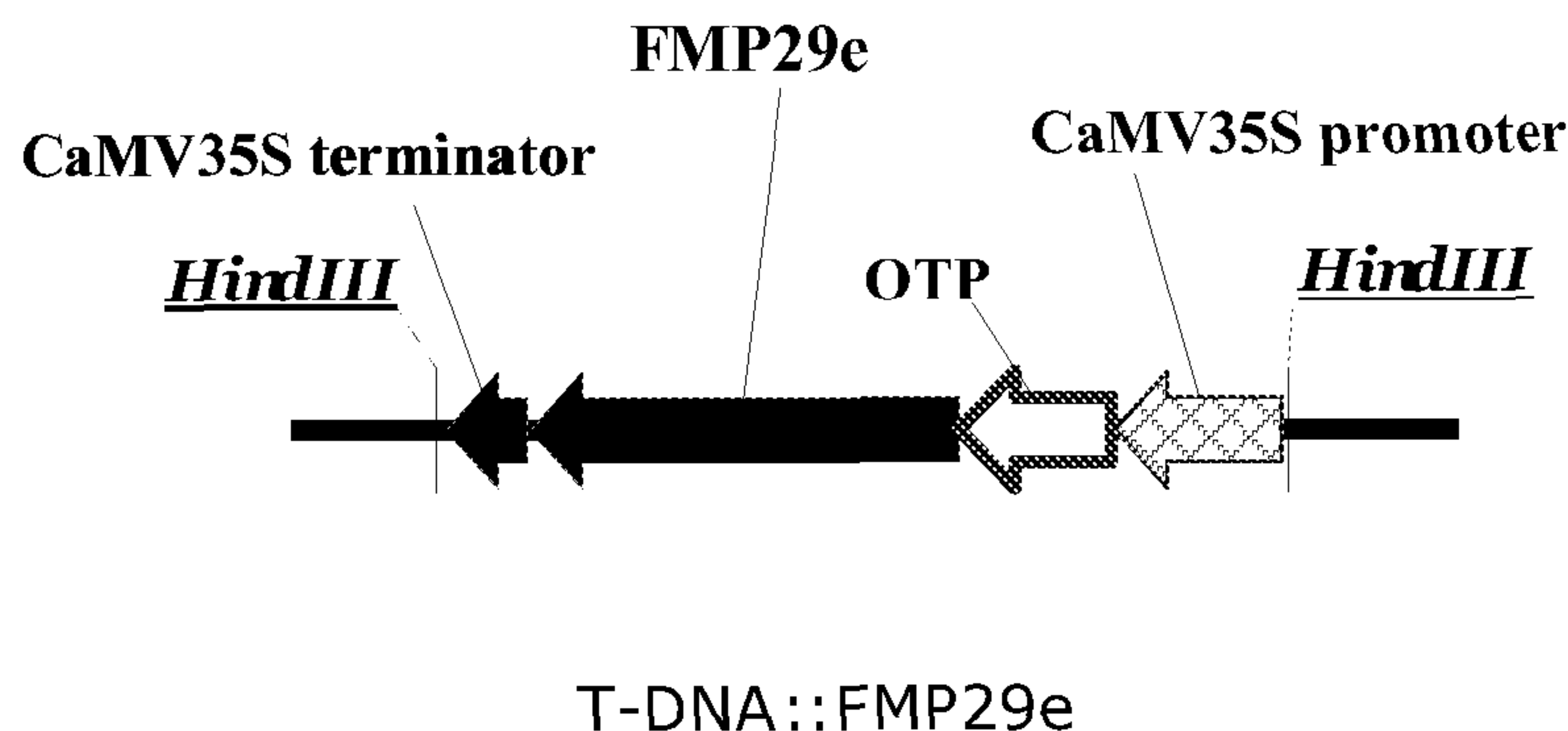




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map of the T-DNA inserted into the tobacco plants.

(57) **Abrégé/Abstract:**

The present invention relates to nucleic acid sequences encoding a hydroxyphenylpyruvate dioxygenase (EC 1.13.1 1.27, abbreviated herein as HPPD) obtained from Euryarchaeota belonging to the family Picrophilaceae, as well as the proteins encoded thereby, and to a chimeric gene which comprises such nucleic acid sequence, and to the use of such nucleic acid sequences, proteins or chimeric genes for obtaining plants which are tolerant to HPPD inhibitor herbicides.

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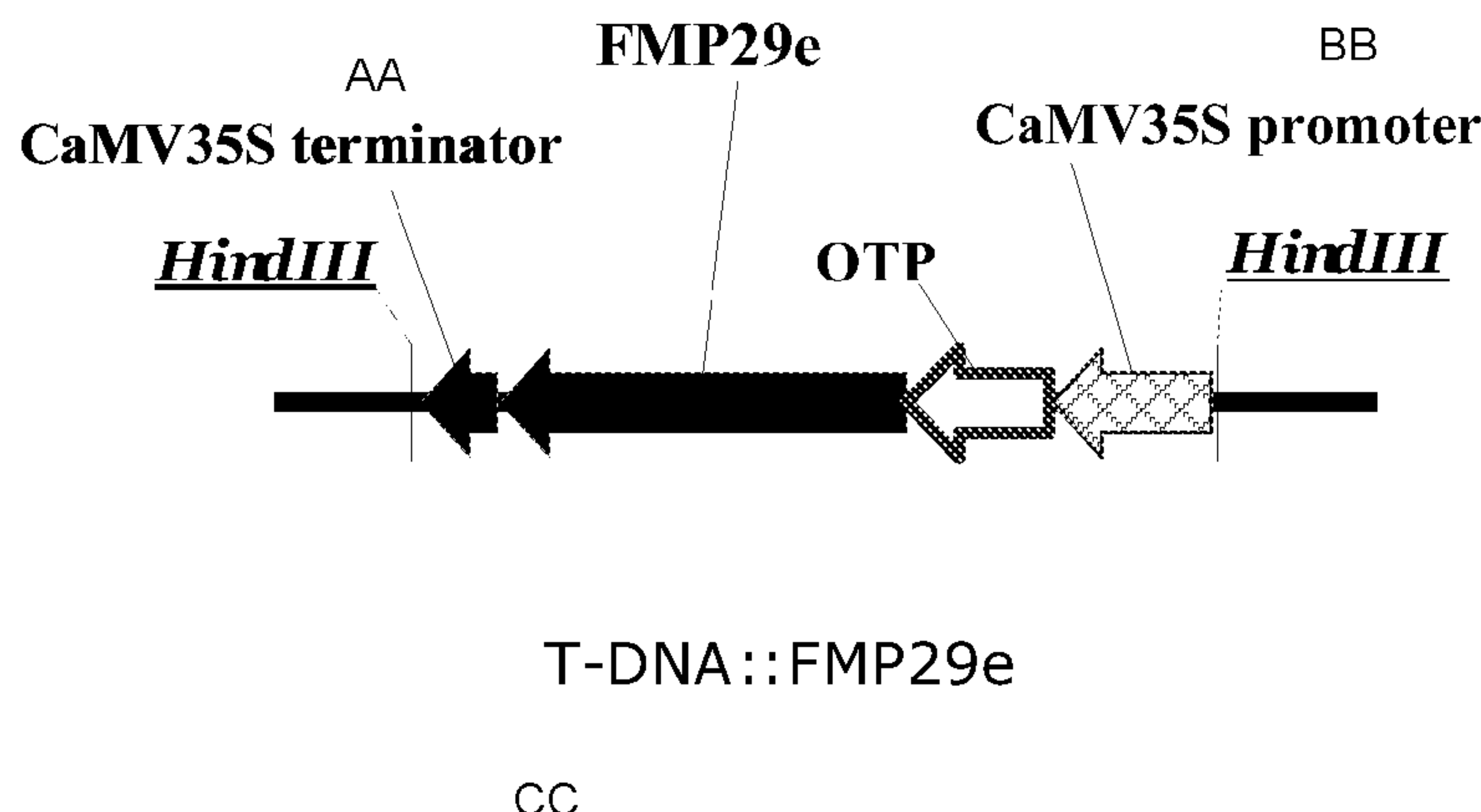


Figure 2: map of the T-DNA inserted into the tobacco plants.

(57) Abstract: The present invention relates to nucleic acid sequences encoding a hydroxyphenylpyruvate dioxygenase (EC 1.13.1 1.27, abbreviated herein as HPPD) obtained from Euryarchaeota belonging to the family Picrophilaceae, as well as the proteins encoded thereby, and to a chimeric gene which comprises such nucleic acid sequence, and to the use of such nucleic acid sequences, proteins or chimeric genes for obtaining plants which are tolerant to HPPD inhibitor herbicides.

Plants tolerant to HPPD inhibitor herbicides.

5

Introduction

The present invention relates to nucleic acid sequences encoding a hydroxyphenylpyruvate dioxygenase (EC 1.13.11.27, abbreviated herein as HPPD) obtained from Euryarchaeota belonging to the family Picrophilaceae, as well as the proteins encoded thereby, and to a chimeric gene which comprises such nucleic acid sequence, and to the use of such nucleic acid sequences, proteins or chimeric genes for obtaining plants which are tolerant to HPPD inhibitor herbicides.

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Background

The HPPDs are enzymes which catalyse the reaction in which para-hydroxyphenylpyruvate (abbreviated herein as HPP), a tyrosine degradation product, is transformed into homogentisate (abbreviated herein as HG), the precursor in plants of tocopherol and plastoquinone (Crouch N.P. et al. (1997) Tetrahedron, 53, 20, 6993-7010, Fritze et al., (2004), Plant Physiology 134:1388-1400). Tocopherol acts as a membrane-associated antioxidant. Plastoquinone, firstly acts as an electron carrier between PSII and the cytochrome b6/f complex and secondly, is a redox cofactor for phytoene desaturase, which is involved in the biosynthesis of carotenoids.

Up to now, more than 700 nucleic acid sequences from various organisms present in NCBI database were annotated as coding for a putative protein having an HPPD domain including the sequence disclosed under the Q6KZ98 accession number given in the UniProtKB/TrEMBL database as well as the YP_024147 accession number given in the NCBI protein database. But for most of those, including the sequence corresponding to the accession number Q6KZ98/YP_024147, it has not

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been proven that the protein would have an HPPD enzymatic activity either in an *in vitro* assay or an *in planta* approach, nor that such HPPD protein can confer herbicide tolerance to HPPD inhibitor herbicides when expressed in a plant. Several HPPD proteins and their primary sequences have been described in the state of the art, in particular the HPPD proteins of bacteria such as *Pseudomonas* (Rüetschi et al., Eur. J. Biochem., 205, 459-466, 1992, WO 96/38567), of plants such as *Arabidopsis* (WO 96/38567, Genebank AF047834), carrot (WO 96/38567, Genebank 87257), *Avena sativa* (WO 02/046387), wheat (WO 02/046387), *Brachiaria platyphylla* (WO 02/046387), *Cenchrus echinatus* (WO 02/046387), *Lolium rigidum* (WO 02/046387), *Festuca arundinacea* (WO 02/046387), *Setaria faberi* (WO 02/046387), *Eleusine indica* (WO 02/046387), *Sorghum* (WO 02/046387), *Coccicoides* (Genebank COITRP), of *Coptis japonica* (WO 06/132270), *Chlamydomonas reinhardtii* (ES 2275365), or of mammals such as mouse or pig.

15

Most plants synthesize tyrosine via arrogenate (Abou-Zeid et al. (1995), Applied Env Microb 41: 1298-1302; Bonner et al., (1995), Plant Cells Physiol. 36, 1013-1022; Byng et al., (1981), Phytochemistry 6: 1289-1292; Connely and Conn (1986), Z. Naturforsch 41c: 69-78; Gaines et al., (1982), Plants 156: 233-240). In these plants, the HPP is derived only from the degradation of tyrosine. On the other hand, in organisms such as the yeast *Saccharomyces cerevisiae* or the bacterium *Escherichia coli*, HPP is a tyrosine precursor, and it is synthesized by the action of an enzyme, prephenate dehydrogenase (hereinafter referred to as PDH), which converts prephenate to HPP (Lingens et al., (1967) European J. Biochem 1: 363-374; Sampathkumar and Morrisson (1982), Bioch Biophys Acta 701: 204-211). In these organisms, the production of HPP is therefore directly connected to the aromatic amino acid biosynthetic pathway (shikimate pathway), and not to the tyrosine degradation pathway.

Inhibition of HPPD leads to uncoupling of photosynthesis, deficiency in accessory light-harvesting pigments and, most importantly, to destruction of chlorophyll by UV-radiation and reactive oxygen species (bleaching) due to the lack of photo protection normally provided by carotenoids (Norris et al. (1995), Plant Cell 7: 2139-
 5 2149). Bleaching of photosynthetically active tissues leads to growth inhibition and plant death.

Some molecules which inhibit HPPD, and which bind specifically to the enzyme in order to inhibit transformation of the HPP into homogentisate, have proven to be
 10 very effective selective herbicides.

At present, most commercially available HPPD inhibitor herbicides belong to one of these four chemical families:

1) the triketones, e.g. sulcotrione [i.e. 2-[2-chloro-4-(methylsulfonyl)benzoyl]-1,3-cyclohexanedione], mesotrione [i.e. 2-[4-(methylsulfonyl)-2-nitrobenzoyl]-1,3-cyclohexanedione]; tembotrione [i.e. 2-[2-chloro-4-(methylsulfonyl)-3-[(2,2,2-trifluoroethoxy)methyl] benzoyl]-1,3-cyclo-hexanedione]; tefuryltrione [i.e. 2-[2-chloro-4-(methylsulfonyl)-3-[[tetrahydro-2-furanyl)methoxy]methyl]benzoyl]-1,3-cyclohexanedione]; bicyclopyrone [i.e. 4-hydroxy-3-[[2-[(2-methoxyethoxy)methyl]-6-(trifluoromethyl)-3-pyridinyl]carbonyl]bicyclo[3.2.1]oct-3-en-2-one] ; Benzobicyclon
 15 [i.e. 3-(2-chloro-4-mesylbenzoyl)-2-phenylthiobicyclo[3.2.1]oct-2-en-4-one]

2) the diketonitriles, e.g. 2-cyano-3-cyclopropyl-1-(2-methylsulphonyl-4-trifluoromethylphenyl)-propane-1,3-dione and 2-cyano-1-[4-(methylsulphonyl)-2-trifluoromethylphenyl]-3-(1-methylcyclopropyl)propane-1,3-dione;
 25

3) the isoxazoles, e.g. isoxaflutole [i.e. (5-cyclopropyl-4-isoxazolyl)[2-(methylsulfonyl)-4-(trifluoromethyl)phenyl]methanone]. In plants, the isoxaflutole is rapidly metabolized in DKN, a diketonitrile compound which exhibits the HPPD
 30 inhibitor property; and

4) the pyrazolates, e.g. topramezone [i.e. [3-(4,5-dihydro-3-isoxazolyl)-2-methyl-4-(methylsulfonyl) phenyl](5-hydroxy-1-methyl-1H-pyrazol-4-yl)methanone], and pyrasulfotole [(5-hydroxy-1,3-dimethylpyrazol-4-yl)(2-mesyl-4-trifluaromethylphenyl)methanone]; pyrazofen [2-[4-(2,4-dichlorobenzoyl)-1,3-dimethylpyrazol-5-yloxy]acetophenone].

These HPPD-inhibiting herbicides can be used against grass and/or broad leaf weeds in crop plants that display metabolic tolerance, such as maize (*Zea mays*) in which they are rapidly degraded (Schulz et al., (1993). FEBS letters, 318, 162-166; Mitchell et al., (2001) Pest Management Science, Vol 57, 120-128; Garcia et al., (2000) Biochem., 39, 7501-7507; Pallett et al., (2001) Pest Management Science, Vol 57, 133-142).. In order to extend the scope of these HPPD-inhibiting herbicides, several efforts have been developed in order to confer to plants, particularly plants without or with an underperforming metabolic tolerance, a tolerance level acceptable under agronomic field conditions.

Besides the attempt of by-passing HPPD-mediated production of homogentisate (US 6,812,010), overexpressing the sensitive enzyme so as to produce quantities of the target enzyme in the plant which are sufficient in relation to the herbicide has been performed (WO96/38567). Overexpression of HPPD resulted in better pre-emergence tolerance to the diketonitrile derivative (DKN) of isoxaflutole (IFT), but tolerance was not sufficient for tolerance to post-emergence treatment (Matringe et al., (2005), Pest Management Science 61: 269-276).

A third strategy was to mutate the HPPD in order to obtain a target enzyme which, while retaining its properties of catalysing the transformation of HPP into homogentisate, is less sensitive to HPPD inhibitors than is the native HPPD before mutation.

This strategy has been successfully applied for the production of plants tolerant to 2-cyano-3-cyclopropyl-1-(2-methylsulphonyl-4-trifluoromethylphenyl)-propane-1,3-dione and to 2-cyano-1-[4-(methylsulphonyl)-2-trifluoromethylphenyl]-3-(1-

methylocyclopropyl)propane-1,3-dione (EP496630), two HPPD-inhibiting herbicides belonging to the diketonitriles family (WO 99/24585). Pro215Leu, Gly336Glu, Gly336Ile, and more particularly Gly336Trp (positions of the mutated amino acid are indicated with reference to the *Pseudomonas* HPPD) were identified as mutations
5 which are responsible for an increased tolerance to pre-emergence treatment with these diketonitrile herbicides without causing an alteration of the activity of the enzyme.

More recently, introduction of a *Pseudomonas* HPPD gene into the plastid genome
10 of tobacco and soybean has shown to be more effective than nuclear transformation, conferring even tolerance to post-emergence application of isoxaflutole (Dufourmantel et al., 2007, Plant Biotechnol J.5(1):118-33).

In WO 04/024928, the inventors have sought to increase the prenylquinone
15 biosynthesis (e.g., synthesis of plastoquinones, tocopherols) in the cells of plants by increasing the flux of the HPP precursor into the cells of these plants. This has been done by connecting the synthesis of said precursor to the "shikimate" pathway by overexpression of a PDH enzyme. They have also noted that the transformation of plants with a gene encoding a PDH enzyme makes it possible to increase the
20 tolerance of said plants to HPPD inhibitors.

In the patent application WO 2009/144079, a nucleic acid sequence encoding a mutated hydroxyphenylpyruvate dioxygenase (HPPD) at position 336 of the *Pseudomonas fluorescens* HPPD protein and its use for obtaining plants which are
25 tolerant to HPPD inhibitor herbicides is disclosed.

In WO 2002/046387, several domains of HPPD proteins originating from plants have been identified that may be relevant to confer tolerance to various HPPD inhibitor herbicides but no *in planta* nor biochemical data have been shown to
30 confirm the impact of the as described domain functions.

In WO 2008/150473, the combination of two distinct tolerance mechanisms – a modified *Avena sativa* gene coding for a mutant HPPD enzyme and a CYP450 Maize monooxygenase (nsf1 gene) – was exemplified in order to obtain an improved tolerance to HPPD inhibitor herbicides, but no data have been disclosed
5 demonstrating the synergistic effects based on the combination of both proteins.

Despite these successes obtained for the development of plants showing tolerance to several HPPD inhibitors herbicides described above, it is still necessary to develop and/or improve the tolerance of plants to newer or to several different
10 HPPD inhibitors, particularly HPPD inhibitors belonging to the classes of the triketones (e.g. sulcotrione, mesotrione, tembotrione, benzobicyclon and bicyclopyrone) and the pyrazolinates (e.g., topramezone and pyrasulfotole).

