Sex ratio distortion in the *Nesolyx thymus* (Hymenoptera: Eulophidae), an ecto-pupal parasitoid of uzify, *Exorista sorbillans* (Diptera: Tachinidae)

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Abstract. The reproductive alterations induced by maternally inherited α-proteo-bacteria *Wolbachia* to their hosts is a well-documented phenomenon. In *Nesolyx thymus*, a gregarious hymenopterous ecto-pupal parasitoid of the uzify, *Exorista sorbillans*, diagnostic PCR assay using specific primers revealed the presence of *Wolbachia*. Following genetic crossing experiments, we observed a female biased sex ratio of 1 : 9.5 at 25°C and 1 : 3 male to female ratio when the populations were exposed to heat shock 33°C for six hours. Furthermore, we found infection polymorphism, where female parasitoids are infected by *Wolbachia* but males are not infected. Infected eggs develop into females, whereas uninfected eggs develop parthenogenetically into males. The results are discussed in the context of the possible mechanism of sex-ratio bias caused by *Wolbachia*.

INTRODUCTION

*Wolbachia* is a genus of maternally inherited intracellular α-proteobacteria which infect a wide range of arthropods and nematodes (Werren, 1997; Braig et al., 1998; Stouthamer et al., 1999), estimated to occur in about 66% of all known insect species (Hilgenboecker et al., 2008). They are abundant intracellular symbionts and have attracted significant attention in terms of their ability to manipulate host reproduction through cytoplasmic incompatibility (CI), feminization, induction of parthenogenesis and male killing (Werren, 1997; Stouthamer et al., 1999). In CI, infected males are incompatible with uninfected females or females infected with some other strains of *Wolbachia*; however, infected females are compatible with infected or uninfected males (Yen & Bar, 1971; O’Neill & Karr, 1990). This is the most commonly observed *Wolbachia*-induced reproductive phenotype in most of insect orders. Feminization is the reproductive phenotype whereby all genetic males are converted into functional females, and is found in some terrestrial isopods as well as one particular species of butterfly (Rigaud et al., 1991; Kageyama et al., 2002). However, in other lepidopteran species such as the corn borer, *Ostrinia scapulalis* (Crambidae), *Wolbachia* causes lethal feminization of genotypic males (Kageyama & Traut, 2004; Sugimoto & Ishikawa, 2012). During parthenogenesis, unfertilized eggs develop into males (arrehe-notoky) or females (thelytoky) (Werren et al., 2008). Each of these reproductive anomalies enhances female production and hence the reproduction of the bacterium and is collectively referred to as “reproductive parasitism” (Werren et al., 2008). The effect of *Wolbachia* depends on a number of factors, including host genetic background, atmospheric temperature, resource quality and host age. These factors directly affect *Wolbachia* densities within its host, which in turn have context specific effects on the respective host populations. Among these, temperature has a remarkable effect upon *Wolbachia*, associated bacteriophage, involved in horizontal gene transfer (Bordenstein & Werren, 2004) and on host population dynamics (Bordenstein & Werren, 2003; Bordenstein & Bordenstein, 2011; Kraaijeveld et al., 2011).

*Nesolyx thymus* (Hymenoptera: Eulophidae) is an ecto-pupal-gregarious parasitoid of the “uzify”, *Exorista sorbillans* (Diptera: Tachinidae), itself an endo-larval parasitoid of the silkworm moth, *Bombyx mori* L. (Lepidoptera: Bombycidae), and accounting for some 8–10% yield loss in India and other silk growing countries. *N. thymus* is often used as a primary biocontrol agent to regulate the populations in nature (Narayanaswamy & Devaih, 1998). Earlier investigations have associated the efficacy of this agent with its phenomenal host searching ability and parasitization capacity (Kumar et al., 1993; Narayanaswamy & Devaiah, 1998). The successful expansion of the parasitoid is based on its ability to tolerate highly variable temperatures ranging from 15 to 35°C (Jyothi et al., 1993). The overall developmental time of the parasitoid from egg to adult is around 16 days, whilst gravid females can produce around 300 offspring during their lifetime. Interestingly, the sex ratio of the parasitoid has been found to vary considerably between studies (ChannaBasavanna et al., 1993; Aruna & Manjunath, 2010). In addition to its use as a biocontrol agent against uzify, *N. thymus* also hyper-parasitizes...
**Bleparipha zebena** (Walker) (Diptera: Tachinidae), a larval endo-parasitoid of the tasar silkworm, *Antheraea mylitta* (Drury) (Saturniidae) and a primary parasitoid of the house fly, *Musca domestica* L. (Muscidae) the blow fly *Chrysomya rufifacies* (Calliphoridae) as well as an unidentified calliphorid fly (Channabasavanna et al., 1993; Narayanaswamy & Devaiah, 1998). During mass rearing, the house fly is used as an inexpensive and easy to rear host (Aruna & Manjunatha, 2010).

