REVIEW

Molecular Identification and Evaluation of *Orius* species (Heteroptera: Anthocoridae) as Biological Control Agents

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Abstract

Orius strigicollis is a predatory bug that attacks small arthropods, such as thrips, and augmentative release of commercial strains has been widely conducted in greenhouses in Japan. The accurate evaluation of its effectiveness is essential for successful biological control programs. However, because *Orius* species occur naturally in the field, it is difficult to discriminate *O. strigicollis* from other *Orius* species. Furthermore, it is necessary to discriminate commercial strains of *O. strigicollis* from field populations. In this review, we introduce two types of molecular procedures: (1) a multiplex polymerase chain reaction technique to identify Japanese *Orius* species, and (2) microsatellite DNA markers that can be used to measure genetic diversity within populations of *O. strigicollis* and to discriminate field and commercial strains. The usefulness of molecular ecological approaches is also discussed.

Discipline: Insect pest / Plant protection

Additional key words: augmentative release, biopesticide, microsatellite DNA marker, natural enemy, polymerase chain reaction

Introduction

Chemical control of agricultural pests always poses a risk of them developing resistance to the pesticides. Therefore, integrated pest management (IPM) is essential for the control of pests, and augmentative release of natural enemies is one of the most important techniques for IPM²⁹. In Japan, natural enemies must be registered as biopesticides when they are provided as commercial products, and 16 species were registered and sold as biopesticides in September, 2008. Although the registered species were exotic in the early stage of IPM, researchers

have focused recently on the indigenous natural enemies to the pests, because such species are innately adapted to the environment in Japan and unexpected nontarget effects can be avoided by using indigenous species²⁸.

The anthocorid bugs, particularly *Orius* spp. (Heteroptera: Anthocoridae), are the most promising biological control agents of various minute pests, such as thrips, aphids and mites¹⁵. Among seven species in the genus distributed in Japan, *O. sauteri* (L.), *O. strigicollis* (Poppius), *O. minutus* (L.), and *O. nagaii* Yasunaga are frequently found in the field^{30–32}. Among these four species, *O. sauteri* has been investigated most often as a biological control agent of thrips, and its effectiveness in suppress-

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ing thrips has been demonstrated both in the field and in greenhouses^{11,13,18,19}. Because of the high incidence of reproductive diapause with short day length^{6,7,9,12,22} and low reproduction and predation rates at low temperatures^{8,18,19}, however, *O. sauteri* is not suitable for use during the winter. In contrast, because of a lower incidence of reproductive diapause^{9,22} and the ease of mass production, *O. strigicollis* was proposed as an effective biological control agent and was registered as a biopesticide in 2001.

In biological control programs, it is essential to select appropriate species and evaluate released strains. These *Orius* species, however, are morphologically similar to each other and difficult to discriminate. Furthermore, released biopesticide strains cannot be distinguished easily from field populations. With the arrival of a vast array of molecular methods, however, it is now possible to develop a system to recognize different species and strains without morphological expertise²⁴. In this article, we review molecular markers by which *Orius* species can be identified and evaluated, and describe our new understanding about the distribution of *Orius* species in Japan.

Molecular identification of *Orius* species and its application

1. Molecular identification techniques

Because several *Orius* species often occur sympatrically²¹, proper identification of each species is necessary for the evaluation of their efficacy in biological control programs. Males are usually identified by their genitalia after dissection, but this process is time-consuming and identifies only adult males. Therefore, a polymerase chain reaction–restriction fragment length polymorphism (PCR-RFLP) method was developed to discriminate five Japanese *Orius* species^{4,17}. This method is complicated, however, because restriction enzyme treatment is required after PCR amplification. Therefore, a multiplex PCR method, allowing simultaneous amplification of several fragments in a single PCR reaction, was developed based on the DNA sequences of nuclear ribosomal DNA by designing species-specific primers for the internal transcribed spacer regions³. Based on tests of 350 individuals of five *Orius* species (*O. minutus*, *O. sauteri*, *O. strigicollis*, *O. nagaii*, and *O. tantillus*), these species could be clearly distinguished by a single multiplex PCR reaction (Fig. 1). This technique is simple and requires no procedures other