Description

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The present invention therefore relates to the generation of transgenic plants containing a gene encoding an HPPD protein obtainable or obtained from an organism belonging to the family of *Picrophilaceae*, and variants or mutants thereof, more especially to a gene from an organism belonging to the genus *Picrophilus*, and
20 variants or mutants thereof, coding for an HPPD enzyme showing the properties of catalysing the conversion of para-hydroxyphenylpyruvate to homogentisate and which plants are less sensitive to HPPD inhibitors than plants not containing any such HPPD encoding transgene.

25 More especially, the present invention therefore relates to the generation of transgenic plants containing a gene obtainable or obtained from an organism belonging to the family of *Picrophilaceae*, especially from the genus *Picrophilus*, more especially obtained from the species *Picrophilus torridus*, and variants or mutants thereof, coding for an HPPD enzyme showing the properties of catalysing
30 the conversion of para-hydroxyphenylpyruvate to homogentisate and which are less sensitive to HPPD inhibitors than plants not containing any such HPPD transgene.

The genes from Picrophilaceae, especially from the genus Picrophilus coding for HPPD proteins were selected as excellent HPPD-inhibitor tolerant candidates due to their high divergences in the amino acids composition at positions relevant for HPPD inhibitor tolerance as determined experimentally and structurally in the HPPD protein compared to the sensitive Arabidopsis HPPD protein which was taken as the HPPD-inhibitor herbicide sensitive reference molecule.

In one embodiment, this invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the Picrophilus HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368, particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6 or 7, preferably SEQ ID No. 6.

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In a further embodiment, the invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the Picrophilus HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368, particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6, 7, preferably SEQ ID No. 6, and in which any amino acids from position 177 to position 368 of SEQ ID No. 4 can be amended by any naturally-occurring amino acid, preferentially it can be any conservative substitution.

25

In a further embodiment, the invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the Picrophilus HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368,

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particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6, 7, preferably SEQ ID No. 6, and having one or more of the following amino acids at the position defined by its number (relating to the number of SEQ ID No. 4) given in brackets, i.e. His(175), Ser(218), Asn(232), Gln(256), His(257), Tyr(286), Gln(321),
 5 Phe(334), Glu(336), Gly(347), and Asn(350).

In a further embodiment, the invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the *Picrophilus* HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least
 10 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368, particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6, 7, preferably SEQ ID No. 6, and at the respective positions given in the second column of Table (i) the originally occurring amino acids can substituted by any of the amino
 15 acids listed in column 3 of Table (i).

Table (i):

Amino acid in SEQ ID No.4	Position in SEQ ID No.4	Substitutions
Val	177	Thr, Cys, Ala, Gly Phe, Tyr, Ile, Val, Ala, Gln, Glu, Asp, Gly, Thr, Ser, Met, Arg,
Leu	201	Lys
Ile	202	Ala, Trp, Leu, Ser, Arg, Lys, His, Asp, Glu, Pro, Gly, Asn
Phe	204	Val, Ile, Ala, Leu, Trp, Met, Gln, His
Leu	216	Met, Val
Lys	219	Ala, Val, Leu, Met, Ile, Arg, Gln, Tyr
Val	221	Leu, Met, Ile, Ala
Lys	222	Ala, Ser, Thr, Val, Arg, Glu, Leu, Ile, Met, His

Ala	353	Glu, Gln, Ser, Val, Phe, Thr
Leu	354	Arg

In a further embodiment, the invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the *Picrophilus* HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368, particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6, 7, preferably SEQ ID No. 6, and at the respective positions given in the second column of Table (ii) the originally occurring amino acids can substituted by any of the amino acids listed in column 3 of Table (ii).

Table (ii):

Amino acid in SEQ ID No.4	Position in SEQ ID No.4	Substitutions
Thr	203	Glu, Ser, Tyr, Phe, His, Gln, Asn, Gly, Leu, Met, Val, Arg, Ile
Val	220	Ala, Thr
Pro	230	Ala, Val, Thr, Asn, Ile,
Leu	280	Met, Ile, Asn
Leu	310	Met
Asn	348	Any except Pro
Gly	349	Ala, Pro, Val, Thr, Met

In a further embodiment, the invention relates to an HPPD protein named herein “the HPPD protein of this invention” or “the *Picrophilus* HPPD protein”, which is an HPPD protein with at least 75 %, at least 80 %, at least 85 %, at least 90 %, at least 95 %; at least 97 %; at least 98 %, or at least 99 % amino acid sequence identity to the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368,

particularly to the amino acid sequence of any one of SEQ ID Nos. 4, 5, 6, 7, preferably SEQ ID No. 6, and at the respective positions given in the second column of Table (iii) the originally occurring amino acids can substituted by any of the amino acids listed in column 3 of Table (iii).

5

Table (iii)

Amino acid in SEQ ID No.4	Position in SEQ ID No.4	Substitutions
Thr	203	Glu, Ser, Arg, Tyr
Val	220	Ala
Pro	230	Ala, Val, Thr
Leu	280	Met
Leu	310	Met
Asn	348	Ile, Ala, Val, Leu, Lys
Gly	349	Ala

10

This includes a protein with amino acids substituted, deleted or added compared to the sequence of SEQ ID No. 4 from amino acid position 2 to amino acid position 368, such as a transit peptide fusion protein, or a protein with amino acid changes in the sequence of SEQ ID No. 4 that retains the enzymatic function of an HPPD protein, and that still confers HPPD tolerance when expressed in plants, preferably HPPD tolerance of comparable range to that conferred by the protein of SEQ ID No. 4. This includes variant or mutant proteins derived from the protein of SEQ ID No. 4, such as any of the proteins of SEQ ID Nos 5, 6 or 7, particularly such mutant or variant which is less sensitive than the host plant's endogenous HPPD to an HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones or

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pyrazolines, preferably such mutant or variant which confers agronomically relevant herbicide tolerance to a host plant expressing it when an HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones and/or pyrazolines, particularly any one of mesotrione, tembotrione, isoxaflutole or bicyclopyrone is applied on such plants, more particularly when applied post-emergence. This also includes a protein comprising an active portion of the sequence of SEQ ID No. 4, which portion confers HPPD inhibitor tolerance when expressed in plants. This includes a protein with substantially the same amino acid sequence as the sequence of SEQ ID No. 4, such as a protein with the amino acid sequence of any one of SEQ ID Nos 4 to 7. This includes isolated proteins as defined below, and also proteins, such as the protein of SEQ ID No. 4 wherein certain amino acids have been replaced by similar amino acids as defined below, preferably conservative amino acid substitutions. Also included herein as HPPD proteins of this invention are HPPD proteins comprising the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to 368, but wherein 1-20, 1-15, 1-10, or 1, 2, 3, 4, 5, 6, 7, 8, or 9 amino acids have been deleted or have been substituted by other amino acids, particularly such protein which retains HPPD enzymatic activity and which confers tolerance to HPPD inhibitor herbicides when expressed in a host plant. Included herein are HPPD proteins encoded by DNA sequences homologous to the DNA sequences of the invention as described below, or HPPD proteins encoded by a DNA sequence which hybridizes to at least a portion (of at least 20-30 nucleotides) of the DNA of SEQ ID No. 1, or which is obtainable using a primer based on SEQ ID No. 1, or HPPD proteins with at least 75 % sequence identity to SEQ ID No. 4 which are encoded by a DNA sequence found in the genome sequence of a microorganism, such as a eukaryotic microorganism, particularly a Euryarchaeota, such as a microorganism of the family Picrophilus. Included herein as an HPPD protein of this invention is a Picrophilus HPPD protein which confers herbicide tolerance to plants when expressed in such plants, wherein such tolerance is to an HPPD inhibitor such as mesotrione, tembotrione, isoxaflutole or bicyclopyrone, particularly such HPPD protein is a Picrophilus torridus HPPD protein, such as a

protein comprising the sequence of SEQ ID No. 4 from amino acid position 2 to 368.

This includes the mutant or variant HPPD proteins as described further below.

The present invention includes and provides an antibody capable of specifically
5 binding a substantially purified protein comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 4, 5, 6, or 7, or derived sequences thereof according to amino acid replacement as disclosed in one or more of tables (i), (ii) or (iii), above.

10 A further aspect of the invention concerns antibodies, single-chain antigen binding molecules, or other proteins that specifically bind to one or more of the protein or peptide molecules of the invention and their homologs, fusions or fragments.
In a particularly preferred embodiment, the antibody specifically binds to a protein having the amino acid sequence set forth in SEQ ID NOs: 4-7 or a fragment thereof,
15 or derived sequences thereof according to amino acid replacement as disclosed in one or more of tables (i), (ii) or (iii), above.

In another embodiment, the antibody specifically binds to a fusion protein comprising an amino acid sequence selected from the amino acid sequence set
20 forth in SEQ ID NOs: 4-7 or a fragment thereof, or derived sequences thereof according to amino acid replacement as disclosed in one or more of tables (i), (ii) or (iii), above.

In another embodiment the antibody specifically binds to a fusion protein comprising
25 an amino acid sequence selected from the amino acid sequence set forth in SEQ ID NOs: 4-7 or a fragment thereof, or derived sequences thereof according to amino acid replacement as disclosed in one or more of tables (i), (ii) or (iii), above.

Antibodies of the invention may be used to quantitatively or qualitatively detect the
30 protein or peptide molecules of the invention, or to detect post translational modifications of the proteins. As used herein, an antibody or peptide is said to

“specifically bind” to a protein or peptide molecule of the invention if such binding is not competitively inhibited by the presence of non-related molecules.

In another embodiment this invention relates to an HPPD nucleic acid or DNA,
5 named herein “the HPPD nucleic acid/DNA of this invention”, which is a nucleic acid or DNA encoding an HPPD of this invention as defined above. This includes a DNA which comprises a nucleotide sequence selected from the group consisting of the sequence of SEQ ID No. 1 from nucleotide position 4 to nucleotide position 1104, the sequence of SEQ ID No. 2 from nucleotide position 25 to nucleotide position
10 1125, or the sequence of SEQ ID No. 3 from nucleotide position 4 to nucleotide position 1500, or which comprises a DNA region which encodes an HPPD, or a DNA which is sufficiently complementary to another DNA so that when it is incubated at a temperature of between 60 and 65° C in 5xSSC (1xSSC (single-strength sodium citrate) means = 0.15M NaCl, 0.015 M trisodium-citrate, 50 mM sodium phosphate
15 pH 7.6), containing 0.1% SDS followed by rinsing at the same temperature with 0.3 5xSSC containing 0.1% SDS, it still hybridizes with a sequence selected from the group consisting of SEQ ID Nos. 1, 2, and 3. When the test and inventive sequences are double stranded the nucleic acid constituting the test sequence preferably has a T_M within 10° C. of that of the sequence selected from the group
20 consisting of SEQ ID Nos 1, 2, and 3. In the case that the test and the sequence selected from the group consisting of SEQ ID Nos. 1, 2, and 3 are mixed together and are denatured simultaneously, the T_M values of the sequences are preferably within 5° C. of each other. More preferably the hybridization is performed under relatively stringent hybridization conditions as defined below. In one embodiment, a
25 denatured test or inventive sequence is preferably first bound to a support and hybridization is effected for a specified period of time at a temperature of between 60 and 65° C. in 5xSSC containing 0.1% SDS followed by rinsing of the support at the same temperature but with 0.1xSSC. Where the hybridization involves a fragment of the sequence selected from the group consisting of SEQ ID Nos. 1, 2,
30 and 3 the hybridization conditions may be less stringent, as will be obvious to the skilled person.

Also included herein as HPPD DNA of this invention, are DNA sequences encoding an HPPD protein of the invention which DNA sequences have been adapted for expression in microorganisms or plants, such as by replacing native codons by codons more preferred in a host cell, or wherein certain restriction sites have been
5 added or removed for ease of cloning, or DNA sequence with a certain number of added, replaced or deleted nucleotides. This also includes isolated DNA sequences and variant, mutant or synthetic DNAs or nucleic acids as described further below.

In a particular embodiment, the *Picrophilus* HPPD DNA of this invention is
10 expressed in plants under the control of a promoter that allows expression of exogenous genes in plants. In a further particular embodiment, at the N-terminus of the so expressed HPPD enzyme a signal peptide, such as a transit peptide is located, preferably a plastid transit peptide, such as a chloroplast transit peptide of about 120 amino acids (about 30 to about 120 amino acids) most preferably a
15 double transit peptide, such as an optimized transit peptide of which the first part is originated from Sunflower (*Helianthus annuus*) and the second part from Zea mays (described in US patent 5,188,642) or a plastid transit peptide of that of the plant ribulose biscarboxylase/ oxygenase small subunit (RuBisCO ssu), where appropriate including a few amino acids of the N-terminal part of the mature
20 RuBisCO ssu (EP 189 707)

In a further particular embodiment, this invention includes a DNA encoding an HPPD protein of this invention which is derived or is obtainable from SEQ ID No. 1 and is optimized for the expression in *E. coli*, such as a codon-optimized DNA, for example
25 a DNA comprising the sequence of SEQ ID No. 2 from nucleotide position 25 to nucleotide position 1125 (including the positions defined).

In a further particular embodiment, this invention includes a DNA encoding an HPPD protein of this invention which is derived from SEQ ID No. 1 and optimized for the
30 expression in plants, such as a codon-optimized DNA, for example a DNA

comprising the sequence of "SEQ ID No. 3 from nucleotide position 400 to nucleotide position 1500 (including the positions defined).

- In a further particular embodiment, the HPPD of the invention, such as the HPPD
- 5 comprising the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to amino acid position 368, or the HPPD comprising the amino acid sequence of any one of SEQ ID Nos. 4 to 7, is less sensitive than the host plant endogenous HPPD to an HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones or pyrazolates, or an HPPD inhibitor herbicide selected from isoxaflutole,
- 10 tembotrione, mesotrione, sulcotrione, pyrasulfotole, topramezone, 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-CF₃phenyl)propane-1,3-dione and 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-2,3 Cl₂ phenyl)propane-1,3-dione, bicyclopyrone, benzobicyclon, tefuryltrione, and pyrazoxyfen.
- 15 In a further particular embodiment, this invention includes a DNA encoding an HPPD protein of this invention which is derived from SEQ ID No. 1 and optimized for the expression in *E. coli*, such as a codon-optimized DNA, for example a DNA comprising the sequence of "SEQ ID No. 2 from nucleotide position 25 to nucleotide position 1125 (including the positions defined) which encodes an HPPD less
- 20 sensitive than the host plant endogenous HPPD to at least one HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones or pyrazolates, preferably to tembotrione, mesotrione, bicyclopyrone, tefuryltrione, isoxaflutole, diketonitrile, pyrasulfotole, topramezone, sulcotrione, pyrazolate and benzofenap.
- 25 In a further particular embodiment, this invention includes a DNA encoding an HPPD protein of this invention which is optimized for the expression in plants, such as a codon-optimized DNA, for example a DNA comprising the sequence of SEQ ID No. 3 from nucleotide position 400 to nucleotide position 1500 (including the positions defined) which encodes an HPPD less sensitive than the host plant
- 30 endogenous HPPD to at least one HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones or pyrazolates, preferably to tembotrione,

mesotrione, bicyclopyrone, tefuryltrione, isoxaflutole, diketonitrile, pyrasulfotole, topramezone, sulcotrione, pyrazolate and benzofenap.

In a further particular embodiment, this invention relates to plants, plant parts, plant
5 cells, and progenies of these plants comprising a DNA encoding an HPPD protein of
this invention which is optimized for the expression in *E. coli*, or is optimized for the
expression in plants such as a codon-optimized DNA, for example a DNA
comprising the sequence of SEQ ID No. 2 from nucleotide position 25 to nucleotide
position 1125 (including the positions defined) or of SEQ ID No. 3 from nucleotide
10 position 400 to nucleotide position 1500 (including the positions defined) which
encodes an HPPD less sensitive than the host plant endogenous HPPD. Such
plants include but are not limited to field crops, fruits and vegetables such as
canola, sunflower, tobacco, sugarbeet, cotton, maize, wheat, barley, rice, sorghum,
tomato, mango, peach, apple, pear, strawberry, banana, melon, potato, carrot,
15 lettuce, cabbage, onion, soya spp, sugar cane, pea, field beans, poplar, grape,
citrus, alfalfa, rye, oats, turf and forage grasses, flax and oilseed rape, and nut
producing plants.

In a more particular embodiment, this invention relates to plants, plant parts, plant
20 cells, and progenies of these plants comprising any of the DNA encoding an HPPD
protein of the invention which is optimized for the expression in *E. coli*, or optimized
for the expression in plants such as a codon-optimized DNA, for example a DNA
comprising the sequence of "SEQ ID No. 2 from nucleotide position 25 to nucleotide
position 1125 (including the positions defined) or of SEQ ID No. 3 from nucleotide
25 position 400 to nucleotide position 1500 (including the positions defined) which
encodes an HPPD less sensitive than the host plant endogenous HPPD and
wherein the plants are selected from the group consisting of canola, sunflower,
tobacco, sugarbeet, cotton, maize, wheat, barley, rice, potato, soya spp, sugar cane,
pea, field beans, poplar, grape, alfalfa, rye, oats, turf and forage grasses, flax and
30 oilseed rape, and nut producing plants, even more preferably such plants are

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selected from the group consisting of soya spp, rice, sugarbeet, wheat, cotton
canola, oilseed rape or maize.

In another particular embodiment, the HPPD protein of the invention comprises the
5 sequence of SEQ ID No. 7 and is less sensitive to an HPPD inhibitor of the class of
triketones (named triketone HPPD inhibitor), such as tembotrione, sulcotrione
mesotrione, bicyclopyrone, tefuryltrione, particularly tembotrione, or of the class
diketonitrile (isoxaflutole) or of the class of pyrazolines (named pyrazoline HPPD
inhibitor), such as pyrasulfotole, pyrazolate, topramezone, benzofenap compared to
10 the endogenous unmutated HPPD of a plant, particularly the host plant wherein
such HPPD of the invention is expressed or is to be expressed.

The enzymatic activity of HPPD proteins can be measured by any method that
makes it possible either to measure the decrease in the amount of the HPP or O₂
15 substrates, or to measure the accumulation of any of the products derived from the
enzymatic reaction, i.e. homogentisate or CO₂. In particular, the HPPD activity can
be measured by means of the method described in Garcia et al. (1997), Biochem. J.
325, 761-769 or Garcia et al. (1999), Plant Physiol. 119, 1507-1516.

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According to the invention, an HPPD inhibitor of the class of triketones (or triketone
HPPD inhibitor) means an HPPD inhibitor having a triketone skeleton. As an
example of such triketone HPPD inhibitor, one can cite the molecules sulcotrione
[i.e. 2-[2-chloro-4-(methylsulfonyl)benzoyl]-1,3-cyclohexanedione], mesotrione [i.e. 2-
25 [4-(methylsulfonyl)-2-nitrobenzoyl]-1,3-cyclohexanedione], and tembotrione [i.e. 2-[2-
chloro-4-(methylsulfonyl)-3-[(2,2,2-trifluoroethoxy)methyl]benzoyl]-1,3-cyclo-
hexanedione], tefuryltrione [i.e. 2-{2-chloro-4-mesyl-3-[(RS)-tetrahydro-2-
furylmethoxymethyl]benzoyl}cyclohexane-1,3-dione], bicyclopyrone [i.e. 4-hydroxy-
3-{2-[(2-methoxyethoxy)methyl]-6-(trifluoromethyl)-3-
30 pyridylcarbonyl}bicyclo[3.2.1]oct-3-en-2-one], benzobicyclon [i.e. 3-(2-chloro-4-
mesylbenzoyl)-2-phenylthiobicyclo[3.2.1]oct-2-en-4-one].

