Cotton bollworm
Helicoverpa armigera

Damage symptoms

Larvae attack all stages of plant growth. In conventional cotton (non-Bt varieties), larval feeding can result in; seedlings being tipped out, chewing damage to squares and small bolls causing them to shed, and chewed holes in maturing bolls, preventing normal development and encouraging boll rot. In any year an average of 15% of Bollgard II area may carry *Helicoverpa* larvae at or above the recommended threshold levels for a short period during peak to late flower. In Bollgard II cotton, chewing damage is mostly confined to fruit and may lead to yield loss.

Sampling

Sample the **egg and larval growth stages** of the pest. The growth stages of the cotton bollworm are defined as:

White egg	WE	pearly white
Brown egg	BE	off-white to brown
Very small larvae	VS	0 mm–3 mm
Small larvae	S	3 mm–7 mm
Medium larvae	M	7 mm–20 mm
Large larvae	L	> 20 mm

Eggs are laid on plant terminals, leaves, stems and the bracts of fruit. Larvae may be found on terminals, the upper or lower surface of leaves, inside squares, flowers and bolls and along stems. Sample the whole plant.

Sample **fruit retention** or fruiting factors once squaring begins, to gauge what level of damage is being caused to the crop.

Sample **key beneficials**. This information will allow thresholds based on the predator to pest ratio to be applied. Collect eggs to check for parasitism by *Trichogramma*.

Frequency

Check at least 2 times/week in both conventional and Bollgard II crops.

Begin cotton bollworm sampling at seedling emergence. Cease sampling when the crop has 30–40% open bolls.

H. armigera larvae (left) have pale hairs compared to darker hairs.on *H.punctigera* larvae (right). (Hugh Brier, DAFF Qld)

Methods

Through the entire season, cotton bollworms are most accurately sampled using visual methods. Check at least 30 plants or 3 separate metres of row for every 50 ha of crop.

Larger samples will give more accurate estimates. Fields are rarely uniform, lush areas often occur in head ditches and these are more attractive to insects. The crop variability within the field may determine the minimum number of sampling points required.

Thresholds

Using eggs as the basis of a threshold can be very misleading as not all eggs hatch. Successful egg hatch has been measured to be 20% early season, 25% mid season and 40% late season. Early in the season eggs are particularly prone to desiccation and being washed or blown from the small plants. Parasitism and predation also reduce survival. *Trichogramma* parasitoids have the potential to reduce egg survival by over 90%. Larval thresholds are also impacted on by beneficial insects. Therefore it is important to assess beneficial insect numbers when making pest control decisions. Fruit retention can also be used to determine whether pests have caused or are at risk of causing economic damage.

Conventional cotton

Helicoverpa spp.

SEEDLING TO FLOWERING	FLOWERING TO CUT-OUT
2 larvae /m or 1 larvae > 8 mm /m	2 larvae /m or 1 larvae > 8 mm /m or 5 brown eggs /m
CUT-OUT TO 15% OPEN BOLLS	**15% TO 40% OPEN BOLLS**
3 larvae /m or 1 larvae > 8 mm /m or 5 brown eggs /m	5 larvae /m or 2 larvae > 8 mm /m or 5 brown eggs /m

Bollgard II cotton

Calculation of spray thresholds in Bollgard II cotton should exclude larvae that are smaller than 3 mm and all eggs. Be sure to objectively assess larval size.

Helicoverpa spp.

SEEDLING TO 40% OPEN BOLLS
2 larvae > 3 mm /m in 2 consecutive checks or 1 larvae > 8 mm /m

Where larvae between 3 mm and 8 mm are observed on Bollgard II cotton, consecutive checks are essential for decision making. *Helicoverpa* spp. must feed in order to ingest the Bt toxin. If the number of 3–8 mm larvae are above threshold on a given check, chances are that a large portion of these will ingest sufficient dose of the toxin and die before the next check.

Using the predator/pest ratio

The predator/pest ratio can be applied in conventional and Bollgard II cotton. The ratio is calculated as:

$$\frac{\text{Total predators*}}{\textit{Helicoverpa}\text{ spp. (eggs + VS + S larvae)}}$$

At least 30 plants or 3 separate metres of row by visual sampling or 20 metres of row by suction sampling is needed in order to use the ratio. The total number of predators must only include the key predator insects (marked with an asterisk in the list below). At least 3 of the key predator species need to be present.

When the predator/pest ratio is 0.5 or higher, the Helicoverpa population should remain below the threshold of 2 larvae/m.

