FRAC Classification of Fungicides

FRAC FUNGICIDE RESISTANCE ACTION COMMITTEE

Fungal control agents by cross resistance pattern and mode of action 2020 (www.frac.info)

A: Nucleic Acids Metabolism

A1: RNA polymerase I
4: PA-fungicides (PhenylAmides)

benalaxyl, benalaxyl-M, furalaxyl, metalaxyl, metalaxyl-M — acylalanines

ofurace — butyrolactones

oxadixyl — oxazolidinones

Group 4

A2: adenosine -deaminase
8: hydroxy (2-amino)-pyrimidines

bupirimate, dimethirimol, ethirimol

Group 8

A3: DNA / RNA synthesis (prop.)
32: heteroaromatics

hymexazole — isoxazoles

octhilinone — isothiazolones

Group 32

A4: DNA topoisomerase type II (gyrase)
31: carboxylic acids

oxolinic acid (bactericide)

Group 31

B: Cytoskeleton and Motor Proteins

B1: ▹ ß-tubulin assembly in mitosis
1: MBC fungicides (= Methyl Benzimidazole Carbamates)

benzimidazoles: benomyl, fuberidazole, thiabendazole, carbendazim

thiophanates: thiophanate, thiophanate-methyl

Group 1

B2: ▹ ß-tubulin assembly in mitosis*
10 N-phenyl carbamates

diethofencarb

* negative cross-resistance to B1

Group 10

B4: cell division (unknown site)
20 phenylureas

pencycuron

Group 20

B3: ▹ß-tubulin assembly in mitosis
22 benzamides and thiazole carboxamides

ethaboxam — ethylamino-thiazole carboxamide

zoxamide — toluamide

Group 22

B5: delocalisation of spectrin-like proteins
43 benzamides

fluopimomide, fluopicolide

Group 43

B6: actin/myosin/fimbrin function
47 cyanoacrylates — # 50 aryl-phenyl-ketones

phenamacril — aminocyanoacrylates

benzophenone, metrafenone — benzoylpyridine

pyriofenone

Group 47 / 50

C: Respiration

C1: complex I NADH Oxido-reductase
39 pyrimidinamines,pyrazole-MET1, quinazoline

diflumetorim — pyrimidinamine

tolfenpyrad — pyrazole-5-carboxamide

fenazaquin, quinazoline

Group 39

C4: complex III
cytochrome bc1(ubiquinone reductase) at Qi site
21 QiI fungicides (Quinone Inside Inhibitors)

cyano-imidazole: cyazofamid

sulfamoyl-triazole: amisulbrom

picolinamide: fenpicoxamid

Group 21

C7: ATP transport (proposed)
38 thiophene- carboxamides

silthiofam

Group 38

C8: inhibition of complex III
cytochrome bc1(ubiquinone reductase) at Qo site stigmatellin binding sub site)
45 QoSI-fungicide (stigmatellin binding)

ametoctradin — triazolo-pyrimidylamine

Group 45

C2: complex II: succinate-dehydrogenase
7 SDHI (Succinate DeHydrogenase Inhibitors)b

fluxapyroxad, penflufen, furametpyr, thifluzamide, isofetamid, fluopyram, oxycarboxin, oxathiin - carboxamides, carboxin, boscalid

pyrazole - 4 - carboxamides: bixafen, phenyl-oxo-ethyl thiophene amide, penthiopyrad, pyridinyl - ethyl - benzamides, pyrazine – carboxamides, isoflucypram, pyraziflumid, N-cyclopropyl-N-Benzyl-pyrazole-carboxamides, pydiflumetofen, N-methoxy-(phenyl-ethyl)-pyrazole-carboxamides

benzovindiflupyr, sedaxane, fluindapyr, inpyrfluxam, flutolanil, benodanil, mepronil, phenyl-benzamides

