

TITLE

BIS(ARYL)CATECHOL DERIVATIVES AS HERBICIDES

FIELD OF THE INVENTION

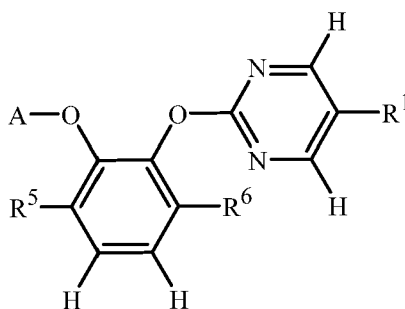
This invention relates to certain 3-cyano-1-pyrimidinyloxy benzene derivatives, their
 5 *N*-oxides, salts and compositions, and methods of their use for controlling undesirable
 vegetation.

BACKGROUND OF THE INVENTION

The control of undesired vegetation is extremely important in achieving high crop
 efficiency. Achievement of selective control of the growth of weeds especially in such
 10 useful crops as rice, soybean, sugar beet, maize, potato, wheat, barley, tomato and plantation
 crops, among others, is very desirable. Unchecked weed growth in such useful crops can
 cause significant reduction in productivity and thereby result in increased costs to the
 consumer. The control of undesired vegetation in noncrop areas is also important. Many
 products are commercially available for these purposes, but the need continues for new
 15 compounds that are more effective, less costly, less toxic, environmentally safer or have
 different sites of action.

SUMMARY OF THE INVENTION

This invention is directed to compounds of Formula 1 (including all stereoisomers),
N-oxides, and salts thereof, agricultural compositions containing them and their use as
 20 herbicides:

**1**

wherein

A is a phenyl ring optionally substituted with up to 4 R^2 ; or a 5- or 6-membered
 heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a
 25 carbon atom, and optionally substituted with up to 4 R^2 ;

R^1 is halogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_4
 alkoxy or $S(O)_mR^3$;

each R^2 is independently halogen, cyano, nitro, SF_5 , CHO, $C(=O)NH_2$, $C(=S)NH_2$,
 SO_2NH_2 , C_1 - C_4 alkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 haloalkyl, C_2 - C_4

haloalkenyl, C₂-C₄ haloalkynyl, C₃-C₆ cycloalkyl, C₃-C₆ halocycloalkyl, C₄-C₈ alkylcycloalkyl, C₄-C₈ cycloalkylalkyl, C₂-C₆ alkylcarbonyl, C₂-C₆ haloalkylcarbonyl, C₂-C₆ alkoxy carbonyl, C₃-C₇ cycloalkylcarbonyl, C₂-C₈ alkylaminocarbonyl, C₃-C₁₀ dialkylaminocarbonyl, C₁-C₄ alkoxy, C₃-C₄ alkenyloxy, C₃-C₄ alkynyloxy, C₁-C₄ haloalkoxy, C₃-C₄ haloalkenyloxy, C₃-C₄ haloalkynyloxy, C₃-C₆ cycloalkoxy, C₃-C₆ halocycloalkoxy, C₄-C₈ cycloalkylalkoxy, C₂-C₆ alkoxyalkyl, C₂-C₆ haloalkoxyalkyl, C₂-C₆ alkoxyhaloalkyl, C₂-C₆ alkoxyalkoxy, C₂-C₄ alkylcarbonyloxy, C₂-C₆ cyanoalkyl, C₂-C₆ cyanoalkoxy, C₁-C₄ hydroxyalkyl, C₂-C₄ alkylthioalkyl, C₁-C₆ alkylamino, C₂-C₆ dialkylamino, S(O)_nR⁴, CH(=NOH), phenyl or pyridinyl; each R³ and R⁴ is independently C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkylamino or C₂-C₆ dialkylamino;

R⁵ is halogen, cyano or C₁-C₂ haloalkyl;

R⁶ is H or F;

m is 0, 1 or 2; and

each n is independently 0, 1 or 2;

provided the compound of Formula 1 is other than 5-bromo-2-[3-bromo-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine, 5-bromo-2-[6-bromo-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine, 5-chloro-2-[3-fluoro-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine, 5-chloro-2-[6-fluoro-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine, 5-chloro-2-[3-methyl-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine or 5-chloro-2-[6-methyl-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine.

More particularly, this invention pertains to a compound of Formula 1 (including all stereoisomers), an *N*-oxide or a salt thereof. This invention also relates to a herbicidal composition comprising a compound of the invention (i.e. in a herbicidally effective amount) and at least one component selected from the group consisting of surfactants, solid diluents and liquid diluents. This invention further relates to a method for controlling the growth of undesired vegetation comprising contacting the vegetation or its environment with a herbicidally effective amount of a compound of the invention (e.g., as a composition described herein).

This invention also includes a herbicidal mixture comprising (a) a compound selected from Formula 1, *N*-oxides, and salts thereof, and (b) at least one additional active ingredient selected from (b1) through (b16) and salts of compounds of (b1) through (b16), as described below.

DETAILS OF THE INVENTION

As used herein, the terms “comprises,” “comprising,” “includes,” “including,” “has,” “having,” “contains,” “containing,” “characterized by” or any other variation thereof, are intended to cover a non-exclusive inclusion, subject to any limitation explicitly indicated.

5 For example, a composition, mixture, process, method, article, or apparatus that comprises a list of elements is not necessarily limited to only those elements but may include other elements not expressly listed or inherent to such composition, mixture, process, method, article, or apparatus.

10 The transitional phrase “consisting of” excludes any element, step, or ingredient not specified. If in the claim, such would close the claim to the inclusion of materials other than those recited except for impurities ordinarily associated therewith. When the phrase “consisting of” appears in a clause of the body of a claim, rather than immediately following the preamble, it limits only the element set forth in that clause; other elements are not excluded from the claim as a whole.

15 The transitional phrase “consisting essentially of” is used to define a composition, method or apparatus that includes materials, steps, features, components, or elements, in addition to those literally disclosed, provided that these additional materials, steps, features, components, or elements do not materially affect the basic and novel characteristic(s) of the claimed invention. The term “consisting essentially of” occupies a middle ground between
20 “comprising” and “consisting of”.

Where applicants have defined an invention or a portion thereof with an open-ended term such as “comprising,” it should be readily understood that (unless otherwise stated) the description should be interpreted to also describe such an invention using the terms “consisting essentially of” or “consisting of.”

25 Further, unless expressly stated to the contrary, “or” refers to an inclusive or and not to an exclusive or. For example, a condition A or B is satisfied by any one of the following: A is true (or present) and B is false (or not present), A is false (or not present) and B is true (or present), and both A and B are true (or present).

Also, the indefinite articles “a” and “an” preceding an element or component of the
30 invention are intended to be nonrestrictive regarding the number of instances (i.e. occurrences) of the element or component. Therefore “a” or “an” should be read to include one or at least one, and the singular word form of the element or component also includes the plural unless the number is obviously meant to be singular.

As referred to herein, the term “seedling”, used either alone or in a combination of
35 words means a young plant developing from the embryo of a seed.

As referred to herein, the term “broadleaf” used either alone or in words such as “broadleaf weed” means dicot or dicotyledon, a term used to describe a group of angiosperms characterized by embryos having two cotyledons.

In the above recitations, the term “alkyl”, used either alone or in compound words such as “alkylthioalkyl” or “haloalkyl” includes straight-chain or branched alkyl, such as, methyl, ethyl, *n*-propyl, *i*-propyl, or the different butyl isomers. “Alkenyl” includes straight-chain or branched alkenes such as ethenyl, 1-propenyl, 2-propenyl, and the different butenyl, pentenyl and hexenyl isomers. “Alkenyl” also includes polyenes such as 1,2-propadienyl and 2,4-hexadienyl. “Alkynyl” includes straight-chain or branched alkynes such as ethynyl, 1-propynyl, 2-propynyl and the different butynyl, pentynyl and hexynyl isomers. “Alkynyl” can also include moieties comprised of multiple triple bonds such as 2,5-hexadiynyl.

“Alkoxy” includes, for example, methoxy, ethoxy, *n*-propyloxy, isopropyloxy and the different butoxy isomers. “Alkoxyalkyl” denotes alkoxy substitution on alkyl. Examples of “alkoxyalkyl” include CH_3OCH_2 , $\text{CH}_3\text{OCH}_2\text{CH}_2$, $\text{CH}_3\text{CH}_2\text{OCH}_2$ and $\text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_2$. “Alkenyloxy” includes straight-chain or branched alkenyloxy moieties. Examples of “alkenyloxy” include $\text{H}_2\text{C}=\text{CHCH}_2\text{O}$, $(\text{CH}_3)\text{CH}=\text{CHCH}_2\text{O}$ and $\text{CH}_2=\text{CHCH}_2\text{CH}_2\text{O}$. “Alkynyloxy” includes straight-chain or branched alkynyloxy moieties. Examples of “alkynyloxy” include $\text{HC}\equiv\text{CCH}_2\text{O}$ and $\text{CH}_3\text{C}\equiv\text{CCH}_2\text{O}$. “Alkylthio” includes branched or straight-chain alkylthio moieties such as methylthio, ethylthio, and the different propylthio and butylthio isomers. “Alkylsulfinyl” includes both enantiomers of an alkylsulfinyl group. Examples of “alkylsulfinyl” include $\text{CH}_3\text{S(O)-}$, $\text{CH}_3\text{CH}_2\text{S(O)-}$, $\text{CH}_3\text{CH}_2\text{CH}_2\text{S(O)-}$, $(\text{CH}_3)_2\text{CHS(O)-}$ and the different butylsulfinyl isomers. Examples of “alkylsulfonyl” include $\text{CH}_3\text{S(O)}_2\text{-}$, $\text{CH}_3\text{CH}_2\text{S(O)}_2\text{-}$, $\text{CH}_3\text{CH}_2\text{CH}_2\text{S(O)}_2\text{-}$, $(\text{CH}_3)_2\text{CHS(O)}_2\text{-}$, and the different butylsulfonyl isomers. “Alkylthioalkyl” denotes alkylthio substitution on alkyl. Examples of “alkylthioalkyl” include CH_3SCH_2 , $\text{CH}_3\text{SCH}_2\text{CH}_2$, $\text{CH}_3\text{CH}_2\text{SCH}_2$ and $\text{CH}_3\text{CH}_2\text{SCH}_2\text{CH}_2$. “Alkylamino”, “dialkylamino”, and the like, are defined analogously to the above examples. “Cyanoalkyl” denotes an alkyl group substituted with one cyano group. Examples of “cyanoalkyl” include NCCH_2 , NCCH_2CH_2 and $\text{CH}_3\text{CH}(\text{CN})\text{CH}_2$.

The term “halogen”, either alone or in compound words such as “haloalkyl”, or when used in descriptions such as “alkyl substituted with halogen” includes fluorine, chlorine, bromine or iodine. Further, when used in compound words such as “haloalkyl”, or when used in descriptions such as “alkyl substituted with halogen” said alkyl may be partially or fully substituted with halogen atoms which may be the same or different. Examples of “haloalkyl” or “alkyl substituted with halogen” include F_3C , ClCH_2 , CF_3CH_2 and CF_3CCl_2 . The terms “haloalkoxy”, “haloalkylthio”, and the like, are defined analogously to the term “haloalkyl”. Examples of “haloalkoxy” include $\text{CF}_3\text{O-}$, $\text{CCl}_3\text{CH}_2\text{O-}$, $\text{HCF}_2\text{CH}_2\text{CH}_2\text{O-}$ and $\text{CF}_3\text{CH}_2\text{O-}$. Examples of “haloalkylthio” include $\text{CCl}_3\text{S-}$, $\text{CF}_3\text{S-}$, $\text{CCl}_3\text{CH}_2\text{S-}$ and $\text{ClCH}_2\text{CH}_2\text{CH}_2\text{S-}$. Examples of “haloalkylsulfinyl” include $\text{CF}_3\text{S(O)-}$, $\text{CCl}_3\text{S(O)-}$, $\text{CF}_3\text{CH}_2\text{S(O)-}$ and $\text{CF}_3\text{CF}_2\text{S(O)-}$. Examples of “haloalkylsulfonyl” include $\text{CF}_3\text{S(O)}_2\text{-}$, $\text{CCl}_3\text{S(O)}_2\text{-}$, $\text{CF}_3\text{CH}_2\text{S(O)}_2\text{-}$ and $\text{CF}_3\text{CF}_2\text{S(O)}_2\text{-}$.

The total number of carbon atoms in a substituent group is indicated by the “C_i-C_j” prefix where i and j are numbers from 1 to 6. For example, C₁-C₄ alkylsulfonyl designates methylsulfonyl through butylsulfonyl; C₂ alkoxyalkyl designates CH₃OCH₂-; C₃ alkoxyalkyl designates, for example, CH₃CH(OCH₃)-, CH₃OCH₂CH₂- or CH₃CH₂OCH₂-; and C₄ alkoxyalkyl designates the various isomers of an alkyl group substituted with an alkoxy group containing a total of four carbon atoms, examples including CH₃CH₂CH₂OCH₂- and CH₃CH₂OCH₂CH₂-.

When a compound is substituted with a substituent bearing a subscript that indicates the number of said substituents can exceed 1, said substituents (when they exceed 1) are independently selected from the group of defined substituents, e.g., (R²)_r in Embodiment 4 wherein r is 0, 1, 2 or 3. When one or more positions on a group are said to be “not substituted” or “unsubstituted”, then hydrogen atoms are attached to take up any free valency.

The term “heterocyclic ring” and “heterocycle” denote a ring in which at least one atom in the ring backbone is not carbon, e.g., nitrogen, oxygen or sulfur. The ring member atoms of the 6-membered heteroaromatic rings forming present substituent A typically consist of carbon atoms and one to three nitrogen atoms. The expression “fully unsaturated” in relation to a ring means that the bonds between the atoms in the ring are single or double bonds according to valence bond theory and furthermore the bonds between the atoms in the ring include as many double bonds as possible without double bonds being cumulative (i.e. no C=C=C, N=C=C, etc.). When a fully unsaturated heterocyclic ring satisfies Hückel’s rule, then said ring is also called a “heteroaromatic ring”. “Aromatic” or “heteroaromatic” according to Hückel’s rule means that each of the ring atoms is essentially in the same plane and has a p-orbital perpendicular to the ring plane, and that (4n + 2) π electrons, where n is a positive integer, are associated with the ring.

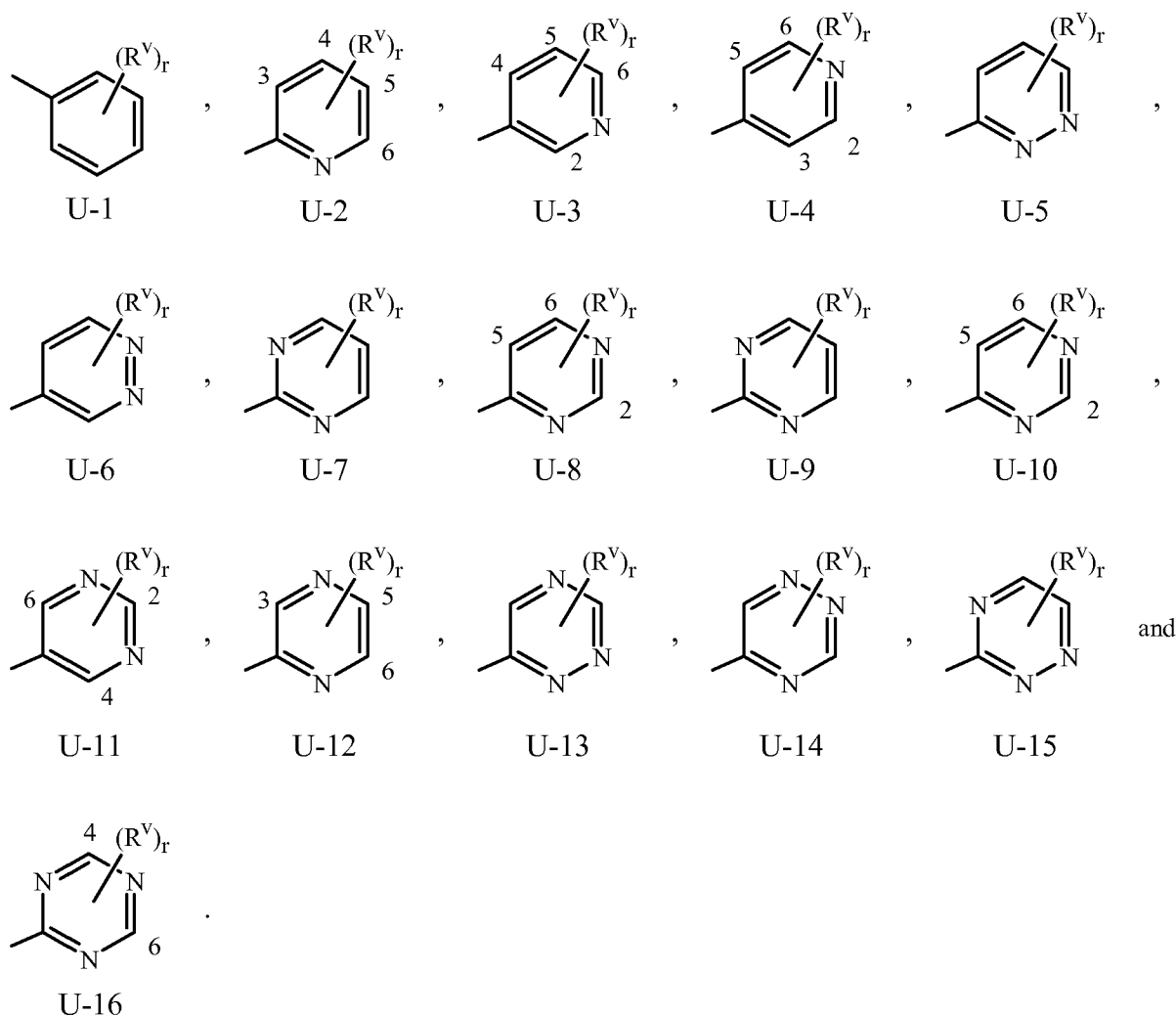
As used herein, the following definitions shall apply unless otherwise indicated. The term “optionally substituted” is used interchangeably with the phrase “unsubstituted or substituted”. Unless otherwise indicated, an optionally substituted group may have a substituent at each substitutable position of the group, and each substitution is independent of the other.

As noted above, A can be (among others) phenyl optionally substituted with one or more substituents selected from a group of substituents as defined in the Summary of the Invention. An example of phenyl optionally substituted with one to five substituents is the ring illustrated as U-1 in Exhibit 1, wherein R^v is R² as defined in the Summary of the Invention for A and r is an integer (from 0 to 4).

As noted above, A can be, among others, a 6-membered heteroaromatic ring, optionally substituted with up to 4 substituents selected from a group of substituents as defined in the Summary of the Invention. When A is a 6-membered nitrogen-containing

heteroaromatic ring, it may be attached to the remainder of Formula 1 through any available carbon ring atom, unless otherwise described. Examples of a 6-membered heteroaromatic ring optionally substituted with up to 4 substituents include the rings U-2 through U-14 illustrated in Exhibit 1 wherein R^V is any substituent as defined in the Summary of the Invention for A (i.e. R^2) and r is an integer from 0 to 4, limited by the number of available positions on each U group.

Exhibit 1



Although R^V groups are shown in the structures U-1 through U-14, it is noted that they do not need to be present since they are optional substituents. Note that when the attachment point between $(R^V)_r$ and the U group is illustrated as floating, $(R^V)_r$ can be attached to any available carbon atom or nitrogen atom of the U group. Preferably R^V substituents are attached to carbon ring atoms. Note that some U groups can only be substituted with less than 4 R^V groups on carbon ring atoms (e.g., U-5 through U-16).

A wide variety of synthetic methods are known in the art to enable preparation of aromatic and nonaromatic heterocyclic rings and ring systems; for extensive reviews see the

eight volume set of *Comprehensive Heterocyclic Chemistry*, A. R. Katritzky and C. W. Rees editors-in-chief, Pergamon Press, Oxford, 1984 and the twelve volume set of *Comprehensive Heterocyclic Chemistry II*, A. R. Katritzky, C. W. Rees and E. F. V. Scriven editors-in-chief, Pergamon Press, Oxford, 1996.

5 Compounds of this invention can exist as one or more stereoisomers. The various stereoisomers include enantiomers, diastereomers, atropisomers and geometric isomers. Stereoisomers are isomers of identical constitution but differing in the arrangement of their atoms in space and include enantiomers, diastereomers, cis-trans isomers (also known as geometric isomers) and atropisomers. Atropisomers result from restricted rotation about
10 single bonds where the rotational barrier is high enough to permit isolation of the isomeric species. One skilled in the art will appreciate that one stereoisomer may be more active and/or may exhibit beneficial effects when enriched relative to the other stereoisomer(s) or when separated from the other stereoisomer(s). Additionally, the skilled artisan knows how to separate, enrich, and/or to selectively prepare said stereoisomers. The compounds of the
15 invention may be present as a mixture of stereoisomers, individual stereoisomers or as an optically active form.

 Compounds of Formula **1** typically exist in more than one form, and Formula **1** thus include all crystalline and non-crystalline forms of the compounds they represent. Non-crystalline forms include embodiments which are solids such as waxes and gums as well as
20 embodiments which are liquids such as solutions and melts. Crystalline forms include embodiments which represent essentially a single crystal type and embodiments which represent a mixture of polymorphs (i.e. different crystalline types). The term “polymorph” refers to a particular crystalline form of a chemical compound that can crystallize in different crystalline forms, these forms having different arrangements and/or conformations of the
25 molecules in the crystal lattice. Although polymorphs can have the same chemical composition, they can also differ in composition due the presence or absence of co-crystallized water or other molecules, which can be weakly or strongly bound in the lattice. Polymorphs can differ in such chemical, physical and biological properties as crystal shape, density, hardness, color, chemical stability, melting point, hygroscopicity, suspensibility,
30 dissolution rate and biological availability. One skilled in the art will appreciate that a polymorph of a compound of Formula **1** can exhibit beneficial effects (e.g., suitability for preparation of useful formulations, improved biological performance) relative to another polymorph or a mixture of polymorphs of the same compound of Formula **1**. Preparation and isolation of a particular polymorph of a compound of Formula **1** can be achieved by
35 methods known to those skilled in the art including, for example, crystallization using selected solvents and temperatures. For a comprehensive discussion of polymorphism see R. Hilfiker, Ed., *Polymorphism in the Pharmaceutical Industry*, Wiley-VCH, Weinheim, 2006.

One skilled in the art will appreciate that not all nitrogen-containing heterocycles can form *N*-oxides since the nitrogen requires an available lone pair for oxidation to the oxide; one skilled in the art will recognize those nitrogen-containing heterocycles which can form *N*-oxides. One skilled in the art will also recognize that tertiary amines can form *N*-oxides.

5 Synthetic methods for the preparation of *N*-oxides of heterocycles and tertiary amines are very well known by one skilled in the art including the oxidation of heterocycles and tertiary amines with peroxy acids such as peracetic and *m*-chloroperbenzoic acid (MCPBA), hydrogen peroxide, alkyl hydroperoxides such as *t*-butyl hydroperoxide, sodium perborate, and dioxiranes such as dimethyldioxirane. These methods for the preparation of *N*-oxides
10 have been extensively described and reviewed in the literature, see for example: T. L. Gilchrist in *Comprehensive Organic Synthesis*, vol. 7, pp 748–750, S. V. Ley, Ed., Pergamon Press; M. Tisler and B. Stanovnik in *Comprehensive Heterocyclic Chemistry*, vol. 3, pp 18–20, A. J. Boulton and A. McKillop, Eds., Pergamon Press; M. R. Grimmett and B. R. T. Keene in *Advances in Heterocyclic Chemistry*, vol. 43, pp 149–161, A. R. Katritzky,
15 Ed., Academic Press; M. Tisler and B. Stanovnik in *Advances in Heterocyclic Chemistry*, vol. 9, pp 285–291, A. R. Katritzky and A. J. Boulton, Eds., Academic Press; and G. W. H. Cheeseman and E. S. G. Werstiuk in *Advances in Heterocyclic Chemistry*, vol. 22, pp 390–392, A. R. Katritzky and A. J. Boulton, Eds., Academic Press.

One skilled in the art recognizes that because in the environment and under
20 physiological conditions salts of chemical compounds are in equilibrium with their corresponding nonsalt forms, salts share the biological utility of the nonsalt forms. Thus a wide variety of salts of a compound of Formula 1 are useful for control of undesired vegetation (i.e. are agriculturally suitable). The salts of a compound of Formula 1 include acid-addition salts with inorganic or organic acids such as hydrobromic, hydrochloric, nitric,
25 phosphoric, sulfuric, acetic, butyric, fumaric, lactic, maleic, malonic, oxalic, propionic, salicylic, tartaric, 4-toluenesulfonic or valeric acids. Accordingly, the present invention comprises compounds selected from Formula 1, *N*-oxides and agriculturally suitable salts thereof.

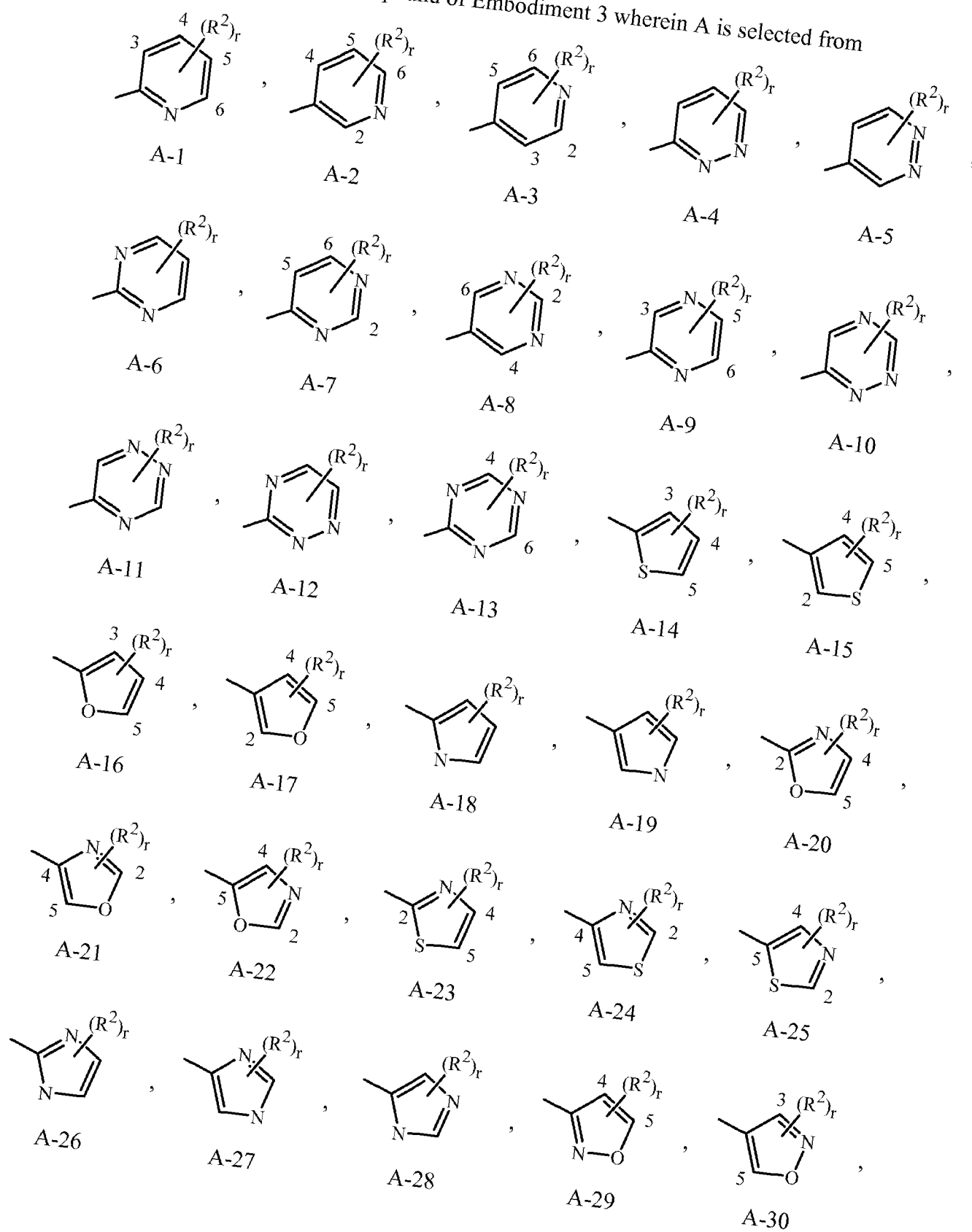
Embodiments of the present invention as described in the Summary of the Invention
30 include (where Formula 1 as used in the following Embodiments includes *N*-oxides and salts thereof):

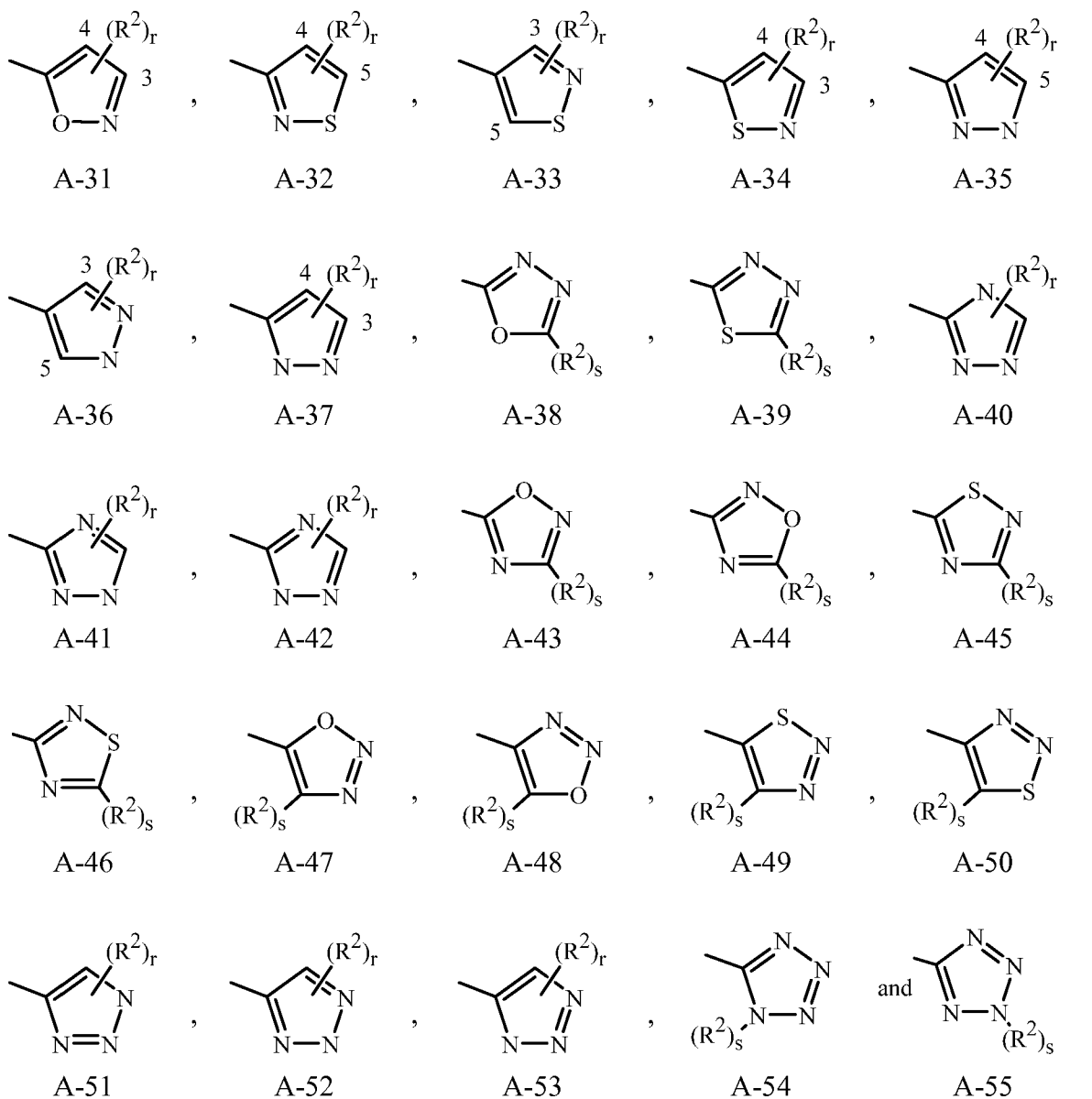
Embodiment 1. A compound of Formula 1 wherein A is a phenyl ring optionally substituted with up to 4 R².

Embodiment 2. A compound of Embodiment 1 wherein A is a phenyl ring optionally
35 substituted with up to 2 R².

Embodiment 3. A compound of Formula 1 wherein A is a 5- or 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R².

Embodiment 4. A compound of Embodiment 3 wherein A is selected from





wherein r is 0, 1, 2 or 3 and s is 0 or 1.

Embodiment 5. A compound of Embodiment 4 wherein A is selected from A-1, A-2, A-4, A-6, A-9, A-10, A-11, A-12 and A-23.

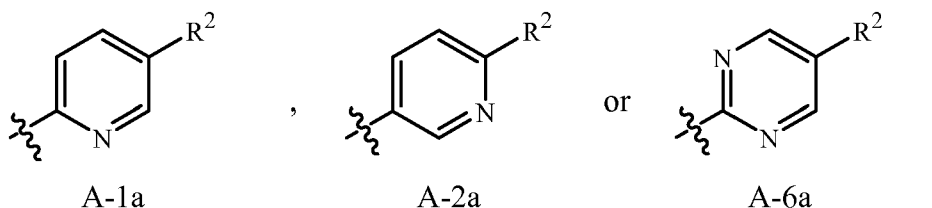
Embodiment 6. A compound of Embodiment 5 wherein A is selected from A-1, A-2 and A-6.

Embodiment 7. A compound of Embodiment 6 wherein A is A-1.

Embodiment 8. A compound of Embodiment 6 wherein A is A-2.

Embodiment 9. A compound of Embodiment 6 wherein A is A-6.

Embodiment 10. A compound of Embodiment 6 wherein A is



Embodiment 11. A compound of Embodiment 10 wherein A is A-1a.

Embodiment 12. A compound of Embodiment 10 wherein A is A-2a.

Embodiment 13. A compound of Embodiment 10 wherein A is A-6a.

Embodiment 14. A compound of Formula 1 or any one of Embodiments 1 through 13
 5 either alone or in combination, wherein R^1 is halogen, C_1 - C_4 alkyl or C_1 - C_4
 haloalkyl.

Embodiment 15. A compound of Embodiment 14 wherein R^1 is halogen.

Embodiment 16. A compound of Embodiment 15 wherein R^1 is F, Cl or Br.

Embodiment 17. A compound of Embodiment 16 wherein R^1 is Cl.

10 Embodiment 18. A compound of Formula 1 or any one of Embodiments 1 through 17
 either alone or in combination, wherein each R^2 is independently halogen, cyano,
 SF_5 , C_1 - C_4 alkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 haloalkyl, C_2 - C_4
 haloalkenyl or C_2 - C_4 haloalkynyl.

Embodiment 19. A compound of Embodiment 18 wherein each R^2 is independently
 15 halogen, C_1 - C_4 alkyl or C_1 - C_4 haloalkyl.

Embodiment 20. A compound of Embodiment 19 wherein each R^2 is independently
 halogen, CH_3 or CF_3 .

Embodiment 21. A compound of Embodiment 20 wherein each R^2 is independently
 halogen.

20 Embodiment 22. A compound of Embodiment 21 wherein each R^2 is independently F,
 Cl or Br.

Embodiment 23. A compound of Formula 1 or any one of Embodiments 1 through 22
 either alone or in combination, wherein R^5 is halogen, cyano, CHF_2 or CF_3 .

