestingly, a study using the B2:GFP system observed that dsRNA-containing VRCs constitute only a part of the structures induced by PVX, which in fact also contain viral ssRNA and coat protein (Monsion et al., 2018). The components and activities harbored within these larger “viral factories” are still largely unknown, but the localization patterns of HSP70-1 and HSP70-3 suggests that these proteins associate not only to replication complexes but also to other entities within viral factories.

Next, we tested the localization of an RNA-binding protein present in our IP MS list that was previously shown to associate to plant virus RNA. This protein, known as Binding to ToMV RNA (BTR1, AT5G04430, 7 counts, Figure 2A), was identified through affinity purification of tagged viral RNA and found in vitro to bind to the 5’ region of the (+) polarity RNA of ToMV, a tobamovirus (Fujisaki and Ishikawa, 2008). In our experimental system, BTR1:tRFP localized to numerous cytoplasmic punctate structures at the cell periphery in non-infected cells (Supplemental Figure S4D). Upon TRV-PDS infection, BTR1 relocalized mainly to B2:GFP-labeled VRCs (Figure 5D), while a fraction was maintained at sites similar to those seen in non-infected cells (Supplemental Figure S4D). At high magnification it is possible to see that BTR1 did not strictly and exclusively colocalize with B2:GFP, but could also be seen in the areas surrounding B2:GFP-labeled dsRNA hotspots (Figure 5D). Similar to HSP70-1 and HSP70-3, it is possible that BTR1 associates not only to VRCs but also to other entities within viral factories.

Proteins not previously linked to infection localize at/near replication complexes
Among the potential TRV replication complex-associated proteins identified through IP, we tested three for which we found no specific function in virus process from the literature: a RING/U-box protein (AT3G45570, 50 counts) and Ribosomal Protein S27a (AT1G23410, 82 counts) and NFD2 (Nuclear Fusion Defective 2 – AT1G24450, 12 counts, Figure 2A).

The RING/U-box protein, which we will refer to as RUP1, belongs to the E3 ubiquitin ligase RBR family. The N-terminal half of the protein is homologous to the RNAse H superfamily, followed on the C-terminal half by a RING-type zinc-finger domain and an IBR (In Between Ring fingers) domain. The Ribosomal Protein 27a, here abbreviated as RP27a, is a small protein of 156 amino acids with a ubiquitin domain N-terminal half and a zinc-binding ribosomal protein superfamily C-terminal section. Importantly, all RP27a peptides detected in B2:GFP co-immunoprecipitated samples belong to the ubiquitin domain, and also match with the protein sequences of UBQ1 through UBQ14 (Supplemental Figure S5). This suggests that ubiquitination of one or more proteins present in the immunoprecipitates that possibly associated to TRV replication complexes.

To investigate which protein/s were ubiquitinated, we searched the mass spectrometry dataset for di-glycine footprints, a hallmark of ubiquitination. Interestingly, the only protein found to contain such a feature was RP27a itself, only on lysine-48 (Supplemental Figure S5), suggesting self-ubiquitination and/or the formation of lysine-48 polyubiquitin chains. Given that no other di-glycine footprint was found in our spectrometry dataset, the proteins targeted by these chains may have been below detection level and remain to be identified. Finally, NFD2 was first identified as a factor involved in karyogamy, the fusion of polar nuclei within the central cell of the female gametophyte prior to fecundation and the fusion of the sperm cells’ nuclei with the egg cell and the central cell upon fecundation (Portereiko et al., 2006). This protein, containing an RNAse III domain, has been also described as RNASE THREE-LIKE 4 (RTL4) (Elvira-Matelot et al., 2016).
In healthy cells, tRFP-tagged RUP1 and RP27a displayed a nucleo-cytoplasmic distribution (Supplemental Figure S6A,B), while NFD2 was essentially found in numerous cytoplasmic bodies (Supplemental Figure S6C). Upon TRV infection, all three proteins were clearly re-localized to or near B2:GFP-labeled dsRNA hotspots (Figure 6, Supplemental Figure S6). More precisely, RUP1 and RP27a showed patterns similar to those observed with HSP70-1, with extensive overlap with B2:GFP as seen from the white color in the merged panels (Figure 6A,B). This suggests that RUP1 and RP27a associate not only with VRCs but also to other entities within viral factories.

Interestingly, NFD2 showed a pattern of localization different from the other proteins tested in this work (Figure 6C, Supplemental Figure S6C). Although localization of NFD2 and B2 seemed mutually exclusive, NFD2 being absent from B2:GFP-labeled structures and vice versa, a continuum between B2:GFP-labeled hotspots and NFD2-labeled structures was observed (Figure 6C), suggesting that NFD2 is intimately linked to TRV-induced subcellular entities and was therefore immunoprecipitated. In addition, the complete localization of NFD2 in close proximity to TRV replication complexes was in stark contrast with the perinuclear and cytoplasmic point-form localization of NFD2 in non-infected plants (Supplemental Figure S6C).

**Knock-out of DRB2 potentiates TRV systemic infection in a Dicer-independent manner**

Next, we tested whether genetic knock-out of the candidate genes analyzed would lead to changes in TRV systemic accumulation. To do so, we acquired A. thaliana lines with T-DNA insertions in the genes of interest: *drb2-1, drb4-1* and *drb2-1 drb4-1* (Curtin et al., 2008), *btr1-1* (Fujisaki and Ishikawa, 2008), SALK_078851 (*rup1*), SALK_093933 (*rp27a*), SALK_088253 (*hsp70*), SALK_135531 (*hsp70-1*) and SAIL_178_E10 (*hsp70-
Supplemental Figure S7). No lines carrying insertions in the annotated 5'UTR or coding sequence of NFD2 were found.

