A whitefly effector Bsp9 targets host immunity regulator WRKY33 to promote performance

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Whiteflies, Bemisia tabaci (Hemiptera), are pests causing economic damage to many crops, capable of transmitting hundreds of plant vector-borne viruses. They are believed to secrete salivary protein effectors that can improve vector colonization and reproductive fitness in host plants. However, little is known about effector biology and the precise mechanism of action of whitefly effectors. Here, we report a functional screening of B. tabaci salivary effector proteins (Bsp) capable of modulating plant innate immunity triggered by plant endogenous pattern peptide Pep1. Four immunity suppressors and two elicitors were identified. Bsp9, the most effective immunity suppressor, was further identified to directly interact with an immunity regulator WRKY33. We provide evidence that Bsp9 may suppress plant immune signalling by interfering with the interaction between WRKY33 and a central regulator in the MAPK cascade. The interference by Bsp9 therefore reduces plant resistance to whitefly by inhibiting activation of WRKY33-regulated immunity-related genes. Further detailed analysis based on transgenic plants found that whitefly effector Bsp9 could promote whitefly preference and performance, increasing virus transmission. This study enriches our knowledge on insect effector biology.

This article is part of the theme issue ‘Biotic signalling sheds light on smart pest management’.

1. Introduction

Piercing/sucking feeding insects cause extensive crop losses directly [1]. Besides ingesting plant sap, phloem-feeding insects, such as whitefly, transmit 80% of known plant vector-borne pathogens and cause huge economic loss indirectly. During feeding, insects secrete salivary protein into plant cells together with virus particles. Although, for plant and animal microbes, it is well established that effectors target host proteins to manipulate host cell processes and promote infection and disease [2], understanding the effector biology of insects is still at a very early stage. So far, only a small number of salivary effectors from aphid species have been identified and a few of them have been confirmed by their functions in plant–insect interactions [3]. Among these insects with identified effectors, the broad host range aphid Myzus persicae is the most researched species. Some of effectors from M. persicae...
and homologues from other aphid species have been successfully identified through genomics. A few of them, such as Mp1 (Pint01) and Mp2 (Pint01), Mp10, Mp42, Mp55, Mp56, Mp57, Mp58 and others, have been functionally characterized to target host plant proteins to modulate immunity for enhancing aphid fitness [3–5]. However, the effectors from other insects, such as plant hopper and whitefly, remain elusive.

Whitefly, Bemisia tabaci (Hemiptera), is a polyphagous insect and a supervector, transmitting more than 300 plant virus species, that is a threat for many crops across the globe [1]. Whitefly might mediate the suppression of plant defences by secreting protein effectors to improve host colonization and reproductive fitness [6,7]. The whitefly-transmitted monopartite geminivirus begomoviruses are frequently associated with pandemic crop diseases such as tomato yellow leaf curl virus (TYLCV) and tomato yellow leaf curl China virus [8]. We have shown that geminiviruses could repress defensive responses in infected plants leading to an improved fitness of their vector, whitefly, thereby promoting vector performance and, in turn, facilitating pathogen spread [9], but it is unknown whether the infestation of whitefly could affect plant immunity, possibly by secreting salivary effectors.

Plants fend off attacks from herbivores and pathogens in various ways, e.g. via physical barriers, volatile or non-volatile compounds, and through induction of defensive responses mainly controlled by phytohormones and innate immunity [10–12]. The phytohormone jasmonate (JA) is known to be indirectly manipulated by viruses to promote whitefly performance [9,13]. Plant innate immunity includes two major types of resistance mechanisms against pathogens and herbivores. The first layer is the pathogen-associated molecular pattern or damage-associated molecular pattern (DAMP)-triggered immunity, so-called pattern-triggered immunity (PTI). The second layer is a more specific effector-triggered immunity (ETI) [14–16]. PTI is a multistep response, which is triggered upon plant pattern recognition receptors recognizing the conserved pathogen molecules or endogenous peptide elicitors such as Pep1–Pep7 family [17,18]. PEPR1 and PEPR2 encode receptors to recognize Pep1 in Arabidopsis. The AtPEP–PEPR system has been reported to be induced by Spodoptera littonis feeding. Thus, Pep peptides function as DAMPs in response to wound- and herbivory-induced stresses [18–20]. Subsequently, many downstream signalling events are initiated, including activation of the mitogen-activated protein kinase (MAPK) cascades and the transcription of defensive genes, especially anti-pest defensin gene (PDF1.2) in Arabidopsis thaliana [21–25].