Over the last 25 years, considerable efforts have been made to curtail the uzifly through chemical, physical and biological methods (Narayanaswamy & Devaiah, 1998). Many parasitoids and microbial agents have been studied (Narayanaswamy & Devaiah, 1998; Puttaraju & Prakash, 2005a, b) and currently, more than 2 species belonging to the families Chalcididae, Diapriidae, Encrytidae, Eulophidae and Pteromalidae have been identified as ecto/endo, larval/pupal or solitary/gregarious parasitoids of this fly. These hymenopterans inflict some 82% of the biocontrol mortality with the remaining achieved through predatory insects and vertebrate predators and some by pathogenic bacteria (Narayanaswamy & Devaiah, 1998; Puttaraju & Prakash, 2005a, b). Because of the prominence of *N. thymus* and the need to improve production, the reproductive behavior of the *N. thymus* and possible interactions with the thelytoky inducing *Wolbachia* endosymbiont were presently examined. Furthermore, since there is no experimental evidence that *Wolbachia* has any effect on sex ratio in the parasitoid, only circumstantial correlation in terms of sex ratio bias, this aspect was also investigated. To this end, we for the first time screened *Wolbachia* infection in *N. thymus* using a diagnostic PCR approach in order to determine whether or not the bacteria do play a role in the reproduction of the parasitoid.

### MATERIAL AND METHODS

#### Collection and rearing of parasitoid *N. thymus*

Around 1,000 parasitoids were collected from the Ramanagara silkworm cocoon market. These were maintained in the laboratory on uzifly pupae at a ratio of approximately 1 parasitoid : 10 hosts. Ten discrete generations were maintained in the laboratory to eliminate any variation in the population due to environmental variations in three 1 l conical flasks, covered with muslin cloth, with ~ 250 parasitoids per flask in the first generation followed by ~ 500–800 parasitoids per flask in subsequent generations. Each flask contained a separate individual population indicated as blocks 1, 2 and 3. Two days after emergence from the host pupae, females in the flask were provided with 250-uzifly pupae per flask for one day and monitored until emergence in separate conical flasks. Populations were maintained in an insect rearing chamber at a temperature of 25 ± 1°C and 80–85% RH and 12L: 12D conditions.

#### Genomic DNA isolation and PCR amplification

Individual male and female parasitoid DNA was extracted using ZR Insect/Tissue DNA kit-STM (Zymo Research, Irvine, CA, USA). Whole insects were ground in ZR Bashing Bead™ lysis tube and homogenized in 600 µl lysis solution, DNA being extracted according to the manufacturer’s protocol, with at least 15 males and females used per population. The dried DNA pellet was re-suspended in 25 µl 10 mM TE buffer, pH 8.0. DNA was quantified using a spectrophotometer (Bio-Rad SmartSpec™ plus) and stored at −20°C for further use.

The polymerase chain reaction (PCR) assay was used to amplify a *Wolbachia* surface protein (WSP) gene using the primer pair: wsp81F 5’-TGG TCC AAC AAG TGA TGA AGA AAC-3’ and wsp891R 5’-AAA AAT TAA ACA CTC TCT CA-3’, which amplify around 630 bp (Braig et al., 1998). This was done in an Eppendorf thermocycler in 20 µl reaction volume containing 1× PCR buffer, 0.2 mM dNTP’s, 2.5 mM MgCl2, and 0.5 unit Taq DNA polymerase (MBI-Fermentas, Amherst, NY, USA). 0.1 µM of each forward and reverse primer, 20 ng of template DNA and a final volume of sterile water to make up a total of 20 µl reaction mixture. PCR conditions were: initial denaturation at 94°C for 3 min followed by 40 cycles with denaturing at 94°C for 1 min, primer annealing at 55°C for 1 min, primer extension at 72°C for 2 min, followed by a final extension step at 72°C for 10 min. For positive and negative controls, uzifly and silkworm DNA was used, respectively (Prakash & Puttaraju, 2007). In addition, an in-sit individual 18s rDNA primer pair amplifying a 555 bp segment of the host genome (18S F1 5’-TTG GAG GCC AAG TCT GGT GC-3’ and 18S R1 5’-ACT TCG GCC GAT CCG TAG CT-3’) were used to determine the quality of the DNA extraction (Wenseleers & Billen, 2000). PCR products were separated on a 1.2% agarose gel run in 1× TBE buffer, pH 8.0 for a length of 5–6 cm at constant 65 volts. The gel was stained with 0.5 µg/ml gel ethidium bromide just prior to casting. A 1 kb standard molecular weight marker (Chromous Biotech™, Bangalore, India) was used to estimate amplified band size.