The predator to pest ratio calculated above does not incorporate parasitoids, particularly Trichogramma, in the calculation. To use both predators and parasitoids, the level of egg parasitism should be deducted from the number of Helicoverpa eggs before the predator to pest ratio is calculated. Levels of egg parasitism can vary greatly from farm to farm, region to region and from season to season. Generally levels decline as the season progresses. Notes on how to monitor egg parasitism levels and how to use the predator/pest ratio refer to page 51.

Key beneficial insects

Predators of eggs – red and blue beetle*, damsel bug*, green lacewing larvae*, brown lacewing*, ants, nightstalking spiders.

Predators of larvae – glossy, brown* and predatory shield bugs, big-eyed bug*, damsel bug*, assassin bug*, red and blue beetle*, brown lacewing*, common brown earwig, lynx, tangleweb and jumping spiders.

Predators of pupae – common brown earwig

Predators of moths – orb-weaver spiders and bats

Parasitoids of eggs – *Trichogramma* spp., *Telenomus* spp.

Parasitoids of larvae – *Microplitis demolitor*, orange caterpillar parasite, two-toned caterpillar parasite

Parasitoids of pupae – banded caterpillar parasite

Selecting an insecticide

The insecticide products registered for the control of *Helicoverpa* spp. in cotton are presented in Table 4 on page 13. The use of more selective insecticide options will help to conserve beneficial insects. Refer to Table 3 on pages 8–9.

Be aware of resistance status and follow IRMS (pages 64–65).

***The total number of predators must only include the key beneficial insects marked by a similar.**

Resistance profile

Conventional cotton

Widespread use of Bollgard II cotton has reduced reliance on chemical insecticides. However large plantings of Bollgard II does not change the overall frequencies of resistance genes in the Helicoverpa population and will not influence the rate at which H. armigera will develop resistance to conventional insecticides if significant selection pressure is imposed. While resistance to indoxacarb (Steward), avermectins (Affirm), rynaxypyr (Altacor) and organophosphates (chlorpyrifos) are low, recent testing has identified that frequencies of resistance to Bifenthrin (SP) have increased to 40%. This means that field failures are now likely for this product. Resistance to general pyrethroids has increased to 90%. Therefore the use of conventional chemistries for control of H. armigera in conventional and Bollgard II crops should be used according to the relevant thresholds and the principles of the IRMS applied to all spray decisions (pages 59–67).

Pupae busting is another key tactic for mitigating resistance risk to all insecticides targeting H. armigera, including Bollgard II. Individuals that have survived seasonal selection by insecticides can be controlled before they have a chance to mate, thereby reducing carryover of resistant insects from one season to the next.

Pupae busting should be a priority post-harvest operation on all cotton farms. The IRMS recommends pupae busting as soon as possible after harvest. For Bollgard II crops, follow the pupae busting directions in the products Resistance Management Plan.

OCCASIONAL DETECTION OF RESISTANCE	WIDESPREAD RESISTANCE
Indoxacarb emamectin benzoate chlorpyrifos (OP)	methomyl/thiodicarb (carbamate) (moderate frequency) general pyrethroids (high frequency) bifenthrin (SP) (moderate frequency)
CROSS RESISTANCE	
H.armigera resistance to Bifenthrin has increased. Field failures are likely.	

Bollgard II cotton

A gene is present in field populations of *H. armigera* that has the potential to confer high-level resistance to Cry1Ac. CSIRO and Monsanto data suggests that this gene occurs at a low frequency which is probably less than 5 in 10,000. It is not cross-resistant to Cry2Ab and in certain environments is largely recessive.

A gene that confers high level resistance to Cry2Ab is also present in field populations of *H. armigera*. This gene does not confer cross-resistance to Cry1Ac. In 2012–13 around 1–2% of the *H. armigera* population carried the Cry2Ab resistance gene. The continued efficacy of Bollgard II has become even more dependent on how the industry manages its refuges and implements the other elements of the resistance management plan (RMP). For further details, including information about recent changes in the frequency of Cry2Ab resistance genes in *H. armigera*, refer to the Preamble to the RMP for Bollgard II on page 68.

Over-wintering habit

The cotton bollworm over-winters in cotton fields as diapausing pupae. These pupae are the major carriers of resistance from one season to the next. The initiation of diapause in the pupae is caused by falling temperatures and shortening day lengths. The proportion of pupae entering diapause increases from 0% in late February to +90% in late April – early May, depending on the region. Across all regions (Central Queensland, Macintyre, Namoi and Macquarie Valleys) diapause is initiated in at least 50% of pupae by the first week in April. Diapause termination is based on rising soil temperatures beginning in mid to late September in most regions. Emergence from diapause usually occurs over a 6 to 8 week period in each valley.