We infected ten plants per genotype and harvested the systemically infected leaves 12 days post-infection (dpi), dividing the plants of each genotype into two equal pools. RNA gel blot analysis of the total RNA from these samples revealed that both the single drb2-1 and double drb2-1 drb4-1 mutants showed markedly increased TRV accumulation in systemic leaves compared to control Col-0 plants (Figure 7A,B). A parallel experiment on inoculated leaves 3 dpi showed that none of the mutants tested affected TRV local accumulation (Supplemental Figure S8). A further experiment confirmed increased TRV accumulation in systemic leaves of drb2-1 (Supplemental Figure S9) and in an independent T-DNA mutant, drb2-2 (Sawano et al., 2017). The drb2-2 mutant showed a milder increase in TRV accumulation compared to the drb2-1 mutant, which we attribute to the fact that in drb2-2 the T-DNA insertion is at the 3’ proximal end of the DRB2 ORF. This likely results only in a partial knock out of the DRB2 protein function, as opposed to the drb2-1 T-DNA insertion at the 5’ end of the gene. These results combined suggest that DRB2 could play an antiviral function with respect to systemic infection by TRV. A moderate increase in TRV accumulation was also observed in systemic leaves of btr1-1 mutants (Figure 7A). Similar effect on ToMV accumulation was reported by Fujisaki and Ishikawa (2008).

Considering that major differences in TRV accumulation between mutant lines and Col-0 control were essentially restricted to drb2-1 and drb2-1 drb4-1 lines, we decided to focus our attention on possible antiviral function of DRB2. Since DRB proteins have been shown by several studies to be involved in small RNA biogenesis (Fukudome et al., 2011; Pélissier et al., 2011; Jakubiec et al., 2012; Montavon et al., 2017; Tschopp et al., 2017) and DRB2 genetic knock-out has been shown to impact accumulation of several microRNAs (Eamens et al., 2012), we decided to analyze the viral siRNA (vsiRNA) present in the drb
mutants described above (Figure 7C). RNA gel blot analysis of small RNAs revealed an increase in vsiRNA in the drb2-1 and drb2-1 drb4-1 mutants analyzed. This most likely reflects the increase in TRV genomic RNAs in these samples (Figure 7A), which are substrates for vsiRNA biogenesis. Moreover, DRB2 knock-out didn’t cause any noticeable changes in the vsiRNA size distributions. These observations, overall, lead us to conclude that knock-out of DRB2 (i) positively impacts TRV RNA and vsiRNA steady-state levels and (ii) does not cause changes in the respective contributions of DCL2, DCL3 and DCL4 to this process. These observations are in line with what has been previously observed for TuMV and TSWV (Curtin et al., 2008). Therefore, the increase of TRV systemic accumulation observed in drb2 mutants is likely not due to impaired dicing activity, a step upstream in the RNA silencing pathway that is normally associated to DRB proteins.

DRB2 over-expression drastically reduces the accumulation of various plant RNA viruses

The absence of DRB2 resulting in increased TRV accumulation (Figure 7), we next tested whether AtDRB2 over-expression could negatively impact infection by TRV and possibly by other distantly-related RNA viruses. To this end, agro-infiltrated N. benthamiana leaves transiently expressing DRB2:tRFP or tRFP were mechanically inoculated with the viruses of interest, and three days after infection leaf disks were collected. RNA gel blot analysis revealed that in tissues infected by TRV-PDS (Figure 8A), Tomato bushy stunt virus (TBSV, Figure 8B), Potato virus X (PVX, Figure 8C) and Grapevine fanleaf virus (GFLV, Figure 8D), over-expression of DRB2:tRFP lead to a dramatic decrease in virus accumulation compared to over-expression of tRFP alone. This effect was particularly prominent for TBSV, PVX and GFLV, despite the presence of comparable amounts of DRB2:tRFP (Figure 8E).
Remarkably, confocal microscopy of B2:GFP-expressing *N. benthamiana* leaves transiently over-expressing DRB2:tRFP and infected with TBSV showed that DRB2:tRFP co-localizes with VRCs (Figure 8F) that are structurally different from those produced upon TRV infection (Figure 4B). To confirm that these are indeed TBSV VRCs, which are known to form on peroxisome membranes (Nagy et al., 2016), we generated a clone to express a tRFP-SKL peroxisome marker (Incarbone et al., 2018). Expression of this marker in B2:GFP-expressing *N. benthamiana* leaves subsequently infected with TBSV reveal that B2-labeled VRCs are indeed localized on the surface of peroxisomes, that in infected conditions appear to group into large multi-peroxisome clusters (Figure 8G). These results clearly show that AtDRB2 localizes to VRCs from different viruses and is a broad-ranged and potent antiviral effector.

**DISCUSSION**

Here, we describe an approach for the identification of VRC-associated proteins through the isolation of replicating viral dsRNA during genuine infection, and validated the localization of most of the candidates through a rapid, robust and simple system. We also showed that one of the proteins we identified as associated to viral dsRNA, DRB2, has antiviral activity against several RNA viruses that belong to different taxonomic groups (Koonin et al., 2020): GFLV (phylum: *Pisoviricota*, class: *Pisoniviricetes*, order: *Picornavirales*, family: *Secoviridae*), TRV (phylum: *Kitrinoviricota*, class: *Alsuviricetes*, order: *Martellivirales*, family: *Virgaviridae*), TBSV (phylum: *Kitrinoviricota*, class: *Tolucaviricetes*, order: *Tolivirales*, family: *Tombusviridae*) and PVX (phylum: *Kitrinoviricota*, class: *Alsuviricetes*, order: *Tymovirales*, family: *Alphaflexiviridae*). Although the proof of concept for our approach to identify VRC-associated proteins is established here only for TRV, it should be compatible with any plant virus as long as it is able to produce dsRNA during its replication.
cycle. Importantly, it does not involve as a prerequisite any modification of viral genomes, the production of infectious clones or the specific tagging of viral protein. Also, considering that the isolation of viral dsRNA and associated proteins is achieved indirectly by anti-GFP antibodies, there is no requirement for virus- or dsRNA-specific antibodies in the process. Hopefully this experimental approach will provide future investigators with a universal tool to successfully explore the proteome associated to the replication complexes of their favorite RNA virus, which can then be studied more in detail to discover the function of VRC-associated proteins and their involvement in the viral life cycle. As hosts, 35S:B2:GFP/A. thaliana (this study) and 35S:B2:GFP/N. benthamiana (Monsion et al., 2018) are compatible with numerous plant virus species. If needed, the systems could be easily adapted to other plant species, as long as they accommodate stable transformation.

