Sogatella furcifera Saliva Mucin-like Protein Is Required for Feeding and Induces Rice Defences

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Abstract: The white-backed planthopper (WBPH), Sogatella furcifera, is one of the most important piercing-sucking pests of rice (Oryza sativa) in Asia. Mucin-like salivary protein (SFMLP) is highly expressed in the salivary glands of WBPH, which plays an important role in WBPH feeding. In this study, WBPH injected with dsSFMLP had difficulty in sucking phloem sap from rice plants, which significantly reduced their food intake, weight, and survival. In contrast, the knockdown of the SFMLP gene had only a marginal effect on the survival of WBPH fed an artificial diet. Further studies showed that silencing SFMLP resulted in the short and single-branched salivary sheaths secretion and less formation of salivary flanges in rice. These data suggest that SFMLP is involved in the formation of the salivary sheath and is essential for feeding in WBPH. Overexpression of the SFMLP gene in rice plants promoted the feeding of WBPH, whereas silencing the gene in rice plants significantly decreased WBPH performance. Additionally, it was found that overexpression of SFMLP in rice plants elicited the signalling pathway of SA (salicylic acid) while suppressing JA (jasmonic acid); in contrast, silencing of the SFMLP gene in rice plants showed the opposite results. This study clarified the function of SFMLP in WBPH feeding as well as mediating rice defences.

Keywords: Sogatella furcifera; mucin-like protein; salivary proteins; plant-insect interaction; rice defense response

1. Introduction

Plants and piercing-sucking pests have been engaged in a coevolutionary arms race in which herbivore saliva plays a significant role. Herbivores secrete salivary effectors into host plant cells to facilitate feeding, whereas plants recognize attackers and activate complex defence responses when attacked by herbivores [1,2].

Effectors produced by herbivores, which are proteins or other molecules, can affect host plant structures and functions [3]. Several effectors have been identified in Nilaparvata lugens (Stål), including β-1,3-glucanase [4], endo-β-1,4-glucanase [5], EF-hang calcium-binding protein [6], and mucin-like protein [7,8]. Activation of the effector aids N. lugens in overcoming the rice cell wall defence and facilitates continuous phloem feeding. Three effectors, i.e., vitellogenin [9], calcium-binding protein [10], and DNase II [11], reported in Laodelphax striatellus (Fallen), can improve its performance and attenuate rice defences when they are secreted into rice plants. In aphids, effectors have also been identified, such as CathB3, C002, Me10, Me23, Mp2, and Mp55, which aid in aphid survival and regulate host defence [12–17].

The defence responses of plants to phloem-sucking insect pests usually induce salicylic acid (SA) accumulation and suppression or only modest induction of jasmonic acid (JA) defences [18,19]. There are several functional genes that have been identified for the biosynthesis of phytohormones in inducing defence responses. For example, the NPR1 (non-expressor of pathogenesis-related genes 1) and ICS1 (isochorismate synthase 1) genes
are reported to be SA synthesis-related genes, and AOS2 (allene oxide synthase 2) and LOX (lipoxygenase) genes are involved in JA biosynthesis [11,20,21]. For instance, the gene Bl56, which is a whitefly salivary protein, promotes whitefly performance on tobacco and induces the SA signalling pathway in host plants [21]. The infestations of whiteflies and aphids on Arabidopsis thaliana reduce or only modestly induce JA defences but induce SA signalling pathway genes [22–24]. Activation of the SA pathway can improve the whitefly performance on tobacco and suppression of the JA signalling pathway [25]. The infestation of the white-backed planthopper (WBPH), i.e., Sogatella furcifera, increases SA-mediated rice plant defences, and the infection of the southern rice black-streaked dwarf virus in the WBPH also can upregulate SA expression levels to a lesser extent [20].

WBPH is a severe phloem-sucking rice pest in Asia, which can cause serious damage by feeding on rice sap and transmitting viruses. During feeding, WBPH stylets penetrate rice cell walls, and their salivary glands secrete both gelling and watery saliva into plant. The gelling saliva can quickly form a complete salivary sheath in the plant, which encases the full length of the stylet and provides mechanical stability and protection for WBPH feeding. In the transcriptome analysis of the WBPH salivary gland, the mucin-like protein gene (SFMLP) was found to be highly expressed in WBPH salivary glands. In insects, several salivary mucin-like proteins have been identified. A mucin-like protein in the Anopheles gambiae salivary glands helps it lubricate its mouthparts [26]. The salivary mucin-like protein of N. lugens plays an important role in the formation of its salivary sheath and feeding on host plants, and it is involved in plant–insect interactions [7,8]. However, the functions of SFMLP in WBPH are largely unknown.