Alternative hosts

Spring host crops include; faba beans, chickpeas, safflower, linseed and canola. Pastures and weed flushes also sustain emerging spring populations. Summer host crops include; soybeans, mungbeans, pigeon pea, sunflower, sorghum and maize. The cotton bollworm will attack flowering crops of sorghum and maize preferentially over most other crop hosts.

Further Information:

CSIRO Narrabri
Sharon Downes: (02) 6799 1576 or 0427 480 967.
Colin Tann: (02) 6799 1557 or 0429 991 501.

Qld DAFF, Toowoomba
Melina Miles: (07) 4688 1369.

NSW DPI, Narrabri
Lisa Bird: (02) 6799 2428.

Native budworm
Helicoverpa punctigera

Damage symptoms

Larvae cause early to mid season damage to terminals, buds, flowers and bolls of conventional cotton (non-Bt varieties) in a similar manner to *H. armigera*.

Sampling

Refer to the section on sampling cotton bollworm on the previous page. It is not possible to visually differentiate the eggs or early larval stages of the native budworm from the cotton bollworm, hence it is appropriate that these pests be sampled as one.

Thresholds

Refer to the section on thresholds for cotton bollworm on the previous page. The thresholds for *Helicoverpa* spp. are based on the assumption of potentially mixed populations of cotton bollworm and native budworm.

Key beneficial insects

Refer to the section on Key Beneficial Insects for the cotton bollworm. These predators and parasitoids also attack the native budworm.

Selecting an insecticide

The insecticide products registered for the control of native budworm in cotton in Australia are presented in Table 4 on page 13. The use of more selective insecticide options will help to conserve beneficial insects. Refer to Table 3 on pages 8–9.

Survival strategies

Resistance profile

Conventional cotton

Resistance to insecticides has only rarely been detected in Australia. In conventional cotton, the tendency for the native budworm to occur in mixed populations with the cotton bollworm often limits insecticide control options to those that are also efficacious on the cotton bollworm.

Bollgard II cotton

A gene is present in field populations of *H. punctigera* that has the potential to confer resistance to Cry1Ac. Research suggests that this gene occurs at a low frequency which is probably less than 1 in 1,000. It is not cross-resistant to Cry2Ab and in certain environments is largely recessive.

A gene that confers high level resistance to Cry2Ab is present in field populations of *H. punctigera*. In 2012–13 around 1–1.5% of the *H. punctigera* population carried a Cry2Ab resistance gene.

The continued efficacy of Bollgard II has become even more dependent on how the industry manages its refuges and implements the other elements of the resistance management plan (RMP). For further details, including information about recent changes in the frequency of Cry2Ab resistance genes in *H. punctigera* refer to the Preamble to the RMP for Bollgard II on page 68.

Over-wintering habit

The native budworm has the capacity to over-winter as pupae, but extensive research conducted in the early 1990s found that it is rarely observed to do so in cotton growing areas. However between 20–50% of overwintering pupae collected from numerous crops and fields in cotton regions during 2007 and 2008 were *H. punctigera* suggesting that this strategy may now be more common. If conditions are favourable during winter, sparse but large populations survive and breed on native host plants in inland (central) Australia. As these winter annuals hay-off in spring, large migrations of moths may fly to cotton growing areas in eastern Australia.

Helicoverpa armigera & *punctigera* moths. (Hugh Brier DAFF Qld)

Alternative hosts

The native budworm is not as closely associated with crop hosts as the cotton bollworm. The host range of the native budworm appears to be restricted to dicotyledonous (broad-leaved) hosts. Spring crop hosts include; faba beans, chickpeas, safflower, linseed and canola. Uncultivated hosts, particularly naturalised medics, are important in the initial buildup of the first spring generation. Summer crop hosts include; soybeans, mungbeans, pigeon pea and sunflower.

