Symbiotic polydnavirus of a parasite manipulates caterpillar and plant immunity

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Obligate symbioses occur when organisms require symbiotic relationships to survive. Some parasitic wasps of caterpillars possess obligate mutualistic viruses called “polydnaviruses.” Along with eggs, wasps inject polydnavirus inside their caterpillar hosts where the hatching larvae develop inside the caterpillar. Polydnaviruses suppress the immune systems of their caterpillar hosts, which enables egg hatch and wasp larval development. It is unknown whether polydnaviruses also manipulate the salivary proteins of the caterpillar, which may affect the elicitation of plant defenses during feeding by the caterpillar. Here, we show that a polydnavirus of the parasitoid Microplitis croceipes, and not the parasitoid larva itself, drives the regulation of salivary enzymes of the caterpillar Helicoverpa zea that are known to elicit tomato plant-defense responses to herbivores. The polydnavirus suppresses glucose oxidase, which is a primary plant-defense elicitor in the saliva of the H. zea caterpillar. By suppressing plant defenses, the polydnavirus allows the caterpillar to grow at a faster rate, thus improving the host suitability for the parasitoid. Remarkably, polydnaviruses manipulate the phenotypes of the wasp, caterpillar, and host plant, demonstrating that polydnaviruses play far more prominent roles in shaping plant–herbivore interactions than ever considered.

herbivore | plant defense | parasitoid | host immunity | phytophyme

Plant–herbivore interactions do not occur in isolation, but are part of a complex, multitrophic network of associated microscopic and macroscopic organisms termed “the phytophyme” (1). Microbes associated with herbivores are one component of the phytophyme, and they may perform important functions in facilitating host use by aiding in digesting plant tissues, detoxifying plant toxins, directly supplying nutrients, or facilitating protection from natural enemies (2). Microbial mediation of plant–herbivore interactions may also occur when microbes directly interfere with the perception of herbivores by plants. Plant-defense induction depends upon the plant’s ability to detect cues associated with herbivory (3, 4). Plants are able to recognize herbivore cues such as touch, wounded, oviposition, and the feeding cues from oral secretions (e.g., saliva and/or regurgitant) (3, 5). However, microbes present in the herbivore’s gut can alter the composition of oral secretions and thus may trigger or suppress plant defensive responses (6, 7).

Another important player in phytophyme interactions is the endoparasitoid wasp of insect herbivores. Some parasitoid species possess obligate mutualistic polydnaviruses (PDVs), which are transferred to their caterpillar hosts when the parasitoid deposits their egg(s) within their hosts (8). PDV genomes are stably integrated in the genomes of parasitoid wasps (8). The infection cycles of PDVs occur between two hosts: PDV particles replicate only in the wasps, but infect tissues (including salivary glands) of, and express viral genes in, their caterpillar hosts (9). PDVs use virulence factors to manipulate the immune systems of their caterpillar hosts to enable the survival of parasitoid eggs and larvae (8, 10, 11). PDVs are associated with parasitic wasps belonging to the Braconidae and Ichneumonidae families, respectively (8).

The ability of PDVs to interfere with the expression of plant defenses has not been reported, but a few investigations indicate that parasitoids can alter plant responses to herbivores. Poelman et al. (12) showed that several parasitoid species of pierid caterpillars differentially elicited defense responses in the host plant Brassica oleracea. It was striking that the species of parasitoid had a stronger effect on the induced plant responses than the identity of the caterpillar host. The parasitoids directly affected the caterpillar’s oral secretions and its subsequent ability to elicit defense responses, but the role of PDVs in mediating possible changes in salivary components was not reported (12). In another study with B. oleracea using the caterpillar Trichoplusia ni and its parasitoid Copidosoma floridanum, parasitized caterpillars induced 1.5 times higher levels of indole glucosinolate defenses in the plant compared with the nonparasitized caterpillars (13). The differential induction in this case was attributed to increased feeding in the parasitized caterpillars (13). In contrast to parasitoids that carry polydnavirus symbionts, this particular parasitic wasp maintains and even enhances the host immune system (14). This wasp species is in the Encyrtidae family, members of which do not possess polydnavirus symbionts.