Expression of double-stranded RNA (dsRNA), which is designed against important insect genes, in transgenic plants has provided protection against pests [27]. In recent years, expression of dsRNA of target genes in transgenic plants has been applied extensively against sap-sucking pests such as aphids [28–30], whiteflies [21,27], and N. lugens [31,32]. These results revealed that the salivary effectors played essential roles on the suppression of host plant immunity and promotion of insect performance. Therefore, these effectors would be the potential targets for phloem-sucking pest control by employing the transgenic plant-mediated RNAi technology.

Here, our results demonstrate that SFMLP is necessary for the formation of salivary sheath and feeding of WBPH on host rice plants. We report that overexpression of SFMLP in rice plants enhances WBPH performance and induces the SA signalling pathway, whereas silencing of SFMLP decreases WBPH performance and impairs the SA signalling pathway. All the results provide valuable aid for developing environmentally friendly, targeted control methods for WBPH.

2. Results
2.1. SFMLP Is Highly Expressed in WBPH Salivary Glands

The full-length cDNA of SFMLP contains a 2160 bp open-reading frame, which encodes 719 amino acid residues. SignalP- and TMHMM-predicted SFMLP constituted the signal peptide, with a likely cleavage site between residues 19 and 20. SFMLP has a high proportion of Ser (22.8%).

2.2. Bioassay of WBPH Injection with dsSFMLP

Double-stranded RNA (dsRNA) of SFMLP was synthesized and injected into 5th instar WBPH nymphs to conduct RNA interference (RNAi) experimental treatment. The results revealed that dsSFMLP showed a very strong silence effect, and it decreased the SFMLP transcript levels significantly (>99%) from the 1st day after treatment compared with the levels of WBPH injection with dsGFP (Figure 1).
The amount of honeydew excreted by WBPH was significantly reduced after silencing of SFMLP, with levels only 62.2% of those with silenced dsGFP (Figure 2A). The body weights of WBPH females and males injected with dsSFMLP both showed significant differences from those injected with dsGFP (Figure 2B). The body weights of new emerging females and males with silenced SFMLP were 86.74% and 85.46% of those of females and males with silenced GFP, respectively (Figure 2B).

Figure 1. Relative expression level of SFMLP in WBPH after injection of dsRNA. dsGFP indicates WBPH injected with GFP-dsRNA; dsSFMLP indicates WBPH injected with SFMLP-dsRNA. The mRNA expression level in the dsGFP group is designated as 1.0. ** indicates statistically significant differences (t-test: \( p < 0.01 \)). Bars, ±SEM.

2.3. The Effects of SFMLP Knockdown on Feeding Behavior

To investigate the role of SFMLP on feeding, we compared the survival rates within 1–10 days for WBPH developed from the 2nd–3rd instar nymphs between feeding on TN1 rice and the artificial diet in Parafilm sachets. The survival rates of WBPH nymphs with silenced SFMLP raised on TN1 rice decreased significantly on the 2nd day after injection, dropped to 20% on the 4th day after injection, and the nymphs died completely on the 9th day after injection compared with the dsGFP control group (Figure 3A). On the contrary,
there was no significant difference in the survival rates within 1–10 days between WBPH developed from the 2nd–3rd instar nymphs with silenced SFMLP and the dsGFP control group raised on the artificial diet (Figure 3B). These results reveal that SFMLP is important to feeding.

![Figure 3](image-url)  
**Figure 3.** Effects of SFMLP knockdown on the survival rates of WBPH. dsGFP indicates WBPH injected with GFP-dsRNA; dsSFMLP indicates WBPH injected with SFMLP-dsRNA. (A) The survival rates of WBPH feeding on TN1; (B) the survival rates of WBPH feeding on artificial diet. *** indicates statistically significant differences (t-test: p < 0.01). Bars, ±SEM.

To study the feeding activities of the WBPH, the electrical penetration graph (EPG) technique was used. There are five types of EPG signals based on the features of typical waves, which represent different insect activities described in materials and methods. The nonpenetration (NP) and xylem phase (N5) times were significantly increased after silencing of SFMLP, whereas the phloem-sap-ingestion phases were reduced significantly (N4-b) (Figure 4). The duration of the pathway phase (PP) or intracellular activity in the phloem (N4-a) have no significant differences between the dsGFP and dsSFMLP treatments (Figure 4).

