



Transmission of *Turnip yellows virus* by *Myzus persicae* Is Reduced by Feeding Aphids on Double-Stranded RNA Targeting the Ephrin Receptor Protein

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Michaël Mulet^{1†}, Baptiste Monsion^{1†}, Sylvaine Boissinot¹, Maryam Rastegar^{1,2}, Sophie Meyer¹, Nicole Bochet¹ and Véronique Brault^{1*}

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*Correspondence:

Véronique Brault
veronique.brault@inra.fr

†These authors have contributed
equally to this work.

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¹SVQV, Université de Strasbourg, Institut National de la Recherche Agronomique, Colmar, France, ²Department of Plant Protection, Shiraz University, Shiraz, Iran

Aphid-transmitted plant viruses are a threat for major crops causing massive economic loss worldwide. Members in the *Luteoviridae* family are transmitted by aphids in a circulative and non-replicative mode. Virions are acquired by aphids when ingesting sap from infected plants and are transported through the gut and the accessory salivary gland (ASG) cells by a transcytosis mechanism relying on virus-specific receptors largely unknown. Once released into the salivary canal, virions are inoculated to plants, together with saliva, during a subsequent feeding. In this paper, we bring *in vivo* evidence that the membrane-bound Ephrin receptor (Eph) is a novel aphid protein involved in the transmission of the *Turnip yellows virus* (TuYV, *Polerovirus* genus, *Luteoviridae* family) by *Myzus persicae*. The minor capsid protein of TuYV, essential for aphid transmission, was able to bind the external domain of Eph in yeast. Feeding *M. persicae* on *in planta*- or *in vitro*-synthesized dsRNA targeting *Eph*-mRNA (dsRNA_{Eph}) did not affect aphid feeding behavior but reduced accumulation of TuYV genomes in the aphid's body. Consequently, TuYV transmission efficiency by the dsRNA_{Eph}-treated aphids was reproducibly inhibited and we brought evidence that Eph is likely involved in intestinal uptake of the virion. The inhibition of virus uptake after dsRNA_{Eph} acquisition was also observed for two other poleroviruses transmitted by *M. persicae*, suggesting a broader role of Eph in polerovirus transmission. Finally, dsRNA_{Eph} acquisition by aphids did not affect nymph production. These results pave the way toward an ecologically safe alternative of insecticide treatments that are used to lower aphid populations and reduce polerovirus damages.

Keywords: polerovirus, virus transmission, virus receptor, RNA interference, transmission inhibition, plant viruses, aphid vector

INTRODUCTION

To circumvent plant immobility, and escape before the plant dies, the majority of plant viruses rely on mobile vectors for their dissemination. Among these vectors, phloem-feeding aphids are by far the most prevalent vectors that can transmit almost half of the insect-borne plant viruses (Hogehout et al., 2008; Dedryver et al., 2010). Different modes of virus transmission have been described. The non-persistent and non-circulative mode of transmission relies on a transient and brief retention of virions at specific sites in the vector's mouthparts, or in close proximity. In contrast, the persistent, circulative and non-propagative mode of transmission requires endocytosis of virions into aphid cells (Ng and Falk, 2006; Hogehout et al., 2008; Ammar el et al., 2009; Blanc et al., 2014; Gray et al., 2014; Drucker and Then, 2015; Whitfield et al., 2015). In the latter mode of transmission, viruses may persist in the aphid's body during the whole insect life with (propagative mode) or without (non-propagative mode) replication.

Members of the *Luteoviridae* family (referred to as luteovirids) are single-stranded RNA positive strand viruses, phloem-limited and strictly transmitted by aphids in a circulative, persistent, and non-propagative mode (Gildow, 1999; Gray and Gildow, 2003; Brault et al., 2007). Luteovirid transmission is highly specific because each virus species is usually transmitted efficiently by only one or a few aphid species (Herrbach, 1999). Luteovirid particles are acquired during the prolonged sap ingestion by aphids while feeding on infected plants. Virions are then transported through the gut cells via a transcytosis mechanism that is thought to be initiated by clathrin-mediated endocytosis (Gildow, 1999; Brault et al., 2007). Virus uptake into intestinal cells occurs either at the posterior midgut, the hindgut or both, depending on the virus species (Brault et al., 2007). Once released into the hemocoel, virions may be protected from degradation by binding to symbionin, an endosymbiotic protein, but this interaction, as well as its implication in luteovirid transmission, remains controversial (van den Heuvel et al., 1994, 1997; Filichkin et al., 1997; Liu et al., 2009; Bouvaine et al., 2011; Cilia et al., 2011). From there, luteovirid particles reach the accessory salivary glands (ASG) and are transported through the ASG cells by a transcytosis mechanism before being released into the salivary canal (Brault et al., 2007). These transcytosis events are suspected to rely on the presence of membrane virus-specific receptors at the gut and ASG levels. The apical plasmalemma of the intestinal cells together with the basal lamina and the basal plasmalemma of the ASG cells have been identified as luteovirid transmission barriers in aphids suggesting that specific interactions between virus structural proteins and cellular partners must exist at these locations to allow virus transmission (Gildow, 1999).

Luteovirid capsids are composed of two structural viral proteins namely the major coat protein (CP of about 22 kDa) and the minor capsid protein (readthrough protein or RT* of about 55 kDa) which is not required for virus particle assembly. Both proteins are involved in aphid transmission. Some mutations in the CP sequence of luteovirids affected aphid transmission without impacting virion formation (Torrance, 1992; Brault

et al., 2003; Kaplan et al., 2007; Doumayrou et al., 2016). Point mutations or deletions in the RT* are deleterious for efficient transport of virions through the gut cells (Gildow et al., 2000; Reinbold et al., 2001) and for virus uptake into the ASG cells (Brault et al., 1995, 2000; Chay et al., 1996; Bruyère et al., 1997; Peter et al., 2008). Moreover, RT* is responsible for luteovirid vector specificity (Brault et al., 2005).

Identifying luteovirid cellular partners, and in particular virus receptors in aphids, is a major challenge that could ultimately result in the development of innovative technologies aimed at inhibiting virus transmission. Up to now only insecticide treatments and aphid- or virus-tolerant or -resistant plants can be deployed to control luteovirid diseases (Walkey and Pink, 1990; Dogimont et al., 1996; Barker and Waterhouse, 1999; Dreyer et al., 2001). The gut membrane protein alanyl aminopeptidase N (APN), was identified previously as a potential receptor of pea enation mosaic virus (PEMV, *Enamovirus* genus, *Luteoviridae* family) in the aphid species *A. pisum* using an array of *in vitro*-based techniques and insect cells experiments (Linz et al., 2015). APN was isolated from a phage display peptide screen and evidence of its role in PEMV transmission by *A. pisum* was provided by competition experiments between the virus and a peptide potentially mimicking the viral determinant binding to the aphid receptor (Liu et al., 2010). Other aphid proteins exhibiting the ability to bind purified luteovirids *in vitro* have been reported but their precise role in virus transmission has not been identified. This includes several proteins extracted from *Myzus persicae* or heads of *Sitobion avenae* which exhibited the capacity to bind to virions of *Turnip yellows virus* (TuYV), previously designated *Beet western yellows virus* (Seddas et al., 2004), or of *Barley yellow dwarf virus* (BYDV) (Li et al., 2001). Another uncharacterized protein located in the ASG cells of *S. avenae* and *Schizaphis graminum* has also been suspected to be involved in luteovirid transmission, as acquisition of antibodies directed against this protein together with BYDV greatly reduced virus transmissibility (Wang and Zhou, 2003). Finally, by coupling quantitative proteomics with aphid genetics, several proteins from *S. graminum*, including a luciferase and a cyclophilin-like protein, were associated with the ability to transmit *Cereal yellow dwarf virus* (CYDV) (Yang et al., 2008). Subsequently, differential gel electrophoresis (DIGE) coupled to mass spectrometry on an F2 population originating from a cross between vector and non-vector biotypes of *S. graminum* exhibiting different barriers to transmission of CYDV-RPV (gut or ASG) revealed eight proteins under-represented in genotypes with a strong gut barrier (Cilia et al., 2011). Genetics studies of BYDV and CYDV transmission by aphids inferred that transmission capacity is a multigenic trait with some of the aphid genes being shared by the two viruses and some others being specific for one virus species (Papura et al., 2002; Dedryver et al., 2005; Burrows et al., 2006, 2007). Importantly, the aphid proteins predicted to function at specific transmission barriers were expressed as two isoforms with distinct charges (Papura et al., 2002; Cilia et al., 2011). Although the aforementioned studies have identified potential luteovirid partners in aphids, they did not link these proteins to the virus aphid-transmission phenotype.