Embodiment 24. A compound of Embodiment 23 wherein R^5 is F, Cl, Br or cyano.

25 Embodiment 25. A compound of Embodiment 23 wherein R^5 is cyano.

Embodiment 26. A compound of Embodiment 24 wherein R^5 is F, Cl or Br.

Embodiment 27. A compound of Embodiment 23 wherein R^5 is cyano, CHF_2 or CF_3 .

Embodiment 28. A compound of Embodiment 23 wherein R^5 is CHF_2 or CF_3 .

Embodiment 29. A compound of Formula 1 or any one of Embodiments 1 through 28
 30 either alone or in combination, wherein R^6 is H.

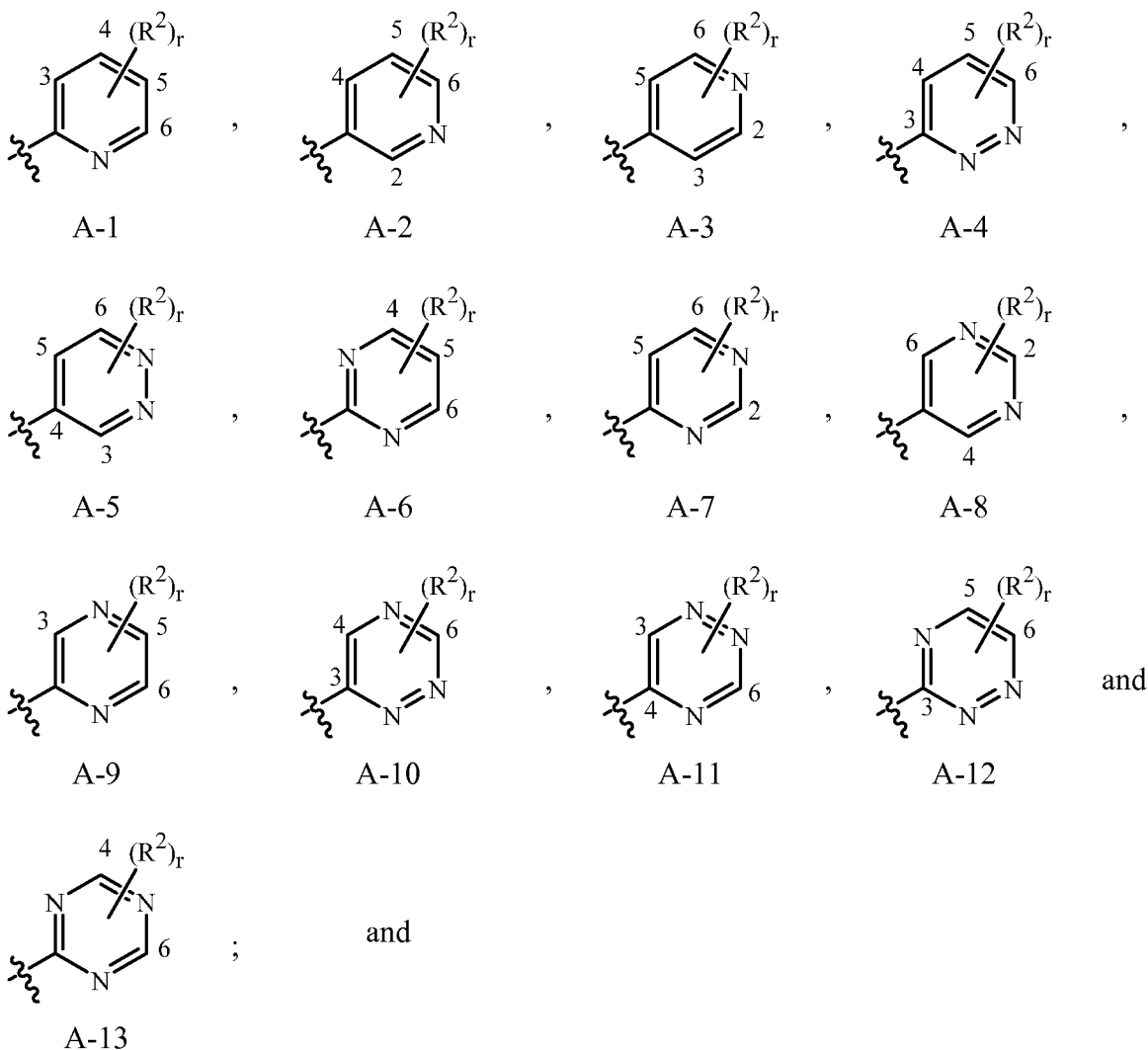
Embodiments of the present invention as described in the Summary of the Invention and Embodiment AAA also include the following:

Embodiment 1P. A compound of Formula 1 wherein A is a phenyl ring optionally substituted with up to 4 R^2 .

5 Embodiment 2P. A compound of Embodiment 1 wherein A is a phenyl ring optionally substituted with up to 2 R^2 .

Embodiment 3P. A compound of Formula 1 wherein A is a 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R^2 .

10 Embodiment 4P. A compound of Embodiment 3 wherein A is selected from



r is 0, 1, 2 or 3.

Embodiment 5P. A compound of Embodiment 4 wherein A is selected from A-1, A-2, A-4, A-6, A-9, A-10, A-11 and A-12.

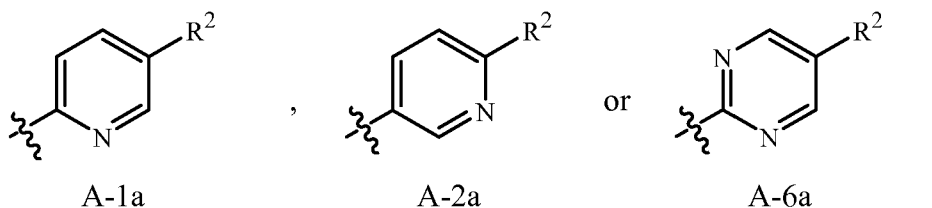
Embodiment 6P. A compound of Embodiment 5 wherein A is selected from A-1, A-2 and A-6.

Embodiment 7P. A compound of Embodiment 6 wherein A is selected from A-1.

Embodiment 8P. A compound of Embodiment 6 wherein A is selected from A-2.

Embodiment 9P. A compound of Embodiment 6 wherein A is selected from A-6.

Embodiment 10P. A compound of Embodiment 6 wherein A is



5 Embodiment 11P. A compound of Embodiment 10 wherein A is A-1a.

Embodiment 12P. A compound of Embodiment 10 wherein A is A-2a.

Embodiment 13P. A compound of Embodiment 10 wherein A is A-6a.

Embodiment 14P. A compound of Formula **1** or any one of Embodiments 1 through 13 wherein R^1 is halogen, C_1-C_4 alkyl or C_1-C_4 haloalkyl.

10 Embodiment 15P. A compound of Embodiment 14 wherein R^1 is halogen.

Embodiment 16P. A compound of Embodiment 15 wherein R^1 is chlorine.

Embodiment 17P. A compound of Formula **1** or any one of Embodiments 1 through 16 wherein each R^2 is independently halogen, C_1-C_4 alkyl or C_1-C_4 haloalkyl.

15 Embodiment 18P. A compound of Embodiment 17 wherein each R^2 is independently halogen, CH_3 or CF_3 .

Embodiment 19P. A compound of Embodiment 18 wherein each R^2 is independently halogen.

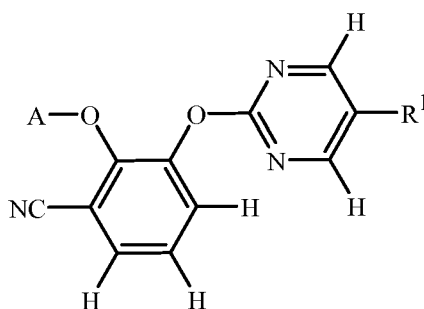
Embodiment 20P. A compound of Embodiment 19 wherein each R^2 is independently F, Cl or Br.

20 Embodiment 21P. A compound of Formula **1** or any one of Embodiments 1 through 20 wherein the phenyl or 6-membered heteroaromatic ring (of A) is substituted with R^2 at the position para to the connection of the ring to the remainder of Formula **1**.

25 Embodiment 22P. A compound of Formula **1** or any one of Embodiments 1 through 21 wherein each R^3 and R^4 is independently C_1-C_4 alkyl.

Also of note is a compound of Formula **1P**

14

**1P**

wherein

A is a phenyl ring optionally substituted with up to 4 R²; or a 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R²;

R¹ is halogen, C₁–C₄ alkyl, C₁–C₄ haloalkyl, C₂–C₆ alkenyl, C₂–C₆ alkynyl, C₁–C₄ alkoxy or S(O)_mR³;

each R² is independently halogen, CHO, C₁–C₄ alkyl, C₁–C₄ haloalkyl, C₁–C₄ cyanoalkyl, C₃–C₆ cycloalkyl, C₂–C₄ alkenyl, C₂–C₄ alkynyl, C₁–C₄ alkoxy, C₃–C₄ alkenyloxy, C₃–C₄ alkynyloxy, C₁–C₄ haloalkoxy, C₁–C₄ hydroxyalkyl, C₂–C₄ alkoxyalkyl, C₂–C₄ alkylthioalkyl, S(O)_nR⁴, C₂–C₆ dialkylamino, CH(=NOH), phenyl or pyridinyl;

each R³ and R⁴ is independently C₁–C₄ alkyl, C₁–C₄ haloalkyl, C₁–C₄ alkylamino or C₂–C₆ dialkylamino;

m is 0, 1 or 2; and

each n is independently 0, 1 or 2.

Embodiments of this invention, including Embodiments 1-29 and 1P-22P above as well as any other embodiments described herein, can be combined in any manner, and the descriptions of variables in the embodiments pertain not only to the a compound of Formula 1 but also to the starting compounds and intermediate compounds useful for preparing the compounds of Formula 1. In addition, embodiments of this invention, including Embodiments 1-29 and 1P-22P above as well as any other embodiments described herein, and any combination thereof, pertain to the compositions and methods of the present invention.

Combination Embodiments illustrated by:

Embodiment AAA. A compound of Formula 1P wherein

A is a phenyl ring optionally substituted with up to 4 R²; or a 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R²;

R^1 is halogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_4 alkoxy or $S(O)_mR^3$;

each R^2 is independently halogen, CHO, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 cyanoalkyl, C_3 - C_6 cycloalkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 alkoxy, C_3 - C_4 alkenyloxy, C_3 - C_4 alkynyloxy, C_1 - C_4 haloalkoxy, C_1 - C_4 hydroxyalkyl, C_2 - C_4 alkoxyalkyl, C_2 - C_4 alkylthioalkyl, $S(O)_nR^4$, C_2 - C_6 dialkylamino, CH(=NOH), phenyl or pyridinyl;

each R^3 and R^4 is independently C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 alkylamino or C_2 - C_6 dialkylamino;

m is 0, 1 or 2; and

each n is independently 0, 1 or 2.

Embodiment AA. A compound of Embodiment AAA or a compound of Formula 1 as described in the Summary of the Invention wherein

A is a phenyl ring optionally substituted with up to 4 R^2 ; or a 5- or 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R^2 ;

R^1 is halogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_4 alkoxy or $S(O)_mR^3$;

each R^2 is independently halogen, cyano, nitro, SF_5 , CHO, $C(=O)NH_2$, $C(=S)NH_2$, SO_2NH_2 , C_1 - C_4 alkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 haloalkyl, C_2 - C_4 haloalkenyl, C_2 - C_4 haloalkynyl, C_3 - C_6 cycloalkyl, C_3 - C_6 halocycloalkyl, C_4 - C_8 alkylcycloalkyl, C_4 - C_8 cycloalkylalkyl, C_2 - C_6 alkylcarbonyl, C_2 - C_6 haloalkylcarbonyl, C_2 - C_6 alkoxy carbonyl, C_3 - C_7 cycloalkylcarbonyl, C_2 - C_8 alkylaminocarbonyl, C_3 - C_{10} dialkylaminocarbonyl, C_1 - C_4 alkoxy, C_3 - C_4 alkenyloxy, C_3 - C_4 alkynyloxy, C_1 - C_4 haloalkoxy, C_3 - C_4 haloalkenyloxy, C_3 - C_4 haloalkynyloxy, C_3 - C_6 cycloalkoxy, C_3 - C_6 halocycloalkoxy, C_4 - C_8 cycloalkylalkoxy, C_2 - C_6 alkoxyalkyl, C_2 - C_6 haloalkoxyalkyl, C_2 - C_6 alkoxyhaloalkyl, C_2 - C_6 alkoxyalkoxy, C_2 - C_4 alkylcarbonyloxy, C_2 - C_6 cyanoalkyl, C_2 - C_6 cyanoalkoxy, C_1 - C_4 hydroxyalkyl, C_2 - C_4 alkylthioalkyl, C_1 - C_6 alkylamino, C_2 - C_6 dialkylamino, $S(O)_nR^4$, CH(=NOH), phenyl or pyridinyl;

each R^3 and R^4 is independently C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 alkylamino or C_2 - C_6 dialkylamino;

R^5 is halogen, cyano or C_1 - C_2 haloalkyl;

R^6 is H or F;

m is 0, 1 or 2; and

each n is independently 0, 1 or 2;

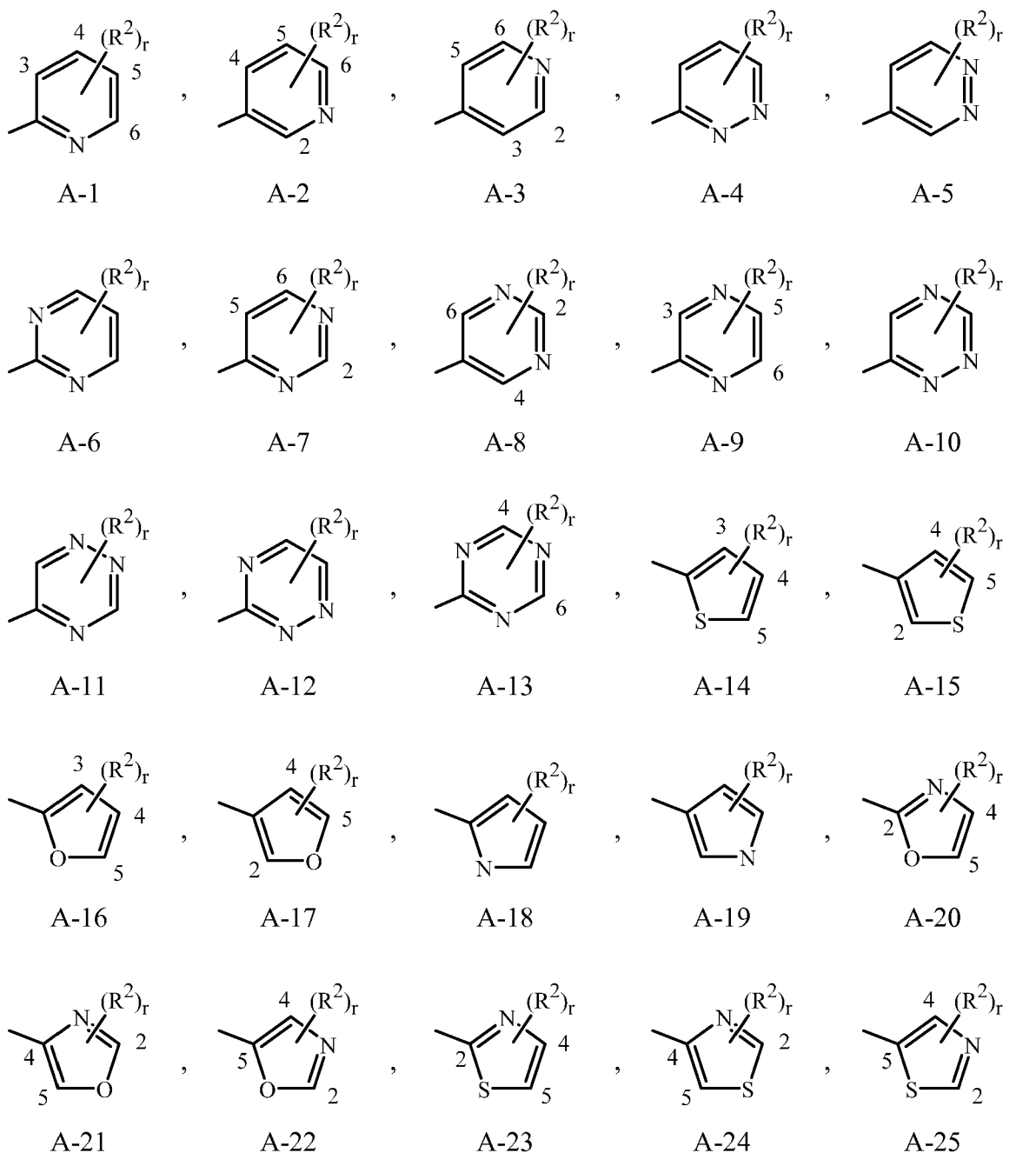
provided the compound of Formula 1 is other than 5-bromo-2-[3-bromo-[2-(5-chloropyridin-2-yloxy)]phenoxy]pyrimidine, 5-bromo-2-[6-bromo-[2-(5-

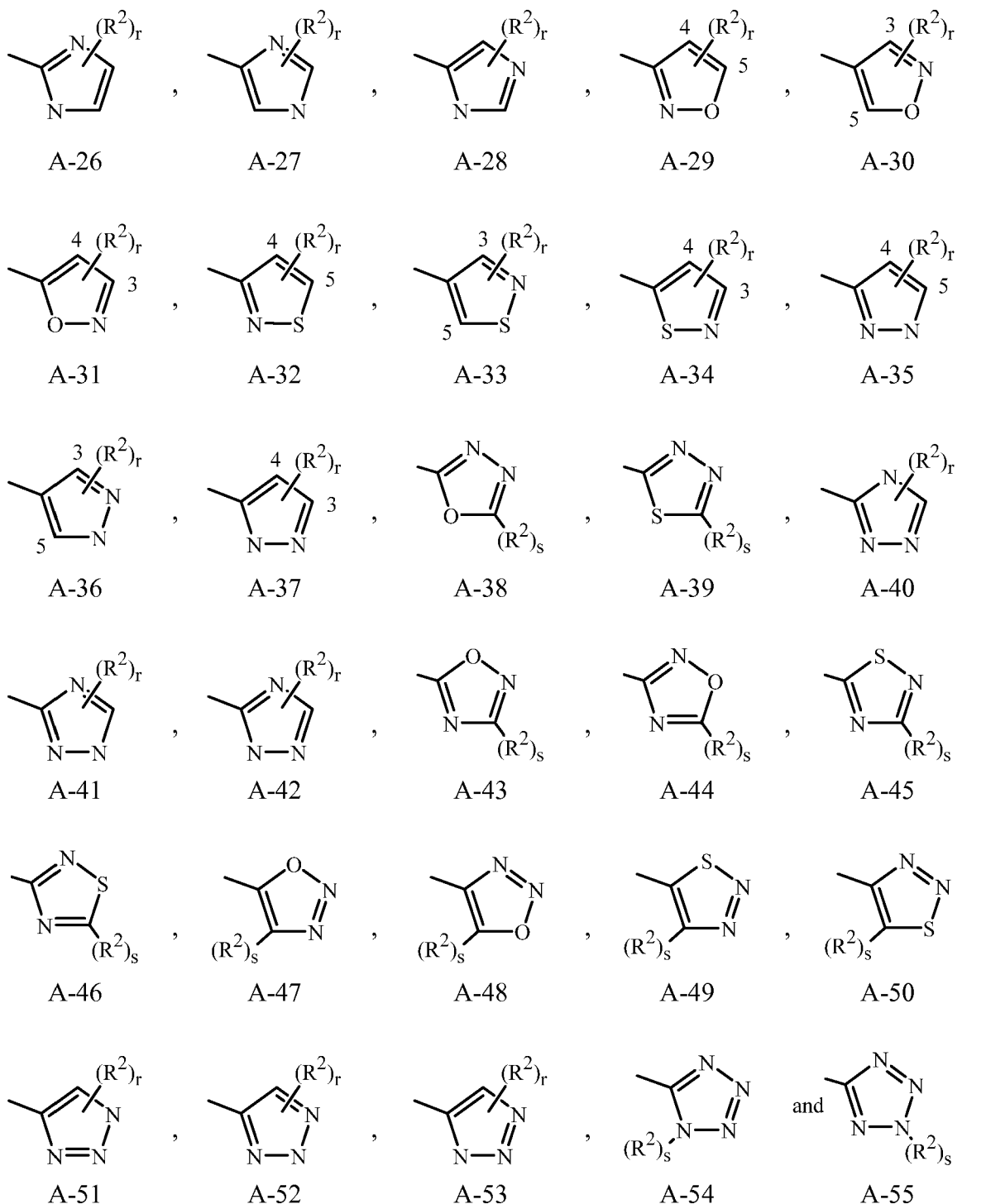
chloropyridin-2-yloxy]phenoxy]pyrimidine, 5-chloro-2-[3-fluoro-[2-(5-chloropyridin-2-yloxy]phenoxy]pyrimidine, 5-chloro-2-[6-fluoro-[2-(5-chloropyridin-2-yloxy]phenoxy]pyrimidine, 5-chloro-2-[3-methyl-[2-(5-chloropyridin-2-yloxy]phenoxy]pyrimidine or 5-chloro-2-[6-methyl-[2-(5-chloropyridin-2-yloxy]phenoxy]pyrimidine.

5

Embodiment A. A compound of Embodiment AA wherein

A is selected from





wherein r is 0, 1, 2 or 3 and s is 0 or 1; and

each R^2 is independently halogen, cyano, SF_5 , C_1 - C_4 alkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 haloalkyl, C_2 - C_4 haloalkenyl or C_2 - C_4 haloalkynyl.

Embodiment AP. A compound of Embodiment AA wherein

- 5 A is a 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R^2 ; and

each R² is independently halogen, C₁–C₄ alkyl or C₁–C₄ haloalkyl.

Embodiment B. A compound of Embodiment A wherein

A is selected from A-1, A-2, A-4, A-6, A-9, A-10, A-11, A-12 and A-23;

R¹ is halogen, C₁–C₄ alkyl or C₁–C₄ haloalkyl; and

5 each R² is independently halogen, C₁–C₄ alkyl or C₁–C₄ haloalkyl.

Embodiment BP. A compound of Embodiment AP wherein

A is selected from A-1, A-2, A-4, A-6, A-9, A-10, A-11 and A-12;

R¹ is halogen; and

each R² is independently halogen, CH₃ or CF₃.

10 Embodiment C. A compound of Embodiment B wherein

A is selected from A-1, A-2 and A-6;

each R² is independently halogen, CH₃ or CF₃;

R⁵ is halogen, cyano, CHF₂ or CF₃; and

R⁶ is H.

15 Embodiment CP. A compound of Embodiment BP wherein

A is selected from A-1, A-2 and A-6.

Embodiment D. A compound of Embodiment C wherein

A is A-6;

R¹ is halogen; and

20 R⁵ is F, Cl, Br or cyano.

Embodiment DP. A compound of Embodiment CP wherein

A is A-6;

R¹ is chlorine; and

each R² is independently halogen.

25 Embodiment E. A compound of Embodiment D wherein

A is A-6a.

Embodiment EP. A compound of Embodiment DP wherein

each R² is independently F, Cl or Br; and

the 6-membered heteroaromatic ring (of A being A-6, i.e. pyrimidinyl ring) is

30 substituted with R² at the position para to the connection of the ring to the remainder of Formula 1.

Specific embodiments include compounds of Formula 1 selected from the group consisting of:

2,3-bis[(5-bromo-2-pyrimidinyl)oxy]benzonitrile (Compound 1);

2,3-bis[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 3);

2,3-bis[(5-fluoro-2-pyrimidinyl)oxy]benzonitrile (Compound 2);

2-[(5-bromo-2-pyrimidinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 4); and

3-[(5-bromo-2-pyrimidinyl)oxy]-2-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 5).

Specific embodiments also include compounds of Formula 1 selected from the group consisting of:

2-[(5-chloro-2-pyridinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 16),
2,2'-[[3-(difluoromethyl)-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine] (Compound 46),
2-[3-bromo-2-[[5-(difluoromethyl)-2-thiazolyl]oxy]phenoxy]-5-chloropyrimidine (Compound 10),
5-chloro-2-[2-fluoro-6-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]phenoxy]pyrimidine (Compound 42),
5-chloro-2-[5-fluoro-6-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]phenoxy]pyrimidine (Compound 43),
5-bromo-2-[2-chloro-6-[(5-chloro-2-pyrimidinyl)oxy]phenoxy]pyrimidine (Compound 17),
5-chloro-2-[5-chloro-6-[(5-fluoro-2-pyrimidinyl)oxy]phenoxy]pyrimidine (Compound 18),
2,2'-[(3,6-difluoro-1,2-phenylene)bis(oxy)]bis[5-fluoropyrimidine] (Compound 29),
5-bromo-2-[2-fluoro-6-[(5-chloro-2-pyrimidinyl)oxy]phenoxy]pyrimidine (Compound 27),
3-[(5-chloro-2-pyrimidinyl)oxy]-2-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]-benzonitrile (Compound 38),
2-[(5-chloro-2-pyrimidinyl)oxy]-3-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]-benzonitrile (Compound 39),
2-[(5-chloro-2-pyrazinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 32),
2,2'-[(3,6-difluoro-1,2-phenylene)bis(oxy)]bis[5-chloropyrimidine] (Compound 34),
2,2'-[[3-fluoro-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine] (Compound 21) and
2,2'-[[3-bromo-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine] (Compound 19).

5 This invention also relates to a method for controlling undesired vegetation comprising applying to the locus of the vegetation herbicidally effective amounts of the compounds of the invention (e.g., as a composition described herein). Of note as embodiments relating to methods of use are those involving the compounds of embodiments described above.

Compounds of the invention are particularly useful for selective control of weeds in crops such as wheat, barley, maize, soybean, sunflower, cotton, oilseed rape and rice, and specialty crops such as sugarcane, citrus, fruit and nut crops.

Also noteworthy as embodiments are herbicidal compositions of the present invention comprising the compounds of embodiments described above.

This invention also includes a herbicidal mixture comprising (a) a compound selected from Formula 1, *N*-oxides, and salts thereof, and (b) at least one additional active ingredient selected from (b1) photosystem II inhibitors, (b2) acetohydroxy acid synthase (AHAS) inhibitors, (b3) acetyl-CoA carboxylase (ACCase) inhibitors, (b4) auxin mimics and (b5) 5-enol-pyruvylshikimate-3-phosphate (EPSP) synthase inhibitors, (b6) photosystem I electron diverters, (b7) protoporphyrinogen oxidase (PPO) inhibitors, (b8) glutamine synthetase (GS) inhibitors, (b9) very long chain fatty acid (VLCFA) elongase inhibitors, (b10) auxin transport inhibitors, (b11) phytoene desaturase (PDS) inhibitors, (b12) 4-hydroxyphenyl-pyruvate dioxygenase (HPPD) inhibitors, (b13) homogentisate solanesyltransferase (HST) inhibitors, (b14) cellulose biosynthesis inhibitors, (b15) other herbicides including mitotic disruptors, organic arsenicals, asulam, bromobutide, cinmethylin, cumyluron, dazomet, difenzoquat, dymron, etobenzanid, flurenol, fosamine, fosamine-ammonium, metam, methyldymron, oleic acid, oxaziclomefone, pelargonic acid and pyributicarb, and (b16) herbicide safeners; and salts of compounds of (b1) through (b16).

“Photosystem II inhibitors” (b1) are chemical compounds that bind to the D-1 protein at the Q_B -binding niche and thus block electron transport from Q_A to Q_B in the chloroplast thylakoid membranes. The electrons blocked from passing through photosystem II are transferred through a series of reactions to form toxic compounds that disrupt cell membranes and cause chloroplast swelling, membrane leakage, and ultimately cellular destruction. The Q_B -binding niche has three different binding sites: binding site A binds the triazines such as atrazine, triazinones such as hexazinone, and uracils such as bromacil, binding site B binds the phenylureas such as diuron, and binding site C binds benzothiadiazoles such as bentazon, nitriles such as bromoxynil and phenyl-pyridazines such as pyridate. Examples of photosystem II inhibitors include ametryn, amicarbazone, atrazine, bentazon, bromacil, bromofenoxim, bromoxynil, chlorbromuron, chloridazon, chlorotoluron, chloroxuron, cumyluron, cyanazine, daimuron, desmedipham, desmetryn, dimefuron, dimethametryn, diuron, ethidimuron, fenuron, fluometuron, hexazinone, ioxynil, isoproturon, isouron, lenacil, linuron, metamitron, methabenzthiazuron, metobromuron, metoxuron, metribuzin, monolinuron, neburon, pentanochlor, phenmedipham, prometon, prometryn, propanil, propazine, pyridafol, pyridate, siduron, simazine, simetryn, tebuthiuron, terbacil, terbumeton, terbuthylazine, terbutryn and trietazine.

“AHAS inhibitors” (b2) are chemical compounds that inhibit acetohydroxy acid synthase (AHAS), also known as acetolactate synthase (ALS), and thus kill plants by inhibiting the production of the branched-chain aliphatic amino acids such as valine, leucine and isoleucine, which are required for protein synthesis and cell growth. Examples of AHAS inhibitors include amidosulfuron, azimsulfuron, bensulfuron-methyl, bispyribac-sodium, cloransulam-methyl, chlorimuron-ethyl, chlorsulfuron, cinosulfuron, cyclosulfamuron, diclosulam, ethametsulfuron-methyl, ethoxysulfuron, flazasulfuron, florasulam, flucarbazone-sodium, flumetsulam, flupyrsulfuron-methyl, flupyrsulfuron-sodium, foramsulfuron, halosulfuron-methyl, imazamethabenz-methyl, imazamox, imazapic, imazapyr, imazaquin, imazethapyr, imazosulfuron, iodosulfuron-methyl (including sodium salt), iofensulfuron (2-iodo-*N*-[[[(4-methoxy-6-methyl-1,3,5-triazin-2-yl)amino]carbonyl]-benzenesulfonamide), mesosulfuron-methyl, metazosulfuron (3-chloro-4-(5,6-dihydro-5-methyl-1,4,2-dioxazin-3-yl)-*N*-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-1-methyl-1*H*-pyrazole-5-sulfonamide), metosulam, metsulfuron-methyl, nicosulfuron, oxasulfuron, penoxsulam, primisulfuron-methyl, propoxycarbazine-sodium, propyrisulfuron (2-chloro-*N*-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-6-propylimidazo[1,2-*b*]pyridazine-3-sulfonamide), prosulfuron, pyrazosulfuron-ethyl, pyribenzoxim, pyriftalid, pyriminobac-methyl, pyriithiobac-sodium, rimsulfuron, sulfometuron-methyl, sulfosulfuron, thiencarbazine, thifensulfuron-methyl, triafamone (*N*-[2-[(4,6-dimethoxy-1,3,5-triazin-2-yl)carbonyl]-6-fluorophenyl]-1,1-difluoro-*N*-methylmethanesulfonamide), triasulfuron, tribenuron-methyl, trifloxysulfuron (including sodium salt), triflusulfuron-methyl and tritosulfuron.

“ACCase inhibitors” (b3) are chemical compounds that inhibit the acetyl-CoA carboxylase enzyme, which is responsible for catalyzing an early step in lipid and fatty acid synthesis in plants. Lipids are essential components of cell membranes, and without them, new cells cannot be produced. The inhibition of acetyl CoA carboxylase and the subsequent lack of lipid production leads to losses in cell membrane integrity, especially in regions of active growth such as meristems. Eventually shoot and rhizome growth ceases, and shoot meristems and rhizome buds begin to die back. Examples of ACCase inhibitors include alloxydim, butoxydim, clethodim, clodinafop, cycloxydim, cyhalofop, diclofop, fenoxaprop, fluazifop, haloxyfop, pinoxaden, profoxydim, propaquizafop, quizalofop, sethoxydim, tepraloxym and tralkoxydim, including resolved forms such as fenoxaprop-P, fluazifop-P, haloxyfop-P and quizalofop-P and ester forms such as clodinafop-propargyl, cyhalofop-butyl, diclofop-methyl and fenoxaprop-P-ethyl.

Auxin is a plant hormone that regulates growth in many plant tissues. “Auxin mimics” (b4) are chemical compounds mimicking the plant growth hormone auxin, thus causing uncontrolled and disorganized growth leading to plant death in susceptible species. Examples of auxin mimics include aminocyclopyrachlor (6-amino-5-chloro-2-cyclopropyl-

4-pyrimidinecarboxylic acid) and its methyl and ethyl esters and its sodium and potassium salts, aminopyralid, benazolin-ethyl, chloramben, clacyfos, clomeprop, clopyralid, dicamba, 2,4-D, 2,4-DB, dichlorprop, fluroxypyr, halauxifen (4-amino-3-chloro-6-(4-chloro-2-fluoro-3-methoxyphenyl)-2-pyridinecarboxylic acid), halauxifen-methyl (methyl 4-amino-3-chloro-6-(4-chloro-2-fluoro-3-methoxyphenyl)-2-pyridinecarboxylate), MCPA, MCPB, mecoprop, picloram, quinclorac, quinmerac, 2,3,6-TBA, triclopyr, and methyl 4-amino-3-chloro-6-(4-chloro-2-fluoro-3-methoxyphenyl)-5-fluoro-2-pyridinecarboxylate.

“EPSP synthase inhibitors” (b5) are chemical compounds that inhibit the enzyme, 5-enol-pyruvylshikimate-3-phosphate synthase, which is involved in the synthesis of aromatic amino acids such as tyrosine, tryptophan and phenylalanine. EPSP inhibitor herbicides are readily absorbed through plant foliage and translocated in the phloem to the growing points. Glyphosate is a relatively nonselective postemergence herbicide that belongs to this group. Glyphosate includes esters and salts such as ammonium, isopropylammonium, potassium, sodium (including sesquisodium) and trimesium (alternatively named sulfosate).

“Photosystem I electron diverters” (b6) are chemical compounds that accept electrons from Photosystem I, and after several cycles, generate hydroxyl radicals. These radicals are extremely reactive and readily destroy unsaturated lipids, including membrane fatty acids and chlorophyll. This destroys cell membrane integrity, so that cells and organelles “leak”, leading to rapid leaf wilting and desiccation, and eventually to plant death. Examples of this second type of photosynthesis inhibitor include diquat and paraquat.

“PPO inhibitors” (b7) are chemical compounds that inhibit the enzyme protoporphyrinogen oxidase, quickly resulting in formation of highly reactive compounds in plants that rupture cell membranes, causing cell fluids to leak out. Examples of PPO inhibitors include acifluorfen-sodium, azafenidin, benzfendizone, bifenox, butafenacil, carfentrazone, carfentrazone-ethyl, chlomethoxyfen, cinidon-ethyl, fluazolate, flufenpyr-ethyl, flumiclorac-pentyl, flumioxazin, fluoroglycofen-ethyl, fluthiacet-methyl, fomesafen, halosafen, lactofen, oxadiargyl, oxadiazon, oxyfluorfen, pentoxazone, proflunazone, pyraclostrobin, pyraflufen-ethyl, saflufenacil, sulfentrazone, thidiazimin, tiafenacil (methyl *N*-[2-[[[2-chloro-5-[3,6-dihydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)-1(2*H*)-pyrimidinyl]-4-fluorophenyl]thio]-1-oxopropyl]-β-alaninate) and 3-[7-fluoro-3,4-dihydro-3-oxo-4-(2-propyn-1-yl)-2*H*-1,4-benzoxazin-6-yl]dihydro-1,5-dimethyl-6-thioxo-1,3,5-triazine-2,4(1*H*,3*H*)-dione.