We have shown that ectopic expression of B2:GFP greatly increases the accumulation of TRV RNA both in A. thaliana and N. benthamiana. Given the activity of the 73 amino-acid double-stranded binding domain of B2 as a VSR (this work), it is tempting to ascribe TRV over-accumulation simply as a consequence of RNA silencing suppression and subsequent enhanced viral replication. While this is probably the case, it cannot be excluded that B2:GFP increases TRV accumulation by RNAi-independent means, such as stabilization of dsRNA or its protection from other host defensive pathways (Li and Wang, 2019).

The drastic effect of B2:GFP on TRV infection can be viewed as a double-edged sword in relation to its use as bait to pull down VRCs. On the one hand, this effect may introduce biases of both quantitative and qualitative nature, such as the unspecific association to VRCs of host proteins that do not play a role during infection in wild-type conditions or changes in the accessibility or protein complement of replicating RNA, for example. On the other hand, the over-accumulation of TRV dsRNA constitutes a real advantage for the study of VRCs. In fact, increased viral replication is in favor of (i) a better immunoprecipitation
efficiency, (ii) an enhanced detection of protein partners by mass spectrometry and (iii) an improved visualization of VRCs with test candidates. While these biases clearly need to be taken into account, we strongly believe that overall this approach has far more advantages than drawbacks.

The abundance of TRV replicase detected in the IPs (624 reads, Figure 2A) is, in our opinion, confirmation of the robustness of the experiment in terms of VRC yield and integrity. The abundant detection of the coat protein suggests that it either plays a direct role in TRV replication or that the VRCs present in the IPs contain not only full-length dsRNA, but also (+)ssRNA that is being encapsidated, possibly during or just after separation from the (-) strand. However, despite the use of detergent during the IPs, it is possible that we have pulled down proteins present on membranes or complexes close to the replication organelles but not actually part of them. Furthermore, B2 can potentially bind dsRNA generated by secondary structures in viral ssRNA as well as dsRNA produced by host RDR proteins. Future work involving B2 IPs combined with RNA sequencing in WT Arabidopsis as well as mutants for RDR genes, will shed further light on the issue.

Remarkably, among the nine candidate A. thaliana proteins detected following B2 IP and tested in this work, only one, HSP70, failed to accumulate in VRCs despite a high spectral count. At this stage, one cannot strictly rule out the possibility that HSP70, in contrast to HSP70-1 and HSP70-3, corresponds to a false positive despite being enriched in the IPs. It is also possible that tRFP could have disrupted the function of the protein or that the A. thaliana HSP70 (AT3G12580) may not be fully functional when expressed in the heterologous host N. benthamiana. It should be noted however that HSP70 has been linked to viral infection in a number of studies (Verchot, 2012; Nagy, 2016) and found to directly bind the viral replicase of at least two viruses (Nishikiori et al., 2006; Serva and Nagy, 2006). Also, Arabidopsis mRNA-interacting proteome includes members of the Hsc70/Hsp70 family.
well in agreement with the RNA-binding functions of the human Hsp70 that was shown to be independent of the protein chaperone activity (Kishor et al., 2017). This raises the possibility that HSP70 family members may associate with viral factories via RNA binding.

All remaining 8 candidates were specifically redistributed upon infection, suggesting involvement of these factors in the viral life cycle. Their localization patterns can be divided into three groups: perfect co-localization (DRB2 and DRB4), partial co-localization (HSP70-1, HSP70-3, BTR1, RUP1 and RP27a) and proximity (NFD2). Perfect co-localization most likely reflects the direct association of DRB2 and DRB4 on replicating dsRNA within the VRCs. This result is in line with the experimentally verified ability of DRB2 and DRB4 to bind dsRNA (Kobayashi et al., 2009; Tschopp et al., 2017) and of DRB4 to bind TYMV dsRNA in vivo (Jakubiec et al., 2012). DRB2 in A. thaliana re-localizes to cytoplasmic punctate bodies upon infection by TuMV, TSWV and TYMV (Barton et al., 2017) and DRB4 re-localizes from nuclei to cytoplasmic VRCs upon TYMV infection (Jakubiec et al., 2012). While DRB4 plays a role in antiviral defense (Qu et al., 2008; Jakubiec et al., 2012) as part of the RNA silencing machinery, the function of DRB2 recruitment to replication complexes remains to be uncovered.

Although additional experiments are required to confirm the direct association of DRB2 and DRB4 to TRV replicating dsRNA and DRB2 to TBSV replicating dsRNA, our data suggest that host proteins including antiviral defense protein such as DRBs may have access to viral dsRNA within replication organelles including TBSV-induced spherules. This potentially questions the suggested function (or at least efficiency) of replication organelles as protective structures against degradation by cellular RNases and detection by putative dsRNA sensors that trigger antiviral responses (Romero-Brey and Bartenschlager, 2014; Nagy et al., 2016; Fernandez de Castro et al., 2017). It is conceivable that B2:GFP and the DRB proteins
gain access to viral dsRNA at early stage of replication organelle morphogenesis before replication complexes become eventually fully protected.

