



Effective resistance management for the neonicotinoids:

**Industry's approach to ensure
the continued efficacy of a key
insecticide class**

IRAC Symposium on Insecticide Sustainability: Neonicotinoids
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Outline

Why effective Resistance Management is essential

What industry is doing:

- Company level
 - Resistance Management Strategy for a Neonicotinoid – A typical responsible company approach to IRM
- Inter-company level
 - Collaborative Industrial Approach to IRM for Neonicotinoids
 - An industry partnership to ensure sustainability
- All-industry level
 - Role of Insecticide Resistance Action Committee (IRAC)



Why Effective Resistance Management is essential

- Sustaining effective commercial life of current insecticides requires intelligent use of presently available compounds
 - Insecticide Resistance Management (IRM)
- For any crop / pest situation, effective IRM requires the availability of a broad range of modes of action
- IRM is made much more difficult by loss of modes of action through resistance development caused by misuse or overuse of insecticides
- We can no longer rely on having a steady stream of new modes of action to circumvent resistance problems.....



Protecting current AIs from resistance is crucial !

- Finding and developing new insecticides is extremely costly & difficult
- We cannot accept losing them to resistance!

Cost of developing and registering a new chemical crop protection AI in year 2000 (\$M)

[Source: ECPA, 2003]

Category	Sub-category	Costs	Total Cost
Research	Chemistry	41	94
	Biology	44	
	Toxicology / Environmental Chemistry	9	
Development	Chemistry	20	79
	Field Trials	25	
	Toxicology	18	
	Environmental Chemistry	16	
Registration			11
Total			184

What industry is doing at a company level



Resistance Management Strategy for a Neonicotinoid

Responsible companies have clear guidelines to manage resistance to Neonicotinoids:

'To prevent the development and spread of resistance, applications of Thiamethoxam and other Neonicotinoid insecticides should be applied in 'block' applications where:

The total duration of the neo-nicotinoid 'blocks' does not exceed more than half of the crop cycle (vegetative period). The duration of a block application should be based on either:

- The generation time of the target pest (Where possible each block should treat a single generation of the target pest)
- The period of insect control that is provided by a single application of the insecticide (Applicable to insects with a short generation time).

Wherever possible it is recommended that the application of an insecticide 'block' should not immediately be followed by an application of an insecticide 'block' of the same chemical class.

Example of Whitefly RM Strategy

Transplanting

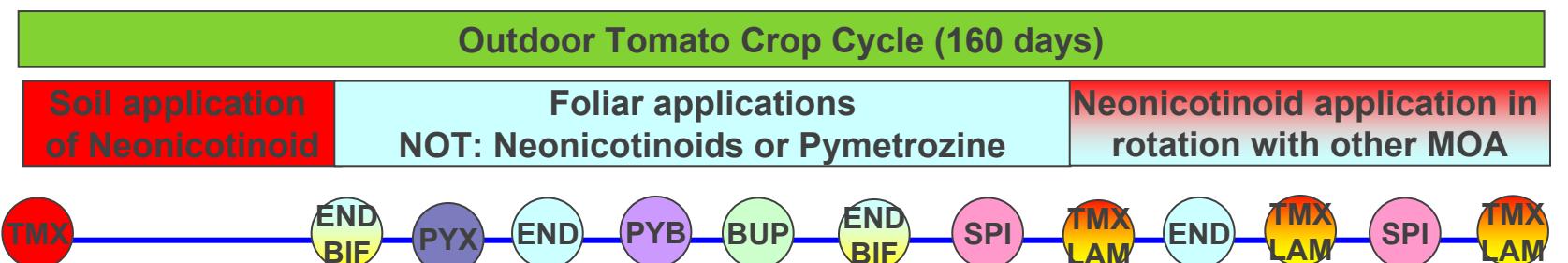
- Soil application of Neonicotinoid provides good control for an extended period - with limited selection for resistance

First foliar applications made after soil application

- Avoid applications of Neonicotinoids or Pymetrozine in this period
- Rotate other available AIs

Foliar applications at end of crop cycle

- Rotate available insecticides with different MoA including Neonicotinoids and Pymetrozine.
- Avoid consecutive applications of same chemical class
- Total number of Neonicotinoid and Pymetrozine applications should be < half crop cycle (based on 15d duration for foliar and 30day for soil)