fenfuram — furan-carboxamides

Group 7

C5: uncouplers of oxidative Phosphorylation # 29

meptyl dinocap, dinitrophenyl crotonates — binapacryl

2,6-dinitro-aniline — fluazinam

Group 29

C6: inhibitors of oxidative phosphorylation, ATP synthase
30 organo tins

fentin acetate, fentin chloride, fentin hydroxide

Group 30

C3: complex III cytochrome bc1 (ubiquinol oxidase) at Qo site (*cyt b gene*)
11 QoI fungicides (Quinone outside Inhibitors)

enoxastrobin, coumoxystrobin, picoxystrobin, pyraoxystrobin — methoxy-acrylates

azoxystrobin, flufenoxystrobin — oximino-acetamides

fenaminstrobin, metominostrobin, orysastrobin, dimoxystrobin

pyrametostrobin, triclopyricarb, pyraclostrobin, pyribencarb — methoxy-carbamates

kresoxim-methyl, oximino-acetates, trifloxystrobin — dihydro-dioxazines

fluoxastrobin, famoxadone, oxazolidine-diones, mandestrobin — methoxy-acetamides

fenamidone — imidazolinones

benzyl-carbamates

metyltetraprole

Group 11 — Group 11A

11A QoI fungicides (Quinone outside Inhibitors); Subgroup A

D: Amino Acid and Protein Synthesis

D1: methionine biosynthesis (cgs gene) (proposed)
9 Anilino-Pyrimidines (AP fungicides)

cyprodinil, mepanipyrim, pyrimethanil

Group 9

D2: protein synthesis (ribosome, termination step)
23 enopyranuronic acid

blasticidin-S

Group 23

D3: protein synthesis (ribosome, initiation step)
24 hexopyranosyl antibiotics

kasugamycin

Group 24

D4: protein synthesis (ribosome, initiation step)
25 glucopyranosyl antibiotics

streptomycin (bactericide)

Group 25

D5: protein synthesis (ribosome, elongation step)
41 tetracycline antibiotics

oxytetracycline (bactericide)

Group 41

E: Signal Transduction

E1: signal transduction (mechanism unknown)
#13 azanaphthalenes

aryloxyquinoline: quinoxyfen

quinazolinone: proquinazid

Group 13

E2: osmotic signal transduction ▹ MAP / histidine- kinase (os-2, HOG1)
12 phenylpyrroles (PP- fungicides)

fenpiclonil

fludioxonil

Group 12

E3: osmotic signal transduction ▹ MAP / histidine kinase (os-1, Daf1)
2 dicarboximides

iprodione, chlozolinate, vinclozolin, procymidone, dimethachlone

Group 2

F: Lipid Synthesis or Transport / Membrane Integrity or Function

F2: phospholipid biosynthesis ▹ methyltransferase
6 phosphorothiolates & dithiolanes

phosphorothiolates: pyrazophos, iprobenfos, edifenphos

dithiolanes: iso-prothiolane

Group 6

F3: cell peroxidation (prop.)
14 aromatic hydrocarbons & heteroaromatics

aromatic hydrocarbons: tecnazene (TCNB), dicloran, quintozene (PCNB), biphenyl, tolclofos-methyl, chloroneb

1,2,4-thiadiazole: etridiazole

Group 14

F4: cell membrane permeability, fatty acids (prop.)
28 carbamates

prothiocarb, propamocarb, iodocarb

Group 28

F7: cell membrane disruption
46 plant extract

Melaleuca alternifolia (Tea Tree Extract)

Plant oils (mixtures) eugenol, geraniol thymol

Group 46

F8: ergosterol binding
48 polyene

natamycin (pimaricin)