Whiteflies are important agricultural pests, but little is known about their effector biology. The goal of this study was to establish an efficient functional screening system of whitefly effectors and to elucidate the precise mechanism of whitefly effectors in the interaction with host and virus. Here, we identify a whitefly salivary protein Bsp9 (whose expression is induced by TYLCV) that can effectively inhibit the plant defence response to whitefly infestation. Bsp9 interacts with a resistance-related transcription factor WRKY33. The plant immune regulation by Bsp9 affects whitefly fitness, thereby leading to a possible enhancement of virus transmission. Our research uncovered how begomoviruses manipulate whitefly effectors to promote virus transmission for worldwide invasion.

2. Material and methods

(a) Plant and insect materials

Tomato (Solanum lycopersicum, Heinz 1706-BG, LA4345) seeds were ordered from Tomato Genetic Resource Center, University of California, Davis, USA, and propagated. Seeds of tomato and Nicotiana benthamiana were grown in a greenhouse at 25°C with a 12 L : 12 D cycle and young seedlings of three to four true leaf stages were used.

Arabidopsis thaliana (ecotype Col-0) was used for Agrobacterium-mediated transformation; the Arabidopsis wrky33 mutant was given by Prof. Jinlong Qiu (Institute of Microbiology, Chinese Academy of Sciences). The plant binary vectors 35S:GUS, 35S:YFP, 35S:Bsp9-YFP, 35S:Bsp9-HA were constructed based on PCR. Plasmids were introduced into Agrobacterium tumefaciens strain EHA105 by electrot transformation and Arabidopsis transformations were performed according to the floral-dipping method [26].

Whiteflies (B. tabaci MEAM1/B) were maintained on tomatoes in a growth chamber at 25°C with a 14 L : 10 D cycle and 65% relative humidity.

(b) Transcriptome sequencing and data analysis

Total RNA was isolated from viruliferous or virus-free whitefly samples using TRIzol reagent (Invitrogen) according to the manufacturer’s instructions and used for library construction and sequencing on Illumina HiSeq X Ten platform at the Annonorad Gene Technology Company (Beijing, China). The de novo assembly of RNA-seq was conducted using the Trinity platform. Illumina sequence data were selectively filtered using SolexaQA to remove read lengths less than 35 bp and low-quality sequence at each nucleotide. Clean reads of Illumina sequence data were mapped by Bowtie2. Raw counts for each predicted gene were calculated as reads per kilobase of exon model per million mapped fragments (FPKM). Based on these statistical analyses, genes with \(p < 0.01\) and log2 (fold change) value of RPKM greater than 1 were considered to be significantly differentially expressed genes (DEGs). DEGs between viruliferous and virus-free whiteflies were identified and mapped to whitefly salivary gland transcriptome using hBLASTn. Finally, we got unigenes upregulated by TYLCV in salivary glands. Trinotate and ORF (open reading frame) Finder were used for performing the functional annotation of unigenes and ORFs. We narrowed the range of the genes by the length of ORFs between 200 and 600 bp. SignAl 4.0 program was used to predict the presence of signal peptides and cleavage sites in the amino acid sequences encoded by the ORFs found in salivary gland ESTs. Subsequently, proteins with a signal peptide were predicted to contain at least one transmembrane domain by TMHMM Server 2.0 and therefore more likely to remain in the membrane of the salivary gland cell during secretion. Besides these, a protein without or with one transmembrane domain included in the part of predicted signal peptide would be considered as a secreted protein, as well as a potential salivary protein.

(c) Virus inoculation

Tomato yellow leaf curl virus (TYLCV-SH2; GenBank accession no. AM282874) was kindly provided by Prof. Xue-Ping Zhou (Institute of Plant Protection, Chinese Academy of Agriculture Sciences, China). Tomato seedlings were inoculated with TYLCV by agro-inoculation [27]. Virus-inoculated tomatoes were cultivated in the growth chamber at 25°C.

(d) Viral DNA measurement

Total genomic DNA was extracted from systemically infected leaves, and viral DNA was detected by real-time PCR with TYLCV-specific primers as well as the Arabidopsis \(\alpha\)-tubulin2

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(At5g62690)-specific primers or tomato tubulin-specific primers as endogenous controls. Three biological replicates were used in this experiment.

(e) Whitefly bioassay
The two-choice experiments with MEAM1 (Middle East-Asia Minor 1) B. tabaci were performed as described previously [9]. The preference of B. tabaci was compared in bioassays between two Arabidopsis genotype lines. Two six-week-old plants of similar size and leaf numbers were placed in a cage (40 × 40 × 40 cm) 20 cm apart. Two hundred adult whiteflies were captured and placed on ice for 1–2 min to temporarily stun them. They were then released from a point equidistant from the two plants. After 20 min whitefly free-choice, the number of whiteflies on each of the two plants was recorded. For one genotype, eight plants were used in each bioassay with three replicates.