### Crossing experiments

Parasitoids were sexed following emergence. Single female and male parasitoids were placed in 100 ml conical flasks covered with muslin cloth and provided with 20 two day old uzifly pupae. The host pupae were replaced every day for five days and parasitized pupae maintained in separate 500 ml conical flasks for each block until the emergence of *N. thymus*, adult parasitoids being sexed each day after emergence. For each block, 20 replicate crosses were maintained, whilst each flask was provided with a 50% honey solution on cotton balls as food for the adult parasitoids (Aruna & Manjunatha, 2010).

#### Heat shock treatment

About 250 one day old parasitoids from each block were exposed to 33°C heat shock treatment for six hours in an incubator. Following heat shock, parasitoids were maintained at 25°C for one more day and crosses then made as shown above with 20 replications from each block.

### Data analysis

Experimental data were pooled and block means, standard deviations and errors calculated. Means were compared by chi-square testing. Since chi-square tests provided only 4 degree of freedom, a correction value of 0.5 was subtracted from the deviation value before squaring (Khan & Khanum, 1994). To test the null hypothesis, the calculated values of chi-square were compared with the table value at 5% and 1% confidence levels.

### RESULTS and DISCUSSION

In all three populations, female *N. thymus* proved PCR positive for *Wolbachia* and yielded the expected band size of around 630 bp, whereas males were negative (Fig. 1). In addition, insect specific 18S rDNA primer yielded an expected band of around 550 bp which confirmed the quality of insect DNA used for amplification of *Wolbachia*. The genetic crossing results revealed that the sex ratio was bi-
assessed towards females, with the number of offspring varying from 83 to 406, and with an average of 217.7. The number of males and females were 20.6 ± 3.5 and 197.1 ± 11.9, respectively, i.e., a ratio of 1:9.5 (Fig. 2) at 25°C, significantly different from a 1:1 ratio at $P$ = 0.01 using a chi-square test ($\chi^2$ = 116.93; $P$ < 0.001).

In addition, temperature treatment for 6 h at 33°C on one-day-old adult parasitoids revealed a reduction in the total offspring number of 23.5% (they produced an average of 166.6 offspring/female) compared to parasitoids reared at 25°C. The sex ratio of offspring of temperature treated parasitoids was on average 40.7 males and 125.9 females (approximately a ratio of 1 male to 3 females) and significantly different from a 1:1 ratio at $P$ = 0.01 ($\chi^2$ = 44.542; $P$ < 0.01). However, the female bias was reduced in heat shock treatment, compared to parasitoids reared at 25°C (Fig. 2). PCR assay confirmed the absence of Wolbachia in male offspring, whose parents had been exposed to heat shock, and thus produced a greater number of males, in turn indicating that temperature had a negative effect on the parasitoid-Wolbachia association. It would be interesting to rear parasitoids at different temperatures for one or more complete generations to quantify the effects of this treatment on Wolbachia and its host. Our hypothesis is that the infected eggs of N. thymus may develop into females, while uninfected eggs develop into males. But we cannot ruled out the possibility of low Wolbachia infection in males which may not be detected through standard PCR approaches.

Earlier Aruna & Manjunath (2010) reported a 1:17 male to female ratio in N. thymus; however, they had not considered the possible presence of Wolbachia and its effects. This could simply be due to arhenotoky, as is typical of Hymenoptera. However, the sex ratio varies between studies which in turn might be due to variation in the level of Wolbachia infection, a possibility these authors seemingly had not considered. High Wolbachia infection rates in females, which facilitates infection of more eggs, leads to the production of more females. In the case of lower Wolbachia levels, asynchronous transmission may result in fewer uninfected eggs, which induces parthenogenetic male production.

Wolbachia is also sometimes known to increase host fitness and fecundity (Dedeine et al., 2001; Puttaraju & Prakash, 2005a,b, 2009). This characteristic is being exploited in several biocontrol programs (e.g. Brelsfoord &...
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