We performed a yeast two-hybrid screen of a total *M. persicae* cDNA library to identify interactions between aphid cellular components and the structural proteins of the *Cucurbit aphid borne yellows virus* (CABYV) (*Polerovirus* genus, *Luteoviridae* family), which is transmitted efficiently by *M. persicae*. The membrane ephrin receptor (Eph) was identified as a potential binding partner of the RT*s of CABYV and TuYV, another polerovirus efficiently transmitted by this aphid species. Functional validation tests were conducted with TuYV in *M. persicae* using techniques based on RNA interference. Feeding aphids on various sources of dsRNA targeting *Eph*-mRNA resulted in reduced internalization of TuYV genomes into the aphid's body and reduced transmission of TuYV, without affecting aphid's fitness. Taken together, these results implicate Eph in the transmission process of TuYV and suggest involvement of this protein in transmission of other poleroviruses by *M. persicae*.

MATERIALS AND METHODS

Aphid Library Construction

Total RNA was isolated from 35 mg of all instars of *M. persicae* using the RNeasy Plant Mini Kit (Qiagen) following the RNeasy Fibrous Tissue Mini Kit protocol. The purification of poly A⁺ RNA from total RNA was performed with the Oligotex[™] mRNA Mini Kit (Qiagen) following the Batch protocol. The reverse transcription was performed starting from 1 μg of mRNA using an Oligo(dT)₂₀ primer with an adaptor extension and following the MMLV High Performance Reverse Transcriptase procedure (EPICENTRE[®] BIOTECHNOLOGIES). The protocol was modified by the addition of a template switching primer (adaptor) after 30 min of incubation for a 3' cDNA extension (Table S1). The cDNA molecules were amplified using a single adaptor primer with the GoldStar[®] DNA polymerase (Eurogentec). The cycles were as follows: 95°C 1 min, 95°C 15 s, 65°C 20 s, 72°C 3 min (23 cycles), and 65°C 20 s, 72°C 6 min. The cDNA fragments were further purified with QIAquick PCR Purification Kit (QIAGEN) and subjected to an over-night digestion at 50°C with the restriction enzyme SfiI. Alongside, the pGADT7 vector (Clontech) was modified by the introduction of SfiI sites using a specific pair of complementary oligonucleotides with NcoI and EcoRI sites (Table S1) leading to pGADT7-SfiI. This plasmid was digested with SfiI before being dephosphorylated using the Thermosensitive Alkaline Phosphatase (Promega). A short run electrophoresis of SfiI-cDNA digestion products was performed on low melting point agarose gel to collect the DNA fragments above 400 bp. DNA fragments were recovered after hot-phenol extraction and DNA precipitation. Insert ligation into pGADT7-SfiI was performed at 16°C for 8 h with a molar ratio vector:insert of 1:30. The ligation mixture was further introduced by electroporation into *Escherichia coli* XL10-Gold Ultracompetent Cells (Stratagene, Agilent Technologies). After streaking the bacteria on LB agar containing ampicillin for 19 h at 37°C, colonies were collected in liquid LB medium complemented with glycerol solution (25% final) and stored at -80°C. The cDNA library was then amplified by inoculating

100 ml of LB containing ampicillin with an aliquot of the glycerol stock and cultivating the bacteria for 3 h at 37°C. Plasmids were then purified using QIAfilter Plasmid Maxi Kit (QIAGEN) and used in the yeast two hybrid screen.

Viral Constructs for the Yeast Two Hybrid Screen

The major (CP) and the minor (RT*) capsid proteins of TuYV (NC_003743) and CABYV (NC_003688) were used in the yeast two hybrid experiments. The CP sequences of TuYV and CABYV were amplified by PCR (Expand High Fidelity PCR System, Roche Applied Science) using appropriate primers (Table S1) from the full-length viral sequences described in Veidt et al. (1992) and Guilley et al. (1994). To clone the RT* sequence of TuYV and CABYV, two overlapping mutagenic oligonucleotides were used in the PCR reaction together with external primers (Table S1) to replace the CP-stop codon by a tyrosine codon in the TuYV and CABYV sequences. The 5'-terminal nucleotide of TuYV- and CABYV-RT* sequence was positioned, respectively, at nt 4793 and nt 4896 on the viral genomes. After digestion with the appropriate restriction enzymes (Promega), CP and RT* from CABYV were purified on column (MSB[®] Spin PCRapace, Invitex GmbH) and cloned downstream the GAL4 DNA binding domain (BD) into the pGBKT7 vector (Clontech). The CP sequence from CABYV was also introduced downstream the GAL4 activation domain (AD) into the pGADT7 vector (Clontech). In addition, CP, RT* from CABYV together with CP and RT* from TuYV were introduced into pLexA-N vector downstream the LexA binding domain (Dualsystems Biotech). Ligations were performed overnight at 16°C in a 10 μL final volume using a molar ratio vector:insert of 1:5 with the T4 DNA Ligase from Promega. Ligation products were introduced by heat-shock into *E. coli* XL10-Gold competent cells.

The recombinant constructs were referred to as pGBKT7-CP_{CA}, pGADT7-CP_{CA}, pGBKT7-RT*_{CA}, pLexAN-CP_{CA}, pLexAN-RT*_{CA}, pLexAN-CP_{Tu}, and pLexAN-RT*_{Tu}. The pGBKT7-derived plasmids were introduced into the Y2HGold yeast strain and the pLexAN-derived plasmids were introduced into the NMY51 yeast strain. Y2HGold and NMY51 yeast strains contain the reporter genes *HIS3* and *ADE2*.

Yeast Two Hybrid Assays

The *M. persicae* cDNA library was screened against the baits following the procedures described in the DUALhunter kit user manual (Dualsystems Biotech). The colonies were plated onto a stringent medium lacking leucine, tryptophan, histidine, and adenine [-LWHA] and cultivated at 28°C for 3–14 days. The colonies developing on the [-LWHA] medium were selected and the recombinant pGADT7 plasmid containing the aphid cDNA was recovered following a yeast DNA extraction method. Briefly, yeast cells from a 2 ml overnight culture were suspended in 100 μl of a buffer (67 mM Potassium Phosphate, pH 7.5) containing 50 units of Lyticase (L2524, Sigma-Aldrich) and incubated 1 h at 37°C before proceeding with the common alkaline lysis *E. coli* plasmid purification. Recombinant plasmids were introduced into *E. coli* to obtain a sufficient amount of plasmids for sequencing.

Aphid Acquisition of dsRNA From Transgenic *A. thaliana* and From *in Vitro*-Synthesized dsRNA

Arabidopsis thaliana expressing a hairpin RNA targeting *Eph* or *LacZ* as a control (Ara:Hp-*Eph* and Ara:Hp-*LacZ*) were described in Mulot et al. (2016) and were grown in an environment-controlled chamber at 23°C day and 20°C night with a 10 h photoperiod as well as Col-0 non-transformed plants. *In vitro*-synthesized dsRNA targeting *Eph* or *LacZ* (dsRNA_{*Eph*} and dsRNA_{*LacZ*}) were obtained as described previously in Mulot et al. (2016).

M. persicae (Sulzer) colonies were reared on pepper (*Capsicum annuum*) at 20°C with a 16 h photoperiod. Aphids were fed on transgenic *A. thaliana* or artificially on *in vitro*-synthesized dsRNA as described in Mulot et al. (2016) except that the acquisition time on the artificial medium containing the dsRNA was extended to 5 days in some experiments and the final dsRNA concentration in the feeding medium was set up to 400 ng/μl in all experiments. When a 5-day acquisition period was performed, the dsRNA-containing medium was replaced after 3 days by a fresh medium containing the dsRNA.

Virus Transmission by *M. persicae*

In the virus transmission experiments, aphids previously fed for 10 days on transgenic *A. thaliana* (Ara:Hp-*Eph* or Ara:Hp-*LacZ*) were transferred for 24 h on purified TuYV prepared as described in Van den Heuvel et al. (1991). The viral concentration was set up at 25 μg/ml in the artificial diet (Bruyère et al., 1997). Aphids fed artificially on dsRNA were either transferred onto purified virus (same set-up as described above) or on TuYV-infected *M. perfoliata* inoculated by agroinfiltration as described in Hipper et al. (2014). After a 24 h acquisition access period of the virus, two potentially viruliferous aphids were transferred on Col-0 test plants for 72 h. After this inoculation access period, some aphids were collected for further analysis (see below) while the remaining aphids were eliminated by an insecticide treatment. The plants were tested by DAS-ELISA 3 weeks later using virus-specific antibodies as described in Bruyère et al. (1997). In this assay, samples from several young leaves were collected on each plant and pooled before grinding.

