

Variation in the Susceptibility of *Drosophila* to Different Entomopathogenic Nematodes

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Entomopathogenic nematodes (EPNs) in the genera *Heterorhabditis* and *Steinernema* are lethal parasites of insects that are of interest as models for understanding parasite-host interactions and as biocontrol agents for insect pests. EPNs harbor a bacterial endosymbiont in their gut that assists in insect killing. EPNs are capable of infecting and killing a wide range of insects, yet how the nematodes and their bacterial endosymbionts interact with the insect immune system is poorly understood. Here, we develop a versatile model system for understanding the insect immune response to parasitic nematode infection that consists of seven species of EPNs as model parasites and five species of *Drosophila* fruit flies as model hosts. We show that the EPN *Steinernema carpocapsae*, which is widely used for insect control, is capable of infecting and killing *D. melanogaster* larvae. *S. carpocapsae* is associated with the bacterium *Xenorhabdus nematophila*, and we show that *X. nematophila* induces expression of a subset of antimicrobial peptide genes and suppresses the melanization response to the nematode. We further show that EPNs vary in their virulence toward *D. melanogaster* and that *Drosophila* species vary in their susceptibilities to EPN infection. Differences in virulence among different EPN-host combinations result from differences in both rates of infection and rates of postinfection survival. Our results establish a powerful model system for understanding mechanisms of host-parasite interactions and the insect immune response to parasitic nematode infection.

Entomopathogenic nematodes (EPNs) of the genera *Steinernema* and *Heterorhabditis* are insect-parasitic nematodes that are phylogenetically distant but share a similar life cycle as a result of convergent evolution (1). EPNs offer numerous advantages as model parasitic nematodes, including small size, short generation time, and amenability to *in vitro* culturing (2). EPN infective larvae are associated with bacterial endosymbionts: *Steinernema* species are associated with bacteria in the genus *Xenorhabdus*, and *Heterorhabditis* species are associated with bacteria in the genus *Photorhabdus* (1). At least some EPNs are capable of infecting the fruit fly *Drosophila melanogaster*, providing a genetically tractable system for understanding the immune response to parasitic nematodes and their bacterial endosymbionts (3–6). However, the insect immune response to EPN infection is poorly understood.

During a particular developmental stage called the infective juvenile (IJ), EPNs infect insects (Fig. 1A). IJs are developmentally arrested, third-stage larvae analogous to the dauer stage of free-living nematodes (7). IJs actively seek out insect hosts using chemosensory cues (8–10) and infect either by entering through natural body openings or by penetrating the insect cuticle (11). IJs harbor their bacterial endosymbiont in their gut and deposit it into the insect upon infection, where it assists the nematode in killing the insect, digesting insect tissues, and inhibiting the growth of other microorganisms (12–14). Following infection, the nematodes reproduce in the insect cadaver and feed on the bacterium-infested tissue until resources are depleted, at which point new IJs form and emerge from the cadaver to search for new hosts (Fig. 1B) (15).

In response to EPN infection, insects mount an innate immune response that involves antimicrobial peptide (AMP) expression as well as activation of the melanization and encapsulation reactions (11). At the same time, the nematodes attempt to evade or suppress the insect immune response through a process that remains poorly understood. Both the nematode and its bacterial endosymbiont appear to inhibit some aspects of AMP production,

melanization, encapsulation, and phagocytosis (11, 16–18). Studies using *D. melanogaster* larvae as a model host for the EPN *Heterorhabditis bacteriophora* and its endosymbiont *Photorhabdus luminescens* have shown that infection induces expression of a large number of immune genes, including AMPs, and that AMP expression is primarily a response to the bacterial endosymbiont rather than to the nematode (4, 19). Infection also stimulates clotting, and clotting mutants show decreased survival in response to *H. bacteriophora*-*P. luminescens* infection (20, 21). However, studies of the immune response of *D. melanogaster* to EPN infection have so far been limited to *H. bacteriophora*-*P. luminescens*, and the extent to which the immune response differs for different EPNs is unclear.

Here, we demonstrate that the distantly related EPN *Steinernema carpocapsae* and its bacterial endosymbiont *Xenorhabdus nematophila* are capable of infecting *D. melanogaster* larvae and are more virulent toward *D. melanogaster* than *H. bacteriophora*-*P. luminescens*. Infection with *S. carpocapsae* symbiont IJs (i.e., IJs harboring *X. nematophila* in their gut) induced expression of a subset of AMP genes. *S. carpocapsae* infection also activated the

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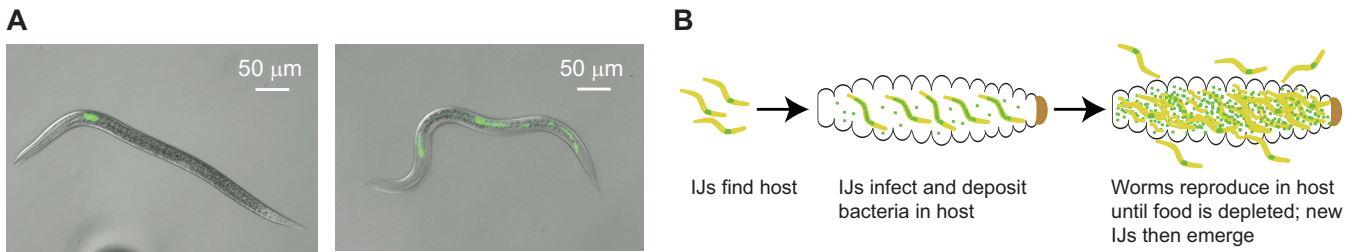


FIG 1 Life cycle of *S. carpocapsae*. (A) Photomicrographs of *S. carpocapsae* infective juveniles (IJs) with GFP-expressing *X. nematophila*. The left frame shows an IJ outside the host, and the right frame shows an IJ that was extracted from a host and that is defecating *X. nematophila*. (B) The life cycle of *S. carpocapsae*. IJs find a host, enter through a natural body opening, and defecate their symbiotic bacteria into the host. The bacteria play an important role in overcoming the host immune system (1). The nematodes develop and reproduce in the insect cadaver until resources are depleted. New IJs then form and exit the cadaver. Green dots represent the bacterial endosymbiont.

melanization pathway, and a higher rate of melanization occurred following infection with axenic IJs than with symbiont IJs, suggesting that *X. nematophila* suppresses the melanization response. Finally, exposure of *D. melanogaster* larvae to seven different EPN species revealed that EPNs vary in their virulence levels toward *D. melanogaster*, and exposure of five different *Drosophila* species to *S. carpocapsae* symbiont IJs revealed that *Drosophila* species vary in their susceptibilities to EPN infection. Our results establish the EPN-*Drosophila* system as a powerful model for investigating the insect immune response to nematode infection.