![Figure 4](image-url)  
**Figure 4.** Effects of SFMLP knockdown on feeding behaviour of WBPH dsGFP indicates WBPH injected with GFP-dsRNA; dsSFMLP indicates WBPH injected with SFMLP-dsRNA. * and ** indicate statistically significant differences (t-test: p < 0.05 and p < 0.01). Bars, ±SEM.

2.4. *SFMLP* Is Necessary for Salivary Sheath Formation

Photographs of salivary gland morphology of the 5th nymphs were taken by a light stereomicroscope (Olympus SZX7). The morphologies of the WBPH salivary glands did not show significant differences between dsGFP and dsSFMLP. However, a colour change in
the A-follicle of the principal gland was found between them. The A-follicle remained dark black in the dsGFP-treated WBPH (Figure 5A), but it appeared light in the dsSFMLP-treated WBPH after paraformaldehyde fixation (Figure 5B).

![Figure 5. Effects of SFMLP knockdown on the morphology of the salivary gland of WBPH. (A) WBPH injected with dsGFP; (B) WBPH injected with dsSFMLP. PG indicates the principal gland. The arrow indicates the A-follicle of the principal gland.](image)

The salivary sheath formation was observed to further investigate the effects of SFMLP on feeding. After feeding on an artificial diet in Parafilm sachets for 1 day, WBPH salivary sheaths were analysed using scanning electron microscopy observation. The photographs revealed that WBPH subjected to dsSFMLP treatment produced significantly shorter and less-branched salivary sheaths than those produced by dsGFP-treated WBPH (Figure 6A–C).

![Figure 6. Effects of SFMLP knockdown on the morphology of the salivary sheath of WBPH (A) and (B) WBPH injected with dsGFP; (C) WBPH injected with dsSFMLP.](image)

We also used scanning electron microscopy to observe the penetration sites of WBPH on rice. On the same area of rice plants, the average number of feeding sites of dsSFMLP-treated WBPH was significantly lower than that of dsGFP-treated WBPH, and the numbers were 714 ± 76 and 314 ± 37, respectively (Figure 7A–D).

![Figure 7. Effects of SFMLP knockdown on the probing trace of WBPH. (A) WBPH injected with dsGFP; (B) WBPH injected with dsSFMLP; (C) enlargement of the probing trace. SS, salivary sheath; SH, stylet hole; SF, salivary flange; (D) the flange number of WBPH injected with dsSFMLP. After feeding on rice for 24 h, the salivary sheaths were collected and observed under SEM. ** indicates statistically significant differences (t-test: p < 0.01). Bars, ±SEM.](image)
2.5. The Influences of SFMLP Silencing and Overexpression Mediated by Transgenic Rice on WBPH Performance

To investigate the potential role of SFMLP as a plant-mediated transgenic target gene on WBPH management, transgenic rice plants with silenced and overexpressed SFMLP were generated. The transgenic (silencing and overexpression) and WT rice plants had no apparent phenotypic differences. The transcript levels of SFMLP were decreased by 52–57% in WBPH feeding on the SFMLP-silenced transgenic plants for 48 h and 64 h compared to those feeding on the WT plants (Figure 8A). In contrast, the transcript levels of SFMLP were increased by 136–310% in WBPH feeding on the SFMLP-overexpressing transgenic lines for 48 h and 64 h compared with those feeding on the WT plants (Figure 8B).

The honeydew excreted by WBPHs feeding on the SFMLP-silenced transgenic plants was decreased by 54.41% for the adults feeding for 72 h (Figure 9A). The body weights of the adults feeding on the SFMLP-silenced transgenic lines for 72 h were decreased by 44.76% (Figure 9B). In contrast, the honeydew excreted by WBPHs feeding on the SFMLP-overexpressing transgenic lines was increased by 324% for the adults feeding for 72 h (Figure 9C). The body weights of the adults feeding on the SFMLP-overexpressing transgenic lines for 72 h were increased by 106% (Figure 9D).

Figure 8. Relative expression levels of SFMLP in WBPHs feeding on transgenic plants. (A) SFMLP expression levels of WBPH feeding on SFMLP-dsRNA-transgenic plants (silence) for 48 h and 64 h. (B) SFMLP expression levels of WBPH feeding on SFMLP-overexpressing transgenic plants (overexpression) for 48 h and 64 h. * indicates statistically significant differences (t-test: \( p < 0.05 \)). Bars, ±SEM.