Eph-mRNA and Viral RNA Accumulation Analysis in Aphids by Real-Time Reverse Transcription Polymerase Chain Reaction (qRT-PCR)

Total RNA was extracted from whole *M. persicae* (20 aphids per sample) as described in Mulot et al. (2016). Total RNA was also extracted from 100 dissected guts using the RNeasy Plant Mini Kit (QIAGEN) as described in Mulot et al. (2016) or alternatively from 35 dissected guts using NucleoSpin® RNA XS (Macherey-Nagel). To evaluate *Eph*-mRNA accumulation, qRT-PCR was performed as in Mulot et al. (2016). As mentioned in Mulot et al. (2016), the relative expression levels were normalized to *Rpl7* and *L27*. To determine the copy number of TuYV genomes internalized into *M. persicae*, total RNA was extracted from whole aphids after transferring them for 3 days on non-infected Col-0

(inoculation access period or IAP) to clear the gut lumen. The viral RNA was converted into cDNA using the reverse primer BPqtR1 and the M-MLV reverse transcriptase kit (Promega). The forward primer BPqtF0 and the reverse primer BPqtR1 (Table S1) were used to amplify by real-time PCR (qRT-PCR) the cDNA corresponding to nts 3694–3830 on TuYV genomic sequence (accession number NC_003743) using the same set-up as described in Mulot et al. (2016). Alongside, viral RNA genomes were extracted from purified virions using the RNeasy Plant Mini Kit (QIAGEN). After quantification at 260 nm (Nanodrop 2000; Thermo Fischer Scientific), the viral RNA was converted to cDNA as described above. Dilution series of 10⁹ to 10⁴ viral cDNA copies obtained from RNA extracted from purified virions were used to calibrate the CFX cyclor and comparison between calibrate standard Ct values and samples Ct values provided an absolute quantification of TuYV genomes.

Aphid Fecundity and Feeding Behavior Tests

Aphid fecundity after feeding on transgenic *A. thaliana* (Ara:Hp-*Eph* or Ara:Hp-*LacZ*) was assessed by depositing individual fourth instars or adults onto these plants for 2 days. After this period, only one nymph was kept on the plant for 10 days to reach the adult stage before being transferred individually onto non-transformed Col-0 plants. Nymph production was recorded after 5 days. Aphid fecundity was also recorded after feeding fourth instars or adults for 5 days on *in vitro*-synthesized dsRNA. Four aphids were then transferred onto each non-transformed Col-0. Nymph production was monitored during 5 days. A Student *t*-test was applied to the values after controlling that the data followed a linear model.

To evaluate the feeding activity of aphids, fourth instars or adults *M. persicae* fed for 5 days on *in vitro*-synthesized dsRNA, were transferred to an artificial feeding medium (MP148, Harrewijn, 1983) for 48 h. Pools of 9–10 aphids were enclosed in individual boxes (10 or 11 boxes per condition) that were internally covered with a pH-indicator paper prepared in 0.2% bromocresol green dissolved in ethanol. The number of honeydew droplets produced by the aphids and which appear as purple dots on the indicator paper was counted manually or evaluated by image analysis (ImageJ). After determining that the data followed a linear model and variance was equal between samples, a Student *t*-test was applied to the values.

RESULTS

The Ephrin Receptor Protein From *M. persicae* Is a Potential Partner of Proliferating Cell Nuclear Antigen (PCNA) Structural Proteins

In order to identify partners of polerovirus particles in the aphid *M. persicae*, we looked for cellular partners of the structural proteins of CABYV which is efficiently transmitted by this aphid species (Lecoq et al., 1992). An aphid cDNA library was obtained from mRNA extracted from whole aphids. The cDNA library was cloned into the pGADT7 vector and the average insert size was about 400 bp. The major and the minor capsid protein

sequences of CABYV (CP_{CA} and RT_{CA}^{*}) were introduced into the pGBKT7 vector and expressed as fusion proteins with the GAL4 DNA binding domain (BD) in the Y2HGold yeast strain in which the *HIS3* and *ADE2* reporter genes are under the control of the GAL4 promoter. Screening the *M. persicae* cDNA library with the baits was performed by introducing the cDNA library into the yeast cells previously transformed with each bait. 6.2×10^6 and 4.6×10^6 double transformed yeast cells were obtained for the CP_{CA} and RT_{CA}^{*} screens, respectively. When plated onto the [-LWHA] medium to select yeast cells in which *in vivo* interactions occurred, 171 and 5 colonies developed for the CP_{CA} and RT_{CA}^{*} screens, respectively. A similar cDNA prey sequence, encoding a 244 amino acid peptide, was found in 4 of 13 colonies analyzed for the CP_{CA} screen and in 3 of the 5 colonies which emerged from the RT_{CA}^{*} screen. When blasted on the *M. persicae* genome (*M. persicae* clone G006 assembly v2, blast server, Aphidbase.com), one scaffold (MYZPE13164 G006 v1.0 000015980) contained the identified sequence which is annotated as ephrin type-B receptor 1-B (LOC111037473) and referred thereafter in the document as Eph (Figure S1).

Ephrin receptors are activated upon binding to their membrane-associated ephrin ligands and plays important roles in developmental processes in mammalian and in pathological diseases like brain and lungs cancers (for review see Pasquale, 2005; Himanen et al., 2007; Genander and Frisen, 2010; Pitulescu and Adams, 2010; Perez White and Getsios, 2014; Kania and Klein, 2016). Interaction of Ephs with ephrin ligands on the surface of neighboring cells triggers Eph kinase-dependent signaling in a bidirectional process. Ephrin receptor family is divided into two subclasses, EphA and EphB, based on amino acid sequence homology and binding affinities to glycosylphosphatidylinositol (GPI)-linked ephrin-A or transmembrane ephrin-B ligands. Interestingly, Eph or ephrin ligand have been shown to display receptor functions for mammalian viruses, bacteria and protozoan parasites (Bonaparte et al., 2005; Lupberger et al., 2011; Kaushansky et al., 2015; Subbarayal et al., 2015).

The extracellular domain of Ephs contains a globular ligand-binding domain and two fibronectin type III repeats (Figure 1). A short transmembrane domain separates the extracellular part from the intracellular cytoplasmic part consisting of the protein kinase domain and a sterile alpha motif domain responsible for Eph clustering (Stapleton et al., 1999) (Figure 1). The identified candidate peptide from the *M. persicae* cDNA library covers the two fibronectin type III repeats (Figure 1).

To address the ability of the plasmid bearing the partial Eph cDNA sequence (referred to hereafter as pGAD-Eph) to activate by itself the transcription of the reporter genes (*HIS3* and *ADE2*), the pGAD-Eph plasmid was introduced together with the empty pGBKT7 vector into Y2HGold yeast strain in which the reporter genes were controlled by the GAL4-responsive promoter. The yeast double-transformed colonies were plated onto medium lacking histidine [-LWH] or histidine and adenine [-LWHA]. Yeast growth was observed on both medium showing the capacity of the Eph partial domain to activate transcription of the reporter genes in the absence of luteovirid CP or RT^{*}

(Figure S2). Interaction of the Eph domain with the GAL4-promoter can be considered as a false positive reaction, but could also mask a true interaction with the viral baits. Considering the function of Eph as human virus receptors, we pursued the yeast two hybrid binding assays and addressed whether the Eph domain was able to interact with another promoter, the LexA promoter. Interestingly, no autoactivation of the transcription of the reporter genes *HIS3* and *ADE2* was observed when the pGAD-Eph and the empty pLexAN plasmids were introduced into the NMY51 yeast strain in which the reporter genes are under the control of the LexA promoter and when the doubled-transformed cells were plated onto [-LWHA] medium for 7 days (Figure S3). A low yeast growth was however observed, in one out of the three colonies, when the growth was extended to 14 days (Figure 2), implying that a low yeast development should not be considered as a true interaction between the prey and baits.