MATERIALS AND METHODS

Nematode strains. The following EPN strains were used: *S. carpocapsae* ALL (8, 9), *H. bacteriophora* Baine (22), *Steinernema glaseri* NC (23), *Steinernema scapterisci* FL (24), *Steinernema riobrave* TX (25), *Heterorhabditis indica* HOM1 (22), and *Steinernema feltiae* SN (26).

***Drosophila* stocks.** Wild-type *D. melanogaster* larvae were from the Canton-S strain. Studies of AMP expression were conducted with strains of *D. melanogaster* containing either an *attacinA::GFP*, *cecropinA1::GFP*, *metchnikowin::GFP*, *drosocin::GFP*, *drosomycin::GFP*, *dipteracin::GFP*, or *defensin::GFP* transgene (where GFP is green fluorescent protein) (27). Wild-type *Drosophila virilis*, *Drosophila simulans*, *Drosophila yakuba*, and *Drosophila pseudoobscura* were stocks 15010-1051.00, 14021-0251.006, 14021-0261.00, and 14011-0121.104 from the *Drosophila* Species Stock Center, respectively. We note that all *D. melanogaster* strains, as well as the wild-type *Drosophila simulans* strain, were confirmed to be infected with *Wolbachia* by PCR using previously described primers (28). However, *Wolbachia* status is unlikely to affect susceptibility to EPNs or their bacterial endosymbionts: although *Wolbachia* infection may protect against some viral infections (29, 30), it does not appear to protect against other types of infections and has little or no effect on AMP expression (29, 31–33).

Bacterial strains. The following bacterial strains were used: wild-type *X. nematophila* HGB800 (34), GFP-expressing *X. nematophila* HGB340 (13), colonization-defective *X. nematophila* HGB777 (35), *Escherichia coli* OP50-GFP, *P. luminescens* TT01-GFP (36), *Photobacterium temperata* NC1-GFP (36), and colonization-defective *P. temperata* NC1-GFP TRN16 (36). *Xenorhabdus* was grown in LB broth containing 0.1% sodium pyruvate, and *Photobacterium* was grown in PP3 broth as previously described (8).

Nematode culturing. To generate symbiont IJs, nematodes were cultured in either the waxworm *Galleria mellonella* (for all species except *S. scapterisci*) or the house cricket *Acheta domestica* (for *S. scapterisci*) as previously described (8, 9). Briefly, five last-instar waxworms or one medium-sized cricket (American Cricket Ranch, Lakeside, CA) was placed in a 5-cm petri dish with a 55-mm Whatman 1 filter paper in the bottom of the dish. Approximately 500 to 1,000 IJs suspended in water were distributed on the filter paper and on the insects. Petri dishes were stored either

at 25°C in the case of *H. bacteriophora* and *S. riobrave* or at room temperature (22 to 23°C) in the case of all other species. *H. bacteriophora* and *S. riobrave* infections were performed at 25°C because these species are found primarily in warm climates (37) and infect insects more efficiently at 25°C than at room temperature. Infections for the other species were performed at room temperature because these species infect more efficiently at room temperature than at 25°C. After ~10 days the insect cadavers were placed on White traps (38) or, in the case of *S. glaseri*, on modified White traps containing plaster of Paris (9). Symbiont IJs were collected from traps within 10 days, stored at 15°C, and tested within 1 month of collection.

To generate axenic *S. carpocapsae* IJs, symbiont *S. carpocapsae* IJs were surface sterilized by incubation in 1% commercial bleach for 5 min. IJs were then rinsed three times in distilled H₂O (dH₂O), incubated in antibiotic solution (10 µg/ml gentamicin, 100 µg/ml streptomycin, 100 µg/ml carbenicillin, and 20 µg/ml kanamycin in dH₂O) for 48 h, and plated onto 1× lipid agar-cholesterol plates (39) (final concentration of cholesterol, 5 mg/liter) containing 0.1% sodium pyruvate and seeded with *X. nematophila* HGB777 bacteria (35). Axenic nematodes were maintained on lipid agar-cholesterol plates seeded with HGB777, and IJs were collected from plates as previously described (8). IJs were incubated in 1% commercial bleach for 5 min and then rinsed three times in dH₂O to surface sterilize them prior to testing. To verify that the IJs were axenic, 5 µl of IJ pellet was plated onto a lipid agar-cholesterol plate and incubated at 25°C. The absence of bacteria on the plate was confirmed after 2 to 3 days.

To generate symbiont *S. carpocapsae* IJs containing GFP-expressing *X. nematophila*, axenic IJs were plated onto lipid agar-cholesterol plates seeded with *X. nematophila* HGB340 bacteria. Nematodes were maintained on lipid agar-cholesterol plates seeded with HGB340, and IJs were collected from plates as previously described (8).

To generate symbiont *H. bacteriophora* IJs containing GFP-expressing *P. luminescens*, symbiont *H. bacteriophora* IJs that had emerged from waxworms were plated onto 1× lipid agar-cholesterol plates (39) containing 0.1% sodium pyruvate seeded with *P. temperata* NC1 TRN16 bacteria (36). IJs were collected from plates as previously described (8) and plated onto 1× lipid agar-cholesterol plates seeded with *P. luminescens* TT01-GFP bacteria (36). Nematodes were maintained on 1× lipid agar-cholesterol plates seeded with TT01-GFP, and IJs were collected from plates as previously described (8). Axenic *H. bacteriophora* IJs were generated as described above for axenic *S. carpocapsae* IJs, except that they were plated onto and maintained on TRN16.