Here we report on the multitrophic role of a symbiotic PDV in mediating the phenotypes of the caterpillar and its host plant using the braconid parasitoid Microplitis croceipes, the host...
noctuid caterpillar *Helicoverpa zea*, and the host plant tomato as an experimental system. The parasitoid uses a PDV, the Bracovirus (McBV), to immunosuppress its caterpillar host *H. zea*, which allows eggs to hatch and the larvae to feed on the hemolymph of the caterpillar (15). In contrast to the study with parasitism in *T. ni*, this parasitoid does not cause hosts to spend more time feeding (16). An earlier study found that *M. croceipes* strongly suppressed protein synthesis in the salivary glands of its host *Heliothis virescens* (17), although effects of caterpillar saliva on host plants were not investigated. Salivary glands are the major source of oral secretions in many caterpillars including *H. zea* (18), thus the ability of PDVs to suppress salivary proteins seems likely based upon previous findings with *M. croceipes*. Notably, glands of *H. virescens* and *H. zea* contain a highly abundant immune-related protein, glucose oxidase (GOX), which also acts as an elicitor of plant defenses during feeding (19, 20). The enzymatic products of GOX are D-glucono-δ-lactone and H₂O₂; the latter product possesses antimicrobial activity and acts as a second messenger for the induction of defense proteins such as polyphenol oxidase and proteinase inhibitors in tomato plants (21).

**Results and Discussion**

To determine if parasitism by *M. croceipes* affects the ability of *H. zea* to induce plant defenses, both parasitized (P) caterpillars and nonparasitized (NP) caterpillars were allowed to feed on tomato leaves for 10 h while the total feeding damage between treatments was kept consistent by restricting caterpillars to feed within a cage. Forty-eight hours later the activities of plant-defense proteins [i.e., trypsin inhibitor (TI) and polyphenol oxidase (PPO)] were assayed on the damaged leaves. P-caterpillars induced significantly lower levels of trypsin inhibitor and PPO activities (Fig. 1) than did the NP-caterpillars. Both protease inhibitors and PPO are known to reduce the growth rate of caterpillars, and of *H. zea* in particular (22–24).

We then examined the transcript levels of defense genes encoding PPO, protease inhibitors, and other defense proteins (at 24 h post feeding) known to be induced by *H. zea* saliva (20, 25) and found that P-caterpillars consistently induced lower levels of transcripts encoding plant defensive proteins such as PPO (PPOB, PPOE, PPOF), threonine deaminase (TD2), and proteinase inhibitors (AspPI, CysPI) than did NP-caterpillars (SI Appendix, Fig. S1). Threonine deaminase degrades the essential amino acid threonine in the guts of caterpillars, thus impairing their growth (26). Transcripts encoding proteins for biosynthesis of toxic secondary metabolites including terpenes (MTSI), phenolics (PAL5), and glycoalkaloids (GAME4) were also attenuated in plants fed on P-caterpillars compared with their nonparasitized counterparts (SI Appendix, Fig. S1).

To determine if caterpillar saliva was responsible for the observed differential plant responses, we applied saliva from P- and NP-caterpillars to plant wounds and then assayed for a subset of the plant-defense genes including PIN2, TD2, and AspPI and the defense protein PPO. Saliva from P-caterpillars induced lower levels of PPO and the three defense genes compared with saliva from NP-caterpillars, indicating that parasitism alters salivary components responsible for eliciting defenses (SI Appendix, Fig. S2). We then measured the effect of parasitism on GOX, which is a highly abundant immunity-related protein in the saliva of *H. zea*, that elicits antiherbivore defenses in tomatoes, including the aforementioned plant-defense genes (20). P-caterpillars had dramatically lower GOX transcript levels than did NP-caterpillars at 2 d post parasitism (Fig. 2A), and GOX enzymatic activity was significantly lower at 4 and 6 d following parasitism (Fig. 2B). The response of tomato to GOX is dose-dependent: leaves treated with increasing levels of GOX triggered higher levels of protease inhibitors (SI Appendix, Fig. S3).

To determine if parasitism by *M. croceipes* is directly responsible for suppressing GOX activity, we purified McBV from the calyx region of the wasp ovaries, injected the virus into caterpillars, and then measured GOX activity and expression after verifying McBV infection by PCR. McBV markedly suppressed GOX enzyme and transcript levels in a similar fashion as parasitism by *M. croceipes* wasps (Fig. 3 and SI Appendix, Fig. S4).

Feeding by caterpillars that were injected with PDV induced significantly lower defense gene expression (e.g., PPOB, CysPI, AspPI, TD2, PIN2) and defense protein activity (TI, PPO) than did the saline-injected caterpillars (Fig. 4), which indicates that McBV is the primary driver in mediating these multitrophic interactions.