Figure 9. Effects on performance for WBPH feeding on transgenic plants. (A) Honeydew and (B) weight gains of WBPH feeding on SFMLP-dsRNA-transgenic plants (silence). (C) Honeydew and (D) weight gain of WBPH feeding on SFMLP-overexpressing transgenic plants (overexpression). ** indicates statistically significant differences (t-test: \( p < 0.01 \)). Bars, ±SEM.
2.6. Transgenic Rice-Mediated Plant Defence Responses

The relative expression levels of the SA marker genes (NPR1 and ICS1) and the JA marker genes (LOX and AOS2) in the silenced and overexpressed transgenic rice plants showed significant differences from those of wild-type plants. In the SFMLP-silenced transgenic rice plants, they showed a lower expression level in SA pathway-NPR1 and a higher expression level in JA pathway-LOX than those in wild-type plants (Figure 10A). In contrast, in SFMLP-overexpressing transgenic rice plants, the SA pathway-NPR1 was expressed at a higher level, and the JA pathway-LOX showed a lower expression level than those in wild-type plants (Figure 10B).

![Figure 10.](image)

**Figure 10.** Relative expression levels of JA marker and SA marker genes in transgenic rice. (A) The expression levels of four genes in SFMLP-dsRNA-transgenic plants (dsSFMLP) and wild-type plants (WT); (B) the expression levels of four genes in SFMLP-overexpressing transgenic plants (SFMLP) and WT plants. * and ** indicate statistically significant differences (t-test: p < 0.05 and p < 0.01). Bars, ±SEM.

After infestation by WBPH, in the wild-type plants, NPR1 was expressed at higher levels than in those that were not infested, and LOX and AOS2 were expressed at lower levels than in those that were not infested (Figure 11A). In the SFMLP-silenced transgenic rice plants, LOX and AOS2 were expressed at higher levels than those in non-infested plants, and NPR1 was expressed at lower levels than those in non-infested plants (Figure 11B). In the SFMLP-overexpressing transgenic rice plants, NPR1 and ICS1 were expressed at higher levels than those in non-infested plants, and LOX was expressed at lower levels than those in non-infested plants (Figure 11C).

![Figure 11.](image)

**Figure 11.** Relative expression levels of JA marker and SA marker genes in rice plants infested by WBPH. (A) The expression levels of genes in the WT plants and WT plants infested by WBPH (WT-infested); (B) the expression levels of genes in the SFMLP-dsRNA-transgenic plants (dsSFMLP) and dsSFMLP plants infested by WBPH (SFMLP-infested); (C) the expression levels of genes in the SFMLP-overexpressing-transgenic plants (SFMLP) and SFMLP plants infested by WBPH (SFMLP-infested). * and ** indicate statistically significant differences (t-test: p < 0.05 and p < 0.01). Bars, ±SEM.
3. Discussion

Mucin-like proteins have been studied to exist in many insects, such as in piercing-sucking planthoppers [8,33], leafhoppers [34], and several mosquito species [35]. The mucin-like proteins of the three major planthoppers, *S. furcifera*, *N. lugens*, and *L. striatellus*, are all rich in Ser with similar proportions (*S. furcifera* 22.8%, GenBank KX670544; *N. lugens* 22.4%, GenBank KY348750; *L. striatellus* 22.51%, GenBank JF502033), which are significantly different from those of *Nasonia vitripennis* (10.44%, GenBank NM_001130052) and *Holotrichia oblit†a* (4.57%, GenBank JF681187). It exhibits close ancestral relationships among the planthopper species compared with other insects. Ser can provide attachment sites for the formation of salivary sheaths. In aphids, the salivary protein SHP was found to aid the solidification of gelling saliva [36–38].

Our previous study revealed that SFMLP was highly expressed in the WBPH salivary gland, and the expression pattern of SFMLP was generally the same as that of mucin-like protein in *N. lugens* [8,33,39]. Mucin-like protein was found to be related to the salivary sheath formation of *N. cincticeps* and *N. lugens*, and feeding was affected significantly after mucin silencing [7,8,34,40]. In our study, the transcription of SFMLP was significantly reduced after its expression was knocked down, which indicates that introduction of double-stranded RNA (dsRNA) of SFMLP into WBPH causes the inhibition or degradation of the complementary mRNA of SFMLP. WBPHs silenced with SFMLP have incomplete, shorter, and less-branched salivary sheaths than those of control WBPH. As a result, the salivary sheaths of SFMLP-RNAi WBPH cannot reach the sieve tube, and WBPH will spend more time in the nonpenetration and pathway phases and less time feeding on the phloem. This decreased the amount of food intake, nymph mass, survival rate, and fecundity of the WBPH. In contrast, silence of SFMLP did not affect the ability of WBPH to feed on an artificial diet. These results suggest that SFMLP is necessary for WBPH to probe the rice plants and obtain the phloem sap. Based on the above results, it can be deduced that SFMLP is an essential component of the WBPH salivary sheath.

