



Insecticide Resistance Action Committee Mode of Action Classification

Group 1: Acetylcholinesterase (AChE) inhibitors (Only major representatives of the groups are shown)

1A Carbamates
Aldicarb, Oxamyl, Methiocarb

1B Organophosphates
Acephate, Fenitrothion, Methamidophos, Chlorpyrifos, Profenofos, Malathion, Triazophos, Dimethoate

Group 2: GABA-gated chloride channel antagonists

2A Cyclodiene Organochlorines
Chlorfenvinphos, Endosulfan

2B Phenylpyrazoles (Fiproles)
Ethiprole, Fipronil

Group 3: Sodium channel modulators (Only major representatives of group 3A are shown)

3A Pyrethroids Pyrethrins
Bifenthrin, Deltamethrin, Cyfluthrin, Cypermethrin, Etofenprox, Permethrin, Lambda-cyhalothrin, Tefluthrin

3B DDT, Methoxychlor
DDT, Methoxychlor

Group 4: Nicotinic acetylcholine receptor (nAChR) competitive modulators

4A Neonicotinoids
Acetaminprid, Imidacloprid, Clothianidin, Thiacloprid, Flupyradifurone

4B Nicotine
Nicotine, Sulfoxalof

4C Sulfoximines
Sulfoximines

4D Butenolides
Thiamethoxam, Flupyradifurone

4E Mesolonic
Triflumazopyrim

Group 5: Nicotinic acetylcholine receptor (nAChR) allosteric modulators site I

5 Spinosyns

Group 6: Glutamate-gated chloride channel (GluCl) allosteric modulators

6 Avermectins & Milbemycins

Group 7: Juvenile hormone mimics

7A Juvenile hormone analogues
Hydroxypropryl, Kinoprene

7B Fenoxycarb, Fenoxycarb, Pyriproxyfen, Pyriproxyfen

Use of Groups and Sub-Groups:

- Alterations, sequences or rotations of compounds between MoA groups reduce selection for target site resistance.
- Applications are arranged into MoA spray windows defined by crop growth stage and pest biology.
- Several sprays of a compound may be possible within each spray window, but successive generations of a pest should not be treated with compounds from the same MoA group.
- Local expert advice should always be followed with regard to spray windows and timing.
- Groups in the classification whose members do not act at a common target site are exempt from the prescription against rotation within the group. These are: Group 8, Group 13 and all the UN groups: UN, UNB, UNE, UNF, UNK, UNP & UNV.
- Sub-groups represent distinct structural classes which are believed to have the same mode of action.
- Sub-groups provide differentiation between compounds that may bind at the same target site but are structurally different enough that risk of metabolic cross-resistance is lower than for close chemical analogs.
- Cross-resistance potential between sub-groups is higher than between groups, so rotation between sub-groups should be considered only when there are no alternatives, and only if cross-resistance does not exist, following consultation with local expert advice. These exceptions are not sustainable, and alternative options should be sought.
- Sub-group 3B: DDT is no longer used in agriculture and therefore this is only applicable for the control of insect vectors of human disease, such as mosquitoes, because of a lack of alternatives.
- Sub-group 10A: Hexythiazox is grouped with clofentezine because they exhibit cross-resistance even though they are structurally distinct. Diflovidazin has been added to this group because it is a close analogue of clofentezine and is expected to have the same mode of action.

Group 8: Miscellaneous non-specific (multi-site) inhibitors

8A Alkyl halides
Methyl bromide

8B Chloropirrin
Chloropirrin

8C Fluorides
Sulfonyl fluoride

8D Borates
Borax

8E Tartar emetic
Tartar emetic

8F Methyl isothiocyanate generators
Dazomet, Metam

Group 9: Chordonal organ TRPV channel modulators

9B Pyridine azomethine derivatives
Pymetrozine

9D Propenones
Pyrifluquinazon, Aldopropen

Group 10: Mite growth inhibitors affecting CHS1

10A Clofentezine, Diflovidazin, Hexythiazox

10B Etoxazole
Etoxazole

Group 11: Microbial disruptors of insect midgut membranes

Different B.t. products that target different insect orders may be used together without compromising their resistance management. Relation between certain specific B.t. microbial products may provide resistance management benefits for some pests. Consult product-specific recommendations.

* Where there are differences among the specific receptors within the midguts of target insects, transgenic crops containing certain combinations of these proteins provide resistance management benefits.