According to the invention, an HPPD of the class of pyrazolines (or pyrazoline HPPD inhibitor) means an HPPD inhibitor having a pyrazole radical. As an example of such pyrazolines HPPD inhibitor, one can cite the molecules topramezone [i.e. [3-(4,5-dihydro-3-isoxazolyl)-2-methyl-4-(methylsulfonyl)phenyl](5-hydroxy-1-methyl-1H-pyrazol-4-yl)methanone] and pyrasulfotole [(5-hydroxy-1,3-dimethylpyrazol-4-yl)(2-methyl-4-trifluoromethylphenyl)methanone].

The present invention also relates to a nucleic acid sequence, particularly an isolated DNA, preferably a plant-expressible chimeric gene, which encodes the Picrophilus HPPD of the invention and adapted sequences thereof.

The present invention also relates to a nucleic acid sequence encoding an HPPD enzyme of this invention which retains its properties of catalysing the conversion of para-hydroxyphenylpyruvate to homogentisate and which is less sensitive to HPPD inhibitors of the class of triketones such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolines such as pyrasulfotole and topramezone, tefuryltrione, bicyclopyrone, benzobicyclon than the endogenous unmutated plant HPPD, and of which the encoded amino acid sequence shows a sequence identity to SEQ ID No. 4 of at least 75 %, 80 %, particularly at least 85%, preferably at least 90%, more preferably at least 95%, even more preferably at least 98% and most preferably at least 99%.

In a more particular embodiment, the nucleic acid sequence of the invention encodes an HPPD enzyme which is less sensitive to an HPPD inhibitor of the class of triketones such as tembotrione, sulcotrione, mesotrione, bicyclopyrone, and tefuryltrione, the class of isoxazoles such as isoxaflutole of the class of pyrazolines (named pyrazoline HPPD inhibitor), such as pyrasulfotole, pyrazolate, topramezone, benzofenap, or the class of diketones such as diketonitrile than the host plant endogenous HPPD.

According to the present invention, a "nucleic acid sequence" is understood as being a nucleotide sequence which can be of the DNA or RNA type, preferably of the DNA type, and in particular double-stranded, whether it be of natural or synthetic origin, in particular a DNA sequence in which the codons which encode the HPPD

5 according to the invention have been optimized in accordance with the host organism in which it is to be expressed (e.g., by replacing codons with those codons more preferred or most preferred in codon usage tables of such host organism or the group to which such host organism belongs, compared to the original or source organism).

10 An "isolated nucleic acid/DNA/protein", as used herein, refers to a nucleic acid/DNA/protein which is not naturally occurring (such as an artificial or synthetic DNA with a different nucleotide sequence than the naturally occurring DNA, or a modified protein) or which is no longer in the natural environment wherein it was originally present, e.g., a DNA coding sequence associated with a heterologous
15 regulatory element (such as a bacterial coding sequence operably linked to a plant-expressible promoter) in a chimeric gene, a DNA transferred into another host cell, such as a transgenic plant cell.

In view of a particular embodiment of the invention and the sought-after solution, i.e.
20 an HPPD which is less sensitive to a triketone, an isoxazole, or pyrazolate HPPD inhibitor, the tolerance level measurement is analyzed using the method extensively described in WO 2009/14407 as described below using a triketone, an isoxazole, or a pyrazolate HPPD inhibitor, particularly an HPPD inhibitor selected from tembotrione, mesotrione, pyrasulfotole, topramezone sulcotrione, bicyclopyrone,
25 diketonitrile, benzofenap, pyrazolate, tefuryltrione.

The terminology DNA or protein "comprising" a certain sequence "X", as used throughout the text, refers to a DNA or protein including or containing at least the sequence "X", so that other nucleotide or amino acid sequences can be included at
30 the 5' (or N-terminal) and/or 3' (or C-terminal) end, e.g. (the nucleotide sequence of) a selectable marker protein, (the nucleotide sequence of) a transit peptide, and/or a

5' leader sequence or a 3' trailer sequence. Similarly, use of the term "comprise", "comprising" or "comprises" throughout the text and the claims of this application should be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of
5 integers or steps.

In one embodiment of the invention, the coding regions encoding HPPD comprise a nucleotide sequence encoding proteins with the amino acid sequences as set forth in SEQ ID Nos 4, 5, 6, and 7 such as the nucleotide sequences of SEQ ID Nos 1, 2,
10 and 3.

However, it will be clear that variants of these nucleotide sequences, including insertions, deletions and substitutions thereof may be also be used to the same effect. Equally, homologues to the mentioned nucleotide sequences from species
15 different from *Picrophilus torridus* can be used.

Variants of the described nucleotide sequence will have a sequence identity which is preferably at least about 70%, 80%, 85% or 90% or at least 95% with identified nucleotide sequences encoding HPPD enzymes such as the ones identified in the
20 sequence listing.

A protein with "substantially the same amino acid sequence" to a protein as described in the invention, as used herein, refers to a protein with at least 90 %, particularly at least 95 %, preferably at least 97 % sequence identity with a protein
25 according to the invention, wherein the percentage sequence identity is determined by using the blosum62 scoring matrix in the GAP program of the Wisconsin package of GCG (Madison, Wisconsin, USA) version 10.0 (GCG defaults used). "Sequence identity", as used throughout this application, when related to proteins, refers to the percentage of identical amino acids using this specified analysis. The "sequence
30 identity", as used herein, when related to DNA sequences, is determined by using

the nwsgapdna scoring matrix in the GAP program of the Wisconsin package of GCG (Madison, Wisconsin, USA) version 10.0 (GCG defaults used).

For the purpose of this invention, the "sequence identity" of two related nucleotide
5 or amino acid sequences, expressed as a percentage, refers to the number of
positions in the two optimally aligned sequences which have identical residues
(x100) divided by the number of positions compared. A gap, i.e. a position in an
alignment where a residue is present in one sequence but not in the other, is
regarded as a position with non-identical residues. The alignment of the two
10 sequences is performed by the Needleman and Wunsch algorithm (Needleman and
Wunsch 1970). The computer-assisted sequence alignment above, can be
conveniently performed using standard software program such as GAP which is part
of the Wisconsin Package Version 10.1 (Genetics Computer Group, Madison,
Wisconsin, USA) using the default scoring matrix with a gap creation penalty of 50
15 and a gap extension penalty of 3.

Nucleotide sequences homologous to the nucleotide sequences encoding an HPPD
enzyme according to the invention may be identified by in silico analysis of genomic
sequence data.

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Homologous nucleotide sequence may also be identified and isolated by
hybridization under stringent conditions using as probes identified nucleotide
sequences encoding HPPD enzymes according to the invention or parts thereof.
Such parts should preferably have a nucleotide sequence comprising at least 40
25 consecutive nucleotides from the coding region of HPPD encoding genes
sequences according to the invention, preferably from the coding region of
SEQ ID No. 1, SEQ ID No. 2 or SEQ ID No. 3. The probes may however comprise
longer regions of nucleotide sequences derived from the HPPD encoding nucleic
acids, such as about 50, 60, 75, 100, 200 or 500 consecutive nucleotides from any
30 of the mentioned HPPD genes. Preferably, the probe should comprise a nucleotide

sequence coding for a highly conserved region which may be identified by aligning the different HPPD proteins.

“Stringent hybridization conditions” as used herein means that hybridization will generally occur if there is at least 95% and preferably at least 97% sequence identity between the probe and the target sequence. Examples of stringent hybridization conditions are overnight incubation in a solution comprising 5xSSC (150 mM NaCl, 15 mM trisodium-citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared carrier DNA such as salmon sperm DNA, followed by washing the hybridization support in 0.1 x SSC at approximately 65 °C, preferably twice for about 10 minutes. Other hybridization and wash conditions are well known and are exemplified in Sambrook et al, Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, NY (1989), particularly chapter 11.

Such variant sequences may also be obtained by DNA amplification using oligonucleotides specific for HPPD genes encoding enzymes as primers, such as but not limited to oligonucleotides comprising about 20 to about 50 consecutive nucleotides selected from the nucleotide sequences of SEQ ID Nos 1, 2, 3 or their complement.

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The invention also encompasses variant HPPD enzymes which are amino acid sequences similar to the HPPD amino acid sequence of SEQ ID No. 4 wherein one or more amino acids have been inserted, deleted or substituted. In the present context, variants of an amino acid sequence refer to those polypeptides, enzymes or proteins which have a similar catalytic activity as the amino acid sequences described herein, notwithstanding any amino acid substitutions, additions or deletions thereto. Preferably the variant amino acid sequence has a sequence identity of at least about 80%, or 85 or 90% or 95% with the amino acid sequence of SEQ ID No. 4. Also preferably, a polypeptide comprising the variant amino acid sequence has HPPD enzymatic activity. Methods to determine HPPD enzymatic

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activity are well known in the art and include assays as extensively described in WO 2009/144079 or in WO 2002/046387.

Substitutions encompass amino acid alterations in which an amino acid is replaced
5 with a different naturally-occurring or a non-conventional amino acid residue. Such substitutions may be classified as "conservative", in which an amino acid residue contained in an HPPD protein of this invention is replaced with another naturally-occurring amino acid of similar character, for example Gly↔Ala, Val↔Ile↔Leu, Asp↔Glu, Lys↔Arg, Asn↔Gln or Phe↔Trp↔Tyr. Substitutions encompassed by
10 the present invention may also be "non-conservative", in which an amino acid residue which is present in an HPPD protein of the invention is substituted with an amino acid with different properties, such as a naturally-occurring amino acid from a different group (eg. substituting a charged or hydrophobic amino acid with alanine. Amino acid substitutions are typically of single residues, but may be of multiple
15 residues, either clustered or dispersed. Amino acid deletions will usually be of the order of about 1-10 amino acid residues, while insertions may be of any length. Deletions and insertions may be made to the N-terminus, the C-terminus or be internal deletions or insertions. Generally, insertions within the amino acid sequence will be smaller than amino- or carboxy-terminal fusions and of the order of 1 to 4
20 amino acid residues. "Similar amino acids", as used herein, refers to amino acids that have similar amino acid side chains, i.e. amino acids that have polar, non-polar or practically neutral side chains. "Non-similar amino acids", as used herein, refers to amino acids that have different amino acid side chains, for example an amino acid with a polar side chain is non-similar to an amino acid with a non-polar side
25 chain. Polar side chains usually tend to be present on the surface of a protein where they can interact with the aqueous environment found in cells ("hydrophilic" amino acids). On the other hand, "non-polar" amino acids tend to reside within the center of the protein where they can interact with similar non-polar neighbours ("hydrophobic" amino acids"). Examples of amino acids that have polar side chains
30 are arginine, asparagine, aspartate, cysteine, glutamine, glutamate, histidine, lysine, serine, and threonine (all hydrophilic, except for cysteine which is hydrophobic).

Examples of amino acids that have non-polar side chains are alanine, glycine, isoleucine, leucine, methionine, phenylalanine, proline, and tryptophan (all hydrophobic, except for glycine which is neutral).

- 5 Also encompassed by the present invention are antibodies which specifically recognize a HPPD enzyme according to the invention.

The invention also relates to the use, in a method for transforming plants, of a nucleic acid which encodes an HPPD according to the invention as a marker gene
10 or as a coding sequence which makes it possible to confer to the plant tolerance to herbicides which are HPPD inhibitors, and the use of HPPD inhibitors on plants comprising a nucleic acid sequence encoding an HPPD according to the invention. In an embodiment of this invention, in such use the HPPD inhibitors are triketones or pyrazolines, preferably tembotrione, mesotrione or sulcotrione, bicyclopyrone,
15 and tefuryltrione. It is, of course, understood that this sequence can also be used in combination with (an) other gene marker(s) and/or sequence(s) which encode(s) one or more protein with useful agricultural properties.

In the commercial production of crops, it is desirable to eliminate under reliable
20 pesticidal management unwanted plants (i.e., "weeds") from a field of crop plants. An ideal treatment would be one which could be applied to an entire field but which would eliminate only the unwanted plants while leaving the crop plants unaffected. One such treatment system would involve the use of crop plants which are tolerant to an herbicide so that when the herbicide is sprayed on a field of herbicide-tolerant
25 crop plants, the crop plants would continue to thrive while non-herbicide-tolerant weeds are killed or severely damaged. Ideally, such treatment systems would take advantage of varying herbicide properties so that weed control could provide the best possible combination of flexibility and economy. For example, individual herbicides have different longevities in the field, and some herbicides persist and
30 are effective for a relatively long time after they are applied to a field while other herbicides are quickly broken down into other and/or non-active compounds. An

ideal treatment system would allow the use of different herbicides so that growers could tailor the choice of herbicides for a particular situation.

While a number of herbicide-tolerant crop plants are presently commercially
5 available, one issue that has arisen for many commercial herbicides and
herbicide/crop combinations is that individual herbicides typically have incomplete
spectrum of activity against common weed species. For most individual herbicides
which have been in use for some time, populations of herbicide resistant weed
species and biotypes have become more prevalent (see, e.g., Tranel and Wright
10 (2002) *Weed Science* 50: 700-712; Owen and Zelaya (2005) *Pest Manag. Sci.* 61:
301-311). Transgenic plants which are resistant to more than one herbicide have
been described (see, e.g., W02005/012515). However, improvements in every
aspect of crop production, weed control options, extension of residual weed control,
and improvement in crop yield are continuously in demand.

15 The HPPD protein or gene of the invention is advantageously combined in plants
with other genes which encode proteins or RNAs that confer useful agronomic
properties to such plants. Among the genes which encode proteins or RNAs that
confer useful agronomic properties on the transformed plants, mention can be made
20 of the DNA sequences encoding proteins which confer tolerance to one or more
herbicides that, according to their chemical structure, differ from HPPD inhibitor
herbicides, and others which confer tolerance to certain insects, those which confer
tolerance to certain diseases, DNAs that encodes RNAs that provide nematode or
insect control, etc...

25 Such genes are in particular described in published PCT Patent Applications WO
91/02071 and WO95/06128.

Among the DNA sequences encoding proteins which confer tolerance to certain
herbicides on the transformed plant cells and plants, mention can be made of a bar
or PAT gene or the *Streptomyces coelicolor* gene described in WO2009/152359
30 which confers tolerance to glufosinate herbicides, a gene encoding a suitable
EPSPS which confers tolerance to herbicides having EPSPS as a target, such as

glyphosate and its salts (US 4,535,060, US 4,769,061, US 5,094,945, US 4,940,835, US 5,188,642, US 4,971,908, US 5,145,783, US 5,310,667, US 5,312,910, US 5,627,061, US 5,633,435), or a gene encoding glyphosate oxydoreductase (US 5,463,175).

- 5 Among the DNA sequences encoding a suitable EPSPS which confer tolerance to the herbicides which have EPSPS as a target, mention will more particularly be made of the gene which encodes a plant EPSPS, in particular maize EPSPS, particularly a maize EPSPS which comprises two mutations, particularly a mutation at amino acid position 102 and a mutation at amino acid position 106
- 10 (WO 2004/074443), and which is described in Patent Application US 6566587, hereinafter named double mutant maize EPSPS or 2mEPSPS, or the gene which encodes an EPSPS isolated from *Agrobacterium* and which is described by SEQ ID No. 2 and SEQ ID No. 3 of US Patent 5,633,435, also named CP4.

- Among the DNA sequences encoding a suitable EPSPS which confer tolerance to
- 15 the herbicides which have EPSPS as a target, mention will more particularly be made of the gene which encodes an EPSPS GRG23 from *Arthrobacter globiformis*, but also the mutants GRG23 ACE1, GRG23 ACE2, or GRG23 ACE3, particularly the mutants or variants of GRG23 as described in WO2008/100353, such as GRG23(ace3)R173K of SEQ ID No. 29 in WO2008/100353.

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In the case of the DNA sequences encoding EPSPS, and more particularly encoding the above genes, the sequence encoding these enzymes is advantageously preceded by a sequence encoding a transit peptide, in particular the "optimized transit peptide" described in US Patent 5,510,471 or 5,633,448.

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In WO 2007/024782, plants being tolerant to glyphosate and at least one ALS (acetolactate synthase) inhibitor are disclosed. More specifically plants containing genes encoding a GAT (Glyphosate-N-Acetyltransferase) polypeptide and a polypeptide conferring resistance to ALS inhibitors are disclosed.

- 30 In US 6855533, transgenic tobacco plants containing mutated *Arabidopsis* ALS/AHAS genes were disclosed.

In US 6,153,401, plants containing genes encoding 2,4-D-monooxygenases conferring tolerance to 2,4-D (2,4-dichlorophenoxyacetic acid) by metabolism are disclosed.

In US 2008/0119361 and US 2008/0120739, plants containing genes encoding
5 Dicamba monooxygenases conferring tolerance to dicamba (3,6-dichloro-2-methoxybenzoic acid) by metabolism are disclosed.

All the above mentioned herbicide tolerance traits can be combined with those performing HPPD tolerance which are subject matter of this invention.

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Among the DNA sequences encoding proteins concerning properties of tolerance to insects, mention will more particularly be made of the Bt proteins widely described in the literature and well known to those skilled in the art. Mention will also be made of proteins extracted from bacteria such as *Photorhabdus* (WO 97/17432 & WO
15 98/08932).

Among such DNA sequences encoding proteins of interest which confer novel properties of tolerance to insects, mention will more particularly be made of the Bt Cry or VIP proteins widely described in the literature and well known to those skilled in the art. These include the Cry1F protein or hybrids derived from a Cry1F protein
20 (e.g., the hybrid Cry1A-Cry1F proteins described in US 6,326,169; US 6,281,016; US 6,218,188, or toxic fragments thereof), the Cry1A-type proteins or toxic fragments thereof, preferably the Cry1Ac protein or hybrids derived from the Cry1Ac protein (e.g., the hybrid Cry1Ab-Cry1Ac protein described in US 5,880,275) or the Cry1Ab or Bt2 protein or insecticidal fragments thereof as described in EP451878,
25 the Cry2Ae, Cry2Af or Cry2Ag proteins as described in WO02/057664 or toxic fragments thereof, the Cry1A.105 protein described in WO 2007/140256 (SEQ ID No. 7) or a toxic fragment thereof, the VIP3Aa19 protein of NCBI accession ABG20428, the VIP3Aa20 protein of NCBI accession ABG20429 (SEQ ID No. 2 in WO 2007/142840), the VIP3A proteins produced in the COT202 or COT203 cotton
30 events (WO 2005/054479 and WO 2005/054480, respectively), the Cry proteins as described in WO01/47952, the VIP3Aa protein or a toxic fragment thereof as

described in Estruch et al. (1996), Proc Natl Acad Sci U S A. 28;93(11):5389-94 and US 6,291,156, the insecticidal proteins from *Xenorhabdus* (as described in WO98/50427), *Serratia* (particularly from *S. entomophila*) or *Photorhabdus* species strains, such as Tc-proteins from *Photorhabdus* as described in WO98/08932 (e.g.,
5 Waterfield et al., 2001, Appl Environ Microbiol. 67(11):5017-24; Ffrench-Constant and Bowen, 2000, Cell Mol Life Sci.; 57(5):828-33). Also any variants or mutants of any one of these proteins differing in some (1-10, preferably 1-5) amino acids from any of the above sequences, particularly the sequence of their toxic fragment, or which are fused to a transit peptide, such as a plastid transit peptide, or another
10 protein or peptide, is included herein.