Therefore, the interactions between CABYV baits and the Eph partial domain were confirmed using the NMY51 yeast strain. The viral structural protein sequences were cloned into the pLexAN yeast vector as fusion proteins with the LexA DNA binding domain (BD). The resulting plasmids referred to as pLexAN-CP_{CA} and pLexAN-RT_{CA}^{*} were introduced into the NMY51 yeast strain together with the pGAD-Eph. Three colonies of the double-transformed yeast cells were plated onto the [-LW] to control yeast growth and on the [-LWHA] stringent medium to select yeast cells in which *in vivo* interaction occurred. We observed that RT_{CA}^{*} was able to interact with the partial domain of Eph since 2 out of 3 colonies developed on the [-LWHA] medium after 14 days of growth. Interaction between the Eph domain and CP_{CA} was less clear as only one out of the 3 colonies grew well on the [-LWHA] medium after 14 days (Figure 2). To control whether Eph could be a potential partner of other poliovirus structural proteins, we tested TuYV, which is also transmitted efficiently by *M. persicae* (Leiser et al., 1992). The TuYV CP and RT^{*} sequences were introduced into the pLexAN vector leading to the pLexAN-CP_{Tu} and pLexAN-RT_{Tu}^{*}. Similarly as described above, the pLexAN recombinant plasmids were introduced into NMY51 yeast cells together with pGAD-Eph and plated onto the [-LWHA] medium. Interestingly, interaction of the Eph domain with the RT_{Tu}^{*} was clearly observed (all the three colonies developed well on the [-LWHA] medium after 14 days of growth) and an interaction with the CP_{Tu} was also suggested since two out of the three colonies grew on the selective medium (Figure 2). The reason for the uneven growth of the three double-transformed colonies on the stringent medium is unknown but is likely due to a tendency of the CP-fusion proteins to self-assemble in yeast (Figure S4) rather than to interact with the prey. No yeast growth was observed on the [-LWHA] medium when the viral pLexAN-derived plasmids were co-transformed with the empty pGAD vector (Figure 2). Binding of Eph domain to the TuYV baits were controlled in an additional experiment (Figure S3). No yeast growth was observed for the control combinations.

In conclusion, we observed an unambiguous interaction between the Eph domain isolated from the *M. persicae* cDNA library and the RT^{*} from TuYV (RT_{Tu}^{*}). In contrast, interaction of Eph domain with the RT^{*} (RT_{CA}^{*}) and the CP from CABYV (CP_{CA}) and from TuYV (CP_{Tu}) was less clear due to uneven or



FIGURE 1 | Ephrin receptor schematic representation. The different boxes represent the following domains: LBD: Ephrin receptor Ligand-Binding Domain; FN3: Fibronectin type-III domain; KIN: Protein kinase domain; SAM: Sterile alpha motif domain. The external (Ext.), transmembrane (Trans.), and internal (Int.) domains are indicated. Numbers above the representation stand for amino acids. The double line represents the amino acid sequence encoded by the cDNA clone identified by the yeast two hybrid (Y2H) screen.

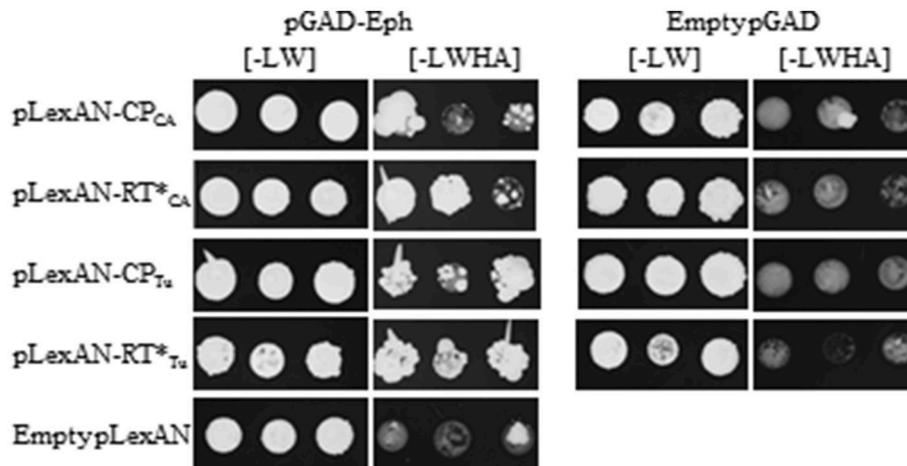


FIGURE 2 | Interaction between CABYV and TuYV structural proteins and Eph partial domain. The yeast strain NMY51 was co-transformed with pGAD-Eph and one of the following constructs: pLexAN-CP_{CA}, pLexAN-RT*_{CA}, pLexAN-CP_{Tu}, pLexAN-RT*_{Tu}, or the empty pLexAN. In parallel, yeast cells were co-transformed with the empty pGAD and one of the viral pLexAN-derived plasmids mentioned above. Three colonies were allowed to grow on a medium lacking leucine and tryptophan [-LW] before being transferred onto a selective medium deprived of leucine, tryptophan, histidine and adenine [-LWHA]. Yeast cells were grown at 28°C for 3 days on [-LW] and for 14 days on [-LWHA] media.

low growth of the double transformed yeast cells on the selective medium.

Feeding *M. persicae* on Transgenic *A. thaliana* Expressing dsRNA_{Eph} or on *in Vitro*-Synthesized dsRNA_{Eph} Reduces Aphid’s Ability to Transmit TuYV

We first analyzed the function of Eph in TuYV transmission by *M. persicae* since a clear interaction was observed in yeast between the Eph domain picked up from the yeast two hybrid screen and RT*_{Tu}. A way to address whether Eph could be involved in TuYV transmission by *M. persicae* is to silence Eph expression in aphids and evaluate the ability of the modified aphids to transmit the virus. We previously compared five different techniques based on the ingestion by aphids of dsRNA molecules targeting *Eph* (Mulot et al., 2016). We selected the two most efficient techniques i.e., feeding aphids (i) on transgenic plants expressing an RNA hairpin of 249 bp corresponding to a central sequence of *Eph* (Hp-Eph) or (ii) on *in vitro*-synthesized

dsRNA of similar sequence. This sequence did not share any sequence identity more than 16 bp with other *M. persicae* expressed genes.

M. persicae were first fed for 10 days on T1 lines of transgenic *A. thaliana* expressing the Hp-Eph. Acquisition of dsRNA_{Eph} and/or siRNA_{Eph} from transgenic plants reproducibly inhibited accumulation of *Eph*-mRNA in whole aphids (53–61% reduction of *Eph*-mRNA accumulation in two independent experiments) when compared with aphids fed on control plants (transgenic *A. thaliana* expressing a 276 bp hairpin RNA targeting the bacterial gene *LacZ*, Hp-LacZ) (Table 1). Depending on the dsRNA sources (Hp-Eph or Hp-LacZ), the aphids were thereafter referred to as dsRNA_{Eph}-treated aphids or dsRNA_{LacZ}-treated aphids. After the dsRNA acquisition from plants, aphids were transferred onto an artificial medium containing purified virus for 24 h. After feeding on virus, the potentially viruliferous dsRNA-treated aphids were then deposited onto Col-0 test plants for virus inoculation and infection of the test plants was assessed by DAS-ELISA 3 weeks later. Interestingly, viruliferous dsRNA_{Eph}-treated aphids

TABLE 1 | Effect of dsRNA_{Eph} acquisition from transgenic *A. thaliana* on TuYV transmission by *M. persicae* and genome internalization in aphids.

Exp.	Aphid feeding source ^a	AAP ^b	Relative fold change Eph-mRNA in whole <i>M. persicae</i> ^c	Source for virus acquisition ^d	nb inf/inoc plants ^e	% of transmission inhibition ^f	Virus genome copies/ μ g RNA internalized into aphids ^g
1	Ara:Hp-Eph (T1) line 1	10 d	0.40 \pm 0.11* (–60%) 1.7E-02 0.45 \pm 0.04* (–55%) 2.6E-02 0.39 \pm 0.16* (–61%) 1.4E-02	TuYV 50 ng/ μ l	6/31* (19%) 9.0E-06	77	3.72 \times 10⁶ \pm 7.33 \times 10⁴* 2.8E-06
2	Ara:Hp-LacZ (T1) line 1 Ara:Hp-Eph (T1) line 1 Ara:Hp-Eph (T1) line 2 Ara:Hp-LacZ (T1) line 1	10 d	1.00 \pm 0.19 0.47 \pm 0.04* (–53%) 2.9E-02 0.42 \pm 0.03* (–58%) 8.6E-04 1.00 \pm 0.02	TuYV 50 ng/ μ l	16/19 (84%) 7/19* (37%) 1.4E-04 7/13* (54%) 8.4E-03 19/20 (95%)	61 43	2.22 \times 10 ⁷ \pm 8.35 \times 10 ⁵ 1.76 \times 10⁷ \pm 1.14 \times 10⁶* 1.5E-06 1.39 \times 10⁷ \pm 6.69 \times 10⁵* 1.3E-06 1.55 \times 10 ⁸ \pm 5.21 \times 10 ⁶

^aTransgenic *A. thaliana* expressing an hairpin RNA targeting *Eph* (Hp-Eph) or LacZ (Hp-LacZ).