Infection of *Drosophila* larvae with EPNs. IJs used to assay survival were grown in waxworms; IJs used to assay infection were grown on GFP-expressing bacteria. IJs were rinsed three times in dH₂O, and 10 µl of water containing 500 IJs was pipetted onto the center of a 5-cm petri dish containing nematode growth medium (NGM). For each trial, 20 third-instar *Drosophila* larvae were rinsed twice in 1× phosphate-buffered saline (PBS) and placed onto the NGM plate containing IJs. *Drosophila*

larvae infected with *H. bacteriophora* and *S. riobrave* were kept in a 25°C incubator; larvae infected with all other strains were kept at room temperature. Different temperatures were used for *H. bacteriophora* and *S. riobrave* because these species are adapted for infection and growth at warmer temperatures than the other species, as described above. Metal rings were placed onto the plate lids as weights to prevent fly larvae from escaping. Infection and survival were scored at 24 and 48 h postexposure to IJs. Melanization was scored at 48 h postexposure to IJs to ensure that a majority of the population had been infected.

To score infection, fly larvae or pupae (in cases where the fly larvae pupated during the course of the experiment) were assayed under an epifluorescence dissecting microscope. IJs grown on GFP-expressing symbiotic bacteria were used to facilitate detection of worms inside the fly host. Fly larvae or pupae were considered infected if worms were visible inside the body. Although worms could be seen inside the host even without the presence of GFP-expressing symbiotic bacteria, worms could be identified more efficiently when they had GFP-expressing symbiotic bacteria. To score survival, fly larvae or pupae were assayed under a dissecting microscope at $\times 50$ magnification. Animals were determined to be alive if they had a visible heartbeat or responded to gentle prodding.

To score melanization, fly larvae or pupae were first scored under a dissecting microscope at $\times 50$ magnification for black spots on the cuticle and then dissected to determine whether they were infected. For each trial, the percentage of infected fly larvae or pupae with visible melanization was quantified. A value of 100% would indicate that all of the infected *D. melanogaster* larvae showed visible melanization; a value of 0% would indicate that none of the infected *D. melanogaster* larvae showed visible melanization.

Infection of *Drosophila* larvae with bacteria. GFP-expressing *X. nematophila*, *E. coli*, or *P. luminescens* cells were used for infection assays; wild-type bacteria were used for all other assays. To generate each assay plate, 100 μ l of a bacterial suspension (for *X. nematophila* and *E. coli*) or 200 μ l of bacterial suspension (for *P. luminescens*) from a 1- or 2-day culture was spread onto a 5-cm plate containing LB supplemented with 100 μ g/ml carbenicillin and 0.1% sodium pyruvate (*X. nematophila*), LB alone (*E. coli*), or 1 \times lipid agar with cholesterol plus 0.1% sodium pyruvate (*P. luminescens*). Plates were incubated at 25°C for 1 to 2 days (*X. nematophila* and *P. luminescens*) or at 37°C overnight (*E. coli*) to create a bacterial lawn. For each trial, 20 second-instar or early-third-instar *D. melanogaster* larvae were rinsed in 1 \times PBS and placed onto a plate containing a bacterial lawn. A second plate containing a bacterial lawn was then secured upside down on top of the first plate to prevent the fly larvae from avoiding the bacteria by crawling onto the plate lid. Survival was scored at 24, 48, and 72 h as described above. To assay infection, fly larvae were washed twice in 1 \times PBS, placed onto unseeded NGM plates, and scored for GFP expression under an epifluorescence dissecting microscope after 24 and 48 h. All flies with visible GFP expression inside the body were scored as GFP positive. The percentages of GFP-positive fly larvae were then calculated. Fly larvae that were GFP negative at 24 h were placed onto new plates seeded with bacteria and scored again at 48 h. No GFP expression was observed in control experiments where fly larvae were not exposed to bacteria.

AMP expression assay. AMP expression was assayed in *D. melanogaster* larvae following infection with symbiont IJs, axenic IJs, or bacteria. For these experiments, both symbiont IJs and axenic IJs were grown *in vitro* on plates containing lawns of *X. nematophila* to eliminate any potential differences in *D. melanogaster* AMP expression resulting from differences in nematode culturing conditions. We used seven different transgenic *D. melanogaster* strains as hosts, each of which expressed a reporter construct in which GFP expression was driven by the promoter of a different AMP gene (27). Details of the transgenic *D. melanogaster* larvae expressing the AMP reporter constructs are described above under “*Drosophila* stocks.” We note that no GFP-expressing *X. nematophila* was present in this experiment; symbiont IJs contained bacteria that did not express GFP, and the bacteria used for bacterial infections also did not

express GFP. Thus, the GFP expression observed in these experiments was from the transgenic *D. melanogaster* hosts and was a reflection of AMP gene expression.

Infection of transgenic *D. melanogaster* larvae was performed as described above; each trial consisted of 20 fly larvae. For the uninfected controls, fly larvae were placed onto either NGM plates without IJs (controls for infection with symbiont or axenic IJs) or LB plates without bacteria (control for infection with *X. nematophila*). AMP expression was scored at 24 h postexposure to IJs. We note that in initial experiments, AMP expression was scored at 8, 24, and 48 h postexposure to IJs. AMP expression levels at 8 h postexposure were low because most fly larvae had not yet been infected, and no difference was observed between AMP expression levels at 24 and 48 h postexposure (data not shown). We therefore focused on the 24-h time point for further experiments.

To score AMP expression, larvae were removed from the plates, rinsed twice in 1 \times PBS, placed onto unseeded NGM plates, and observed using the GFP filter of an epifluorescence dissecting microscope. For the *attacin*, *cecropin*, *defensin*, *drosocin*, and *metchnikowin* genes, fly larvae were scored as GFP positive if any GFP expression was observed. For the *diptericin* and *drosomyacin* genes, fly larvae were scored as GFP positive if diffuse GFP expression was observed because small spots of GFP expression were often observed under normal culturing conditions. The percentage of fly larvae expressing GFP was then calculated. We note that we previously validated our visual GFP scoring method for these reporter lines by comparing expression data obtained by visual scoring versus quantification in ImageJ and determining that expression data obtained by both methods were consistent (4). To control for GFP expression not due to infection with IJs or bacteria, the percentage of GFP-positive fly larvae obtained from the uninfected control experiments was subtracted from the percentage of GFP-positive fly larvae obtained from the infection experiments. Thus, Fig. 3 reports the background-subtracted values for the percentage of fly larvae that express the indicated AMP reporter construct.