Further Information:

CSIRO Entomology, Narrabri
Sharon Downes: (02) 6799 1576 or 0427 480 967
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TABLE 4: Control of *Helicoverpa spp.*				
Active ingredient	**Concentration and formulation**	**Application rate of product**	***H. armigera* resistance present**	**Comments**
Cotton bollworm, *Helicoverpa armigera*, and native budworm, *Helicoverpa punctigera*				
Abamectin	18 g/L EC	0.3 or 0.6 L/ha	No	For the control of *Helicoverpa punctigera* only. Use the higher rate alone or the lower rate with a suitable mixing partner. Do not use more than twice in one season – see IRMS.[#]
Alpha-cypermethrin	100 g/L EC 250 g/L SC	0.3, 0.4 or 0.5 L/ha 0.12, 0.16, or 0.2 L/ha	Yes	Use low rate for eggs or newly hatched larvae. Use higher rates for higher egg pressure or larger larvae.
Amitraz	200 g/L EC	2.0 L/ha		Apply as an ovicide with larvicide when eggs or very small larvae are detected. May suppress mites.
Amorphous silica	450 g/L SC	2.5–5.0 L/ha		Apply during egg lay to egg hatch. Best results are obtained from two sequential applications 6–7 days apart.
Bacillus thuringiensis	Btk SC	0.5–4.0 L/ha	No*	Use alone or with mixtures. Refer to relevant label for details. *See RMP preamble page 74
Beta-cyfluthrin	25 g/L EC	0.46–0.8 L/ha	Yes	Can be mixed with mineral spraying oil for ULV applications or with water for EC applications.
Bifenthrin	100 g/L EC 250g/L EC	0.6–0.8 L/ha 0.24–0.32L/ha	Yes	Time spray to coincide with egg hatch. DO NOT apply to larvae >5 cm.[#]
Cyfluthrin	50 g/L EC	0.6 L./ha or 0.8 L/ha	Yes	Application should be timed to coincide with egg hatch.
Cypermethrin	200 g/L EC 250 g/L EC 260 g/L EC	.3–0.70 L/ha 0.3–0.5 L/ha 0.29–0.48 L/ha	Yes	See label for higher rate situations.
Deltamethrin	5.5 g/L ULV 27.5 g/L EC	2.5–3.5 L/ha 0.5–0.7 L/ha	Yes	Use low rate as ovicide and high rates for small to medium larvae.[#]
Emamectin benzoate	17 g/L EC	0.55–0.7 L/ha	No	Apply at or just prior to hatching. Use non-ionic surfactant as per label.[#]
Esfenvalerate	50 g/L EC	0.5–0.7 L/ha	Yes	Use low rate when larvae are small and pressure is low.[#]
Gamma -cyhalothrin	150 g/L CS	0.05 or 0.06, 0.07 L/ha	Yes	Ovicidal rate. Apply higher rate when egg lay is heavy and/or *H.punctigera* >10mm and/or *H.armigera* <5mm.[#]
Indoxacarb	150 g/L EC	0.65 or 0.85 L/ha	No	Refer to label for rate selection criteria. Compatible with amitraz.#
Lambda-cyhalothrin	250 g/L ME	0.06, 0.07 or 0.085 L/ha	Yes	Ovicidal rate. Use low rate for newly hatched larvae.[#]
Helicoverpa NPV	2000 M-Obs/mL LC 5x109 M-Obs/mL LC	0.5 L/ha 0.5 L/ha		Alone or with compatible larvicide. See label for details. Target application to coincide with egg hatching.
Magnet		0.5L/100 m row (10–50 cm bands) in 72 m or 36 m		Use including insecticides as per label instructions
Methomyl	225 g/L SL	0.5–1.0 L/ha 1.8–2.4 L/ha	Yes	Ovicidal rate. Larvicidal rate. Higher rate of larvicidal rate may cause reddening of foliage, if excessive use an alternative. Do not apply during periods of plant stress.[#]
Paraffinic oil	792 g/L	2% or 2L/100L of water		Use a minimum of 80L/ha of water. Apply only by ground rig before crop closure.
Piperonyl butoxide	800 g/L EC	0.3–0.4 L/ha		Use as a synergist when applying synthetic pyrethroids. See label.
Chlorantraniliprole	350 g/kg WDG	0.090 or 0.150 g/ha + non ionic surfactant @ 125 gai/100 L	No	Target brown eggs or hatchling to 2nd instar larvae before they become entrenched in squares, flowers and bolls. Use high rate where the potential is for >3.5 larvae/m and to achieve longer residual control.
Thiodicarb	375 g/L SC 800 g/L WG	0.5–1.0L/ha + Larvacide 2.0-2.5L/ha 0.235–0.470 kg/ha + Larvacide 0.940–1.2kg/ha	Yes	This product has ovicidal and larvicidal activity. See label for details.[#]

#See label for instructions to minimise impact on bees.