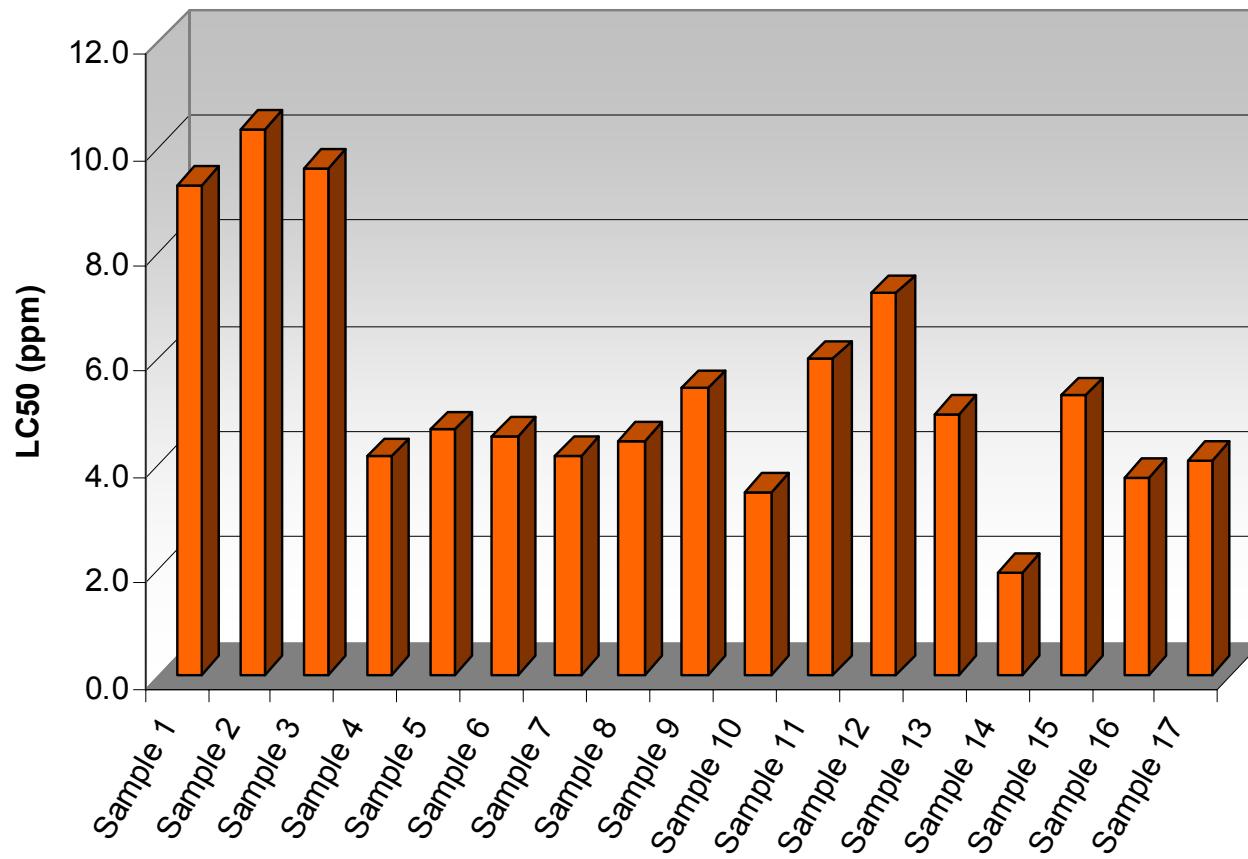


Baselines and Resistance Monitoring

- Establishment of clear baselines followed by appropriate susceptibility monitoring ensures we know current status of susceptibility to key insecticides
 - **Timely awareness of any emerging problems**
- As indicated in Ralf Nauen's presentation each company establishes baselines for its products e.g. :
 - **Thiamethoxam against Soybean Aphid**
 - **Emamectin benzoate against Grape Berry Moth**
- And undertakes regular susceptibility monitoring for key at-risk pests e.g. :
 - **Cotton Aphid, Peach Potato Aphid, Glasshouse Whitefly, etc**
 - **Diamondback Moth, Fall Armyworm, Cotton Bollworm, etc**
 - **Colorado Potato Beetle, Pollen Beetle, Spider Mites, etc**

Example of Susceptibility Monitoring

- Adult Western Corn Rootworm ‘baseline’ susceptibility to thiamethoxam
- Independent samples collected from USA Corn Belt.



Sponsorship of Resistance Research

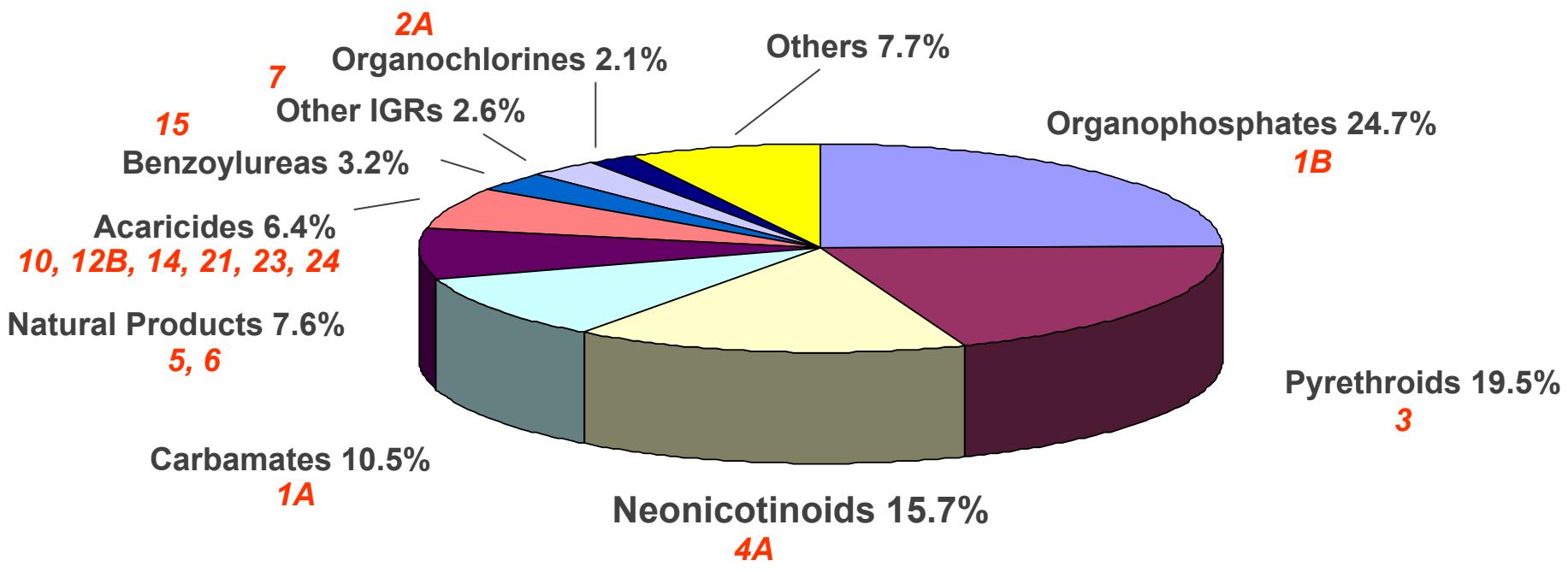
- Sponsorship of various research projects on resistance to Neonicotinoids including:
- ‘Characterisation and management of Neonicotinoid resistance in *Bemisia tabaci*’
- At Rothamsted Research, UK (Ian Denholm, Kevin Gorman)