“GS inhibitors” (b8) are chemical compounds that inhibit the activity of the glutamine synthetase enzyme, which plants use to convert ammonia into glutamine. Consequently, ammonia accumulates and glutamine levels decrease. Plant damage probably occurs due to the combined effects of ammonia toxicity and deficiency of amino acids required for other metabolic processes. The GS inhibitors include glufosinate and its esters and salts such as

glufosinate-ammonium and other phosphinothricin derivatives, glufosinate-P ((2*S*)-2-amino-4-(hydroxymethylphosphinyl)butanoic acid) and bilanaphos.

“VLCFA elongase inhibitors” (b9) are herbicides having a wide variety of chemical structures, which inhibit the elongase. Elongase is one of the enzymes located in or near chloroplasts which are involved in biosynthesis of VLCFAs. In plants, very-long-chain fatty acids are the main constituents of hydrophobic polymers that prevent desiccation at the leaf surface and provide stability to pollen grains. Such herbicides include acetochlor, alachlor, anilofos, butachlor, cafenstrole, dimethachlor, dimethenamid, diphenamid, fenoxasulfone (3-[[[(2,5-dichloro-4-ethoxyphenyl)methyl]sulfonyl]-4,5-dihydro-5,5-dimethylisoxazole), fentrazamide, flufenacet, indanofan, mefenacet, metazachlor, metolachlor, naproanilide, napropamide, napropamide-M ((2*R*)-*N,N*-diethyl-2-(1-naphthalenyloxy)propanamide), pethoxamid, piperophos, pretilachlor, propachlor, propisochlor, pyroxasulfone, and thenylchlor, including resolved forms such as *S*-metolachlor and chloroacetamides and oxyacetamides.

“Auxin transport inhibitors” (b10) are chemical substances that inhibit auxin transport in plants, such as by binding with an auxin-carrier protein. Examples of auxin transport inhibitors include diflufenzopyr, naptalam (also known as *N*-(1-naphthyl)phthalamic acid and 2-[(1-naphthalenylamino)carbonyl]benzoic acid).

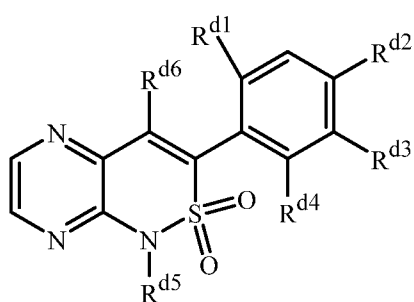
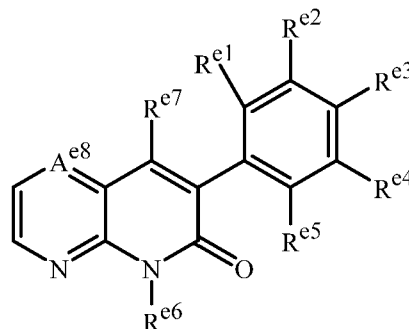
“PDS inhibitors” (b11) are chemical compounds that inhibit carotenoid biosynthesis pathway at the phytoene desaturase step. Examples of PDS inhibitors include beflubutamid, diflufenican, fluridone, flurochloridone, flurtamone norflurzon and picolinafen.

“HPPD inhibitors” (b12) are chemical substances that inhibit the biosynthesis of synthesis of 4-hydroxyphenyl-pyruvate dioxygenase. Examples of HPPD inhibitors include benzobicyclon, benzofenap, bicyclopyrone (4-hydroxy-3-[[2-[(2-methoxyethoxy)methyl]-6-(trifluoromethyl)-3-pyridinyl]carbonyl]bicyclo[3.2.1]oct-3-en-2-one), fenquinotrine (2-[[8-chloro-3,4-dihydro-4-(4-methoxyphenyl)-3-oxo-2-quinoxaliny]carbonyl]-1,3-cyclohexanedione), isoxachlortole, isoxaflutole, mesotrione, pyrasulfotole, pyrazolynate, pyrazoxyfen, sulcotrione, tefuryltrione, tembotrione, topramezone, 5-chloro-3-[(2-hydroxy-6-oxo-1-cyclohexen-1-yl)carbonyl]-1-(4-methoxyphenyl)-2(1*H*)-quinoxalinone, 4-(2,6-diethyl-4-methylphenyl)-5-hydroxy-2,6-dimethyl-3(2*H*)-pyridazinone, 4-(4-fluorophenyl)-6-[(2-hydroxy-6-oxo-1-cyclohexen-1-yl)carbonyl]-2-methyl-1,2,4-triazine-3,5(2*H*,4*H*)-dione, 5-[(2-hydroxy-6-oxo-1-cyclohexen-1-yl)carbonyl]-2-(3-methoxyphenyl)-3-(3-methoxypropyl)-4(3*H*)-pyrimidinone, 2-methyl-*N*-(4-methyl-1,2,5-oxadiazol-3-yl)-3-(methylsulfinyl)-4-(trifluoromethyl)benzamide and 2-methyl-3-(methylsulfonyl)-*N*-(1-methyl-1*H*-tetrazol-5-yl)-4-(trifluoromethyl)benzamide.

“HST inhibitors” (b13) disrupt a plant’s ability to convert homogentisate to 2-methyl-6-solanyl-1,4-benzoquinone, thereby disrupting carotenoid biosynthesis. Examples of HST inhibitors include haloxydine, pyriclor, 3-(2-chloro-3,6-difluorophenyl)-4-

hydroxy-1-methyl-1,5-naphthyridin-2(1*H*)-one, 7-(3,5-dichloro-4-pyridinyl)-5-(2,2-difluoroethyl)-8-hydroxypyrido[2,3-*b*]pyrazin-6(5*H*)-one and 4-(2,6-diethyl-4-methylphenyl)-5-hydroxy-2,6-dimethyl-3(2*H*)-pyridazinone.

HST inhibitors also include compounds of Formulae **A** and **B**.

**A****B**

wherein R^{d1} is H, Cl or CF_3 ; R^{d2} is H, Cl or Br; R^{d3} is H or Cl; R^{d4} is H, Cl or CF_3 ; R^{d5} is CH_3 , CH_2CH_3 or CH_2CHF_2 ; and R^{d6} is OH, or $-OC(=O)-i\text{-Pr}$; and R^{e1} is H, F, Cl, CH_3 or CH_2CH_3 ; R^{e2} is H or CF_3 ; R^{e3} is H, CH_3 or CH_2CH_3 ; R^{e4} is H, F or Br; R^{e5} is Cl, CH_3 , CF_3 , OCF_3 or CH_2CH_3 ; R^{e6} is H, CH_3 , CH_2CHF_2 or $C\equiv CH$; R^{e7} is OH, $-OC(=O)Et$, $-OC(=O)-i\text{-Pr}$ or $-OC(=O)-t\text{-Bu}$; and A^{e8} is N or CH.

“Cellulose biosynthesis inhibitors” (b14) inhibit the biosynthesis of cellulose in certain plants. They are most effective when applied preemergence or early postemergence on young or rapidly growing plants. Examples of cellulose biosynthesis inhibitors include chlorthiamid, dichlobenil, flupoxam, indaziflam (*N*²-[(1*R*,2*S*)-2,3-dihydro-2,6-dimethyl-1*H*-inden-1-yl]-6-(1-fluoroethyl)-1,3,5-triazine-2,4-diamine), isoxaben and triaziflam.

“Other herbicides” (b15) include herbicides that act through a variety of different modes of action such as mitotic disruptors (e.g., flamprop-M-methyl and flamprop-M-isopropyl), organic arsenicals (e.g., DSMA, and MSMA), 7,8-dihydropteroate synthase inhibitors, chloroplast isoprenoid synthesis inhibitors and cell-wall biosynthesis inhibitors. Other herbicides include those herbicides having unknown modes of action or do not fall into a specific category listed in (b1) through (b14) or act through a combination of modes of action listed above. Examples of other herbicides include aclonifen, asulam, amitrole, bromobutide, cinmethylin, clomazone, cumyluron, cyclopyrimorate (6-chloro-3-(2-cyclopropyl-6-methylphenoxy)-4-pyridazinyl 4-morpholinecarboxylate), daimuron, difenzoquat, etobenzanid, fluometuron, flurenol, fosamine, fosamine-ammonium, dazomet, dymron, ipfencarbazone (1-(2,4-dichlorophenyl)-*N*-(2,4-difluorophenyl)-1,5-dihydro-*N*-(1-methylethyl)-5-oxo-4*H*-1,2,4-triazole-4-carboxamide), metam, methyldymron, oleic acid, oxaziclomefone, pelargonic acid, pyributicarb and 5-[[[(2,6-difluorophenyl)methoxy]methyl]-4,5-dihydro-5-methyl-3-(3-methyl-2-thienyl)isoxazole].

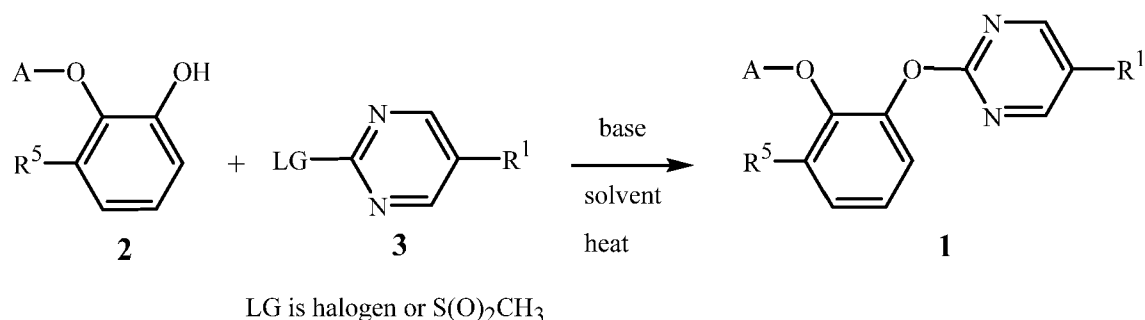
“Herbicide safeners” (b16) are substances added to a herbicide formulation to eliminate or reduce phytotoxic effects of the herbicide to certain crops. These compounds protect crops from injury by herbicides but typically do not prevent the herbicide from controlling undesired vegetation. Examples of herbicide safeners include but are not limited to benoxacor, cloquintocet-mexyl, cumyluron, cyometrinil, cyprosulfamide, daimuron, dichlormid, dicyclonon, dimepiperate, fenchlorazole-ethyl, fenclorim, flurazole, fluxofenim, furilazole, isoxadifen-ethyl, mefenpyr-diethyl, mephenate, methoxyphenone, naphthalic anhydride, oxabetrinil, *N*-(aminocarbonyl)-2-methylbenzenesulfonamide and *N*-(aminocarbonyl)-2-fluorobenzenesulfonamide, 1-bromo-4-[(chloromethyl)sulfonyl]benzene, 2-(dichloromethyl)-2-methyl-1,3-dioxolane (MG 191), 4-(dichloroacetyl)-1-oxa-4-azospiro[4.5]decane (MON 4660).

An embodiment of the present invention is a herbicidal mixture comprising (a) a compound of Formula 1, and (b) at least one additional active ingredient selected from (b1) photosystem II inhibitors, (b2) acetohydroxy acid synthase (AHAS) inhibitors, (b4) auxin mimics, (b5) 5-enol-pyruvylshikimate-3-phosphate (EPSP) synthase inhibitors, (b7) protoporphyrinogen oxidase (PPO) inhibitors, (b9) very long chain fatty acid (VLCFA) elongase inhibitors and (b12) 4-hydroxyphenyl-pyruvate dioxygenase (HPPD) inhibitors.

The compounds of Formula 1 can be prepared by general methods known in the art of synthetic organic chemistry. One or more of the following methods and variations as described in Schemes 1–8 can be used to prepare the compounds of Formula 1. The definitions of R¹ and A in the compounds of Formulae 1–14 below are as defined above in the Summary of the Invention unless otherwise noted. Formula 1a is a subset of Formula 1, and all substituents for Formula 1a are as defined above for Formula 1 unless otherwise noted. Formulae 5a and 5b are subsets of Formula 5, and all substituents for Formulae 5a and 5b are as defined for Formula 5 unless otherwise noted.

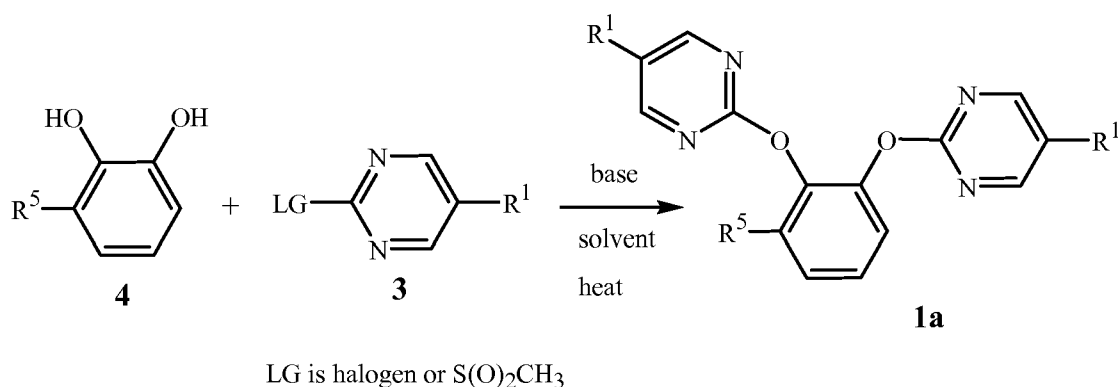
As shown in Scheme 1, a compound of Formula 1 (wherein R⁶ is H) can be prepared by nucleophilic substitution by heating a phenolic intermediate of Formula 2 in a suitable solvent, such as acetonitrile, tetrahydrofuran or *N,N*-dimethylformamide, in the presence of a base, such as potassium or cesium carbonate, with a compound of Formula 3 (where LG is a nucleophilic reaction leaving group, i.e. nucleofuge, such as halogen or S(O)₂CH₃). The reaction is typically conducted at temperatures ranging from 50 to 110 °C.

Scheme 1



As shown in Scheme 2, a compound of Formula **1** (i.e. Formula **1** wherein A is 5- R^1 -pyrimidin-2-yl) can also be prepared by coupling the compound of Formula **4** with two equivalents of a compound of Formula **3** under the same conditions as described for Scheme 1. The method of Scheme 2 is illustrated by Synthesis Example 1.

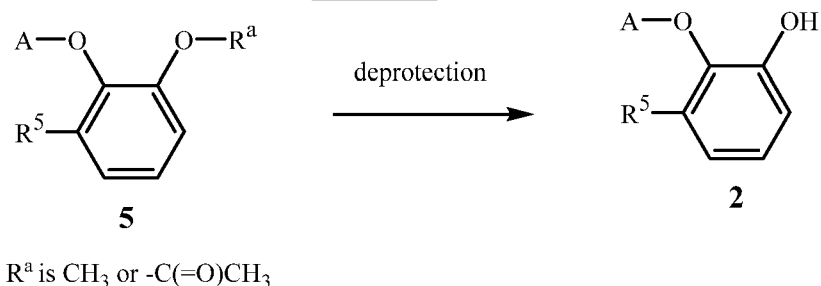
Scheme 2



As shown in Scheme 3, a compound of Formula **2** can be prepared by deprotection of a compound of Formula **5** wherein R^a is CH_3 or $-(C=O)CH_3$ with a suitable deprotecting agent. A suitable deprotecting agent for methoxy in a compound of Formula **5** (i.e. R^a is CH_3), such as BBr_3 , $AlCl_3$, Me_3SiI and HBr in acetic acid, can be used in the presence of solvents such as toluene, dichloromethane and dichloroethane at a temperature ranging from -80 to 120 °C. A suitable deprotecting agent for acetoxy in a compound Formula **5** (i.e. R^a is $-C(=O)CH_3$), such as potassium carbonate in methanol or ammonium acetate in aqueous methanol at room temperature can be used as discussed in Das et al., *Tetrahedron* **2003**, 59, 1049–1054 and methods cited therein. Alternatively, a compound of Formula **5** wherein R^a is $-C(=O)CH_3$ can be combined with Amberlyst® 15 in methanol (as discussed in Das et al. *Tet. Lett.* **2003**, 44, 5465–5468) or combined with sodium acetate in ethanol (as discussed in T. Narender et al. *Synthetic Communications* **2009**, 39(11), 1949–1956) to obtain a compound of Formula **2**. Other useful phenolic protecting groups suitable for use in preparing a compound of Formula **2** can be found in Greene, T. W.; Wuts, P. G. M. *Protective Groups in Organic Synthesis*, 4th ed.; Wiley: Hoboken, New Jersey, 2012).

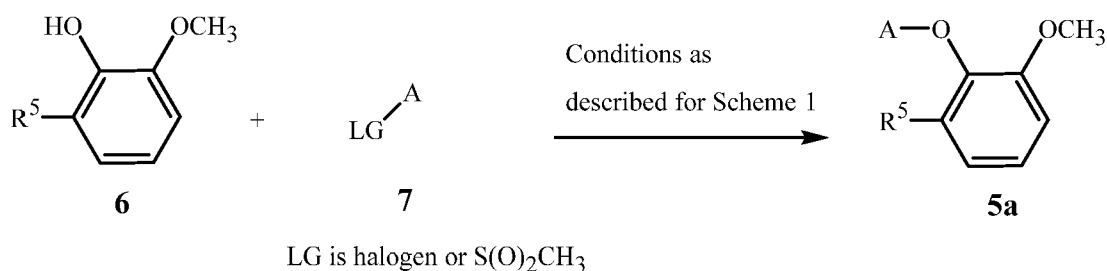
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Scheme 3

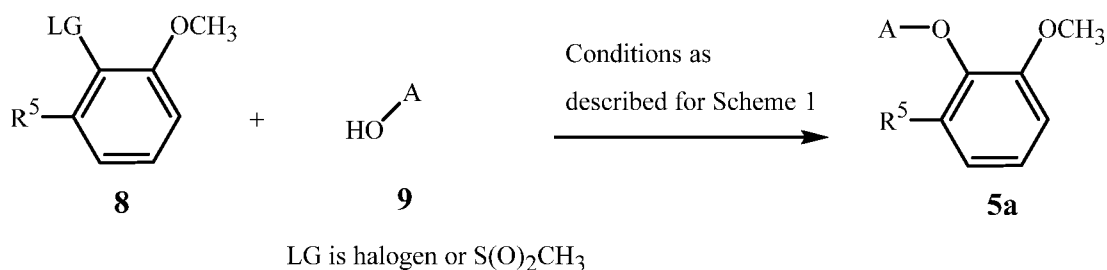


An intermediate of Formula **5a** (i.e. Formula **5** wherein R^a is CH_3) can be prepared by a variety of methods known to one skilled in the art. As shown in Scheme 4 and Scheme 5 by selecting appropriate coupling partners, e.g., compounds of Formulae **6** and **7** or compounds of Formulae **8** and **9**, compounds of Formula **5a** can be obtained by simple substitution using the conditions described for Scheme 1.

Scheme 4

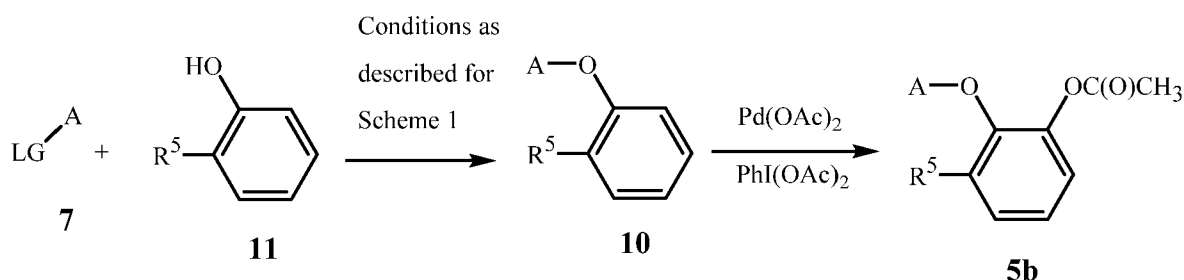


Scheme 5



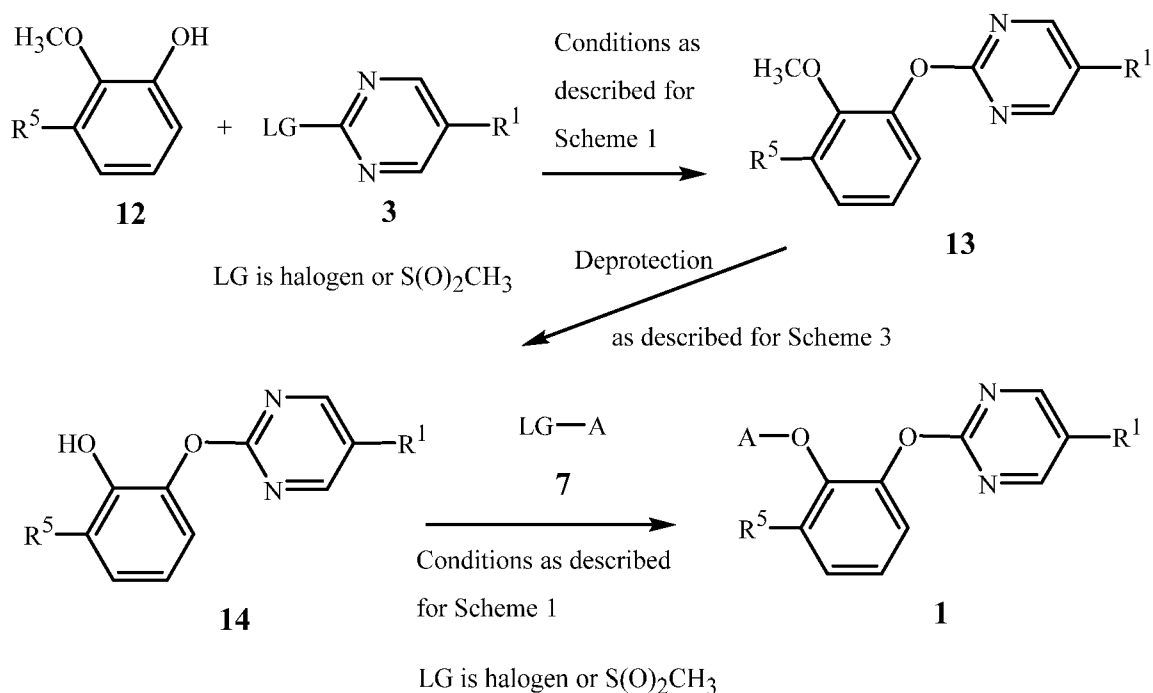
As shown in Scheme 6, a compound of Formula **5b** (i.e. Formula **5** wherein R^a is $-\text{C}(=\text{O})\text{CH}_3$) can be prepared from an intermediate of Formula **10** by “C-H activation”, utilizing palladium(II) acetate and (diacetoxyiodo)benzene. Typical procedures for this type of reaction are described, for example, in *J. Org. Chem.* **2009**, *74*, 7203. The intermediate of Formula **10** can be prepared by nucleophilic substitution reaction of a compound of Formula **7** with a compound of Formula **11** under the conditions described for Scheme 1.

Scheme 6



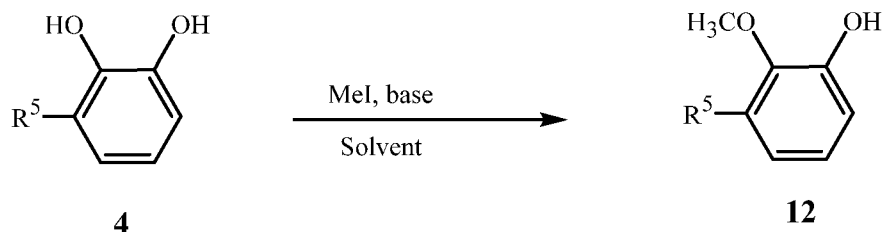
One skilled in the art will recognize that a compound of Formula 1 can also be constructed using the sequence shown in Scheme 7. In the first step of this method, the methoxyphenol of Formula 12 is reacted with a compound of Formula 3 using reaction conditions as described for Scheme 1 to provide the methoxyphenyl ether of Formula 13. In the next step, the methyl group is removed using deprotection conditions described for Scheme 3 to provide the phenol ether of Formula 14, which in the last step is reacted with a compound of Formula 7 using reaction conditions as described for Scheme 1 to provide the compound of Formula 1. The first step of the method of Scheme 7 is illustrated by Step A of Synthesis Example 2. The second step of the method of Scheme 7 is illustrated by Step B of Synthesis Example 2. The final step of the method of Scheme 7 is illustrated by Step C of Synthesis Example 2.

Scheme 7



As shown in Scheme 8, the compound of Formula 12 can be prepared by selective methylation of the compound of Formula 4.

Scheme 8



Compounds of Formulae **3**, **4**, **6**, **7**, **8** and **9** can be synthesized according to general methods known in the art of synthetic organic chemistry. Furthermore, some of the starting materials, such as the compound of Formula **4**, are commercially available.

It is recognized by one skilled in the art that various functional groups can be converted into others to provide different compounds of Formula **1**. For a valuable resource that illustrates the interconversion of functional groups in a simple and straightforward fashion, see Larock, R. C., *Comprehensive Organic Transformations: A Guide to Functional Group Preparations*, 2nd Ed., Wiley-VCH, New York, 1999. For example, intermediates for the preparation of compounds of Formula **1** may contain aromatic nitro groups, which can be reduced to amino groups, and then be converted via reactions well known in the art such as the Sandmeyer reaction, to various halides, providing compounds of Formula **1**. The above reactions can also in many cases be performed in alternate order.

It is recognized that some reagents and reaction conditions described above for preparing compounds of Formula **1** may not be compatible with certain functionalities present in the intermediates. In these instances, the incorporation of protection/deprotection sequences or functional group interconversions into the synthesis will aid in obtaining the desired products. The use and choice of the protecting groups will be apparent to one skilled in chemical synthesis (see, for example, Greene, T. W.; Wuts, P. G. M. *Protective Groups in Organic Synthesis*, 2nd ed.; Wiley: New York, 1991). One skilled in the art will recognize that, in some cases, after the introduction of a given reagent as depicted in any individual scheme, it may be necessary to perform additional routine synthetic steps not described in detail to complete the synthesis of compounds of Formula **1**. One skilled in the art will also recognize that it may be necessary to perform a combination of the steps illustrated in the above schemes in an order other than that implied by the particular presented to prepare the compounds of Formula **1**.

One skilled in the art will also recognize that compounds of Formula **1** and the intermediates described herein can be subjected to various electrophilic, nucleophilic, radical, organometallic, oxidation, and reduction reactions to add substituents or modify existing substituents.

Without further elaboration, it is believed that one skilled in the art using the preceding description can utilize the present invention to its fullest extent. The following non-limiting

Examples are illustrative of the invention. Steps in the following Examples illustrate a procedure for each step in an overall synthetic transformation, and the starting material for each step may not have necessarily been prepared by a particular preparative run whose procedure is described in other Examples or Steps. Percentages are by weight except for chromatographic solvent mixtures or where otherwise indicated. Parts and percentages for chromatographic solvent mixtures are by volume unless otherwise indicated. ¹H NMR spectra are reported in ppm downfield from tetramethylsilane in CDCl₃ unless otherwise noted; “s” means singlet, “d” means doublet, “dd” means doublet of doublets, and “m” means multiplet. Mass spectra (MS) are reported as the molecular weight of the highest isotopic abundance parent ion (M+1) formed by addition of H⁺ (molecular weight of 1) to the molecule, or (M–1) formed by the loss of H⁺ (molecular weight of 1) from the molecule, observed by using liquid chromatography coupled to a mass spectrometer (LCMS) using either atmospheric pressure chemical ionization (AP⁺), where “amu” stands for unified atomic mass units.

SYNTHESIS EXAMPLE 1

Preparation of 2,3-bis[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 3)

2,3-Dihydroxybenzonitrile (270 mg, 2 mmol) and 2,5-dichloropyrimidine (655 mg, 4.4 mmol) were combined in *N,N*-dimethylformamide (6 mL) under a nitrogen atmosphere. Powdered potassium carbonate (1.2 g, 8.8 mmol) was added, and the resulting mixture was heated at 100 °C for 8 h. The reaction mixture was cooled and diluted with water and ethyl acetate. The aqueous layer was separated and extracted with ethyl acetate (3×). The combined organic layers were washed with brine, dried (MgSO₄), filtered and concentrated under reduced pressure. The residue was purified by medium pressure liquid chromatography on silica gel, eluted with 0 to 15% ethyl acetate in hexanes, to yield the title product, a compound of the present invention, as a solid (640 mg).

¹H NMR (400 MHz, CDCl₃) δ 7.42–7.49 (m, 1H), 7.57 (dd, *J*=8.31, 1.47 Hz, 1H), 7.65 (dd, *J*=7.83, 1.96 Hz, 1H), 8.42 (m, 4H).

SYNTHESIS EXAMPLE 2

Preparation of 2-[(5-bromo-2-pyrimidinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile (Compound 4)

Step A: Preparation of 3-[(5-chloro-2-pyrimidinyl)oxy]-2-methoxybenzonitrile

3-Hydroxy-2-methoxybenzonitrile (730 mg, 4.9 mmol) and 2,5-dichloropyrimidine (803 mg, 5.4 mmol) were combined in acetonitrile (10 mL) under a nitrogen atmosphere. Powdered potassium carbonate (1.48 g, 10.7 mmol) was added, and the resulting mixture was heated at 80 °C for 1 h. The reaction mixture was cooled and concentrated under reduced pressure. The residue was purified by medium pressure liquid chromatography on silica gel eluted with 0 to 20% ethyl acetate in hexanes to yield the title compound (1 g).

MS(AP⁺) 262 amu (M+1).

Step B: Preparation of 3-[(5-chloro-2-pyrimidinyl)oxy]-2-hydroxybenzonitrile

3-[(5-Chloro-2-pyrimidinyl)oxy]-2-methoxybenzonitrile (i.e. the product of Step A) (1.00 g, 3.82 mmol) was dissolved in dichloromethane (5 mL) and cooled to 0 °C. Then boron tribromide (1 M in CH₂Cl₂, 19.1 mL, 19.1 mmol) was added to the solution, and the mixture was stirred at room temperature for 3 h. The reaction mixture was quenched by adding saturated aqueous NaHCO₃ solution at 0 °C. The aqueous layer was separated and extracted with dichloromethane (2×). The combined organic layers were washed with brine, dried (MgSO₄), filtered and concentrated under reduced pressure. The residue was used in the next step without further purification.

MS(AP⁺) 246 amu (M-1).

Step C: Preparation of 2-[(5-bromo-2-pyrimidinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)-oxy]benzonitrile

3-[(5-Chloro-2-pyrimidinyl)oxy]-2-hydroxybenzonitrile (i.e. the product of Step B) (150 mg, 0.6 mmol) and 2-chloro-5-bromopyrimidine (128 mg, 0.66 mmol) were combined in *N,N*-dimethylformamide (6 mL) under a nitrogen atmosphere. Powdered potassium carbonate (182 mg, 1.32 mmol) was added, and the resulting mixture was heated at 80 °C for 12 h. The reaction mixture was cooled and diluted with water and ethyl acetate. The aqueous layer was separated and extracted with ethyl acetate (3×). The combined organic layers were washed with brine, dried (MgSO₄), filtered and concentrated under reduced pressure. The residue was purified by medium pressure liquid chromatography on silica gel, eluted with 0 to 20% ethyl acetate in hexanes, to yield the title product, a compound of the present invention, as a solid (70 mg).

¹H NMR (400 MHz, CDCl₃) δ 7.45 (m, 1H), 7.57 (m, 1H), 7.65 (dd, *J*=7.83, 1.47 Hz, 1H), 8.42 (m, 2H), 8.51 (m, 2H).

SYNTHESIS EXAMPLE 3

Preparation of 5-chloro-2-[5-fluoro-2-[5-(trifluoromethyl)pyridin-2-yl]oxy]phenoxy]pyrimidine, (Compound 45)

Step A: Preparation of 2-(2-fluorophenoxy)-5-(trifluoromethyl)pyridine

A solution of 2-fluorophenol (0.94 g, 8.39 mmoles) in DMF (20 mL) was stirred under a nitrogen atmosphere. Powdered potassium carbonate (2.9 g, 20.98 mmoles) was added followed by 2-chloro-5-(trifluoromethyl)pyridine (1.6 g, 8.82 mmoles) and copper(I) chloride 0.42 g, 4.2 mmoles). The reaction mixture was heated at 110 °C for 2 hours and allowed to cool to room temperature overnight. The mixture was diluted with deionized water and diethyl ether, partitioned, and the aqueous phase was extracted twice with diethyl ether. The combined organic phases were washed twice with saturated aqueous ethylenediamine tetraacetic acid solution, once with 1N aqueous sodium hydroxide and once

with saturated aqueous sodium chloride solution. The organic phases were then dried over magnesium sulfate and concentrated to give the title compound (1.98 g) as an oil.

^1H NMR (400 MHz, CDCl_3) δ 8.41 (s, 1H), 7.94(d, 1H), 7.18-7.25 (m, 4H), 7.08 (d, 1H).

Step B: Preparation of 3-fluoro-2-[[5-(trifluoromethyl)-2-pyridinyl]oxy]phenol 1-acetate

A solution of 2-(2-fluorophenoxy)-5-(trifluoromethyl)pyridine (i.e. the product of Step A) (2.0 g, 7.78 mmol) in a 1:1 mix of acetic anhydride and acetic acid (26 mL each) was treated with diacetoxyiodobenzene (5.0 g, 15.56 mmol) and palladium acetate (0.08 g, 0.38 mmol). The reaction mixture was then heated at 100 °C for 4 hours and allowed to cool to room temperature overnight. The mixture was diluted with toluene and concentrated under vacuum. The residue was partitioned between ethyl acetate and saturated aqueous sodium bicarbonate solution. The phases were separated and the aqueous phase was extracted with ethyl acetate. The combined organic phases were washed with saturated aqueous sodium chloride solution, dried over magnesium sulfate, and concentrated to an oil (3 g). The crude oil was purified by flash column chromatography with a 40 gram Isco MPLC silica gel column using 0-30% ethyl acetate/hexanes gradient to give the title compound (1.38 g) as an oil.

^1H NMR (400 MHz, CDCl_3) δ 8.40 (s, 1H), 7.94 (d, 1H), 7.25 (m, 1H), 7.11 (m, 2H), 7.04 (d, 1H), 2.16 (s, 3H).

Step C: Preparation of 3-fluoro-2-[[5-(trifluoromethyl)-2-pyridinyl]oxy]phenol

A solution of 3-fluoro-2-[[5-(trifluoromethyl)-2-pyridinyl]oxy]phenol 1-acetate (i.e. the product of Step B) (1.36 g, 4.31 mmol) in 32 mL of methanol and 10 mL of deionized water was treated with ammonium acetate (2.66 g, 34.5 mmol) then stirred at room temperature overnight. The mixture was treated with additional ammonium acetate (1 g) and stirred at room temperature for another 24 hours. The reaction mixture was concentrated under vacuum, partitioned between ethyl acetate and water and the phases separated. The aqueous phase was extracted with ethyl acetate and the combined organic phases were washed with saturated aqueous sodium chloride solution, dried over magnesium sulfate and concentrated. The crude oil was purified by flash column chromatography with a 12 gram Isco MPLC silica gel column using 0-10% ethyl acetate/hexanes gradient to give the title compound (0.39 g) as an oil.

^1H NMR (400 MHz, CDCl_3) δ 8.46 (s, 1H), 7.99 (d, 1H), 7.22 (d, 1H), 7.12 (m, 1H), 6.89 (d, 1H), 6.77 (m, 1H), 6.29 (s, 1H).