Fig. 1. Electrophoretogram of multiplex polymerase chain reaction for five *Orius* species and three prey species

PCR products of a single insect were applied to each lane. Only *O. minutus* showed PCR products longer than 800 bp. *Orius tantillus* showed one band with a length of ca. 500 bp. *Orius strigicollis*, *O. sauteri* and *O. nagaii* showed two bands, one longer than 600 bp, and a shorter band that depended on the species: longer than 400 bp (*O. strigicollis*), between 400 and 200 bp (*O. sauteri*) and shorter than 200 bp (*O. nagaii*). Prey species did not show any bands. M: molecular weight marker; 1-4: *O. minutus*; 5-9: *O. strigicollis*; 10-16: *O. sauteri*; 17-18: *O. nagaii*; 19: *O. tantillus*; 20: *Thrips palmi*; 21: *Ephestia kuehniella*; 22: *Tetranychus kanzawai*; 23: no template DNA³.

than PCR and subsequent electrophoresis, thus reducing the cost of identifying species and increasing the number of specimens that can be identified.

2. Geographic distribution of *Orius strigicollis*

Among the four major *Orius* species, only the distribution of *O. strigicollis* is limited to southwestern Japan. However, their distribution areas and the factors limiting them have not been surveyed in detail. By using molecular discrimination techniques as well as morphological dissection, we identified *Orius* species collected on flowers of the tall goldenrod, *Solidago altissima* L., in Kanto district, east-central Japan, which is the northern limit of the distribution of *O. strigicollis*²³. The factor determining this northern limit of *O. strigicollis* is the low temperature in winter²³; the species is distributed in areas where the daily minimum temperature in January is higher than -1°C on average and the number of days with a minimum temperature lower than 0°C was fewer than 50 days. Due

to this limitation, *O. strigicollis* is distributed only in the coastal and urban areas in Kanto district.

In autumn from 2002 to 2007, we collected and identified *Orius* species from tall goldenrod flowers throughout southwestern Japan (Fig. 2). *O. strigicollis* was found only in coastal and urban areas. Interestingly, *O. strigicollis* is the dominant species in southern Kyushu but it is rarely distributed in northern Kyushu, where this species should be able to survive based on the temperature in winter²³. Because *O. strigicollis* is distributed in the Okinawan islands, Taiwan and southern China as well³¹, the species may have originated in the south of Japan or more southern countries, with its distribution spreading northward with recent climate warming²³.

Conservation biological control, which manipulates the environments in agro-ecosystems to enhance natural enemies¹⁴, has recently attracted a great deal of attention in Japan. The detailed information on the distribution of *Orius* species described here will be necessary for the

Fig. 2. Species composition of *Orius* collected from flowers of tall goldenrod, *Solidago altissima*, in autumn from 2002 to 2007

The numbers of *Orius* adults identified are shown in circles. *Orius strigicollis* was found only in coastal and urban areas in southwestern Japan. Arrows and the white square indicate field populations and the experimental sites (shown in Fig. 6).

Fig. 3. Species composition of *Orius* collected from the various flowers

The composition was significantly different among the height of the flowers on which *Orius* species were collected (high: higher than 200 cm above the ground; mid: between 20 and 200 cm; low: less than 20 cm; Fisher's exact test; $p < 0.0001$).

▨ *O. minutus*, ■ *O. strigicollis*, □ *O. sauteri*, ▤ *O. nagaii*

proper conservation of these species and the management of their habitat.