The present invention also relates to a chimeric gene (or expression cassette) which comprises a coding sequence as well as heterologous regulatory elements, at the 5' and/or 3' position, at least at the 5' position, which are able to function in a host
15 organism, in particular plant cells or plants, with the coding sequence containing at least one nucleic acid sequence which encodes an HPPD as previously defined.

In a particular embodiment, the present invention relates to a chimeric gene as previously described, wherein the host organism is selected from bacteria, yeast,
20 *Pichia*, fungi, baculovirus, in vitro cells, protoplasts, plant cells, plants, plant parts, and plant seeds thereof.

In another particular embodiment, the present invention relates to a chimeric gene as previously described, wherein the chimeric gene contains in the 5' position of the
25 nucleic acid sequence which encodes a HPPD according to the invention, a nucleic acid sequence which encodes a plant transit peptide, with this sequence being arranged between the promoter region and the sequence encoding the HPPD according to the invention so as to permit expression of a transit peptide/HPPD fusion protein.

In a further particular embodiment, the present invention relates to the use of HPPD inhibitor herbicides on plants, plant parts, or plant seeds comprising HPPD tolerant gene according to the invention, or to the use of HPPD inhibitor herbicides on soil where such plants, plant parts or seeds are to be grown or sown, either alone or in combination with one or more other known herbicides acting in a different matter to HPPD inhibitors. In a more particular embodiment, the employed HPPD inhibitor herbicide is selected from the group consisting of triketones (named triketone HPPD inhibitor), such as tembotrione, sulcotrione mesotrione, bicyclopyrone, tefuryltrione, particularly tembotrione, of the class diketone such as diketonitrile of the class of isoxazoles such as isoxaflutole or of the class of pyrazolates (named pyrazolate HPPD inhibitor), such as pyrasulfotole, pyrazolate, topramezone, benzofenap, even more specifically present invention relates to the application of tembotrione, mesotrione, diketonitrile, bicyclopyrone, tefuryltrione, benzofenap, pyrasulfotole, pyrazolate and sulcotrione to such HPPD inhibitor tolerant plants, plant parts or plant seeds.

As a regulatory sequence which functions as a promoter in plant cells and plants, use may be made of any promoter sequence of a gene which is naturally expressed in plants, in particular a promoter which is expressed especially in the leaves of plants, such as for example "constitutive" promoters of bacterial, viral or plant origin, or "light-dependent" promoters, such as that of a plant ribulose-biscarboxylase/oxygenase (RuBisCO) small subunit gene, or any suitable known promoter-expressible which may be used. Among the promoters of plant origin, mention will be made of the histone promoters as described in EP 0 507 698 A1, the rice actin promoter (US 5,641,876), or a plant ubiquitin promoter (US 5,510,474). Among the promoters of a plant virus gene, mention will be made of that of the cauliflower mosaic virus (CaMV 19S or 35S, Sanders et al. (1987), Nucleic Acids Res. 15(4):1543-58.), the circovirus (AU 689 311) or the Cassava vein mosaic virus (CsVMV, US 7,053,205).

In one embodiment of this invention, a promoter sequence specific for particular regions or tissues of plants can be used to express the HPPD proteins of the invention, such as promoters specific for seeds (Datla, R. et al., 1997, Biotechnology Ann. Rev. 3, 269-296), especially the napin promoter (EP 255 378 A1), the phaseolin promoter, the glutenin promoter, the helianthinin promoter (WO 92/17580), the albumin promoter (WO 98/45460), the oleosin promoter (WO 98/45461), the SAT1 promoter or the SAT3 promoter (PCT/US98/06978).

Use may also be made of an inducible promoter advantageously chosen from the phenylalanine ammonia lyase (PAL), HMG-CoA reductase (HMG), chitinase, glucanase, proteinase inhibitor (PI), PR1 family gene, nopaline synthase (nos) and vspB promoters (US 5 670 349, Table 3), the HMG2 promoter (US 5 670 349), the apple beta-galactosidase (ABG1) promoter and the apple aminocyclopropane carboxylate synthase (ACC synthase) promoter (WO 98/45445).

According to the invention, use may also be made, in combination with the promoter, of other regulatory sequences, which are located between the promoter and the coding sequence, such as transcription activators ("enhancers"), for instance the translation activator of the tobacco mosaic virus (TMV) described in Application WO 87/07644, or of the tobacco etch virus (TEV) described by Carrington & Freed 1990, J. Virol. 64: 1590-1597, for example, or introns such as the adh1 intron of maize or intron 1 of rice actin.

In a further particular embodiment, the gene of the invention is present in plants in multiple, preferably two copies, each of these controlled by a different plant expressible promoter.

In a further particular embodiment, the chimeric gene of the invention can be combined with any further chimeric gene coding for an HPPD protein, preferably these different genes are controlled by different regulatory elements being active in plants.

In a further particular embodiment, the chimeric gene of the invention can be combined with a CYP450 Maize monooxygenase (nsf1 gene) gene being under the control of an identical or different plant expressible promoter.

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As a regulatory terminator or polyadenylation sequence, use may be made of any corresponding sequence of bacterial origin, such as for example the nos terminator of *Agrobacterium tumefaciens*, of viral origin, such as for example the CaMV 35S
10 terminator, or of plant origin, such as for example a histone terminator as described in published Patent Application EP 0 633 317 A1.

The term "gene", as used herein refers to a DNA coding region flanked by 5' and/or 3' regulatory sequences allowing a RNA to be transcribed which can be translated to
15 a protein, typically comprising at least a promoter region. A "chimeric gene", when referring to an HPPD encoding DNA of this invention, refers to an HPPD encoding DNA sequence having 5' and/or 3' regulatory sequences different from the naturally occurring Euryarchaeota 5' and/or 3' regulatory sequences which drive the expression of the HPPD protein in its native host cell (also referred to as
20 "heterologous promoter" or "heterologous regulatory sequences").

The terms "DNA/protein comprising the sequence X" and "DNA/protein with the sequence comprising sequence X", as used herein, refer to a DNA or protein including or containing at least the sequence X in their nucleotide or amino acid sequence, so that other nucleotide or amino acid sequences can be included at the
25 5' (or N-terminal) and/or 3' (or C-terminal) end, e.g., a N-terminal transit or signal peptide. The term "comprising", as used herein, is open-ended language in the meaning of "including", meaning that other elements than those specifically recited can also be present. The term "consisting of", as used herein, is closed-ended language, i.e., only those elements specifically recited are present. The term "DNA
30 encoding a protein comprising sequence X", as used herein, refers to a DNA comprising a coding sequence which after transcription and translation results in a

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protein containing at least amino acid sequence X. A DNA encoding a protein need not be a naturally occurring DNA, and can be a semi-synthetic, fully synthetic or artificial DNA and can include introns and 5' and/or 3' flanking regions. The term "nucleotide sequence", as used herein, refers to the sequence of a DNA or RNA molecule, which can be in single- or double-stranded form.

HPPD proteins according to the invention may be equipped with a signal peptide according to procedures known in the art, see, e.g., published PCT patent application WO 96/10083, or they can be replaced by another peptide such as a chloroplast transit peptide (e.g., Van Den Broeck et al., 1985, Nature 313, 358, or a modified chloroplast transit peptide of US patent 5, 510,471) causing transport of the protein to the chloroplasts, by a secretory signal peptide or a peptide targeting the protein to other plastids, mitochondria, the ER, or another organelle, or it can be replaced by a methionine amino acid or by a methionine-alanine dipeptide. Signal sequences for targeting to intracellular organelles or for secretion outside the plant cell or to the cell wall are found in naturally targeted or secreted proteins, preferably those described by Klösgen et al. (1989, Mol. Gen. Genet. 217, 155-161), Klösgen and Weil (1991, Mol. Gen. Genet. 225, 297-304), Neuhaus & Rogers (1998, Plant Mol. Biol. 38, 127-144), Bih et al. (1999, J. Biol. Chem. 274, 22884-22894), Morris et al. (1999, Biochem. Biophys. Res. Commun. 255, 328-333), Hesse et al. (1989, EMBO J. 8 2453-2461), Tavladoraki et al. (1998, FEBS Lett. 426, 62-66), Terashima et al. (1999, Appl. Microbiol. Biotechnol. 52, 516-523), Park et al. (1997, J. Biol. Chem. 272, 6876-6881), Shcherban et al. (1995, Proc. Natl. Acad. Sci USA 92, 9245-9249), particularly the signal peptide sequences from targeted or secreted proteins of corn, cotton, soybean, or rice. A DNA sequence encoding such a plant signal peptide can be inserted in the chimeric gene encoding the HPPD protein for expression in plants

Unless otherwise stated in the examples, all procedures for making and manipulating recombinant DNA are carried out by the standard procedures described in Sambrook et al., Molecular Cloning - A Laboratory Manual, Second

- Ed., Cold Spring Harbor Laboratory Press, NY (1989), and in Volumes 1 and 2 of Ausubel et al. (1994) Current Protocols in Molecular Biology, Current Protocols, USA. Standard materials and methods for plant molecular biology work are described in Plant Molecular Biology Labfax (1993) by R.R.D. Croy, jointly published
- 5 by BIOS Scientific Publications Ltd (UK) and Blackwell Scientific Publications (UK). Procedures for PCR technology can be found in "PCR protocols: a guide to methods and applications", Edited by M.A. Innis, D.H. Gelfand, J.J. Sninsky and T.J. White (Academic Press, Inc., 1990).
- 10 The terms "tolerance", "tolerant" or "less sensitive" are interchangeable used and mean the relative levels of inherent tolerance of the HPPD screened according to a visible indicator phenotype of the strain or plant transformed with a nucleic acid comprising the gene coding for the respective HPPD protein in the presence of different concentrations of the various HPPD inhibitors.. Dose responses and
- 15 relative shifts in dose responses associated with these indicator phenotypes (formation of brown colour, growth inhibition, bleaching, herbicidal effect etc) are conveniently expressed in terms, for example, of GR50 (concentration for 50% reduction of growth) or MIC (minimum inhibitory concentration) values where increases in values correspond to increases in inherent tolerance of the expressed
- 20 HPPD, in the normal manner based upon plant damage, meristematic bleaching symptoms etc. at a range of different concentrations of herbicides. These data can be expressed in terms of, for example, GR50 values derived from dose/response curves having "dose" plotted on the x-axis and "percentage kill", "herbicidal effect", "numbers of emerging green plants" etc. plotted on the y-axis where increased
- 25 GR50 values correspond to increased levels of inherent tolerance of the expressed HPPD. Herbicides can suitably be applied pre-emergence or post emergence. Likewise, tolerance level of the nucleic acid or gene encoding an HPPD protein according to the invention, or the HPPD protein of the invention is screened via transgenesis, regeneration, breeding and spray testing of a test plant such as
- 30 tobacco, or a crop plant such as soybean or cotton and according to these results, such plants are at least 2-4x more tolerant to HPPD inhibitors like tembotrione,

mesotrione, diketonitrile and/or bicyclopyrone, than plants that do not contain any exogenous gene encoding an HPPD protein, or than plants that contain a gene comprising an *Arabidopsis thaliana* HPPD-encoding DNA, under control of the same promoter as the HPPD DNA of the invention.

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"Host organism" or "host" is understood as being any unicellular or multicellular heterologous organism into which the nucleic acid or chimeric gene according to the invention can be introduced for the purpose of producing HPPD according to the invention. These organisms are, in particular, bacteria, for example *E. coli*, yeasts, in particular of the genera *Saccharomyces* or *Kluyveromyces*, *Pichia*, fungi, in particular *Aspergillus*, a baculovirus or, preferably, plant cells and plants.

"Plant cell" is understood, according to the invention, as being any cell which is derived from or found in a plant and which is able to form or is part of undifferentiated tissues, such as calli, differentiated tissues such as embryos, parts of plants, plants or seeds. This includes protoplasts and pollen, cultivated plants cells or protoplasts grown in vitro, and plant cells that can regenerate into a complete plant.

"Plant" is understood, according to the invention, as being any differentiated multicellular organism which is capable of photosynthesis, in particular a monocotyledonous or dicotyledonous organism, more especially cultivated plants which are or are not intended for animal or human nutrition, such as maize or corn, wheat, *Brassica spp.* plants such as *Brassica napus* or *Brassica juncea*, soya spp, rice, sugarcane, beetroot, tobacco, cotton, vegetable plants such as cucumber, leek, carrot, tomato, lettuce, peppers, melon, watermelon, etc. Transgenic plants, as used herein, refer to plants comprising a foreign or heterologous gene stably inserted in their genome.

In one embodiment the invention relates to the transformation of plants. Any promoter sequence of a gene which is expressed naturally in plants, or any hybrid

or combination of promoter elements of genes expressed naturally in plants, including *Agrobacterium* or plant virus promoters, or any promoter which is suitable for controlling the transcription of a herbicide tolerance gene in plants, can be used as the promoter sequence in the plants of the invention (named "plant-expressible promoter" herein). Examples of such suitable plant-expressible promoters are described above. In one embodiment of this invention, such plant-expressible promoters are operably-linked to a coding sequence encoding an HPPD protein of the invention to form a chimeric HPPD gene of this invention.

10 According to the invention, it is also possible to use, in combination with the promoter regulatory sequence, other regulatory sequences which are located between the promoter and the coding sequence, such as intron sequences, or transcription activators (enhancers). Examples of such suitable regulatory sequences are described above.

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Any corresponding sequence of bacterial or viral origin, such as the nos terminator from *Agrobacterium tumefaciens*, or of plant origin, such as a histone terminator as described in application EP 0 633 317 A1, may be used as transcription termination (and polyadenylation) regulatory sequence.

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In one particular embodiment of the invention, a nucleic acid sequence which encodes a transit peptide is employed 5' (upstream) of the nucleic acid sequence encoding the exogenous HPPD according to the invention, with this transit peptide sequence being arranged between the promoter region and the sequence encoding the exogenous HPPD so as to permit expression of a transit peptide-HPPD fusion protein, such as the protein of SEQ ID No. 6 or SEQ ID No. 7. The transit peptide makes it possible to direct the HPPD into the plastids, more especially the chloroplasts, with the fusion protein being cleaved between the transit peptide and the HPPD protein of the invention when the latter enters the plastid. The transit peptide may be a single peptide, such as an EPSPS transit peptide (described in US patent 5,188,642) or a transit peptide of the plant ribulose biphosphate

carboxylase/ oxygenase small subunit (RuBisCO ssu), where appropriate, including a few amino acids of the N-terminal part of the mature RuBisCO ssu (EP 189 707 A1), or else may be a fusion of several transit peptides such as a transit peptide which comprises a first plant transit peptide which is fused to a part of the N-terminal
5 sequence of a mature protein having a plastid location, with this part in turn being fused to a second plant transit peptide as described in patent EP 508 909 A1, and, more especially, the optimized transit peptide which comprises a transit peptide of the sunflower RuBisCO ssu fused to 22 amino acids of the N-terminal end of the maize RuBisCO ssu, in turn fused to the transit peptide of the maize RuBisCO ssu,
10 as described, with its coding sequence, in patent EP 508 909 A1.

The present invention also relates to the transit peptide-HPPD fusion protein and a nucleic acid or plant-expressible chimeric gene encoding such fusion protein, wherein the two elements of this fusion protein are as defined above.

15 The present invention also relates to a cloning, transformation and/or expression vector, which vector contains at least one chimeric gene as defined above. In addition to the above chimeric gene, this vector can contain an origin of replication. This vector can be a plasmid or plasmid portion, a cosmid, or a bacteriophage or a virus which has been transformed by introducing the chimeric gene according to the
20 invention. Transformation vectors are well known to the skilled person and widely described in the literature. The transformation vector which can be used, in particular, for transforming plant cells or plants may be a virus, which can be employed for transforming plant cells or plants and which additionally contains its own replication and expression elements. According to the invention, the vector for
25 transforming plant cells or plants is preferably a plasmid, such as a disarmed *Agrobacterium* Ti plasmid.

The present invention also relates to the host organisms, in particular plant cells, seeds or plants, which comprise a chimeric gene which comprises a sequence
30 encoding an HPPD protein of the invention, such as a protein comprising the amino acid sequence of SEQ ID Nos 4, 5, 6, or 7 as defined above, and the use of the

plants or seeds of the invention in a field to grow a crop and harvest a plant product, e.g., soya spp, rice, wheat, barley or corn grains or cotton bolls, where in one embodiment said use involves the application of an HPPD inhibitor herbicide to such plants to control weeds. In one embodiment of this invention, in such use the HPPD
5 inhibitors are triketones or pyrazolates, preferably tembotrione, mesotrione, topramezone or sulcotrione, bicyclopyrone, pyrasulfotole, pyrazolate, benzofenap and tefuryltrione, particularly tembotrione.

Therefore, the present invention relates to a host organism, in particular a plant cell,
10 seed, or plant, characterized in that it contains at least one HPPD chimeric gene as described above, or at least an HPPD nucleic acid sequence as previously described.

In a particular embodiment, the present invention relates to a plant cell or plant
15 characterized in that it contains at least a nucleic acid sequence which encodes an HPPD protein of this invention which retain its properties of catalysing the conversion of para-hydroxyphenylpyruvate to homogentisate and which makes this plant more tolerant than plants of the same species not comprising such HPPD protein of the present invention, particularly to triketones, or pyrazolates,
20 preferably tembotrione, mesotrione, topramezone or sulcotrione, bicyclopyrone, pyrasulfotole, pyrazolate, benzofenap and tefuryltrione, particularly tembotrione and such plants containing the HPPD of the invention have an agronomically acceptable tolerance to an HPPD inhibitor herbicide particularly to triketones, or pyrazolates, preferably tembotrione, mesotrione, topramezone or sulcotrione, bicyclopyrone,
25 pyrasulfotole, pyrazolate, benzofenap and tefuryltrione, particularly tembotrione.

In another particular embodiment, the present invention relates to a plant cell or plant characterized in that it contains at least a nucleic acid sequence which encodes an HPPD of this invention which retain its properties of catalysing the
30 conversion of para-hydroxyphenylpyruvate to homogentisate and which is less sensitive to an HPPD inhibitor than the host plant endogenous HPPD, such as the

HPPD from *Arabidopsis thaliana*, particularly the HPPD comprising the amino acid sequence of SEQ ID No. 11 (from the amino acid position 126 to the amino acid position 568), or comprising the amino acid sequence of SEQ ID No.11 or SEQ ID No. 12 (from the amino acid position 134 to the amino acid position 575).

5

In a particular embodiment, the present invention relates to a host plant cell, seed or host plant characterized in that it contains at least a nucleic acid sequence which encodes an HPPD of the invention as defined herein, wherein the HPPD of the invention is less sensitive than the host plant endogenous HPPD to an HPPD inhibitor herbicide of the class of isoxazoles, diketonitriles, triketones or pyrazolines more especially from isoxaflutole, tembotrione, mesotrione, sulcotrione, pyrasulfotole, bicyclopyrone, tefuryltrione, topramezone, 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-CF₃phenyl)propane-1,3-dione and 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-2,3 Cl₂ phenyl)propane-1,3-dione, even more particularly tembotrione, mesotrione, diketonitrile, bicyclopyrone, topramezone, pyrazolate, benzofenap, sulcotrione, tefuryltrione, and pyrasulfotole, most particularly tembotrione, mesotrione and bicyclopyrone.