Group 48

F9: lipid homeostasis and transfer/storage
49 OSBPI Oxysterol binding protein homologue inhibition

oxathiapiprolin

fluoxapiprolin

Piperidinyl thiazole - isoxazoline

Group 49

I: Melanin Synthesis in Cell Wall

I1: reductase in melanin biosynthesis
16.1 Melanin Biosynthesis Inhibitors: Reductase (MBI-R)

isobenzo-furanone: fthalide

pyrrolo-quinolione: pyroquilon

triazolobenzo-thiazole: tricyclazole

Group 16.1

I2: dehydratase in melanin biosynthesis
16.2 Melanin Biosynthesis Inhibitors: Dehydratase (MBI-D)

carboxamide: diclocymet

cyclopropane carboxamide: carpropamid

propionamide: fenoxanil

Group 16.2

I3: polyketide synthase in melanin biosynthesis
16.3 Melanin Biosynthesis Inhibitors: Polyketide synthase (MBI-P)

trifluoroethyl-carbamate: tolprocarb

Group 16.3

G: Sterol Biosynthesis in Membranes

G1: C14-demethylase in sterol biosynthesis (*erg11/cyp51*)
3 DMI-fungicides (DeMethylation Inhibitors)
(SBI : Class I)

triazoles: etaconazole, azaconazole, ipconazole, fenbuconazole, tebuconazole, bitertanol, metconazole, fluquinconazole, tetraconazole, bromuconazole, myclobutanil, flusilazole, triadimefon, cyproconazole, penconazole, triadimenol, flutriafol, propiconazole, difenoconazole, hexaconazole, triticonazole, diniconazole, simeconazole, imibenconazole, mefentrifluconazole, epoxiconazole, prothioconazole

piperazines: triforine

pyridines: pyrifenox, pyrisoxazole

pyrimidines: nuarimol, fenarimol

imidazoles: imazalil, triflumizole, pefurazoate, oxpoconazole, prochloraz

triazolinthione

Group 3

G2:Δ14-reductase and Δ8→ Δ7-isomerase in sterol biosynthesis (*erg2, erg 24*)
5 Amines("Morpholines") (SBI : Class II)

piperidines: fenpropidin, piperalin

spiroketal-amines: spiroxamine

aldimorph, fenpropimorph, tridemorph (x = 10 to 13; x = 12 – 70%), dodemorph

morpholines

Group 5

G3:3-keto reductase in C4-de-methylation (*erg27*)
17 (KRI fungicides KetoReductase Inhibitors) (SBI : Class III)

hydroxyanilides: fenhexamid

amino-pyrazolinones: fenpyrazamine

Group 17

G4: squalene epoxidase in sterol biosynthesis (*erg1*)
18 (SBI : Class IV)

allylamines: terbinafine, naftifine

thiocarbamates: pyributicarb (herbicide)

Group 18

H: Cell Wall Biosynthesis

H4: chitin synthase
19 Polyoxins

Peptidyl Pyrimidine nucleoside: Polyoxin

Group 19

H5: cellulose synthase
40 Carboxylic Acid Amides (CAA fungicides)

cinnamic acid amides: dimethomorph, flumorph, pyrimorph

valinamide carbamates: benthiavalicarb, iprovalicarb, valifenalate

mandelic acid amides: mandipropamid

Group 40

P: Host Plant Defence Induction

P1: salicylate related
#P01 benzothiadiazole BTH

acibenzolar-S-methyl

Group P01

P2: salicylate related
#P02 benzothiazole

probenazole

Group P02

P3: salicylate related
#P03 thiadiazole carboxamide

isotianil, tiadinil

Group P03

P4: polysaccharide elicitors

laminarin — #P04 polysaccharide

Group P04

P5: anthraquinone elicitors

Reynoutria sachalinensis (Giant Knotweed Extract) — #P05 plant extract

Group P05

P6: microbial elicitors
#P06

Bacillus mycoides (Isolate J) — Bacterial *Bacillus* spp.

Cell walls of *Saccharomyces cerevisiae* Strain LAS117 — Fungal *Saccharomyces* spp.