To determine if attenuated plant defenses benefited caterpillar growth, we fed P-caterpillars leaves from plants that were previously treated with saliva from NP- and P-caterpillars. The growth rate of P-caterpillars was higher when fed on leaves treated with saliva from P-caterpillars compared with leaves treated with saliva from NP-caterpillars (SI Appendix, Fig. S5). These results suggest that the improved growth of the P-caterpillars would also benefit parasitoid growth by increasing their host resources/suitability (27). In *H. zea* it has been shown that diets that stress the caterpillar host to the greatest extent have the greatest negative effects on parasitoids (28). Parasitoid
be toxic to an endoparasitoid larva when ingested by its *H. zea* host (32) and to cause disruption of pupation (33), indicating that PDVs may be important and ubiquitous mediators of the interactions of *H. zea* with its host plants. Remarkably, *McBV* is able to manipulate phenotypes across multiple trophic levels including the phenotypes of a parasitoid, herbivore, and host plant. This finding reveals an important dimension of the symbiotic role and ecological benefits of the PDVs to their parasitoid hosts; by indirectly manipulating the host plant, PDVs potentially benefit the parasitoid larva growing inside the caterpillar host and ultimately their own fitness. The importance of PDVs as mediators of interactions within the phytobiome may be of very broad occurrence due to the estimated 50,000 braconid and 14,000 ichneumonid species harboring PDVs, which in turn infect thousands of insect herbivore species (8). In certain contexts, there may be costs associated with PDV manipulation of host plant defenses. The benefits to parasitoid fitness provided by PDVs could be mitigated in circumstances where hyperparasitoids may more easily locate their parasitoid hosts through changes in herbivore-induced plant volatiles (36). Nevertheless, these studies reveal that PDVs play far more prominent roles in shaping plant–herbivore interactions across multiple trophic levels than previously understood. Our findings indicate a unique aspect of how PDVs benefit their parasitoid hosts through suppression of both the caterpillar and the host plant defense or immune systems.

**Materials and Methods**

**Insect Colonies.** *H. zea* eggs were purchased from Frontier Agricultural Sciences, and the colony was maintained in our laboratory for multiple generations. Larvae were fed an artificial diet (37) and reared individually until pupation. Pupa were collected and placed in a plastic container (15 (diameter) cm × 28 (height) cm) through adult emergence, and sugar solution (10%) was provided as food for adults.

The *M. croceipes* colony was obtained from Henry Fadamiro, Auburn University, Auburn, AL, and maintained in our laboratory according to established protocols (38). At the last day of the second instar stage (head capsule slippage stage), *H. zea* larvae were offered to one female parasitoid

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**Fig. 2.** GOX transcript levels and enzyme activity at different time points after parasitism. Values are untransformed mean ± SEM. Different letters indicate significant differences between treatments. Treatments include P- and NP-caterpillars. Relative expression of GOX (A) was measured 1, 2, and 6 d after parasitism: ANOVA, α = 0.05; followed by Student’s t test; day 1, n = 5–8, F(1, 11) = 0.0225, P = 0.882; day 2, n = 11–14, F(1, 23) = 11.29, P = 0.003; day 6, n = 6, F(1, 10) = 0.8836, P = 0.37. GOX activity (B) was measured at 0, 2, 4, and 6 d after parasitism: ANOVA, α = 0.05; followed by Student’s t test; day 0, n = 12–15, F(1, 25) = 0.69, P = 0.42; day 2, n = 15, F(1, 28) = 0.74, P = 0.4; day 4, n = 9–11, F(1, 18) = 4.75, P = 0.04; day 6, n = 15, F(1, 28) = 5.57, P = 0.027.

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**Fig. 3.** Effects of parasitism and polydnavirus on salivary GOX activity of caterpillars. Treatments include saline (S), 0.1 FE and P-treated caterpillars. Values are untransformed mean ± SEM. Different letters indicate significant differences between treatments: ANOVA followed by LSD test, α = 0.05; n = 13–16, F(2, 41) = 6.33, P = 0.004. P, parasitized caterpillars; S, Pringle’s saline-injected caterpillars; 0.1 FE, caterpillars injected with 0.1 FE purified *McBV*.
Caterpillars were removed immediately following a single oviposition by the parasitoid to avoid multiple attacks by the parasitoid. Parasitized caterpillars were fed an artificial diet until parasitoid larval egression. Both insect species were reared in a growth incubator (25 ± 2°C, 16 h light:8 h dark). Parasitoid larvae typically require 10 d to fully develop, egress, and spin the pupal cocoon (on the seventh day of the fourth instar stage H. zea). Adult wasps were sexed using the length of antenna and presence of an ovipositor (38). Following pupal eclosion, each female was maintained with several males in a container [9.5 (diameter) cm × 6 (height) cm] and fed on cotton saturated with 30% honey solution. Parasitized caterpillars in all experiments were used 6 d after parasitism (the third day of the fourth instar stage), and nonparasitized caterpillars were used on the second day of the fourth instar stage (there are 3 d in the fourth instar stage) unless otherwise noted.