When feeding, herbivores secrete salivary effectors into host plant cells to elicit or suppress plant defence responses [1,2]. For instance, infestation by *N. lugens* can increase the levels of H$_2$O$_2$, SA, and ethylene but not JA in rice plants [41]. Mucin-like protein of *N. lugens* triggered rice defence responses associated with the JA signalling pathway revealed by the callose deposition maker [8] although endo-$\beta$-1,4-glucanase secreted by *N. lugens* did not elicit the rice defence-related signal molecules, including SA, JA, and jasmonoyl-isoleucine (JA-Ile) [5]. Wang et al. [42] found that the infestation of WBPH gravid females significantly increased the levels of JA, JA-Ile, and H$_2$O$_2$ and reduced the level of ethylene in rice although WBPH nymph infestation had no effect on these signal molecules. After infection by WBPH infected with the southern rice black-streaked dwarf virus, rice plant defences can partially be elicited, which is related to the smaller increase in SA and greater decrease in JA than nonviruliferous WBPH-infested plants [20]. In our study, after infestation by WBPH in rice, the relative expression levels of the SA marker gene NPR1 were higher, and the JA marker genes LOX and AOS2 were lower than those in wild-type plants. Overexpression of the Bt56 gene in tobacco elicits the SA-signalling pathway and promotes the performance of the whitefly, whereas silencing the whitefly Bt56 gene significantly decreases its feeding on host plants and makes it to lose the ability to induce the SA pathway [21]. Due to the salivary effector Bt56 being an SA elicitor, it promoted the whitefly performance on tobacco.

In this study, we obtained SFMLP-overexpressing and SFMLP-silenced transgenic plants to investigate their control effects on WBPH. For the overexpressed and silenced transgenic rice plants, the relative expression levels of the SA-signalling pathway and JA-signalling pathway genes showed the opposite results. WBPH infestation or not, overexpression of the SFMLP gene in planta elicits the SA signalling pathway corresponding to the upregulation of NPR1 or ICS1, and silencing of the SFMLP gene suppresses the SA signalling pathway corresponding to the downregulation of NPR1. Therefore, from the above results, we propose that SFMLP triggers rice defence responses by eliciting
SA and suppressing the JA signalling pathway. Furthermore, transgenic plant-mediated SFMLP RNAi decreased WBPH performance on rice plants and enhanced rice resistance to WBPH. In contrast, overexpression of SFMLP in rice promoted the feeding of WBPH on rice plants and impaired rice defences to WBPH. From the above results, we propose that overexpression of SFMLP in rice promoted WBPH performance because it activated the SA-signalling pathway. On the contrary, silence of SFMLP in rice decreased WBPH performance due to the repression of the SA-signalling pathway.

Host-induced gene silencing (HIGS) has great potential for effective control pest by generating transgenic plants carrying double-stranded RNAs (dsRNAs)-targeted genes against essential pest genes, and this technology is considered to the promising alternative to chemical pesticides [43]. The silence of LeECPI in L. striatellus decreases its fecundity and survival rates, and rice-mediated LeECPI knockdown enhanced rice resistance to L. striatellus [10]. When Vg of L. striatellus is overexpressed in rice plants, it can significantly hinder hydrogen peroxide (H2O2) accumulation and promote insect performance [9]. The fecundity of aphids was reduced after feeding on transgenic plants that were mediated by RNAi of salivary effector genes (Mpc002, MpPIntO1, Mp55, MpPIntO2) [14,17,28,29,44]. Therefore, these effectors might be the potential plant-mediated RNAi targets for aphid control.

In summary, our results reveal that SFMLP plays dual roles as a component in WBPH sheath formation and activating the SA signalling pathway of rice defence responses. Furthermore, silencing or overexpressing SFMLP genes via transgenic rice plants can significantly influence WBPH performance in rice plants. This work not only deepens our knowledge of the physiological role of SFMLP and its effects on understanding plant-herbivore interactions but also provides a feasible strategy to control WBPH infestation by applying SFMLP-silenced transgenic rice under field conditions.