In another particular embodiment, the present invention relates to a plant cell or plant characterized in that it contains at least a nucleic acid sequence encoding an HPPD of the invention as previously described, and in addition a chimeric gene comprising a plant-expressible promoter as described above, operably-linked to a nucleic acid sequence encoding a PDH (prephenate dehydrogenase) enzyme (US 2005/0257283).

25

The present invention also relates to the plants which contain transformed cells, in particular the plants which are regenerated from the transformed cells, and progeny plants or seeds thereof, comprising the chimeric HPPD gene of the invention. The regeneration can be obtained by any appropriate method, with the method depending on the nature of the species, as described, for example, in the above references. The following patents and patent applications may be cited, in particular,

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with regard to the methods for transforming plant cells and regenerating plants:

US 4,459,355, US 4,536,475, US 5,464,763, US 5,177,010, US 5,187,073,
EP 267,159 A1, EP 604 662 A1, EP 672 752 A1, US 4,945,050, US 5,036,006,
US 5,100,792, US 5,371,014, US 5,478,744, US 5,179,022, US 5,565,346,
5 US 5,484,956, US 5,508,468, US 5,538,877, US 5,554,798, US 5,489,520,
US 5,510,318, US 5,204,253, US 5,405,765, EP 442 174 A1, EP 486 233 A1,
EP 486 234 A1, EP 539 563 A1, EP 674 725 A1, WO 91/02071 and WO 95/06128.

The present invention also relates to the transgenic plants or part thereof, which are
10 derived by cultivating and/or crossing the above transgenic plants, and to the seeds
of the transgenic plants, comprising the HPPD chimeric gene of the invention.

The present invention also relates to the end products such as the meal or oil which
are obtained from the plants, part thereof, or seeds of the invention.

15

The transformed plants which can be obtained in accordance with the invention can
be of the monocotyledonous type, such as wheat, barley, sugarcane, rice, onion,
and corn or maize, or of the dicotyledonous type, such as tobacco, soya spp, alfalfa
Brassica spp. plants such as oilseed rape, cotton, sugarbeet clover, vegetables, etc.

20

The invention relates to a method for transforming host organisms, in particular
plant cells or plants, by integrating in such organisms at least one nucleic acid
sequence or one chimeric gene as previously defined, wherein it is possible to
obtain the transformation by any appropriate known means, which means are amply
25 described in the specialist literature and, in particular, the references cited in the
present application, e.g., by using the vector according to the invention.

One transformation method in accordance with this invention comprises bombarding
cells, protoplasts or tissues with solid or liquid particles to which DNA is attached, or
30 containing DNA. Another transformation method comprises using, as mean for
transfer into the plant, a chimeric gene which is inserted into an *Agrobacterium*

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tumefaciens Ti plasmid or an *Agrobacterium rhizogenes* Ri plasmid. Other methods may be used, such as microinjection or electroporation or otherwise direct gene transfer using PEG. The skilled person can select any appropriate method for transforming the host organism of choice, in particular the plant cell or the plant. As examples, the technology for soybean transformation has been extensively described in the examples 1 to 3 disclosed in EP 1186666 A1. For rice, *Agrobacterium*-mediated transformation (Hiei et al., 1994 Plant J 6:271-282, and Hiei et al., 1997 Plant Mol Biol. 35:205-21), electroporation (US 5,641,664 and US 5,679,558), or bombardment (Christou et al., 1991, Biotechnology 9:957) could be performed. A suitable technology for transformation of monocotyledonous plants, and particularly rice, is described in WO 92/09696. For cotton, *Agrobacterium*-mediated transformation (Gould J.H. and Magallanes-Cedeno M., 1998 Plant Molecular Biology reporter, 16:1-10 and Zapata C, 1999, Theoretical Applied Genetics, 98(2):1432-2242), polybrene and/or treatment-mediated transformation (Sawahel W.A., 2001, - Plant Molecular Biology reporter, 19:377a-377f) have been described.

In a particular embodiment of the invention, the HPPD of the invention is targeted into the chloroplast. This may be done by fusing a nucleic acid sequence which encodes a transit peptide to the nucleic acid sequence encoding the HPPD protein of the invention to obtain a nucleic acid encoding a fusion protein as described above.

Alternatively, the HPPD of the invention may be expressed directly in the plastids, such as the chloroplasts, using transformation of the plastid, such as the chloroplast genome. A suitable method comprises the bombardment of plant cells or tissue by solid particles coated with the DNA or liquid particles comprising the DNA, and integration of the introduced gene encoding the protein of the invention by homologous recombination. Suitable vectors and selection systems are known to the person skilled in the art. An example of means and methods which can be used

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for such integration into the chloroplast genome of tobacco plants is given in WO 06/108830.

The present invention also relates to a method for obtaining a plant to an HPPD inhibitor, characterized in that the plant is transformed with a chimeric HPPD gene of the invention as previously described.

Therefore, the present invention also relates to a method for obtaining a plant tolerant to an HPPD inhibitor, characterized in that the plant contains a chimeric HPPD gene of the invention which comprises a coding sequence as well as a heterologous regulatory element in the 5' and optionally in the 3' positions, which are able to function in a host organism, characterized in that the coding sequence comprises at least a nucleic acid sequence defining a gene encoding an HPPD of the invention as previously described.

15

In one embodiment of this invention, the HPPD inhibitor in the above method is a triketone or pyrazolate herbicide, preferably tembotrione, mesotrione, bicyclopyrone, tefuryltrione pyrasulfotole, pyrazolate, diketonitrile, benzofenap, or sulcotrione, particularly tembotrione.

20

According to this invention, a method for obtaining a plant tolerant to an HPPD inhibitor as described above is also provided, characterized in that a plant is obtained comprising a first transgene which is a chimeric HPPD gene of the invention, and a second transgene, which is a chimeric gene comprising a plant-expressible promoter operably-linked to a nucleic acid encoding a PDH (prephenate dehydrogenase) enzyme. A plant comprising such two transgenes can be obtained by transforming a plant with one transgene, and then re-transforming this transgenic plant with the second transgene, or by transforming a plant with the two transgenes simultaneously (in the same or in 2 different transforming DNAs or vectors), or by crossing a plant comprising the first transgene with a plant comprising the second transgene, as is well known in the art.

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The invention also relates to a method for selectively removing weeds or preventing the germination of weeds in a field to be planted with plants or to be sown with seeds, or in a plant crop, by application of an HPPD inhibitor to such field or plant
5 crop, in particular an HPPD inhibitor herbicide as previously defined, which method is characterized in that this HPPD inhibitor herbicide is applied to plants which have been transformed in accordance with the invention, either before sowing the crop (hereinafter named pre-planting application), before emergence of the crop (hereinafter named pre-emergence application), or after emergence of the crop
10 (hereinafter named post-emergence application).

The invention also relates to a method for controlling in an area or a field which contains transformed seeds as previously described in the present invention, which method comprises applying, to the said area of the field, a dose of an HPPD
15 inhibitor herbicide which is toxic for the said weeds, without significantly affecting the seeds or plants which contain the HPPD nucleic acid or the chimeric HPPD gene of the invention as previously described in the present invention.

The present invention also relates to a method for cultivating the plants which have
20 been transformed with a chimeric gene according to the invention, which method comprises planting seeds comprising a chimeric gene of the invention, in an area of a field which is appropriate for cultivating the said plants, and in applying, if weeds are present, a dose, which is toxic for the weeds, of a herbicide whose target is the above-defined HPPD to the said area of the said field, without significantly affecting
25 the said transformed seeds or the said transformed plants, and in then harvesting the cultivated plants or plant parts when they reach the desired stage of maturity and, where appropriate, in separating the seeds from the harvested plants.

In the above methods, the herbicide whose target is the HPPD enzyme can be
30 applied in accordance with the invention, either before sowing the crop, before the crop emerges or after the crop emerges.

The present invention also relates to a process for obtaining oil, particularly soya spp, corn, or cotton oil, or meal, comprising growing a crop, particularly a soya spp crop, expressing an HPPD protein of the invention optionally treating such crop with
5 an HPPD inhibitor herbicide, harvesting the grains and milling the grains to make meal and extract the oil. Also the seeds or grains, either whole, broken or crushed, comprising the chimeric gene of the invention are part of this invention.

Therefore, the present invention relates to a method for obtaining oil or meal
10 comprising growing a transformed plant as described above, optionally treating such plant with an HPPD inhibitor herbicide, harvesting the grains and milling the grains to make meal and extract the oil.

Further provided in this invention, are the above methods involving an HPPD
15 inhibitor herbicide selected from isoxaflutole, tembotrione, mesotrione, pyrasulfotole, sulcotrione, bicyclopyrone, tefuryltrione, topramezone, 2-cyano-3-cyclopropyl-1-(2-methylsulphonyl-4-trifluoromethylphenyl)-propane-1,3-dione and to 2-cyano-1-[4-(methylsulphonyl)-2-trifluoromethylphenyl]-3-(1-methylcyclopropyl)propane-1,3-dione.

20

Also provided herein are the above methods of the invention involving an HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone,
25 tefuryltrione and mesotrione, more particularly tembotrione.

Within the meaning of the present invention, "herbicide" is understood as being a herbicidally active substance on its own or such a substance which is combined with an additive which alters its efficacy, such as, for example, an agent which increases
30 its activity (a synergistic agent) or which limits its activity (a safener). It is of course to be understood that, for their application in practice, the above herbicides are

combined, in a manner which is known per se, with the formulation adjuvants which are customarily employed in agricultural chemistry.

HPPD inhibitor herbicides like those of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione, have an outstanding herbicidal activity against a broad spectrum of economically important monocotyledonous and dicotyledonous annual harmful plants. The active substances also act efficiently on perennial harmful plants which produce shoots from rhizomes, wood stocks or other perennial organs and which are difficult to control.

The present invention therefore also relates to a method of controlling undesired plants or for regulating the growth of plants in crops of plants comprising an HPPD according to the invention, where one or more HPPD inhibitor herbicides of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione are applied to the plants (for example harmful plants such as monocotyledonous or dicotyledonous weeds or undesired crop plants), to the seeds (for example grains, seeds or vegetative propagules such as tubers or shoot parts with buds) or to the area on which the plants grow (for example the area under cultivation). In this context, an HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione can be applied for example pre-planting (if appropriate also by incorporation into the soil), pre-emergence or post-emergence. Examples of individual representatives of the monocotyledonous and dicotyledonous weeds which can be controlled with an HPPD inhibitor herbicide of the class of triketones,

such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolinates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione are hereby mentioned, without this mentioning being
5 intended as a limitation to certain species only:

- Monocotyledonous harmful plants of the genera: Aegilops, Agropyron, Agrostis, Alopecurus, Apera, Avena, Brachiaria, Bromus, Cenchrus, Commelina, Cynodon, Cyperus, Dactyloctenium, Digitaria, Echinochloa,
10 Eleocharis, Eleusine, Eragrostis, Eriochloa, Festuca, Fimbristylis, Heteranthera, Imperata, Ischaemum, Leptochloa, Lolium, Monochoria, Panicum, Paspalum, Phalaris, Phleum, Poa, Rottboellia, Sagittaria, Scirpus, Setaria, Sorghum.
- 15 Dicotyledonous weeds of the genera: Abutilon, Amaranthus, Ambrosia, Anoda, Anthemis, Aphanes, Artemisia, Atriplex, Bellis, Bidens, Capsella, Carduus, Cassia, Centaurea, Chenopodium, Cirsium, Convolvulus, Datura, Desmodium, Emex, Erysimum, Euphorbia, Galeopsis, Galinsoga, Galium, Hibiscus, Ipomoea, Kochia, Lamium, Lepidium, Lindernia, Matricaria, Mentha,
20 Mercurialis, Mullugo, Myosotis, Papaver, Pharbitis, Plantago, Polygonum, Portulaca, Ranunculus, Raphanus, Rorippa, Rotala, Rumex, Salsola, Senecio, Sesbania, Sida, Sinapis, Solanum, Sonchus, Sphenoclea, Stellaria, Taraxacum, Thlaspi, Trifolium, Urtica, Veronica, Viola, Xanthium.
- 25 In transgenic crops according to the invention, comprising an HPPD protein, DNA or chimeric gene according invention and which may also show one more further herbicide resistances against herbicides that differ from HPPD inhibitor herbicides, the use of HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolinates, such as pyrasulfotole
30 and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione in

economically important transgenic crops of useful plants and ornamentals, for example of cereals such as wheat, barley, rye, oats, sorghum and millet, rice and maize or else crops of sugar beet, cotton, soya spp, oil seed rape, potato, tomato, peas and other vegetables is preferred.

5

As it relates to plant properties other than the tolerance to HPPD inhibitor herbicides as described in the present invention, conventional ways of generating novel plants which, in comparison with existing plants, have modified properties are, for example, traditional breeding methods and the generation of mutants. Alternatively, novel

10 plants with modified properties can be generated with the aid of recombinant methods (see, for example, EP-A-0221044 A1, EP-A-0131624 A1). For example, the following have been described in several cases:

- recombinant modifications of crop plants for the purposes of modifying the starch synthesized in the plants (for example WO 92/11376, WO 92/14827,
15 WO 91/19806),
- transgenic crop plants which are resistant to certain herbicides of the glufosinate type (cf., for example, EP-A-0242236, EP-A-242246) or of the glyphosate type (WO 92/00377) or of the sulfonylurea type (EP-A-0257993, US-A-5013659),
- 20 - transgenic crop plants, for example corn, cotton or soya spp, which are capable of producing *Bacillus thuringiensis* toxins (Bt toxins), or hybrids or mutants thereof, which make the plants resistant to certain pests (EP-A-0193259),
- transgenic crop plants with a modified fatty acid composition (WO 91/13972),
- 25 - genetically modified crop plants with novel constituents or secondary metabolites, for example novel phytoalexins, which bring about an increased disease resistance (EPA 309862, EPA0464461),
- genetically modified plants with reduced photorespiration which feature higher yields and higher stress tolerance (EPA 0305398),
- 30 - transgenic crop plants which produce pharmaceutically or diagnostically important proteins ("molecular pharming"),

- transgenic crop plants which are distinguished by higher yields or better quality,
 - transgenic crop plants which are distinguished by a combination of novel properties such as a combination of the abovementioned novel properties
- 5 ("gene stacking").

A large number of molecular-biological techniques by means of which novel transgenic plants with modified properties can be generated are known in principle; see, for example, I. Potrykus and G. Spangenberg (eds.) Gene Transfer to Plants,
10 Springer Lab Manual (1995), Springer Verlag Berlin, Heidelberg, or Christou, "Trends in Plant Science" 1 (1996) 423-431).

To carry out such recombinant manipulations, it is possible to introduce nucleic acid molecules into plasmids, which permit a mutagenesis or sequence modification by
15 recombination of DNA sequences. For example, base substitutions can be carried out, part-sequences can be removed, or natural or synthetic sequences may be added with the aid of standard methods. To link the DNA fragments with one another, it is possible to add adapters or linkers to the fragments; see, for example, Sambrook et al., 1989, Molecular Cloning, A Laboratory Manual, 2. ed., Cold Spring
20 Harbor Laboratory Press, Cold Spring Harbor, NY; or Winnacker "Gene und Klon", VCH Weinheim 2. ed., 1996

The generation of plant cells with a reduced activity for a gene product can be achieved for example by the expression of at least one corresponding antisense
25 RNA, a sense RNA for achieving a cosuppression effect, or a combination of both an antisense and sense RNA forming a double-stranded silencing RNA molecule (RNAi), or by the expression of at least one correspondingly constructed ribozyme, which specifically cleaves transcripts of the abovementioned gene product. To do this, it is possible firstly to use DNA molecules which comprise all of the coding
30 sequence of a gene product, including any flanking sequences which may be present, or else DNA molecules which only comprise parts of the coding sequence,

it being necessary for these parts to be long enough to bring about an antisense effect in the cells. It is also possible to use DNA sequences which have a high degree of homology with the coding sequences of a gene product, but which are not entirely identical.

5

When expressing nucleic acid molecules in plants, the obtained protein may be localized in any compartment of the plant cell. In order to achieve localization in a particular compartment, however, it is possible for example to link the coding region to DNA sequences which ensure the localization in a specific compartment. Such sequences are known to the skilled person (see, for example, Braun et al., EMBO J. 11 (1992), 3219-3227; Wolter et al., Proc. Natl. Acad. Sci. USA 85 (1988), 846-850; Sonnewald et al., Plant J. 1 (1991), 95-106). However, the nucleic acid molecules can also be expressed in the organelles of the plant cells.

15 The transgenic plant cells can be regenerated by known techniques to give intact plants. In principle, the transgenic plants may be plants of any plant species, including monocotyledonous or dicotyledonous plants.

Thus, transgenic plants can be obtained which - in addition to the chimeric HPPD gene of the invention - have modified properties as the result of overexpression, suppression or inhibition of homologous (= natural) genes or gene sequences or expression of heterologous (= foreign) genes or gene sequences.

On the plants, plant cells or seeds of the invention, it is preferred to employ the HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolinates, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione in transgenic crops which are also resistant to growth regulators such as, for example, 2,4-D or dicamba, or against herbicides which inhibit essential plant enzymes, for example acetolactate synthases (ALS), EPSP synthases, or glutamine synthases

30

(GS), or against herbicides from the group of the sulfonylureas, glyphosate, or glufosinate and analogous active substances.

The invention therefore also relates to the use of herbicides applied to this HPPD
5 tolerant plants according to the invention for controlling harmful plants (i.e. weeds)
which also extends to transgenic crop plants comprising a second or more herbicide
resistance(s) beside the resistance against HPPD inhibitor herbicide of the class of
triketones, such as tembotrione, sulcotrione and mesotrione, of the class of
isoxazoles such as isoxaflutole or of the class of pyrazolates, such as
10 pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione,
topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly
tembotrione.

HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione
15 and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and
topramezone, particularly selected from tembotrione, sulcotrione, topramezone,
bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione can be
employed in the customary preparations in the form of wettable powders,
emulsifiable concentrates, sprayable solutions, dusts or granules.

20 HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione
and mesotrione, or of the class of pyrazolates, such as pyrasulfotole and
topramezone, particularly selected from tembotrione, sulcotrione, topramezone,
bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione can be
25 formulated in various ways, depending on the prevailing biological and/or physico-
chemical parameters. Examples of possible formulations are: wettable powders
(WP), water-soluble powders (SP), water-soluble concentrates, emulsifiable
concentrates (EC), emulsions (EW), such as oil-in-water and water-in-oil emulsions,
sprayable solutions, suspension concentrates (SC), oil- or water-based dispersions,
30 oil-miscible solutions, capsule suspensions (CS), dusts (DP), seed-dressing
products, granules for application by broadcasting and on the soil, granules (GR) in

the form of microgranules, spray granules, coated granules and adsorption granules, water-dispersible granules (WG), water-soluble granules (SG), ULV formulations, microcapsules and waxes.

- 5 These individual types of formulation are known in principle and are described, for example, in: Winnacker-Küchler, "Chemische Technologie" [Chemical technology], volume 7, C. Hanser Verlag Munich, 4th Ed. 1986; Wade van Valkenburg, "Pesticide Formulations", Marcel Dekker, N.Y., 1973; K. Martens, "Spray Drying" Handbook, 3rd Ed. 1979, G. Goodwin Ltd. London.