Plants. Tomato (Solanum lycopersicum cv. Betterboy) seeds were germinated in potting soil (Sunshine Mix4 Aggregate Plus, Sungrow Horticulture) and grown in a greenhouse (16 h light:8 h dark) at Pennsylvania State University. Seedlings were transferred to pots (10 × 10 × 9 cm) 2 wk after germination and watered daily. Three grams of fertilizer (Osmocote, 15–9–12) was applied on the top of the soil to promote plant growth. Plants with five to six fully expanded leaves (4–5 wk old) were used in all experiments.

Caterpillar Feeding and Plant-Defense Responses. To determine how parasitized caterpillars affect plant-defense responses, plant-defense–related gene expression and protein activities were examined in this study. Plants were separated into three treatment groups: P-caterpillars feeding, NP-caterpillars feeding, and control plants (C) without caterpillars. One P- or NP-caterpillar was placed in a clip cage on the third (from bottom) terminal leaflet of each tomato plant. This method restricts caterpillars to consuming a similar amount of leaf tissues (3.15 cm²) at a particular site during a prescribed time period. In the control treatment, an empty cage was placed on the tomato plant. Leaf cages were removed when the caterpillar consumed the entire leaf tissue inside the cage within 10 h. Twenty-four hours after placing the caterpillars on the plants, 100 mg of the third terminal leaflet was collected for RNA extraction. Leaf samples were ground (Geno/Grinder 2000, Spex Sample Prep) with liquid nitrogen. RNA was extracted with TRizol reagent (Ambion). One microgram of RNA was used to synthesize cDNA with the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems) in a PCR thermal cycler (GeneAmp PCR System 9700) (7). Gene primers of defense genes used for quantitative real-time PCR (qRT-PCR) analysis were designed by Primer Quest Software (Applied Biosystems) and listed (SI Appendix, Table S2). qRT-PCR analysis was performed by the 7500 Fast Real-Time PCR System (Applied Biosystems) with the FastStart Universal SYBR Green PCR Probe Master (Roche). Actin (ACT) and ubiquitin (UB) genes were used as reference genes in tomato plants. Target genes’ relative expressions were compared with intact control (C) by using the 2−ΔΔct method (39).
Fourty-eight hours after treatments, 50 mg of the third terminal leaflet was collected for PPO and Ti activity assays. Leaf samples were frozen with liquid nitrogen and ground with the Genogrinder. Immediately, 1.25 mL of phosphate buffer (0.1 M, pH 7) with 5% polyvinylpyrrolidone (PVP) (Alfa Aesar 41631) was added, and samples were vortexed and incubated on ice for 5 min. Supernatant was collected after centrifugation (4 °C, 11,000 × g, 10 min). To measure PPO activity, 5 μL of each sample was added to 200 μL caffeic acid (3 mM; Sigma C6026), and the change in absorbance at 450 nm was recorded in the Spectrmax 190 plate reader (Molecular Devices) (22).

For trypsin inhibition analysis, samples were powdered by the Genogrinder, and then 1.25 mL of extraction buffer (0.046 M Tris and 0.0115 M CaCl2, pH 8.1) with 5% PVP was added. The supernatant (4 °C, 11,000 × g, 10 min) was collected for Ti activity measurement. Ten microliters of sample was mixed with 10 μL of trypsin (20 μg/mL; Sigma T1426) and 80 μL of extraction buffer. Ten minutes later, 100 μL of p-toluene-sulfonyl-arginine methyl ester (0.002 M; Sigma T4626) was added, and absorbance values were read at 247 nm (7). Percentage inhibition was calculated by comparing the activity of trypsin and extraction buffer alone.