For the bacterial infection experiment, plates were observed at 24 and 48 h postexposure to *X. nematophila*. On plates where some or all of the fly larvae had burrowed into the agar by the 24-h time point, all larvae on that plate were transferred to a new *X. nematophila* plate to ensure that the fly larvae remained in contact with the bacteria for the duration of the experiment.

Examining the time course for infection and survival following EPN exposure. For each trial, ~ 20 fly larvae were exposed to symbiont IJs containing GFP-expressing *X. nematophila* or *P. temperata*. IJs were used within 1 week of collection. Infection was scored at 2, 4, 6, 8, 24, and 48 h postexposure to IJs and was visualized using the GFP filter on an epifluorescence microscope. At each time point, larvae were rinsed twice in 1 \times PBS and placed onto unseeded NGM plates prior to scoring. After scoring, infected larvae were placed onto unseeded NGM plates that did not contain IJs, while uninfected larvae were placed back onto the original plate containing IJs so that they could be scored for infection at later time points. To examine postinfection survival, all larvae infected by 8 h were scored for survival at 24 and 48 h. Trials in which fewer than three fly larvae became infected by 8 h were not included in the analysis. To assay long-term survival, animals were scored for infection and survival as described above, except that any animals still alive at 48 h were placed onto new unseeded NGM plates and monitored for survival to adulthood.

Statistical analysis. Statistical analysis was performed using GraphPad Instat or Prism software. Standard statistical tests were used for all experiments, as described in the figure legends. All statistical comparisons are described in the relevant figure legends and supplemental tables. The value for sample size (*n*) used for statistical analysis refers to the number of trials performed for each treatment, condition, or genotype; each trial consisted of ~ 20 fly larvae.

RESULTS AND DISCUSSION

***S. carpocapsae* infects and kills *D. melanogaster*.** *S. carpocapsae* has a wide geographical distribution and a broad host range and is

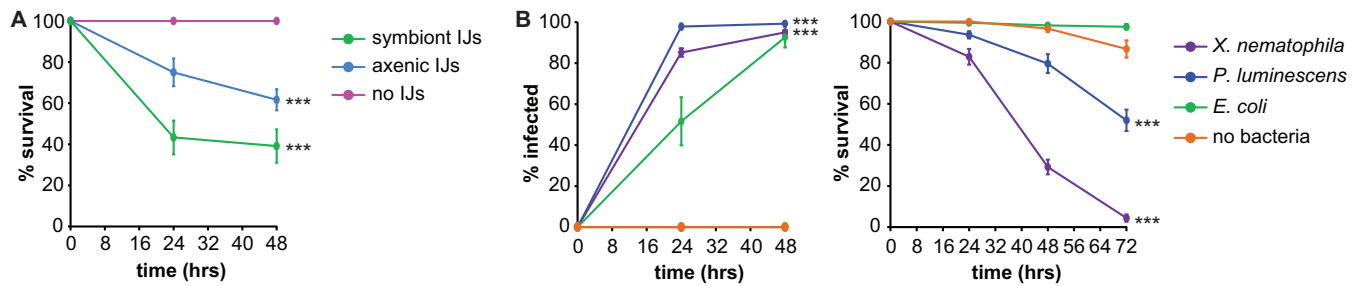


FIG 2 *S. carpocapsae* and its bacterial endosymbiont *X. nematophila* are pathogenic toward *D. melanogaster*. (A) Survival of *D. melanogaster* larvae exposed to symbiont *S. carpocapsae* IJs, axenic *S. carpocapsae* IJs, or no IJs. All three survival curves are significantly different ($P < 0.0001$, log rank test). ***, $P < 0.001$ relative to the no-IJ control (log rank test with Bonferroni correction; $n = 6$ trials for each condition). (B) Infection (left) and survival (right) of *D. melanogaster* larvae exposed to either *X. nematophila*, *P. luminescens*, *E. coli*, or no bacteria. All three species of bacteria successfully infected *D. melanogaster* larvae although infection rates were significantly different for each species ($P < 0.01$, log rank test). ***, $P < 0.001$ for *X. nematophila* and *P. luminescens* relative to *E. coli* (log rank test with Bonferroni correction). The survival curve for fly larvae exposed to *E. coli* was not significantly different from the survival curve for the no-bacteria control (right graph); all other survival curves were significantly different from each other ($P < 0.001$, log rank test with Bonferroni correction). ***, $P < 0.001$ relative to *E. coli* and the no-bacteria control (log rank test with Bonferroni correction). The no-bacteria control shown in the graph was performed on LB plates; a no-bacteria control was also performed on lipid-agar plates (for comparison to *P. luminescens*), and results were not significantly different from those of the control on LB plates ($P = 0.2728$, log rank test; $n = 5$ to 9 trials for each condition). For all graphs, the x axis refers to time postexposure, and error bars represent standard errors of the means. In some cases, error bars are too small to be visible.

used as a biocontrol agent for numerous insect pests (40). To determine whether *S. carpocapsae* is pathogenic for *D. melanogaster*, we used an infection assay in which we exposed third-instar fly larvae to symbiont IJs. Third-instar larvae were used for this assay because EPNs typically infect late-stage insect larvae (41). We scored infection and survival at 24 and 48 h postexposure since EPNs generally kill hosts within 48 h (41). We found that approximately 60% of the fly larvae exposed to symbiont IJs died within 24 h (Fig. 2A). Thus, *S. carpocapsae* is capable of infecting and killing *D. melanogaster* larvae. To determine whether pathogenicity was conferred primarily by *S. carpocapsae* or *X. nematophila*, we exposed fly larvae to axenic IJs. We found that axenic IJs were also capable of infecting and killing *D. melanogaster* larvae, although with less efficiency than symbiont IJs (Fig. 2A). Thus, *S. carpocapsae* IJs are pathogenic for *D. melanogaster* even in the absence of *X. nematophila*, consistent with previous studies of *S. carpocapsae* infection in larger insects, such as *Galleria mellonella* (42).