What industry is doing at an inter-company level



Global Insecticide sales



Total Global Insecticide Sales 2003 = \$6,650 million

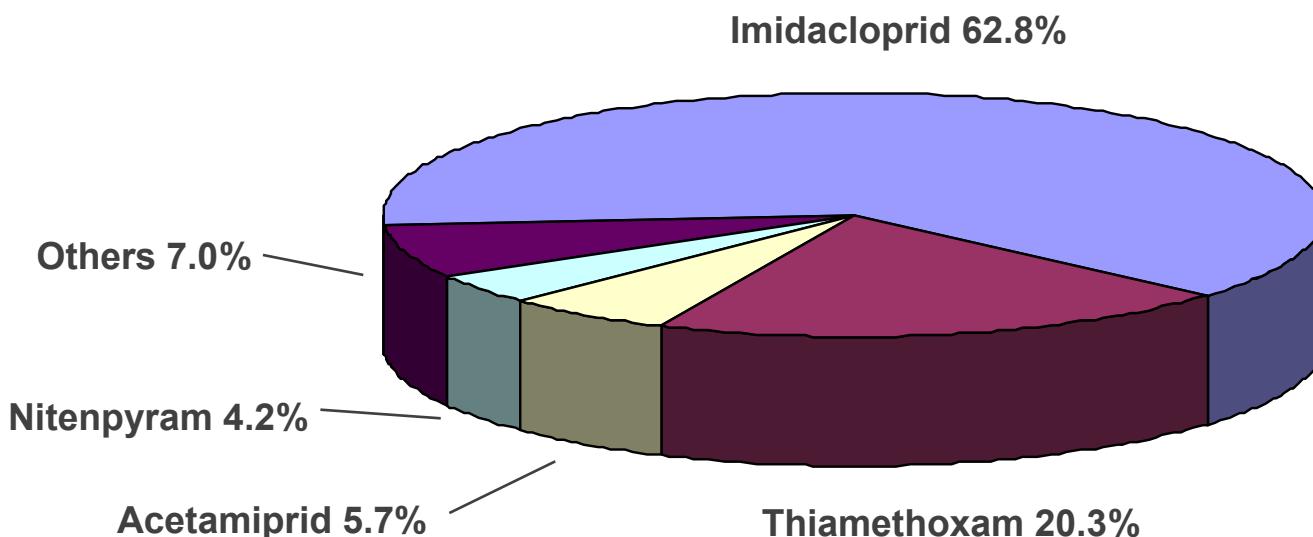
[Source Phillips McDougall, November 2004]

[Natural products includes avermectins and spinosyns]

IRAC Mode of Action Classification Number

Neonicotinoids now 3rd largest group of insecticides based on global sales

Global Neonicotinoid sales



Global Neonicotinoid Sales 2003 = \$1,059 million

[Source Phillips McDougall, November 2004]

IRAC Mode of Action Classification Number 4A

Imidacloprid and Thiamethoxam account for >80% of market

Bayer CropScience and Syngenta – Collaborative Neonicotinoid Stewardship Project

- Together, sales of Bayer CropScience and Syngenta Neonicotinoid products total c. \$930M – 88% of Neonicotinoids
 - Bayer CropScience: Imidacloprid, Thiacloprid, Clothianidin
 - Syngenta: Thiamethoxam
- Two companies uniquely placed to take a lead in developing and delivering coherent and effective IRM strategies at both a global and local level
- March 2005: Bayer CropScience and Syngenta agreed a joint global stewardship project to manage Neonicotinoids, focussing on key at-risk pests
- Other [Japanese] companies invited to join – but declined



Bayer CropScience / Syngenta – Collaborative Neonicotinoid Stewardship Project

- Aim: To intensify the implementation of IRM by working together through contacts at country level
- Proposal to develop local IRM recommendations based on the two company global IRM recommendations for Neonicotinoids
 - Syngenta: Maximal exposure to Neonicotinoids is limited to 50% of the cropping cycle
 - Bayer CropScience: Neonicotinoids are limited to maximum of 3 applications per pest species and crop cycle
- Although slightly different, in practice expected in most situations that time periods covered by neonicotinoids are similar for both company recommendations
- Agreed that local guidelines can be stricter than the global guidelines, but should not be more flexible

Bayer CropScience and Syngenta - Collaborative Neonicotinoid Stewardship Project – Targets

Agreed first list of key at-risk targets for co-operation:

Country	Target	Crop
Mexico	Whitefly	Vegetables
Guatemala	Whitefly	Melons & Tomatoes
India	Brown planthopper	Rice
Brazil	Whitefly	Beans
Morocco	Whitefly	Covered vegetables
Italy	Whitefly	Covered vegetables
Turkey	Whitefly	Covered vegetables
Japan	Thrips	Vegetables

Bayer CropScience and Syngenta - Collaborative Neonicotinoid Stewardship Project

Key actions:

- Establish local contacts between nominated persons from both companies and set up joint meetings
- Establish agreed local IRM Neonicotinoid strategies taking into account the general guidelines of both companies
 - Agree and adopt positioning in high risk crops
 - Adapt for local conditions, positioning and availability of products
- Involve local regulatory authorities and encourage them to take ownership
- Involve key local academic groups and influencers
- Involve local IRAC groups
- Approach other local neonicotinoid companies
- Check possibility of implementing a labelling scheme (similar to US)



BCS and Syngenta sponsor UK Neonicotinoid LINK project

- **Bayer CropScience and Syngenta are jointly sponsoring [with others] :**

UK Department of Environment, Food and Rural Affairs, SA-LINK Project: Sustainability of Neonicotinoid Insecticides

- **To assess and manage increasing risk of resistance to Neonicotinoids in *Myzus persicae* (and other aphids)**
- **Risk arises from rapid recent increase in uses of neonicotinoids in multiple host crops for *M. persicae***
- **Co-ordinating laboratory Rothamsted Research (Dr Ian Denholm et al)**

What industry is doing at an all-industry level



Insecticide Resistance Action Committee - IRAC

- IRAC formed in 1984 to provide a co-ordinated industry response to the development of resistance in insect and mite pests
- Currently 6 IRAC International members:

BASF

DuPont

FMC

Bayer CropScience

Dow AgroSciences

Syngenta

IRAC Objectives:

- Facilitate communication and education on insecticide and acaricide resistance
- Promote the development of resistance management strategies in crop protection and vector control to maintain efficacy and support sustainable agriculture and improved public health

IRAC, US - Members

- AMVAC**
- Arysta LifeScience**
- BASF**
- Bayer CropScience**
- Cerexagri**
- Chemtura**
- Dow AgroSciences**
- DuPont**
- FMC**
- Mitsui**
- Monsanto**
- Nisso America**
- Syngenta**
- Valent**

www.irac-online.org

IRAC Country Groups

- IRAC Country groups deal with key resistance issues at local level – supported by IRAC International
 - Multi-company cooperation and coordinated IRM
 - IRAC Country groups often include additional companies not involved in IRAC International
 - Country groups may include or involve others from Academia and Research Institutes and Regulatory bodies (IRAG groups)
- Current IRAC Country groups:
 - IRAC Australia (AIRMG)
 - IRAC Brazil
 - IRAC India
 - IRAC South Africa
 - IRAC Spain
 - IRAC US
 - IRAC China (proposed)



IRAC Mode of Action Classification



Insecticide Resistance Action Committee
www.irac-online.org

IRAC Mode of Action Classification

Fully revised & re-issued, September 2005

Version: 5.1

The IRAC Mode of Action (MoA) classification provides farmers, growers, advisors, extension staff, consultants and crop protection professionals with a guide to the selection of insecticides or acaricides for use in an effective and sustainable insecticide or acaricide resistance management (IRM) strategy. In addition to presenting the MoA classification, this document outlines the background to, and purposes of, the classification list and provides guidance on how it is used for IRM purposes. The list is reviewed and re-issued at intervals as required.