Step D: Preparation of 5-chloro-2-[5-fluoro-[2-[5-(trifluoromethyl)pyridin-2-yl]oxy]phenoxy]pyrimidine

A solution of 3-fluoro-2-[[5-(trifluoromethyl)-2-pyridinyl]oxy]phenol (i.e. the product of Step C) (0.16 g, 0.585 mmol) in 2 mL of acetonitrile was treated with powdered potassium carbonate (0.2 g, 1.45 mmol) and 2,5-dichloropyrimidine (0.07 g, 0.47 mmol). The reaction mixture was heated at 80 °C for 4.5 hours. The mixture was cooled, diluted with de-ionized water and ethyl acetate and the phases separated. The aqueous phase was extracted with ethyl acetate and the combined organic phases were washed with saturated aqueous sodium chloride solution, dried over magnesium sulfate and concentrated under vacuum. The crude oil was purified by flash column chromatography with a 12 gram Isco MPLC silica gel column using 0-10% ethyl acetate/hexanes gradient to give the title compound, a compound of the present invention, as an oil (0.20 g).

¹H NMR (400 MHz, CDCl₃) δ 8.41 (s, 2H), 8.34 (s, 1H), 7.85 (m, 1H), 7.31 (m, 1H), 7.13 (m, 2H), 6.93 (d, 1H).

SYNTHESIS EXAMPLE 4

Preparation of 2-[3-bromo-2-[[5-(difluoromethyl)-2-thiazolyl]oxy]phenoxy]-5-chloropyrimidine (Compound 10)

Step A: Preparation of 2-(2-bromo-6-methoxyphenoxy)-5-thiazolecarboxaldehyde

To a solution of 2-bromo-6-methoxyphenol (5.0 g, 24.63 mmol) in *N,N'*-dimethylformamide (50 mL) was added potassium carbonate (6.8 g, 48.6 mmol) and 2-chloro-5-thiazolecarboxaldehyde (3.6 g, 24.63 mmol) at 0 °C. The reaction mixture was stirred at ambient temperature for 12 hours. The reaction mixture was poured into water (100 mL) and extracted with ethyl acetate (3 x 100 mL). The combined organic phases were washed with water followed by saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate and concentrated under reduced pressure. The crude residue was purified by column chromatography using ethyl acetate: hexanes (1:5) to afford the title compound (5.2 g) as pale yellow solid.

¹H NMR (400 MHz, CDCl₃) δ 3.80 (s, 3H), 6.98-7.00 (dd, 1H), 7.16-7.20 (t, 1H), 7.23-7.26 (dd, 1H), 7.87 (s, 1H), 9.84 (s, 1H).

Step B: Preparation of 2-(2-bromo-6-hydroxyphenoxy)-5-thiazolecarboxaldehyde

To a solution of 2-(2-bromo-6-methoxyphenoxy)-5-thiazolecarboxaldehyde (i.e. the product of Step A) (2.0 g, 6.36 mmol) in dichloromethane (20 mL) was added a 1 M solution of boron tribromide in dichloromethane (12.7 mL, 12.73 mmol) at 0 °C. The reaction mixture was stirred at ambient temperature for 5 hours. The reaction mixture was poured into ice water (30 mL) and extracted with dichloromethane (50 mL). The combined organic phases were washed with saturated sodium bicarbonate solution (20 mL) and water (20 mL),

dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The crude residue was purified by column chromatography using ethyl acetate: hexanes (1: 4) to afford the title compound (0.9 g) as pale yellow solid.

Mass spectrum = 299.8

- 5 Step C: Preparation of 2-[2-bromo-6-(5-chloro-2-pyrimidinyl)oxy]phenoxy]-5-thiazolecarboxaldehyde
-

To a solution of 2-(2-bromo-6-hydroxyphenoxy)-5-thiazolecarboxaldehyde (0.7 g, 2.33 mmol) (i.e. the product of Step B) in *N,N'*-dimethylformamide (10 mL) was added potassium carbonate (0.64 g, 4.66 mmol) and 5-chloro-2-(methylsulfonyl)pyrimidine (0.45 g, 2.33 mmol). The reaction mixture was stirred at 50 °C for 16 hours. The reaction mixture was poured into water (50 mL) and extracted with ethyl acetate (3 x 50mL). The combined organic phases were washed with water followed by saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate and concentrated under reduced pressure to afford the crude title compound. The crude compound was directly used for next step.

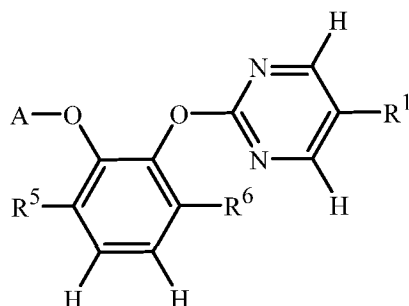
- 15 Step D: Preparation of 2-[3-bromo-2-[[5-(difluoromethyl)-2-thiazolyl]oxy]phenoxy]-5-chloropyrimidine
-

To a solution of 2-[2-bromo-6-(5-chloro-2-pyrimidinyl)oxy]phenoxy]-5-thiazolecarboxaldehyde (i.e. the product of Step C) (0.55 g, 1.33 mmol) in dichloromethane (10 mL) was added diethylaminosulfur trifluoride (0.5mL, 4.01 mmol) at 0 °C and the reaction mixture was stirred at ambient temperature for 16 hours. The reaction mixture was diluted with dichloromethane (50 mL) and washed with water (20 mL). The organic phase was separated, dried over anhydrous sodium sulfate and concentrated under reduced pressure. The crude residue was purified by column chromatography using ethyl acetate: hexanes (1: 4) to afford the title compound, a compound of the present invention, (30 mg) as off white solid.

¹H NMR (400 MHz, DMSO-*d*₆) δ 7.13 -7.40 (t, CHF₂), 7.42-7.46 (t, 1H), 7.54-7.61 (m, 2H), 7.73 -7.76(dd, 1H), 8.77 (s, 2H).

By the procedures described herein together with methods known in the art, the following compounds of Tables 1–15 can be prepared. The following abbreviations are used in the Tables which follow: Me means methyl, Et means ethyl, Pr means propyl, *i*-Pr means isopropyl, Ph means phenyl, OMe means methoxy, CN means cyano, NO₂ means nitro, and S(O)₂Me means methylsulfonyl. In the names of some of the heterocycles forming substituent A the locant number is alternatively inserted before “yl”, instead of prefixing the heterocycle name. For example, “pyridin-2-yl” means the same as “2-pyridinyl”, and “pyrimidin-5-yl” means the same as “5-pyrimidinyl”.

Table 1



R^1 is F, R^5 is Cl and R^6 is H

A	A	A
pyridin-2-yl	3-Me-pyridin-4-yl	2-OMe-pyrimidin-5-yl
5-F-pyridin-2-yl	pyridazin-3-yl	2-CF ₃ -pyrimidin-5-yl
5-Cl-pyridin-2-yl	6-F-pyridazin-3-yl	2-OCF ₃ -pyrimidin-5-yl
5-Br-pyridin-2-yl	6-Cl-pyridazin-3-yl	2-CHF ₂ -pyrimidin-5-yl
5-I-pyridin-2-yl	6-Br-pyridazin-3-yl	2-CN-pyrimidin-5-yl
5-Me-pyridin-2-yl	6-I-pyridazin-3-yl	pyrazin-2-yl
5-OMe-pyridin-2-yl	6-Me-pyridazin-3-yl	5-F-pyrazin-2-yl
5-CF ₃ -pyridin-2-yl	6-OMe-pyridazin-3-yl	5-Cl-pyrazin-2-yl
5-OCF ₃ -pyridin-2-yl	6-CF ₃ -pyridazin-3-yl	5-Me-pyrazin-2-yl
5-CHF ₂ -pyridin-2-yl	6-OCF ₃ -pyridazin-3-yl	5-CF ₃ -pyrazin-2-yl
5-CN-pyridin-2-yl	5-F-pyridazin-3-yl	6-F-pyrazin-2-yl
5-CHO-pyridin-2-yl	5-Cl-pyridazin-3-yl	6-Cl-pyrazin-2-yl
5-S(O) ₂ Me-pyridin-2-yl	5-Br-pyridazin-3-yl	6-Me-pyrazin-2-yl
5-NO ₂ -pyridin-2-yl	5-I-pyridazin-3-yl	6-CF ₃ -pyrazin-2-yl
4-F-pyridin-2-yl	5-Me-pyridazin-3-yl	3-Cl-pyrazin-2-yl
4-Cl-pyridin-2-yl	5-OMe-pyridazin-3-yl	3-Me-pyrazin-2-yl
4-Br-pyridin-2-yl	5-CF ₃ -pyridazin-3-yl	1,2,4-triazin-6-yl
4-I-pyridin-2-yl	5-OCF ₃ -pyridazin-3-yl	3-F-1,2,4-triazin-6-yl
4-Me-pyridin-2-yl	4-Cl-pyridazin-3-yl	3-Cl-1,2,4-triazin-6-yl
4-OMe-pyridin-2-yl	pyridazin-4-yl	3-Me-1,2,4-triazin-6-yl
4-CF ₃ -pyridin-2-yl	6-F-pyridazin-4-yl	3-CF ₃ -1,2,4-triazin-6-yl
4-OCF ₃ -pyridin-2-yl	6-Cl-pyridazin-4-yl	5-F-1,2,4-triazin-6-yl
4-CHF ₂ -pyridin-2-yl	6-Br-pyridazin-4-yl	5-Cl-1,2,4-triazin-6-yl
4-CN-pyridin-2-yl	6-I-pyridazin-4-yl	5-Me-1,2,4-triazin-6-yl
4-CHO-pyridin-2-yl	6-Me-pyridazin-4-yl	5-CF ₃ -1,2,4-triazin-6-yl
4-S(O) ₂ Me-pyridin-2-yl	6-OMe-pyridazin-4-yl	1,2,4-triazin-5-yl
4-NO ₂ -pyridin-2-yl	6-CF ₃ -pyridazin-4-yl	3-F-1,2,4-triazin-5-yl

A	A	A
3-F-pyridin-2-yl	6-OCF ₃ -pyridazin-4-yl	3-Cl-1,2,4-triazin-5-yl
3-Cl-pyridin-2-yl	5-Cl-pyridazin-4-yl	3-Me-1,2,4-triazin-5-yl
3-Br-pyridin-2-yl	3-Cl-pyridazin-4-yl	3-CF ₃ -1,2,4-triazin-5-yl
3-I-pyridin-2-yl	pyrimidin-2-yl	6-F-1,2,4-triazin-5-yl
3-Me-pyridin-2-yl	5-F-pyrimidin-2-yl	6-Cl-1,2,4-triazin-5-yl
3-OMe-pyridin-2-yl	5-Cl-pyrimidin-2-yl	6-Me-1,2,4-triazin-5-yl
3-CF ₃ -pyridin-2-yl	5-Br-pyrimidin-2-yl	6-CF ₃ -1,2,4-triazin-5-yl
3-OCF ₃ -pyridin-2-yl	5-I-pyrimidin-2-yl	1,2,4-triazin-3-yl
6-F-pyridin-2-yl	5-F-pyrimidin-2-yl	6-F-1,2,4-triazin-3-yl
6-Cl-pyridin-2-yl	5-Cl-pyrimidin-2-yl	6-Cl-1,2,4-triazin-3-yl
6-Br-pyridin-2-yl	5-Br-pyrimidin-2-yl	6-Br-1,2,4-triazin-3-yl
6-I-pyridin-2-yl	5-I-pyrimidin-2-yl	6-I-1,2,4-triazin-3-yl
6-Me-pyridin-2-yl	5-Me-pyrimidin-2-yl	6-Me-1,2,4-triazin-3-yl
6-OMe-pyridin-2-yl	5-OMe-pyrimidin-2-yl	6-CF ₃ -1,2,4-triazin-3-yl
6-CF ₃ -pyridin-2-yl	5-CF ₃ -pyrimidin-2-yl	6-OCF ₃ -1,2,4-triazin-3-yl
6-OCF ₃ -pyridin-2-yl	5-OCF ₃ -pyrimidin-2-yl	6-CN-1,2,4-triazin-3-yl
pyridin-3-yl	5-CHF ₂ -pyrimidin-2-yl	1,3,5-triazin-2-yl
6-F-pyridin-3-yl	5-CN-pyrimidin-2-yl	4-Cl-1,3,5-triazin-2-yl
6-Cl-pyridin-3-yl	5-CHO-pyrimidin-2-yl	4-Me-1,3,5-triazin-2-yl
6-Br-pyridin-3-yl	5-S(O) ₂ Me-pyrimidin-2-yl	4-CF ₃ -1,3,5-triazin-2-yl
6-I-pyridin-3-yl	5-NO ₂ -pyrimidin-2-yl	Ph
6-Me-pyridin-3-yl	4-Cl-pyrimidin-2-yl	4-F-Ph
6-OMe-pyridin-3-yl	4-Me-pyrimidin-2-yl	4-Cl-Ph
6-CF ₃ -pyridin-3-yl	4,6-di-Cl-pyrimidin-2-yl	4-Br-Ph
6-OCF ₃ -pyridin-3-yl	4,6-di-Me-pyrimidin-2-yl	4-I-Ph
5-F-pyridin-3-yl	pyrimidin-4-yl	4-Me-Ph
5-Cl-pyridin-3-yl	6-F-pyrimidin-4-yl	4-CF ₃ -Ph
5-Br-pyridin-3-yl	6-Cl-pyrimidin-4-yl	4-OCF ₃ -Ph
5-I-pyridin-3-yl	6-Br-pyrimidin-4-yl	4-CN-Ph
5-Me-pyridin-3-yl	6-I-pyrimidin-4-yl	3-F-Ph
5-OMe-pyridin-3-yl	6-Me-pyrimidin-4-yl	3-Cl-Ph
5-CF ₃ -pyridin-3-yl	6-OMe-pyrimidin-4-yl	3-Br-Ph
5-OCF ₃ -pyridin-3-yl	6-CF ₃ -pyrimidin-4-yl	3-I-Ph
4-Cl-pyridin-3-yl	6-OCF ₃ -pyrimidin-4-yl	3-Me-Ph
2-Cl-pyridin-3-yl	6-CHF ₂ -pyrimidin-4-yl	3-CF ₃ -Ph
pyridin-4-yl	6-CN-pyrimidin-4-yl	3-OCF ₃ -Ph
2-F-pyridin-4-yl	5-Cl-pyrimidin-4-yl	3-CN-Ph

A	A	A
2-Cl-pyridin-4-yl	5-Me-pyrimidin-4-yl	3,5-di-F-Ph
2-Br-pyridin-4-yl	2-Me-pyrimidin-4-yl	3,5-di-Cl-Ph
2-I-pyridin-4-yl	pyrimidin-5-yl	3,5-di-Br-Ph
2-Me-pyridin-4-yl	2-F-pyrimidin-5-yl	3,5-di-I-Ph
2-OMe-pyridin-4-yl	2-Cl-pyrimidin-5-yl	3,5-di-Me-Ph
2-CF ₃ -pyridin-4-yl	2-Br-pyrimidin-5-yl	3,5-di-CF ₃ -Ph
2-OCF ₃ -pyridin-4-yl	2-I-pyrimidin-5-yl	3,5-di-OCF ₃ -Ph
3-Cl-pyridin-4-yl	2-Me-pyrimidin-5-yl	3,5-di-CN-Ph
5-CF ₃ -thiazol-2-yl	4-I-oxazol-2-yl	5-CF ₃ -furan-2-yl
5-CHF ₂ -thiazol-2-yl	4-OMe-oxazol-2-yl	5-CHF ₂ -furan-2-yl
5-CH ₃ -thiazol-2-yl	4-OCF ₃ -oxazol-2-y	5-CH ₃ -furan-2-yl
5-Cl-thiazol-2-yl	4-CN-oxazol-2-yl	5-Cl-furan-2-yl
5-Br-thiazol-2-yl	5-CF ₃ -thiophene-2-yl	5-Br-furan-2-yl
5-I-thiazol-2-yl	5-CHF ₂ -thiophene-2-yl	5-I-furan-2-yl
5-OMe-thiazol-2-yl	5-CH ₃ -thiophene-2-yl	5-OMe-furan-2-yl
5-OCF ₃ -thiazol-2-y	5-Cl-thiophene-2-yl	5-OCF ₃ -furan-2-y
5-CN-thiazol-2-yl	5-Br-thiophene-2-yl	5-CN-furan-2-yl
4-CF ₃ -thiazol-2-yl	5-I-thiophene-2-yl	2-CF ₃ -thiazol-4-yl
4-CHF ₂ -thiazol-2-yl	5-OMe-thiophene-2-yl	2-CHF ₂ -thiazol-4-yl
4-CH ₃ -thiazol-2-yl	5-OCF ₃ -thiophene-2-y	2-CH ₃ -thiazol-4-yl
4-Cl-thiazol-2-yl	5-CN-thiophene-2-yl	2-Cl-thiazol-4-yl
4-Br-thiazol-2-yl	4-CF ₃ -thiophene-2-yl	2-Br-thiazol-4-yl
4-I-thiazol-2-yl	4-CHF ₂ -thiophene-2-yl	2-I-thiazol-4-yl
4-OMe-thiazol-2-yl	4-CH ₃ -thiophene-2-yl	2-OMe-thiazol-4-yl
4-OCF ₃ -thiazol-2-y	4-Cl-thiophene-2-yl	2-OCF ₃ -thiazol-4-y
4-CN-thiazol-2-yl	4-Br-thiophene-2-yl	2-CN-thiazol-4-yl
5-CF ₃ -oxazol-2-yl	4-I-thiophene-2-yl	2-CF ₃ -thiazol-5-yl
5-CHF ₂ -oxazol-2-yl	4-OMe-thiophene-2-yl	2-CHF ₂ -thiazol-5-yl
5-CH ₃ -oxazol-2-yl	4-OCF ₃ -thiophene-2-y	2-CH ₃ -thiazol-5-yl
5-Cl-oxazol-2-yl	4-CN-thiophene-2-yl	2-Cl-thiazol-5-yl
5-Br-oxazol-2-yl	5-CF ₃ -thiophene-5-yl	2-Br-thiazol-5-yl
5-I-oxazol-2-yl	5-CHF ₂ -thiophene-5-yl	2-I-thiazol-5-yl
5-OMe-oxazol-2-yl	5-CH ₃ -thiophene-5-yl	2-OMe-thiazol-5-yl
5-OCF ₃ -oxazol-2-y	5-Cl-thiophene-5-yl	2-OCF ₃ -thiazol-5-y
5-CN-oxazol-2-yl	5-Br-thiophene-5-yl	2-CN-thiazol-5-yl
4-CF ₃ -oxazol-2-yl	5-I-thiophene-5-yl	4-CF ₃ -imidazol-2-yl
4-CHF ₂ -oxazol-2-yl	5-OMe-thiophene-5-yl	3-CF ₃ -1,2,4-oxadiazol-5-yl

A	A	A
4-CH ₃ -oxazol-2-yl	5-OCF ₃ -thiophene-5-yl	3-CF ₃ -1,2,4-thiadiazol-5-yl
4-Cl-oxazol-2-yl	5-CN-thiophene-5-yl	
4-Br-oxazol-2-yl		

The present disclosure also includes Tables 2 through 165. Each Table is constructed in the same manner as Table 1 above, except that the row heading in Table 1 (i.e. “R¹ is F, R⁵ is Cl and R⁶ is H”) is replaced with the respective row heading shown below. For example, the first entry in Table 2 is a compound of Formula 1 wherein R¹ is F, R⁵ is F, R⁶ is H and A is pyridin-2-yl. Tables 3 through 165 are constructed similarly.

5

Header Row				Header Row			
Table	R ¹	R ⁵	R ⁶	Table	R ¹	R ⁵	R ⁶
2	F	F	H	84	CF ₃	CF ₂	H
3	F	F	F	85	CF ₃	CF ₃	H
4	F	Br	H	86	CF ₃	CHF ₂	H
5	F	I	H	87	CF ₃	CH ₂ CF ₃	H
6	F	CN	H	88	CF ₃	CF ₂ CF ₃	H
7	F	CF ₂	H	89	CHF ₂	F	H
8	F	CF ₃	H	90	CHF ₂	F	F
9	F	CHF ₂	H	91	CHF ₂	Cl	H
10	F	CH ₂ CF ₃	H	92	CHF ₂	Br	H
11	F	CF ₂ CF ₃	H	93	CHF ₂	I	H
12	Cl	F	H	94	CHF ₂	CN	H
13	Cl	F	F	95	CHF ₂	CF ₂	H
14	Cl	Cl	H	96	CHF ₂	CF ₃	H
15	Cl	Br	H	97	CHF ₂	CHF ₂	H
16	Cl	I	H	98	CHF ₂	CH ₂ CF ₃	H
17	Cl	CN	H	99	CHF ₂	CF ₂ CF ₃	H
18	Cl	CF ₂	H	100	CH ₂ F	F	H
19	Cl	CF ₃	H	101	CH ₂ F	F	F
20	Cl	CHF ₂	H	102	CH ₂ F	Cl	H
21	Cl	CH ₂ CF ₃	H	103	CH ₂ F	Br	H
22	Cl	CF ₂ CF ₃	H	104	CH ₂ F	I	H
23	Br	F	H	105	CH ₂ F	CN	H
24	Br	F	F	106	CH ₂ F	CF ₂	H
25	Br	Cl	H	107	CH ₂ F	CF ₃	H
26	Br	Br	H	108	CH ₂ F	CHF ₂	H
27	Br	I	H	109	CH ₂ F	CH ₂ CF ₃	H
28	Br	CN	H	110	CH ₂ F	CF ₂ CF ₃	H

Table	Header Row			Table	Header Row		
	R ¹	R ⁵	R ⁶		R ¹	R ⁵	R ⁶
29	Br	CF ₂	H	111	CH ₂ CF ₃	F	H
30	Br	CF ₃	H	112	CH ₂ CF ₃	F	F
31	Br	CHF ₂	H	113	CH ₂ CF ₃	Cl	H
32	Br	CH ₂ CF ₃	H	114	CH ₂ CF ₃	Br	H
33	Br	CF ₂ CF ₃	H	115	CH ₂ CF ₃	I	H
34	I	F	H	116	CH ₂ CF ₃	CN	H
35	I	F	F	117	CH ₂ CF ₃	CF ₂	H
36	I	Cl	H	118	CH ₂ CF ₃	CF ₃	H
37	I	Br	H	119	CH ₂ CF ₃	CHF ₂	H
38	I	I	H	120	CH ₂ CF ₃	CH ₂ CF ₃	H
39	I	CN	H	121	CH ₂ CF ₃	CF ₂ CF ₃	H
40	I	CF ₂	H	122	C≡CH	F	H
41	I	CF ₃	H	123	C≡CH	F	F
42	I	CHF ₂	H	124	C≡CH	Cl	H
43	I	CH ₂ CF ₃	H	125	C≡CH	Br	H
44	I	CF ₂ CF ₃	H	126	C≡CH	I	H
45	Me	F	H	127	C≡CH	CN	H
46	Me	F	F	128	C≡CH	CF ₂	H
47	Me	Cl	H	129	C≡CH	CF ₃	H
48	Me	Br	H	130	C≡CH	CHF ₂	H
49	Me	I	H	131	C≡CH	CH ₂ CF ₃	H
50	Me	CN	H	132	C≡CH	CF ₂ CF ₃	H
51	Me	CF ₂	H	133	OMe	F	H
52	Me	CF ₃	H	134	OMe	F	F
53	Me	CHF ₂	H	135	OMe	Cl	H
54	Me	CH ₂ CF ₃	H	136	OMe	Br	H
55	Me	CF ₂ CF ₃	H	137	OMe	I	H
56	Et	F	H	138	OMe	CN	H
57	Et	F	F	139	OMe	CF ₂	H
58	Et	Cl	H	140	OMe	CF ₃	H
59	Et	Br	H	141	OMe	CHF ₂	H
60	Et	I	H	142	OMe	CH ₂ CF ₃	H
61	Et	CN	H	143	OMe	CF ₂ CF ₃	H
62	Et	CF ₂	H	144	OEt	F	H
63	Et	CF ₃	H	145	OEt	F	F
64	Et	CHF ₂	H	146	OEt	Cl	H

Table	Header Row			Table	Header Row		
	R ¹	R ⁵	R ⁶		R ¹	R ⁵	R ⁶
65	Et	CH ₂ CF ₃	H	147	OEt	Br	H
66	Et	CF ₂ CF ₃	H	148	OEt	I	H
67	i-Pr	F	H	149	OEt	CN	H
68	i-Pr	F	F	150	OEt	CF ₂	H
69	i-Pr	Cl	H	151	OEt	CF ₃	H
70	i-Pr	Br	H	152	OEt	CHF ₂	H
71	i-Pr	I	H	153	OEt	CH ₂ CF ₃	H
72	i-Pr	CN	H	154	OEt	CF ₂ CF ₃	H
73	i-Pr	CF ₂	H	155	SO ₂ Me	F	H
74	i-Pr	CF ₃	H	156	SO ₂ Me	F	F
75	i-Pr	CHF ₂	H	157	SO ₂ Me	Cl	H
76	i-Pr	CH ₂ CF ₃	H	158	SO ₂ Me	Br	H
77	i-Pr	CF ₂ CF ₃	H	159	SO ₂ Me	I	H
78	CF ₃	F	H	160	SO ₂ Me	CN	H
79	CF ₃	F	F	161	SO ₂ Me	CF ₂	H
80	CF ₃	Cl	H	162	SO ₂ Me	CF ₃	H
81	CF ₃	Br	H	163	SO ₂ Me	CHF ₂	H
82	CF ₃	I	H	164	SO ₂ Me	CH ₂ CF ₃	H
83	CF ₃	CN	H	165	SO ₂ Me	CF ₂ CF ₃	H

Formulation/Utility

A compound of this invention will generally be used as a herbicidal active ingredient in a composition, i.e. formulation, with at least one additional component selected from the group consisting of surfactants, solid diluents and liquid diluents, which serves as a carrier.

- 5 The formulation or composition ingredients are selected to be consistent with the physical properties of the active ingredient, mode of application and environmental factors such as soil type, moisture and temperature.

Useful formulations include both liquid and solid compositions. Liquid compositions include solutions (including emulsifiable concentrates), suspensions, emulsions (including microemulsions, oil-in-water emulsions, flowable concentrates and/or suspoemulsions) and the like, which optionally can be thickened into gels. The general types of aqueous liquid compositions are soluble concentrate, suspension concentrate, capsule suspension, concentrated emulsion, microemulsion, oil-in-water emulsion, flowable concentrate and suspo-emulsion. The general types of nonaqueous liquid compositions are emulsifiable concentrate, microemulsifiable concentrate, dispersible concentrate and oil dispersion.

The general types of solid compositions are dusts, powders, granules, pellets, prills, pastilles, tablets, filled films (including seed coatings) and the like, which can be water-dispersible (“wetable”) or water-soluble. Films and coatings formed from film-forming solutions or flowable suspensions are particularly useful for seed treatment. Active ingredient can be (micro)encapsulated and further formed into a suspension or solid formulation; alternatively the entire formulation of active ingredient can be encapsulated (or “overcoated”). Encapsulation can control or delay release of the active ingredient. An emulsifiable granule combines the advantages of both an emulsifiable concentrate formulation and a dry granular formulation. High-strength compositions are primarily used as intermediates for further formulation.

Sprayable formulations are typically extended in a suitable medium before spraying. Such liquid and solid formulations are formulated to be readily diluted in the spray medium, usually water, but occasionally another suitable medium like an aromatic or paraffinic hydrocarbon or vegetable oil. Spray volumes can range from about one to several thousand liters per hectare, but more typically are in the range from about ten to several hundred liters per hectare. Sprayable formulations can be tank mixed with water or another suitable medium for foliar treatment by aerial or ground application, or for application to the growing medium of the plant. Liquid and dry formulations can be metered directly into drip irrigation systems or metered into the furrow during planting.

The formulations will typically contain effective amounts of active ingredient, diluent and surfactant within the following approximate ranges which add up to 100 percent by weight.

	Weight Percent		
	<u>Active Ingredient</u>	<u>Diluent</u>	<u>Surfactant</u>
Water-Dispersible and Water-soluble Granules, Tablets and Powders	0.001–90	0–99.999	0–15
Oil Dispersions, Suspensions, Emulsions, Solutions (including Emulsifiable Concentrates)	1–50	40–99	0–50
Dusts	1–25	70–99	0–5
Granules and Pellets	0.001–99	5–99.999	0–15
High Strength Compositions	90–99	0–10	0–2

Solid diluents include, for example, clays such as bentonite, montmorillonite, attapulgite and kaolin, gypsum, cellulose, titanium dioxide, zinc oxide, starch, dextrin, sugars (e.g., lactose, sucrose), silica, talc, mica, diatomaceous earth, urea, calcium carbonate, sodium carbonate and bicarbonate, and sodium sulfate. Typical solid diluents are described

in Watkins et al., *Handbook of Insecticide Dust Diluents and Carriers*, 2nd Ed., Dorland Books, Caldwell, New Jersey.

Liquid diluents include, for example, water, *N,N*-dimethylalkanamides (e.g., *N,N*-dimethylformamide), limonene, dimethyl sulfoxide, *N*-alkylpyrrolidones (e.g., *N*-methylpyrrolidinone), alkyl phosphates (e.g., triethyl phosphate), ethylene glycol, triethylene glycol, propylene glycol, dipropylene glycol, polypropylene glycol, propylene carbonate, butylene carbonate, paraffins (e.g., white mineral oils, normal paraffins, isoparaffins), alkylbenzenes, alkylnaphthalenes, glycerine, glycerol triacetate, sorbitol, aromatic hydrocarbons, dearomatized aliphatics, alkylbenzenes, alkylnaphthalenes, ketones such as cyclohexanone, 2-heptanone, isophorone and 4-hydroxy-4-methyl-2-pentanone, acetates such as isoamyl acetate, hexyl acetate, heptyl acetate, octyl acetate, nonyl acetate, tridecyl acetate and isobornyl acetate, other esters such as alkylated lactate esters, dibasic esters, alkyl and aryl benzoates and γ -butyrolactone, and alcohols, which can be linear, branched, saturated or unsaturated, such as methanol, ethanol, *n*-propanol, isopropyl alcohol, *n*-butanol, isobutyl alcohol, *n*-hexanol, 2-ethylhexanol, *n*-octanol, decanol, isodecyl alcohol, isooctadecanol, cetyl alcohol, lauryl alcohol, tridecyl alcohol, oleyl alcohol, cyclohexanol, tetrahydrofurfuryl alcohol, diacetone alcohol, cresol and benzyl alcohol. Liquid diluents also include glycerol esters of saturated and unsaturated fatty acids (typically C₆–C₂₂), such as plant seed and fruit oils (e.g., oils of olive, castor, linseed, sesame, corn (maize), peanut, sunflower, grapeseed, safflower, cottonseed, soybean, rapeseed, coconut and palm kernel), animal-sourced fats (e.g., beef tallow, pork tallow, lard, cod liver oil, fish oil), and mixtures thereof. Liquid diluents also include alkylated fatty acids (e.g., methylated, ethylated, butylated) wherein the fatty acids may be obtained by hydrolysis of glycerol esters from plant and animal sources, and can be purified by distillation. Typical liquid diluents are described in Marsden, *Solvents Guide*, 2nd Ed., Interscience, New York, 1950.

The solid and liquid compositions of the present invention often include one or more surfactants. When added to a liquid, surfactants (also known as “surface-active agents”) generally modify, most often reduce, the surface tension of the liquid. Depending on the nature of the hydrophilic and lipophilic groups in a surfactant molecule, surfactants can be useful as wetting agents, dispersants, emulsifiers or defoaming agents.

Surfactants can be classified as nonionic, anionic or cationic. Nonionic surfactants useful for the present compositions include, but are not limited to: alcohol alkoxylates such as alcohol alkoxylates based on natural and synthetic alcohols (which may be branched or linear) and prepared from the alcohols and ethylene oxide, propylene oxide, butylene oxide or mixtures thereof; amine ethoxylates, alkanolamides and ethoxylated alkanolamides; alkoxylated triglycerides such as ethoxylated soybean, castor and rapeseed oils; alkylphenol alkoxylates such as octylphenol ethoxylates, nonylphenol ethoxylates, dinonyl phenol

ethoxylates and dodecyl phenol ethoxylates (prepared from the phenols and ethylene oxide, propylene oxide, butylene oxide or mixtures thereof); block polymers prepared from ethylene oxide or propylene oxide and reverse block polymers where the terminal blocks are prepared from propylene oxide; ethoxylated fatty acids; ethoxylated fatty esters and oils; 5 ethoxylated methyl esters; ethoxylated tristyrylphenol (including those prepared from ethylene oxide, propylene oxide, butylene oxide or mixtures thereof); fatty acid esters, glycerol esters, lanolin-based derivatives, polyethoxylate esters such as polyethoxylated sorbitan fatty acid esters, polyethoxylated sorbitol fatty acid esters and polyethoxylated glycerol fatty acid esters; other sorbitan derivatives such as sorbitan esters; polymeric 10 surfactants such as random copolymers, block copolymers, alkyd peg (polyethylene glycol) resins, graft or comb polymers and star polymers; polyethylene glycols (pegs); polyethylene glycol fatty acid esters; silicone-based surfactants; and sugar-derivatives such as sucrose esters, alkyl polyglycosides and alkyl polysaccharides.

Useful anionic surfactants include, but are not limited to: alkylaryl sulfonic acids and 15 their salts; carboxylated alcohol or alkylphenol ethoxylates; diphenyl sulfonate derivatives; lignin and lignin derivatives such as lignosulfonates; maleic or succinic acids or their anhydrides; olefin sulfonates; phosphate esters such as phosphate esters of alcohol alkoxylates, phosphate esters of alkylphenol alkoxylates and phosphate esters of styryl phenol ethoxylates; protein-based surfactants; sarcosine derivatives; styryl phenol ether 20 sulfate; sulfates and sulfonates of oils and fatty acids; sulfates and sulfonates of ethoxylated alkylphenols; sulfates of alcohols; sulfates of ethoxylated alcohols; sulfonates of amines and amides such as *N,N*-alkyltaurates; sulfonates of benzene, cumene, toluene, xylene, and dodecyl and tridecylbenzenes; sulfonates of condensed naphthalenes; sulfonates of naphthalene and alkyl naphthalene; sulfonates of fractionated petroleum; sulfosuccinamates; 25 and sulfosuccinates and their derivatives such as dialkyl sulfosuccinate salts.

Useful cationic surfactants include, but are not limited to: amides and ethoxylated amides; amines such as *N*-alkyl propanediamines, tripropylenetriamines and dipropylenetetramines, and ethoxylated amines, ethoxylated diamines and propoxylated amines (prepared from the amines and ethylene oxide, propylene oxide, butylene oxide or 30 mixtures thereof); amine salts such as amine acetates and diamine salts; quaternary ammonium salts such as quaternary salts, ethoxylated quaternary salts and diquaternary salts; and amine oxides such as alkyldimethylamine oxides and bis-(2-hydroxyethyl)-alkylamine oxides.

Also useful for the present compositions are mixtures of nonionic and anionic 35 surfactants or mixtures of nonionic and cationic surfactants. Nonionic, anionic and cationic surfactants and their recommended uses are disclosed in a variety of published references including *McCutcheon's Emulsifiers and Detergents*, annual American and International Editions published by McCutcheon's Division, The Manufacturing Confectioner Publishing

Co.; Sisely and Wood, *Encyclopedia of Surface Active Agents*, Chemical Publ. Co., Inc., New York, 1964; and A. S. Davidson and B. Milwidsky, *Synthetic Detergents*, Seventh Edition, John Wiley and Sons, New York, 1987.