The MEMA1 whitefly oviposition experiment was performed as described in Li et al. [9]. Each cohort of three male and female adult whiteflies was released into a leaf-clip cage that enclosed a single leaf of a six-week-old plant. Whitefly eggs on each leaf were counted after infestation for 10 days using a microscope. Eight plants of each line were used in the experiment. The MEMA1 whitefly nympha development experiment was then performed as described in Li et al. [9]. Each of 16 female adults were released into a leaf-clip cage that enclosed a single leaf of a six-week-old plant. The number of nymphs on the Arabidopsis leaves was counted after 20 days using a microscope. Eight biological replicates were conducted in this experiment.

(f) Whitefly infestation
To investigate the effect of whitefly infestation on virus accumulation, three-week-old tomato plants were first infected by TYLCV for 14 days with the same virus load, then infested by MEMA1 whiteflies. After 3 days of whitefly infestation, all whiteflies were gently removed from the plants. TYLCV-infected tomato plants without whitefly infestation were used as control check (CK). Tomato plant samples were taken after one week of whitefly infestation.

For gene expression in whitefly-infested plants, leaves of healthy Arabidopsis plants were placed inside leaf-clip cages. Fifty adult whiteflies were captured and released into each cupped leaf. Leaf samples were collected after whitefly feeding at the indicated time points.

For the detection of Bsp9 secretion from whitefly to tomato, 500 adult whiteflies were released into a leaf-clip cage that enclosed a single leaf of a four-week-old plant and infested for 72 h before plant sampling.

For the detection of virus transmission by whitefly, virus-free adult whiteflies were placed on TYLCV-infected tomato plants for 48 h of virus acquisition. Thirty viruliferous whiteflies were then captured and released onto a three-week-old Arabidopsis plant grown on Murashige and Skoog medium and enclosed in a leaf-clip cage. After 72 h of whitefly-to-plant virus transmission, whiteflies were then gently removed and plant samples were collected to isolate individual DNA.

(g) Quantitative RT-PCR
Total DNA of tomato leaves was isolated by the CTAB method (cetyltrimethyl ammonium bromide). Total RNA of whitefly was extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions. Total RNA of tomato leaves was isolated by plant RNA purification reagent (Invitrogen, Carlsbad, CA, USA) [28]. Total RNA of 800 ng for each sample was used for reverse transcription with TransScript One-Step gDNA Removal and followed by cDNA synthesis (Synthesis SuperMix, Transgen, China). Quantitative PCR was performed on a Bio-Rad CFX96 real-time PCR system with ThunderbirdTM SYBR qPCR mix (TOYOBO). Four independent biological samples were analysed for each experiment and three independent experiments were performed and similar results were observed. The primers used for mRNA detection of target genes by real-time PCR are listed in electronic supplementary material, table S1.

(h) Luciferase activity assay
Arabidopsis PDF1.2 promoter: luciferase was used as a reporter construct. Candidate whitefly salivary protein genes were driven by Cauliflower mosaic virus (CaMV) 35S promoter: genes as effector constructs. Leaves of N. benthamiana were agroinfiltrated with the indicated agrobacterium stains which contain individual constructs. Infiltrated leaves were harvested after 2 days’ treatment and the luciferase activity was quantified by a microplate reader.

Synthetic Pep1 (1 μM) peptide was used as an elicitor to activate plant immunity for 3 h before sampling. Each treatment was repeated eight times in one experiment. The experiment was repeated twice with similar results. The fold of luciferase activation by Pep1 was calculated against control group without Pep1 treatment. 35S:YFP was used as the vector control.

For luciferase complementation-based protein interaction assay, Agrobacterium carrying the indicated constructs were infiltrated into N. benthamiana leaves and the luciferase imaging assays were performed 48 h after infiltration [29].

(i) Yeast two-hybrid analysis
For this, the Arabidopsis Mate and Plate Library was used (Clontech). Full-length protein of Bsp9 was cloned into the pGBT9 vector to generate BD-Bsp9 as a bait vector and putative interaction was screened by following the manufacturer’s protocol (Matchmaker Gold Yeast Two-Hybrid System, Clontech). To further confirm the interaction between Bsp9 and WRKY33, the yeast strain Y2H Gold was co-transformed with BD-Bsp9 and AD-WRKY33 constructs and plated on SD-Leu-Trp selective dropout medium. Colonies were transferred onto SD-Leu-Trp-His plates to verify positive clones. The binding domain vector (BD) pGBK7 and activation domain vector (AD) pGADT7 were used as negative controls.

(j) Bimolecular fluorescence complementation assay
All constructs were transferred into Agrobacterium C58C1 competent cells. The bimolecular fluorescence complementation (BiFC) assay was performed, as previously described in Sun et al. [30]. Agrobacterial cells containing indicated constructs were infiltrated into three-week-old N. benthamiana leaves. Fluorescence was observed owing to the complementation of Bsp9 fused with the cEYFP and WRKY33 fused with nEYFP. Images of fluorescence were taken by confocal microscopy (Leica SP8) after 48 h incubation. Plant nuclei were stained with DAPI (4′,6-diamidino-2-phenylindole) infiltrated into leaves 30 min prior to detection.