What is resistance

Resistance to insecticides may be defined as '*a heritable change in the sensitivity of a pest population that is reflected in the repeated failure of a product to achieve the expected level of control when used according to the label recommendation for that pest species*' (IRAC). This definition differs slightly from others in the literature, but IRAC believes it represents the most accurate, practical definition of relevance to farmers and growers. Resistance arises through the over-use or mis-use of an insecticide or acaricide against a pest species and results in the selection of resistant forms of the pest and the consequent evolution of populations that are resistant to that insecticide or acaricide.

MoA, Target-site resistance and Cross-resistance

In the majority of cases, not only does resistance render the selecting compound ineffective but it often also confers cross-resistance to other chemically related compounds. This is

- **Definitive scheme developed and endorsed by IRAC in consultation with key researchers**
- **Worldwide distribution**
- **All current insecticides allocated to a Mode of Action group or sub-group**
- **A key tool for selection of insecticides in effective IRM programs**
- **Updated as required**
- **Latest version Sept 2005**

IRAC Mode of Action Classification

IRAC Mode of Action Classification			Version: 5.1
IRAC Mode of Action Classification v5, September 2005 1			
Main Group and Primary Site of Action	Chemical Sub-group or exemplifying Active Ingredient	Active Ingredients	
1 Acetylcholine esterase inhibitors	1A Carbamates	Adicarb, Alanycarb, Bendiocarb, Benfuracarb, Butocarboxim, Butoxycarboxim, Carbaryl, Carbutoran, Carbosulfan, Ethofencarb, Fenobucarb, Formetanate, Furathiocarb, Isopropcarb, Methiocarb, Methomyl, Metolcarb, Oxamyl, Pimicarb, Propoxur, Thiodicarb, Thiofanox, Trimethacarb, XMC, Xylylcarb	
	Triazemate	Triazemate	
	1B Organophosphates	Acephate, Acamiphos, Acinphos-ethyl, Acinphos-methyl, Cadusafos, Chlorethoxyfos, Chlorfenvinphos, Chlormephos, Chlorynphos, Chlorynphos-methyl, Coumaphos, Cyanophos, Demeton-S-methyl, Diazinon, Disulfoton/ DDVP, Disotophos, Dimethoate, Dimethylvinphos, Disulfoton, EPN, Ethon, Ethoprophos, Famphur, Fenamiphos, Fenitrothion, Fenthion, Fosthiazate, Heptenophos, Isotenphos, Isopropyl O-methoxyaminothiophosphonyl) salicylate, Isoxathion, Malathion, Mecarban, Methamiphos, Methidation, Mevinphos, Monocrotophos, Naled, Omethoate, Oxydemeton-methyl, Parathion-methyl, Phenothate, Phorate, Phosalone, Phosmet, Phos�anidon, Phoxim, Primiphos-ethyl, Profenofos, Propetamphos, Prothifos, Pyraclofos, Pyridaphenthion, Quinalphos, Sulfotep, Tebupirimfos, Temephos, Terbufos, Tetraclorvinphos, Thiometon, Triazophos, Trichlorfon, Vamidothion	
2 GABA-gated chloride channel antagonists	2A Cyclodiene organochlorines	Chlordane, Endosulfan, gamma-HCH (Lindane)	
	2B Phenylpyrazoles (Fiproles)	Ethiprole, Fipronil	
3 Sodium channel modulators	DDT	DDT	
	Methoxychlor	Methoxychlor	
	Pyrethroids	Aerinathrin, Allethrin, d-cis-trans Allethrin, dtrans Allethrin,	

Example pages

Main Group and Primary Site of Action	Chemical Sub-group or exemplifying Active Ingredient	Active Ingredients
4 Insect Acetylcholine receptor agonists / agonists	4A Neonicotinoids	Acetamiprid, Clothianidin, Dinotefuran, Imidacloprid, Nititenpyram, Thiacloprid, Thiamethoxam
	4B Nicotine	Nicotine
	4C Bensultap	Bensultap
5 Insect Acetylcholine receptor agonists (not group 4)	Cartap hydrochloride	Cartap hydrochloride
	Nereistoxin analogues	Thiocyclam, Thiosultap-sodium
	Spinosyns	Spinosad
6 Nerve channel blockers		
	Avermectins, Milbemycins	Abamectin, Enamectin benzoate, Milbemectin
7 Juvenile hormone mimics		
	7A Juvenile hormone analogues	Hydroprene, Kinoprene, Methoprene
	7B Fenoxycarb	Fenoxycarb
8 Compounds of unknown or non-specific mode of action (fumigants)	7C Pyriproxyfen	Pyriproxyfen
	8A Alkyl halides	Methyl bromide and other alkyl halides
	8B Chlormecidin	

Using the IRAC MoA Classification

- The IRAC Mode of action scheme is central to developing effective IRM strategies
- Sequences, rotations or alternations of different MoA groups help prevent or delay resistance, or deal with existing resistance problems
- Modify locally to take account of known metabolic mechanisms conferring cross-resistance between MoA groups or insect populations with multiple resistances
- In the absence of any information, intelligent sequences of MoA groups will always reduce selection pressures and help prevent or delay resistance, and help regain susceptibility
- IRAC strongly supports MoA labeling schemes – e.g. US, Australia
 - And IRAC campaigns for wider use of such schemes
- Use of symbols and colors for MoA groups can help e.g. Brazil

IRAC MoA Posters and Wallchart

IRAC Insecticide Resistance Action Committee

Insecticide Mode of Action Classification:
A key to effective insecticide resistance management

CropLife INTERNATIONAL

IRAC website: www.irac-online.org

Introduction
IRAC promotes the use of a Mode of Action (MoA) classification of insecticides as the basis for effective and sustainable insecticide resistance management (IRM). Insecticides are grouped into specific groups based on their target site. Reviewed and revised periodically, the IRAC MoA classification list provides farmers, growers, advisors, extension staff, consultants and crop protection professionals with a guide to the selection of insecticides or arboricides in IRM programs. Effective IRM of this type preserves the utility and diversity of available insecticides and arboricides. A selection of MoA groups is shown below.