Compositions of this invention may also contain formulation auxiliaries and additives, known to those skilled in the art as formulation aids (some of which may be considered to also function as solid diluents, liquid diluents or surfactants). Such formulation auxiliaries and additives may control: pH (buffers), foaming during processing (antifoams such as polyorganosiloxanes), sedimentation of active ingredients (suspending agents), viscosity (thixotropic thickeners), in-container microbial growth (antimicrobials), product freezing (antifreezes), color (dyes/pigment dispersions), wash-off (film formers or stickers), evaporation (evaporation retardants), and other formulation attributes. Film formers include, for example, polyvinyl acetates, polyvinyl acetate copolymers, polyvinylpyrrolidone-vinyl acetate copolymer, polyvinyl alcohols, polyvinyl alcohol copolymers and waxes. Examples of formulation auxiliaries and additives include those listed in *McCutcheon's Volume 2: Functional Materials*, annual International and North American editions published by McCutcheon's Division, The Manufacturing Confectioner Publishing Co.; and PCT Publication WO 03/024222.

The compound of Formula 1 and any other active ingredients are typically incorporated into the present compositions by dissolving the active ingredient in a solvent or by grinding in a liquid or dry diluent. Solutions, including emulsifiable concentrates, can be prepared by simply mixing the ingredients. If the solvent of a liquid composition intended for use as an emulsifiable concentrate is water-immiscible, an emulsifier is typically added to emulsify the active-containing solvent upon dilution with water. Active ingredient slurries, with particle diameters of up to 2,000 μm can be wet milled using media mills to obtain particles with average diameters below 3 μm . Aqueous slurries can be made into finished suspension concentrates (see, for example, U.S. 3,060,084) or further processed by spray drying to form water-dispersible granules. Dry formulations usually require dry milling processes, which produce average particle diameters in the 2 to 10 μm range. Dusts and powders can be prepared by blending and usually grinding (such as with a hammer mill or fluid-energy mill). Granules and pellets can be prepared by spraying the active material upon preformed granular carriers or by agglomeration techniques. See Browning, "Agglomeration", *Chemical Engineering*, December 4, 1967, pp 147-48, *Perry's Chemical Engineer's Handbook*, 4th Ed., McGraw-Hill, New York, 1963, pages 8-57 and following, and WO 91/13546. Pellets can be prepared as described in U.S. 4,172,714. Water-dispersible and water-soluble granules can be prepared as taught in U.S. 4,144,050, U.S. 3,920,442 and DE 3,246,493. Tablets can be prepared as taught in U.S. 5,180,587, U.S. 5,232,701 and U.S. 5,208,030. Films can be prepared as taught in GB 2,095,558 and U.S. 3,299,566.

For further information regarding the art of formulation, see T. S. Woods, "The Formulator's Toolbox – Product Forms for Modern Agriculture" in *Pesticide Chemistry and Bioscience, The Food–Environment Challenge*, T. Brooks and T. R. Roberts, Eds., Proceedings of the 9th International Congress on Pesticide Chemistry, The Royal Society of Chemistry, Cambridge, 1999, pp. 120–133. See also U.S. 3,235,361, Col. 6, line 16 through Col. 7, line 19 and Examples 10–41; U.S. 3,309,192, Col. 5, line 43 through Col. 7, line 62 and Examples 8, 12, 15, 39, 41, 52, 53, 58, 132, 138–140, 162–164, 166, 167 and 169–182; U.S. 2,891,855, Col. 3, line 66 through Col. 5, line 17 and Examples 1–4; Klingman, *Weed Control as a Science*, John Wiley and Sons, Inc., New York, 1961, pp 81–96; Hance et al., *Weed Control Handbook*, 8th Ed., Blackwell Scientific Publications, Oxford, 1989; and *Developments in formulation technology*, PJB Publications, Richmond, UK, 2000.

In the following Examples, all percentages are by weight and all formulations are prepared in conventional ways. Compound numbers refer to compounds in Index Table A. Without further elaboration, it is believed that one skilled in the art using the preceding description can utilize the present invention to its fullest extent. The following non-limiting Examples are illustrative of the invention. Percentages are by weight except where otherwise indicated.

Example A

High Strength Concentrate

Compound 3	98.5%
silica aerogel	0.5%
synthetic amorphous fine silica	1.0%

Example B

Wettable Powder

Compound 1	65.0%
dodecylphenol polyethylene glycol ether	2.0%
sodium ligninsulfonate	4.0%
sodium silicoaluminate	6.0%
montmorillonite (calcined)	23.0%

Example C

Granule

Compound 2	10.0%
attapulgit granules (low volatile matter, 0.71/0.30 mm; U.S.S. No. 25–50 sieves)	90.0%

Example D

Extruded Pellet

Compound 4	25.0%
anhydrous sodium sulfate	10.0%

crude calcium ligninsulfonate	5.0%
sodium alkyl naphthalenesulfonate	1.0%
calcium/magnesium bentonite	59.0%

Example EEmulsifiable Concentrate

Compound 3	10.0%
polyoxyethylene sorbitol hexoleate	20.0%
C ₆ -C ₁₀ fatty acid methyl ester	70.0%

Example FMicroemulsion

Compound 4	5.0%
polyvinylpyrrolidone-vinyl acetate copolymer	30.0%
alkylpolyglycoside	30.0%
glyceryl monooleate	15.0%
water	20.0%

Example GSuspension Concentrate

Compound 3	35%
butyl polyoxyethylene/polypropylene block copolymer	4.0%
stearic acid/polyethylene glycol copolymer	1.0%
styrene acrylic polymer	1.0%
xanthan gum	0.1%
propylene glycol	5.0%
silicone based defoamer	0.1%
1,2-benzisothiazolin-3-one	0.1%
water	53.7%

Example HEmulsion in Water

Compound 4	10.0%
butyl polyoxyethylene/polypropylene block copolymer	4.0%
stearic acid/polyethylene glycol copolymer	1.0%
styrene acrylic polymer	1.0%
xanthan gum	0.1%
propylene glycol	5.0%
silicone based defoamer	0.1%
1,2-benzisothiazolin-3-one	0.1%
aromatic petroleum based hydrocarbon	20.0%
water	58.7%

Example IOil Dispersion

Compound 5	25%
polyoxyethylene sorbitol hexaoleate	15%
organically modified bentonite clay	2.5%
fatty acid methyl ester	57.5%

Test results indicate that the compounds of the present invention are highly active preemergent and/or postemergent herbicides and/or plant growth regulants. The compounds of the invention generally show highest activity for postemergence weed control (i.e. applied after weed seedlings emerge from the soil) and preemergence weed control (i.e. applied before weed seedlings emerge from the soil). Many of them have utility for broad-spectrum pre- and/or postemergence weed control in areas where complete control of all vegetation is desired such as around fuel storage tanks, industrial storage areas, parking lots, drive-in theaters, air fields, river banks, irrigation and other waterways, around billboards and highway and railroad structures. Many of the compounds of this invention, by virtue of selective metabolism in crops versus weeds, or by selective activity at the locus of physiological inhibition in crops and weeds, or by selective placement on or within the environment of a mixture of crops and weeds, are useful for the selective control of grass and broadleaf weeds within a crop/weed mixture. One skilled in the art will recognize that the preferred combination of these selectivity factors within a compound or group of compounds can readily be determined by performing routine biological and/or biochemical assays. Compounds of this invention may show tolerance to important agronomic crops including, but is not limited to, alfalfa, barley, cotton, wheat, rape, sugar beets, corn (maize), sorghum, soybeans, rice, oats, peanuts, vegetables, tomato, potato, perennial plantation crops including coffee, cocoa, oil palm, rubber, sugarcane, citrus, grapes, fruit trees, nut trees, banana, plantain, pineapple, hops, tea and forests such as eucalyptus and conifers (e.g., loblolly pine), and turf species (e.g., Kentucky bluegrass, St. Augustine grass, Kentucky fescue and Bermuda grass). Compounds of this invention can be used in crops genetically transformed or bred to incorporate resistance to herbicides, express proteins toxic to invertebrate pests (such as *Bacillus thuringiensis* toxin), and/or express other useful traits. Those skilled in the art will appreciate that not all compounds are equally effective against all weeds. Alternatively, the subject compounds are useful to modify plant growth.

As the compounds of the invention have (both preemergent and postemergent herbicidal) activity, to control undesired vegetation by killing or injuring the vegetation or reducing its growth, the compounds can be usefully applied by a variety of methods involving contacting a herbicidally effective amount of a compound of the invention, or a composition comprising said compound and at least one of a surfactant, a solid diluent or a

liquid diluent, to the foliage or other part of the undesired vegetation or to the environment of the undesired vegetation such as the soil or water in which the undesired vegetation is growing or which surrounds the seed or other propagule of the undesired vegetation.

A herbicidally effective amount of the compounds of this invention is determined by a number of factors. These factors include: formulation selected, method of application, amount and type of vegetation present, growing conditions, etc. In general, a herbicidally effective amount of compounds of this invention is about 0.001 to 20 kg/ha with a preferred range of about 0.004 to 1 kg/ha. One skilled in the art can easily determine the herbicidally effective amount necessary for the desired level of weed control.

In one common embodiment, a compound of the invention is applied, typically in a formulated composition, to a locus comprising desired vegetation (e.g., crops) and undesired vegetation (i.e. weeds), both of which may be seeds, seedlings and/or larger plants, in contact with a growth medium (e.g., soil). In this locus, a composition comprising a compound of the invention can be directly applied to a plant or a part thereof, particularly of the undesired vegetation, and/or to the growth medium in contact with the plant.

Plant varieties and cultivars of the desired vegetation in the locus treated with a compound of the invention can be obtained by conventional propagation and breeding methods or by genetic engineering methods. Genetically modified plants (transgenic plants) are those in which a heterologous gene (transgene) has been stably integrated into the plant's genome. A transgene that is defined by its particular location in the plant genome is called a transformation or transgenic event.

Genetically modified plant cultivars in the locus which can be treated according to the invention include those that are resistant against one or more biotic stresses (pests such as nematodes, insects, mites, fungi, etc.) or abiotic stresses (drought, cold temperature, soil salinity, etc.), or that contain other desirable characteristics. Plants can be genetically modified to exhibit traits of, for example, herbicide tolerance, insect-resistance, modified oil profiles or drought tolerance. Useful genetically modified plants containing single gene transformation events or combinations of transformation events are listed in Exhibit C. Additional information for the genetic modifications listed in Exhibit C can be obtained from publicly available databases maintained, for example, by the U.S. Department of Agriculture.

The following abbreviations, T1 through T37, are used in Exhibit C for traits. "tol." means "tolerance". A hyphen "-" means the entry is not available.

Trait	Description	Trait	Description	Trait	Description
T1	Glyphosate tolerance	T15	Cold tolerance	T27	High tryptophan
T2	High lauric acid oil	T16	Imidazolinone herbicide tol.	T28	Erect leaves semidwarf
T3	Glufosinate tolerance	T17	Modified alpha-amylase	T29	Semidwarf
T4	Phytate breakdown	T18	Pollination control	T30	Low iron tolerance

T5	Oxynil tolerance	T19	2,4-D tolerance	T31	Modified oil/fatty acid
T6	Disease resistance	T20	Increased lysine	T32	HPPD tolerance
T7	Insect resistance	T21	Drought tolerance	T33	High oil
T9	Modified flower color	T22	Delayed ripening/senescence	T34	Aryloxyalkanoate tol.
T11	ALS herbicide tol.	T23	Modified product quality	T35	Mesotrione tolerance
T12	Dicamba tolerance	T24	High cellulose	T36	Reduced nicotine
T13	Anti-allergy	T25	Modified starch/carbohydrate	T37	Modified product
T14	Salt tolerance	T26	Insect & disease resistance		

Exhibit C

Crop	Event Name	Event Code	Trait(s)	Gene(s)
Alfalfa	J101	MON-00101-8	T1	cp4 epsps (aroA:CP4)
Alfalfa	J163	MON-00163-7	T1	cp4 epsps (aroA:CP4)
Canola*	23-18-17 (Event 18)	CGN-89465-2	T2	te
Canola*	23-198 (Event 23)	CGN-89465-2	T2	te
Canola*	61061	DP-061061-7	T1	gat4621
Canola*	73496	DP-073496-4	T1	gat4621
Canola*	GT200 (RT200)	MON-89249-2	T1	cp4 epsps (aroA:CP4); goxv247
Canola*	GT73 (RT73)	MON-00073-7	T1	cp4 epsps (aroA:CP4); goxv247
Canola*	HCN10 (Topas 19/2)	-	T3	bar
Canola*	HCN28 (T45)	ACS-BN008-2	T3	pat (syn)
Canola*	HCN92 (Topas 19/2)	ACS-BN007-1	T3	bar
Canola*	MON88302	MON-88302-9	T1	cp4 epsps (aroA:CP4)
Canola*	MPS961	-	T4	phyA
Canola*	MPS962	-	T4	phyA
Canola*	MPS963	-	T4	phyA
Canola*	MPS964	-	T4	phyA
Canola*	MPS965	-	T4	phyA
Canola*	MS1 (B91-4)	ACS-BN004-7	T3	bar
Canola*	MS8	ACS-BN005-8	T3	bar
Canola*	OXY-235	ACS-BN011-5	T5	bxn
Canola*	PHY14	-	T3	bar
Canola*	PHY23	-	T3	bar
Canola*	PHY35	-	T3	bar
Canola*	PHY36	-	T3	bar
Canola*	RF1 (B93-101)	ACS-BN001-4	T3	bar
Canola*	RF2 (B94-2)	ACS-BN002-5	T3	bar
Canola*	RF3	ACS-BN003-	T3	bar

Crop	Event Name	Event Code	Trait(s)	Gene(s)
		6		
Bean	EMBRAPA 5.1	EMB-PV051-1	T6	ac1 (sense and antisense)
Brinjal #	EE-1	-	T7	cry1Ac
Cotton	19-51a	DD-Ø1951A-7	T11	S4-HrA
Cotton	281-24-236	DAS-24236-5	T3,T7	pat (syn); cry1F
Cotton	3006-210-23	DAS-21Ø23-5	T3,T7	pat (syn); cry1Ac
Cotton	31707	-	T5,T7	bxn; cry1Ac
Cotton	31803	-	T5,T7	bxn; cry1Ac
Cotton	31807	-	T5,T7	bxn; cry1Ac
Cotton	31808	-	T5,T7	bxn; cry1Ac
Cotton	42317	-	T5,T7	bxn; cry1Ac
Cotton	BNLA-601	-	T7	cry1Ac
Cotton	BXN10211	BXN10211-9	T5	bxn; cry1Ac
Cotton	BXN10215	BXN10215-4	T5	bxn; cry1Ac
Cotton	BXN10222	BXN10222-2	T5	bxn; cry1Ac
Cotton	BXN10224	BXN10224-4	T5	bxn; cry1Ac
Cotton	COT102	SYN-IR102-7	T7	vip3A(a)
Cotton	COT67B	SYN-IR67B-1	T7	cry1Ab
Cotton	COT202	-	T7	vip3A
Cotton	Event 1	-	T7	cry1Ac
Cotton	GMF Cry1A	GTL- GMF311-7	T7	cry1Ab-Ac
Cotton	GHB119	BCS-GH005-8	T7	cry2Ae
Cotton	GHB614	BCS-GH002-5	T1	2mepsps
Cotton	GK12	-	T7	cry1Ab-Ac
Cotton	LLCotton25	ACS-GH001-3	T3	bar
Cotton	MLS 9124	-	T7	cry1C
Cotton	MON1076	MON-89924-2	T7	cry1Ac
Cotton	MON1445	MON-01445-2	T1	cp4 epsps (aroA:CP4)
Cotton	MON15985	MON-15985-7	T7	cry1Ac; cry2Ab2
Cotton	MON1698	MON-89383-1	T7	cp4 epsps (aroA:CP4)
Cotton	MON531	MON-00531-6	T7	cry1Ac
Cotton	MON757	MON-00757-7	T7	cry1Ac
Cotton	MON88913	MON-88913-8	T1	cp4 epsps (aroA:CP4)
Cotton	Nqwe Chi 6 Bt	-	T7	-
Cotton	SKG321	-	T7	cry1A; CpTI
Cotton	T303-3	BCS-GH003-6	T3,T7	cry1Ab; bar
Cotton	T304-40	BCS-GH004-7	T3,T7	cry1Ab; bar
Cotton	CE43-67B	-	T7	cry1Ab
Cotton	CE46-02A	-	T7	cry1Ab
Cotton	CE44-69D	-	T7	cry1Ab

Crop	Event Name	Event Code	Trait(s)	Gene(s)
Cotton	1143-14A	-	T7	cry1Ab
Cotton	1143-51B	-	T7	cry1Ab
Cotton	T342-142	-	T7	cry1Ab
Cotton	PV-GHGT07 (1445)	-	T1	cp4 epsps (aroA:CP4)
Cotton	EE-GH3	-	T1	mepsps
Cotton	EE-GH5	-	T7	cry1Ab
Cotton	MON88701	MON-88701-3	T3,T12	Modified dmo; bar
Cotton	OsCr11	-	T13	Modified Cry j
Flax	FP967	CDC-FL001-2	T11	als
Lentil	RH44	-	T16	als
Maize	3272	SYN-E3272-5	T17	amy797E
Maize	5307	SYN-05307-1	T7	ecry3.1Ab
Maize	59122	DAS-59122-7	T3,T7	cry34Ab1; cry35Ab1; pat
Maize	676	PH-000676-7	T3,T18	pat; dam
Maize	678	PH-000678-9	T3,T18	pat; dam
Maize	680	PH-000680-2	T3,T18	pat; dam
Maize	98140	DP-098140-6	T1,T11	gat4621; zm-hra
Maize	Bt10	-	T3,T7	cry1Ab; pat
Maize	Bt176 (176)	SYN-EV176-9	T3,T7	cry1Ab; bar
Maize	BVLA430101	-	T4	phyA2
Maize	CBH-351	ACS-ZM004-3	T3,T7	cry9C; bar
Maize	DAS40278-9	DAS40278-9	T19	aad-1
Maize	DBT418	DKB-89614-9	T3,T7	cry1Ac; pinII; bar
Maize	DLL25 (B16)	DKB-89790-5	T3	bar
Maize	GA21	MON-00021-9	T1	mepsps
Maize	GG25	-	T1	mepsps
Maize	GJ11	-	T1	mepsps
Maize	Fl117	-	T1	mepsps
Maize	GAT-ZM1	-	T3	pat
Maize	LY038	REN-00038-3	T20	cordapA
Maize	MIR162	SYN-IR162-4	T7	vip3Aa20
Maize	MIR604	SYN-IR604-5	T7	mcry3A
Maize	MON801 (MON80100)	MON801	T1,T7	cry1Ab; cp4 epsps (aroA:CP4); goxv247
Maize	MON802	MON-80200-7	T1,T7	cry1Ab; cp4 epsps (aroA:CP4); goxv247
Maize	MON809	PH-MON-809-2	T1,T7	cry1Ab; cp4 epsps (aroA:CP4); goxv247
Maize	MON810	MON-00810-6	T1,T7	cry1Ab; cp4 epsps (aroA:CP4); goxv247
Maize	MON832	-	T1	cp4 epsps (aroA:CP4); goxv247
Maize	MON863	MON-00863-5	T7	cry3Bb1
Maize	MON87427	MON-87427-7	T1	cp4 epsps (aroA:CP4)

Crop	Event Name	Event Code	Trait(s)	Gene(s)
Maize	MON87460	MON-87460-4	T21	cspB
Maize	MON88017	MON-88017-3	T1,T7	cry3Bb1; cp4 epsps (aroA:CP4)
Maize	MON89034	MON-89034-3	T7	cry2Ab2; cry1A.105
Maize	MS3	ACS-ZM001-9	T3,T18	bar; barnase
Maize	MS6	ACS-ZM005-4	T3,T18	bar; barnase
Maize	NK603	MON-00603-6	T1	cp4 epsps (aroA:CP4)
Maize	T14	ACS-ZM002-1	T3	pat (syn)
Maize	T25	ACS-ZM003-2	T3	pat (syn)
Maize	TC1507	DAS-01507-1	T3,T7	cry1Fa2; pat
Maize	TC6275	DAS-06275-8	T3,T7	mocry1F; bar
Maize	VIP1034	-	T3,T7	vip3A; pat
Maize	43A47	DP-043A47-3	T3,T7	cry1F; cry34Ab1; cry35Ab1; pat
Maize	40416	DP-040416-8	T3,T7	cry1F; cry34Ab1; cry35Ab1; pat
Maize	32316	DP-032316-8	T3,T7	cry1F; cry34Ab1; cry35Ab1; pat
Maize	4114	DP-004114-3	T3,T7	cry1F; cry34Ab1; cry35Ab1; pat
Melon	Melon A	-	T22	sam-k
Melon	Melon B	-	T22	sam-k
Papaya	55-1	CUH-CP551-8	T6	prsv cp
Papaya	63-1	CUH-CP631-7	T6	prsv cp
Papaya	Huanong No. 1	-	T6	prsv rep
Papaya	X17-2	UFL-X17CP-6	T6	prsv cp
Plum	C-5	ARS-PLMC5-6	T6	ppv cp
Canola**	ZSR500	-	T1	cp4 epsps (aroA:CP4); goxv247
Canola**	ZSR502	-	T1	cp4 epsps (aroA:CP4); goxv247
Canola**	ZSR503	-	T1	cp4 epsps (aroA:CP4); goxv247
Rice	7Cp#242-95-7	-	T13	7crp
Rice	7Cp#10	-	T13	7crp
Rice	GM Shanyou 63	-	T7	cry1Ab; cry1Ac
Rice	Huahui-1/TT51-1	-	T7	cry1Ab; cry1Ac
Rice	LLRICE06	ACS-OS001-4	T3	bar
Rice	LLRICE601	BCS-OS003-7	T3	bar
Rice	LLRICE62	ACS-OS002-5	T3	bar
Rice	Tarom molaii + cry1Ab	-	T7	cry1Ab (truncated)
Rice	GAT-OS2	-	T3	bar
Rice	GAT-OS3	-	T3	bar
Rice	PE-7	-	T7	Cry1Ac
Rice	7Cp#10	-	T13	7crp
Rice	KPD627-8	-	T27	OASA1D
Rice	KPD722-4	-	T27	OASA1D
Rice	KA317	-	T27	OASA1D

Crop	Event Name	Event Code	Trait(s)	Gene(s)
Rice	HW5	-	T27	OASA1D
Rice	HW1	-	T27	OASA1D
Rice	B-4-1-18	-	T28	Δ OsBRI1
Rice	G-3-3-22	-	T29	OSGA2ox1
Rice	AD77	-	T6	DEF
Rice	AD51	-	T6	DEF
Rice	AD48	-	T6	DEF
Rice	AD41	-	T6	DEF
Rice	13pNasNa800725atAprt1	-	T30	HvNAS1; HvNAAT-A; APRT
Rice	13pAprt1	-	T30	APRT
Rice	gHvNAS1-gHvNAAT-1	-	T30	HvNAS1; HvNAAT-A; HvNAAT-B
Rice	gHvIDS3-1	-	T30	HvIDS3
Rice	gHvNAAT1	-	T30	HvNAAT-A; HvNAAT-B
Rice	gHvNAS1-1	-	T30	HvNAS1
Rice	NIA-OS006-4	-	T6	WRKY45
Rice	NIA-OS005-3	-	T6	WRKY45
Rice	NIA-OS004-2	-	T6	WRKY45
Rice	NIA-OS003-1	-	T6	WRKY45
Rice	NIA-OS002-9	-	T6	WRKY45
Rice	NIA-OS001-8	-	T6	WRKY45
Rice	OsCr11	-	T13	Modified Cry j
Rice	17053	-	T1	cp4 epsps (aroA:CP4)
Rice	17314	-	T1	cp4 epsps (aroA:CP4)
Rose	WKS82 / 130-4-1	IFD-52401-4	T9	5AT; bp40 (f3'5'h)
Rose	WKS92 / 130-9-1	IFD-52901-9	T9	5AT; bp40 (f3'5'h)
Soybean	260-05 (G94-1, G94-19, G168)	-	T9	gm-fad2-1 (silencing locus)
Soybean	A2704-12	ACS-GM005-3	T3	pat
Soybean	A2704-21	ACS-GM004-2	T3	pat
Soybean	A5547-127	ACS-GM006-4	T3	pat
Soybean	A5547-35	ACS-GM008-6	T3	pat
Soybean	CV127	BPS-CV127-9	T16	csr1-2
Soybean	DAS68416-4	DAS68416-4	T3	pat
Soybean	DP305423	DP-305423-1	T11,T31	gm-fad2-1 (silencing locus); gm-hra
Soybean	DP356043	DP-356043-5	T1,T31	gm-fad2-1 (silencing locus); gat4601
Soybean	FG72	MST-FG072-3	T32,T1	2mepsps; hppdPF W336
Soybean	GTS 40-3-2 (40-3-2)	MON-04032-6	T1	cp4 epsps (aroA:CP4)
Soybean	GU262	ACS-GM003-1	T3	pat

Crop	Event Name	Event Code	Trait(s)	Gene(s)
Soybean	MON87701	MON-87701-2	T7	cry1Ac
Soybean	MON87705	MON-87705-6	T1,T31	fatb1-A (sense & antisense); fad2-1A (sense & antisense); cp4 epsps (aroA:CP4)
Soybean	MON87708	MON-87708-9	T1,T12	dmo; cp4 epsps (aroA:CP4)
Soybean	MON87769	MON-87769-7	T1,T31	Pj.D6D; Nc.Fad3; cp4 epsps (aroA:CP4)
Soybean	MON89788	MON-89788-1	T1	cp4 epsps (aroA:CP4)
Soybean	W62	ACS-GM002-9	T3	bar
Soybean	W98	ACS-GM001-8	T3	bar
Soybean	MON87754	MON-87754-1	T33	dgat2A
Soybean	DAS21606	DAS-21606	T34,T3	Modified aad-12; pat
Soybean	DAS44406	DAS-44406-6	T1,T3,T34	Modified aad-12; 2mepsps; pat
Soybean	SYHT04R	SYN-0004R-8	T35	Modified avhppd
Soybean	9582.814.19.1	-	T3,T7	cry1Ac, cry1F, PAT
Squash	CZW3	SEM-ØCZW3-2	T6	cmv cp, zymv cp, wmv cp
Squash	ZW20	SEM-ØZW20-7	T6	zymv cp, wmv cp
Sugar Beet	GTSB77 (T9100152)	SY-GTSB77-8	T1	cp4 epsps (aroA:CP4); goxv247
Sugar Beet	H7-1	KM-000H71-4	T1	cp4 epsps (aroA:CP4)
Sugar Beet	T120-7	ACS-BV001-3	T3	pat
Sugar Beet	T227-1	-	T1	cp4 epsps (aroA:CP4)
Sugarcane	NXI-1T	-	T21	EcbetA
Sunflower	X81359	-	T16	als
Pepper	PK-SP01	-	T6	cmv cp
Tobacco	C/F/93/08-02	-	T5	bxn
Tobacco	Vector 21-41	-	T36	NtQPT1 (antisense)
Sunflower	X81359	-	T16	als
Wheat	MON71800	MON-71800-3	T1	cp4 epsps (aroA:CP4)

* Argentine (*Brassica napus*), ** Polish (*B. rapa*), # Eggplant

Although most typically, compounds of the invention are used to control undesired vegetation, contact of desired vegetation in the treated locus with compounds of the invention may result in super-additive or synergistic effects with genetic traits in the desired vegetation, including traits incorporated through genetic modification. For example, resistance to phytophagous insect pests or plant diseases, tolerance to biotic/abiotic stresses or storage stability may be greater than expected from the genetic traits in the desired vegetation.

An embodiment of the present invention is a method for controlling the growth of undesired vegetation in genetically modified plants that exhibit traits of glyphosate tolerance, glufosinate tolerance, ALS herbicide tolerance, dicamba tolerance, imidazolinone herbicide

tolerance, 2,4-D tolerance, HPPD tolerance and mesotrione tolerance, comprising contacting the vegetation or its environment with a herbicidally effective amount of a compound of Formula 1.

Compounds of this invention can also be mixed with one or more other biologically active compounds or agents including herbicides, herbicide safeners, fungicides, insecticides, nematocides, bactericides, acaricides, growth regulators such as insect molting inhibitors and rooting stimulants, chemosterilants, semiochemicals, repellents, attractants, pheromones, feeding stimulants, plant nutrients, other biologically active compounds or entomopathogenic bacteria, virus or fungi to form a multi-component pesticide giving an even broader spectrum of agricultural protection. Mixtures of the compounds of the invention with other herbicides can broaden the spectrum of activity against additional weed species, and suppress the proliferation of any resistant biotypes. Thus the present invention also pertains to a composition comprising a compound of Formula 1 (in a herbicidally effective amount) and at least one additional biologically active compound or agent (in a biologically effective amount) and can further comprise at least one of a surfactant, a solid diluent or a liquid diluent. The other biologically active compounds or agents can be formulated in compositions comprising at least one of a surfactant, solid or liquid diluent. For mixtures of the present invention, one or more other biologically active compounds or agents can be formulated together with a compound of Formula 1, to form a premix, or one or more other biologically active compounds or agents can be formulated separately from the compound of Formula 1, and the formulations combined together before application (e.g., in a spray tank) or, alternatively, applied in succession.

A mixture of one or more of the following herbicides with a compound of this invention may be particularly useful for weed control: acetochlor, acifluorfen and its sodium salt, aclonifen, acrolein (2-propenal), alachlor, alloxymid, ametryn, amicarbazone, amidosulfuron, aminocyclopyrachlor and its esters (e.g., methyl, ethyl) and salts (e.g., sodium, potassium), aminopyralid, amitrole, ammonium sulfamate, anilofos, asulam, atrazine, azimsulfuron, beflubutamid, benazolin, benazolin-ethyl, bencarbazone, benfluralin, benfuresate, bensulfuron-methyl, bensulide, bentazone, benzobicyclon, benzofenap, bicyclopyrone, bifenox, bilanafos, bispyribac and its sodium salt, bromacil, bromobutide, bromofenoxim, bromoxynil, bromoxynil octanoate, butachlor, butafenacil, butamifos, butralin, butoxydim, butylate, cafenstrole, carbetamide, carfentrazone-ethyl, catechin, chlomethoxyfen, chloramben, chlorbromuron, chlorflurenol-methyl, chloridazon, chlorimuron-ethyl, chlorotoluron, chlorpropham, chlorsulfuron, chlorthal-dimethyl, chlorthiamid, cinidon-ethyl, cinmethylin, cinosulfuron, clacyfos, clefoxydim, clethodim, clodinafop-propargyl, clomazone, clomeprop, clopyralid, clopyralid-olamine, cloransulam-methyl, cumyluron, cyanazine, cycloate, cyclopyrimorate, cyclosulfamuron, cycloxydim, cyhalofop-butyl, 2,4-D and its butotyl, butyl, isoctyl and isopropyl esters and its

dimethylammonium, diolamine and trolamine salts, daimuron, dalapon, dalapon-sodium, dazomet, 2,4-DB and its dimethylammonium, potassium and sodium salts, desmedipham, desmetryn, dicamba and its diglycolammonium, dimethylammonium, potassium and sodium salts, dichlobenil, dichlorprop, diclofop-methyl, diclosulam, difenzoquat metilsulfate, 5 diflufenican, diflufenzopyr, dimefuron, dimepiperate, dimethachlor, dimethametryn, dimethenamid, dimethenamid-P, dimethipin, dimethylarsinic acid and its sodium salt, dinitramine, dinoterb, diphenamid, diquat dibromide, dithiopyr, diuron, DNOC, endothal, EPTC, esprocarb, ethalfluralin, ethametsulfuron-methyl, ethiozin, ethofumesate, ethoxyfen, ethoxysulfuron, etobenzanid, fenoxaprop-ethyl, fenoxaprop-P-ethyl, fenoxasulfone, 10 fenquinotriene, fentrazamide, fenuron, fenuron-TCA, flamprop-methyl, flamprop-M-isopropyl, flamprop-M-methyl, flazasulfuron, florasulam, fluazifop-butyl, fluazifop-P-butyl, fluazolate, flucarbazone, flucetosulfuron, fluchloralin, flufenacet, flufenpyr, flufenpyr-ethyl, flumetsulam, flumiclorac-pentyl, flumioxazin, fluometuron, fluoroglycofen-ethyl, flupoxam, flupyrsulfuron-methyl and its sodium salt, flurenol, 15 flurenol-butyl, fluridone, flurochloridone, fluroxypyr, flurtamone, fluthiacet-methyl, fomesafen, foramsulfuron, fosamine-ammonium, glufosinate, glufosinate-ammonium, glufosinate-P, glyphosate and its salts such as ammonium, isopropylammonium, potassium, sodium (including sesquisodium) and trimesium (alternatively named sulfosate), halauxifen, halauxifen-methyl, halosulfuron-methyl, haloxyfop-ethyl, haloxyfop-methyl, hexazinone, 20 imazamethabenz-methyl, imazamox, imazapic, imazapyr, imazaquin, imazaquin-ammonium, imazethapyr, imazethapyr-ammonium, imazosulfuron, indanofan, indaziflam, iofensulfuron, iodosulfuron-methyl, ioxynil, ioxynil octanoate, ioxynil-sodium, ipfencarbazone, isoproturon, isouron, isoxaben, isoxaflutole, isoxachlortole, lactofen, lenacil, linuron, maleic hydrazide, MCPA and its salts (e.g., MCPA-dimethylammonium, MCPA-potassium and 25 MCPA-sodium, esters (e.g., MCPA-2-ethylhexyl, MCPA-butyl) and thioesters (e.g., MCPA-thioethyl), MCPB and its salts (e.g., MCPB-sodium) and esters (e.g., MCPB-ethyl), mecoprop, mecoprop-P, mefenacet, mefluidide, mesosulfuron-methyl, mesotrione, metam-sodium, metamifop, metamitron, metazachlor, metazosulfuron, methabenzthiazuron, methylarsonic acid and its calcium, monoammonium, monosodium and disodium salts, 30 methylmymron, metobenzuron, metobromuron, metolachlor, S-metolachlor, metosulam, metoxuron, metribuzin, metsulfuron-methyl, molinate, monolinuron, naproanilide, napropamide, napropamide-M, naptalam, neburon, nicosulfuron, norflurazon, orbencarb, orthosulfamuron, oryzalin, oxadiargyl, oxadiazon, oxasulfuron, oxaziclomefone, oxyfluorfen, paraquat dichloride, pebulate, pelargonic acid, pendimethalin, penoxsulam, 35 pentanochlor, pentoxazone, perfluidone, pethoxamid, pethoxyamid, phenmedipham, picloram, picloram-potassium, picolinafen, pinoxaden, piperophos, pretilachlor, primisulfuron-methyl, prodiamine, profoxydim, prometon, prometryn, propachlor, propanil, propaquizafop, propazine, propham, propisochlor, propoxycarbazone, propyrisulfuron,

propyzamide, prosulfocarb, prosulfuron, pyraclonil, pyraflufen-ethyl, pyrasulfotole, pyrazogyl, pyrazolynate, pyrazoxyfen, pyrazosulfuron-ethyl, pyribenzoxim, pyributicarb, pyridate, pyriftalid, pyriminobac-methyl, pyrimisulfan, pyrithiobac, pyrithiobac-sodium, pyroxasulfone, pyroxsulam, quinclorac, quinmerac, quinoclamine, quizalofop-ethyl, 5 quizalofop-P-ethyl, quizalofop-P-tefuryl, rimsulfuron, saflufenacil, sethoxydim, siduron, simazine, simetryn, sulcotrione, sulfentrazone, sulfometuron-methyl, sulfosulfuron, 2,3,6-TBA, TCA, TCA-sodium, tebutam, tebuthiuron, tefuryltrione, tembotrione, tepraloxym, terbacil, terbutryn, terbuthylazine, terbutryn, thenylchlor, thiazopyr, thiencarbazone, thifensulfuron-methyl, thiobencarb, tiafenacil, tiocarbazil, topramezone, tralkoxydim, 10 tri-allate, triafamone, triasulfuron, triaziflam, tribenuron-methyl, triclopyr, triclopyr-butyl, triclopyr-triethylammonium, tridiphane, trietazine, trifloxysulfuron, trifluralin, triflusulfuron-methyl, tritosulfuron, vernolate, 3-(2-chloro-3,6-difluorophenyl)-4-hydroxy-1-methyl-1,5-naphthyridin-2(1*H*)-one, 5-chloro-3-[(2-hydroxy-6-oxo-1-cyclohexen-1-yl)carbonyl]-1-(4-methoxyphenyl)-2(1*H*)-quinoxalinone, 2-chloro-*N*-(1-methyl-1*H*-tetrazol-5-yl)-6-(trifluoromethyl)-3-pyridinecarboxamide, 7-(3,5-dichloro-4-pyridinyl)-5-(2,2-difluoroethyl)-8-hydroxypyrido[2,3-*b*]pyrazin-6(5*H*)-one, 4-(2,6-diethyl-4-methylphenyl)-5-hydroxy-2,6-dimethyl-3(2*H*)-pyridazinone, 5-[(2,6-difluorophenyl)methoxy]methyl]-4,5-dihydro-5-methyl-3-(3-methyl-2-thienyl)isoxazole (previously methioxolin), 3-[7-fluoro-3,4-dihydro-3-oxo-4-(2-propyn-1-yl)-2*H*-1,4-benzoxazin-6-yl]dihydro-1,5-dimethyl-6-thioxo- 20 1,3,5-triazine-2,4(1*H*,3*H*)-dione, 4-(4-fluorophenyl)-6-[(2-hydroxy-6-oxo-1-cyclohexen-1-yl)carbonyl]-2-methyl-1,2,4-triazine-3,5(2*H*,4*H*)-dione, methyl 4-amino-3-chloro-6-(4-chloro-2-fluoro-3-methoxyphenyl)-5-fluoro-2-pyridinecarboxylate, 2-methyl-3-(methylsulfonyl)-*N*-(1-methyl-1*H*-tetrazol-5-yl)-4-(trifluoromethyl)benzamide and 2-methyl-*N*-(4-methyl-1,2,5-oxadiazol-3-yl)-3-(methylsulfinyl)-4-(trifluoromethyl)benzamide. Other 25 herbicides also include bioherbicides such as *Alternaria destruens* Simmons, *Colletotrichum gloeosporioides* (Penz.) Penz. & Sacc., *Drechslera monoceras* (MTB-951), *Myrothecium verrucaria* (Albertini & Schweinitz) Ditmar: Fries, *Phytophthora palmivora* (Butl.) Butl. and *Puccinia thlaspeos* Schub.