We then examined the pathogenicity of *X. nematophila* in the absence of its nematode vector by exposing fly larvae to agar plates containing lawns of *X. nematophila*. For comparison, we also exposed fly larvae to lawns of *P. luminescens* and *E. coli*. We found that exposure to all three bacteria resulted in infection of fly larvae, as determined by counting the number of GFP-positive fly larvae postexposure to GFP-labeled bacteria (Fig. 2B, left graph). *X. nematophila* was pathogenic for *D. melanogaster* larvae: 95% of fly larvae exposed to *X. nematophila* were dead by 72 h (Fig. 2B, right graph). *X. nematophila* was significantly more virulent than *P. luminescens*, which killed only approximately 50% of fly larvae by 72 h (Fig. 2B). In contrast, *E. coli* was not pathogenic for *D. melanogaster* larvae. These results are consistent with a previous study which found that *X. nematophila* is more virulent than *P. luminescens* when injected into *D. melanogaster* adults (43). Taken together, these results suggest that both *P. luminescens* and *X. nematophila* are pathogenic for *D. melanogaster* larvae but differ in their virulence levels.

Infection of fly larvae with *X. nematophila* most likely occurred by ingestion since GFP-expressing bacteria appeared to localize initially to the digestive tract (see Fig. S1 in the supplemental ma-

terial). This is consistent with our previous observations of exposure of fly larvae to *Photorhabdus* bacteria (4). Susceptibility to *X. nematophila* is not likely to be the result of exposure to external toxins, since nonfeeding third-instar larvae did not become infected with *X. nematophila* in our assay. However, we cannot exclude the possibility that *X. nematophila* secretes toxins that have external effects on second- and early-third-instar larvae but not older third-instar larvae.

Infection induces expression of antimicrobial peptide (AMP) genes. A major component of the insect innate immune response is AMP production by the fat body, a structure similar to the mammalian liver and adipose tissue (44). Studies of a number of insects, including the cecropia moth *Hyalophora cecropia*, the beet armyworm *Spodoptera exigua*, and the tobacco hornworm *Manduca sexta* have shown that EPN infection can induce expression of AMP genes and that both the nematode and the bacteria can suppress AMP activity (45–49). We previously showed that infection of *D. melanogaster* larvae with *H. bacteriophora* symbiont IJs resulted in expression of four AMP genes, *attacin*, *dipteracin*, *drosomycin*, and *metchnikowin*, and that this expression was a specific response to *P. luminescens* (4). Similar results were subsequently observed for infection of *M. sexta* with *H. bacteriophora* symbiont IJs, thus validating *D. melanogaster* as a model for other insect hosts (45). The AMP genes *dipteracin* and *drosomycin* have also been shown to be upregulated following direct injection of either *P. luminescens* or *X. nematophila* into *D. melanogaster* adults (43). However, the AMP response of *D. melanogaster* to infection with *S. carpocapsae* symbiont IJs had not yet been examined, and the extent to which AMP expression is induced by EPNs versus their bacterial endosymbionts remains unclear (4, 6, 45).

To determine whether infection of *D. melanogaster* larvae with *S. carpocapsae*-*X. nematophila* induces AMP expression, we exposed both second-instar and third-instar fly larvae to symbiont IJs and monitored AMP expression at 24 h postexposure. To monitor AMP expression, we used seven different transgenic fly lines, each of which contained a reporter construct that expressed GFP under the control of a different AMP gene promoter. For each transgenic line, AMP expression was determined by scoring the fly larvae or pupae (in cases where the fly larvae pupated during the

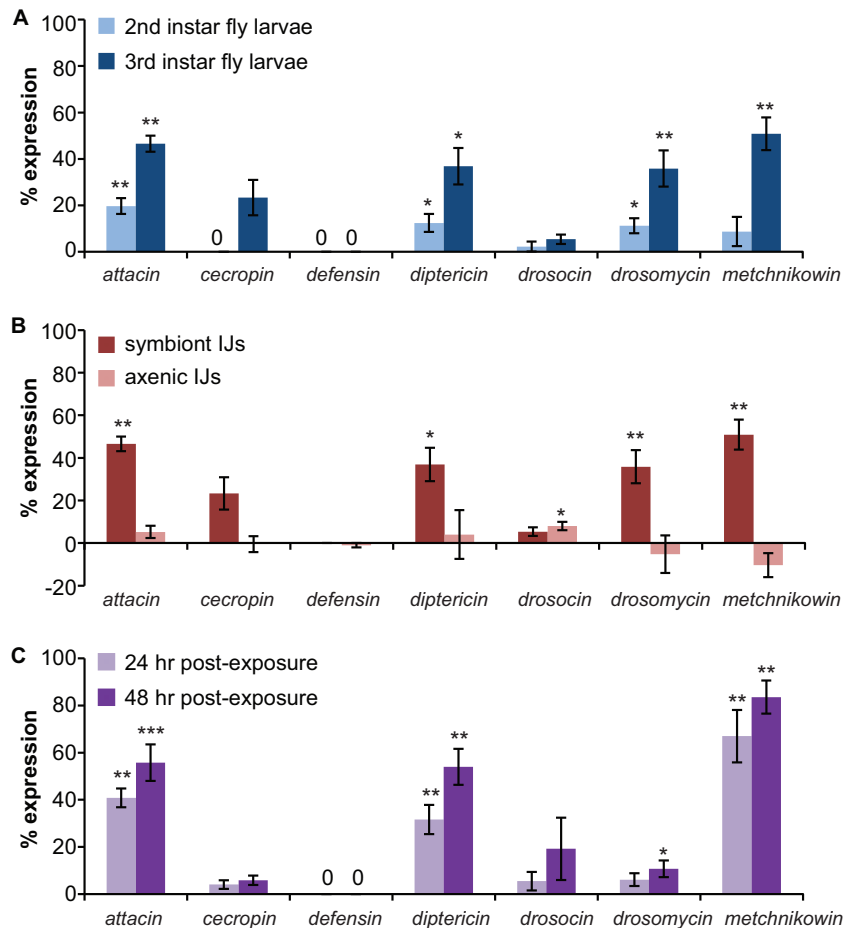


FIG 3 Infection of *D. melanogaster* larvae with symbiotic *S. carpocapsae* IJs or *X. nematophila* induces a humoral immune response. (A) AMP expression following infection with symbiotic *S. carpocapsae* IJs. Infection of second- or third-instar *D. melanogaster* larvae with symbiotic *S. carpocapsae* IJs induces expression of a subset of AMP genes. (B) AMP expression following infection with symbiotic versus axenic *S. carpocapsae* IJs. Infection of third-instar *D. melanogaster* larvae with symbiotic *S. carpocapsae* IJs results in AMP expression, while infection with axenic *S. carpocapsae* IJs results in little or no AMP expression. Data for symbiotic IJ exposure are from panel A. For panels A and B, AMP expression was examined at 24 h postexposure to IJs using transgenic fly larvae containing reporter constructs in which an AMP gene promoter was used to drive expression of GFP (27). (C) Infection of second-instar *D. melanogaster* larvae with *X. nematophila* results in AMP expression. AMP expression was examined at 24 and 48 h postexposure to bacteria using the same transgenic fly larvae as used in the experiments described in panels A and B. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$, unpaired *t* test or Mann-Whitney test (infected larvae versus uninfected control larvae of the same genotype; $n = 5$ to 12 trials for each condition). For all graphs, error bars represent standard errors of the means.

