Synergy between Cyt1Aa and Cry Toxins

Bti is highly efficient and specific against mosquito and black fly larvae.3 Most importantly, no resistance has been observed in nature after about 30 years of extensive use worldwide.4 Different activities and modes of action of its four major toxins form a lethal combination against larvae of all mosquito species tested.5 Resistance is not selected due to synergy among Bti components, mostly the low-toxic, non-specific Cyt1Aa. High synergy levels affected by Cyt1Aa were observed by Crickmore6 and Wirth,7 the latter also demonstrated that Cyt1Aa prevented selection of resistant mosquitoes.8

The question raised was whether a combination of anti-Lepidopteran toxins with Cyt1Aa imitates this rare advantage of Bti. To partially answer this question, two genes were cloned for expression in Escherichia coli, cry1Ac (from Bt ssp. kurstaki) and cry1Ca (from Bt ssp. aizawai), with and without cyt1Aa, and tested against three pests, Helicoverpa armigera, Pectinophora gossypiella and Spodoptera littoralis.9 Co-expression of all three genes, and with p20 encoding an accessory protein (for reasons beyond the scope of this report), indeed synergized toxicity against H. armigera but antagonized it against P. gossypiella. Moreover, very high toxicity against S. littoralis and huge synergy value between the two tested Cry’s were found without Cyt1Aa and P20. Thus, one cannot predict which gene combination would be useful in pest control, and each idea must experimentally be tested separately.

Cyanobacteria to Deliver Bti Toxins against Mosquitoes

Several disadvantages hamper the use of Bti: in nature, it does not proliferate whereas the toxins disappear by sinking and adsorption to silt particles and are inactivated by sunlight.3 To achieve a real biological control, the vector must multiply in the same niche as its target. Photo-synthetic cyanobacteria have several additional features that render them excellent candidates to control mosquito larvae;10 their floating capacity avoids sinking and adsorption to silt hence keeps them in the same zone...
Figure 1. *Anabaena siamensis* is ingested and digested by *Aedes aegypti* larvae. A second instar larva of *A. aegypti* ingests (A) *Anabaena siamensis* (B), excreting from its anus, (C) leaving behind digested filaments and intact heterocysts (D).

Cyt1Aa synergizes the Cry’s in *Anabaena* as well, and the toxic activity is protected from sunlight inactivation. Our attempts to circumvent the disadvantages of Bti by using the nitrogen-fixing, filamentous cyanobacterium *Anabaena* PCC 7120 were quite successful. The first condition for cyanobacteria to control mosquito larvae, feeding them, was demonstrated (Fig. 1): the closely related species *Anabaena siamensis* is ingested and digested by larvae of *Aedes aegypti*, similarly to *Anabaena* PCC 7120 (not shown).

In cooperation with the Sanger Institute, we sequenced pBtoxis, the 128 kb plasmid of Bti that harbors all the genetic information necessary for mosquito larvicidal activity. All 15 possible combinations (2^4 - 1) of four genes, three encoding the toxins Cry4Aa, Cry11Aa and Cyt1Aa and the accessory P20, were cloned for expression in *E. coli*. This protocol of cloning gene combinations under identical promoters allowed comparisons of toxicities and synergy levels among the toxins in vivo. Cyt1Aa was indeed found to synergize Cry4Aa and Cry11Aa hence is anticipated to reduce the likelihood of selection for resistance in the target organisms.

The most toxic combinations were appropriately moved into *Anabaena* PCC 7120 and confirmed our working hypotheses: Cyt1Aa synergizes the Cry’s in *Anabaena* as well, and the toxic activity is protected from sunlight inactivation in semi-field conditions. They were about seven-fold more effective than a commercial preparation of Bti itself. One of our future plans, adding the last major Cry gene cry4Ba to this battery, would improve this bio-control agent.

**Environmental Considerations**

For various reasons, field tests to release living genetically engineered microorganisms are not yet allowed worldwide. One justifiable reason that has been demanded by The European Council Directive, namely the use of markers that confer resistance to “clinically used” antibiotics must be phased out! Drug resistance markers must be removed from transgenic clones before they are even to be considered for release in nature. Release to the environment of *Anabaena* transgenic clones such as ours that were derived by selection of antibiotic resistance markers requires marker-free strains. The most elegant way to achieve this goal is by site-specific recombination.

The site-specific recombination system of the λ-like coliphage HK022, which has been implemented in *Arabidopsis* plants and in human cells, is designed to remove these genes from the...
Anabaena genome. The responsible enzyme, Integrase (Int) catalyzes site-specific integration and excision of DNA provided that the recombination target sites attP + attB or attR + attL, respectively, are available. In human cells Int is active on the extra-chromosomal level with plasmids as well as on the chromosomal level, in both cis and trans orientations.