Compounds of this invention can also be used in combination with plant growth 30 regulators such as aviglycine, *N*-(phenylmethyl)-1*H*-purin-6-amine, epocholeone, gibberellic acid, gibberellin A₄ and A₇, harpin protein, mepiquat chloride, prohexadione calcium, prohydrojasmon, sodium nitrophenolate and trinexapac-methyl, and plant growth modifying organisms such as *Bacillus cereus* strain BP01.

General references for agricultural protectants (i.e. herbicides, herbicide safeners, 35 insecticides, fungicides, nematocides, acaricides and biological agents) include *The Pesticide Manual, 13th Edition*, C. D. S. Tomlin, Ed., British Crop Protection Council, Farnham, Surrey, U.K., 2003 and *The BioPesticide Manual, 2nd Edition*, L. G. Copping, Ed., British Crop Protection Council, Farnham, Surrey, U.K., 2001.

For embodiments where one or more of these various mixing partners are used, active ingredients are often applied at an application rate between one-half and the full application rate specified on product labels for use of the active ingredient alone. The amounts are listed in references such as *The Pesticide Manual* and *The BioPesticide Manual*. The weight ratio of these various mixing partners (in total) to the compound of Formula 1 is typically between about 1:3000 and about 3000:1. Of note are weight ratios between about 1:300 and about 300:1 (for example ratios between about 1:30 and about 30:1). One skilled in the art can easily determine through simple experimentation the biologically effective amounts of active ingredients necessary for the desired spectrum of biological activity. It will be evident that including these additional components may expand the spectrum of weeds controlled beyond the spectrum controlled by the compound of Formula 1 alone.

In certain instances, combinations of a compound of this invention with other biologically active (particularly herbicidal) compounds or agents (i.e. active ingredients) can result in a greater-than-additive (i.e. synergistic) effect on weeds and/or a less-than-additive effect (i.e. safening) on crops or other desirable plants. Reducing the quantity of active ingredients released in the environment while ensuring effective pest control is always desirable. Ability to use greater amounts of active ingredients to provide more effective weed control without excessive crop injury is also desirable. When synergism of herbicidal active ingredients occurs on weeds at application rates giving agronomically satisfactory levels of weed control, such combinations can be advantageous for reducing crop production cost and decreasing environmental load. When safening of herbicidal active ingredients occurs on crops, such combinations can be advantageous for increasing crop protection by reducing weed competition.

Of note is a combination of a compound of the invention with at least one other herbicidal active ingredient. Of particular note is such a combination where the other herbicidal active ingredient has different site of action from the compound of the invention. In certain instances, a combination with at least one other herbicidal active ingredient having a similar spectrum of control but a different site of action will be particularly advantageous for resistance management. Thus, a composition of the present invention can further comprise (in a herbicidally effective amount) at least one additional herbicidal active ingredient having a similar spectrum of control but a different site of action.

Compounds of this invention can also be used in combination with herbicide safeners such as allidochlor, benoxacor, cloquintocet-mexyl, cumyluron, cyometrinil, cyprosulfonamide, daimuron, dichlormid, dicyclonon, dietholate, dimepiperate, fenchlorazole-ethyl, fenclorim, flurazole, fluxofenim, furilazole, isoxadifen-ethyl, mefenpyr-diethyl, mephenate, methoxyphenone naphthalic anhydride (1,8-naphthalic anhydride), oxabetrinil, *N*-(aminocarbonyl)-2-methylbenzenesulfonamide, *N*-(aminocarbonyl)-2-fluorobenzenesulfonamide, 1-bromo-4-[(chloromethyl)sulfonyl]benzene (BCS), 4-

(dichloroacetyl)-1-oxa-4-azospiro[4.5]decane (MON 4660), 2-(dichloromethyl)-2-methyl-1,3-dioxolane (MG 191), ethyl 1,6-dihydro-1-(2-methoxyphenyl)-6-oxo-2-phenyl-5-pyrimidinecarboxylate, 2-hydroxy-*N,N*-dimethyl-6-(trifluoromethyl)pyridine-3-carboxamide, and 3-oxo-1-cyclohexen-1-yl 1-(3,4-dimethylphenyl)-1,6-dihydro-6-oxo-2-phenyl-5-pyrimidinecarboxylate to increase safety to certain crops. Antidotally effective amounts of the herbicide safeners can be applied at the same time as the compounds of this invention, or applied as seed treatments. Therefore an aspect of the present invention relates to a herbicidal mixture comprising a compound of this invention and an antidotally effective amount of a herbicide safener. Seed treatment is particularly useful for selective weed control, because it physically restricts antidoting to the crop plants. Therefore a particularly useful embodiment of the present invention is a method for selectively controlling the growth of undesired vegetation in a crop comprising contacting the locus of the crop with a herbicidally effective amount of a compound of this invention wherein seed from which the crop is grown is treated with an antidotally effective amount of safener. Antidotally effective amounts of safeners can be easily determined by one skilled in the art through simple experimentation.

Of note is a composition comprising a compound of the invention (in a herbicidally effective amount), at least one additional active ingredient selected from the group consisting of other herbicides and herbicide safeners (in an effective amount), and at least one component selected from the group consisting of surfactants, solid diluents and liquid diluents.

Preferred for better control of undesired vegetation (e.g., lower use rate such as from synergism, broader spectrum of weeds controlled, or enhanced crop safety) or for preventing the development of resistant weeds are mixtures of a compound of this invention with a herbicide selected from the group 2,4-D, acetochlor, alachlor, atrazine, bromoxynil, bentazon, bicyclopyrone, carfentrazone-ethyl, cloransulam-methyl, dicamba, dimethenamid-p, florasulam, flufenacet, flumioxazin, flupyr-sulfuron-methyl, fluroxypyr-methyl, glyphosate, halauxifen-methyl, isoxaflutole, MCPA, mesotrione, metolachlor, metsulfuron-methyl, nicosulfuron, pyrasulfotole, pyroxasulfone, pyroxsulam, rimsulfuron, saflufenacil, tembotrione, thifensulfuron-methyl, topramazine and tribenuron.

Table A1 lists specific combinations of a Component (a) with Component (b) illustrative of the mixtures, compositions and methods of the present invention. Compound No. (Compound Number) (i.e. Compound 1) in the Component (a) column is identified in Index Table A. The second column of Table A1 lists the specific Component (b) compound (e.g., "2,4-D" in the first line). The third, fourth and fifth columns of Table A1 lists ranges of weight ratios for rates at which the Component (a) compound is typically applied to a field-grown crop relative to Component (b) (i.e. (a):(b)). Thus, for example, the first line of Table A1 specifically discloses the combination of Component (a) (i.e. Compound 1 in

Index Table A) with 2,4-D is typically applied in a weight ratio between 1:192 – 6:1. The remaining lines of Table A1 are to be construed similarly.

TABLE A1

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	2,4-D	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Acetochlor	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Acifluorfen	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Aclonifen	1:857 – 2:1	1:285 – 1:3	1:107 – 1:12
1	Alachlor	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Ametryn	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Amicarbazone	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Amidosulfuron	1:6 – 168:1	1:2 – 56:1	1:1 – 11:1
1	Aminocyclopyrachlor	1:48 – 24:1	1:16 – 8:1	1:6 – 2:1
1	Aminopyralid	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Amitrole	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Anilofos	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Asulam	1:960 – 2:1	1:320 – 1:3	1:120 – 1:14
1	Atrazine	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Azimsulfuron	1:6 – 168:1	1:2 – 56:1	1:1 – 11:1
1	Beflubutamid	1:342 – 4:1	1:114 – 2:1	1:42 – 1:5
1	Benfuresate	1:617 – 2:1	1:205 – 1:2	1:77 – 1:9
1	Bensulfuron-methyl	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Bentazone	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Benzobicyclon	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Benzofenap	1:257 – 5:1	1:85 – 2:1	1:32 – 1:4
1	Bicyclopyrone	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Bifenox	1:257 – 5:1	1:85 – 2:1	1:32 – 1:4
1	Bispyribac-sodium	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Bromacil	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Bromobutide	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Bromoxynil	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Butachlor	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Butafenacil	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Butylate	1:1542 – 1:2	1:514 – 1:5	1:192 – 1:22
1	Carfenstrole	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Carfentrazone-ethyl	1:128 – 9:1	1:42 – 3:1	1:16 – 1:2

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Chlorimuron-ethyl	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Chlorotoluron	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Chlorsulfuron	1:6 – 168:1	1:2 – 56:1	1:1 – 11:1
1	Cincosulfuron	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Cinidon-ethyl	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Cinmethylin	1:34 – 34:1	1:11 – 12:1	1:4 – 3:1
1	Clacyfos	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Clethodim	1:48 – 24:1	1:16 – 8:1	1:6 – 2:1
1	Clodinafop-propargyl	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Clomazone	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Clomeprop	1:171 – 7:1	1:57 – 3:1	1:21 – 1:3
1	Clopyralid	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Cloransulam-methyl	1:12 – 96:1	1:4 – 32:1	1:1 – 6:1
1	Cumyluron	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Cyanazine	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Cyclopyrimorate	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Cyclosulfamuron	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Cycloxydim	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Cyhalofop	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Daimuron	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Desmedipham	1:322 – 4:1	1:107 – 2:1	1:40 – 1:5
1	Dicamba	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Dichlobenil	1:1371 – 1:2	1:457 – 1:4	1:171 – 1:20
1	Dichlorprop	1:925 – 2:1	1:308 – 1:3	1:115 – 1:13
1	Diclofop-methyl	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Diclosulam	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Difenzoquat	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Diffenican	1:857 – 2:1	1:285 – 1:3	1:107 – 1:12
1	Diffenzopyr	1:12 – 96:1	1:4 – 32:1	1:1 – 6:1
1	Dimethachlor	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Dimethametryn	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Dimethenamid-P	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Dithiopyr	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Diuron	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	EPTC	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Esprocarb	1:1371 – 1:2	1:457 – 1:4	1:171 – 1:20
1	Ethalfuralin	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Ethametsulfuron-methyl	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Ethoxyfen	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Ethoxysulfuron	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Etobenzanid	1:257 – 5:1	1:85 – 2:1	1:32 – 1:4
1	Fenoxaprop-ethyl	1:120 – 10:1	1:40 – 4:1	1:15 – 1:2
1	Fenoxasulfone	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Fenquinotrione	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Fentrazamide	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Flazasulfuron	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Florasulam	1:2 – 420:1	1:1 – 140:1	2:1 – 27:1
1	Fluazifop-butyl	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Flucarbazone	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Flucetosulfuron	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Flufenacet	1:257 – 5:1	1:85 – 2:1	1:32 – 1:4
1	Flumetsulam	1:24 – 48:1	1:8 – 16:1	1:3 – 3:1
1	Flumiclorac-pentyl	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Flumioxazin	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Fluometuron	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Flupyrsulfuron-methyl	1:3 – 336:1	1:1 – 112:1	2:1 – 21:1
1	Fluridone	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Fluroxypyr	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Flurtamone	1:857 – 2:1	1:285 – 1:3	1:107 – 1:12
1	Fluthiacet-methyl	1:48 – 42:1	1:16 – 14:1	1:3 – 3:1
1	Fomesafen	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Foramsulfuron	1:13 – 84:1	1:4 – 28:1	1:1 – 6:1
1	Glufosinate	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Glyphosate	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Halosulfuron-methyl	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Halauxifen	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Halauxifen methyl	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Haloxypop-methyl	1:34 – 34:1	1:11 – 12:1	1:4 – 3:1
1	Hexazinone	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Imazamox	1:13 – 84:1	1:4 – 28:1	1:1 – 6:1

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Imazapic	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Imazapyr	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Imazaquin	1:34 – 34:1	1:11 – 12:1	1:4 – 3:1
1	Imazethabenz-methyl	1:171 – 7:1	1:57 – 3:1	1:21 – 1:3
1	Imazethapyr	1:24 – 48:1	1:8 – 16:1	1:3 – 3:1
1	Imazosulfuron	1:27 – 42:1	1:9 – 14:1	1:3 – 3:1
1	Indanofan	1:342 – 4:1	1:114 – 2:1	1:42 – 1:5
1	Indaziflam	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Iodosulfuron-methyl	1:3 – 336:1	1:1 – 112:1	2:1 – 21:1
1	Ioxynil	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Ipfencarbazone	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Isoproturon	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Isoxaben	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Isoxaflutole	1:60 – 20:1	1:20 – 7:1	1:7 – 2:1
1	Lactofen	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Lenacil	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Linuron	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	MCPA	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	MCPB	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Mecoprop	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Mefenacet	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Mefluidide	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Mesosulfuron-methyl	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1
1	Mesotrione	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Metamifop	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Metazachlor	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Metazosulfuron	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Methabenzthiazuron	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Metolachlor	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Metosulam	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Metribuzin	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Metsulfuron-methyl	1:2 – 560:1	1:1 – 187:1	3:1 – 35:1
1	Molinate	1:1028 – 2:1	1:342 – 1:3	1:128 – 1:15
1	Napropamide	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Napropamide-M	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Naptalam	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Nicosulfuron	1:12 – 96:1	1:4 – 32:1	1:1 – 6:1
1	Norflurazon	1:1152 – 1:1	1:384 – 1:3	1:144 – 1:16
1	Orbencarb	1:1371 – 1:2	1:457 – 1:4	1:171 – 1:20
1	Orthosulfamuron	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Oryzalin	1:514 – 3:1	1:171 – 1:2	1:64 – 1:8
1	Oxadiargyl	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Oxadiazon	1:548 – 3:1	1:182 – 1:2	1:68 – 1:8
1	Oxasulfuron	1:27 – 42:1	1:9 – 14:1	1:3 – 3:1
1	Oxaziclomefone	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Oxyfluorfen	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Paraquat	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Pendimethalin	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Penoxsulam	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Pentoxamid	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Pentoxazone	1:102 – 12:1	1:34 – 4:1	1:12 – 1:2
1	Phenmedipham	1:102 – 12:1	1:34 – 4:1	1:12 – 1:2
1	Picloram	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Picolinafen	1:34 – 34:1	1:11 – 12:1	1:4 – 3:1
1	Pinoxaden	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Pretilachlor	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Primisulfuron-methyl	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Prodiamine	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Profoxydim	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Prometryn	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Propachlor	1:1152 – 1:1	1:384 – 1:3	1:144 – 1:16
1	Propanil	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Propaquizafop	1:48 – 24:1	1:16 – 8:1	1:6 – 2:1
1	Propoxycarbazone	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Propyrisulfuron	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Propyzamide	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Prosulfocarb	1:1200 – 1:2	1:400 – 1:4	1:150 – 1:17
1	Prosulfuron	1:6 – 168:1	1:2 – 56:1	1:1 – 11:1
1	Pyraclonil	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Pyraflufen-ethyl	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Pyrasulfotole	1:13 – 84:1	1:4 – 28:1	1:1 – 6:1
1	Pyrazolynate	1:857 – 2:1	1:285 – 1:3	1:107 – 1:12
1	Pyrazosulfuron-ethyl	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Pyrazoxyfen	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1
1	Pyribenzoxim	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Pyributicarb	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Pyridate	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Pyriftalid	1:10 – 112:1	1:3 – 38:1	1:1 – 7:1
1	Pyriminobac-methyl	1:20 – 56:1	1:6 – 19:1	1:2 – 4:1
1	Pyrimisulfan	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Pyrithiobac	1:24 – 48:1	1:8 – 16:1	1:3 – 3:1
1	Pyroxasulfone	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Pyroxsulam	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1
1	Quinclorac	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Quizalofop-ethyl	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Rimsulfuron	1:13 – 84:1	1:4 – 28:1	1:1 – 6:1
1	Saflufenacil	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Sethoxydim	1:96 – 12:1	1:32 – 4:1	1:12 – 1:2
1	Simazine	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Sulcotrione	1:120 – 10:1	1:40 – 4:1	1:15 – 1:2
1	Sulfentrazone	1:147 – 8:1	1:49 – 3:1	1:18 – 1:3
1	Sulfometuron-methyl	1:34 – 34:1	1:11 – 12:1	1:4 – 3:1
1	Sulfosulfuron	1:8 – 135:1	1:2 – 45:1	1:1 – 9:1
1	Tebuthiuron	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Tefuryltrione	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1
1	Tembotrione	1:31 – 37:1	1:10 – 13:1	1:3 – 3:1
1	Tepraloxydim	1:25 – 45:1	1:8 – 15:1	1:3 – 3:1
1	Terbacil	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Terbuthylazine	1:857 – 2:1	1:285 – 1:3	1:107 – 1:12
1	Terbutryn	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Thenylchlor	1:85 – 14:1	1:28 – 5:1	1:10 – 1:2
1	Thiazopyr	1:384 – 3:1	1:128 – 1:1	1:48 – 1:6
1	Thiencarbazone	1:3 – 336:1	1:1 – 112:1	2:1 – 21:1
1	Thifensulfuron-methyl	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1
1	Tiafenacil	1:42 – 27:1	1:14 – 9:1	1:5 – 2:1

Component (a) (Compound No.)	Component (b)	Typical Weight Ratio	More Typical Weight Ratio	Most Typical Weight Ratio
1	Thiobencarb	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Topramzone	1:6 – 168:1	1:2 – 56:1	1:1 – 11:1
1	Tralkoxydim	1:68 – 17:1	1:22 – 6:1	1:8 – 2:1
1	Triallate	1:768 – 2:1	1:256 – 1:2	1:96 – 1:11
1	Triasulfuron	1:5 – 224:1	1:1 – 75:1	1:1 – 14:1
1	Triaziflam	1:171 – 7:1	1:57 – 3:1	1:21 – 1:3
1	Tribenuron-methyl	1:3 – 336:1	1:1 – 112:1	2:1 – 21:1
1	Triclopyr	1:192 – 6:1	1:64 – 2:1	1:24 – 1:3
1	Trifloxysulfuron	1:2 – 420:1	1:1 – 140:1	2:1 – 27:1
1	Trifluralin	1:288 – 4:1	1:96 – 2:1	1:36 – 1:4
1	Triflusulfuron-methyl	1:17 – 68:1	1:5 – 23:1	1:2 – 5:1
1	Tritosulfuron	1:13 – 84:1	1:4 – 28:1	1:1 – 6:1

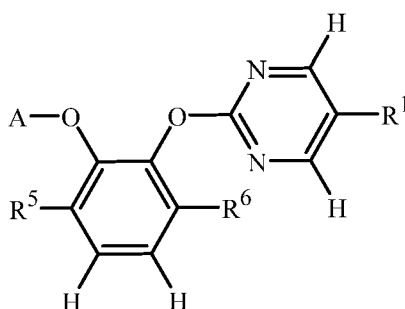
Table A2 is constructed the same as Table A1 above except that entries below the “Component (a)” column heading are replaced with the respective Component (a) Column Entry shown below. Compound No. in the Component (a) column is identified in Index Table A. Thus, for example, in Table A2 the entries below the “Component (a)” column heading all recite “Compound 2” (i.e. Compound 2 identified in Index Table A), and the first line below the column headings in Table A2 specifically discloses a mixture of Compound 2 with 2,4-D. Tables A3 through A5 are constructed similarly.

<u>Table Number</u>	<u>Component (a) Column Entries</u>	<u>Table Number</u>	<u>Component (a) Column Entries</u>
A2	Compound 2	A12	Compound 27
A3	Compound 3	A13	Compound 29
A4	Compound 4	A14	Compound 32
A5	Compound 5	A15	Compound 34
A6	Compound 10	A16	Compound 38
A7	Compound 16	A17	Compound 39
A8	Compound 17	A18	Compound 42
A9	Compound 18	A19	Compound 43
A10	Compound 19	A20	Compound 46
A11	Compound 21		

The compounds of the present invention are useful for the control of weed species that are resistant to herbicides with the AHAS-inhibitor or (b2) [chemical compound that inhibits acetohydroxy acid synthase (AHAS), also known as acetolactate synthase (ALS)] mode of action.

The following Tests demonstrate the control efficacy of the compounds of this invention against specific weeds. The weed control afforded by the compounds is not limited, however, to these species. See Index Table A for compound descriptions. The following abbreviations are used in the Index Table which follow: *t* is tertiary, *s* is secondary, *n* is normal, *i* is iso, *c* is cyclo, Me is methyl, Et is ethyl, Pr is propyl, *i*-Pr is isopropyl, Bu is butyl, *c*-Pr is cyclopropyl, *t*-Bu is *tert*-butyl, Ph is phenyl, OMe is methoxy, OEt is ethoxy, SMe is methylthio and -CN is cyano. The abbreviation “Cmpd. No.” stands for “Compound Number”. The abbreviation “Ex.” stands for “Example” and is followed by a number indicating in which example the compound is prepared. Mass spectra are reported with an estimated precision within ± 0.5 Da as the molecular weight of the highest isotopic abundance parent ion (M+1) formed by addition of H⁺ (molecular weight of 1) to the molecule. The presence of molecular ions containing one or more higher atomic weight isotopes of lower abundance (e.g., ³⁷Cl, ⁸¹Br) is not reported. The alternate molecular ion peaks (e.g., M+2 or M+4) that occur with compounds containing multiple halogens are not reported. The reported M+1 peaks were observed by mass spectrometry using atmospheric pressure chemical ionization (AP⁺) or electrospray ionization (ESI).

INDEX TABLE A



Cmpd. No.	A	R ¹	R ⁵	R ⁶	M.S.(AP+) or m.p.
1	5-bromo-2-pyrimidinyl	Br	CN	H	450
2	5-fluoro-2-pyrimidinyl	F	CN	H	328
3	5-chloro-2-pyrimidinyl	Cl	CN	H	361
4	5-bromo-2-pyrimidinyl	Cl	CN	H	405
5	5-chloro-2-pyrimidinyl	Br	CN	H	405
6	5-(CF ₃)-2-thiazolyl	Cl	Br	H	145-148
7	3-pyridinyl	Br	CN	H	370
8	3-pyridinyl	Cl	CN	H	325
9	5-bromo-2-pyridinyl	F	CN	H	387
10	5-(CHF ₂)-2-thiazolyl	Cl	Br	H	113-116
11	5-chloro-2-pyridinyl	Br	CN	H	405

Cmpd. No.	A	R ¹	R ⁵	R ⁶	M.S.(AP+)
					or m.p.
12	5-bromo-2-pyridinyl	Br	CN	H	449
13	6-(CF ₃)-3-pyridinyl	Cl	CN	H	393
14	5-(CF ₃)-2-pyridinyl	Cl	CN	H	393
15	5-bromo-2-pyridinyl	Cl	CN	H	405
16	5-chloro-2-pyridinyl	Cl	CN	H	359
17	5-bromo-2-pyrimidinyl	Cl	Cl	H	166-169
18	5-fluoro-2-pyrimidinyl	Cl	Cl	H	353
19	5-chloro-2-pyrimidinyl	Cl	Br	H	415
20	6-chloro-3-pyridinyl	Cl	F	H	352
21	5-chloro-2-pyrimidinyl	Cl	F	H	354
22	5-fluoro-2-pyrimidinyl	Cl	Br	H	113-116
23	5-fluoro-2-pyrimidinyl	Cl	F	H	141-145
24	5-(CF ₃)-2-pyrimidinyl	Cl	Br	H	447
25	5-(CF ₃)-2-pyrimidinyl	Cl	F	F	405
26	6-(CF ₃)-3-pyridazinyl	Cl	CN	H	91-95
27	5-bromo-2-pyrimidinyl	Cl	F	H	145-148
28	5-fluoro-2-pyrimidinyl	Cl	F	F	94-98
29	5-fluoro-2-pyrimidinyl	F	F	F	158-161
30	5-(CF ₃)-2-pyrimidinyl	CF ₃	F	F	74-80
31	5-bromo-2-pyrimidinyl	Br	F	F	166-170
32	5-chloro-2-pyrazinyl	Cl	CN	H	361
33	4-(OCF ₃)-phenyl	Cl	CN	H	76-80
34	5-chloro-2-pyrimidinyl	Cl	F	F	158-161
35	5-(CF ₃)-2-pyrimidinyl	CF ₃	CN	H	428
36	4-chlorophenyl	Cl	CN	H	172-175
37	3-chlorophenyl	Cl	CN	H	122-126
38	5-(CF ₃)-2-pyrimidinyl	Cl	CN	H	101-103
39	5-chloro-2-pyrimidinyl	CF ₃	CN	H	109-112
40	2-chlorophenyl	Cl	CN	H	120-123
41	5-chloro-2-pyrimidinyl	Br	CHF ₂	H	431
42	5-chloro-2-pyrimidinyl	CF ₃	F	H	387
43	5-(CF ₃)-2-pyrimidinyl	Cl	F	H	387
44	5-chloro-2-pyrimidinyl	Cl	CH ₂ F	H	367
45	5-(CF ₃)-2-pyridinyl	Cl	F	H	386
46	5-chloro-2-pyrimidinyl	Cl	CHF ₂	H	385

BIOLOGICAL EXAMPLES OF THE INVENTIONTEST A

Seeds of plant species selected from barnyardgrass (*Echinochloa crus-galli*), kochia (*Kochia scoparia*), ragweed (common ragweed, *Ambrosia elatior*), Italian ryegrass (*Lolium multiflorum*), large (Lg) crabgrass (*Digitaria sanguinalis*), giant foxtail (*Setaria faberii*), morningglory (*Ipomoea* spp.), pigweed (*Amaranthus retroflexus*), velvetleaf (*Abutilon theophrasti*), wheat (*Triticum aestivum*), and corn (*Zea mays*) were planted into a blend of loam soil and sand and treated preemergence with a directed soil spray using test chemicals formulated in a non-phytotoxic solvent mixture which included a surfactant.

At the same time, plants selected from these crop and weed species and also blackgrass (*Alopecurus myosuroides*), and galium (catchweed bedstraw, *Galium aparine*) were planted in pots containing the same blend of loam soil and sand and treated with postemergence applications of test chemicals formulated in the same manner. Plants ranged in height from 2 to 10 cm and were in the one- to two-leaf stage for the postemergence treatment. Treated plants and untreated controls were maintained in a greenhouse for approximately 10 days, after which time all treated plants were compared to untreated controls and visually evaluated for injury. Plant response ratings, summarized in Table A, are based on a 0 to 100 scale where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Table A		Compounds											
1000 g ai/ha		17	18	20	22	23	24	27					
Postemergence													
Barnyardgrass		100	80	90	90	90	90	90					
Blackgrass		100	100	60	70	80	70	80					
Corn		100	50	50	70	50	50	40					
Foxtail, Giant		100	100	90	90	90	90	90					
Galium		100	100	90	90	90	90	90					
Kochia		100	100	90	100	90	90	90					
Pigweed		100	100	100	100	100	100	100					
Ragweed		90	20	60	90	90	60	80					
Ryegrass, Italian		100	100	40	90	90	80	90					
Wheat		100	50	50	70	60	50	50					
Table A		Compounds											
500 g ai/ha		1	2	3	4	11	13	19	21	42	43	46	
Postemergence													
Barnyardgrass		90	90	90	100	100	0	100	90	100	100	100	
Blackgrass		-	-	-	100	100	0	-	-	100	100	100	
Corn		80	90	80	100	100	0	60	60	100	100	100	

70

5	Crabgrass, Large	100	100	90	-	-	-	90	90	-	-	-			
	Foxtail, Giant	90	100	90	100	100	0	90	90	100	100	100			
	Galium	-	-	-	100	100	0	-	-	100	100	100			
	Kochia	-	-	-	100	100	0	-	-	100	100	60			
	Morningglory	30	90	100	-	-	-	100	100	-	-	-			
	Pigweed	100	100	100	100	100	40	100	100	100	100	100			
	Ragweed	-	-	-	100	10	0	-	-	100	100	100			
	Ryegrass, Italian	-	-	-	100	90	0	-	-	100	100	100			
	Velvetleaf	90	90	100	-	-	-	100	100	-	-	-			
10	Wheat	60	90	80	100	90	0	60	60	100	100	100			
	Table A	Compounds													
	125 g ai/ha	1	2	3	4	5	6	7	8	9	10	11	12	13	14
	Postemergence														
15	Barnyardgrass	70	20	90	100	90	100	0	10	50	20	80	20	0	0
	Blackgrass	-	-	-	100	90	40	0	0	80	70	90	50	0	30
	Corn	30	0	50	80	80	10	10	10	10	10	30	20	0	0
	Crabgrass, Large	60	30	90	-	-	-	-	-	-	-	-	-	-	-
	Foxtail, Giant	70	40	90	100	90	60	0	0	70	20	95	40	0	20
20	Galium	-	-	-	100	90	100	10	20	100	100	100	50	0	50
	Kochia	-	-	-	100	100	100	10	30	100	80	95	20	0	0
	Morningglory	30	60	100	-	-	-	-	-	-	-	-	-	-	-
	Pigweed	100	90	100	100	100	90	20	20	100	100	100	100	10	50
	Ragweed	-	-	-	100	40	20	0	0	70	30	10	0	0	0
25	Ryegrass, Italian	-	-	-	100	50	50	0	0	70	60	25	0	0	0
	Velvetleaf	70	70	100	-	-	-	-	-	-	-	-	-	-	-
	Wheat	30	20	60	50	40	10	0	0	40	0	50	20	0	0
	Table A	Compounds													
	125 g ai/ha	15	16	19	21	25	26	28	29	30	31	32	33	34	35
	Postemergence														
30	Barnyardgrass	20	70	70	80	60	20	0	80	0	20	100	10	50	40
	Blackgrass	70	70	-	-	70	30	20	90	30	30	70	10	70	50
	Corn	10	20	50	50	30	20	0	90	0	30	50	10	10	30
	Crabgrass, Large	-	-	70	80	-	-	-	-	-	-	-	-	-	-
	Foxtail, Giant	90	80	80	90	50	30	20	60	10	40	90	20	70	20
35	Galium	90	90	-	-	90	50	70	100	70	100	90	30	90	70
	Kochia	90	100	-	-	70	60	70	100	0	80	90	40	70	90
	Morningglory	-	-	70	90	-	-	-	-	-	-	-	-	-	-
	Pigweed	100	100	100	100	100	80	60	90	70	90	100	80	80	100

71

	Ragweed	10	30	-	-	30	30	30	60	0	0	30	10	50	20
	Ryegrass, Italian	50	80	-	-	50	0	0	20	0	30	50	10	50	30
	Velvetleaf	-	-	100	90	-	-	-	-	-	-	-	-	-	-
	Wheat	10	20	50	50	30	10	0	80	0	30	30	0	30	0
5	Table A	Compounds													
	125 g ai/ha	36	37	38	39	40	41	42	43	44	45	46			
	Postemergence														
	Barnyardgrass	30	0	90	90	0	80	100	90	90	0	90			
	Blackgrass	40	0	90	100	0	80	90	90	80	30	90			
10	Corn	20	0	90	100	0	50	80	70	70	30	90			
	Crabgrass, Large	-	-	-	-	-	-	-	-	-	-	-			
	Foxtail, Giant	50	0	100	100	0	90	100	90	90	0	90			
	Galium	100	0	100	100	10	80	100	100	80	70	100			
	Kochia	100	0	100	100	0	90	100	100	80	80	100			
15	Morningglory	-	-	-	-	-	-	-	-	-	-	-			
	Pigweed	100	0	100	100	30	100	100	100	70	20	100			
	Ragweed	20	0	70	60	0	50	70	70	70	20	100			
	Ryegrass, Italian	10	0	90	90	0	60	100	80	30	0	90			
	Velvetleaf	-	-	-	-	-	-	-	-	-	-	-			
20	Wheat	10	0	90	90	0	20	80	80	80	20	50			
	Table A	Compounds													
	31 g ai/ha	5	6	7	8	9	10	11	12	14	15	16	25	26	28
	Postemergence														
	Barnyardgrass	20	10	0	0	0	0	0	0	0	0	10	10	0	0
25	Blackgrass	40	10	0	0	30	10	10	0	0	0	50	60	0	0
	Corn	20	0	0	0	0	0	10	0	0	10	0	10	0	0
	Foxtail, Giant	10	10	0	0	10	0	0	0	0	20	70	30	10	0
	Galium	50	70	10	10	70	50	70	10	40	50	90	80	30	20
	Kochia	70	70	0	10	60	20	40	0	0	40	50	50	40	30
30	Pigweed	70	50	10	10	50	90	100	10	0	100	100	90	50	50
	Ragweed	20	0	0	0	30	0	0	0	0	10	10	20	30	10
	Ryegrass, Italian	20	0	0	0	20	0	0	0	0	0	40	0	0	0
	Wheat	20	0	0	0	0	0	20	0	0	0	0	0	0	0
	Table A	Compounds													
35	31 g ai/ha	29	30	31	32	33	34	35	36	37	38	39	40	41	44
	Postemergence														
	Barnyardgrass	20	0	10	40	0	0	0	10	0	90	90	0	40	20
	Blackgrass	80	0	20	60	0	60	10	20	0	40	80	0	40	40