^bAcquisition Access Period for dsRNA acquisition; d = days.

^cRelative fold change of Eph-mRNA accumulation \pm standard deviation of triplicates. In brackets the level of expression compared to aphids fed on control conditions (*A. thaliana* expressing dsRNA targeting LacZ). Each result corresponds to one pool of 20 aphids. * and bold characters indicate significant differences in accumulation of Eph-mRNA (Student t-test, $p < 0.05$; p-values are indicated in italics).

^dVirus acquisition was 24h and virus inoculation 7 days on Col-0 test plants. Five viruliferous aphids were deposited per test plant.

^eNumber of plants positive by DAS-ELISA/total number of plants inoculated with aphids. * and bold characters indicate significant differences in the number of plants infected between both conditions (Fisher's exact test, $p < 0.05$; p-values are indicated in italics). In brackets, the percentage of infected plants.

^fPercentage of TuYV transmission inhibition when using dsRNA_{Eph}-treated aphids when compared to dsRNA_{LacZ}-treated aphids. The percentage of infected plants for the LacZ control condition was considered as 100%.

^gTuYV genome copies analyzed by qRT-PCR in whole aphids (pool of 20 aphids) collected after the 4 days of the inoculation access period on the test plants. * and bold characters indicate significant difference in the amount of viral genomes between the two conditions (Student t-test, $p < 0.05$; p-values are indicated in italics).

transmitted TuYV with a significantly lower efficiency than dsRNA_{LacZ}-treated aphids (Table 1). Moreover, the lower ability of the viruliferous dsRNA_{Eph}-treated aphids to transmit TuYV was correlated with a statistically reduced accumulation of the viral genomes (6–11 times less) in the dsRNA_{Eph}-treated aphids when compared to dsRNA_{LacZ}-treated aphids (Table 1).

To confirm these results, *M. persicae* were fed artificially on *in vitro*-synthesized dsRNA targeting *Eph*, since this technique reduces *Eph*-mRNA accumulation in the aphid gut (Mulot et al., 2016). In the first two experiments, using an experimental setup (72 h AAP on dsRNA-Eph at 200 or 400 ng/ μ l) described in Mulot et al. (2016), we observed a reduction in the accumulation of *Eph*-mRNA in the aphid gut (53 and 20% inhibition of *Eph*-mRNA accumulation in guts) (Table 2, Exp. 1 & 2). However, no reduction in TuYV transmission efficiency was observed after feeding the dsRNA_{Eph}-treated aphids on purified virus (Table 2, Exp. 1 and 2). While maintaining the dsRNA concentration in the artificial feeding medium at 400 ng/ μ l, the acquisition time was then extended to 5 days and the virus transmission assay was performed as before. In three independent experiments (Table 2, Exp. 3–5), a significant reduction in TuYV transmission efficiency (from 38 to 81% of reduction) by the dsRNA_{Eph}-treated aphids was observed. Again, the reduction in the virus transmission efficiency was correlated with a significant lower TuYV genome accumulation in the dsRNA_{Eph}-treated aphids after gut clearing (Table 2, Exp. 4 & 5). In these two experiments, TuYV accumulated about 4 times less in the dsRNA_{Eph}-treated aphids compared to dsRNA_{LacZ}-treated aphids. In order to see whether the nature of the virus source could impact the virus transmission efficiency of the dsRNA_{Eph}-treated aphids, virus acquisition was performed on TuYV-infected *Montia perfoliata*. Again, TuYV transmission rate was reduced by 50 and 47% when using the dsRNA_{Eph}-treated aphids (Table 2, Exp. 5 & 6) although the difference in virus transmission was not statistically significant compared with dsRNA_{LacZ}-treated aphids. Nevertheless, this reduction in the TuYV transmission efficiency by the dsRNA_{Eph}-treated aphids correlated with a statistically significant decrease of viral genomes internalized (1.3- and 2.3-fold fewer viral genomes in dsRNA_{Eph}-treated aphids than in dsRNA_{LacZ}-treated aphids) (Table 2, Exp. 5 & 6).

Surprisingly, the virus transmission reduction observed in the five experiments (Table 2, Exp. 3–6) was not always correlated with a reduction of *Eph*-mRNA accumulation in the gut cells and, in two experiments, a higher accumulation of *Eph*-mRNA was observed in the dsRNA_{Eph}-treated aphids compared to dsRNA_{LacZ}-treated aphids (Table 2, Exp. 3 & 5). To evaluate whether the artificial feeding step of aphids on an artificial medium for 5 days could affect *Eph* expression stability, we compared *Eph*-mRNA accumulation in the digestive tubes of aphids fed for 5 days on artificial medium with those of aphids fed on *C. annuum* (plant species used to rear *M. persicae*). Unexpectedly, accumulation of *Eph*-mRNA (normalized to the expression of the two housekeeping genes *L27* and *Rpl7*) in gut cells varied significantly when the aphids were fed on artificial medium (Figures 3A, S5). Expression of *L27* and *Rpl7* was however stable in similar conditions (Figure 3B). In contrast,

TABLE 2 | Effect of *in vitro*-synthesized dsRNA_{Eph} acquisition on TuYV transmission by *M. persicae* and genome internalization in aphids.

Exp.	Aphid feeding source ^a	AAp ^b	Relative fold change Eph-mRNA in <i>M. persicae</i> guts ^c	Source for virus acquisition ^e	nb int/inoc plants ^f	% of transmission inhibition ^g	Virus genome copies/ μ g RNA internalized into aphids ^h
1	dsRNA _{Eph}	72 h	0.47 ± 0.02* (-53%) ^d 1.0E-05	TuYV 25 ng/ μ l	18/20 (90%) 6.6E-01	no	nd
2	dsRNA _{LacZ}	72 h	1.00 ± 0.03	TuYV 25 ng/ μ l	16/20 (80%)	no	nd
	dsRNA _{Eph}		0.80 ± 0.03* (-20%) 1.5E-02		44/69 (64%) 3.2E-01		
3	dsRNA _{LacZ}	5 d	1.00 ± 0.05	TuYV 25 ng/ μ l	43/79 (54%)	38%	nd
	dsRNA _{Eph}		2.78 ± 0.10* (+178%) 4.2E-03		15/30* (50%) 1.8E-02		
4	dsRNA _{LacZ}	5 d	1.00 ± 0.06	TuYV 25 ng/ μ l	24/30 (80%)	67%	3.65 × 10⁶ ± 1.25 × 10⁵ 3.1E-06
	dsRNA _{Eph}		0.20 ± 0.00* (-80%) 2.8E-06		10/39* (26%) 1.9E-05		
5 ⁱ	dsRNA _{LacZ}	5 d	1.00 ± 0.02	TuYV 25 ng/ μ l	27/35 (77%)	81%	1.48 × 10 ⁷ ± 5.03 × 10 ⁵ 1.90 × 10⁶ ± 1.52 × 10⁴* 9.4E-06
	dsRNA _{Eph}		1.44 ± 0.04* (+44%) 2.3E-04		3/30* (10%) 5.0E-04		
6	dsRNA _{LacZ}	5 d	1.00 ± 0.02	Inf. TuYV <i>M. perfoliata</i>	15/28 (54%)	50%	8.90 × 10 ⁶ ± 4.30 × 10 ⁵ 1.76 × 10⁶ ± 4.68 × 10⁴* 3.1E-05
	dsRNA _{Eph}		1.44 ± 0.04* (+44%) 2.3E-04		5/30 (17%) 2.3E-01		
6	dsRNA _{LacZ}	5 d	1.00 ± 0.02	Inf. TuYV <i>M. perfoliata</i>	10/30 (33%)	47%	2.36 × 10 ⁶ ± 1.69 × 10 ⁵ 1.81 × 10⁶ ± 4.38 × 10⁴* 4.0E-05
	dsRNA _{Eph}		0.91 ± 0.02 (-9%) 1.6E-01		9/33 (27%) 7.7E-01		
	dsRNA _{LacZ}		1.00 ± 0.05		17/33 (52%)		4.22 × 10 ⁶ ± 2.08 × 10 ⁵

^aThe aphid feeding source were *in vitro* dsRNA targeting Eph (dsRNA_{Eph}) or LacZ (dsRNA_{LacZ}) at a concentration of 400 ng/ μ l in the artificial medium except in Exp. 1 where the concentration was 200 ng/ μ l.