course of the experiment) for GFP expression at 24 h postexposure to IJs and quantifying the percentage of fly larvae or pupae expressing GFP. We found that exposure of third-instar fly larvae to symbiotic IJs induced significant expression of four AMP genes: *attacin*, *dipteracin*, *drosomyacin*, and *metchnikowin* (Fig. 3A). Thus, the same subset of AMP genes is induced by exposure to *S. carpocapsae* symbiotic IJs and *H. bacteriophora* symbiotic IJs (4). The percentage of animals showing AMP expression was higher for third-instar larvae exposed to symbiotic IJs than for second-instar larvae exposed to symbiotic IJs (Fig. 3A), most likely because symbiotic IJs were more effective at killing third-instar than second-instar larvae (see Fig. S2 in the supplemental material).

To determine whether AMP expression is a response to the nematode or the bacteria, we first exposed fly larvae to axenic IJs. Third-instar fly larvae were used for this experiment since a higher rate of AMP expression was observed with third-instar larvae than with second-instar larvae (Fig. 3A). We found that whereas fly larvae exposed to symbiotic IJs showed AMP ex-

pression, fly larvae exposed to axenic IJs showed little or no AMP expression (Fig. 3B). Thus, the AMP response observed upon infection with symbiotic IJs is not observed upon infection with axenic IJs.

We then exposed fly larvae to bacteria alone by placing fly larvae on a plate containing a lawn of *X. nematophila*. Second-instar or early-third-instar larvae were used for these experiments since older third-instar larvae did not become infected with bacteria in this assay. Exposure to *X. nematophila* induced expression of the same four AMP genes that were induced by infection with symbiotic IJs (Fig. 3C). The routes of infection differ for fly larvae exposed to symbiotic IJs and bacteria, and we cannot exclude the possibility that AMP expression might vary based on the route of infection. However, our results suggest that AMP expression is primarily a response to *X. nematophila* rather than *S. carpocapsae*.

We note that for comparison of AMP expression following infection of *D. melanogaster* larvae with either symbiotic IJs or axenic IJs, we used nematodes grown *in vitro* on plates containing

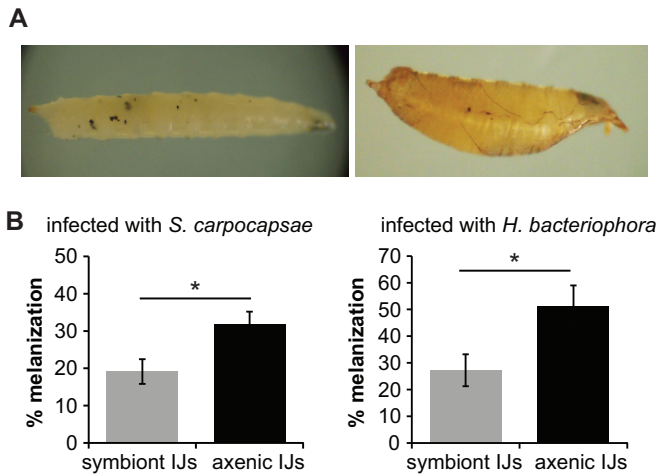


FIG 4 *X. nematophila* and *P. luminescens* partially suppress the melanization response of *D. melanogaster*. (A) Melanization from *S. carpocapsae* infection. The panel shows a representative *D. melanogaster* larva (left) and pupa (right) melanized by axenic *S. carpocapsae* IJs. (B) *X. nematophila* and *P. luminescens* inhibit melanization. Infection of *D. melanogaster* larvae with axenic *S. carpocapsae* (left graph) or *H. bacteriophora* (right graph) IJs resulted in a higher percentage of melanized fly larvae or pupae than infection with symbiont IJs. *, $P < 0.05$, unpaired t test. No melanization was observed in fly larvae not exposed to IJs. Melanization was scored at 48 h postexposure to IJs. Error bars represent standard errors of the means ($n = 7$ to 8 trials).

lawns of *X. nematophila* rather than nematodes grown in waxworms (see Materials and Methods). Growing nematodes *in vitro* was necessary to obtain axenic IJs, and thus both symbiont IJs and axenic IJs were grown *in vitro* for these experiments so that differences in AMP expression between axenic IJs and symbiont IJs could not be attributed to differences in nematode culturing conditions. However, we also directly tested whether AMP expression levels in fly larvae differed following infection with IJs cultured *in vitro* versus *in vivo*. We compared AMP expression in third-instar fly larvae infected with symbiont IJs grown on plates of *X. nematophila* and symbiont IJs grown in waxworms. No significant differences in AMP expression levels were observed following infection with IJs cultured *in vitro* versus *in vivo* (see Fig. S3 in the supplemental material), suggesting that the *D. melanogaster* im-

mune response to *S. carpocapsae* symbiont IJs is similar, regardless of whether the IJs are cultured *in vitro* versus *in vivo*.

***X. nematophila* and *P. luminescens* suppress the melanization response of *D. melanogaster*.** Melanization is a cellular immune response of arthropods that results in melanin production at the wound site and that contributes to pathogen killing and wound healing (11, 50). Previous studies have shown that EPN infection of some insects results in rapid melanization and encapsulation of IJs although in permissive hosts IJs can escape the capsule and kill the insect (51–53). Both *Xenorhabdus* and *Photorhabdus* produce specific inhibitors of phospholipase A₂, a key component of the melanization and encapsulation reactions, suggesting that the bacterial endosymbionts of EPNs promote nematode survival by suppressing these reactions (54, 55). To test whether infection with *S. carpocapsae*-*X. nematophila* or *H. bacteriophora*-*P. luminescens* activates the melanization response, we exposed fly larvae to either symbiont or axenic IJs and scored infected larvae for the presence of visible melanin spots (Fig. 4A). We found that both symbiont and axenic IJs induced melanization but that axenic IJs induced a higher rate of melanization than symbiont IJs (Fig. 4B). These results suggest that *X. nematophila* and *P. luminescens* facilitate the killing of *D. melanogaster* larvae by partially suppressing the melanization response.