	Corn	20	0	20	30	0	0	0	20	0	40	60	0	10	20
	Foxtail, Giant	30	0	20	50	10	30	0	20	0	100	100	0	30	10
	Galium	100	60	70	90	10	80	40	90	0	80	100	0	60	70
	Kochia	80	0	20	90	20	50	60	80	0	100	100	0	50	50
5	Pigweed	90	60	90	100	30	70	70	90	0	100	100	0	100	30
	Ragweed	40	0	0	10	0	30	10	10	0	40	40	0	30	20
	Ryegrass, Italian	20	0	10	10	0	10	0	0	0	50	40	0	30	20
	Wheat	30	0	20	0	0	0	0	0	0	20	10	0	0	20
	Table A	Compound													
10	31 g ai/ha	45													
	Postemergence														
	Barnyardgrass	0													
	Blackgrass	0													
	Corn	0													
15	Foxtail, Giant	0													
	Galium	60													
	Kochia	20													
	Pigweed	20													
	Ragweed	0													
20	Ryegrass, Italian	0													
	Wheat	0													
	Table A	Compounds													
	1000 g ai/ha	17	18	20	22	23	24	27							
	Preemergence														
25	Barnyardgrass	100	100	100	100	100	100	100							
	Foxtail, Giant	100	100	100	100	100	100	100							
	Kochia	100	100	90	100	100	90	100							
	Pigweed	100	100	100	100	100	100	100							
	Ragweed	100	30	40	90	100	70	90							
30	Ryegrass, Italian	100	70	70	90	80	80	70							
	Table A	Compounds													
	500 g ai/ha	1	2	3	4	11	13	19	21	42	43	46			
	Preemergence														
	Barnyardgrass	90	100	100	100	100	0	90	100	100	100	100			
35	Corn	60	60	70	-	-	-	60	60	-	-	-			
	Crabgrass, Large	100	100	100	-	-	-	100	100	-	-	-			
	Foxtail, Giant	100	100	100	100	100	0	100	100	100	100	100			
	Kochia	-	-	-	100	100	0	-	-	100	100	100			

73

5	Morningglory	100	100	100	-	-	-	100	100	-	-	-				
	Pigweed	100	100	100	100	100	0	100	100	100	100	100				
	Ragweed	-	-	-	100	10	0	-	-	100	100	100				
	Ryegrass, Italian	-	-	-	100	90	0	-	-	100	100	80				
	Velvetleaf	100	100	100	-	-	-	90	90	-	-	-				
	Wheat	60	30	80	-	-	-	70	60	-	-	-				
	Table A	Compounds														
	125 g ai/ha	1	2	3	4	5	6	7	8	9	10	11	12	13	14	
10	Preemergence															
	Barnyardgrass	70	20	100	70	80	70	0	0	90	30	80	30	0	0	
	Corn	20	20	60	-	-	-	-	-	-	-	-	-	-	-	
	Crabgrass, Large	100	100	100	-	-	-	-	-	-	-	-	-	-	-	
	Foxtail, Giant	90	100	100	100	100	100	0	0	100	90	100	70	0	0	
15	Kochia	-	-	-	90	100	80	0	0	100	50	75	20	0	0	
	Morningglory	30	50	100	-	-	-	-	-	-	-	-	-	-	-	
	Pigweed	100	100	100	100	100	90	0	10	100	100	100	100	0	80	
	Ragweed	-	-	-	80	20	0	0	0	10	0	0	0	0	0	
	Ryegrass, Italian	-	-	-	20	60	20	0	0	0	40	10	0	0	0	
20	Velvetleaf	70	40	100	-	-	-	-	-	-	-	-	-	-	-	
	Wheat	20	20	30	-	-	-	-	-	-	-	-	-	-	-	
	Table A	Compounds														
		125 g ai/ha	15	16	19	21	25	26	28	29	30	31	32	33	34	35
		Preemergence														
25	Barnyardgrass	60	70	80	90	70	20	10	100	10	70	90	10	80	50	
	Corn	-	-	10	40	-	-	-	-	-	-	-	-	-	-	
	Crabgrass, Large	-	-	100	100	-	-	-	-	-	-	-	-	-	-	
	Foxtail, Giant	100	100	100	100	90	60	40	100	50	90	100	40	90	90	
	Kochia	100	90	-	-	100	30	20	100	20	90	100	0	90	60	
30	Morningglory	-	-	70	90	-	-	-	-	-	-	-	-	-	-	
	Pigweed	100	100	100	100	100	100	80	100	100	100	100	30	100	100	
	Ragweed	0	0	-	-	0	20	0	90	0	20	30	0	10	10	
	Ryegrass, Italian	10	20	-	-	40	0	0	30	0	0	60	0	50	10	
	Velvetleaf	-	-	80	80	-	-	-	-	-	-	-	-	-	-	
35	Wheat	-	-	40	60	-	-	-	-	-	-	-	-	-	-	
	Table A	Compounds														
		125 g ai/ha	36	37	38	39	40	41	42	43	44	45	46			
		Preemergence														
		Barnyardgrass	40	0	100	100	0	80	100	100	100	30	90			

[illegible]

TEST B

Plant species in the flooded paddy test selected from rice (*Oryza sativa*), sedge, umbrella (small-flower umbrella sedge, *Cyperus difformis*), ducksalad (*Heteranthera limosa*), and barnyardgrass (*Echinochloa crus-galli*) were grown to the 2-leaf stage for testing. At time of treatment, test pots were flooded to 3 cm above the soil surface, treated by application of test compounds directly to the paddy water, and then maintained at that water depth for the duration of the test. Treated plants and controls were maintained in a greenhouse for 13 to 15 days, after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in Table B, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Table B

Compounds

250 g ai/ha	1	2	3	4	5	10	11	13	14	15	16	17	18	19
Flood														
Barnyardgrass	40	20	20	0	35	0	0	0	0	0	20	20	50	15
Ducksalad	90	70	90	95	100	0	100	0	30	75	90	100	100	95
Rice	0	15	35	0	0	0	0	0	15	0	40	35	80	25
Sedge, Umbrella	95	60	90	80	75	0	85	0	0	90	95	100	100	85

Table B

Compounds

250 g ai/ha	20	21	22	23	24	25	26	27	29	30	31	32	33	34
Flood														
Barnyardgrass	20	35	45	60	50	55	0	50	90	0	0	70	0	0
Ducksalad	95	100	100	90	100	90	70	80	90	0	50	75	0	30
Rice	15	40	45	60	90	20	0	60	85	0	0	40	0	0
Sedge, Umbrella	90	90	98	80	100	75	60	85	75	0	50	90	0	30

Table B

Compounds

250 g ai/ha	35	36	37	38	39	40	41	42	43	44	45	46
Flood												
Barnyardgrass	0	0	0	85	50	0	0	100	85	40	0	0
Ducksalad	60	65	0	100	100	0	30	100	100	95	75	80
Rice	40	0	0	45	50	0	0	80	60	40	0	10
Sedge, Umbrella	60	95	0	100	100	0	40	100	90	75	70	65

TEST C

Seeds of plant species selected from blackgrass (*Alopecurus myosuroides*), Italian ryegrass (*Lolium multiflorum*), winter wheat (*Triticum aestivum*), galium (catchweed bedstraw, *Galium aparine*), corn (*Zea mays*), large (Lg) crabgrass (*Digitaria sanguinalis*), giant foxtail (*Setaria faberii*), johnsongrass (*Sorghum halepense*), lambsquarters

(*Chenopodium album*), morningglory (*Ipomoea coccinea*), yellow nutsedge (*Cyperus esculentus*), pigweed (*Amaranthus retroflexus*), ragweed (common ragweed, *Ambrosia elatior*), soybean (*Glycine max*), barnyardgrass (*Echinochloa crus-galli*), oilseed rape (*Brassica napus*), waterhemp (common waterhemp, *Amaranthus rudis*), and velvetleaf (*Abutilon theophrasti*) were planted into a blend of loam soil and sand and treated preemergence with test chemicals formulated in a non-phytotoxic solvent mixture which included a surfactant.

At the same time, plants selected from these crop and weed species and also chickweed (common chickweed, *Stellaria media*), kochia (*Kochia scoparia*), and wild oat (*Avena fatua*), were planted in pots containing Redi-Earth[®] planting medium (Scotts Company, 14111 Scottslawn Road, Marysville, Ohio 43041) comprising sphagnum peat moss, vermiculite, wetting agent and starter nutrients and treated with postemergence applications of test chemicals formulated in the same manner. Plants ranged in height from 2 to 18 cm (1- to 4-leaf stage) for postemergence treatments. Treated plants and controls were maintained in a greenhouse for 13 to 15 days, after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in Table C, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Plant species in the flooded paddy test consisted of rice (*Oryza sativa*), sedge, umbrella (small-flower umbrella sedge, *Cyperus difformis*), ducksalad (*Heteranthera limosa*), and barnyardgrass (*Echinochloa crus-galli*) grown to the 2-leaf stage for testing. At time of treatment, test pots were flooded to 3 cm above the soil surface, treated by application of test compounds directly to the paddy water, and then maintained at that water depth for the duration of the test. Treated plants and controls were maintained in a greenhouse for 13 to 15 days, after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in Table C, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Table C

Compounds

250 g ai/ha	3	17	18	20	22	23	27
Postemergence							
Barnyardgrass	60	10	10	10	10	25	5
Blackgrass	30	15	45	15	10	30	5
Chickweed	95	85	95	80	80	60	80
Corn	75	15	55	15	10	10	25
Crabgrass, Large	85	10	70	20	20	30	20
Foxtail, Giant	60	40	45	15	20	35	10
Galium	95	95	95	95	95	95	95
Johnsongrass	50	10	10	5	5	5	10

5	Kochia	95	95	95	98	90	90	95							
	Lambsquarters	100	95	95	80	98	95	95							
	Morningglory	98	70	95	65	80	95	90							
	Nutsedge, Yellow	35	20	10	10	10	5	10							
	Oat, Wild	35	15	30	10	15	5	15							
	Oilseed Rape	85	40	80	70	35	45	60							
	Pigweed	100	98	98	98	100	100	98							
	Ragweed	85	55	95	60	90	100	80							
	Ryegrass, Italian	35	15	40	15	5	35	5							
10	Soybean	95	90	95	65	0	95	90							
	Velvetleaf	80	70	85	55	70	90	65							
	Waterhemp	98	98	98	95	95	98	100							
	Wheat	15	0	15	5	0	5	0							
Table C															
15	Compounds														
	125 g ai/ha	3	4	16	17	18	20	21	22	23	27	29	32	34	36
	Postemergence														
	Barnyardgrass	20	5	20	10	5	10	5	15	10	5	15	30	10	20
	Blackgrass	30	10	10	15	5	5	70	5	5	5	20	10	5	30
20	Chickweed	90	80	80	60	80	45	100	55	60	70	65	85	95	100
	Corn	-	10	15	20	20	20	5	10	10	20	5	25	5	20
	Crabgrass, Large	15	10	20	10	55	10	15	25	10	10	10	30	5	20
	Foxtail, Giant	15	35	35	10	25	15	20	10	10	5	20	20	25	10
	Galium	95	95	95	90	95	70	90	90	95	95	98	95	85	95
25	Johnsongrass	15	5	10	5	5	5	5	5	5	5	20	40	10	35
	Kochia	95	100	95	95	85	90	98	85	80	85	95	90	90	100
	Lambsquarters	95	90	90	90	95	75	90	95	90	80	85	90	80	100
	Morningglory	90	95	95	85	85	65	98	85	85	70	95	95	95	70
	Nutsedge, Yellow	5	10	15	5	10	10	5	10	5	10	0	20	10	15
30	Oat, Wild	35	10	5	5	5	10	40	10	0	10	35	30	10	35
	Oilseed Rape	70	60	50	80	50	80	80	60	50	55	90	85	70	98
	Pigweed	90	95	100	98	95	90	95	90	95	95	90	95	90	100
	Ragweed	60	85	70	60	90	45	95	90	75	60	80	80	60	75
	Ryegrass, Italian	5	5	15	5	0	10	25	5	0	5	20	30	0	30
35	Soybean	95	70	55	90	90	65	95	90	90	90	85	85	40	45
	Velvetleaf	70	60	70	40	70	35	70	70	75	65	80	75	65	80
	Waterhemp	85	70	90	95	95	90	65	95	90	90	95	95	90	98
	Wheat	10	5	0	0	0	5	5	0	0	0	0	0	0	5
	Table C														
Compounds															

78

	125 g ai/ha	38	39	42	43	46										
	Postemergence															
	Barnyardgrass	35	25	20	35	5										
	Blackgrass	15	30	80	95	5										
5	Chickweed	85	90	95	100	75										
	Corn	15	20	25	5	10										
	Crabgrass, Large	30	10	40	40	10										
	Foxtail, Giant	25	20	35	35	25										
	Galium	90	90	100	90	65										
10	Johnsongrass	10	10	40	45	5										
	Kochia	95	100	100	100	100										
	Lambsquarters	90	85	95	95	98										
	Morningglory	90	85	85	80	90										
	Nutsedge, Yellow	10	5	5	5	5										
15	Oat, Wild	10	15	60	60	10										
	Oilseed Rape	5	40	85	50	65										
	Pigweed	90	95	98	98	85										
	Ragweed	50	85	65	80	95										
	Ryegrass, Italian	5	10	35	15	10										
20	Soybean	80	70	85	70	80										
	Velvetleaf	60	60	85	85	65										
	Waterhemp	95	95	95	90	5										
	Wheat	0	0	10	5	10										
	Table C	Compounds														
25	62 g ai/ha	3	4	16	17	18	20	21	22	23	25	27	29	32	34	
	Postemergence															
	Barnyardgrass	10	5	15	5	5	5	5	5	10	10	5	5	20	10	
	Blackgrass	20	0	5	5	5	10	20	5	5	5	5	5	5	0	
	Chickweed	85	50	80	70	55	40	60	50	50	60	40	50	80	80	
30	Corn	35	5	10	20	5	15	5	10	5	5	15	5	25	5	
	Crabgrass, Large	10	5	15	5	5	5	5	15	5	5	5	5	20	5	
	Foxtail, Giant	15	45	20	15	5	20	20	5	5	5	5	5	15	5	
	Galium	70	60	70	80	90	50	85	90	90	80	90	90	95	85	
	Johnsongrass	15	5	10	5	10	5	5	0	0	5	5	20	10	10	
35	Kochia	90	80	90	90	50	90	90	60	70	75	80	90	90	90	
	Lambsquarters	95	80	75	70	90	75	98	75	80	75	65	85	85	70	
	Morningglory	90	85	90	90	70	60	70	70	75	15	75	90	95	90	
	Nutsedge, Yellow	0	10	10	5	10	5	5	5	0	10	5	0	20	10	

	Oat, Wild	30	10	5	0	0	10	0	5	0	15	5	30	30	5
	Oilseed Rape	60	45	80	50	35	70	60	40	30	45	50	70	85	70
	Pigweed	80	80	85	95	90	80	85	85	80	95	90	95	90	90
	Ragweed	55	75	50	45	75	20	60	35	70	25	35	75	80	60
5	Ryegrass, Italian	5	0	5	0	0	5	0	0	0	5	0	35	10	0
	Soybean	90	70	45	80	85	70	95	75	65	45	75	75	85	85
	Velvetleaf	70	50	60	35	60	30	65	65	70	30	35	75	60	60
	Waterhemp	75	75	95	95	85	90	50	85	75	75	75	90	85	80
	Wheat	0	5	5	0	0	0	0	0	0	0	0	0	0	0
10	Table C	Compounds													
	62 g ai/ha	36	38	39	42	43	46								
	Postemergence														
	Barnyardgrass	15	20	15	10	10	5								
	Blackgrass	30	10	15	20	75	0								
15	Chickweed	80	70	90	95	70	60								
	Corn	20	15	10	10	5	5								
	Crabgrass, Large	10	10	10	15	10	10								
	Foxtail, Giant	10	10	10	35	15	50								
	Galium	95	90	80	100	85	60								
20	Johnsongrass	35	5	10	10	10	5								
	Kochia	95	90	100	100	100	95								
	Lambsquarters	98	80	80	90	90	85								
	Morningglory	65	85	90	75	80	85								
	Nutsedge, Yellow	10	5	5	5	0	5								
25	Oat, Wild	20	10	10	40	20	5								
	Oilseed Rape	90	25	40	70	65	50								
	Pigweed	98	90	90	98	90	70								
	Ragweed	40	70	60	70	65	65								
	Ryegrass, Italian	10	5	5	10	10	5								
30	Soybean	50	50	95	65	75	75								
	Velvetleaf	70	55	60	70	75	55								
	Waterhemp	100	85	85	95	80	70								
	Wheat	0	0	5	0	0	5								
	Table C	Compounds													
35	31 g ai/ha	3	4	16	17	18	20	21	22	23	25	27	29	32	34
	Postemergence														
	Barnyardgrass	5	5	15	5	5	5	5	5	5	5	5	0	20	5
	Blackgrass	20	0	5	5	0	5	5	5	0	5	5	0	5	0

5	Ryegrass, Italian	5	5	0	0	0	5								
	Soybean	25	85	60	55	55	65								
	Velvetleaf	45	65	55	60	55	25								
	Waterhemp	90	75	75	90	70	50								
	Wheat	0	0	0	0	0	5								
	Table C	Compounds													
16 g ai/ha	4	16	21	25	29	32	34	36	38	39	42	43	46		
10	Postemergence														
	Barnyardgrass	5	10	5	5	0	10	5	10	10	5	5	5	0	
	Blackgrass	0	5	5	0	0	5	0	5	0	5	10	0	0	
	Chickweed	20	25	50	20	45	70	50	70	50	60	20	40	35	
	Corn	5	10	0	5	0	20	5	10	10	5	5	5	5	
	Crabgrass, Large	5	10	5	0	0	5	0	5	5	5	10	5	5	
15	Foxtail, Giant	40	10	5	0	0	20	5	5	5	10	30	5	35	
	Galium	55	60	70	55	85	80	80	80	70	60	75	35	60	
	Johnsongrass	5	5	5	5	0	10	5	20	5	5	10	0	5	
	Kochia	60	90	85	50	80	90	60	90	90	90	90	90	70	
	Lambsquarters	30	65	15	80	75	80	55	90	65	75	80	60	55	
	Morningglory	70	85	75	10	70	85	70	70	85	5	60	45	65	
20	Nutsedge, Yellow	5	5	10	0	0	5	0	5	0	0	0	0	0	
	Oat, Wild	5	5	0	5	0	20	0	20	5	5	5	0	5	
	Oilseed Rape	10	50	50	35	60	80	45	65	5	5	15	30	5	
	Pigweed	70	85	75	80	75	80	70	90	65	60	85	70	70	
	Ragweed	30	25	40	10	65	60	50	15	55	10	30	25	10	
	Ryegrass, Italian	0	5	0	5	0	0	0	0	0	0	0	0	0	
25	Soybean	30	65	55	45	70	65	35	10	65	75	40	35	40	
	Velvetleaf	25	40	65	40	70	70	35	65	45	60	60	70	15	
	Waterhemp	35	85	30	60	70	75	60	90	75	70	70	65	65	
	Wheat	0	0	0	5	0	0	0	0	0	0	0	0	0	
	Table C	Compound					Table C					Compound			
	8 g ai/ha	25					8 g ai/ha					25			
	Postemergence	Postemergence													
	Barnyardgrass	5					Nutsedge, Yellow						0		
	Blackgrass	0					Oat, Wild						5		
	Chickweed	0					Oilseed Rape						30		
	Corn	5					Pigweed						75		
	Crabgrass, Large	0					Ragweed						0		
	Foxtail, Giant	0					Ryegrass, Italian						0		

Table C Compound

8 g ai/ha 25

Postemergence

Galium 50

Johnsongrass 0

Kochia 0

Lambsquarters 60

Morningglory 20

Table C

Compounds

250 g ai/ha 3 17 18 22 23 24 27

Preemergence

Barnyardgrass 100 100 100 100 100 100 100

5 Blackgrass 90 95 95 95 95 95 95

Corn 90 80 95 95 95 65 75

Crabgrass, Large 100 100 100 100 100 100 100

Foxtail, Giant 100 100 100 100 100 100 100

Galium 95 100 100 100 100 100 100

10 Johnsongrass 100 90 100 98 100 100 100

Lambsquarters 100 100 100 98 98 98 100

Morningglory 100 98 100 100 98 95 95

Nutsedge, Yellow 75 80 85 90 85 60 75

Oilseed Rape 95 98 100 100 100 100 100

15 Pigweed 100 100 100 100 100 100 100

Ragweed 100 85 95 95 100 80 85

Ryegrass, Italian 90 95 95 90 90 95 95

Soybean 95 70 90 95 95 85 90

Velvetleaf 100 100 100 100 100 90 100

20 Waterhemp 100 100 100 100 100 100 100

Wheat 90 85 85 90 90 90 85

Table C

Compounds

125 g ai/ha 3 4 5 15 17 18 19 21 22 23 24 27 29 32

Preemergence

25 Barnyardgrass 100 100 98 98 100 100 100 100 100 100 100 100 100 100

Blackgrass 90 95 70 80 95 95 95 90 90 95 95 90 90 95

Corn 85 35 75 40 55 60 90 98 80 70 45 60 65 65

Crabgrass, Large 100 100 100 100 100 100 100 100 100 100 100 100 98 100

Foxtail, Giant 100 100 100 100 100 100 100 100 100 100 100 100 98 100

30 Galium 90 100 100 100 100 100 100 100 100 100 100 100 100 95

83

5	Johnsongrass	100	98	80	-	85	90	98	98	70	90	80	75	85	100	
	Lambsquarters	100	100	100	95	95	95	100	98	100	100	98	100	90	100	
	Morningglory	100	98	95	70	98	70	100	98	95	98	80	98	85	90	
	Nutsedge, Yellow	55	55	60	25	55	25	55	85	60	60	30	55	40	75	
	Oilseed Rape	95	100	100	100	100	100	100	100	100	100	100	100	95	95	
	Pigweed	100	100	100	100	100	100	100	100	100	100	100	100	100	100	
	Ragweed	90	70	100	40	55	95	60	85	90	100	30	70	95	50	
	Ryegrass, Italian	90	70	60	60	90	85	90	90	80	85	95	80	60	90	
	Soybean	85	95	85	60	45	90	80	98	90	90	50	-	90	90	
10	Velvetleaf	100	100	100	100	100	100	100	100	100	100	90	100	98	85	
	Waterhemp	100	100	100	100	100	100	100	100	100	100	100	100	100	100	
	Wheat	90	5	35	40	40	70	85	90	20	50	60	40	60	5	
Table C		Compounds														
15	125 g ai/ha	34	38	39	42	43	46									
	Preemergence															
	Barnyardgrass	98	100	100	100	100	100									
	Blackgrass	70	90	90	90	95	95									
	Corn	40	65	85	85	90	90									
20	Crabgrass, Large	100	100	100	100	100	100									
	Foxtail, Giant	100	100	100	100	100	100									
	Galium	98	100	100	98	100	100									
	Johnsongrass	98	100	100	100	100	100									
25	Lambsquarters	98	98	95	100	100	100									
	Morningglory	90	100	100	100	100	100									
	Nutsedge, Yellow	5	85	85	75	60	80									
	Oilseed Rape	90	100	100	100	100	100									
	Pigweed	100	100	100	100	100	100									
	Ragweed	40	90	85	90	90	100									
	Ryegrass, Italian	35	90	90	95	95	100									
	Soybean	55	90	95	95	90	95									
	Velvetleaf	80	100	100	100	100	100									
30	Waterhemp	100	100	100	100	100	100									
	Wheat	10	35	60	90	90	50									
	Table C		Compounds													
	35	62 g ai/ha	3	4	5	15	17	18	19	21	22	23	24	27	29	32
Preemergence																
Barnyardgrass		100	85	85	70	85	100	100	100	95	100	100	100	98	98	
	Blackgrass	90	95	40	10	90	90	90	90	60	85	95	60	90	90	

[illegible]

85

	31 g ai/ha	3	4	5	15	17	18	19	21	22	23	24	27	29	32
	Preemergence														
	Barnyardgrass	80	60	60	25	20	75	55	85	60	75	75	55	75	85
	Blackgrass	70	30	30	0	60	15	40	30	10	65	60	15	90	70
5	Corn	20	5	25	0	0	10	25	25	10	5	5	10	15	60
	Crabgrass, Large	100	100	98	100	100	100	100	100	100	100	100	98	95	98
	Foxtail, Giant	100	98	80	80	95	98	100	100	100	95	98	80	80	98
	Galium	90	30	50	100	90	100	100	100	98	100	98	100	100	95
	Johnsongrass	45	15	40	-	10	50	50	50	10	40	40	25	70	85
10	Lambsquarters	100	95	100	80	85	85	100	100	75	100	75	80	85	90
	Morningglory	45	35	60	5	40	10	45	80	0	45	25	5	75	10
	Nutsedge, Yellow	5	0	5	0	0	0	5	5	0	0	10	0	0	35
	Oilseed Rape	90	75	70	98	85	100	85	100	85	100	80	85	98	95
	Pigweed	100	100	100	100	100	100	100	100	100	100	100	100	100	100
15	Ragweed	90	10	20	30	0	85	30	85	55	85	10	20	85	20
	Ryegrass, Italian	5	5	5	5	20	30	50	50	5	5	55	10	15	50
	Soybean	40	30	45	30	5	60	50	90	60	60	-	40	85	45
	Velvetleaf	80	45	90	75	65	85	98	100	70	100	40	60	98	40
	Waterhemp	100	100	100	100	100	100	100	100	100	100	100	100	100	100
20	Wheat	5	0	0	0	0	0	25	40	0	5	15	0	5	0
	Table C	Compounds													
	31 g ai/ha	34	38	39	42	43	46								
	Preemergence														
	Barnyardgrass	55	85	95	90	85	70								
25	Blackgrass	0	35	50	65	50	70								
	Corn	5	30	40	5	20	25								
	Crabgrass, Large	85	100	100	100	100	100								
	Foxtail, Giant	98	95	95	100	100	100								
	Galium	95	90	90	95	95	85								
30	Johnsongrass	55	85	65	70	75	40								
	Lambsquarters	90	90	90	100	100	100								
	Morningglory	20	80	80	85	85	90								
	Nutsedge, Yellow	0	20	20	10	0	35								
	Oilseed Rape	90	100	90	100	100	90								
35	Pigweed	100	100	98	100	100	100								
	Ragweed	0	25	40	55	80	25								
	Ryegrass, Italian	5	10	20	70	50	25								
	Soybean	20	55	65	50	45	40								

[illegible]

Table C	Compounds
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62 g ai/ha	1	4	20
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Flood			
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Barnyardgrass	0	0	0
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5 Ducksalad	0	40	0
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Rice	0	0	0
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Sedge, Umbrella	30	0	0
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Table C	Compounds
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31 g ai/ha	1	4	20
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10 Flood			
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Barnyardgrass	0	0	0
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Ducksalad	0	0	0
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Rice	0	0	0
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Sedge, Umbrella	0	0	0
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15 TEST D

Seeds of plant species selected from bluegrass (annual bluegrass, *Poa annua*), blackgrass (*Alopecurus myosuroides*), canarygrass (*Phalaris minor*), chickweed (common chickweed, *Stellaria media*), cutleaf geranium (*Geranium dissectum*), galium (catchweed bedstraw, *Galium aparine*), downy brome grass (*Bromus tectorum*), field poppy (*Papaver rhoeas*), field violet (*Viola arvensis*), green foxtail (*Setaria viridis*), deadnettle (henbit deadnettle, *Lamium amplexicaule*), Italian ryegrass (*Lolium multiflorum*), kochia (*Kochia scoparia*), lambsquarters (*Chenopodium album*), oilseed rape (*Brassica napus*), pigweed (*Amaranthus retroflexus*), chamomile (scentless chamomile, *Matricaria inodora*), Russian thistle (*Salsola kali*), speedwell (bird's-eye speedwell, *Veronica persica*), spring barley (*Hordeum vulgare*), spring wheat (*Triticum aestivum*), wild buckwheat (*Polygonum convolvulus*), wild mustard (*Sinapis arvensis*), wild oat (*Avena fatua*), wild radish (*Raphanus raphanistrum*), windgrass (*Apera spica-venti*), winter barley (*Hordeum vulgare*), and winter wheat (*Triticum aestivum*) were planted into a silt loam soil and treated preemergence with test chemicals formulated in a non-phytotoxic solvent mixture which included a surfactant.

At the same time, these species were planted in pots containing Redi-Earth[®] planting medium (Scotts Company, 14111 Scottslawn Road, Marysville, Ohio 43041) comprising sphagnum peat moss, vermiculite, wetting agent and starter nutrients and treated with postemergence applications of the test chemicals formulated in the same manner. Plants ranged in height from 2 to 18 cm (1- to 4-leaf stage). Treated plants and controls were maintained in a controlled growth environment for 14 to 21 days after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in

Table D, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Table D		Compounds				
125 g ai/ha		3	22	23	24	42
5	Postemergence					
	Barley, Spring	20	20	10	25	30
	Barley, Winter	15	15	5	25	15
	Blackgrass	20	40	55	45	75
	Bluegrass	30	40	35	40	70
10	Bromegrass, Downy	25	25	25	25	60
	Buckwheat, Wild	100	80	75	95	100
	Canarygrass	20	25	55	45	75
	Chamomile	35	5	5	15	15
	Chickweed	100	75	80	85	90
15	Deadnettle	80	90	70	85	75
	Field Poppy	70	70	80	95	100
	Field Violet	100	75	70	98	95
	Foxtail, Green	40	30	20	30	70
	Galium	100	85	85	90	100
20	Geranium, Cutleaf	–	60	65	65	–
	Kochia	100	70	50	95	100
	Lambsquarters	100	70	60	75	95
	Mustard, Wild	85	75	95	85	100
	Oat, Wild	20	45	35	35	70
25	Oilseed Rape	90	80	95	80	95
	Pigweed	100	95	90	100	100
	Radish, Wild	85	90	85	100	90
	Russian Thistle	–	15	15	55	70
	Ryegrass, Italian	15	35	25	35	65
30	Speedwell	100	90	100	90	80
	Wheat, Spring	15	15	10	20	15
	Wheat, Winter	15	10	0	15	20
	Windgrass	30	35	20	35	60
Table D		Compounds				
35	62 g ai/ha	3	22	23	24	42
	Postemergence					
	Barley, Spring	15	15	5	20	20
	Barley, Winter	10	10	5	20	10

	Blackgrass	15	35	35	35	60
	Bluegrass	25	15	15	35	60
	Bromegrass, Downy	25	20	20	20	40
	Buckwheat, Wild	95	70	75	75	95
5	Canarygrass	15	20	25	45	70
	Chamomile	20	0	0	10	5
	Chickweed	80	75	70	75	85
	Deadnettle	70	65	55	75	60
	Field Poppy	55	70	75	90	100
10	Field Violet	100	70	60	85	90
	Foxtail, Green	20	25	15	20	60
	Galium	100	75	85	85	90
	Geranium, Cutleaf	-	55	45	45	-
	Kochia	100	60	30	80	100
15	Lambsquarters	100	75	60	60	90
	Mustard, Wild	65	55	65	70	100
	Oat, Wild	15	35	20	25	40
	Oilseed Rape	60	70	75	75	85
	Pigweed	100	75	70	98	100
20	Radish, Wild	75	90	80	95	85
	Russian Thistle	-	10	10	30	25
	Ryegrass, Italian	10	25	10	25	60
	Speedwell	100	75	100	80	80
	Wheat, Spring	5	10	5	20	5
25	Wheat, Winter	10	5	0	15	10
	Windgrass	25	25	15	25	50
	Table D	Compounds				
	31 g ai/ha	3	22	23	24	42
	Postemergence					
30	Barley, Spring	10	10	5	20	15
	Barley, Winter	5	10	5	15	5
	Blackgrass	10	30	20	30	55
	Bluegrass	20	5	10	20	50
	Bromegrass, Downy	20	10	10	20	35
35	Buckwheat, Wild	90	65	70	60	80
	Canarygrass	10	10	20	35	65
	Chamomile	0	0	0	5	0
	Chickweed	70	75	70	70	65

	Deadnettle	70	50	20	70	40
	Field Poppy	40	65	70	80	75
	Field Violet	100	40	60	75	65
	Foxtail, Green	15	15	10	15	15
5	Galium	95	60	70	70	85
	Geranium, Cutleaf	-	55	45	30	-
	Kochia	100	45	20	65	100
	Lambsquarters	80	40	30	45	85
	Mustard, Wild	60	40	35	80	70
10	Oat, Wild	15	25	15	20	45
	Oilseed Rape	55	45	65	70	75
	Pigweed	100	80	60	95	100
	Radish, Wild	60	35	80	75	75
	Russian Thistle	-	5	5	25	10
15	Ryegrass, Italian	10	20	5	20	50
	Speedwell	85	40	75	60	50
	Wheat, Spring	0	5	5	15	5
	Wheat, Winter	10	5	0	10	5
	Windgrass	15	10	10	15	40
20	Table D	Compounds				
	16 g ai/ha	3	22	23	24	42
	Postemergence					
	Barley, Spring	5	5	0	15	10
	Barley, Winter	0	5	0	10	5
25	Blackgrass	5	20	10	25	20
	Bluegrass	15	0	5	15	50
	Bromegrass, Downy	10	5	5	15	30
	Buckwheat, Wild	85	55	50	60	75
	Canarygrass	5	10	10	30	35
30	Chamomile	0	0	0	5	0
	Chickweed	50	30	60	60	65
	Deadnettle	60	50	10	55	35
	Field Poppy	35	55	50	70	55
	Field Violet	90	25	15	65	65
35	Foxtail, Green	10	10	10	15	10
	Galium	75	70	65	40	70
	Geranium, Cutleaf	-	45	25	20	-
	Kochia	85	40	15	40	85

	Lambsquarters	70	50	10	35	70
	Mustard, Wild	45	35	25	50	70
	Oat, Wild	15	10	15	15	25
	Oilseed Rape	40	15	30	65	75
5	Pigweed	100	75	55	85	100
	Radish, Wild	50	25	25	75	80
	Russian Thistle	-	0	0	10	10
	Ryegrass, Italian	0	15	0	15	20
	Speedwell	85	30	40	35	40
10	Wheat, Spring	0	5	0	10	0
	Wheat, Winter	0	0	0	10	0
	Windgrass	10	5	10	10	25
	Table D	Compounds				
	125 g ai/ha	3	22	23	24	42
15	Preemergence					
	Barley, Spring	40	65	75	85	65
	Barley, Winter	70	40	70	70	80
	Blackgrass	70	55	85	55	75
	Bluegrass	85	90	100	35	90
20	Bromegrass, Downy	15	35	70	45	65
	Buckwheat, Wild	100	85	100	100	100
	Canarygrass	80	55	85	45	100
	Chamomile	85	100	100	100	-
	Chickweed	100	100	100	100	100
25	Deadnettle	100	100	95	100	100
	Field Poppy	100	100	98	100	-
	Field Violet	100	95	100	100	100
	Foxtail, Green	100	40	100	85	100
	Galium	100	100	100	70	100
30	Geranium, Cutleaf	-	100	100	100	-
	Kochia	100	70	100	100	100
	Lambsquarters	100	100	100	100	100
	Mustard, Wild	100	100	100	100	100
	Oat, Wild	50	25	98	75	95
35	Oilseed Rape	100	90	100	80	100
	Pigweed	100	100	100	100	100
	Radish, Wild	100	80	100	70	100
	Russian Thistle	-	5	5	80	55

	Ryegrass, Italian	40	65	75	65	70
	Speedwell	100	100	100	100	-
	Wheat, Spring	20	35	45	40	65
	Wheat, Winter	10	40	70	10	50
5	Windgrass	80	70	100	85	100
	Table D	Compounds				
	62 g ai/ha	3	22	23	24	42
	Preemergence					
	Barley, Spring	0	15	30	30	15
10	Barley, Winter	20	5	30	25	15
	Blackgrass	35	35	70	35	35
	Bluegrass	25	60	80	25	45
	Bromegrass, Downy	5	30	35	35	50
	Buckwheat, Wild	75	15	100	20	100
15	Canarygrass	20	35	80	35	35
	Chamomile	35	55	75	100	-
	Chickweed	100	100	100	100	100
	Deadnettle	100	85	80	100	100
	Field Poppy	100	100	90	100	-
20	Field Violet	100	95	95	75	100
	Foxtail, Green	100	40	55	75	100
	Galium	100	75	95	85	100
	Geranium, Cutleaf	-	70	100	90	-
	Kochia	100	20	60	60	100
25	Lambsquarters	100	100	100	100	100
	Mustard, Wild	95	100	100	55	85
	Oat, Wild	0	20	70	25	50
	Oilseed Rape	35	70	98	20	100
	Pigweed	100	100	100	100	100
30	Radish, Wild	100	70	90	45	85
	Russian Thistle	-	5	5	35	15
	Ryegrass, Italian	50	25	35	25	35
	Speedwell	100	100	100	100	-
	Wheat, Spring	10	15	30	0	15
35	Wheat, Winter	5	5	10	0	0
	Windgrass	50	25	98	25	90
	Table D	Compounds				
	31 g ai/ha	3	22	23	24	42

	Preemergence					
	Barley, Spring	0	5	5	20	35
	Barley, Winter	0	0	0	15	0
	Blackgrass	10	20	30	25	10
5	Bluegrass	0	20	65	20	0
	Bromegrass, Downy	0	25	25	35	10
	Buckwheat, Wild	30	10	50	-	0
	Canarygrass	5	25	65	30	35
	Chamomile	20	40	75	100	-
10	Chickweed	100	25	100	100	100
	Deadnettle	95	60	65	60	85
	Field Poppy	100	95	80	100	-
	Field Violet	100	75	25	20	100
	Foxtail, Green	60	35	5	35	60
15	Galium	100	65	75	25	70
	Geranium, Cutleaf	-	35	100	75	-
	Kochia	100	10	60	25	65
	Lambsquarters	95	85	100	100	100
	Mustard, Wild	60	30	100	35	65
20	Oat, Wild	0	10	15	15	5
	Oilseed Rape	25	40	85	15	70
	Pigweed	100	70	100	100	100
	Radish, Wild	80	55	70	20	-
	Russian Thistle	-	0	0	0	20
25	Ryegrass, Italian	0	15	10	15	0
	Speedwell	100	90	100	100	-
	Wheat, Spring	0	5	5	0	0
	Wheat, Winter	0	0	5	0	0
	Windgrass	15	20	75	15	60
30	Table D	Compounds				
	16 g ai/ha	3	22	23	24	42
	Preemergence					
	Barley, Spring	0	0	0	0	0
	Barley, Winter	0	0	0	10	0
35	Blackgrass	0	5	20	15	0
	Bluegrass	0	0	20	15	10
	Bromegrass, Downy	0	15	5	20	0
	Buckwheat, Wild	0	10	40	10	0

94

	Canarygrass	0	10	60	25	15
	Chamomile	10	30	20	55	-
	Chickweed	100	10	50	10	70
	Deadnettle	70	20	15	30	70
5	Field Poppy	90	65	65	75	-
	Field Violet	100	15	25	20	25
	Foxtail, Green	10	15	0	10	10
	Galium	20	60	70	0	65
	Geranium, Cutleaf	-	25	25	75	-
10	Kochia	100	0	15	0	35
	Lambsquarters	10	20	25	100	85
	Mustard, Wild	15	20	98	15	70
	Oat, Wild	0	5	5	0	5
	Oilseed Rape	10	30	80	0	50
15	Pigweed	100	75	98	60	100
	Radish, Wild	30	45	25	0	25
	Russian Thistle	-	0	0	0	0
	Ryegrass, Italian	0	10	5	10	0
	Speedwell	70	85	100	35	-
20	Wheat, Spring	0	0	0	0	0
	Wheat, Winter	0	0	0	0	0
	Windgrass	10	15	20	5	15

TEST E

Seeds of plant species selected from corn (*Zea mays*), soybean (*Glycine max*),
25 velvetleaf (*Abutilon theophrasti*), cocklebur (common cocklebur, *Xanthium strumarium*),
lambsquarters (*Chenopodium album*), wild poinsettia (*Euphorbia heterophylla*), palmer
pigweed (*Amaranthus palmeri*), waterhemp (common waterhemp, *Amaranthus rudis*),
surinam grass (*Brachiaria decumbens*), large (Lg) crabgrass (*Digitaria sanguinalis*),
Brazilian crabgrass (*Digitaria horizontalis*), fall panicum (*Panicum dichotomiflorum*), giant
30 foxtail (*Setaria faberii*), green foxtail (*Setaria viridis*), goosegrass (*Eleusine indica*),
johnsongrass (*Sorghum halepense*), ragweed (common ragweed, *Ambrosia elatior*),
barnyardgrass (*Echinochloa crus-galli*), sandbur (southern sandbur, *Cenchrus echinatus*),
arrowleaf sida (*Sida rhombifolia*), Italian ryegrass (*Lolium multiflorum*), dayflower (Virginia
(VA) dayflower, *Commelina virginica*), field bindweed (*Convolvulus arvensis*),
35 morningglory (*Ipomoea coccinea*), nightshade (eastern black nightshade, *Solanum
ptycanthum*), kochia (*Kochia scoparia*), yellow nutsedge (*Cyperus esculentus*), horseweed
(*Conyza canadensis*), and hairy beggarticks (*Bidens pilosa*), were planted into a silt loam

soil and treated preemergence with test chemicals formulated in a non-phytotoxic solvent mixture which included a surfactant.