For competitive inhibition assay, A. tumefaciens strains containing expression vectors for cEYFP-MPK6 + nEYFP-WRKY33 and 35S:Bsp9 were co-injected into N. benthamiana leaf cells and kept in the dark for 2 days. The control was co-injected with the same volume MMA buffer as 35S:Bsp9. Co-expression with the same volume 35S:GL5 was used as a negative control. Fluorescence intensity was measured by ImageJ.

(k) Antibody preparation
The DNA fragment of Bsp9 was cloned into pET-28a (+) vector to generate 6×His-Bsp9 fusion construct. His-Bsp9 protein was purified using Ni-nitrilotriacetate (Ni-NTA) agarose (Qiagen)
analysis with anti-Bsp9 polyclonal antibody. Bsp9 protein was detected by Western blot gels and transferred to a polyvinylidene difluoride membrane of total protein were separated on 10% SDS–polyacrylamide gels and transferred to a polyvinylidene difluoride membrane (Millipore) [30]. Bsp9 protein was detected by Western blot analysis with anti-Bsp9 polyclonal antibody.

(l) Protein extraction and Western blot
Protein was extracted from plants with extraction buffer (20 mM Tris–HCl pH 8.0, 100 mM NaCl, 10 mM MgCl2, 50 mM DTT, 0.5 mM PMSF and protease inhibitor cocktail). Equal amounts of total protein were separated on 10% SDS–polyacrylamide gels and transferred to a polyvinylidene difluoride membrane (Millipore) [30]. Bsp9 protein was detected by Western blot analysis with anti-Bsp9 polyclonal antibody.

(m) Data analysis
Differences in TYLCV accumulation levels, gene expression levels, whitefly performance and relative fluorescence intensity and relative luciferase activity were determined using Student’s t-tests for comparing two treatments or two lines. Differences in whitefly choice between different lines were analysed by non-parametric Wilcoxon-matched pair tests (with two dependent samples). All tests were carried out with Excel and GraphPad Prism.

(n) Accession numbers
Sequence data from this work can be found in Genebank/EMBL or the Arabidopsis Information Resource (www.Arabidopsis.org) under the following accession numbers: TYLCV-SH2 (AM282874), AtPDF1.2 (AT5G44420), AtPEPR1 (AT1G73080), AtWRKY33 (AT2G38470), AtTPS10 (AT2G24210), AtTPS14 (AT1G61680), AtTPS15 (AT3G14520), AtTPS20 (AT5G48110), MED whitefly Bsp9 (MH744980) and MEAM1 whitefly Bsp9 (MH744981).

3. Results
(a) Whitefly infestation increases the accumulation of tomato yellow leaf curl virus
We inoculated tomato with TYLCV, a begomovirus transmitted by an invasive MEAM1 whitefly, and examined the symptoms of infected plants. TYLCV-infected tomatoes showed mild yellowish symptoms at 14 days after inoculation while using the agroinfiltration method alone. Interestingly, TYLCV-infected tomato plants, followed by infestation of MEAM1 whiteflies for 3 days, exhibited more obvious yellowish and upward curling leaf symptoms at 7 days after infestation of whitefly compared with mock control (figure 1a).

(b) Comparative transcriptomic screening of whitefly salivary effectors
One hypothesis for the enhancement of TYLCV pathogenesis by the infestation of whitefly is that TYLCV could induce accumulation of whitefly salivary effectors and, in turn, inhibit host defence against virus. To identify candidate whitefly salivary effectors induced by TYLCV, we firstly undertook a comparative transcriptomic analysis of viruliferous whitefly versus virus-free whitefly (electronic supplementary material, figure S1). High-quality total RNA samples of viruliferous whitefly and virus-free MEAM1 whitefly were extracted, and de novo sequencing was conducted with the Illumina sequencing platform. Through filtering the adaptors and low-quality sequences, assembling by Trinity and mapping by Bowtie2, we finally got 86 428 unigenes based on the raw data of scaffolds. Among them, expression levels of 53 353 unigenes were induced by TYLCV at least twofold. Comparing with the whitefly salivary gland dataset, only 1.4% of TYLCV-induced unigenes (778/53 353) were found in salivary glands [31]. Taking into consideration that most effectors are putative secreted proteins which possess a suitable opening reading frame size, an N-terminal signal peptide and have no transmembrane regions, finally, we cloned 10 full-length cDNA of TYLCV-induced candidate salivary effector proteins for further functional characterizations (electronic supplementary material, table S2).