Virulence differs for different EPN-*Drosophila* combinations. To examine the versatility of the fruit fly-EPN model system, we compared the ability of symbiont IJs from seven EPN species—*S. carpocapsae*, *S. scapterisci*, *S. riobrave*, *S. glaseri*, *S. feltiae*, *H. bacteriophora*, and *H. indica*—to infect and kill *D. melanogaster* larvae. These EPN species differ dramatically in their host ranges: *S. carpocapsae* and *S. feltiae* have broad host ranges that include insects from multiple orders, *S. scapterisci* has a narrow host range that is limited to orthopterans, and the other species have intermediate host ranges (24, 56–59). *S. feltiae* was also recently shown to be virulent toward *D. melanogaster* (3). We exposed *D. melanogaster* larvae to symbiont IJs of the different EPN species and scored survival at 24 and 48 h. We found that virulence differed greatly among species: *S. scapterisci* and *S. riobrave* were not virulent toward *D. melanogaster* larvae, *S. carpocapsae* and *S. feltiae* were highly virulent, and the other species displayed intermediate virulence (Fig. 5A). Thus, EPNs vary in their virulence toward *D. melanogaster* larvae.

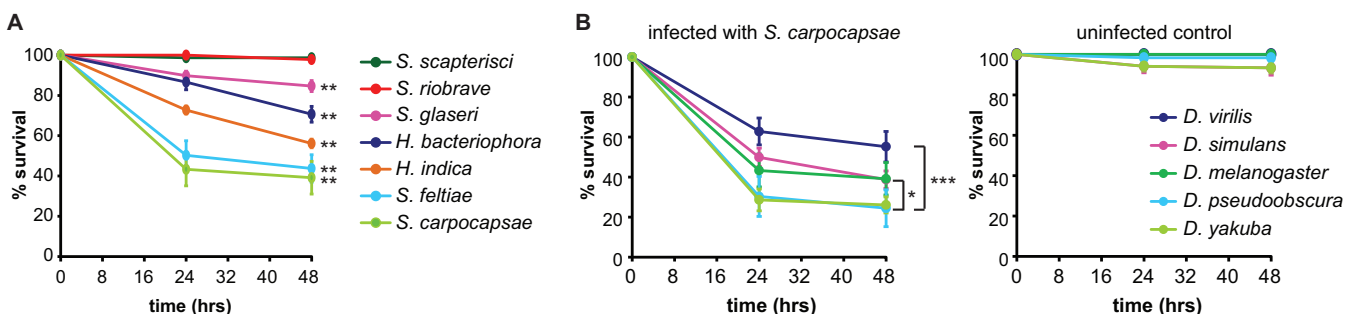


FIG 5 Species specificity of *Drosophila*-EPN interactions. (A) Infection of *D. melanogaster* with EPNs. EPNs vary in their virulence toward *D. melanogaster* larvae. Fly larvae were infected with symbiont IJs. Survival curves are significantly different ($P < 0.0001$, log rank test). Posttest results for each pairwise comparison are shown in Table S1 in the supplemental material. **, $P < 0.01$ relative to *S. scapterisci* (log rank test with Bonferroni correction; $n = 6$ to 8 trials for each condition). (B) *Drosophila* species vary in their susceptibility to *S. carpocapsae* symbiont IJ infection. Survival curves (left graph) are significantly different ($P < 0.01$, log rank test). Posttest results for each pairwise comparison are shown in Table S2 in the supplemental material. ***, $P < 0.001$ for *D. virilis* versus *D. pseudoobscura* and *D. virilis* versus *D. yakuba*; *, $P < 0.05$ for *D. pseudoobscura* versus *D. simulans*. The right graph shows survival curves of uninfected control larvae ($n = 6$ to 7 trials for each condition). For all graphs, the x axis refers to time postexposure, and error bars represent standard errors of the means.