^bAcquisition Access Period for dsRNA acquisition; h = hours; d = days.

^cRelative fold change of Eph-mRNA accumulation ± standard deviation of triplicates. In brackets the level of expression compared to aphids fed on control conditions (dsRNA_{LacZ}). Each result corresponds to one pool of 100 aphid guts. * and bold characters indicate significant differences between aphids fed on dsRNA_{LacZ} and aphids fed on dsRNA_{Eph} (Student t-test, p < 0.05; p-values are indicated in italics).

^dEph-mRNA accumulation from Mulot et al. (2016) (Table 2E, Exp. 1).

^eAfter feeding on dsRNA, aphids were first transferred onto purified virus or infected *M. perfoliata* for 24 h before being transferred onto Col-0 test plants for 72 h.

^fNumber of plants positive by DAS-ELISA/total number of plants inoculated with aphids. * and bold characters indicate significant differences in the number of plants infected between both conditions (Fisher's exact test, p < 0.05; p-values are indicated in italics). In brackets, the percentage of infected plants.

^gPercentage of TuYV transmission inhibition when using dsRNA_{Eph}-treated aphids when compared to dsRNA_{LacZ}-treated aphids. no = no reduction of the virus transmission efficiency. The percentage of infected plants for the LacZ control condition was considered as 100%.

^hTuYV genome copies analyzed by qRT-PCR in whole aphids (pool of 20 aphids) collected after the 4 days of the inoculation access period on the test plants.

ⁱIn Exp. 5, after dsRNA acquisition, the aphids were split into two batches; one acquired TuYV from purified virus and the other from TuYV-infected plants.

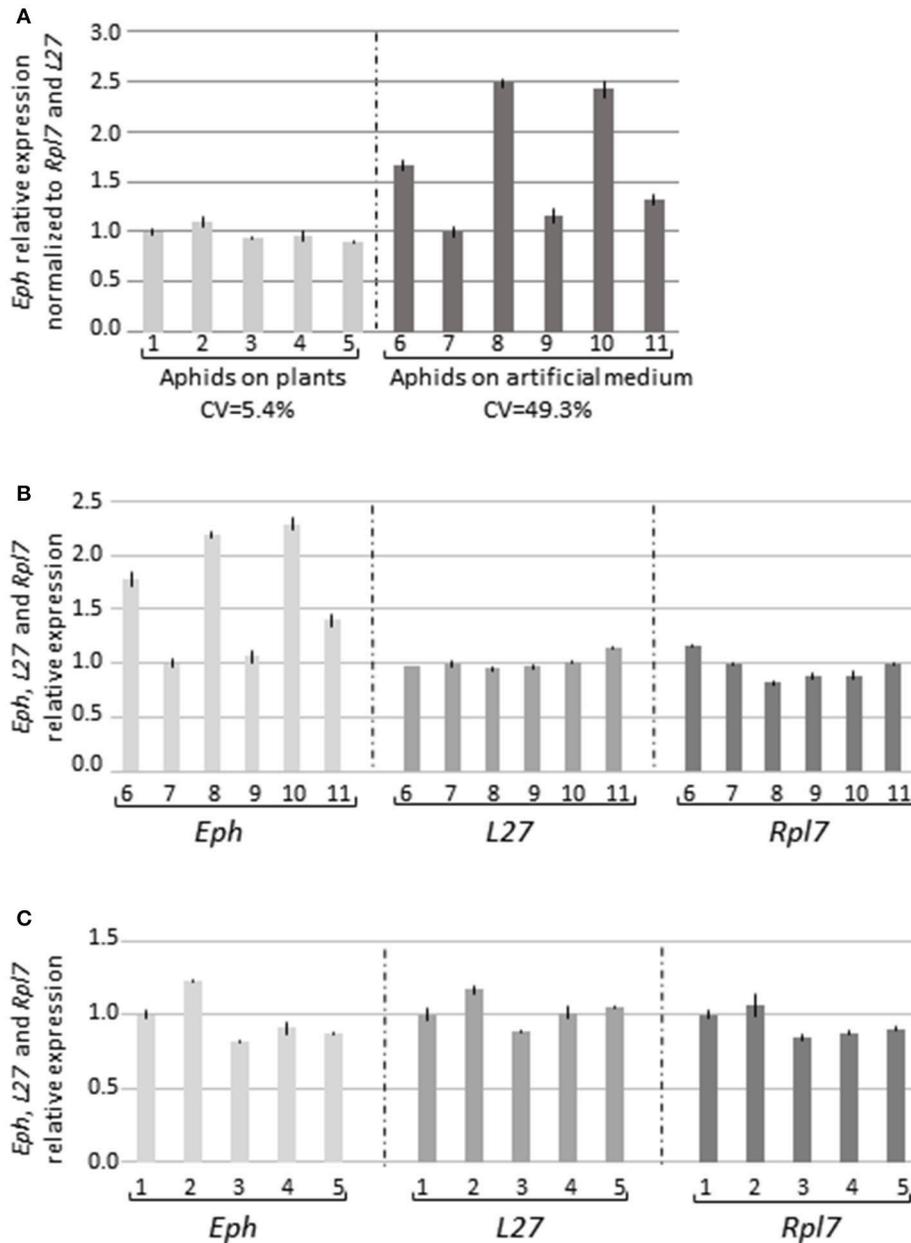


FIGURE 3 | *Eph*-mRNA accumulation in guts from aphids fed on plants or on artificial medium. **(A)** Guts (35 per sample) were dissected from *M. persicae* reared on pepper plants (five samples) or fed during 5 days on the artificial medium MP148 (Harrewijn, 1983) (six samples). The data represent the relative expression of *Eph* in each sample normalized to the accumulation of the two reference genes *L27* and *Rpl7* ± standard deviation of triplicates. The first sample for each condition was arbitrarily fixed to 1. CV: coefficient of variation. *Eph*, *L27*, and *Rpl7* expression without normalization in aphids fed on an artificial medium **(B)** or on plants **(C)**. Similar numbers referred to the same biological samples.

expression of *Eph* as well as *L27* and *Rpl7* was stable in gut samples collected from aphids fed on plants (Figures 3A,C, S5). The high variation of *Eph*-mRNA accumulation in aphids fed on artificial medium may account for our inability to reproducibly observe a reduction of *Eph*-mRNA accumulation in aphids fed during 5 days on an artificial diet containing the dsRNA_{Eph}.

Feeding *M. persicae* on *in Vitro*-synthesized dsRNA_{Eph} Impacts Internalization of Other Poleroviruses Into the Aphid's Body

The specific reduction of virus transmission after acquisition of dsRNA_{Eph} was evaluated with two other poleroviruses transmitted by *M. persicae* (Lecoq et al., 1992; Stevens et al.,

TABLE 3 | Effect of *in vitro*-synthesized dsRNA_{Eph} acquisition on BMVYV and CABYV transmission by *M. persicae* and genome internalization in aphids.

Exp.	Aphid feeding source ^a	Source for virus acquisition ^b	nb inf/inoc plants ^c	% of transmission inhibition ^d	Virus genome copies/ μ g RNA internalized into aphids ^e
1	dsRNA _{Eph}	BMVYV 50 ng/ μ l	10/32 (31%) <i>1.0E-01</i>	38%	$1.28 \times 10^6 \pm 1.25 \times 10^5$ <i>2.9E-03</i>
	dsRNA _{LacZ}		16/32 (50%)		$2.30 \times 10^6 \pm 2.41 \times 10^5$
2	dsRNA _{Eph}	BMVYV 50 ng/ μ l	11/28 (39%) <i>1.4E-01</i>	31%	$3.02 \times 10^6 \pm 0.52 \times 10^5$ <i>7.8E-05</i>
	dsRNA _{LacZ}		16/28 (57%)		$4.15 \times 10^6 \pm 1.06 \times 10^5$
3	dsRNA _{Eph}	CABYV 100 ng/ μ l	1/11 (9%) ^f <i>2.9E-01</i>	67%	$6.02 \times 10^6 \pm 4.46 \times 10^5$ <i>2.7E-03</i>
	dsRNA _{LacZ}		3/11 (27%)		$8.11 \times 10^6 \pm 3.14 \times 10^5$

^aThe aphid feeding source were *in vitro* dsRNA targeting Eph (dsRNA_{Eph}) or LacZ (dsRNA_{LacZ}) at a concentration of 400 ng/ μ l in the artificial medium and the AAP was fixed to 5 days.