(c) Whitefly salivary protein Bsp9 suppresses plant immune response
As a functional analysis platform for whitefly salivary proteins has rarely been reported, we sought to establish a suitable screening and reporter system to identify whitefly effectors. We observed that the expression of a PTI membrane receptor PEPR1 was rapidly induced upon whitefly infestation in Arabidopsis (figure 2a). Furthermore, another
downstream defensive marker gene *Arabidopsis* PDF1.2 was also highly induced upon whitefly infestation or the treatment with Pep1 polypeptide, a ligand for PEPR1/2 receptors (figure 2b; electronic supplementary material, figure S3). These results indicated that Pep1 treatment can mimic the stimulation of whitefly infestation on plants and the expression level of PDF1.2 may represent the level of plant immune response to whitefly. Accordingly, we developed a novel system consisting of an effector and a reporter system together to functionally identify a whitefly salivary immunity regulator (as shown in figure 2c). Each of these two plasmids was transformed into *A. tumefaciens* and co-inoculated leaves of *N. benthamiana* by agroinfiltration. By measuring the activity of the PDF1.2 promoter upon co-expression of tested protein, it is convenient for high-throughput screening of whitefly salivary effectors which can repress plant immune response induced by Pep1. As expected, we found that Pep1 treatment stimulates activity of the PDF1.2 promoter (figure 2d). Interestingly, four salivary effectors repressed the Pep1-triggered activation of PDF1.2 promoter. By contrast, two salivary effectors improved this activity. The preliminary functions of all tested salivary proteins are listed in electronic supplementary material, table S2. Among those, the degree of immune repression by Bsp9 was the highest (figure 2d). Thus, we chose Bsp9 for a detailed downstream analysis on the mechanism of its suppression of PTI.

**Figure 2.** Functional characterization of TYLCV-induced whitefly salivary effectors. (a) Relative expression level of *PEPR1* that encoded a receptor to recognize DAMPs upon infestation of whitefly in *Arabidopsis*. Three-week-old Col-0 plants were infested with whiteflies for the indicated time. Values are mean ± s.d. (n = 4). (b) Relative induction of anti-herbivory marker gene PDF1.2 upon infestation of whitefly or Pep1 treatment. Three-week-old Col-0 plants were infested with whiteflies or 1 μM DAMP molecular peptide Pep1 for the indicated time. Values are mean ± s.e. (n = 4). (c) Schematic diagram of whitefly salivary effectors’ functional screening system showing the effector and luciferase reporter constructs used in (d). The *Arabidopsis* PDF1.2 promoter-driven luciferase was used as a reporter. *CaMV 35S* promoter-driven whitefly salivary protein genes were used as effector constructs. (d) Whitefly (*Bemisia tabaci*) salivary proteins (Bsp1–Bsp10) were screened for their ability to affect DAMP-induced plant immunity on *Nicotiana benthamiana* leaf. Immunity activator Pep1 (1 μM) was spread for 3 h before sampling. The fold increase of luciferase activation by Pep1 was calculated against a control group without Pep1 treatment. *CaMV 35S:*YFP was used as a control. Asterisks indicate significant differences in fold increase of luciferase activation between control and candidate saliva protein genes. Bars represent means ± s.e (n = 8) (*p < 0.05; **p < 0.01; ***p < 0.001; Student’s t-test for all the experiments). (e) Detection of Bsp9 protein in plants. Tomato plants were infested with whitefly for 72 h and all whiteflies in leaves were removed before sampling. Bsp9 protein was detected by Western blot using polyclonal antibody anti-Bsp9. Tomato leaves without whitefly infestation were used as a negative control, and whitefly total protein was used as a positive control. Stained gel bands of the large subunit of Rubisco (rbcL) were used as a loading control. (Online version in colour.)

**Bsp9 secretes from whitefly into plant cells**

*Bsp9* encodes a small protein with 112 amino acid residues and a molecular weight of 12.4 kDa (electronic supplementary material, table S3). The N-terminal of Bsp9 protein contains a 25 amino acid residue signal peptide with a
indeed detected in tomato leaves infested by whitefly, as anti-Bsp9 polyclonal antibody. The Bsp9 protein was protein in whitefly-infested tomato by immunoblot using secreted from whitefly into plant cells, we detected Bsp9 is rich in a leaf cells (electronic supplementary material, N. benthamiana expressed and highly accumulated in the cytoplasm of localization of Bsp9 in plants, Bsp9-YFP fusion protein was out whitefly feeding. To further check the subcellular expected, there is no detectable signal in tomato leaves with-out whitefly feeding. To further check the subcellular localization of Bsp9 in plants, Bsp9-YFP fusion protein was expressed and highly accumulated in the cytoplasm of N. benthamiana leaf cells (electronic supplementary material, figure S4a,b). The secondary structure of Bsp9 mature protein is rich in α-helix. To confirm whether Bsp9 protein can be secreted from whitefly into plant cells, we detected Bsp9 protein in whitefly-infested tomato by immunoblot using anti-Bsp9 polyclonal antibody. The Bsp9 protein was indeed detected in tomato leaves infested by whitefly, as well as protein extracted from adult whitefly (figure 2c). As expected, there is no detectable signal in tomato leaves without whitefly feeding. To further check the subcellular localization of Bsp9 in plants, Bsp9-YFP fusion protein was expressed and highly accumulated in the cytoplasm of N. benthamiana leaf cells (electronic supplementary material, figure S4c).