4. Materials and Methods
4.1. Plant Materials and Insects
Taichung Native 1 (TN1, a WBPH-susceptible rice cultivar), Kitaake (KIT, a WBPH-susceptible rice cultivar), and transgenic KIT lines with SFMLP overexpression or silencing were used in this work. All rice plants were grown within 100-mesh insect-proof cages (80 × 80 × 80 cm) in a greenhouse (30 ± 5 °C, 15 L: 9 D). WBPH colonies were originally obtained from rice fields in Xing’an County (Latitude 25.61° N, Longitude 110.67° E), Guilin City, Guangxi Zhuang Autonomous Region, China. The population was maintained on TN1 rice seedlings in a climate-controlled room at 27 ± 1 °C with a relative humidity (RH) of 75 ± 10% and a photoperiod of 16:8 (L:D).

4.2. RNAi Experiment
SFMLP in watery saliva of WBPH was identified by shotgun LC–MS/MS analysis, and the sequence of SFMLP from our transcriptomic analysis was similar to the full length of the WBPH mucin-like gene deposited in GenBank (accession number KX670544). In our previous study, SFMLP transcripts of WBPH were detected at higher levels in the salivary gland than in the head, gut, testes, ovary, and remaining body [39]. A 524 bp SFMLP fragment and a 594 bp control gene GFP fragment were amplified with primers harbouring a T7 promoter sequence (Table 1). dsSFMLP and dsGFP were synthesized using the HiScribe™ T7 Quick High Yield RNA Synthesis Kit (New England Biolabs, Ipswich, MA, USA) according to the manufacturer’s protocol. The dsRNA was resuspended in nuclease-free water to a concentration of 4 μg/μL and stored at −80 °C until use.

The 2nd or 5th instar WBPH nymphs (for details, see descriptions of different experiments) were anaesthetized with carbon dioxide for approximately 20 s and placed on 1% (w/v) agarose plates with their abdomens facing up for injection. A 25 nL (for the 2nd instar nymph) and 50 nL (for the 5th instar nymph) volume of dsSFMLP or dsGFP was injected into each insect at the junction between the prothorax and mesothorax using a microprocessor-controlled Nanolitre 2010 injector (World Precision Instruments). After injection, the injected WBPH were reared on rice cv TN1 plants for the following experiments.
Table 1. The primers used in this study.

| Primers          | Primer Sequence (5′-3′)                              |
|------------------|------------------------------------------------------|
| qPCR             |                                                     |
| mucin-F          | TCATCTTCTGGGTGGCATTC                                  |
| mucin-R          | GGGGGGTGTGAAAATAGTA                                  |
| L9-F             | CAAGATGAGAGCCGTGTA                                   |
| L9-R             | CGAGTTGGAACAGCATGAC                                  |
| L10-F            | GCGACCTCATGGGCTGCA                                   |
| L10-R            | CACTCTAGCCACTGTC                                |
| SA-NPR1-F        | TTTCCGATGGAGGCAAGAG                                  |
| SA-NPR1-R        | GCTGTCAATCCGAGCTAAGTGTT                             |
| JA-LOX-F         | GCAATCCCAAACAGCACATC                                  |
| JA-LOX-R         | AATAAGATTTGGGAGTACATA                                 |
| RNAi             |                                                     |
| dsmucin-T7-F     | TAATACGACCTCATATAGGGTACTAC              |
| dsmucin-T7-R     | AAAACCTCTCCAAATC                                    |
| dsGFP-T7-F       | TAAATACGACTCATATAGGGA                              |
| dsGFP-T7-R       | AGAAAGACTTTTCACTGG                                   |
| dsGFP-T7-R       | TAAATACGACTCATATAGGGGATGCAAA                      |
| dsGFP-T7-R       | CCGATCCACTTC                                       |
| Expression in plant |                                                   |
| Overexpression-Mucin-F | CCG gatata ATGAGGT-                          |
| Overexpression-Mucin-R | GTTCTCGATGTATCG                                |
| dsGFP-T7-R       | TAAGAAGCACTTTTCACTGG                             |
| dsGFP-T7-R       | GGATCTTCTCAGTACGTGCG                               |
| dsGFP-T7-R       | CTCGAGccataagctcgtacagtacgc                      |

4.3. Bioassay of WBPH Injected with dsRNA

To measure the survival rates of WBPH, the 2nd instar nymphs injected with SFMLP or GFP dsRNA were allowed to feed on TN1 plants or artificial diet. In the survival rate tests on rice plants, 20 WBPH nymphs were transferred to a glass tube (5 cm diameter × 40 cm height) with 1-month-old TN1 rice plants grown as described above, and the number of surviving nymphs was recorded daily for 10 days. For the survival rate tests on the artificial diet, a transparent plastic bottle (3 cm diameter × 5 cm height) was used as the feeding chamber, and the top lid was covered with sachets formed from two layers of stretched Parafilm (4 times the original area) containing 200 µL artificial diet with 2.5% sucrose solution. A 2% (w/v) agarose plate was placed at the bottom of the bottle to maintain humidity. Holes (6 mm in diameter) were made on the bottle body for ventilation. The artificial diet was changed once every 24 h. Twenty nymphs injected with SFMLP or GFP dsRNA were released into each bottle, and the number of surviving nymphs was recorded daily for 10 days. The survival rate tests on rice plants and on the artificial diet were both repeated five times.