Caterpillar saliva and Plant-Defense Responses. To determine if saliva is responsible for the differences that we observed in plant-defense responses from feeding caterpillars, caterpillars were parasitized as described above. Labial salivary glands were collected from P- and NP-caterpillars. Glands were homogenized in phosphate buffer (0.1 M, pH 7), and the homogenate was collected after centrifugation (4 °C, 7,500 × g, 10 min). Protein in the supernatant was quantified by Bradford assay (40), and all samples were diluted to 1 μg/μL. The third terminal leaflets of tomato plants were treated with mechanical wounding and application of 15 μL of salivary gland supernatant from parasitized and nonparasitized caterpillars, respectively. The third group was the intact control plant without any treatment. For gene expression experiments, samples of the wounded leaf were collected 24 h after treatment. PPO was analyzed 48 h after treatment.

Caterpillar Salivary Glucose Oxidase Transcript Levels and Enzyme Activities. To determine how parasitism affects GOX gene expression and enzyme activity, time-course experiments were conducted for P- and NP-caterpillars. Due to the very small size of younger caterpillars and feasibility of dissections, third instar stage H. zea caterpillars were used for these two experiments.

For GOX gene expression, on the last day of the third instar stage, H. zea larvae were parasitized by M. croceipes. Salivary glands were collected from P-caterpillars 1, 2, and 6 d after parasitism. Because NP-caterpillars develop faster, NP-caterpillar salivary glands were collected at the same developmental stage as the P-caterpillar treatment. Caterpillar salivary gland RNA extraction and cDNA synthesis were as described above. H. zea glucose oxidase (GOX) gene expression was tested by qRT-PCR analysis, and actin (ACT) was used as the reference gene. Gene relative expression was calculated as described above.

For GOX enzyme activity, on the last day of the third instar stage, H. zea larvae were parasitized by M. croceipes. P-caterpillar salivary glands were collected at 0, 2, 4, and 6 d after parasitism. NP-caterpillar salivary glands were collected at the same developmental stage as caterpillars. GOX enzyme activity was analyzed as described (41).

GOX Dosage Effect on Plant-Induced Defense Response. To determine the effect of GOX dosage on plant-defense response, tomato plants were treated with varying concentrations of GOX from Aspergillus niger (Sigma-Aldrich) based upon a previous estimate of levels of GOX secretion by H. zea (37). The youngest fully expanded terminal leaf was mechanically wounded, and immediately 20 μL of GOX was diluted in 0.1 M phosphate buffer (pH 7.2). After 48 h, 50-μg samples of tissue were collected from the wounded leaf, frozen in liquid nitrogen, and then stored at −80 °C until further analysis. Leaf tissue was analyzed for trypsin inhibitor activity as previously described.

Caterpillar Host Performance. To evaluate the effect of plant-defense responses on P-caterpillar performance, we performed a larval relative growth rate experiment. There were three groups in the experiment: caterpillars feeding on plants treated with saliva from P-caterpillars, NP-caterpillars, and intact control plants (C). Plants were treated with labial salivary gland homogenate as described above. Forty-eight hours after treatment, the treated leaf was collected for bioassay. Third instar stage H. zea of similar body size were selected and parasitized by M. croceipes. Caterpillars were weighed and placed on the treated tomato leaves in plastic cups lined with 2% agar to keep leaves moist. Twenty-four hours later, caterpillars were reweighed and relative growth rate was calculated as follows: (final weight − initial weight)/average weight × no. of days).

Parasitoid Performance. To determine if different levels of induced plant-defense responses caused by parasitized and nonparasitized caterpillars in- fluence parasitoid development, we conducted a parasitoid performance experiment. Plants were treated with labial salivary gland homogenate as described above. Forty-eight hours after treatment, the treated leaf was collected and placed in a plastic cup lined with 2% agar to keep leaves moist. Third instar stage H. zea larvae of similar body size were parasitized by M. croceipes and fed in one of two treatments: P-caterpillar–treated plants or NP-caterpillar–treated plants (total n = 54; six replicates of nine individuals per treatment). Leaves were changed every other day to keep food fresh until parasitoid larvae spin the pupa cocoon. Larval duration, cocoon weight, pupal duration, larval mortality, cocoon formation failure rate, adult emergence rate, and survival rate were recorded. Cocoon weight was measured 2 d after cocoon formation, and adult emergence rate was calculated 30 d after cocoon formation. For the percentage of larval mortality, percentage of cocoon formation failure rate, percentage of adult emergence, and percentage of total survival, data were calculated from six replicates with n = 9 individuals per treatment.