Moultion & Metamorphosis
Group 18: Enzyme agonist / disruptor
Deltamethrin (e.g. Tefuthrotoide)
Group 7: Juvenile hormone mimics
JH analogues, Fenprocarb, Pyriproxyfen, etc.

Midgut
Group 77: Microbial disruptors of insect midgut membranes
Toxin produced by the bacterium *Serratia chrysomelae* (St); Et
Spartanidin & Organon metabolite I
Group 12: Juvenile hormone mimics in the midgut (specific cuticular tannins sub-group)

Nervous System
Groups 10 & 15: Organophosphate (AChE) inhibitors
Carbamates and Organonothiocarb
Group 1: Acetylcholinesterase inhibitors
Group 2: GABA-gated chloride channel antagonists
Group 3: Cholinergic receptor agonists
Group 4: Nicotinic Acetylcholine Receptor Agonists (not A)
Group 5: Chloride Channel activators

Metabolic Processes
Many group 1 acting on a wide range of metabolic processes including:
• Disruption of oxidase phosphorylation, disruption of ATP
Deltamethrin & Organon metabolite I
Group 12: Urethane's of oxidase phosphorylation via disruption of H₊ proton gradient - Chlorthalide

Cuticle Synthesis
Groups 15: Allomimic inhibitors or other inhibitors of cuticle synthesis
Benzothiuron (Lepidoptera and others), Buprofezin (Homoptera), etc.

Use MoA of Action wisely for good IRM!

Insecticide Mode of Action Classification:
A Key to Effective Insecticide Resistance Management in Whiteflies

Introduction
IRAC has developed a Mode of Action (MoA) classification for whiteflies. It promotes the use of the MoA basis for insecticide resistance management (IRM) in whiteflies. The IRAC MoA classification is a key to effective insecticide resistance management in whiteflies. It provides sustainable and effective IRM. This ensures that selection from compounds in the same MoA group is minimized. Applications are often arranged into MoA spray windows or blocks that are defined by the stage of crop development and the biology of the pest(s) of concern. Local expert advice should always be followed with regard to spray windows and timings. Several sprays of a compound may be possible within each spray window but it is generally essential to ensure that successive generations of the pest are not treated with compounds from the same MoA group. Metabolic resistance mechanisms may give cross-resistance between MoA groups, and where this is known to occur, the above advice must be modified accordingly.

Effective IRM strategies: Alterations or sequences of MoA
Minimize the selection for resistance from any one type of insecticide or arboricide. In practice, alternative sequences or combinations of different MoA groups provide sustainable and effective IRM. This ensures that selection from compounds in the same MoA group is minimized. Applications are often arranged into MoA spray windows or blocks that are defined by the stage of crop development and the biology of the pest(s) of concern. Local expert advice should always be followed with regard to spray windows and timings. Several sprays of a compound may be possible within each spray window but it is generally essential to ensure that successive generations of the pest are not treated with compounds from the same MoA group. Metabolic resistance mechanisms may give cross-resistance between MoA groups, and where this is known to occur, the above advice must be modified accordingly.

Non-specific MoA
Group 9: Compounds of non-specific mode of action

Insecticides acting on the nervous system
The nervous system is the target for many current insecticides, but within this system so many target sites, insecticides with specific mode of action and their targets:

- Acetylcholinesterases (AChE) inhibitors:** Carbamate (C) and Organonothiocarb (ON) act as inhibitor of AChE nerve synapses. This results in hyperactivity in the nervous system
- Group 2: GABA-gated chloride channel antagonists:** Cycloheximide (CH) and Organonothiocarb (ON) act as inhibitor of GABA-gated chloride channel. When CH appears in the nervous system, it blocks the GABA-gated chloride channel, which causes hyperactivity in the nervous system. When ON appears in the nervous system, it blocks the GABA-gated chloride channel, which causes hyperactivity in the nervous system.
- Group 3: Sodium channel modulators:** Sodium channels are involved in the propagation of action potential. Cycloheximide (CH) and Organonothiocarb (ON) act as sodium channel modulator. When CH appears in the nervous system, it blocks the sodium channel, which causes hyperactivity and nerve block.
- Group 4: Acetylcholine receptor agonists:** The receptor activity (RA) act as agonist of acetylcholine at the post-synaptic membrane. Organonothiocarb (ON) has the receptor activity (RA) and when ON appears in the nervous system, it stimulates the receptor and hyperactivity.

Insecticides desribing cuticle synthesis (Type 1)
The cuticle is a protective layer on the insect's body. It is made up of the lipid cycle and insectocerine which interact with this process to help the molting cycle lead to death of the insect.

- Group 5: Components of unknown action / Pyrethrins:** Pyrethrins (P) has a non-selective mode of action. It appears to involve a selective inhibition of the nervous system. These compounds inhibit cholinesterases in a number of insects including whiteflies.
- Insecticides acting as feeding blockers:**
Group 9: Components of unknown action / Pyrethrins: Pyrethrins (P) has a non-selective mode of action. It appears to involve a selective inhibition of the nervous system. These compounds inhibit cholinesterases in a number of insects including whiteflies.

Insecticide classes for whitefly control
IRAC lists 20 modes of action groups (including sub-groups), 10 of them are currently used in whitefly control.