While the precise molecular pathways linking DRB2 to VRCs remain to be uncovered, we have shown through genetic ablation and over-expression that this protein is a key element in the host’s restriction of viral systemic infection. We have also shown that the antiviral activity of DRB2 likely does not involve Dicer function, since viral siRNA production remains unchanged upon knock-out of DRB2. Our data, however, does not rule out a possible involvement of DRB2 in steps of the RNA interference pathway that are downstream of Dicer processing. Whatever the molecular mode of action of Arabidopsis DRB2, our over-expression experiments have shown that heightened production of this protein in planta drastically reduces the accumulation of viruses belonging to various families. In contrast with our observations, a recent study has shown that the DRB2B protein of N. benthamiana, when over-expressed, strongly increased PVX accumulation (Fatyol et al., 2020). While in this study no other virus species were tested, these differences between A. thaliana DRB2 and N. benthamiana DRB2B suggest that the functions of these gene products in relation to RNA viruses are likely not conserved. However, it is also possible that another of the DRB2 homoeologs present in the allotetraploid N. benthamiana genome, and not tested in the aforementioned study, could have the antiviral effect observed for A. thaliana DRB2. In light of our results, we believe that further study of Arabidopsis DRB2 and its use as a biotech tool in crop defense against viral infection hold substantial potential.

The pattern of partial co-localization, observed for HSP70-1, HSP70-3, BTR1, RUP1 and RP27a/Ubiquitin, consisted in the localization at the B2-labeled VRCs per se, as well as features in close proximity, generally designated as “viral factories”. In the case of PVX, the dsRNA-containing replication complexes reside within larger viral factories harboring other viral proteins and viral ssRNA (Monsion et al., 2018). In general, these viral factories are
most likely the hub for a plethora of viral activities beyond RNA replication \textit{sensu stricto}, such as translation, encapsidation, etc…, and which likely require specific host-encoded proteins. Our work suggests that HSP70-1, HSP70-3, BTR1, RUP1 and RP27a/UBiquitin may play such functions during replication of TRV and possibly other viruses. Indeed, HSP70-3 and BTR1 have been shown to interact with TuMV replicase (Dufresne et al., 2008) and ToMV ssRNA (Fujisaki and Ishikawa, 2008), respectively. Similarly, ubiquitin and the ubiquitin pathway have been shown in a number of studies to play important roles in plant virus life cycle, both pro-viral and anti-viral, the details of which are exhaustively reviewed in (Alcaide-Loridan and Jupin, 2012; Verchot, 2016). Concerning NFD2, the pattern of proximity suggests that this protein may indirectly be involved in viral factory function without direct association with viral dsRNA \textit{per se}. The fact that genetic knock-out of most of these factors did not lead to drastic changes in viral RNA accumulation (with the notable exception of DRB2) does not rule out their involvement in viral functions despite their localization to VRCs. They could act redundantly with other proteins, or could affect parameters that do not perturb viral RNA accumulation, to name but a few possibilities. The genetic dissection of the roles played by the proteins here identified, through experiments including IP and mass spectrometry of tagged alleles of these factors in different genetic backgrounds, is outside the scope of this manuscript and will be addressed in further studies.

Finally, we have previously shown that B2:RFP can be used to mark \textit{Drosophila C virus} (DCV) replication complexes in insect cells \textit{in vivo} (Monsion et al., 2018), suggesting that B2:GFP could be used as a tool to pull down replication complexes of RNA viruses in organisms other than plants. Since dsRNA is a key replication intermediate of all RNA viruses, we believe the further development and adaptation of this tool also holds great potential for the investigation of RNA viruses infecting humans.
MATERIALS AND METHODS

Plant material and growth conditions

Transgenic 35S:B2:GFP/Nicotiana benthamiana plants were previously described (Monsion et al., 2018). Transgenic Arabidopsis thaliana plants (Col-0 line and genetic backgrounds including mutants of the core antiviral Dicer-Like genes, dcl2-1, dcl4-2 and triple dcl2-1 dcl3-1 dcl4-2 lines (Deleris et al., 2006) expressing 35S:B2:GFP were generated using same plasmid (pEAQΔP19-B2:GFP) and agrobacteria described in (Monsion et al., 2018), following floral dip transformation (Harrison et al., 2006) with addition of Plant Preservative Mixture (Plant Cell Technology) at 2 ml/L in Murashige and Skoog (MS) medium. Individual Arabidopsis transformed lines were self-pollinated to generate (F3) plants homozygous for the transgene. T-DNA insertion in the genes of interest were confirmed in the SALK/SAIL mutant lines through standard PCR-based genotyping, using primers listed in Supplemental Table S1. Refer to Supplemental Figure S7 for T-DNA insertion sites and results of PCR-based genotyping.

N. benthamiana plants were grown in a greenhouse at 22-18°C, 16-h/8-h light/dark photoperiod, while A. thaliana were grown in a neon-lit growth chamber at 22-18°C, 12-h/12-h light/dark photoperiod.

Golden Gate pEAQΔP19 vector construction

Binary vector pEAQΔP19-GG was obtained by, (i) removing 3 SapI restriction sites present in pEAQ-HT (Sainsbury et al., 2009), (ii) inserting a Golden Gate cassette (similar to Gateway without AttR1/2) with SapI sites at extremities and (iii) removing P19. Two silent substitutions into Neomycin phosphotransferase (nptII) gene and one substitution near the origin of replication (ColE1) were produced by PCR mutagenesis using Phusion polymerase
in GC buffer supplemented with 5% DMSO (primers in Supplemental Table S1, n°595-596 and 638-641) in order to obtain plasmid pEAQ-HT-ΔSapI. A Golden Gate cassette amplicon (pEAQ-HT as matrix, see primer n°589+642 Supplemental Table S1) was inserted via AgeI/XhoI restriction sites in pEAQ-HT-ΔSapI. Finally, P19 was excised by double restriction EcoNI/SgsI (FD1304, FD1894, Thermo Scientific), extremities were filled in with Klenow fragment (EP0051, Thermo Scientific), supplemented with dNTPs, followed by a ligation step and transformation in E. coli (ccdB Survival strain, Invitrogen).