Experiments with M. croceipes Polydnavirus (McBV). McBV was purified from the calyx region of the ovaries of M. croceipes (SI Appendix, Fig. S5) following an established protocol (42) with slight modifications. Briefly, female M. croceipes were chilled on ice for 15 min and rinsed in autoclaved water before dissection. Ovaries were collected from 30 females into a 1.7-ml microtube and then homogenized with 100 μL of Tris buffer (0.5 M, pH 6.8). Crude calyx extract was collected after centrifugation (4 °C, 800 × g, 15 min). One hundred microliters of Tris buffer was added to the pellet for a second extraction. The two extracts were pooled and centrifuged (4 °C, 12,000 × g, 5 min). The supernatant was applied to a sucrose gradient (40–70% sucrose) and centrifuged in a swinging bucket rotor (4 °C, 24,000 × g, 30 min). After centrifugation, the PDVs were visible as a blue band in the sucrose gradient. The band was collected with a micropipette, and PDVs were pelleted by centrifugation (4 °C, 49,000 × g, 45 min). The resulting pellet was resuspended in Pringle’s saline (pH 7.39) and adjusted to a suitable concentration for injection. Virus collected from each female was defined as one female equivalent (FE). For injection, 5 μL of Pringle’s saline (5) or 5 μL of 0.02 FE of PDV (0.1 FE) was injected into last-day third instar stage H. zea caterpillars using a microinjector (Burkard PAX 100) with a 32-gauge hypodermic needle. All processes, including PDV collection, purification, and injection were finished within 4.5 h. The same stage of H. zea caterpillars was offered to M. croceipes females for parasitism (P) treatment. Two days after injection or parasitism, labial glands were collected from a subset of each treatment. To confirm virus expression in the labial glands of injected caterpillars, the PDV gene ORPHMS was amplified after RNA extraction and cDNA synthesis (as described above). The primers used to amplify the putative region of the McBV segment are shown in SI Appendix, Table S2. All caterpillars (n = 6) that were injected with McBV showed viral gene expression in the labial glands (SI Appendix, Fig. S7). Six days after parasitism (third day of the fifth instar stage and second day of the fifth instar stage of S- or McBV-treated H. zea caterpillars) were caged on plants for measuring plant-defense gene expression and PDV and PDV host interactions as described above. To test the effect of virus on salivary enzyme activity, labial glands were collected from injected caterpillars 6 d post injection, and GOX activity was assayed as described above. To determine if McBV is the factor reducing H. zea salivary gland GOX gene expression, a GOX relative expression experiment was conducted. McBV were collected and purified as described above. There were three groups treated: S-injected, 0.1 FE McBV-injected, and parasitized caterpillars. Caterpillar salivary glands were collected 2 d after treatment. Salivary gland RNA extraction, cDNA synthesis, and qRT-PCR analysis were as described above.

Confocal Laser-Scanning Microscopy of Female M. croceipes Reproductive Organs. Female adult wasps were chilled on ice and dissected in 0.1 M monobasic phosphate buffer. After removing the female ovipositor, the attached reproductive organs were fixed (2.5% glutaraldehyde, 0.1 M phosphate buffer, and 0.05 g/mL sucrose) in room temperature. The tissues were placed on microscope slides with glycerol and then examined with an Olympus FV10i Confocal Laser Scanning Microscope (CLSM) using the following excitation wavelengths: 405, 473, and 559 nm with emission wavelengths 430, 520, and 600 nm, respectively. Images were processed using FIJI–ImageJ.

Statistical Analyses. Data were transformed as needed to obtain a normal distribution and to address residuals with heterogeneity of variance; SAS 9.4 (SAS Institute) was used for all analyses. Plant-defense responses (gene
expression and TI and PPO activities) and caterpillar performance bioassays were analyzed using one-way ANOVA (Proc GLM), followed by means comparisons using Tukey's Least Significant Difference (LSD) test (significance level, P < 0.05). Insect salivary gland (GOX) gene expression and GOX enzyme activity (time course) were compared using Student's t test. GOX concentration on TI activity was analyzed by one-way ANOVA (Proc GLM), followed by means comparisons using Tukey's LSD test (significance level, P < 0.05). Parasitoid performance (larval duration, cocoon weight, pupal duration, and percentage of larval mortality, cocoon formation adult emergence, and total survival) were compared using Student's t test. MdBV injection experiments (plant-defense responses, insect GOX gene expression, and enzyme activity) were analyzed using one-way ANOVA (Proc GLM), followed by means comparisons using Tukey's LSD test (significance level, P < 0.05).

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