3. Vertical distribution of *Orius* species

We collected *Orius* adults from flowers of a variety of plants and identified them by male genitalia or molecular techniques. Although *O. sauteri*, *O. strigicollis*, *O. minutus*, and *O. nagaii* can be found on a variety of plants, the microhabitats of each species seem to be different (Fig. 3). *Orius sauteri* adults were abundant on flowers at less than 20 cm above the ground, such as white clover (*Trifolium repens* L.). *Orius minutus* was mainly found on flowers higher than 200 cm above the ground, such as chestnut (*Castanea crenata* Sieb. et Zucc.) and apple (*Malus pumila* Mill.) trees, which was in accord with the previous observations³¹. *Orius strigicollis* tended to inhabit flowers at intermediate heights above the ground between 20 and 200 cm. The few *O. nagaii* individuals were found only on low flowers. The preference of these four species seems to have no relation to plants or prey species.

To clarify whether *O. sauteri*, *O. strigicollis* and *O. minutus* showed a differential distribution by height on the same plant species, we investigated the species composition at different heights on mulberry (*Morus alba* L.) trees. A mulberry field (ca. 4,000 m²) was divided into four sections; the heights of trees were adjusted to high (180–190 cm) in two sections and low (70–80 cm) in the other sections by trimming. *Orius* individuals found within 15 cm of the top of newly growing shoots were collected and identified. The distribution of the three *Orius* species differed significantly among heights (Fig. 4).

Fig. 4. Differential distribution of three *Orius* species at the different heights (high: 180–190 cm; low: 70–80 cm) of *Morus alba*

The distributions of the species were significantly different among heights (χ^2 -test; $p < 0.01$).

These findings allow the proposal of new usages of *Orius* as biological control agents. For example, *O. sauteri* has a good possibility for biological control on low-height plants, and *O. minutus* would be a good candidate to control fruit tree pests. In fact, *O. minutus* was shown to control two-spotted spider mites, *Tetranychus urticae* Koch, in apple orchards²⁵. These two species can play

important roles in the situations where *O. strigicollis* cannot control pests effectively.

Intraspecific DNA markers of *Orius strigicollis*

1. Microsatellite DNA markers

Orius strigicollis is now widely used in Japan as a biological control agent in greenhouses²⁹. Because *O. strigicollis* is common in the field of coastal and urban areas, it is difficult to judge the effectiveness of released strains in controlling pests. To discriminate among intraspecific strains of *O. strigicollis*, we developed six microsatellite DNA markers that distinguish field and commercial populations².

When genetic diversity within each field population collected on the tall goldenrod were tested, the inbreeding coefficient²⁷ (F_{IS}) was significantly different from zero, whereas biopesticide populations did not show such deviation. This indicated that the field populations of *O. strigicollis* would generally show heterozygote deficits. *Orius* adults appear to immigrate into tall goldenrod habitats in late autumn, attracted by herbivorous insects crowding on the flowers²³. Thus, the heterozygote deficits might result from population genetic structures, such as the Wahlund effect.

Genetic differentiation among populations was estimated by the fixation index²⁷ (F_{ST}) values. Field populations collected in geographically close areas showed no genetic differentiation. In contrast, two biopesticide strains were markedly differentiated genetically from any field populations, revealing the high genetic specificity of these strains. Specific alleles for biopesticide populations would be very useful for distinguishing released biopesticide from field populations. No such specific alleles were found, however, probably because the commercial *O. strigicollis* populations were derived from somewhere in Japan.

Biopesticide populations were also characterized by significantly less allelic richness than that of field populations (Fig. 5). The biopesticide populations might have lost genetic diversity through bottleneck effects and genet-

ic drift during laboratory rearing, as well as through possible selection during mass production. The importance of quality control in mass-production procedures of biological control agents has been recognized, and the genetic structure of laboratory populations is one of the most important factors¹. When several strains are reared, contamination of different strains should be avoided. Some genetic effects such as bottleneck and genetic drift often reduce or alter the attributes of reared populations. By quantifying genetic composition within populations, such deterioration can be easily monitored.