Cloning of candidate genes

Candidate genes were amplified from A. thaliana genomic DNA with primers designed to contain SapI restriction sites compatible with Golden Gate cloning (Engler et al., 2009) and adapters necessary for ligation to an N-terminal or C-terminal tag (primer list in Supplemental Table S1). In the case of genes containing SapI restriction sites, silent mutations were introduced to remove these sites through overlap PCR. In parallel, tRFP was amplified with primers designed to contain SapI restriction sites, adapters for ligation to the N- or C-terminal end of the candidate gene and a peptide linker (GGGSGGG amino acid sequence) between tRFP and the candidate gene. tRFP-SKL was generated by adding the bases to encode the SKL tripeptide in the reverse primer before the stop codon. PCR products were purified from agarose gel and used in a Golden Gate reaction containing the candidate gene, tRFP, binary vector pEAQΔP19-GG, SapI (R0569L, New England Biolabs), CutSmart buffer (New England Biolabs), T4 DNA ligase 5U/µl (EL0011, Thermo Scientific) and 0.5 or 1 mM ATP (R0441, Thermo Scientific). Golden Gate reaction cycling: 10 cycles of 37°C 10 min, 18°C 10 min ; 18°C 50 min, 50°C 10 min, 80°C 10 min. Following transformation in E. coli (TOP10 strain, Invitrogen), purification and sequencing, plasmids were transformed into A. tumefaciens strain GV3101. pEAQΔP19-B2:RFP served as matrix to generate plasmid
pEAQΔP19-B2mut:RFP as described in (Monsion et al., 2018)(primers n°631-632, Supplemental Table S1).

**Plant inoculation, infection and sampling**

For fluorescence microscopy experiments, leaves of 5- to 6-week-old 35S:B2:GFP/N. benthamiana were infiltrated with A. tumefaciens GV3101 carrying plasmid pEAQΔP19 containing the tagged gene of interest, at absorbance$_{600nm}$ ($A_{600}$) of 0.2. Prior to inoculation, bacteria were incubated in 10 mM MES pH 5.6, 10 mM MgCl$_2$, 200 μM acetosyringone for 1 hour. TRV infection was initiated upon agro-infection with bacteria carrying plasmids expressing the two viral genomic RNAs (Liu et al., 2002), at $A_{600}$ 0.01 each. This method of virus delivery was chosen because it results in homogenous and ubiquitous infection throughout the inoculated tissue. 3-4 days post-inoculation, leaf disks of 5 mm in diameter were collected, vacuum-infiltrated with water and the abaxial side of the disks observed by confocal microscopy (see below). A. thaliana infection for B2:GFP IP experiments was carried out as for N. benthamiana, with the difference that A. tumefaciens was induced by incubating 5-6 hours in induction medium (10.5 g/L K$_2$HPO$_4$, 4.5 g/L KH$_2$PO$_4$, 1 g/L (NH$_4$)$_2$SO$_4$, 0.5 g/L sodium citrate, 0.1 g/L MgSO$_4$, 0.4% glycerol, 0.1 g/L MES, 200 μM acetosyringone), and bacteria used at $A_{600}$ 0.5 each.

For evaluation of TRV local and systemic accumulation in Arabidopsis mutants, sap rub inoculation was used. Briefly, Col-0 leaves systemically infected with TRV-PDS were ground in liquid N$_2$, homogenized in 50 mM sodium phosphate buffer pH 7.5, cleared by centrifugation for 2 min at 2000g, transferred to a new tube and kept on ice until inoculation. This was carried out by sprinkling celite on 4-5 leaves/plant, dipping a cue tip in the inoculum and gently rubbing it onto the leaves. These were then rinsed with water. Systemically infected leaves (the upper 7-8 leaves of each plant) were harvested 12-13 dpi. Samples used
for IP experiments consisted of pools of at least 10 plants, while samples for assessment of
TRV accumulation consisted of pools of 4-5 plants each. Sampled leaves were immediately
frozen in liquid N\textsubscript{2} and thoroughly ground to a powder in a mortar to ensure homogenous
representation in the subsequent analyses.

For the experiments shown in Figure 8, virus infection was carried out by rub
inoculation: the day following agro-infiltration with 35S::RFP, 35S::RFP-SKL or
35S::DRB2::RFP, the abaxial side of the infiltrated leaves was mechanically inoculated. The
inoculum was obtained by grinding frozen \textit{N. benthamiana} tissues infected with TBSV, TRV-
PDS, PVX-GFP or GFLV in 50 mM sodium phosphate buffered at pH 7 (except for TBSV, at
pH 5.8). For confocal microscopy, samples were prepared as described above. For virus
accumulation assessment, a total of 50-60 leaf disks of 5 mm were taken from inoculated
leaves (3 plants/condition, 4 leaves/plant), frozen and pulverized.

\textbf{Immunoprecipitation}

Immunoprecipitations were performed as previously described (Incarbone et al., 2017), with
minor modifications. 0.15 g of young rosette leaves were ground in liquid nitrogen,
homogenized in a chilled mortar with 1 ml lysis buffer (50 mM Tris–HCl, pH 8, 50 mM
NaCl, 1\% Triton X-100) containing 1 tablet/50 ml of protease inhibitor cocktail (Roche),
transferred to a tube and incubated for 15 min at 4°C on a wheel. Cell debris was removed by
two successive centrifugations at 12000g for 10 min at 4°C, after which an aliquot of
supernatant was taken as input fraction. The remaining extract was incubated with magnetic
microbeads coated with monoclonal anti-GFP antibodies (\textmu MACS purification system,
Miltenyi Biotech, catalog number #130-091-125) at 4°C for 20 min. Sample was then passed
through M column (MACS purification system, Miltenyi Biotech) and an aliquot of the flow-
through fraction was taken. The M column was then washed 2 times with 500 \mu l of lysis
buffer and 1 time with 100 μl of washing buffer (20 mM Tris–HCl, pH 7.5). The beads and associated immune complexes were recovered by removing the M column from the magnetic stand and passing 1 ml Tri Reagent (for subsequent RNA analysis – see dedicated section) or 200 μl hot 1X Laemmli buffer (for protein analysis – see dedicated section). 4X Laemmli buffer was added to input and flow-through fractions before protein denaturation for 5 min at 95°C.