transmembrane domain (electronic supplementary material, figure S4a,b). The secondary structure of Bsp9 mature protein is rich in α-helix. To confirm whether Bsp9 protein can be secreted from whitefly into plant cells, we detected Bsp9 protein in whitefly-infested tomato by immunoblot using anti-Bsp9 polyclonal antibody. The Bsp9 protein was indeed detected in tomato leaves infested by whitefly, as well as protein extracted from adult whitefly (figure 2c). As expected, there is no detectable signal in tomato leaves without whitefly feeding. To further check the subcellular localization of Bsp9 in plants, Bsp9-YFP fusion protein was expressed and highly accumulated in the cytoplasm of N. benthamiana leaf cells (electronic supplementary material, figure S4c).

(e) Whitefly salivary protein Bsp9 interacts with
Arabidopsis transcription factor WRKY33

To investigate the molecular mechanism of how whitefly salivary protein Bsp9 promotes TYLCV accumulation in plants, we sought to identify Bsp9-targeted host factor(s). Therefore, a yeast two-hybrid experiment was conducted to screen an Arabidopsis cDNA library by using Bsp9 as the bait, and an immunity-related transcription factor AtWRKY33 was found as a putative positive interactor. Yeast transformants carrying AD-WRK33 and BD-Bsp9 were able to grow on SD-Leu-Trp-His selection plates with 2 mM 3-amino-1,2,4-triazole, whereas yeast transformants carrying AD and BD-Bsp9 constructs were unable to do so (figure 3a). To confirm the interaction between WRKY33 and Bsp9 proteins in vivo, we performed a BiFC assay in N. benthamiana. The N-terminus of the yellow fluorescent protein was fused in-frame to WRKY33 (nEYFP-WRK33) and C-terminus YFP was fused to Bsp9 (cEYFP-Bsp9). The constructs were transiently expressed in N. benthamiana leaf cells by Agrobacterium co-infiltration. A direct interaction between WRKY33 and Bsp9 was observed in the form of cytoplasmic speckles, which altered the nucleus localization of WRKY33 as a functional transcription factor to regulate downstream defensive gene expression (figure 3b). No fluorescence was detected when cEYFP-Bsp9 or nEYFP-WRK33 was co-expressed with nEYFP or cEYFP as a negative control. A split-luciferase complementation assay was further performed to confirm the interaction in vivo. Agrobacterium carrying the constructs of nLUC-Bsp9 and cLUC-WRK33 were infiltrated into N. benthamiana leaves, and the intensity of luminescence was increased only by the combination of nLUC-Bsp9 and cLUC-WRK33 compared with the vector control (figure 3c). Taken together, these results consistently prove that Bsp9 interacts with WRKY33.

(f) Bsp9 may disrupt the interaction between MPK6 and WRKY33

WRKY33 is an essential transcription factor in response to the attack of pathogens, but how it regulates immunity against whitefly and begomovirus is unknown. Bsp9 interacts with WRKY33 in the cytoplasmic bodies, raising a possibility that Bsp9 competes with MPK3 or MPK6 for the interaction with WRKY33.

A modified BiFC competitive protein-binding assay was used to test this hypothesis. Agrobacterium tumefaciens strains containing expression vectors for fusion proteins of MPK6 and WRKY33, together with 35S:Bsp9, were co-injected into N. benthamiana leaf cells. Yellow fluorescence was observed owing to the interaction between cEYFP-MPK6 and nEYFP-WRK33. Co-expression with a 35S:β-glucuronidase (35S:GUS) was used as a negative control. The interaction strength of MPK6-WRK33 as indicated by EYFP fluorescence intensity
was significantly decreased after the addition of Bsp9 protein (figure 4a). A negative control of GUS co-expression did not affect the interaction between MPK6 and WRKY33. These results demonstrate that Bsp9 interferes with the interaction between MPK6 and WRKY33, and, therefore, might disrupt a signal transduction event.