11A *Bacillus thuringiensis*
Bacillus thuringiensis and the insecticidal proteins produced B.t. israelensis, B.t. aizawa, B.t. kurstaki, B.t. tenebrionis B.t. crop protiens *
Cry1Ab, Cry1Ac, Cry1Fa, Cry1Aa, Cry1A10a, Cry2Ab, Vip3A, mCry3A, Cry3Ab, Cry3Bb, Cry34/35A1/Cry35A1

11B *Bacillus sphaericus*
Bacillus sphaericus

Group 12: Inhibitors of mitochondrial ATP synthase

12A Diafenthiuron
Diafenthiuron

12B Organotin miticides
Cyhexatin, Fenbutatin oxide, Propargite

12D Tetradifon
Tetradifon

Group 13: Uncouplers of oxidative phosphorylation via disruption of proton gradient

13 Pyrethroids, Dinitrophenols, Sulfuramid, DNOC

Group 14: Nicotinic acetylcholine receptor (nAChR) channel blockers

14 Nereistoxin analogues
Nereistoxin, Bursultox, Carip, Thiocyclam, Thiocyclam, Thiocyclam, Thiocyclam

Group 15: Inhibitors of chitin biosynthesis affecting CHS1

15 Benzoylureas
Diflubenzuron, Flufenoxuron, Lufenuron, Novalonur, Tebufenozuron

Group 16: Inhibitors of chitin biosynthesis, type 1

16 Buprofezin
Buprofezin

Group 17: Moulting disruptors, Dipteran

17 Cyromazine
Cyromazine

Group 18: Ecdysone receptor agonists

18 Diacylhydrazines
Chromafenozide, Halofenozide, Methoxyfenozide, Tebufenozide

Group 19: Octopamine receptor agonists

19 Amitraz
Amitraz

Group 20: Mitochondrial complex III electron transport inhibitors

20A Hydrarmethylinon
Hydrarmethylinon

20B Acequinocyl
Acequinocyl

20C Fluacrypyrim
Fluacrypyrim

20D Bifenazate
Bifenazate

Group 21: Mitochondrial complex I electron transport inhibitors

21B Rotenone
Rotenone

21A METI acaricides and insecticides
Fenpyrostate, Pyridaben, Tolfenpyrad, Tabufenpyrad

Group 22: Voltage-dependent sodium channel blockers

22A Oxadiazines
Indoxacarb, Oxadiazines

22B Semicarbazones
Metalfurzone

Group 23: Inhibitors of acetyl CoA carboxylase

23 Tetrone & Tetramic acid derivatives
Spirodicifen, Spiromesifen, Spirotetramat

Group 24: Mitochondrial complex IV electron transport inhibitors

AIP
Aluminum phosphide

Zn3P2
Zinc phosphide

Ca3P2
Calcium phosphide

24A Phosphides
Phosphine

24B Cyanides
Cyanide salts

Group 25: Mitochondrial complex II electron transport inhibitors

25A beta-Ketonitrile derivatives
Cyenoprafen, Cyflumetofen

25B Carboxanilides
Pyflubumide

Group 28: Ryanodine receptor modulators

R-Cl
Chlorantraniliprole R-Cl, Cyantraniliprole R-Cl

R-CN
Cyclaniliprole

28 Diamides
Flubendamide

29: Chordonal organ modulators - undefined target site

29 Floricamid
Floricamid

30 GABA-gated chloride channel allosteric modulators

30 Meta-diamides & Isoxazolines
Brotufenilide, Fluxametamide

31 Baculoviruses

Cydia pomonella GV, Thaumalobla leucostola GV, Granuloviruses & Nucleopolydoviruses

Anticarsa gemmatilis MNPV, Anticarsa arimigera NPV

32 Nicotinic Acetylcholine Receptor (nAChR) Allosteric Modulators - Site II

GS-omega/kappa HXDX-Hv Ia peptide

Group UN: Compounds of unknown or uncertain mode of action

S
CaSx (Lime sulfur)

Pyridalyl

Diocofol

Bromopropylate

Azadirachtin

Benzoximate

Chromethalon

Mancozeb

UNB Bacterial agents (non-Bt) of unknown or uncertain MoA

Burkholderia spp, Wolbachia piparitis (Zap)

UNE Botanical essence including synthetic, extracts and unrefined oils with unknown or uncertain MoA

Chenopodium ambricosides near ambricosides extract, Fatty acid monoesters with glycerol or propargenol, Neem oil

UNF Fungal agents of unknown or uncertain MoA

Beauveria bassiana strains, Metarhizium anisopliae strain F52, Phaeogenozia fuliginosa, Aploka strain 97

UNM Non-specific mechanical disruptors

Diatomaceous earth

Key to Targeted Physiology

- Nerve & Muscle
- Growth & Development
- Respiration
- Midgut
- Unknown or Non-specific

Poster Notes:

- Groups 26 and 27 are unassigned.
- The poster is for educational purposes only. Information presented is accurate to the best of our knowledge at the time of publication, but IRAC or its member companies cannot accept responsibility for how this information is used or interpreted. Advice should always be sought from local experts or advisors, and health and safety recommendations followed.
- In some cases only representative compounds in Groups are shown where indicated.
- Please visit www.ircac-online.org for the complete IRAC classification.

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