Expression of lacZ in *Anabaena* PCC 7120 was designed to demonstrate the Int-catalyzed excisive recombination reaction, whether located on a plasmid or on the chromosome. A plasmid pMVO carrying the four Bti toxin genes was constructed such that its antibiotic resistance marker *nptII* can be excised with Int (Fig. 2). This plasmid was introduced into the *Anabaena* chromosome by homologous recombination after conjugation using neomycin selection. The excision of *nptII* along with additional unnecessary DNA (*luxAB*) out of the resultant mosquito larvicedal transgenic *Anabaena* is underway.

How Can Maize Control Mosquitoes?

Pollen of maize (*Zea mays*) provide complete food source for *Anopheles arabiensis* larvae, which is the reason for the sharp rise in malaria prevalence in Africa during blooming seasons. Vast quantities of maize pollen accumulate on the surface of nearby puddles, enhancing development of mosquito larvae in breeding sites that lie within 50–60 meters range. Moreover, maize pollen is phagostimulant for mosquito larvae. Maize is therefore engineered to express a decapeptide (*YDPAPPPPPP*) that exerts its effects against a relatively narrow range of targets, starves the larvae to death by blocking TMOF (Trypsin Modulating Oostastic Hormone) of *A. aegypti*. This hormone, an unblocked decapeptide (YDPAPPPPPP) that exerts its effects against a relatively narrow range of targets, starves the larvae to death by blocking TMOF (Trypsin Modulating Oostastic Hormone) of *A. aegypti*. This hormone, an unblocked decapeptide (YDPAPPPPPP) that exerts its effects against a relatively narrow range of targets, starves the larvae to death by blocking TMOF (Trypsin Modulating Oostastic Hormone) of *A. aegypti*. This hormone, an unblocked decapeptide (YDPAPPPPPP) that exerts its effects against a relatively narrow range of targets, starves the larvae to death by blocking translation of trypsin-like mRNA in the midgut. Since starved larvae are 6–35-fold more sensitive to Bti toxins than are fed larvae, TMOF is anticipated to synergize the Bti toxins in suppressing larval densities around fields of maize genetically modified appropriately. Continuous anti-vector coverage for an entire village is likely to be achieved with a few patches of transgenic maize producing larvicideal pollen. Transformed plantlets with *cry11Aa-tmfA*, *cry4Aa-tmfA* and *cry1Aa-tmfA* have been generated, moved to the greenhouse (to be published elsewhere), and additional gene combinations are currently being prepared.

Anti-Coleopteran Active Genes to Control Capnodis ssp.

Our recently-embarked project is to discover Bt genes that will control a pest prevalent in countries surrounding the Mediterranean. Three ssp. of the flat-headed borer Capnodis, *Capnodis tenebrionis*, *C. carbonaria* and *C. cariosa*, kill trees of cultivated stone-fruits. The larvae destroy the root systems of almond, apricot, cherry, nectarine, peach, plum and pistachio. Tree mortality and economic losses are reported from all Southern-European and Mediterranean countries. Since natural occurring arthropod enemies of Capnodis are rare, growers use intensively organophosphates or carbamates onto the foliage or the stem and the surrounding soil. The populations of Capnodis increase in areas where they had been considered minor pests few decades ago. Development of environmentally friendly measures to control them is thus highly important.

As the first step to achieve this goal, an artificial diet was recently developed, and is currently exploited to screen Bt strains for toxicity against them. Of a battery of 215 field isolates, 38 that were found to include at least one gene encoding anti-Coleopteran Cry toxin are being bioassayed. The genes from the best isolates will be cloned for expression in the roots of the target trees.

Concluding Remarks

Use of environment friendly and cost effective alternatives to chemical pesticides improves health and safety, enhances crop output and lowers levels of pollution. Toxins of entomopathogenic bacteria have become leading bio-pesticides to control populations of insect pests and vectors transmitting severe human diseases. Implementation of innovative ideas to exploit molecular methods and interactions between organisms, together with considering various ecological aspects, is likely to become hallmark of future generations.

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Figure 2. Plasmid pMVO, designed for excision by Int via site-specific recombination between the *attR* and *attL* sites. The 23.4 kb pMVO carries the four Bti toxin genes (*cry4A, cry11A, p20* [twice] and *cry1A*) and antibiotic resistance marker *nptII*. Two promoters were introduced, *Pemh* (of the photosystem II’s D1) and *P<sub>T7</sub>* (T7 phage early promoter) at the denoted positions. *nptII*, neomycin/kanamycin resistance gene; *orf all3924*, a PCR amplified sequence encoding a probable penicillin amidase (see in http://bacteria.kazusa.or.jp/cyanobase/index.html).
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