To assess the effect of SFMLP knockdown on WBPH feeding, the 5th instar nymphs at 1 day after the injection of SFMLP or GFP dsRNA were starved for 1 h, and one nymph was transferred into a pre-weighed small Parafilm bag (4 × 4 cm), which was affixed to the rice stems. The nymph was transferred out after feeding for 24 h, and the Parafilm sachet was reweighed. The weight difference of the Parafilm bag was defined as the WBPH honeydew secretion. The experiment was replicated 25 times.

To investigate the influence of SFMLP knockdown on WBPH weight gain, 5th instar nymphs at 1 day after the injection of SFMLP or GFP dsRNA were weighed and allowed to feed on rice plants until new brachypterous adults emerged. The weight differences of the female and male adults (within 1 day of emergence) with the 5th instar nymphs were defined as female and male adult weight gain. The experiment was replicated 25 times.
4.4. EPG Recording of WBPH Feeding Behavior

The feeding behaviour of the WBPH was recorded using a Giga-8 direct current electrical penetration graph (DC-EPG) amplifier with a $10^9$-$\Omega$ input resistance in a Faraday cage (manufactured by Wageningen University, Wageningen, The Netherlands). The 5th instar WBPH nymphs at 1 day after the injection of SFMLP or GFP dsRNA were starved for 1 h before the EPG recordings. The WBPH attachment and amplifier connection methods were the same as those described by Lei et al. [45]. Each insect was continuously recorded for 6 h, and 10 to 15 replications were recorded in each treatment. Five distinctive output waveforms were interpreted with reference to the studies by Khan and Saxena [46] and Seo et al. [47], including NP for nonpenetration, PP (N1 + N2 + N3) for the pathway phase (including penetration initiation, salivation and stylet movement, and extracellular activity near the phloem), N4-a for intracellular activity in the phloem, N4-b for phloem sap ingestion, and N5 for the xylem phase. The EPG parameters were calculated for each recording with reference to Sarria et al. [48] and Lei et al. [45].

4.5. Observation of Salivary Glands and Salivary Sheaths of WBPH Injected with dsRNA

To assess the effects of dsRNA injection on the morphology of salivary glands, 5th instar WBPH nymphs at 1 day after the injection of SFMLP or GFP dsRNA were allowed to feed on TN1 rice until the emergence of adults. The salivary glands of adults within 1 day of emergence were dissected using micro forceps (Shanghai Medical Instruments Ltd., Corp., Shanghai, China) in $1 \times$ phosphate buffered solution and transferred into 4% paraformaldehyde buffer for fixation for 30 min. The morphology of the salivary gland was observed under an anatomical lens (Leica Microsystems GmbH, Wentzler, Germany). The experiment was replicated 10 times.

To explore the effects of dsRNA injection on the morphology of salivary sheaths, the 5th instar WBPH nymphs injected with SFMLP or GFP dsRNA were allowed to feed on TN1 rice for 1 day, and 30 nymphs were transferred into the feeding chamber, which was a transparent plastic bottle (5 cm long and 3 cm in diameter) with the top lid covered with sachets formed from two layers of stretched Parafilm (4 times the original area) containing 200 $\mu$L 2.5% sucrose solution. A 2% (w/v) agarose plate was placed at the bottom to maintain humidity, and holes (6 mm in diameter) were made on the bottle body for ventilation. After feeding for 1 day, the inner Parafilm layer was washed three times using $1 \times$ phosphate-buffered solution and dried in a vacuum desiccator for morphological observation using scanning electron microscopy. The experiment was replicated three times.

To observe the feeding frequencies, the 5th instar WBPH nymphs injected with SFMLP or GFP dsRNA were allowed to feed on TN1 rice for 1 day, and 30 nymphs were transferred into the feeding chamber, which was a plastic tube (4 cm diameter and 6 cm long) affixed to the base of a rice stem and covered by sponges at both ends, leaving an approximately 4 cm long feeding space inside. After feeding for 1 day, the feeding site was collected and fixed with 4% glutaraldehyde for 48 h and then 1% osmium tetroxide for 3 h. After washing and dehydration, the feeding sites were observed using scanning electron microscopy. The experiment was replicated three times.