^bAfter feeding on dsRNA, aphids were first transferred onto purified virus for 24 h before being transferred onto Col-0 test plants for 72 h.

^cNumber of plants positive by DAS-ELISA/total number of plants inoculated with aphids. In brackets, the percentage of infected plants. *p*-values from the Fisher's exact test are indicated in italics.

^dPercentage of TuYV transmission inhibition when using dsRNA_{Eph}-treated aphids when compared to dsRNA_{LacZ}-treated aphids. The percentage of infected plants for the LacZ control condition was considered as 100%.

^eBMVYV or CABYV genome copies analyzed by qRT-PCR in whole aphids (6 pools of 20 aphids in Exp. 1, 3 pools of 20 aphids in Exp. 2 & 3) collected after the 4 days of inoculation access period on test plants. * and bold characters indicate significant difference in the amount of viral genomes between the two conditions (Student *t*-test, *p* < 0.05; *p*-values are indicated in italics).

^fIn this experiment, four viruliferous aphids (instead of two in the other experiments) were transferred on each test plant for virus inoculation.

2005), *Beet mild yellowing virus* (BMVYV) and CABYV. After *in vitro* acquisition of dsRNA, dsRNA_{Eph}-treated aphids transmitted less efficiently BMVYV in two independent experiments (38 and 31% transmission inhibition in Exp. 1 and 2, respectively, **Table 3**), but the transmission inhibition was not statistically significant when compared to the control condition using dsRNA_{LacZ}-treated aphids. However, a statistically significant decrease of BMVYV genomes internalized into the dsRNA_{Eph}-treated aphids was measured (1.8 and 1.4 times less viral genomes in dsRNA_{Eph}-treated aphids than in dsRNA_{LacZ}-treated aphids; Exp. 1 and 2, respectively, **Table 3**). When CABYV was used as a virus source, the number of infected plants was particularly low, which makes it difficult to draw a clear conclusion on the effect of dsRNA_{Eph} acquisition on the CABYV transmission efficiency (**Table 3**, Exp. 3). Such low transmission efficiency of CABYV from purified virus has previously been observed in our laboratory even with high concentrations of virus in the artificial medium (V. Brault, unpublished), and could be explained by instability of the particles after the purification procedure. Nevertheless, a moderate, but still significant, reduction of the number of CABYV genomes internalized was observed in the dsRNA_{Eph}-treated aphids (**Table 3**, Exp. 3). These results suggest that Eph could have a broader role in poleroviruses transmission by *M. persicae*.

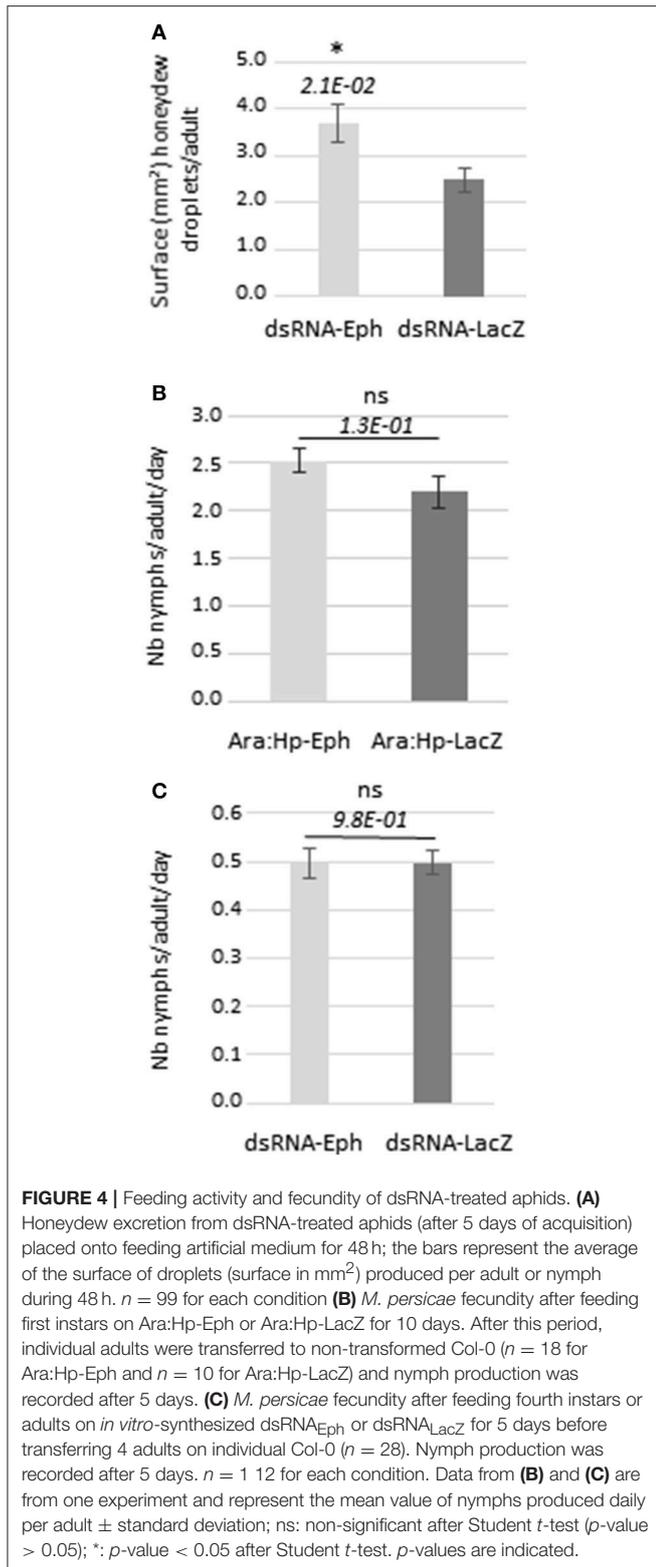
The Inhibition of TuYV Transmission by dsRNA_{Eph}-Treated Aphids Is Not Due to a Reduction in Feeding Activity

The lower virus transmission efficiency of dsRNA_{Eph}-treated aphids could be due to a reduced feeding activity on the virus source. Since the majority of the virus transmission

experiments presented in this manuscript (9 out of 11 in total) were performed using purified virus as the virus source, we measured the feeding activity of the dsRNA-treated aphids when placed onto the artificial diet. Electropetrography could not be developed to measure the feeding phases of aphids because this technique is not adapted to evaluate sustained ingestion activity of aphids from an artificial medium (Tjallingii, 1985). Therefore, we measured honeydew excretion of the dsRNA_{Eph}-treated aphids after transferring them onto a fresh artificial medium for 48 h. The aphid feeding activity on plants infected with luteovirids has been previously correlated with the efficiency of virus transmission (Sylvester, 1967; Van den Heuvel and Peters, 1990). The surface area covered by honeydew droplets produced by dsRNA_{Eph}-treated aphids was slightly higher, than the one secreted by dsRNA_{LacZ}-treated aphids (**Figures 4A, S6**). In another experiment, no difference in honeydew excretion was observed between dsRNA_{Eph}- and dsRNA_{LacZ}-treated aphids (**Figure S7**). These results show that the reduction of TuYV transmission by dsRNA_{Eph}-treated aphids is not correlated with a lower feeding activity of these aphids on the artificial medium and therefore not linked to a reduced ingestion of virus particles.

Fecundity of dsRNA_{Eph}-Treated Aphids Is Not Affected by Ingestion of dsRNA and/or siRNA From Transgenic *A. thaliana*:Hp-Eph or From *in Vitro*-synthesized dsRNA_{Eph}

In order to investigate the impact of dsRNA and/or siRNA acquisition from transgenic plants or from *in vitro*-synthesized dsRNA on aphid's fecundity, nymph production by the dsRNA-treated aphids was analyzed. Feeding aphids during 10 days on the transgenic plants (Ara:Hp-Eph or Ara:Hp-LacZ) (**Figure 4B**)



or during 5 days on the dsRNA targeting *Eph*-mRNA or *LacZ*-mRNA (Figure 4C) did not affect aphid's fecundity. These results suggest that acquisition of dsRNA or siRNA from either source did not significantly impact aphid physiology. A low fecundity

of aphids fed during 5 days on the artificial medium containing the dsRNA can be observed (Figure 4C). The artificial medium composition may affect aphid fecundity but the amount of dsRNA ingested by aphids from the artificial diet may also have an impact on the aphid physiology.