10

- The formulation auxiliaries required, such as inert materials, surfactants, solvents and further additives, are also known and are described, for example, in: Watkins, "Handbook of Insecticide Dust Diluents and Carriers", 2nd Ed., Darland Books, Caldwell N.J., H.v. Olphen, "Introduction to Clay Colloid Chemistry"; 2nd Ed., J. Wiley & Sons, N.Y.; C. Marsden, "Solvents Guide"; 2nd Ed., Interscience, N.Y. 1963; McCutcheon's "Detergents and Emulsifiers Annual", MC Publ. Corp., Ridgewood N.J.; Sisley and Wood, "Encyclopedia of Surface Active Agents", Chem. Publ. Co. Inc., N.Y. 1964; Schönfeldt, "Grenzflächenaktive Äthylenoxidaddukte" [Interface-active ethylene oxide adducts], Wiss. Verlagsgesell., Stuttgart 1976;
- 15
- 20 Winnacker-Küchler, "Chemische Technologie" [Chemical technology], volume 7, C. Hanser Verlag Munich, 4th Ed. 1986.

- Based on these formulations, it is also possible to prepare combinations with other pesticidally active substances such as, for example, insecticides, acaricides,
- 25 herbicides, fungicides, and with safeners, fertilizers and/or growth regulators, for example in the form of a ready mix or a tank mix.

- Wettable powders are preparations which are uniformly dispersible in water and which, besides the active substance, also comprise ionic and/or nonionic surfactants (wetters, dispersers), for example polyoxyethylated alkylphenols, polyoxyethylated
- 30 fatty alcohols, polyoxyethylated fatty amines, fatty alcohol polyglycol ether sulfates,

alkanesulfonates, alkylbenzenesulfonates, sodium lignosulfonate, sodium 2,2'-dinaphthylmethane-6,6'-disulfonate, sodium dibutyl-naphthalenesulfonate or else sodium oleoylmethyltaurate, besides a diluent or inert substance. To prepare the wettable powders, the herbicidally active substances are ground finely, for example
5 in customary apparatuses such as hammer mills, blower mills and air-jet mills, and mixed with the formulation auxiliaries, either simultaneously or subsequently.

Emulsifiable concentrates are prepared by dissolving the active substance in an organic solvent, for example butanol, cyclohexanone, dimethylformamide, xylene or
10 else higher-boiling aromatics or hydrocarbons or mixtures of the organic solvents with addition of one or more ionic and/or nonionic surfactants (emulsifiers).
Examples of emulsifiers which may be used are: calcium alkylarylsulfonates such as calcium dodecylbenzenesulfonate, or nonionic emulsifiers such as fatty acid polyglycol esters, alkylaryl polyglycol ethers, fatty alcohol polyglycol ethers,
15 propylene oxide/ethylene oxide condensates, alkyl polyethers, sorbitan esters such as, for example, sorbitan fatty acid esters or polyoxyethylene sorbitan esters such as, for example, polyoxyethylene sorbitan fatty acid esters.

Dusts are obtained by grinding the active substance with finely divided solid
20 materials such as, for example, talcum, natural clays such as kaolin, bentonite and pyrophyllite, or diatomaceous earth.

Suspension concentrates can be water- or oil-based. They can be prepared for example by wet-grinding by means of commercially available bead mills, if
25 appropriate with addition of surfactants as already listed above for example in the case of the other formulation types.

Emulsions, for example oil-in-water emulsions (EW), can be prepared for example by means of stirrers, colloid mills and/or static mixers using aqueous organic
30 solvents and, if appropriate, surfactants, as have already been mentioned for example above for the other formulation types.

Granules can be prepared either by spraying the active substance onto adsorptive, granulated inert material, or by applying active substance concentrates to the surface of carriers such as sand, kaolinites or granulated inert material with the aid of stickers, for example polyvinyl alcohol, sodium polyacrylate or else mineral oils. Suitable active substances can also be granulated in the manner which is customary for the production of fertilizer granules, if desired as a mixture with fertilizers.

Water-dispersible granules are generally prepared by customary methods such as spray drying, fluidized-bed granulation, disk granulation, mixing with high-speed stirrers, and extrusion without solid inert material.

To prepare disk granules, fluidized-bed granules, extruder granules and spray granules, see, for example, methods in "Spray-Drying Handbook" 3rd ed. 1979, G. Goodwin Ltd., London; J.E. Browning, "Agglomeration", Chemical and Engineering 1967, pages 147 et seq.; "Perry's Chemical Engineer's Handbook", 5th Ed., McGraw-Hill, New York 1973, p. 8-57.

For further details of the formulation of crop protection products see, for example, G.C. Klingman, "Weed Control as a Science", John Wiley and Sons, Inc., New York, 1961, pages 81-96 and J.D. Freyer, S.A. Evans, "Weed Control Handbook", 5th Ed., Blackwell Scientific Publications, Oxford, 1968, pages 101-103.

As a rule, the agrochemical preparations comprise from 0.1 to 99% by weight, in particular from 0.1 to 95% by weight, of compounds according to the invention. In wettable powders, the active substance concentration is, for example, approximately 10 to 90% by weight, the remainder to 100% by weight being composed of customary formulation constituents. In the case of emulsifiable concentrates, the active substance concentration can amount to approximately 1 to 90, preferably 5 to 80% by weight. Formulations in the form of dusts comprise from 1 to 30% by weight of active substance, preferably in most cases from 5 to 20% by

weight of active substance, and sprayable solutions comprise approximately from 0.05 to 80, preferably from 2 to 50% by weight of active substance. In the case of water-dispersible granules, the active substance content depends partly on whether the active compound is in liquid or solid form, and on the granulation auxiliaries, fillers and the like which are being used. In the case of the water-dispersible granules, for example, the active substance content is between 1 and 95% by weight, preferably between 10 and 80% by weight.

In addition, the active substance formulations mentioned comprise, if appropriate, the auxiliaries which are conventional in each case, such as stickers, wetters, dispersants, emulsifiers, penetrations, preservatives, antifreeze agents, solvents, fillers, carriers, colorants, antifoams, evaporation inhibitors, and pH and viscosity regulators.

Based on these formulations, it is also possible to prepare combinations of an HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolines, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione with other pesticidally active substances such as, for example, insecticides, acaricides, herbicides, fungicides, and with safeners, fertilizers and/or growth regulators, for example in the form of a ready mix or a tank mix to be applied to HPPD tolerant plants according to the invention.

Active substances which can be applied to HPPD tolerant plants according to the present invention in combination with HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolines, such as pyrasulfotole and topramezone, particularly selected from tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione in mixed formulations or in the tank mix are, for example, known active substances which are based on the inhibition of, for

example, acetolactate synthase, acetyl-CoA carboxylase, cellulose synthase, enolpyruvylshikimate-3-phosphate synthase, glutamine synthetase, p-hydroxyphenylpyruvate dioxygenase, phytoene desaturase, photosystem I, photosystem II, protoporphyrinogen oxidase, as are described in, for example,

5 Weed Research 26 (1986) 441-445 or "The Pesticide Manual", 14th edition, The British Crop Protection Council and the Royal Soc. of Chemistry, 2003 and the literature cited therein. Known herbicides or plant growth regulators which can be combined with the compounds according to the invention are, for example, the following active substances (the compounds are either designated by the common

10 name according to the International Organization for Standardization (ISO) or by a chemical name, if appropriate together with the code number) and always comprise all use forms such as acids, salts, esters and isomers such as stereoisomers and optical isomers. In this context, one and in some cases also several use forms are mentioned by way of example:

15 acetochlor, acibenzolar, acibenzolar-S-methyl, acifluorfen, acifluorfen-sodium, aclonifen, alachlor, allidochlor, alloxydim, alloxydim-sodium, ametryne, amicarbazone, amidochlor, amidosulfuron, aminocyclopyrachlor, aminopyralid, amitrole, ammonium sulfamate, ancymidol, anilofos, asulam, atrazine, azafenidin,

20 azimsulfuron, aziprotryne, BAH-043, BAS-140H, BAS-693H, BAS-714H, BAS-762H, BAS-776H, BAS-800H, beflubutamid, benazolin, benazolin-ethyl, bencarbazon, benfluralin, benfuresate, bensulide, bensulfuron-methyl, bentazone, benzfendizone, benzobicyclon, benzofenap, benzofluor, benzoylprop, bifenox, bilanafos, bilanafos-sodium, bispyribac, bispyribac-sodium, bromacil, bromobutide, bromofenoxim,

25 bromoxynil, bromuron, buminafos, busoxinone, butachlor, butafenacil, butamifos, butenachlor, butralin, butroxydim, butylate, cafenstrole, carbetamide, carfentrazone, carfentrazone-ethyl, chlomethoxyfen, chloramben, chlorazifop, chlorazifop-butyl, chlorbromuron, chlorbufam, chlorfenac, chlorfenac-sodium, chlorfenprop, chlorflurenol, chlorflurenol-methyl, chloridazon, chlorimuron, chlorimuron-ethyl,

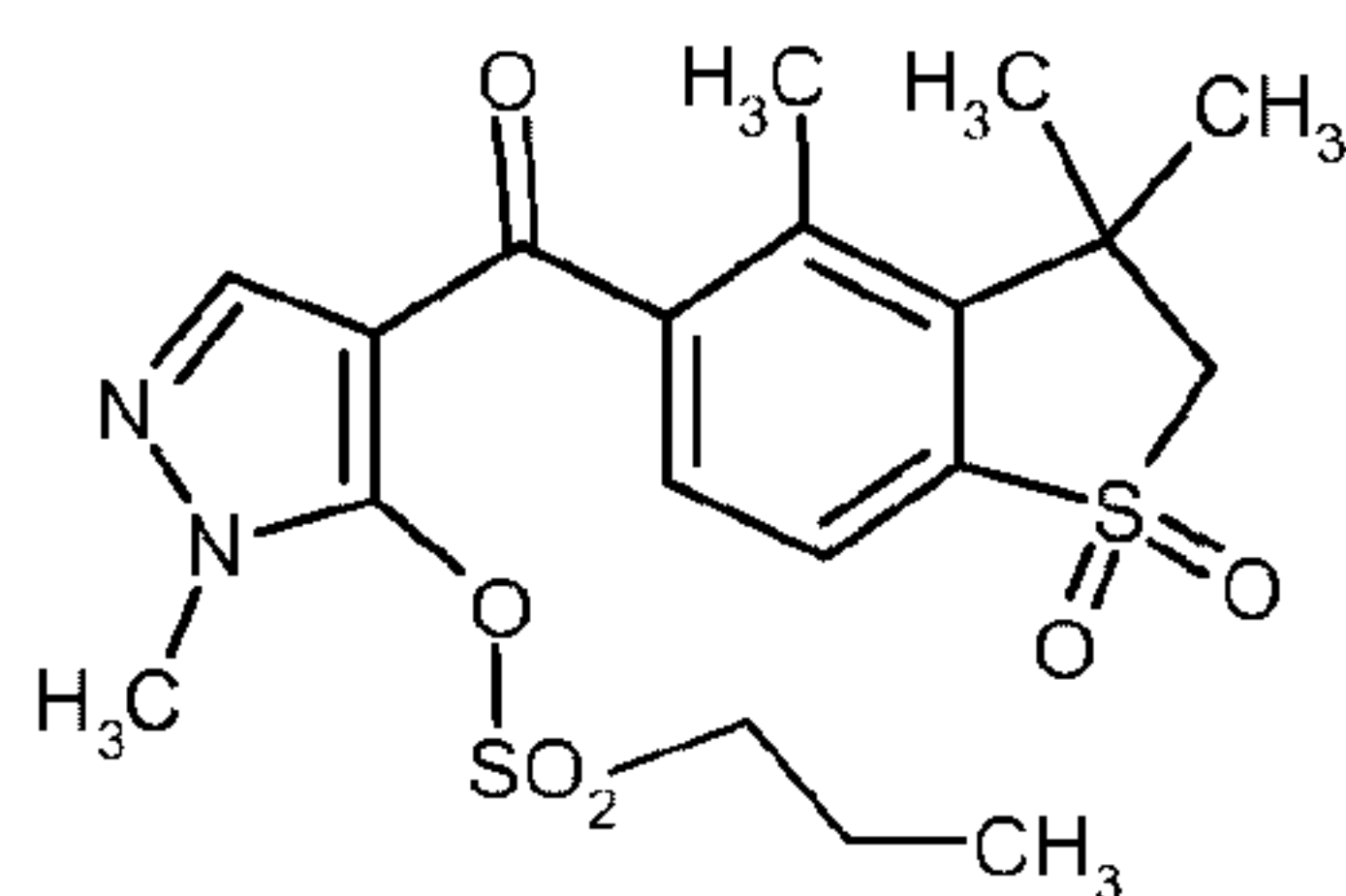
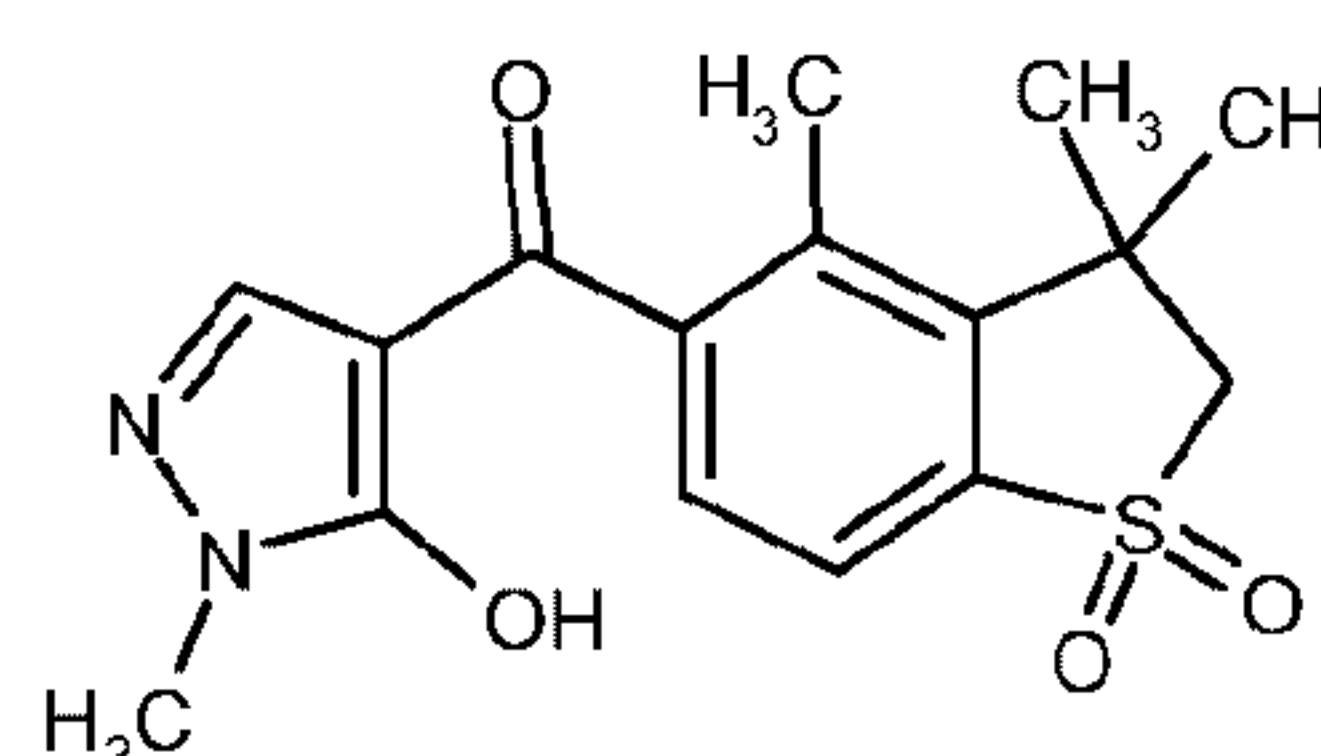
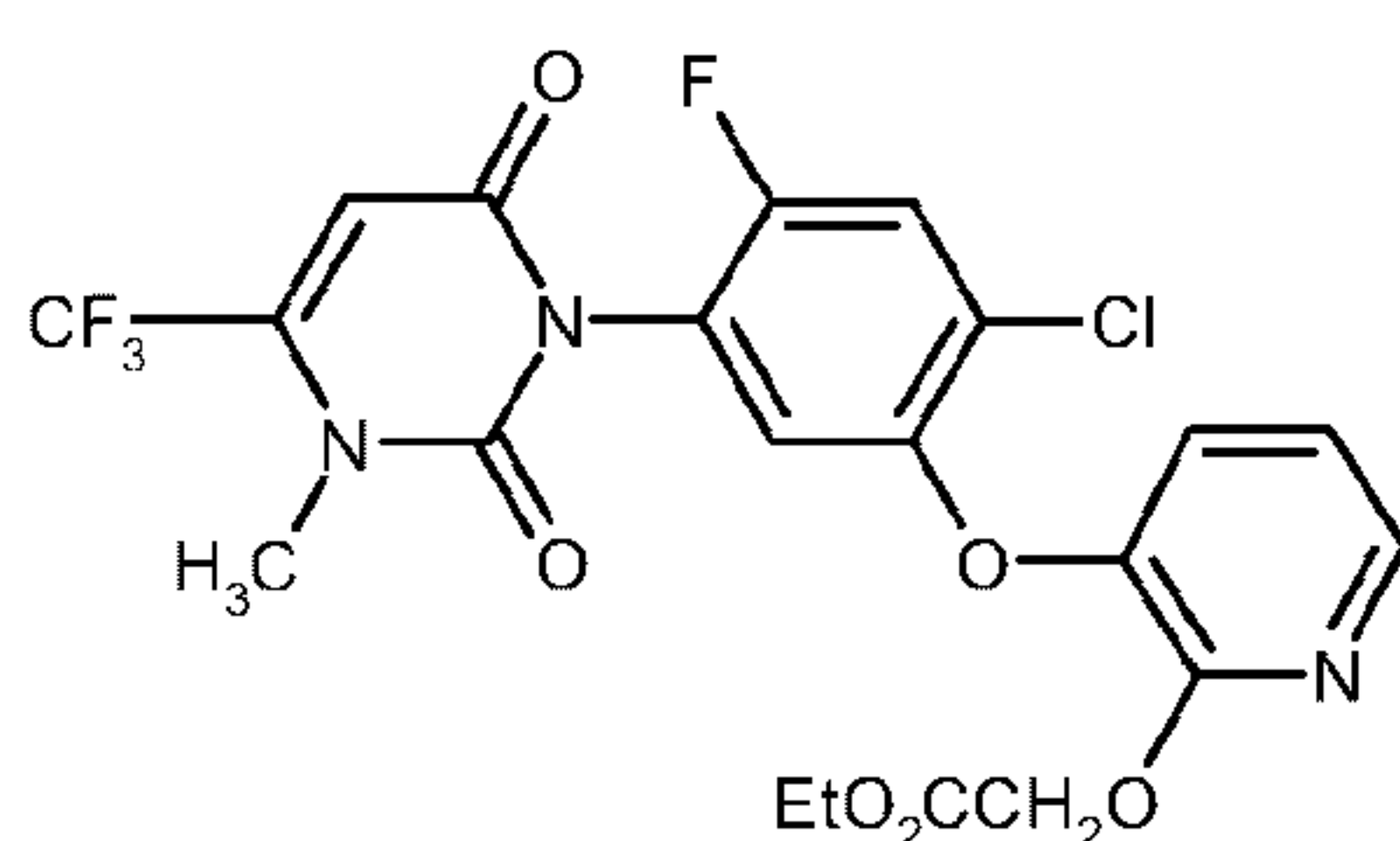
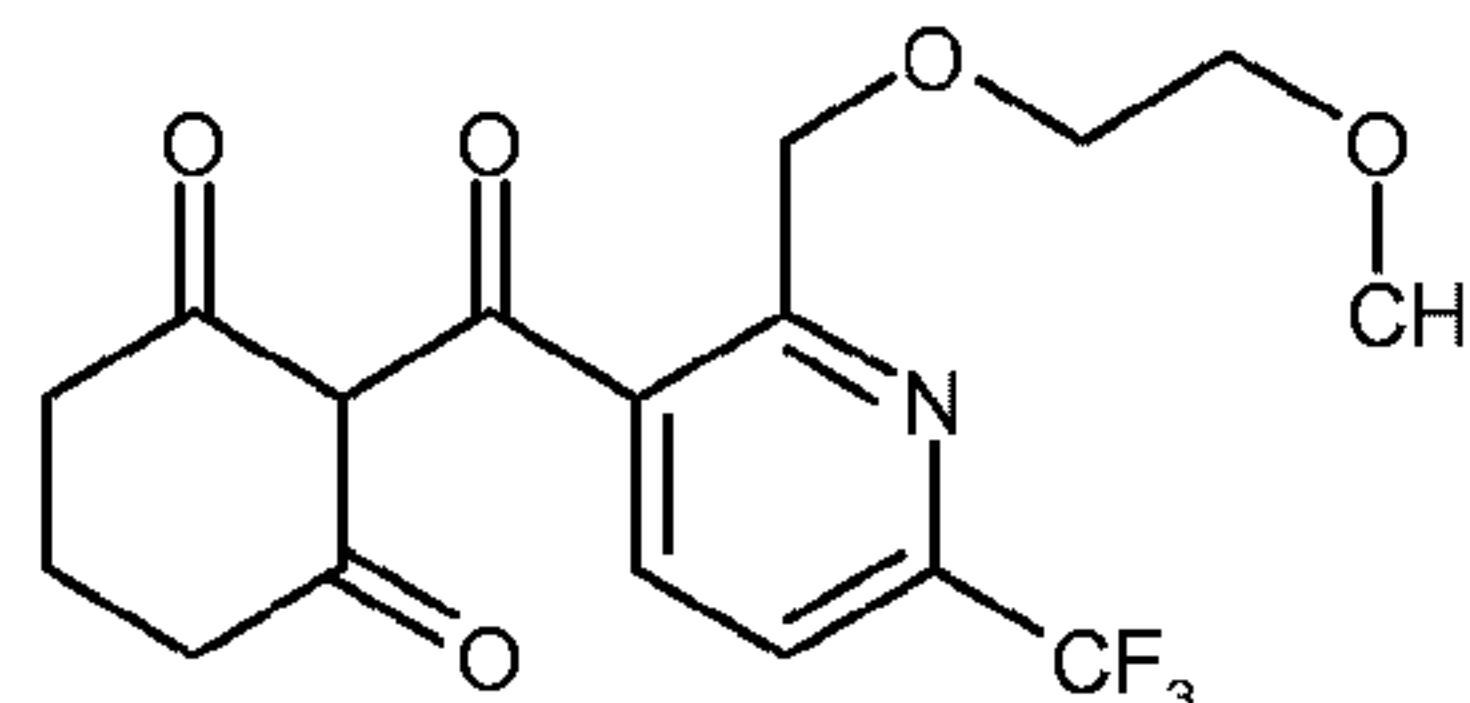
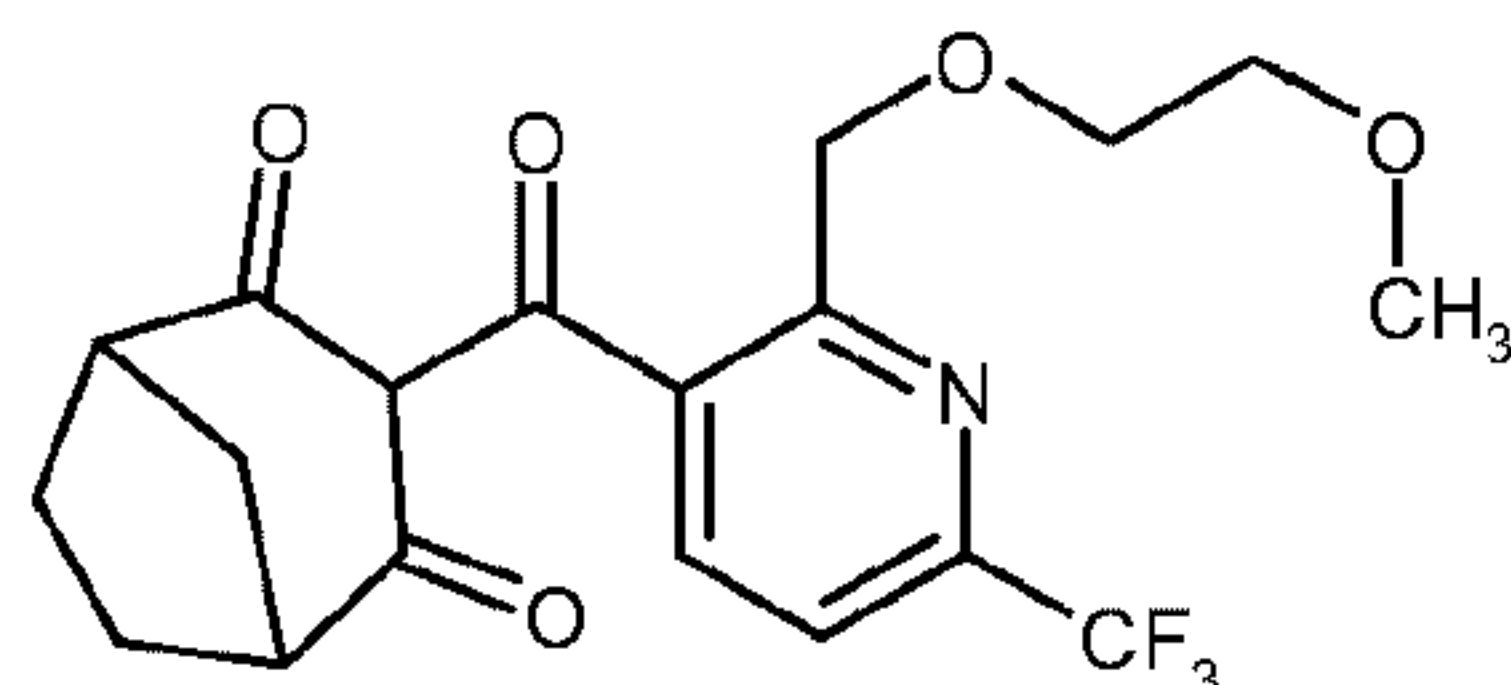
30 chlormequat-chloride, chlornitrofen, chlorophthalim, chlorthal-dimethyl, chlorotoluron, chlorsulfuron, cinidon, cinidon-ethyl, cinmethylin, cinosulfuron,