4.6. Transgenic Rice Plant Development and the WBPH Bioassay

To obtain the SFMLP-silenced transgenic rice lines, the 250 bp cDNA fragment was inserted into a Gateway pENTR/D-TOPO cloning vector, which carried two recombination sites for the LR Clonase reaction. Then, the fragment derived from a target gene was transferred into a pANDA destination vector to generate a hairpin RNAi. To obtain the SFMLP-overexpressing transgenic rice lines, the full CDS region containing a 2160 bp cDNA fragment was inserted into the PRHV cloning vector. The final RNAi and overexpression vectors were introduced into Agrobacterium tumefaciens (strain EHA105) employing the heat-shock method. The construct was transformed into the rice plants of the variety KIT via an Agrobacterium tumefaciens-mediated method. Transgenic rice plants, either overexpressing or silencing SFMLP, were developed by the Institute of Crop Sciences, Chinese Academy
of Agricultural Sciences. Integration of the SFMLP fragment in transgenic rice lines was determined via PCR and quantitative real-time PCR (RT–qPCR) (Figures S1 and S2). The primers used for dsMucin detection are listed in Table 1. The target plants were screened according to the above methods for subsequent experiments.

To investigate the SFMLP expression level of WBPH feeding on silenced and overexpression transgenic rice plants, single 1-month-old rice plants (transgenic rice lines and wild-type plants) were planted in one plastic pot (8 cm diameter and 27 cm height) and maintained as described above. A total of ten 5th instar WBPH nymphs were released into pots and transferred out after feeding for 48 h or 64 h for RNA isolation and cDNA synthesis. The cDNA samples were subjected to RT–qPCR to detect the expression level of the SFMLP gene. The experiment was repeated three times.

To assess the effect of SFMLP silencing or overexpression in rice plants on WBPH performance, 1-month-old rice plants (transgenic rice lines and wild-type plants) were planted in plastic pots (8 cm diameter and 13.5 cm height). After starvation for 1 h, the adults (within 1 day of emergence) were weighed, and then, one adult was transferred into a pre-weighed small Parafilm bag (4 × 4 cm), which was affixed to the rice stems. The adult was transferred out after feeding for 72 h, and the adult and Parafilm sachets were reweighed. The differences in the two weights for Parafilm bags and adults were defined as WBPH honeydew secretion and weight gain, respectively. The experiment was replicated 25 times.

4.7. Analysis of the Relative Expression of Phytohormone-Related Genes

To quantify the transcript levels of phytohormone-related genes in leaf sheaths of transgenic and wild-type rice plants infested by WBPH, 1-month-old rice plants were planted in plastic pots (8 cm diameter and 27 cm height). After starvation for 1 h, a total of twenty-five 5th instar nymphs were released into each pot and maintained as described above. Rice stems (5–10 cm) from the middle parts of the leaf sheath were collected from each pot, and then, all samples were ground in liquid nitrogen for RNA isolation and cDNA synthesis. The experiment was repeated four times.

The transcript levels of SA- and JA-related genes in rice leaf sheaths were quantified using RT–qPCR for the three treatment groups (Table 1). The extraction of total RNA from the leaf sheath samples and the synthesis of cDNA were described by Li et al., 2016 [40]. qPCR was performed using the Bester SybrGreen qPCR mastermix (DBI Bioscience, Germany) with the 7500 Sequence Detection System (Applied Biosystems, Foster City, CA, USA). Amplification reactions were performed in a 20 µL final volume containing 10 µL of TB Green qPCR premix Ex Taq II, 0.8 µL of forward primer (10 µM) and reverse primer (10 µM) pairs (Table 1), 0.4 µL of Rox II, 5 µL of cDNA (100 ng/µL), and 3 µL of sterilized H2O. The reaction conditions were as follows: 95 °C for 30 s followed by 40 cycles of 5 s at 95 °C and 34 s at 60 °C, followed by melt curve stages. Blank controls without template were included in each experiment. qPCR was performed for three samples, and each sample had three technical repeats. The comparative $2^{-\Delta\Delta CT}$ method was used to calculate the relative gene expression levels in different samples [49].

4.8. Statistical Analysis

All the results were based on the mean values excluding the morphology figure. The statistical significance of differences between treatments and control groups was assessed by Student’s t-test (* $p < 0.05$, ** $p < 0.01$). The error bars represent the means ± standard errors (SE). All analyses were conducted in IBM SPSS Statistics version 22.0 (SPSS Inc., Chicago, IL, USA).

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