MoA	MoA Class	Group	Sub-group	Example
1: Carbamate herbicides	1: Carbamate	1:1		Organochlorines
2: GABA-gated chloride channel antagonists	2: GABA-gated chloride channel	2:1		Pyrethrins
3: Sodium channel modulators	3: Sodium channel	3:1		Organonothiocarb
4: Acetylcholine receptor agonists	4: Acetylcholine receptor agonists	4:1		Organonothiocarb
5: Components of unknown action / Pyrethrins	5: Components of unknown action / Pyrethrins	5:1		Pyrethrins
6: Carbamate insecticides	6: Carbamate	6:1		Organochlorines
7: Organonothiocarb herbicides	7: Organonothiocarb	7:1		Organonothiocarb
8: Organonothiocarb insecticides	8: Organonothiocarb	8:1		Organonothiocarb
9: Components of unknown action / Pyrethrins	9: Components of unknown action / Pyrethrins	9:1		Pyrethrins
10: Organonothiocarb herbicides	10: Organonothiocarb	10:1		Organonothiocarb
11: Organonothiocarb insecticides	11: Organonothiocarb	11:1		Organonothiocarb
12: Organonothiocarb insecticides	12: Organonothiocarb	12:1		Organonothiocarb
13: Organonothiocarb insecticides	13: Organonothiocarb	13:1		Organonothiocarb
14: Organonothiocarb insecticides	14: Organonothiocarb	14:1		Organonothiocarb
15: Organonothiocarb insecticides	15: Organonothiocarb	15:1		Organonothiocarb
16: Inhibitors of chitin biosynthesis	16: Chitin	16:1		Organonothiocarb
17: Moulting disruptor	17: Moulting	17:1		Organonothiocarb
18: Organonothiocarb insecticides	18: Organonothiocarb	18:1		Organonothiocarb
19: Organonothiocarb insecticides	19: Organonothiocarb	19:1		Organonothiocarb
20: Organonothiocarb insecticides	20: Organonothiocarb	20:1		Organonothiocarb
21: Coupling info / electron transport inhibitors	21: Coupling info / electron transport inhibitors	21:1		Organonothiocarb
22: Voltage Dependent Sodium Channel blockers	22: VDS	22:1		Organonothiocarb
23: Inhibitors of lipid in the waxy layer	23: Waxy	23:1		Organonothiocarb
24: Mitochondrial complex IV electron transport inhibitors	24: Mitochondrial	24:1		Organonothiocarb
25: Neuronal inhibitors (unknown mode of action)	25: Neuronal	25:1		Organonothiocarb
26: Acetylcholinesterase inhibitors	26: AChE	26:1		Organonothiocarb
27: P450 monooxygenase inhibitors	27: P450	27:1		Organonothiocarb
28: Benzene ring hydroxylases	28: Benzene ring hydroxylases	28:1		Organonothiocarb
29: Synergists	29: Synergists	29:1		Organonothiocarb
30: Miscellaneous non-specific (multi-site) inhibitors	30: Miscellaneous	30:1		Organonothiocarb

Conference Posters

IRAC Mode of Action Wallchart: MoA Classification Groups & Structures

A1 printed versions available

More information on the IRAC® Resistance Action Committee and the Mode of Action Classification is available from www.irac-online.org or from Alan Porter email: porter@spin.org

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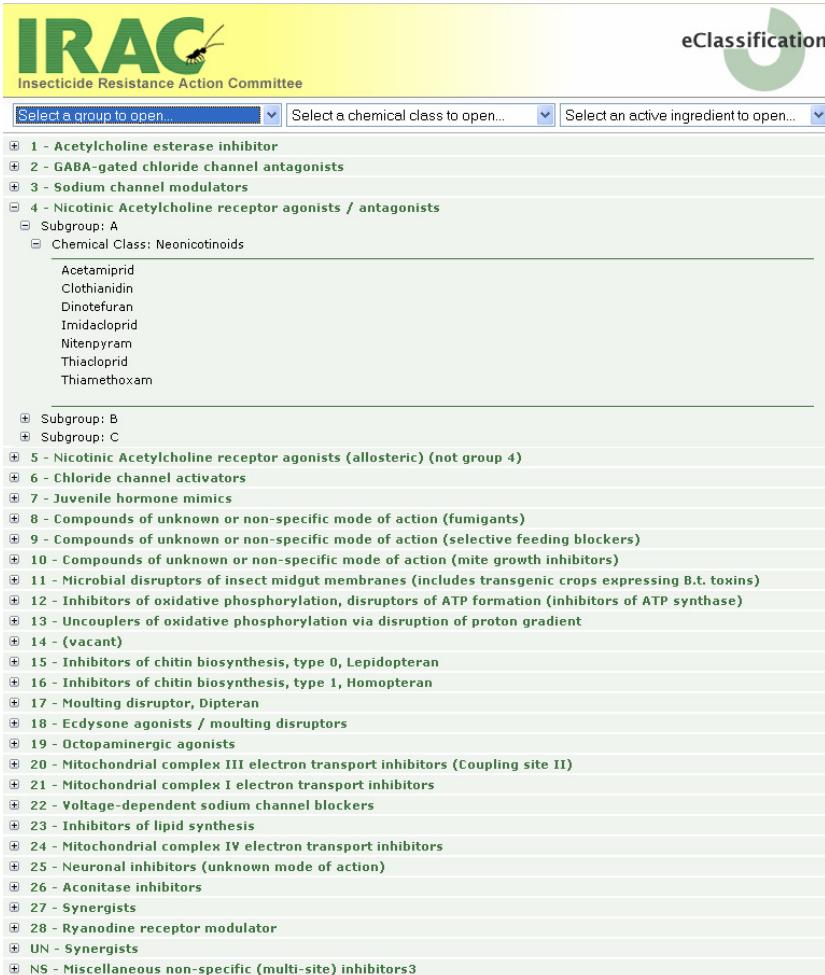
* Only major representatives of the groups are shown

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syngenta

eClassification – interactive MoA

New interactive online tool www.irac-online.org

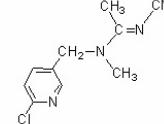


Select a group to open...	Select a chemical class to open...	Select an active ingredient to open...
(vacant)	Select a chemical class to open...	Select an active ingredient to open...
Acetylcholine esterase inhibitor	Avermectins, Milbemycins	Abamectin
Aconitase inhibitors	Benzoylureas	Acephate
Chloride channel activators	Carbamates	Acequinocyl
Compounds of unknown or non-specific	Cyclodiene organochlorines	Acetamiprid
Compounds of unknown or non-specific	Esterase inhibitors	Acrinathrin
Compounds of unknown or non-specific	Fipronil (or Phenylpyrazoles)	Alencyarb
Ecdysone agonists / moult disruptor	Juvenile hormone analogues	Aldicarb
GABA-gated chloride channel antagonist	METI acaricides, Rotenone	Allethrin
Inhibitors of chitin biosynthesis, type 0, I	Methyl bromide	Aluminum phosphide
Inhibitors of chitin biosynthesis, type 1, I	Neonicotinoids	Aminocarb
Inhibitors of lipid synthesis	Organophosphates	Amitraz
Inhibitors of oxidative phosphorylation,	Organotin miticides	Azadirachtin
Juvenile hormone mimics	P450 monooxygenase inhibitors	Azamethiphos
Microbial disruptors of insect midgut me	Pyrethroids	Azinphos-ethyl
Miscellaneous non-specific (multi-site)	Tetranic acid derivatives	Azinphos-methyl
Mitochondrial complex I electron trans		Azocyclotin
Mitochondrial complex III electron trans		B.t. var. aizawai
Mitochondrial complex IV electron trans		B.t. var. israelensis
Moult disruptor, Dipteron		B.t. var. kurstaki
Neuronal inhibitors (unknown mode of a		B.t. var. sphæericus
Nicotinic Acetylcholine receptor agonis		B.t. var. tenebrionicus
Nicotinic Acetylcholine receptor agonis		Bendiocarb
Octopaminergic agonists		Benfurcarb
Ryanodine receptor modulator		Bensulfate
Sodium channel modulators		Benzoximate
Synergists		Bifenazate
Synergists		Bifenthrin
Uncouplers of oxidative phosphorylatic		Bioallethrin
Voltage-dependent sodium channel blo		Bioallethrin S-cyclopentenyl