clethodim, clodinafop clodinafop-propargyl, clofencet, clomazone, clomeprop,
 cloprop, clopyralid, cloransulam, cloransulam-methyl, cumyluron, cyanamide,
 cyanazine, cyclanilide, cycloate, cyclosulfamuron, cycloxydim, cycluron, cyhalofop,
 cyhalofop-butyl, cyperquat, cyprazine, cyprazole, 2,4-D, 2,4-DB, daimuron/dymron,
 5 dalapon, daminozide, dazomet, n-decanol, desmedipham, desmetryn, detosyl-
 pyrazolate (DTP), di-allate, dicamba, dichlobenil, dichlorprop, dichlorprop-P,
 diclofop, diclofop-methyl, diclofop-P-methyl, diclosulam, diethatyl, diethatyl-ethyl,
 difenoxuron, difenzoquat, diflufenican, diflufenzopyr, diflufenzopyr-sodium,
 dimefuron, dikegulac-sodium, dimefuron, dimepiperate, dimethachlor,
 10 dimethametryn, dimethenamid, dimethenamid-P, dimethipin, dimetrasulfuron,
 dinitramine, dinoseb, dinoterb, diphenamid, dipropetryn, diquat, diquat-dibromide,
 dithiopyr, diuron, DNOC, eglinazine-ethyl, endothal, EPTC, esprocarb, ethalfluralin,
 ethametsulfuron-methyl, ethephon, ethidimuron, ethiozin, ethofumesate, ethoxyfen,
 ethoxyfen-ethyl, ethoxysulfuron, etobenzanid, F-5331, i.e. N-[2-chloro-4-fluoro-5-[4-
 15 (3-fluoro-propyl)-4,5-dihydro-5-oxo-1H-tetrazol-1-yl]-phenyl]ethanesulfonamide,
 fenoprop, fenoxaprop, fenoxaprop-P, fenoxaprop-ethyl, fenoxaprop-P-ethyl,
 fentrazamide, fenuron, flamprop, flamprop-M-isopropyl, flamprop-M-methyl,
 flazasulfuron, florasulam, fluazifop, fluazifop-P, fluazifop-butyl, fluazifop-P-butyl,
 fluazolate, flucarbazone, flucarbazone-sodium, flucetosulfuron, fluchloralin,
 20 flufenacet (thiaflumamide), flufenpyr, flufenpyr-ethyl, flumetralin, flumetsulam,
 flumiclorac, flumiclorac-pentyl, flumioxazin, flumipropyn, fluometuron, fluorodifen,
 fluoroglycofen, fluoroglycofen-ethyl, flupoxam, flupropacil, flupropanate,
 flupyrsulfuron, flupyrsulfuron-methyl-sodium, flurenol, flurenol-butyl, fluridone,
 flurochloridone, fluroxypyr, fluroxypyr-meptyl, flurprimidol, flurtamone, fluthiacet,
 25 fluthiacet-methyl, fluthiamide, fomesafen, foramsulfuron, forchlorfenuron, fosamine,
 furyloxyfen, gibberellic acid, glufosinate, L-glufosinate, L-glufosinate-ammonium,
 glufosinate-ammonium, glyphosate, glyphosate-isopropylammonium, H-9201,
 halosafen, halosulfuron, halosulfuron-methyl, haloxyfop, haloxyfop-P, haloxyfop-
 ethoxyethyl, haloxyfop-P-ethoxyethyl, haloxyfop-methyl, haloxyfop-P-methyl,
 30 hexazinone, HNPC-9908, HOK-201, HW-02, imazamethabenz, imazamethabenz-
 methyl, imazamox, imazapic, imazapyr, imazaquin, imazethapyr, imazosulfuron,

inabenfide, indanofan, indoleacetic acid (IAA), 4-indol-3-ylbutyric acid (IBA),
 iodosulfuron, iodosulfuron-methyl-sodium, ioxynil, isocarbamid, isopropalin,
 isoproturon, isouron, isoxaben, isoxachlortole, isoxaflutole, isoxapyrifop, KUH-043,
 KUH-071, karbutilate, ketospiradox, lactofen, lenacil, linuron, maleic hydrazide,
 5 MCPA, MCPB, MCPB-methyl, -ethyl and -sodium, mecoprop, mecoprop-sodium,
 mecoprop-butotyl, mecoprop-P-butotyl, mecoprop-P-dimethylammonium, mecoprop-
 P-2-ethylhexyl, mecoprop-P-potassium, mefenacet, mefluidide, mepiquat-chloride,
 mesosulfuron, mesosulfuron-methyl, methabenzthiazuron, metam, metamifop,
 metamitron, metazachlor, methazole, methoxyphenone, methyldymron,
 10 1-methylcyclopropene, methyl isothiocyanate, metobenzuron, metobenzuron,
 metobromuron, metolachlor, S-metolachlor, metosulam, metoxuron, metribuzin,
 metsulfuron, metsulfuron-methyl, molinate, monalide, monocarbamide,
 monocarbamide dihydrogen sulfate, monolinuron, monosulfuron, monuron, MT 128,
 MT-5950, i.e. N-[3-chloro-4-(1-methylethyl)-phenyl]-2-methylpentanamide, NGGC-
 15 011, naproanilide, napropamide, naptalam, NC-310, i.e. 4-(2,4-dichlorobenzoyl)-
 1-methyl-5-benzyloxypyrazole, neburon, nicosulfuron, nipyraclufen, nitralin, nitrofen,
 nitrophenolat-sodium (isomer mixture), nitrofluorfen, nonanoic acid, norflurazon,
 orbencarb, orthosulfamuron, oryzalin, oxadiargyl, oxadiazon, oxasulfuron,
 oxaziclomefone, oxyfluorfen, paclobutrazole, paraquat, paraquat dichloride,
 20 pelargonic acid (nonanoic acid), pendimethalin, pendralin, penoxsulam,
 pentanochlor, pentoxazone, perfluidone, pethoxamid, phenisopham,
 phenmedipham, phenmedipham-ethyl, picloram, picolinafen, pinoxaden, piperophos,
 pirifenop, pirifenop-butyl, pretilachlor, primisulfuron, primisulfuron-methyl,
 probenazole, profluazol, procyazine, prodiamine, prifluraline, profoxydim,
 25 prohexadione, prohexadione-calcium, prohydrojasmon, prometon, prometryn,
 propachlor, propanil, propaquizafop, propazine, propham, propisochlor,
 propoxycarbazone, propoxycarbazone-sodium, propyzamide, prosulfalin,
 prosulfocarb, prosulfuron, prynachlor, pyraclonil, pyraflufen, pyraflufen-ethyl,
 pyrazolynate (pyrazolate), pyrazosulfuron-ethyl, pyrazoxyfen, pyribambenz,
 30 pyribambenz-isopropyl, pyribenzoxim, pyributicarb, pyridafol, pyridate, pyriftalid,
 pyriminobac, pyriminobac-methyl, pyrimisulfan, pyriothiobac, pyriothiobac-sodium,

pyroxasulfone, pyroxsulam, quinclorac, quinmerac, quinclamine, quizalofop, quizalofop-ethyl, quizalofop-P, quizalofop-P-ethyl, quizalofop-P-tefuryl, rimsulfuron, saflufenacil, secbumeton, sethoxydim, siduron, simazine, simetryn, SN-106279, sulf-
 5 allate (CDEC), sulfentrazone, sulfometuron, sulfometuron-methyl, sulfosate (glyphosate-trimesium), sulfosulfuron, SYN-523, SYP-249, SYP-298, SYP-300, tebutam, tebuthiuron, tecnazene, tepraloxymid, terbacil, terbucarb, terbuchlor, terbumeton, terbuthylazine, terbutryne, TH-547, thenylchlor, thiafluamide, thiazafluron, thiazopyr, thidiazimin, thidiazuron, thiencarbazone, thiencarbazone-methyl, thifensulfuron, thifensulfuron-methyl, thiobencarb, tiocarbazil, tralkoxydim,
 10 tri-allate, triasulfuron, triaziflam, triazofenamide, tribenuron, tribenuron-methyl, trichloroacetic acid (TCA), triclopyr, tridiphane, trietazine, trifloxysulfuron, trifloxysulfuron-sodium, trifluralin, triflusulfuron, triflusulfuron-methyl, trimeturon, trinexapac, trinexapac-ethyl, tritosulfuron, tsitodef, uniconazole, uniconazole-P, vernolate, ZJ-0166, ZJ-0270, ZJ-0543, ZJ-0862 and the following compounds

15

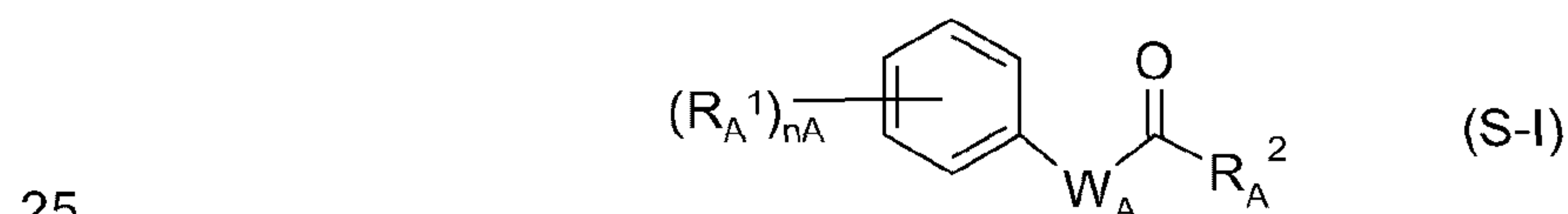


The application rate required of the HPPD inhibitor herbicide of the class of triketones, such as tembotrione, sulcotrione and mesotrione, or of the class of pyrazolines, such as pyrasulfotole and topramezone, particularly selected from
 5 tembotrione, sulcotrione, topramezone, bicyclopyrone, tefuryltrione and mesotrione, more particularly tembotrione to be applied to areas where HPPD tolerant plants according to the present invention are growing varies as a function of the external conditions such as temperature, humidity, the nature of the herbicide used and the like. It can vary within wide limits, for example between 0.001 and 1.0 kg/ha and
 10 more of active substance, but it is preferably between 0.005 and 750 g/ha.

In case of combined applications of HPPD inhibitor herbicides with herbicides that differ from HPPD inhibitor herbicides to the HPPD tolerant plants according to the present invention, these mixtures may cause crop injury, based on the presence of
 15 the non HPPD inhibitor herbicides. In order to reduce/eliminate such crop injuries, appropriate safeners may be added. These safeners, which are employed in antidotically active amounts, reduce the phytotoxic side effects of herbicides/pesticides used, for example in economically important crops, such as cereals (wheat, barley, rye, corn, rice, millet), alfalfa, sugar beet, sugarcane, oilseed
 20 rape, cotton and soya spp., preferably corn, cotton, sugarbeet, or soya spp.