At the same time, plants from these crop and weed species and also waterhemp_RES1, (ALS & Triazine resistant common waterhemp, *Amaranthus rudis*), and waterhemp_RES2, (ALS & HPPD resistant common waterhemp, *Amaranthus rudis*) were planted in pots containing Redi-Earth[®] planting medium (Scotts Company, 14111 Scottslawn Road, Marysville, Ohio 43041) comprising sphagnum peat moss, vermiculite, wetting agent and starter nutrients were treated with postemergence applications of test chemicals formulated in the same manner. Plants ranged in height from 2 to 18 cm for postemergence treatments (1- to 4-leaf stage). Treated plants and controls were maintained in a greenhouse for 14 to 21 days, after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in Table E, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

Table E	Compounds				Table E	Compounds		
125 g ai/ha	36	38	42	43	62 g ai/ha	36	38	42
Postemergence					Postemergence			
Arrowleaf Sida	85	90	98	95	Arrowleaf Sida	80	90	95
Barnyardgrass	50	20	20	15	Barnyardgrass	30	10	10
Beggarticks	40	50	70	60	Beggarticks	30	40	60
Corn	25	20	15	15	Corn	20	15	10
Crabgrass, Brazil	30	20	30	40	Crabgrass, Brazil	30	20	20
Dayflower, VA	70	60	75	70	Dayflower, VA	60	60	70
Field Bindweed	85	85	70	70	Field Bindweed	65	80	60
Horseweed	10	40	10	10	Horseweed	5	20	10
Kochia	95	98	95	95	Kochia	100	100	95
Panicum, Fall	25	–	30	50	Panicum, Fall	15	10	20
Pigweed, Palmer	85	90	100	90	Pigweed, Palmer	85	60	95
Poinsettia, Wild	90	–	70	75	Poinsettia, Wild	70	–	70
Ragweed	50	50	60	60	Ragweed	35	40	50
Ryegrass, Italian	30	15	40	20	Ryegrass, Italian	10	10	20
Sandbur	25	20	10	10	Sandbur	20	10	0
Soybean	50	98	95	95	Soybean	70	95	95
Waterhemp	95	95	95	95	Waterhemp	95	95	90
Waterhemp_RES1	100	95	90	80	Waterhemp_RES1	95	90	75
Waterhemp_RES2	90	95	95	80	Waterhemp_RES2	80	90	85

Table E	Compounds				Table E	Compounds		
31 g ai/ha	36	38	42	43	16 g ai/ha	36	38	42

Postemergence					Postemergence				
Arrowleaf Sida	70	80	85	80	Arrowleaf Sida	70	60	75	
Barnyardgrass	25	10	0	0	Barnyardgrass	20	0	0	
Beggarticks	35	40	60	50	Beggarticks	30	35	50	
Corn	20	0	10	5	Corn	10	0	5	
Crabgrass, Brazil	20	15	10	25	Crabgrass, Brazil	20	10	10	
Dayflower, VA	50	50	50	50	Dayflower, VA	20	40	35	
Field Bindweed	50	65	50	55	Field Bindweed	40	60	40	
Horseweed	0	10	0	0	Horseweed	0	5	0	
Kochia	95	80	80	75	Kochia	90	80	70	
Panicum, Fall	25	10	15	40	Panicum, Fall	20	10	10	
Pigweed, Palmer	85	75	85	80	Pigweed, Palmer	75	60	80	
Poinsettia, Wild	60	-	60	60	Poinsettia, Wild	50	-	50	
Ragweed	25	20	25	50	Ragweed	15	20	10	
Ryegrass, Italian	15	5	10	0	Ryegrass, Italian	10	0	0	
Sandbur	20	0	0	0	Sandbur	10	0	0	
Soybean	50	80	60	70	Soybean	60	70	50	
Waterhemp	90	90	85	65	Waterhemp	95	85	75	
Waterhemp_RES1	95	90	75	65	Waterhemp_RES1	98	80	60	
Waterhemp_RES2	90	85	65	65	Waterhemp_RES2	60	75	50	

Table E

Compounds

125 g ai/ha 19 21 34 38 42 43

Preemergence

5	Arrowleaf Sida	100	100	100	100	100	100
	Barnyardgrass	70	100	60	90	100	100
	Beggarticks	0	0	75	35	0	0
	Cocklebur	0	50	-	-	-	-
	Corn	50	50	0	50	65	50
10	Crabgrass, Brazil	100	100	98	100	100	100
	Crabgrass, Large	100	100	98	100	100	100
	Dayflower, VA	10	95	90	60	95	95
	Field Bindweed	10	40	0	65	85	50
	Foxtail, Giant	100	100	80	100	100	100
15	Foxtail, Green	100	100	70	100	100	100
	Goosegrass	98	98	80	100	98	100
	Horseweed	-	-	0	98	-	-
	Johnsongrass	100	100	-	95	100	100
	Kochia	100	100	100	100	100	100

	Lambsquarters	100	100	100	100	100	100
	Morningglory	65	80	65	35	95	95
	Nightshade	98	100	95	100	98	100
	Nutsedge, Yellow	20	15	0	65	40	25
5	Panicum, Fall	100	100	100	100	100	100
	Pigweed, Palmer	100	100	100	100	100	90
	Poinsettia, Wild	70	60	50	-	100	100
	Ragweed	50	98	35	70	100	85
	Ryegrass, Italian	70	70	70	80	98	98
10	Sandbur	20	50	70	80	65	100
	Soybean	50	90	60	80	100	65
	Surinam Grass	10	50	10	40	85	80
	Velvetleaf	100	100	100	100	100	100
	Waterhemp	100	100	100	100	100	100
15	Table E	Compounds					
	62 g ai/ha	19	21	34	38	42	
	Preemergence						
	Arrowleaf Sida	100	100	98	80	100	
	Barnyardgrass	20	65	20	70	70	
20	Beggarticks	0	0	70	0	0	
	Cocklebur	0	40	-	-	-	
	Corn	20	10	0	35	35	
	Crabgrass, Brazil	100	100	95	100	100	
	Crabgrass, Large	100	95	98	100	100	
25	Dayflower, VA	5	25	20	20	90	
	Field Bindweed	0	15	0	95	40	
	Foxtail, Giant	100	100	85	100	100	
	Foxtail, Green	98	70	65	100	95	
	Goosegrass	90	75	5	90	90	
30	Horseweed	-	-	0	100	-	
	Johnsongrass	50	65	-	100	95	
	Kochia	98	75	100	100	100	
	Lambsquarters	100	100	90	100	100	
	Morningglory	5	90	50	50	70	
35	Nightshade	65	98	80	98	98	
	Nutsedge, Yellow	0	0	0	20	35	
	Panicum, Fall	98	100	98	100	100	
	Pigweed, Palmer	100	100	70	98	80	

98

	Poinsettia, Wild	15	50	20	-	80
	Ragweed	10	50	25	60	65
	Ryegrass, Italian	65	35	35	65	80
	Sandbur	0	15	10	35	35
5	Soybean	0	65	25	95	35
	Surinam Grass	10	0	-	15	40
	Velvetleaf	50	100	100	85	100
	Waterhemp	100	100	100	98	100
	Table E	Compounds				
10	31 g ai/ha	19	21	34	38	42 43
	Preemergence					
	Arrowleaf Sida	40	100	80	50	98 100
	Barnyardgrass	0	20	0	10	50 20
	Beggarticks	0	0	60	0	0 0
15	Cocklebur	-	35	-	-	- -
	Corn	0	0	0	0	0 0
	Crabgrass, Brazil	40	65	35	98	95 95
	Crabgrass, Large	80	75	25	75	98 98
	Dayflower, VA	0	0	30	0	20 25
20	Field Bindweed	0	0	0	70	20 20
	Foxtail, Giant	95	90	50	70	80 85
	Foxtail, Green	60	50	10	30	80 70
	Goosegrass	75	50	0	50	50 50
	Horseweed	-	-	0	0	0 -
25	Johnsongrass	0	30	-	0	80 90
	Kochia	75	50	95	98	90 98
	Lambsquarters	100	40	70	100	100 95
	Morningglory	0	50	20	0	35 5
	Nightshade	0	90	50	95	70 98
30	Nutsedge, Yellow	0	0	0	0	0 0
	Panicum, Fall	85	75	80	98	98 95
	Pigweed, Palmer	70	60	50	100	70 75
	Poinsettia, Wild	0	10	10	-	20 65
	Ragweed	0	50	0	40	20 60
35	Ryegrass, Italian	0	0	0	25	40 40
	Sandbur	0	0	0	0	5 15
	Soybean	0	25	10	15	35 30
	Surinam Grass	0	10	0	0	0 25

	Velvetleaf	0	70	100	65	80	70
	Waterhemp	65	80	100	98	100	90
	Table E	Compounds					
	16 g ai/ha	19	21	34	38	42	
5	Preemergence						
	Arrowleaf Sida	65	90	60	60	100	
	Barnyardgrass	0	15	0	0	5	
	Beggarticks	0	0	50	0	0	
	Cocklebur	0	-	-	-	-	
10	Corn	0	0	0	0	0	
	Crabgrass, Brazil	0	0	0	80	35	
	Crabgrass, Large	20	50	0	50	80	
	Dayflower, VA	0	0	0	0	0	
	Field Bindweed	0	0	0	35	25	
15	Foxtail, Giant	0	5	0	5	50	
	Foxtail, Green	0	15	0	5	35	
	Goosegrass	5	10	0	30	0	
	Horseweed	-	-	0	0	-	
	Johnsongrass	0	0	-	0	80	
20	Kochia	35	20	0	80	80	
	Lambsquarters	60	0	0	98	100	
	Morningglory	0	10	-	0	0	
	Nightshade	0	80	50	5	80	
	Nutsedge, Yellow	0	0	0	0	0	
25	Panicum, Fall	100	35	50	0	0	
	Pigweed, Palmer	0	0	0	60	60	
	Poinsettia, Wild	0	0	0	-	20	
	Ragweed	0	0	0	40	0	
	Ryegrass, Italian	0	0	0	0	20	
30	Sandbur	0	0	0	0	0	
	Soybean	0	0	0	15	0	
	Surinam Grass	0	0	0	0	0	
	Velvetleaf	0	80	30	20	75	
	Waterhemp	65	65	100	50	98	

35 TEST F

Seeds of plant species selected from bermudagrass (*Cynodon dactylon*), Surinam grass (*Brachiaria decumbens*), large (Lg) crabgrass (*Digitaria sanguinalis*), crabgrass, naked

(naked crabgrass, *Digitaria nuda*), foxtail, green (green foxtail, *Setaria viridis*), johnsongrass (*Sorghum halepense*), kochia (*Kochia scoparia*), morningglory (pitted morningglory, *Ipomoea lacunosa*), nutsedge, purple (purple nutsedge, *Cyperus rotundus*), ragweed (common ragweed, *Ambrosia elatior*), mustard, black (black mustard, *Brassica nigra*), guineagrass (*Panicum maximum*), dallisgrass (*Paspalum dilatatum*), barnyardgrass (*Echinochloa crus-galli*), sandbur (southern sandbur, *Cenchrus echinatus*), sowthistle (common sowthistle, *Sonchus oleraceus*), Italian ryegrass (*Lolium multiflorum*), signalgrass (broadleaf signalgrass, *Brachiaria platyphylla*), dayflower (Virginia (VA) dayflower, *Commelina virginica*), bluegrass (annual bluegrass, *Poa annua*), quackgrass (*Elytrigia repens*), mallow (common mallow, *Malva sylvestris*), buckwheat, wild (wild buckwheat, *Polygonum convolvulus*), leafy spurge (*Euphorbia esula*), chickweed (common chickweed, *Stellaria media*), wild poinsettia (*Euphorbia heterophylla*), and pigweed (*Amaranthus retroflexus*) were planted into a blend of loam soil and sand and treated preemergence with test a chemical formulated in a non-phytotoxic solvent mixture which included a surfactant.

Treated plants and controls were maintained in a greenhouse for 21 days, after which time all species were compared to controls and visually evaluated. Plant response ratings, summarized in Table F, are based on a scale of 0 to 100 where 0 is no effect and 100 is complete control. A dash (–) response means no test result.

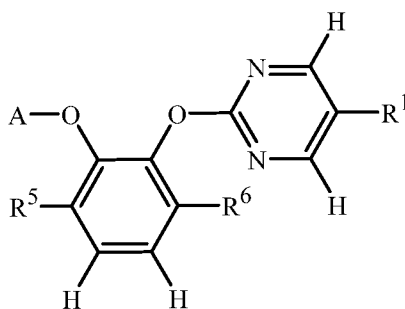
Table F	Compound	Table F	Compound	Table F	Compound
250 g ai/ha	3	125 g ai/ha	3	62 g ai/ha	3
Preemergence		Preemergence		Preemergence	
Barnyardgrass	100	Barnyardgrass	100	Barnyardgrass	98
Bermudagrass	100	Bermudagrass	100	Bermudagrass	100
Bluegrass	100	Bluegrass	100	Bluegrass	80
Buckwheat, Wild	100	Buckwheat, Wild	100	Buckwheat, Wild	100
Chickweed	100	Chickweed	100	Chickweed	100
Crabgrass, Large	100	Crabgrass, Large	100	Crabgrass, Large	100
Crabgrass, Naked	100	Crabgrass, Naked	100	Crabgrass, Naked	100
Dallisgrass	100	Dallisgrass	100	Dallisgrass	100
Dayflower, VA	100	Dayflower, VA	100	Dayflower, VA	98
Foxtail, Green	100	Foxtail, Green	100	Foxtail, Green	100
Guineagrass	100	Guineagrass	100	Guineagrass	100
Johnsongrass	100	Johnsongrass	100	Johnsongrass	75
Kochia	100	Kochia	100	Kochia	100
Leafy Spurge	98	Leafy Spurge	100	Leafy Spurge	98
Mallow	100	Mallow	100	Mallow	100
Morningglory, Pitt	100	Morningglory, Pitt	100	Morningglory, Pitt	100
Mustard, Black	100	Mustard, Black	100	Mustard, Black	100

Table F	Compound	Table F	Compound	Table F	Compound
250 g ai/ha	3	125 g ai/ha	3	62 g ai/ha	3
Preemergence		Preemergence		Preemergence	
Nutsedge, Purple	85	Nutsedge, Purple	60	Nutsedge, Purple	40
Pigweed	100	Pigweed	100	Pigweed	100
Poinsettia, Wild	100	Poinsettia, Wild	95	Poinsettia, Wild	95
Quackgrass	98	Quackgrass	90	Quackgrass	65
Ragweed	100	Ragweed	100	Ragweed	85
Ryegrass, Italian	100	Ryegrass, Italian	100	Ryegrass, Italian	95
Sandbur	100	Sandbur	100	Sandbur	35
Signalgrass	100	Signalgrass	100	Signalgrass	100
Sowthistle	100	Sowthistle	100	Sowthistle	100
Surinam Grass	100	Surinam Grass	100	Surinam Grass	75

CLAIMS

What is claimed is:

1. A compound selected from Formula 1, *N*-oxides and salts thereof,



1

5 wherein

A is a phenyl ring optionally substituted with up to 4 R²; or a 5- or 6-membered heteroaromatic ring, the ring bonded to the remainder of Formula 1 through a carbon atom, and optionally substituted with up to 4 R²;

10 R¹ is halogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₄ alkoxy or S(O)_mR³;

each R² is independently halogen, cyano, nitro, SF₅, CHO, C(=O)NH₂, C(=S)NH₂, SO₂NH₂, C₁-C₄ alkyl, C₂-C₄ alkenyl, C₂-C₄ alkynyl, C₁-C₄ haloalkyl, C₂-C₄ haloalkenyl, C₂-C₄ haloalkynyl, C₃-C₆ cycloalkyl, C₃-C₆ halocycloalkyl, C₄-C₈ alkylcycloalkyl, C₄-C₈ cycloalkylalkyl, C₂-C₆ alkylcarbonyl, C₂-C₆ haloalkylcarbonyl, C₂-C₆ alkoxy carbonyl, C₃-C₇ cycloalkylcarbonyl, C₂-C₈ alkylaminocarbonyl, C₃-C₁₀ dialkylaminocarbonyl, C₁-C₄ alkoxy, C₃-C₄ alkenyloxy, C₃-C₄ alkynyloxy, C₁-C₄ haloalkoxy, C₃-C₄ haloalkenyloxy, C₃-C₄ haloalkynyloxy, C₃-C₆ cycloalkoxy, C₃-C₆ halocycloalkoxy, C₄-C₈ cycloalkylalkoxy, C₂-C₆ alkoxyalkyl, C₂-C₆ haloalkoxyalkyl, C₂-C₆ alkoxyhaloalkyl, C₂-C₆ alkoxyalkoxy, C₂-C₄ alkylcarbonyloxy, C₂-C₆ cyanoalkyl, C₂-C₆ cyanoalkoxy, C₁-C₄ hydroxyalkyl, C₂-C₄ alkylthioalkyl, C₁-C₆ alkylamino, C₂-C₆ dialkylamino, S(O)_nR⁴, CH(=NOH), phenyl or pyridinyl; each R³ and R⁴ is independently C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkylamino or C₂-C₆ dialkylamino;

25 R⁵ is halogen, cyano or C₁-C₂ haloalkyl;

R⁶ is H or F;

m is 0, 1 or 2; and

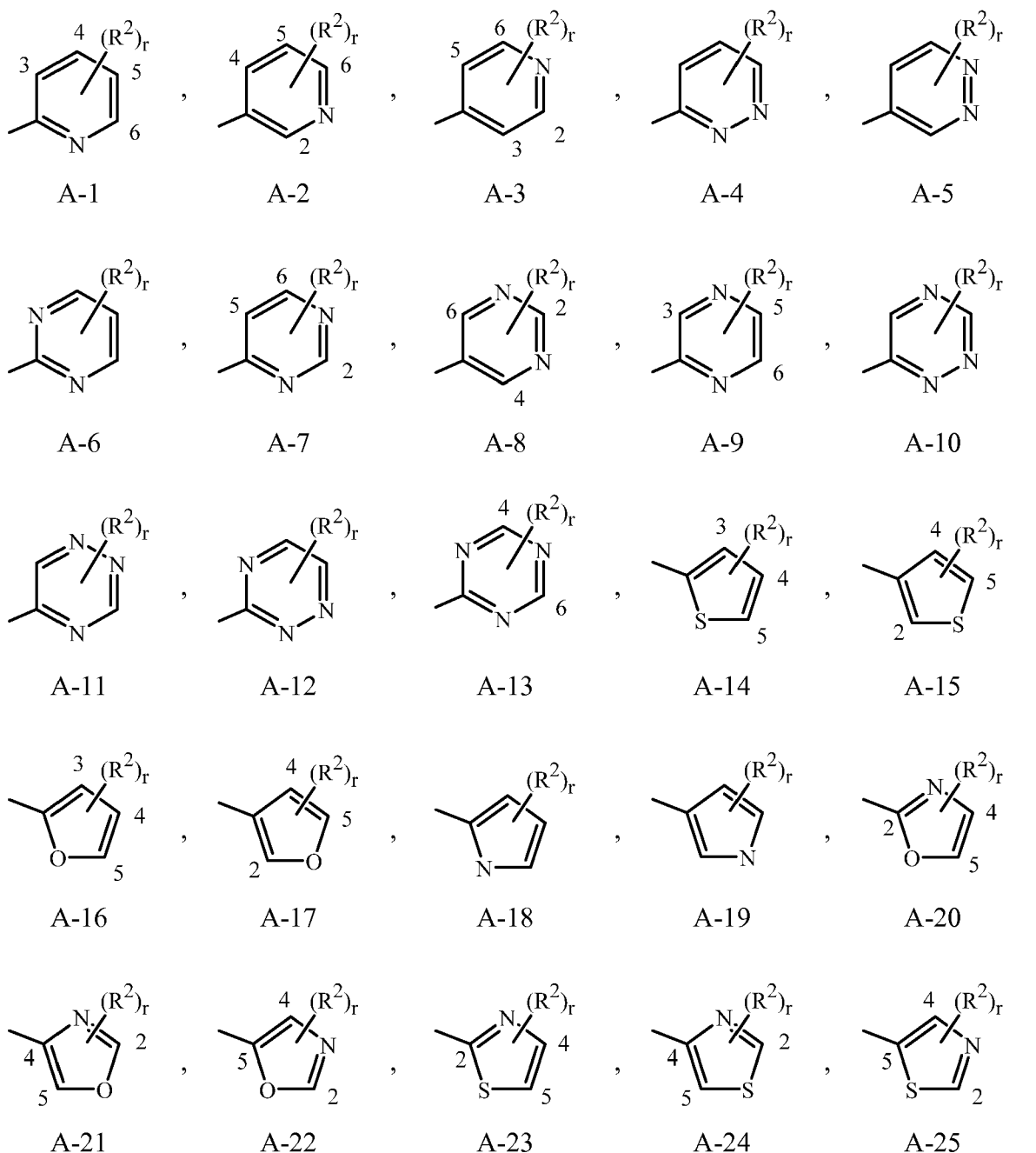
each n is independently 0, 1 or 2;

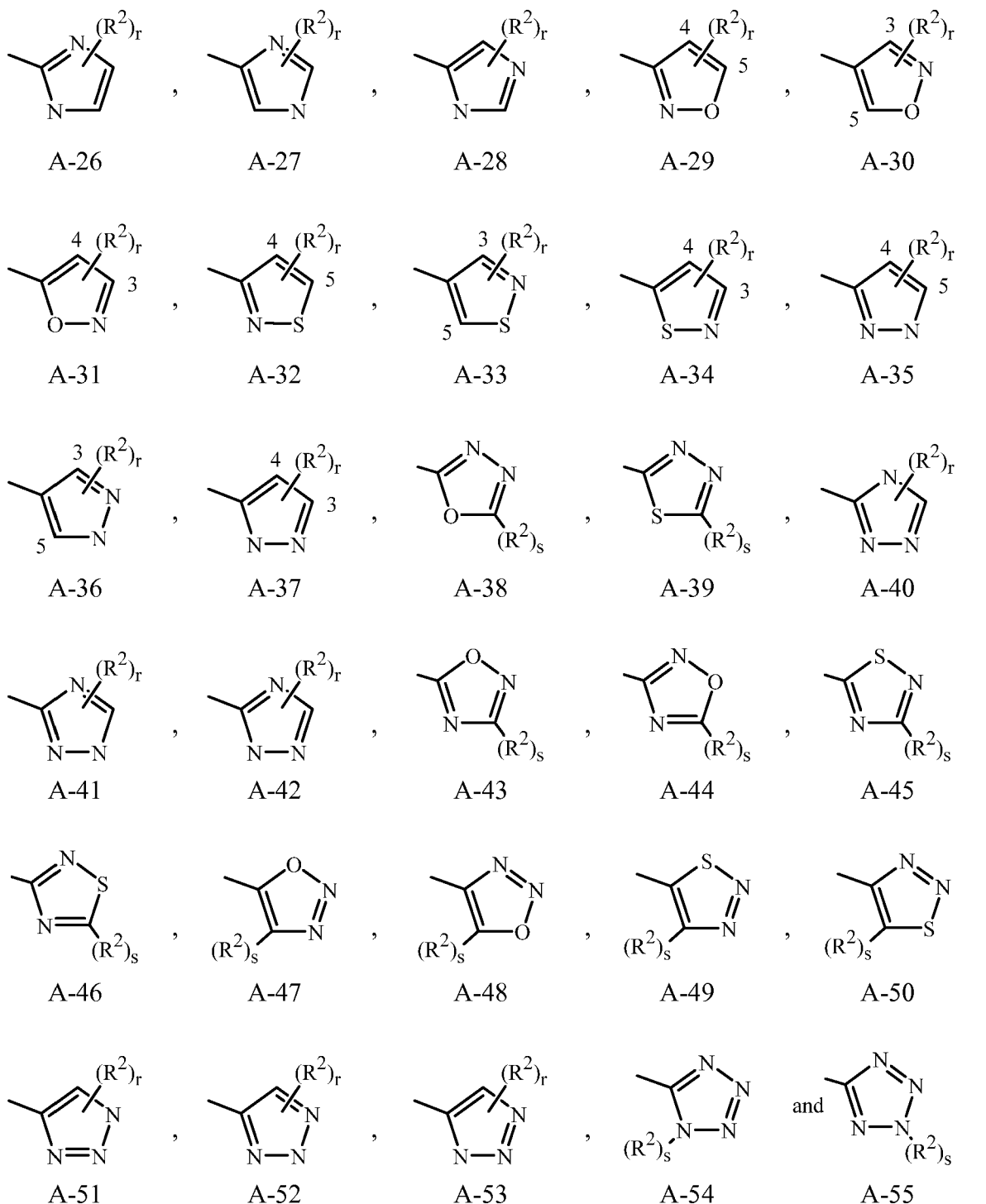
5

provided the compound of Formula 1 is other than 5-bromo-2-[3-bromo-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine, 5-bromo-2-[6-bromo-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine, 5-chloro-2-[3-fluoro-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine, 5-chloro-2-[6-fluoro-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine, 5-chloro-2-[3-methyl-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine or 5-chloro-2-[6-methyl-[2-(5-chloropyridin-2-yloxy)phenoxy]pyrimidine.

2. A compound of Claim 1 wherein

A is selected from





wherein r is 0, 1, 2 or 3 and s is 0 or 1; and

each R^2 is independently halogen, cyano, SF_5 , C_1 - C_4 alkyl, C_2 - C_4 alkenyl, C_2 - C_4 alkynyl, C_1 - C_4 haloalkyl, C_2 - C_4 haloalkenyl or C_2 - C_4 haloalkynyl.

3. A compound of Claim 2 wherein

5 A is selected from A-1, A-2, A-4, A-6, A-9, A-10, A-11, A-12 and A-23;

R^1 is halogen, C_1 - C_4 alkyl or C_1 - C_4 haloalkyl; and

each R² is independently halogen, C₁–C₄ alkyl or C₁–C₄ haloalkyl.

4. A compound of Claim 3 wherein

A is selected from A-1, A-2 and A-6;

each R² is independently halogen, CH₃ or CF₃;

5 R⁵ is halogen, cyano, CHF₂ or CF₃; and

R⁶ is H.

5. A compound of Claim 4 wherein

A is A-6;

R¹ is halogen; and

10 R⁵ is F, Cl, Br or cyano.

6. A compound of Claim 5 wherein

A is A-6a.

7. A compound of Claim 1 selected from the group consisting of

2,3-bis[(5-bromo-2-pyrimidinyl)oxy]benzonitrile,

2,3-bis[(5-chloro-2-pyrimidinyl)oxy]benzonitrile,

2,3-bis[(5-fluoro-2-pyrimidinyl)oxy]benzonitrile,

2-[(5-bromo-2-pyrimidinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile,

3-[(5-bromo-2-pyrimidinyl)oxy]-2-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile,

2-[(5-chloro-2-pyridinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile,

2,2'-[[3-(difluoromethyl)-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine],

2-[3-bromo-2-[[5-(difluoromethyl)-2-thiazolyl]oxy]phenoxy]-5-chloropyrimidine,

5-chloro-2-[2-fluoro-6-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]phenoxy]-

pyrimidine,

5-chloro-2-[5-fluoro-6-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]phenoxy]-

pyrimidine,

5-bromo-2-[2-chloro-6-[(5-chloro-2-pyrimidinyl)oxy]phenoxy]pyrimidine,

5-chloro-2-[5-chloro-6-[(5-fluoro-2-pyrimidinyl)oxy]phenoxy]pyrimidine,

2,2'-[(3,6-difluoro-1,2-phenylene)bis(oxy)]bis[5-fluoropyrimidine],

5-bromo-2-[2-fluoro-6-[(5-chloro-2-pyrimidinyl)oxy]phenoxy]pyrimidine,

3-[(5-chloro-2-pyrimidinyl)oxy]-2-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]-

benzonitrile,

2-[(5-chloro-2-pyrimidinyl)oxy]-3-[[5-(trifluoromethyl)-2-pyrimidinyl]oxy]-

benzonitrile,

2-[(5-chloro-2-pyrazinyl)oxy]-3-[(5-chloro-2-pyrimidinyl)oxy]benzonitrile,

2,2'-[(3,6-difluoro-1,2-phenylene)bis(oxy)]bis[5-chloropyrimidine],

2,2'-[[3-fluoro-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine] and

2,2'-[[3-bromo-1,2-phenylene]bis(oxy)]bis[5-chloropyrimidine].

8. A herbicidal composition comprising a compound of Claim 1 and at least one component selected from the group consisting of surfactants, solid diluents and liquid diluents.

9. A herbicidal composition comprising a compound of Claim 1, at least one
5 additional active ingredient selected from the group consisting of other herbicides and herbicide safeners, and at least one component selected from the group consisting of surfactants, solid diluents and liquid diluents.

10. A herbicidal mixture comprising (a) a compound of Claim 1, and (b) at least one
10 additional active ingredient selected from (b1) through (b16) and salts of compounds of (b1) through (b16).

11. A herbicidal mixture comprising (a) a compound of Claim 1, and (b) at least one
additional active ingredient selected from (b1) photosystem II inhibitors, (b2) acetohydroxy
acid synthase (AHAS) inhibitors, (b4) auxin mimics, (b5) 5-enol-pyruvylshikimate-3-
phosphate (EPSP) synthase inhibitors, (b7) protoporphyrinogen oxidase (PPO) inhibitors,
15 (b9) very long chain fatty acid (VLCFA) elongase inhibitors and (b12)
4-hydroxyphenyl-pyruvate dioxygenase (HPPD) inhibitors.

12. A herbicidal mixture comprising (a) a compound of Claim 1, and (b) at least one
additional active ingredient selected from the group consisting of 2,4-D, acetochlor, alachlor,
atrazine, bromoxynil, bentazon, bicyclopyrone, carfentrazone-ethyl, cloransulam-methyl,
20 dicamba, dimethenamid-p, florasulam, flufenacet, flumioxazin, flupyrsulfuron-methyl,
fluroxypyr-meptyl, glyphosate, halauxifen-methyl, isoxaflutole, MCPA, mesotrione,
metolachlor, metsulfuron-methyl, nicosulfuron, pyrasulfotole, pyroxsulfone, pyroxsulam,
rimsulfuron, saflufenacil, tembotrione, thifensulfuron-methyl, topramazine and tribenuron.

13. A method for controlling the growth of undesired vegetation comprising
25 contacting the vegetation or its environment with a herbicidally effective amount of a
compound of Claim 1.

14. A method for controlling the growth of undesired vegetation in a genetically
modified plants that exhibit traits of glyphosate tolerance, glufosinate tolerance, ALS
herbicide tolerance, dicamba tolerance, imidazolinone herbicide tolerance, 2,4-D tolerance,
30 HPPD tolerance and mesotrione tolerance, comprising contacting the vegetation or its
environment with a herbicidally effective amount of a compound of Claim 1.