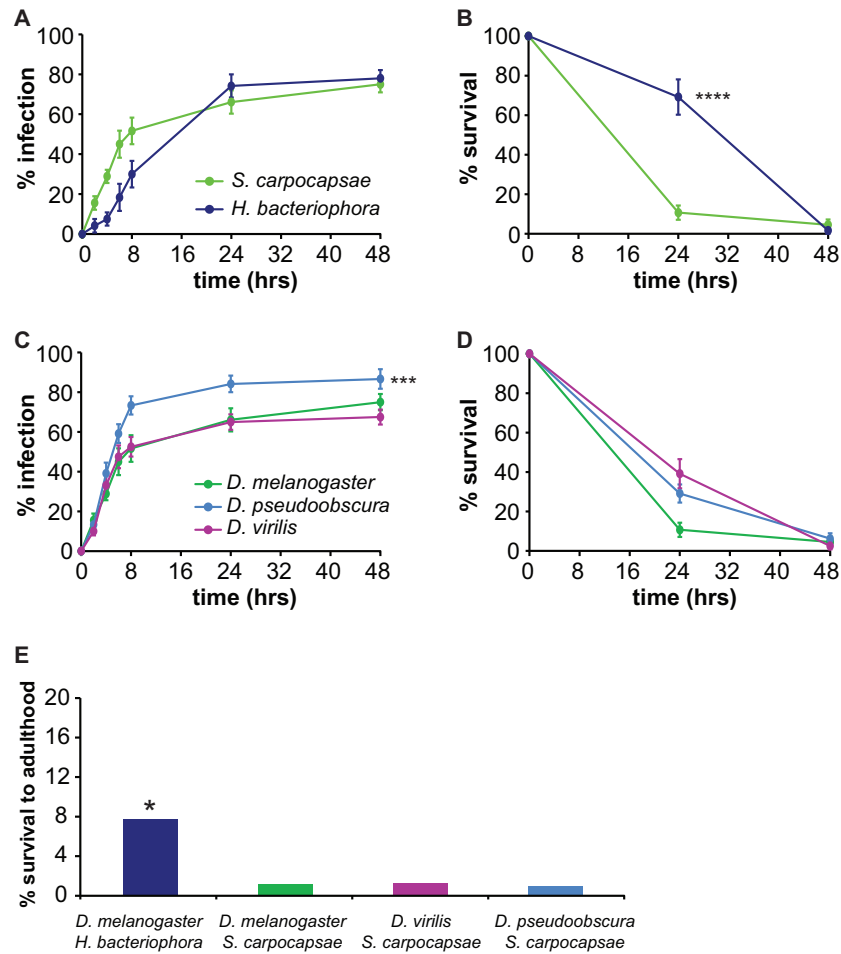


FIG 6 Virulence of EPNs for *Drosophila* species. (A) *D. melanogaster* infection. *S. carpocapsae* and *H. bacteriophora* symbiont IJs infect *D. melanogaster* larvae at the same rate ($P > 0.05$, log rank test; $n = 6$ to 9 trials for each condition). (B) Survival of *D. melanogaster*. *S. carpocapsae* symbiont IJs kill *D. melanogaster* larvae more rapidly than *H. bacteriophora* symbiont IJs. ****, $P < 0.0001$, log rank test. Survival was scored for fly larvae that became infected within 8 h of exposure to IJs ($n = 4$ to 9 trials for each condition). (C) Infection with *S. carpocapsae*. *S. carpocapsae* symbiont IJs infect *D. pseudoobscura* larvae more rapidly than *D. melanogaster* and *D. virilis* larvae. ***, $P < 0.001$ for *D. pseudoobscura* relative to *D. melanogaster* and *D. virilis* (log rank test with Bonferroni correction; $n = 6$ to 9 trials for each condition). (D) Survival from *S. carpocapsae* infection. *S. carpocapsae* symbiont IJs kill *D. melanogaster*, *D. pseudoobscura*, and *D. virilis* larvae at the same rate ($P > 0.05$, log rank test; $n = 6$ to 9 trials for each condition). Note that panels B and D show survival rates only of infected fly larvae rather than of all fly larvae exposed to EPNs. (E) Long-term survival of EPN-infected fly larvae. *D. melanogaster* larvae infected with *H. bacteriophora* symbiont IJs show a higher rate of long-term survival than the other *Drosophila*-EPN combinations. *, $P < 0.05$, chi-square test ($n = 52$ to 104 fly larvae for each condition). For graphs in panels A to D, error bars show standard errors of the means.

We then examined the ability of *S. carpocapsae*, one of the most virulent EPNs for *D. melanogaster*, to infect and kill four phylogenetically and ecologically diverse *Drosophila* species: *D. virilis*, *D. simulans*, *D. pseudoobscura*, and *D. yakuba*. We found that *S. carpocapsae* symbiont IJs were capable of infecting and killing all *Drosophila* species tested (Fig. 5B). However, survival rates following exposure to *S. carpocapsae* symbiont IJs varied across species, with *D. virilis* showing the highest survival rate and *D. pseudoobscura* and *D. yakuba* showing the lowest survival rates (Fig. 5B). Thus, *Drosophila* species vary in their susceptibility to EPN infection.

Studies of larger insects, such as the Japanese beetle *Popillia japonica*, the house cricket *Acheta domestica*, and the Colorado potato beetle *Leptinotarsa decemlineata*, have suggested that differences in virulence among EPNs can be attributed to differences in the abilities of EPNs to infect different hosts as well as differences in the host immune response to infection (51–53). To in-

vestigate the cause of differences in survival rates among *Drosophila* species exposed to different EPNs, we examined infection rates and postinfection survival rates of selected *Drosophila*-EPN combinations. Infection rates were examined by using IJs containing GFP-expressing endosymbionts to facilitate detection of IJs within the host. Fly larvae were considered infected if nematodes were visible inside the body. We found that different *Drosophila*-EPN combinations varied in both the rates at which the fly larvae became infected and the rates at which they succumbed to the infection (Fig. 6). For example, although *S. carpocapsae* symbiont IJs and *H. bacteriophora* symbiont IJs infected *D. melanogaster* at the same rate, *S. carpocapsae* symbiont IJs killed *D. melanogaster* more rapidly than *H. bacteriophora* symbiont IJs (Fig. 6A and B). In contrast, *S. carpocapsae* symbiont IJs infected *D. pseudoobscura* more rapidly than *D. melanogaster* or *D. virilis*, but all three fly species succumbed to infection at the same rate (Fig. 6C and D). Thus, the *Drosophila*-EPN model system can be used to study

differences in both nematode infectivity and the host immune response to nematode infection.

We also assayed the long-term survival of EPN-infected *Drosophila* larvae by exposing fly larvae to symbiont IJs, separating out all fly larvae that became infected by 8 h postexposure to IJs, and monitoring their survival until death or adulthood. We found that *Drosophila* larvae were capable of surviving EPN infection at low levels (Fig. 6E). Moreover, survival rates varied for different EPN species: the long-term survival rate was 1% for *D. melanogaster*, *D. virilis*, and *D. pseudoobscura* infected with *S. carpocapsae* but 8% for *D. melanogaster* infected with *H. bacteriophora* (Fig. 6E). Thus, *Drosophila* larvae are more successful at overcoming some EPN infections than others. Whether long-term survival occurs because nematodes exit the host shortly after infection or because the host immune system overcomes the infection remains to be determined.

Conclusions. Our results demonstrate that both *S. carpocapsae* and its bacterial endosymbiont *X. nematophila* are pathogenic for *D. melanogaster* larvae. We also show that EPN species vary in their virulence toward *D. melanogaster* and that *Drosophila* species vary in their susceptibility to EPN infection. These differences in virulence reflect differences in both the rates at which fly larvae become infected with EPNs and the rates at which infected fly larvae succumb to the infection. All five of the *Drosophila* species tested have sequenced genomes (60), and six of the seven nematode species tested have sequenced or nearly sequenced genomes (61, 62). A comparison of *Drosophila* genomes revealed that many immune-related genes evolve more rapidly than other genes and identified numerous species-specific differences in immune-related genes, including copy number differences in AMP genes (63). Our results establish a versatile model system for investigating at a genome-wide level how genetic differences contribute to the diverse immune responses of insects to parasitic nematode infection.

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