RNA extraction and analysis

RNA from total and immunoprecipitated fractions was performed with Tri-Reagent (Sigma) according to manufacturer’s instructions. Briefly, 0.2 g tissue were ground in liquid nitrogen and homogenized in 1 ml Tri-Reagent, 400 μl of chloroform were added, and sample was thoroughly shaken for 2 min. After 10 min spin at 13000 rpm, 4°C, supernatant was added to at least 1 vol isopropanol (and 1.5 μl glycogen in the case of immunoprecipitated samples - IP) and incubated 1 hour on ice (O/N for IP). After 15 min spin at 13000 rpm, 4°C (30 min for IP), pellet was washed in 80% ethanol, dried and resuspended in water. RNA was analyzed by RNA gel blot (denaturing agarose gel to detect high molecular weight RNA, denaturing PAGE to detect low molecular weight RNA) and dsRNA-protein blot (native agarose gel to detect long double-stranded RNA). In the RNA gel blot, miRNAs were detected through DNA oligonucleotides labeled with γ-32P-ATP using T4 PNK (see Supplemental Table S1). TRV genomic and subgenomic RNAs were detected in the same way, with an oligonucleotide complementary to a part of the 3’UTR sequence common to RNA1 and RNA2. The same was done for TBSV. TRV-PDS-derived siRNAs were detected through PCR-amplified A. thaliana PDS sequence labeled by random priming reactions in the presence of α-32P-dCTP. The same was done to detect PVX and GFLV RNA in the RNA gel.
In the dsRNA-protein gel blot, dsRNA were detected with recombinant Strep-Tagged FHV B2, as previously described (Monsion et al., 2018).

Quantitative RT-PCR was performed in the following manner: RNA samples were treated with DNase (Thermofisher TURBO DNA-free kit) and cDNA was generated with random primers (Thermofisher RevertAid First Strand cDNA synthesis kit), according to manufacturer’s instructions. qPCR was then performed with Roche FastStart Essential DNA Green Master kit on a LightCycler 96, in technical triplicates for each sample and primer combination. Data was analyzed with standard ΔΔCt normalization.

Protein extraction and analysis

Proteins from total fractions were extracted as previously described (Hurkman and Tanaka, 1986). Immunoprecipitated proteins for mass spectrometry analysis were isolated as described above, then denatured 5 min at 95°C. Immunoprecipitated proteins from RNA IP were obtained by collecting the phenolic phase following Tri-reagent/chloroform extraction, adding 3-4 vol acetone and incubating at -20°C O/N. After centrifugation (13000 rpm, 15 min, 4°C) pellet was washed in 80% acetone and resuspended in 1X Laemmli. Proteins were resolved by SDS-PAGE and electro-blotted onto Immobilion-P membrane. This was incubated with the appropriate antibodies (anti-GFP polyclonal antibody and anti-tRFP antibody, Evrogen, reference # AB233) and revealed with Roche LumiLight ECL kit after incubation with secondary antibody.

Mass spectrometry analysis and data processing

Proteins were digested with sequencing-grade trypsin (Promega) and analyzed by nanoLC-MS/MS on a TripleTOF 5600 mass spectrometer (Sciex, USA) as described previously (Chicher et al., 2015). Data were searched against the TAIR v.10 database with a decoy
strategy (27281 protein forward sequences). Peptides were identified with Mascot algorithm (version 2.5, Matrix Science, London, UK) and data were further imported into Proline v1.4 software (http://proline.profiproteomics.fr/). Proteins were validated on Mascot pretty rank equal to 1, and 1% FDR on both peptide spectrum matches (PSM score) and protein sets (Protein Set score). The total number of MS/MS fragmentation spectra was used to quantify each protein from at least three independent biological replicates. A statistical analysis based on spectral counts was performed using a homemade R package as described in (Lange et al., 2019). The R package uses a negative binomial GLM model based on EdgeR (Robinson et al., 2010) and calculates, for each identified protein, a fold-change, a p-value and an adjusted p-value corrected using Benjamini-Hochberg method.

Confocal laser scanning microscopy
Observations of leaf disks were carried out using Zeiss LSM700 and LSM780 laser scanning confocal microscopes. eGFP was excited at 488 nm, while tRFP was excited at 561 nm. Image processing was performed using ImageJ/FIJI, while figure panels were assembled with Adobe Photoshop.

ACCESSION NUMBERS

_BTR1_: AT5G04430; _DRB2_: AT2G28380; _DRB4_: AT3G62800; _HSP70_: AT3G12580;

_HSP70-1 (or HSC70-1)_: AT5G02500; _HSP70-3 (or HSC70-3)_: AT3G09440; _NFD2_: AT1G24450; _RP27a_: AT1G23410; _RUP1_: AT3G45570; TRV RdRP: Q9J942; TRV CP: Q88897; TRV 16k: Q77JX3

All data are available in the manuscript and in Supplementary files. Raw data from gels and blots can be found with the blotting source data file, additional confocal acquisitions for each
candidate tested can be found in the microscopy source data file. These are available at the following https://doi.org/10.5281/zenodo.5159940

SUPPLEMENTAL DATA

Supplemental Figure S1: validation of B2 VSR activity and characterization of B2 Arabidopsis lines.