(g) WRKY33 is essential for Arabidopsis anti-whitefly resistance

Considering the reduced interaction between MPK6 and WRKY33 due to the interference of Bsp9, we hypothesized that Bsp9 may affect the WRKY33-mediated immune response. However, whether WRKY33 can mediate plant innate immunity against whitefly is still unclear. To test the regulation of WRKY33 in response to whitefly, we performed a whitefly two-choice preference assay. Electronic supplementary material, figure S5a shows that more whiteflies were attracted to wrky33 plants compared with Col-0 plants. Our previous work has indicated that suppression of JA-regulated repellent terpene biosynthesis in Arabidopsis makes the host more attractive to the whitefly vector [9], and thus we examined the expressions of Arabidopsis Terpene Synthase (TPS) genes such as TPS10, TPS14, TPS18 and TPS20. As expected, the expression levels of TPS10, TPS18 and TPS20 in the wrky33 mutant were greatly reduced compared with those of Col-0 plants (electronic supplementary material, figure S5b). In addition to the increased whitefly attraction, more eggs were laid by whiteflies on wrky33 mutants than on Col-0 plants (figure 4c). Furthermore, late
fourth whitefly nymphs were much more prevalent on \textit{wrky33} than on Col-0 plants (figure 4f). That whiteflies preferred and performed better on \textit{wrky33} plants indicates that WRKY33 is essential for plant resistance against whitefly. Taken together, our results imply that Bsp9 inhibits WRKY33-mediated anti-whitefly resistance.

(h) Bsp9 increases whitefly performance and transmission of TYLCV

Previous studies demonstrated that plants pre-infested with TYLCV-viruliferous whitefly could attract more vector insects [32]. Transient expression of Bsp9 inhibits WRKY33-mediated resistance, but whether the suppression of Bsp9 affects whitefly preference for better virus transmission is unknown. Thus, we generated transgenic plants overexpressing Bsp9. Intriguingly, we found that 35S:Bsp9-HA stable transgenic lines were more attractive to whiteflies compared with 35S:HA vector control plants (electronic supplementary material, figure S5c). Besides increased whitefly attraction, daily number of eggs laid per female whitefly on 35S:Bsp9-HA plants showed a significant increase compared with that of vector control plants, indicating that Bsp9 suppresses plant resistance against whitefly (figure 4e). The promoted whitefly preference and performance raised the conjecture of whether the manipulation of whitefly behaviours by Bsp9 can eventually affect the viral transmission efficiency from whitefly to plants. To test this, we detected the amount of whitefly virus transmission into \textit{Arabidopsis} plants which were fed on by TYLCV-viruliferous whiteflies for 72 h, at which time point no viral transcription could be detected in plants (electronic supplementary material, figure S6). Results showed that TYLCV titre in 35S:Bsp9-HA plants was threefold higher than that in vector control plants (figure 4f). Consistent with the promoted whitefly performance on the \textit{wrky33} mutant, the amount of virus transmission to \textit{wrky33} plants increased fourfold compared to that of vector control plants. Based on the above results, we conclude that the whitefly salivary protein Bsp9 suppresses WRKY33-mediated immunity to increase whitefly preference, performance and, in turn, eventually increases virus transmission.

4. Discussion

How have whiteflies become a successful supervector, able to transmit 300 species of viruses? [1] In this report, we provide a new layer of insight, in which a virus-induced whitefly salivary effector Bsp9 benefits both vector and virus. We have provided several lines of evidence that Bsp9 is a critical salivary effector in enhancing virus transmission. First, Bsp9 can effectively suppress the plant immune response activated by whitefly infestation (figure 2). Second, Bsp9 can be secreted into plants and interacts with the transcription factor WRKY33 (figures 2e and 3; electronic supplementary material, figure S4). Third, Bsp9 alters the localization of WRKY33 and affects the interaction between WRKY33 and MPK6, interfering with the WRKY33-mediated plant innate immunity against pathogen and insects (figures 3b and 4). Fourth, Bsp9 can effectively increase whitefly preference, performance and also TYLCV transmission (figure 4; electronic supplementary material, figure S5). Fifth, genomic analysis shows that Bsp9 is highly conserved in two invasive species of whitefly, MEAM1 and MED (electronic supplementary material, table S3). The coding region of the Bsp9 gene is highly conserved in these two whitefly species and there is only one non-synonymous mutation, indicating the critical role of Bsp9 for whitefly fitness on host plant adaption. Meanwhile, the high mutation rates in intron and untranslated regions suggest that a possible transcriptional level regulation may play an important role for whiteflies in response to biotic stresses, e.g. TYLCV infection in this report. Nevertheless, it is essential to clone Bsp9 homologues from native whiteflies and other Aleyrodoidea for understanding its role in whitefly invasion and begomoviral diseases pandemic in the world. With the rapid development of CRISPR/CAS-based single-base gene-editing technology [33], it is very interesting to look at the significance of single nucleotide differences of Bsp9 for the tripartite interactions of whitefly–plant–virus and environmental stress responses between MEAM1 and MED whiteflies.