Group P06

P7: phosphonates
#P07 phosphonates

fosetyl-al — ethyl-phosphonates

H_3PO_3 — Phosphorous acid

phosphorous acid

Group P07

NC : Not Classified

Mineral oils, organic oils, inorganic salts material of biological origin

M: Chemicals with Multi-Site Activity

M01: inorganic (electrophiles)
copper preparations — Cu

Group M01

M02: inorganic (electrophiles)
Sulphur — S

Group M02

M05: chloronitriles (unspecified mechanism)
chlorothalonil

Group M05

M08: triazines (unspecified mechanism)
anilazine

Group M08

M09: anthraquinones (electrophiles)
dithianon

Group M09

M12: thiocarbamates (electrophiles)
methasulfocarb

Group M12

M03: dithiocarbamates & relatives (electrophiles)
febram, mancozeb, zinc thiazole, propineb, metiram, zineb, maneb, thiram, ziram

Group M03

M04: phthalimides (electrophiles)
captan, captafol, folpet

Group M04

M06: sulphamides (electrophiles)
dichlofluanid, tolylfluanid

Group M06

M10: quinoxalines (electrophiles)
chinomethionat

Group M10

M11: maleimides (electrophiles)
fluoroimide

Group M11

M07: bis-guanidines (membrane disruptors, detergents)
mixture of iminoctadine and other polyamines — guazatine

iminoctadine

Group M07

Unknown Mode of Action

•Temporary status; information on mode of action and / or resistance risk is still uncertain

cymoxanil — Group 27 — cyanoacetamide-oxime

tecloftalam — Group 34 — phthalamic acid

triazoxide — Group 35 — benzotriazines

flusulfamide — Group 36 — benzene-sulfonamides

diclomezine — Group 37 — pyridazinones

cyflufenamid — Group 06 — phenyl acetamides

flutianil — Group U13 — cyano-methylene thiazolidine

ferimzone — Group U14 — pyrimidinone-hydrazones

tebufloquin — Group U16 — 4-quinolyl acetate

picarbutrazox — Group U17 — tetrazolyloxime

validamycin — Group U18 — glucopyranosyl antibiotic

dodine — Group U12 — guanidines

BM: Biologicals with Multiple Modes of Action

BM 02: microbial (living microbes or extract, metabolites)

Trichoderma atroviride (strain SC1); *Trichoderma atroviride* (strain I-1237); *Trichoderma atroviride* (strain T34) — fungal *Trichoderma* spp.

Pseudomonas chlororaphis (strain AFS009) — bacterial *Pseudomonas* spp.

Bacillus subtilis (strain AFS032321); *Bacillus amyloliquefaciens* (strain QST713 strain FZB24 strain MBI600 strain D747 strain F727) — bacterial *Bacillus* spp.

Trichoderma atroviride (strain LU132); *Streptomyces lydicus* (strain WYEC108); *Streptomyces griseovirides* (strain K61) — bacterial *Streptomyces* spp.

Clonostachys rosea (strain CR-7); *Glocladium catenulatum* (strain J1446) — fungal *Clonostachys* spp.

BM 01: plant extract

Extract from the cotyledons of lupine plantlets (BLAD) — polypeptide lectin

Extract from *Swinglea glutinosa* — phenols, sesquiterpenes, triterpenoids, coumarins

Mode of action of fungicides based on the FRAC fungicide list (published at www.frac.info/publications)

The groupings can change when new results become known. Yellow zones represent accepted mode of action groups or in cases where resistance is known, cross resistance groups. In spite of this general classification, cases of incomplete or even lacking cross resistance are known within some groups with some pathogens.

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Legend:

C: Respiration

C2: inhibition of complex II: ▹ succinate-dehydrogenase
7 SDHI (Succinate dehydrogenase inhibitors)

thiazole carboxamides

- mode of action group
- sub-group
- ▹ target site of action (where known) or putative target site (=prop.)
- FRAC code no. (#) and group name
- chemical (sub-) group

FRAC Mode of Action Poster © This version was updated in February 2020

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