Drop down menus

>>options

Cas No	135410-20-7
Common Name	Acetamiprid
Chemical Class	Neonicotinoids
Primary Site of Action	Nicotinic Acetylcholine receptor agonists / antagonists
MOA Group Number	4A
Relevant Pests	Control of Hemiptera, especially aphids, Thysanoptera and Lepidoptera
Relevant Crops	Wide range of crops, especially vegetables, fruit and tea
Use Patterns	Soil or foliar
Comments	



IRAC Website – www.irac-online.org

IRAC
Insecticide Resistance Action Committee

Resistance Management for Sustainable Agriculture & improved Public Health

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About IRAC Resources IRAC International Country Groups Growers

Resistance Management from IRAC

The Insecticide Resistance Action Committee (IRAC) is an inter company group formed in 1984 to provide insecticide and acaricide resistance management strategies to help reduce the development of resistance in insect and mite pests. IRAC believes that Resistance Management should be an integral part of Integrated Pest Management and provides for Sustainable Agriculture and Improved Public Health.

We would encourage you to further explore the site using the drop-down menus at the top of the page or via the other links. Further background on IRAC, its Constitution and Mission along with the IRAC newsletter eConnection can be located under the About IRAC heading. The majority of the IRAC documents can be located under Resources. This includes Mode of Action Classification, Test Methods, Resistance Management Guidelines, Posters and Publications along with other Educational and Regulatory Material. The headings, IRAC International and IRAC Country Groups, provide information on these different Teams some of which is only available to Team Members. The final drop-down menu provides Grower information in the form of links to Publications, Associations and Magazines. We hope you find the site a useful resource.

Links to New Resistance Management Information

- [General Principles of Insecticide RM from IRAC](#)
- [IRAC Test Methods - Overview and Summary](#)
- [IRAC Paper and Poster on MOA at ICE Meeting Brisbane, August 2004](#)
- [Neonicotinoid IRM Guidelines, September 2004](#)
- [New IRAC Logo and Branding material - September 2004](#)
- [EWSN Meeting, Croatia 5-9th October 2004 - Whitemfly MOA Poster](#)
- [IRAC eConnection - Issues 5, December 2004](#)
- [IRAC Intl. Conference Call - Action Log, January 2005 \(Members only\)](#)
- [Website Traffic and Activity Report March 2005](#)
- [Latest IRAC eConnection - Issues 6, April 2005](#)
- [IRAC US Meeting Minutes, April 2005 \(Members only\)](#)
- [IRAC Intl. Spring Meeting Minutes, Florence, April 2005 \(Members only\)](#)
- [Work on resistance in Italy on Codling Moth and the new Italian group GIRIF](#)
- [JUST RELEASED - New IRAC MoA Classification Scheme \(ver 4.2, May 2005\)](#)
- [IRAC Website - Update and Review, May 2005](#)

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CropLife INTERNATIONAL

- **IRAC's key communication vehicle**
- **Av. 150 hits, 30 visits a day (09.05)**
- **Accessed by over 80 countries**
- **Search terms - #1 'IRAC', #2 'Insecticide Resistance'**
- **190 pages including public, protected & utility pages**
- **160 docs. & 92 image files**
- **Information on IRAC, Mode of Action, advice on IRM**
- **Education modules**
- **Resources incl. key papers, documents, posters**
- **Links for growers**
- **IRAC Country group information**

IRAC Susceptibility Test Methods



Insecticide Resistance Action Committee
www.irac-online.org
Resistance Management for Sustainable Agriculture and Improved Public Health

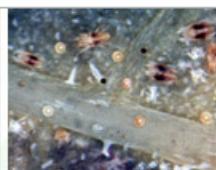
IRAC Susceptibility Test Methods Series

Version: 2

Method No: 3

Details:

Method:	No: 3
Status:	Approved
Species:	<i>Panonychus ulmi</i> <i>Tetranychus cinnabarinus</i>
Species Stage	<i>P. ulmi</i> (summer eggs) <i>T. cinnabarinus</i> (eggs)
Product Class:	clofentezine hexythiazox tetradifon
Comments:	None



Photograph Courtesy of Whitney Cranshaw, Colorado State University, www.csu.edu

Description:

Materials:

Petri dishes (9-cm diameter), filter paper to fit Petri dishes, cotton wool, *untreated* apple or plum leaves, small scissors, small forceps, fine pointed brush or cocktail stick, beakers or glass jars (ca. 100-ml capacity) for test liquids, 1-ml disposable plastic syringes for liquids for weighing balance for solids, hand lens (minimum 10 x) or binocular microscope, maximum-minimum thermometer.

Methods:

- Cut square sections about 1.5 x 1.5 cm from chemically untreated apple or plum leaves. Use young leaves, but not before they are fully expanded. Leaves must be in good condition. Use a minimum of four replicates (leaf sections) per treatment.
- Place these sections, upper surface uppermost, on a sheet of moist filter paper on moist cotton wool in open Petri dishes.
- Collect apple leaves with adult mites, and with the fine pointed brush or cocktail stick transfer 10 – 15 females onto each leaf section. Maintain at a minimum temperature of 20°C, minimum photoperiod 16 h and a high light intensity, but not in direct sunlight.
- After 24 h, check that the female mites have laid eggs. Aim for at least 20 eggs per leaf section. If there are not enough eggs, leave for a further 24 h. Do not leave longer than 48 h.
- When sufficient egg numbers have been obtained, remove the mites with the fine pointed brush or cocktail stick. Record the time when this is done.
- Prepare appropriate test dilutions of formulations in water. The use of a wetter is not recommended.