2. Post-release evaluation

The effectiveness of *O. strigicollis* as a biological control agent released in a greenhouse was evaluated by microsatellite markers²⁰. Eggplants were planted in October 2002, and the population dynamics of *Orius* and thrips were monitored until June 2003. A commercial strain of *O. strigicollis* was released three times in the greenhouse where an IPM program was conducted. In contrast to the insecticide-sprayed greenhouse, in which thrips could not be controlled after March, the density of thrips was at a low level in the IPM greenhouse from October to June. After the harvest, *Orius* collected in the greenhouses and on weeds outside were identified by the multiplex PCR method (Table 1). In the IPM greenhouse, almost all the *Orius* individuals were *O. strigicollis*, and in the insecticide-sprayed greenhouse the dominant species was also *O. strigicollis*. Outside of the greenhouses, however, the most dominant species was *O. nagaii*, followed by *O. sauteri*. In this case, the *O. strigicollis* collected in the IPM greenhouse should be descendants of the released strain; however, it is also possible that wild *O. strigicollis* intruded into the greenhouses. We compared the microsatellite allele frequencies of *O. strigicollis* collected from three experimental sites (the IPM greenhouse, insecticide-sprayed greenhouse and weedy area), four field populations in this district (indicated by arrows in Fig. 2), and a released biopesticide population. The three experimental populations were genetically very close to the biopesticide population, whereas the field populations were significantly different

Table 1. Species composition of *Orius* collected at three experimental sites, where a biopesticide strain was released in a greenhouse to control thrips on eggplants²⁰

Site	<i>O. strigicollis</i>	<i>O. sauteri</i>	<i>O. nagaii</i>	n*
IPM greenhouse	91.7%	8.3%	0.0%	24
Insecticide-sprayed greenhouse	79.4%	17.6%	2.9%	34
Weedy area	29.0%	32.3%	38.7%	31

*n: Total number of *Orius* adults collected.

Fig. 5. Example of allele frequencies of a microsatellite marker in *Orius strigicollis*

Allele frequencies of *OstMS009* in two fields and two commercial populations are shown. The number of alleles detected was greater in the field populations than in the biopesticide populations. In a biopesticide Oristar-A® (Sumitomo Chemical, Co. Ltd.), no polymorphisms were detected in this locus².

Fig. 6. Pairwise fixation index (F_{ST}) among *Orius strigicollis* populations

A: among field populations (collection sites are shown in Fig. 2); B: between field and experimental sites; C: between field populations and a biopesticide population; D: among experimental sites; E: between experimental sites and a biopesticide population. When F_{ST} is zero, the populations compared are considered to be genetically identical²⁰.

(Fig. 6). Therefore, the *Orius* strain collected in the IPM greenhouse, which successfully controlled thrips, was not a field strain but the released biopesticide strain. These results indicated that the preservation of released *Orius* by reducing insecticide spray was important in controlling thrips in the IPM program.

Conclusion and future remarks

Biological control programs often fail because of the lack of understanding about the fundamental biology of the biological control agents or its target pests, including the population genetic structure, dynamics, overwintering, reproduction, and host preference¹⁶. Identification of species is the first step in understanding such factors, but this is sometimes difficult due to the agents' small size. The advent of PCR, however, has allowed amplification of molecular markers from small amounts of DNA, even from a single insect egg.

Molecular markers can also help us to evaluate the effect of biological control agents at the intraspecific level. For example, although small arthropods cannot be observed directly during a long-term cultivation period, inherited genetic markers can enable us to monitor them over generations. Microsatellite DNA markers have been widely employed in molecular ecological studies because their high polymorphism makes the markers suitable for studying intraspecific variation. However, microsatellite markers are difficult to isolate from the arthropod genome,

so that the molecular ecological approach has not been established in biological control. Nevertheless, through the recent development of isolation techniques³³, microsatellite markers have been developed for biological control agents, especially in parasitoid wasps^{5,10,26}.

By using molecular markers, the uncertain factors that previously caused difficulties in biological control programs can be measured quantitatively. We hope that the molecular ecological approach reviewed here will be developed and used more widely for the further development of IPM programs.

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