Supplemental Figure S2: immunoblot validation of immunoprecipitates analysed by mass spectrometry.

Supplemental Figure S3: mass spectrometry data analysis using additional controls.

Supplemental Figure S4: lower magnification microscopy of samples in Figure 5.

Supplemental Figure S5: analysis of ubiquitin detected by mass spectrometry.

Supplemental Figure S6: lower magnification microscopy of samples in Figure 6.

Supplemental Figure S7: validation of T-DNA mutants.

Supplemental Figure S8: TRV accumulation in inoculated leaves.

Supplemental Figure S9: TRV accumulation in two drb2 mutants.

Supplemental Table S1: List of primers and probes used in this study.

Supplemental Data Set S1: List of proteins detected by mass spectrometry in GFP pull-downs from 35S:GFP/Col-0 and 35S:B2:GFP/Col-0 plants infected with TRV-PDS.

Supplemental Data Set S2: List of proteins detected by mass spectrometry in GFP pull-downs from 35S:GFP/Col-0 and 35S:B2:GFP/Col-0 plants in both non-infected and TRV-infected conditions.

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**COMPETING INTERESTS**

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

**AUTHOR CONTRIBUTIONS**
Study conception and design: M.I. and C.R.; generation of transgenic A. thaliana lines: B.M.; genotyping: M.I. and E.V.; immunoprecipitation experiments: M.I.; RNA and protein extraction, northern and north-western blotting: M.I. and M.C.; western blotting: M.I., M.C. and H.S.; mass spectrometry: L.K.; statistical analysis of co-IP data: L.K. and H.S.; molecular cloning: M.I., B.M. and V.P.; recombinant B2 production: V.P.; plant inoculation and infection: M.I., M.C. and E.V.; laser confocal microscopy: M.I.; data analysis: M.I., M.C., B.M., L.K., H.S., V.P., P.D. P.G., C.R. writing: M.I. and C.R.; supervision: C.R.; funding acquisition: C.R., P.G. and P.D.

FIGURE LEGENDS

Figure 1: Immunoprecipitation of B2:GFP allows the isolation of TRV dsRNA in vivo. (A) Photos of A. thaliana Col-0 and 35S:B2:GFP/Col-0 plants 13 days post-infection with TRV-PDS. (B) Confocal microscopy analysis of non-infected (n.i. - top left) and TRV-PDS systemically-infected leaves of 35S:B2:GFP/Col-0 plants. On the right, higher magnification (63x) images of TRV replication complexes from the same tissues as those visible at lower magnification (20x) on the left middle and bottom. (C) RNA gel blot analysis of high molecular weight RNA from total fractions of TRV-PDS-infected wild-type or 35S:B2:GFP N. benthamiana (left), and from total (middle) and anti-GFP immunoprecipitated (right) fractions from infected 35S:GFP and 35S:B2:GFP/Col-0 A. thaliana. (D) DsRNA-protein gel blot analysis of native-state high molecular weight RNA from samples described in (C). EtBr staining was used as loading control in (C) and (D). Note the absence of detectable levels of dsRNA species in the non-infected WT plants in agreement with previous similar analyses (Monsion et al., 2018). (E) RNA gel blot analysis of low molecular weight RNA from samples described in (C). The probes were applied sequentially on the same membrane in successive rounds of probing and stripping. snU6 and miR159 were used as loading controls.
Immunoblot analysis of proteins from the same immunoprecipitation experiment analyzed in (C).

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Figure 3: tRFP does not re-localize to TRV replication complexes. Laser confocal microscopy of 35S:B2:GFP/>N. benthamiana leaves transiently expressing 35S:tRFP. (A) Acquisition from non-infected leaf disks (20x objective). Scale bars indicate 50 μm. (B)
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**Figure 4:** *A. thaliana* double-stranded RNA-binding proteins localize at TRV replication complexes. Laser confocal microscopy on 35S:drb2::GFP/N. benthamiana leaves transiently expressing 35S:DRB2::tRFP (A,B) or 35S:DRB4::tRFP (C,D). (A) Acquisitions with 20x objective of non-infected (top) and TRV-PDS-infected (bottom) leaf disks expressing DRB2::tRFP. Scale bars indicate 50 μm. (B) Acquisitions with 63x objective of TRV-PDS-infected leaf disks of tissue described in (A). Scale bars indicate 10 μm. (C,D) As in (A,B), but from tissue expressing DRB4::tRFP.

**Figure 5:** Proteins previously implicated in viral life cycle localize at or near the replication complexes. Laser confocal microscopy on 35S:drb2::GFP/N. benthamiana TRV-PDS-infected leaf disks transiently expressing (A) 35S:HSP70::tRFP, (B) 35S:HSP70-1::tRFP, (C) 35S:HSP70-3::tRFP, (D) 35S:BTR1::tRFP. Acquisitions in (A): 20x objective, scale bars indicate 50 μm. Acquisitions in (B, C, D): 63x objective, scale bars indicate 10 μm.

**Figure 6:** Localization of previously undescribed proteins at or near the replication complexes. Laser confocal microscopy (63x objective) on 35S:drb2::GFP/N. benthamiana TRV-PDS-infected leaf disks transiently expressing (A) 35S:rup1::tRFP, (B) 35S:tRFP:rp27a, (C) 35S:tRFP:nfd2. Scale bars indicate 10 μm.

**Figure 7:** Knock-out of DRB2 causes increased systemic accumulation of TRV in Arabidopsis, through a mechanism independent from small RNA biogenesis. (A,B) RNA
gel blot analysis of RNA from systemically infected leaves of Arabidopsis knock-out lines infected with TRV-PDS, 12 days post-infection (dpi). Previously published mutants are indicated with their current name, while the others are indicated with their SALK/SAIL nomenclature. Each sample is a pool of 4-5 plants, and two samples were analyzed per genotype (1 and 2), per time point. EtBr (A) or methylene blue (B) staining were used as loading control. (C) PAGE RNA gel blot analysis of small RNA from the corresponding samples in (B). snU6 and miR159 were used as loading controls. n.i.: non-infected.