DISCUSSION

To be transmitted by aphids luteovirids acquired in the phloem of infected plants must cross several epithelia at the gut and salivary gland levels in the vector before being inoculated into a plant during feeding. Virion transport through the epithelia requires virus particle recognition by specific receptors (Mercer et al., 2010). In the present study, we identified Eph as a potential receptor of TuYV in *M. persicae*. Part of the external domain of Eph, corresponding to the fibronectin type III repeats, was able to bind in yeast to the minor capsid protein (RT*) of TuYV, which is strictly required for aphid transmission (Brault et al., 1995). RT* protein is involved in the transcytosis of TuYV through the posterior midgut cells and strictly required for virus internalization into the ASG cells (Brault et al., 1995; Reinbold et al., 2001). We showed that feeding *M. persicae* with siRNA/dsRNA targeting *Eph*-mRNA, produced *in planta* or *in vitro*, prior to TuYV acquisition, consistently reduced TuYV accumulation in aphids and subsequently virus transmission to plants. Since siRNA/dsRNA ingestion by aphids did not affect aphid feeding behavior, we concluded that ingestion of siRNA/dsRNA targeting *Eph*-mRNA likely affected TuYV acquisition by *M. persicae*. A lower accumulation of two other poleroviruses, BMYV and CABYV, in similarly treated aphids, was also observed, suggesting a broader implication of Eph in the internalization of poleroviruses into *M. persicae*.

Virus transmission reduction was sometimes correlated with a reduction in the accumulation of *Eph*-mRNA in aphids, in particular after feeding aphids on transgenic plants expressing the dsRNA_{Eph} during 10 days. However, we observed *Eph*-mRNA instability in aphids fed on an artificial medium. This instability could be potentially intensified, in these non-natural conditions, by fluctuations in dsRNA ingestion along the acquisition period, or alternatively, by dsRNA stability that could be altered during this 5-day period. Nevertheless, it is important to mention that virus transmission inhibition was always linked to a reduction of virus accumulation in aphids. Although we show that Eph is involved in polerovirus transmission by *M. persicae*, we cannot conclude from our experiments whether Eph is acting at the gut, at the ASG, or at both levels in *M. persicae*. Nevertheless, considering that oral acquisition of dsRNA targeting Eph preferentially affects gene expression in the gut level (Mulot et al., 2016), and reduces virus accumulation into the aphid's body, it is likely that Eph, at least, is acting at the gut level.

Ephrin receptors are good candidates to be involved in polerovirus transmission by *M. persicae*. In most of the cases, Ephs and their ephrin ligands control a wide array of cell-to-cell interactions in mammals without involving internalization of both proteins. In contrast, in some instances the interaction between Ephs and ligands results in the endocytosis of the

complex (Marston et al., 2003; Zimmer et al., 2003; Pitulescu and Adams, 2010), a phenomenon that clearly resembles poliovirus internalization into aphid cells. Moreover, there is evidence that Eph receptors can also be activated by soluble ephrin ligands present in the environment (Alford et al., 2010). In this regard, it is interesting to point out that Eph receptors are involved in human and simian virus uptake. Eph A2 was identified as a host co-factor for Hepatitis C virus entry in liver cells (Lupberger et al., 2011). Internalization of this enveloped virus by clathrin-dependent endocytosis requires several cell-surface molecules, some thought to be essential receptors while others facilitating virus uptake (von Hahn and Rice, 2008). Another human virus, the *Kaposi's sarcoma-associated herpesvirus* (KSHV) also relies on the presence of Eph A2 to enter epithelial cells, but by micropinocytosis rather than endocytosis (Chakraborty et al., 2012; Hahn et al., 2012). Finally, the *Rhesus monkey rhadinovirus*, a closely relative to KSHV, use a wide array of Ephs to be endocytosed into monkey endothelial cells (Hahn and Desrosiers, 2013).

It is also interesting to mention that several Eph receptors are known to bind to caveolin-1, a protein involved in endocytosis of non-enveloped viruses like simian virus 40 (Pelkmans et al., 2001), echovirus 1 (Marjomaki et al., 2002), and *Junonia coenia* densovirus (Wang et al., 2013). Although luteovirids are thought to be internalized into aphid cells by clathrin-mediated endocytosis, an alternative route for virus uptake based on caveolae is still conceivable (Gildow, 1999; Brault et al., 2007). Indeed, compared to clathrin-mediated endocytosis, caveolae entry results in the formation of vesicles that do not enter the traditional acidic endosome/lysosome system (Thomsen et al., 2002). By avoiding acidification, virus internalization into caveolae might be beneficial for non-replicating viruses, like polioviruses. In the light of these results, it could be interesting to reassess the mechanism of poliovirus internalization into aphid cells by targeting Cav-1 expression in aphids by RNA interference or by using caveolin-specific inhibitors (Rejman et al., 2005).

The ephrin type-B receptor 1-B could be the second aphid protein identified as a potential luteovirid receptor. APN was previously shown to be involved in PEMV (*Enamovirus* genus) internalization into *A. pisum* (Liu et al., 2010; Linz et al., 2015). Here, we bring evidence that Eph is another aphid protein involved in poliovirus acquisition and transmission by *M. persicae*. Determining whether these two proteins act in concert in both aphid species, or are specific for one aphid species, will be a challenge for future studies. As already mentioned, implication of several proteins in the internalization process of mammalian viruses into cells is more likely a general mechanism: viral surface components must first bind to attachment factors on the cell surface before interacting with receptors that drive reactions leading to entry (Mercer et al., 2010; Grove and Marsh, 2011; Cossart and Helenius, 2014). One extreme example is HCV which has been shown to require about ten different molecules for cell entry (Grove and Marsh, 2011). Interestingly, some of them are responsible for virus non-specific attachment on the cell surface while interaction with the liver specific intercellular adhesion molecule-3-grabbing non-integrin (L-SIGN) is thought to confer tissue tropism *in vivo* (Gardner et al., 2003). Analyzing

Eph-mRNA distribution along the digestive tube of *M. persicae* will show whether this protein is responsible for TuYV gut tropism at the posterior midgut.

In this study, we showed that acquisition by aphids of dsRNA molecules targeting *Eph* can reduce TuYV transmission. Aphid survival and fecundity were not affected by *Eph*-mRNA targeting. Eph may therefore be an ecologically safe target to reduce luteovirids impacts by inhibiting their dispersion by aphids. The dsRNA_{Eph} molecules could be expressed in transgenic plants (as in this study), but could also alternately be sprayed on cultures. This innovative delivery system has been assayed on different aphid species by aerosolizing siRNA targeting a carotene dehydrogenase and a branched chain-amino acid transaminase (Thairu et al., 2017). A moderate inhibition of gene expression was observed but the effect varied upon the targeted gene and the aphid species. Before applying this technology to inhibit expression of *Eph* in aphids, additional experiments are required to address dsRNA_{Eph} stability in the environment and efficacy when aerosolized on aphids. However, at this point, it is tempting to make a parallel with the new strategies that are developed to curtail viral human diseases, and in particular Human immunodeficiency virus infection. Indeed, among the therapies to inhibit Human immunodeficiency virus cell entry, a simultaneous knock down of the CCR5 co-receptor by small RNA hairpins together with the expression of an antiviral fusion inhibitor peptide is under a clinical trial (Wolstein et al., 2014; Hutter et al., 2015).

The results presented in this paper pave the way toward a better comprehension of the molecular mechanisms governing polioviruses transmission by aphids. Expression and localization of the Eph in aphid species differing in their ability to transmit polioviruses together with the identification of Eph viral ligands and cellular partners need to be addressed in the future.

AUTHOR CONTRIBUTIONS

VB, BM, MM, and SB: designed the experiments; MM, BM, SB, MR, SM, and NB: performed the experiments; VB, BM, MM, and SB wrote the paper.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fmicb.2018.00457/full#supplementary-material>

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Conflict of Interest Statement: 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.

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