For further information please contact: Alan Porter, IRAC International Coordinator
www.irac-online.org, email: a.porter@irac.org

IRAC Susceptibility Test Methods Series

Method No: 3

Version: 2

recommended.

- Agitate test liquids and then dip the leaf sections for 5 sec. Dip equal number of control leaf sections in water only.
- Record the number of eggs per leaf section.
- Return leaf sections to Petri dishes and maintain in conditions described above. Record maximum and minimum temperatures. Moisten cotton wool daily.
- Using a hand lens or binocular microscope observe leaf sections daily until there has been complete (or nearly complete) hatch on the untreated (water only) leaf sections. Record numbers of un-hatched eggs on treated and untreated leaf sections.
- Express results as percentage mortality and correct for untreated mortality using Abbott's formula. Untreated mortality should be recorded.

Percautions & Notes:

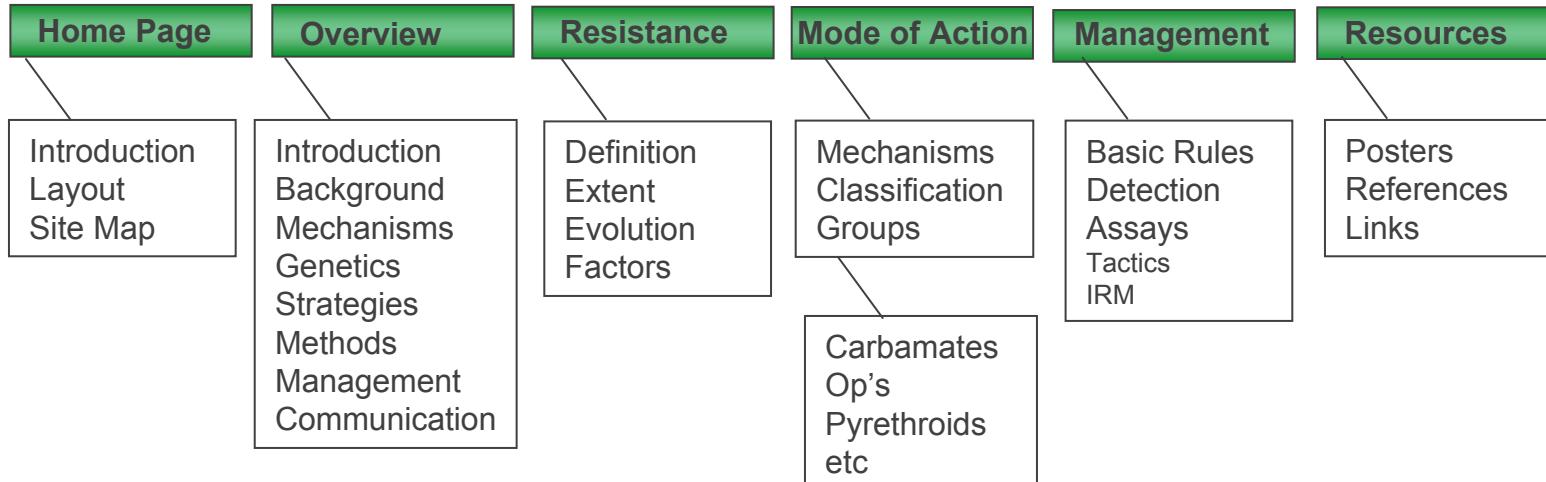
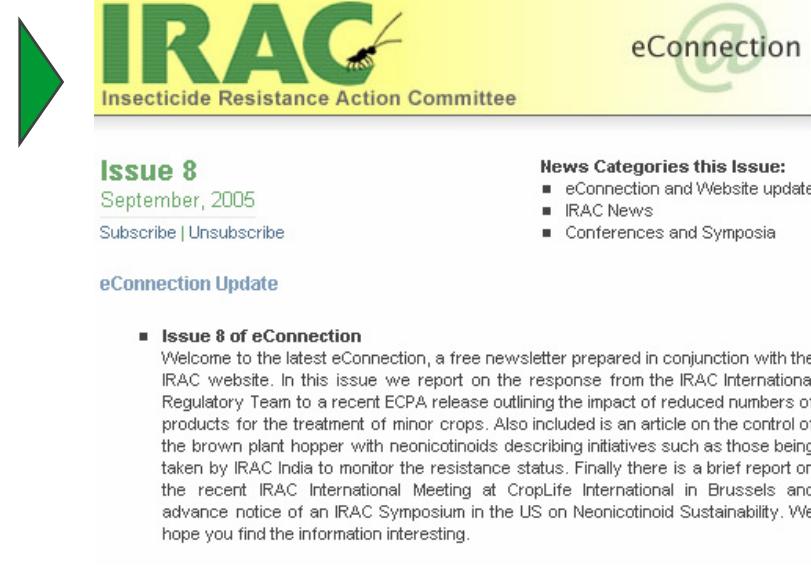
If the lids are left off, the leaf sections may dry out and, unless the cotton wool can be moistened at least daily, the test may be invalidated by excessive control mortality. In such circumstances, the method may have to be modified to suit the local conditions, e.g. use lids with holes cut in them to reduce water loss without creating a condensation problem.

For *Tetranychus cinnabarinus* which lies mainly on the lower leaf surface, the leaf sections may need to be placed lower surface uppermost. Leaves of kidney beans are particularly suitable.

- IRAC provides validated test methods of proven ability to detect changes in susceptibility
- IRAC currently has a program to update the methods
- Intention to include methods for all pests in new EPPO guidelines
- New methods will include biochemical and molecular methods

Communication & Education IRAC Activities

- **eConnection** – Free electronic IRAC newsletter raises awareness of resistance issues, existence of country groups, raises profile of IRAC
- **eLearning** - extensive, interactive web-based education and training modules being developed



Conclusions

- The major Neonicotinoid companies undertake extensive research to understand factors influencing the effectiveness of their compounds
- There is a large body of ongoing work to maintain awareness of susceptibility in key at-risk pests
- Key companies like Bayer CropScience and Syngenta are collaborating at a local level to harmonise their guidelines for IRM for the neonicotinoids
- IRAC works for the industry to promote awareness of and solutions to resistance
 - Communication and education on IRM are vital
 - IRAC provides key resources such as the MoA scheme, methodologies, IRM advice to help manage resistance
 - IRAC country groups work to tackle local problems