While microbial pathogen effectors have been extensively studied for a long time, a number of functional approaches to identify effectors secreted by insects have only recently attracted attention [3–5]. However, the majority of the salivary gland transcripts encode small proteins, which lack sequence similarity to function-annotated proteins. Transcriptomic and proteomic analyses of aphid salivary glands or aphid saliva, combined in some cases with RNAi and plant stable overexpression approaches, have revealed the presence of potential effector proteins, such as Mp1 (PIntO1) and Mp2 (PIntO1), Mp10, Mp42, Mp55, Mp56, Mp57 and Mp58 in aphids [3,34–36]. However, most of the functional studies of insect effectors are designed on the basis of plant–pathogen interaction systems. For example, in most of these analyses, host immune responses are triggered by applying a pathogen-derived elicitor such as flg22 or elf18 [17], like aphid effector Mp10, which was functionally identified by the suppression of the oxidative burst induced by the bacterial elicitor flg22 [4]. But, this does not reflect the nature of plant–insect interaction. In the interaction between insects and plants upon feeding damage, DAMPs (damage-associated molecular patterns) are more likely to reflect the real situation of plant–insect interaction. In our study, we found that a DAMP Pep1 treatment could mimic the whitefly infestation on a plant, and the expression level of \textit{PDF1.2} may reflect changes of plant defence specific to whitefly. We developed a novel screening platform and successfully identified four candidate effectors, from the invasive MEAM1 whitefly, which can suppress Pep1-induced expression of \textit{PDF1.2}. Among them, Bsp9, which has the strongest suppression efficacy of plant immune responses, has been demonstrated as a functionally characterized whitefly salivary protein able to disrupt WRKY33-mediated immune signalling. Nevertheless, in the future, it will be promising to check Bsp9 knockout/down whitefly to look for other roles in whitefly biology and whitefly–plant interaction. Also, it will be interesting to check the effect of \textit{wrky33} mutation on the Bsp9 overexpression lines to further confirm whether WRKY33 is the major target of Bsp9.

Previously published studies show that WRKY33 is a pathogen-inducible transcription factor, whose expression is regulated by the MPK3/MPK6 cascade. In \textit{Arabidopsis}, WRKY33 functions in response to PTI signalling and also reprogramming of the expression of camalexin biosynthetic genes. WRKY33 expression is regulated by the MAPK cascades [22]. Previous studies found that MAPKs can be
activated by whitefly-mediated inoculation of TYLCV [37,38].

MPK3/MPK6 can phosphorylate WRKY33, the latter then binds to not only its own promoter but also the PAD3 promoter and positively regulates the expression of its bound genes. WRKY33 binds camalexin biosynthetic genes which encode proteins to produce defensive metabolites for pathogen resistance in Arabidopsis [39,40]. WRKY33 drives the metabolic flow to camalexin production challenged by pathogens. The interaction between WRKY33 and MPK3/6 in the nucleus cells plays an important role in the MAPK–WRKY33 cascade. WRKY33 is subject to post-translational modification by MAPKs and the expression of plant defensive genes. In begomovirus-infected whiteflies, whitefly saliva proteins can be induced by TYLCV and are secreted into plants during feeding. One of whitefly effectors Bsp9 can effectively suppress the expression of marker gene PDF1.2. Bsp9 interacts with plant transcription factor WRKY33 and interferes with the interaction of WRKY33 and MPK6, and therefore, suppresses the WRKY33-induced immune response. Thus, begomovirus might manipulate saliva effectors to suppress host immune responses to benefit whitefly fitness and virus spread.

However, the detailed mechanism of how TYLCV promotes the transcription of whitefly Bsp9 is still a subject of study in the laboratory.

5. Conclusion

In summary, we identify a new strategy for vector-borne viruses to conquer host immunity. In a healthy plant, whitefly feeding activates the PTI which triggers the activation of MAPKs and the expression of plant defensive genes. In begomovirus-infected whiteflies, whitefly saliva proteins can be induced by TYLCV and are secreted into plants during feeding. One of whitefly effectors Bsp9 can effectively suppress the expression of marker gene PDF1.2. Bsp9 interacts with plant transcription factor WRKY33 and interferes with the interaction of WRKY33 and MPK6, and therefore, suppresses the WRKY33-induced immune response. Thus, begomovirus might manipulate saliva effectors to suppress host immune responses to benefit whitefly fitness and virus spread.

The datasets supporting this article have been uploaded as part of the electronic supplementary material.

Authors’ contributions. J.Y. and N.W. conceived of the study and designed the experiments and drafted the manuscript. Experiments were mainly carried out by N.W., P.Z. and Y.M. The data were analysed by P.Z., X.H. and J.J. The manuscript was written by J.Y. and N.W. with contributions from R.F. and C.Z., Y.S. and X.Y. participated in the whitefly feeding and molecular analysis. Y.Z. provided the MEAM1 whitefly colony and the draft genomic sequence of MED species. All authors contributed to the analysis and interpretation of data, revised the manuscripts and gave final approval for publication.

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