Figure 8: Over-expression of DRB2 in N. benthamiana leaves drastically reduces accumulation of a wide range of RNA viruses. (A) Northern blot analysis of RNA from N. benthamiana leaf disks 4 days after transient transformation with 35S:tRFP or 35S:DRB2:tRFP and 3 dpi with TRV-PDS (except for n.i.: non-infected). Each sample is a pool of 40-50 leaf disks from 4-5 leaves. In the case of the virus-infected leaves, two samples were analyzed per condition (indicated with 1 and 2). Methylene blue staining of the membrane was used as loading control. (B) As in (A), but after rub-inoculation with tomato bushy stunt virus (TBSV). Two independent biological replicates are shown with either methylene blue or EtBr staining of the membranes as loading control. (C) As in (A), but after rub-inoculation with Potato virus X (PVX). (D) As in (A), but after rub-inoculation with Grapevine fanleaf virus (GFLV). (E) Immunoblot analysis on protein extracts from the samples analyzed in (A-D), to detect DRB2:tRFP (top arrowhead) and tRFP (bottom arrowhead). Coomassie blue staining was used as loading control. Source data is available with the Blotting Source Data. (F) Laser confocal microscopy acquisitions of B2-labeled TBSV replication complexes, from 35S:B2:GFP/N. benthamiana plants transiently expressing DRB2:tRFP and infected with TBSV. Scale bars indicate 50 (top) or 10 μm (middle and bottom). (G) As in (F), but from plants (non-infected in the top row, TBSV-
infected in the rest) transiently expressing the peroxisome marker tRFP-SKL. Scale bars indicate 50 (top two acquisitions) or 10 μm (bottom two acquisitions).

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Figure 4: *A. thaliana* double-stranded RNA-binding proteins localize at TRV replication complexes. Laser confocal microscopy on 35S:B2:GFP/N. benthamiana leaves transiently expressing 35S:DRB2:tRFP (A,B) or 35S:DRB4:tRFP (C,D). (A) Acquisitions with 20x objective of non-infected (top) and TRV-PDS-infected (bottom) leaf disks expressing DRB2:tRFP. Scale bars indicate 50 μm. (B) Acquisitions with 63x objective of TRV-PDS-infected leaf disks of tissue described in (A). Scale bars indicate 10 μm. (C,D) As in (A,B), but from tissue expressing DRB4:tRFP.
Figure 5: Proteins previously implicated in viral life cycle localize at or near the replication complexes. Laser confocal microscopy on 35S:B2:GFP/N. benthamiana TRV-PDS-infected leaf disks transiently expressing (A) 35S:HSP70:ΔRFP, (B) 35S:HSP70-1:ΔRFP, (C) 35S:HSP70-3:ΔRFP, (D) 35S:BTR1:ΔRFP. Acquisitions in (A): 20x objective, scale bars indicate 50 μm. Acquisitions in (B, C, D): 63x objective, scale bars indicate 10 μm.
Figure 6: Localization of previously undescribed proteins at or near the replication complexes. Laser confocal microscopy (63x objective) on 35S:B2:GFP/N. benthamiana TRV-PDS-infected leaf disks transiently expressing (A) 35S:RUP1:tRFP, (B) 35S:tRFP:RP27a, (C) 35S:tRFP:NFD2. Scale bars indicate 10 μm.
Figure 7: Knock-out of DRB2 causes increased systemic accumulation of TRV in Arabidopsis, through a mechanism independent from small RNA biogenesis. (A,B) Northern blot analysis of RNA from systemically infected leaves of Arabidopsis knock-out lines infected with TRV-PDS, 12 days post-infection (dpi). Previously published mutants are indicated with their current name, while the others are indicated with their SALK/SAIL nomenclature. Each sample is a pool of 4-5 plants, and two samples were analyzed per genotype (1 and 2), per time point. EtBr (A) or methylene blue (B) staining were used as loading control. (C) PAGE northern blot analysis of small RNA from the corresponding samples in (B). snU6 and miR159 were used as loading controls. N.i.: non-infected.
Figure 8: Over-expression of DRB2 in *N. benthamiana* leaves drastically reduces accumulation of a wide range of RNA viruses. (A) Northern blot analysis of RNA from *N. benthamiana* leaf disks 4 days after transient transformation with 35S::RFP or 35S::DRB2::RFP and 3 dpi with TRV-PDS (except for n.i.: non-infected). Each sample is a pool of 40-50 leaf disks from 4-5 leaves. In the case of the virus-infected leaves, two samples were analyzed per condition (indicated with 1 and 2). Methylene blue staining of the membrane was used as loading control. (B) As in (A), but after rub-inoculation with tomato bushy stunt virus (TBSV). Two independent biological replicates are shown with either methylene blue or EtBr staining of the membranes as loading control. (C) As in
(A), but after rub-inoculation with Potato virus X (PVX). (D) As in (A), but after rub-inoculation with Grapevine fanleaf virus (GFLV). (E) Western blot analysis on protein extracts from the samples analyzed in (A-D), to detect DRB2:tRFP (top arrowhead) and tRFP (bottom arrowhead). Coomassie blue staining was used as loading control. Source data is available with the Blotting Source Data. (F) Laser confocal microscopy acquisitions of B2-labeled TBSV replication complexes, from 35S:B2:GFP/N. benthamiana plants transiently expressing DRB2:tRFP and infected with TBSV. Scale bars indicate 50 (top) or 10 μm (middle and bottom). (G) As in (F), but from plants (non-infected in the top row, TBSV-infected in the rest) transiently expressing the peroxisome marker tRFP-SKL. Scale bars indicate 50 (top two acquisitions) or 10 μm (bottom two acquisitions).
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