The safeners are preferably selected from the group consisting of:

A) compounds of the formula (S-I)

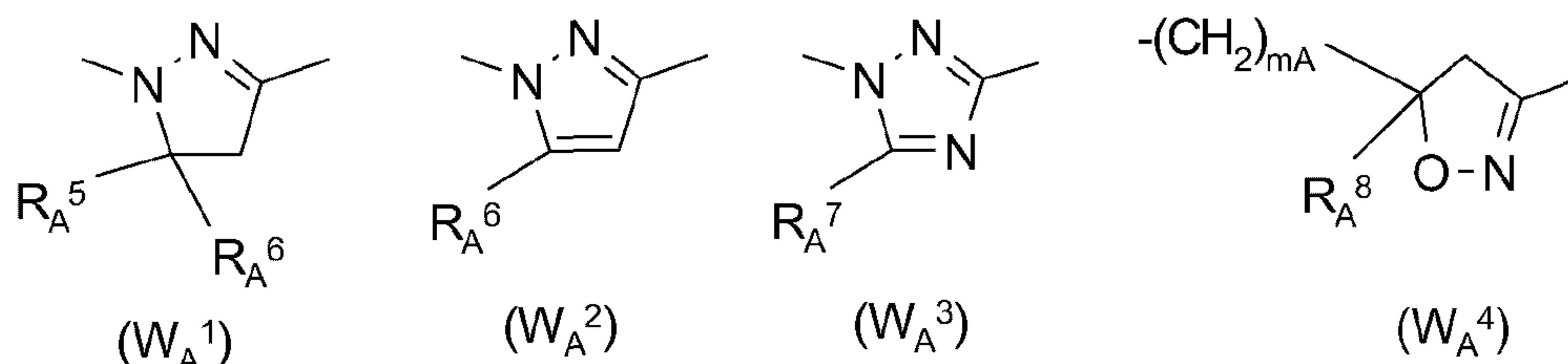


where the symbols and indices have the following meanings:

n_A is a natural number from 0 to 5, preferably from 0 to 3;

R_A^1 is halogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, nitro or (C₁-C₄)-haloalkyl;

W_A is an unsubstituted or substituted divalent heterocyclic radical from the group consisting of partially unsaturated or aromatic five-membered heterocycles having 1 to 3 hetero ring atoms of the type N or O, where at least one nitrogen atom and at most one oxygen atom is present in the ring, preferably a radical from the group consisting of (W_A^1) to (W_A^4),



m_A is 0 or 1;

R_A^2 is OR_A^3 , SR_A^3 or $NR_A^3R_A^4$ or a saturated or unsaturated 3- to 7-membered heterocycle having at least one nitrogen atom and up to 3 heteroatoms, preferably from the group consisting of O and S, which is attached via the nitrogen atom to the carbonyl group in (S-I) and which is unsubstituted or substituted by radicals from the group consisting of (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy and optionally substituted phenyl, preferably a radical of the formula OR_A^3 , NHR_A^4 or $N(CH_3)_2$, in particular of the formula OR_A^3 ;

R_A^3 is hydrogen or an unsubstituted or substituted aliphatic hydrocarbon radical having preferably a total of 1 to 18 carbon atoms;

R_A^4 is hydrogen, (C₁-C₆)-alkyl, (C₁-C₆)-alkoxy or substituted or unsubstituted phenyl;

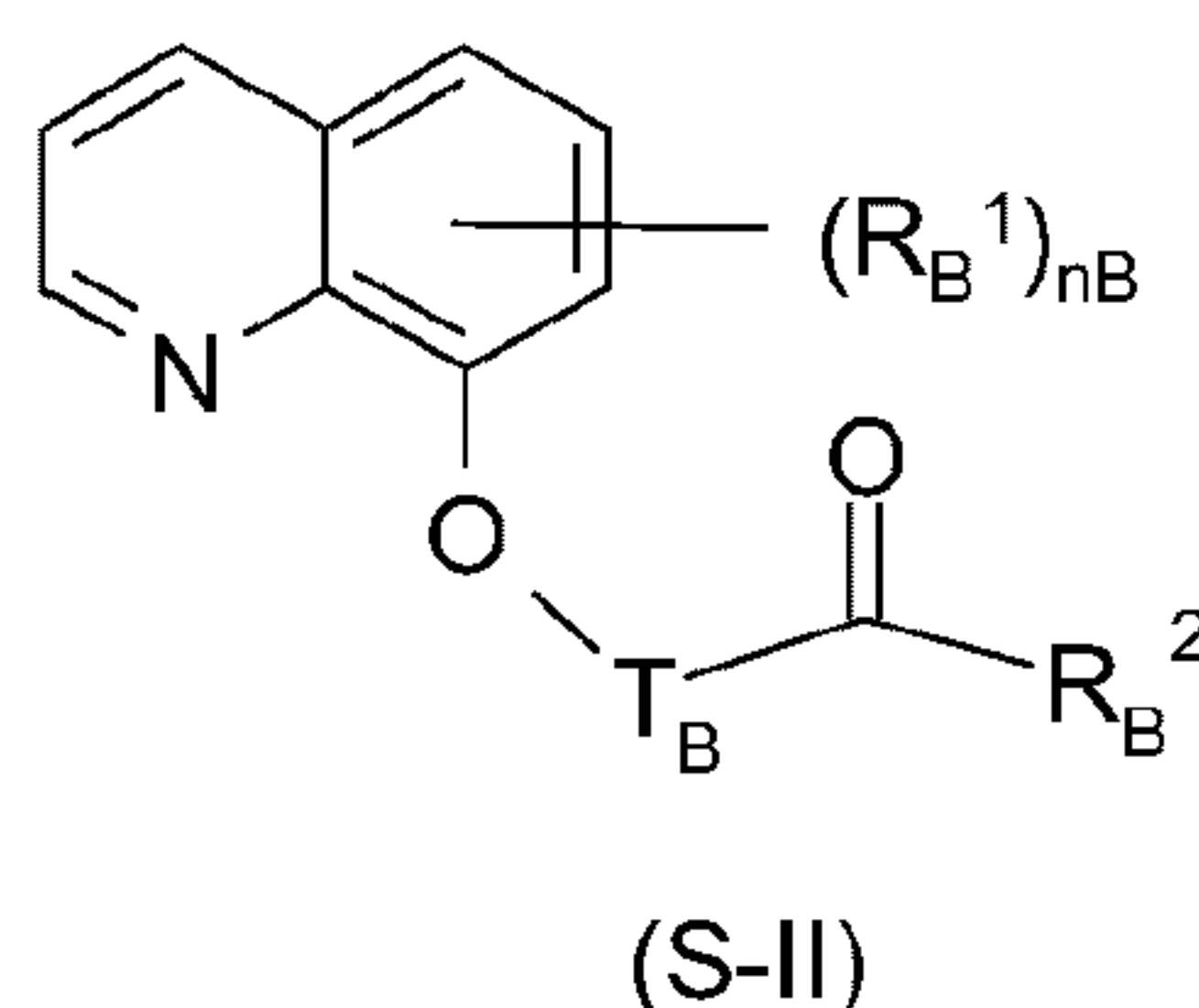
R_A^5 is H, (C₁-C₈)-alkyl, (C₁-C₈)-haloalkyl, (C₁-C₄)-alkoxy-(C₁-C₈)-alkyl, cyano or $COOR_A^9$ where R_A^9 is hydrogen, (C₁-C₈)-alkyl, (C₁-C₈)-haloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₁-C₆)-hydroxyalkyl, (C₃-C₁₂)-cycloalkyl or tri-(C₁-C₄)-alkylsilyl;

R_A^6 , R_A^7 , R_A^8 are identical or different and are hydrogen, (C₁-C₈)-alkyl, (C₁-C₈)-haloalkyl, (C₃-C₁₂)-cycloalkyl or substituted or unsubstituted phenyl;

preferably:

- a) compounds of the type of the dichlorophenylpyrazoline-3-carboxylic acid, preferably compounds such as ethyl 1-(2,4-dichlorophenyl)-5-(ethoxycarbonyl)-5-methyl-2-pyrazoline-3-carboxylate (S1-1) ("mefenpyr-diethyl", see Pestic. Man.), and related compounds, as described in WO 91/07874;
- 5 b) derivatives of dichlorophenylpyrazolecarboxylic acid, preferably compounds such as ethyl 1-(2,4-dichlorophenyl)-5-methylpyrazole-3-carboxylate (S1-2), ethyl 1-(2,4-dichlorophenyl)-5-isopropylpyrazole-3-carboxylate (S1-3), ethyl 1-(2,4-dichlorophenyl)-5-(1,1-dimethylethyl)pyrazole-3-carboxylate (S1-4), ethyl 1-(2,4-dichlorophenyl)-5-phenylpyrazole-3-carboxylate (S1-5) and related
- 10 compounds, as described in EP-A-333 131 and EP-A-269 806;
- c) compounds of the type of the triazolecarboxylic acids, preferably compounds such as fenchlorazole(-ethyl ester), i.e. ethyl 1-(2,4-dichlorophenyl)-5-trichloromethyl-(1H)-1,2,4-triazole-3-carboxylate (S1-6), and related compounds, as described in EP-A-174 562 and EP-A-346 620;
- 15 d) compounds of the type of the 5-benzyl- or 5-phenyl-2-isoxazoline-3-carboxylic acid or the 5,5-diphenyl-2-isoxazoline-3-carboxylic acid, preferably compounds such as ethyl 5-(2,4-dichlorobenzyl)-2-isoxazoline-3-carboxylate (S1-7) or ethyl 5-phenyl-2-isoxazoline-3-carboxylate (S1-8) and related compounds, as described
- 20 in WO 91/08202, or ethyl 5,5-diphenyl-2-isoxazolinecarboxylate (S1-9) ("isoxadifen-ethyl") or n-propyl 5,5-diphenyl-2-isoxazolinecarboxylate (S1-10) or ethyl 5-(4-fluorophenyl)-5-phenyl-2-isoxazoline-3-carboxylate (S1-11), as described in the patent application WO-A-95/07897.

- 25 B) Quinoline derivatives of the formula (S-II)



where the symbols and indices have the following meanings:

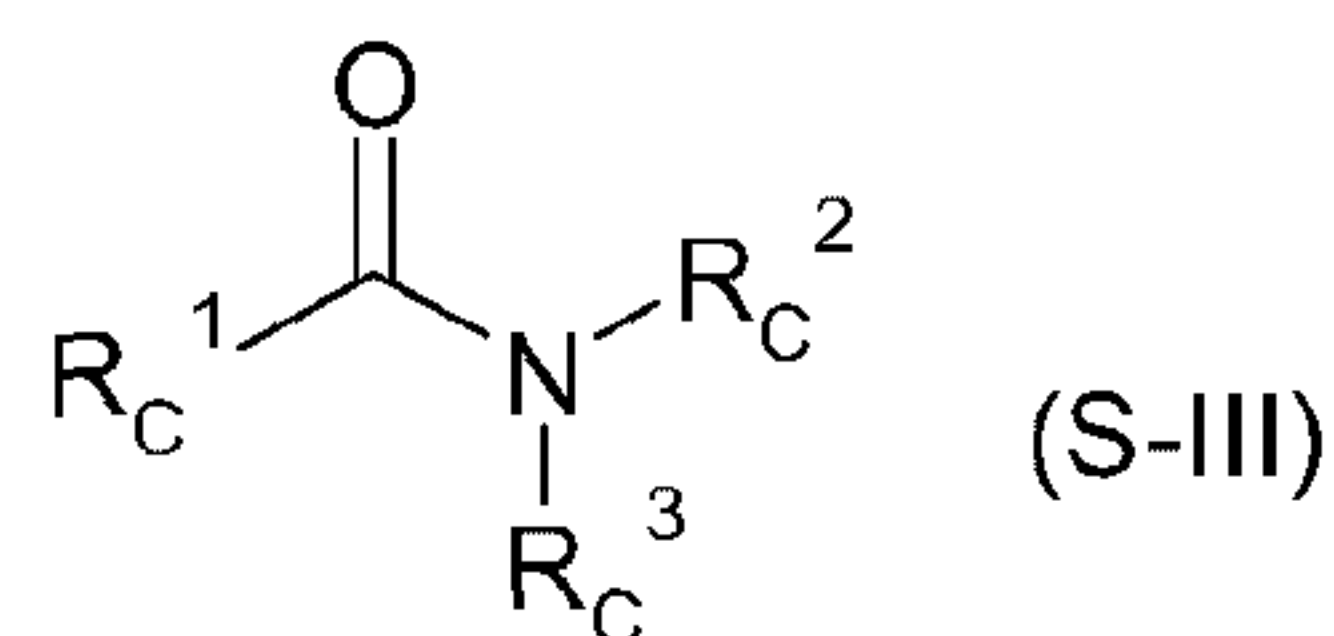
- R_B^1 is halogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, nitro or (C₁-C₄)-haloalkyl;
 n_B is a natural number from 0 to 5, preferably from 0 to 3;
 R_B^2 OR_B^3 , SR_B^3 or $NR_B^3R_B^4$ or a saturated
 5 or unsaturated 3- to 7-membered heterocycle having at least one nitrogen atom and up to 3 heteroatoms, preferably from the group consisting of O and S, which is attached via the nitrogen atom to the carbonyl group in (S-II) and is unsubstituted or substituted by radicals from the group consisting of (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy or optionally substituted phenyl, preferably a radical of the formula OR_B^3 , NHR_B^4 or
 10 $N(CH_3)_2$, in particular of the formula OR_B^3 ;
 R_B^3 is hydrogen or an unsubstituted or substituted aliphatic hydrocarbon radical having preferably a total of 1 to 18 carbon atoms;
 R_B^4 is hydrogen, (C₁-C₆)-alkyl, (C₁-C₆)-alkoxy or substituted or unsubstituted phenyl;
 15 T_B is a (C₁- or C₂)-alkanediyl chain which is unsubstituted or substituted by one or two (C₁-C₄)-alkyl radicals or by [(C₁-C₃)-alkoxy]carbonyl;

preferably:

- a) compounds of the type of the 8-quinolinoxyacetic acid (S2), preferably
 20 1-methylhexyl (5-chloro-8-quinolinoxy)acetate (common name "cloquintocet-mexyl" (S2-1) (see Pestic. Man.),
 1,3-dimethylbut-1-yl (5-chloro-8-quinolinoxy)acetate (S2-2),
 4-allyloxybutyl (5-chloro-8-quinolinoxy)acetate (S2-3),
 1-allyloxyprop-2-yl (5-chloro-8-quinolinoxy)acetate- (S2-4),
 25 ethyl (5-chloro-8-quinolinoxy)acetate (S2-5),
 methyl (5-chloro-8-quinolinoxy)acetate (S2-6),
 allyl (5-chloro-8-quinolinoxy)acetate (S2-7),
 2-(2-propylideneiminoxy)-1-ethyl (5-chloro-8-quinolinoxy)acetate (S2-8), 2-oxoprop-
 1-yl (5-chloro-8-quinolinoxy)acetate (S2-9) and related compounds, as described in
 30 EP-A-86 750, EP-A-94 349 and EP-A-191 736 or EP-A-0 492 366, and also their hydrates and salts, as described in WO-A-2002/034048.

- b) Compounds of the type of the (5-chloro-8-quinolinoxy)malonic acid, preferably compounds such as diethyl (5-chloro-8-quinolinoxy)malonate, diallyl (5-chloro-8-quinolinoxy)malonate, methyl ethyl (5-chloro-8-quinolinoxy)malonate and
5 related compounds, as described in EP-A-0 582 198.

C) Compounds of the formula (S-III)



where the symbols and indices have the following meanings:

- 10 R_C^1 is (C₁-C₄)-alkyl, (C₁-C₄)-haloalkyl, (C₂-C₄)-alkenyl, (C₂-C₄)-haloalkenyl, (C₃-C₇)-cycloalkyl, preferably dichloromethyl;
- R_C^2 , R_C^3 are identical or different and are hydrogen, (C₁-C₄)-alkyl, (C₂-C₄)-alkenyl, (C₂-C₄)-alkynyl, (C₁-C₄)-haloalkyl, (C₂-C₄)-haloalkenyl, (C₁-C₄)-alkylcarbamoyl- (C₁-C₄)-alkyl, (C₂-C₄)-alkenylcarbamoyl-(C₁-C₄)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl,
15 dioxolanyl-(C₁-C₄)-alkyl, thiazolyl, furyl, furylalkyl, thienyl, piperidyl, substituted or unsubstituted phenyl, or R_C^2 and R_C^3 together form a substituted or unsubstituted heterocyclic ring,
preferably an oxazolidine, thiazolidine, piperidine, morpholine, hexahydropyrimidine or benzoxazine ring;
- 20 preferably:
- active compounds of the type of the dichloroacetamides which are frequently used as pre-emergence safener (soil-acting safeners), such as, for example,
"dichlormid" (see Pestic.Man.) (= N,N-diallyl-2,2-dichloroacetamide),
25 "R-29148" (= 3-dichloroacetyl-2,2,5-trimethyl-1,3-oxazolidine from Stauffer),
"R-28725" (= 3-dichloroacetyl-2,2,-dimethyl-1,3-oxazolidine from Stauffer),
"benoxacor" (see Pestic. Man.) (= 4-dichloroacetyl-3,4-dihydro-3-methyl-2H-1,4-benzoxazine),
"PPG-1292" (= N-allyl-N-[(1,3-dioxolan-2-yl)methyl]dichloroacetamide from PPG

Industries),

"DKA-24" (= N-allyl-N-[(allylaminocarbonyl)methyl]dichloroacetamide from Sagro-Chem),

"AD-67" or "MON 4660" (= 3-dichloroacetyl-1-oxa-3-aza-spiro[4,5]decane from

5 Nitrokemia or Monsanto),

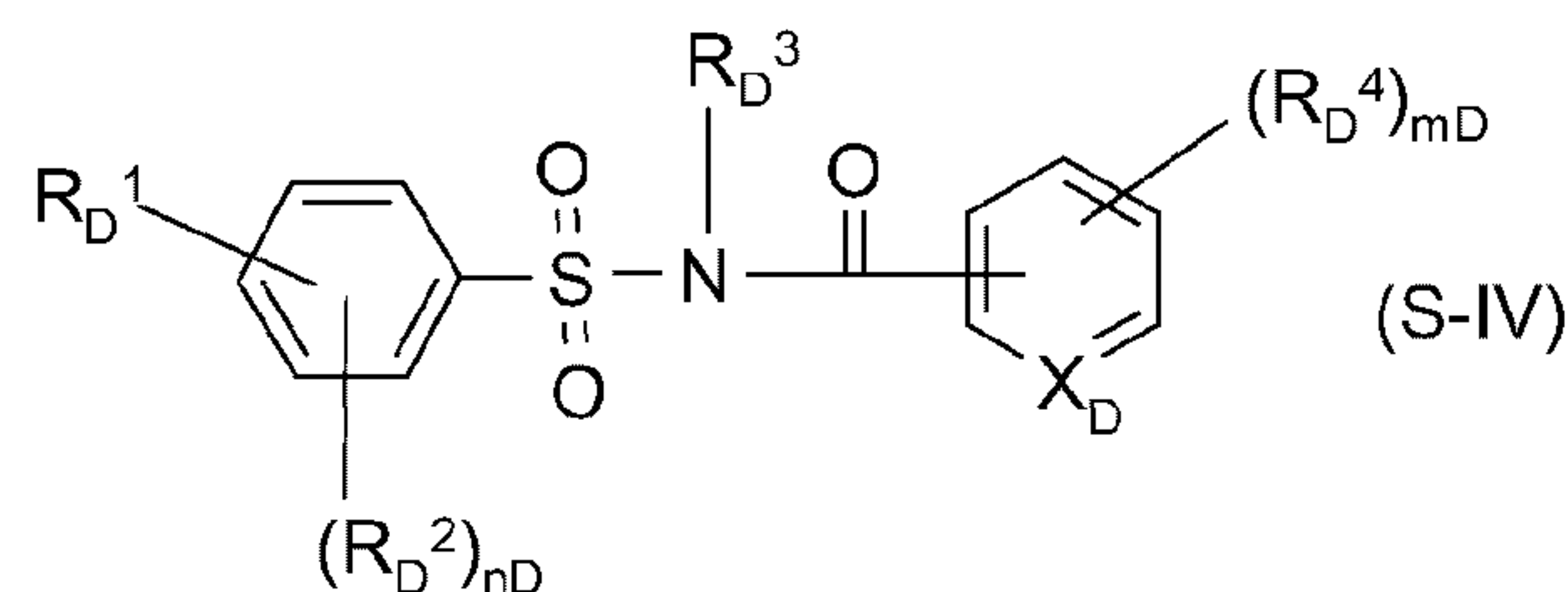
"TI-35" (= 1-dichloroacetylazepane from TRI-Chemical RT)

"diclonon" (dicyclonone) or "BAS145138" or "LAB145138" (= 3-dichloroacetyl-2,5,5-trimethyl-1,3-diazabicyclo[4.3.0]nonane from BASF) and

"furilazole" or "MON 13900" (see Pestic. Man.) (= (RS)-3-dichloroacetyl-5-(2-furyl)-

10 2,2-dimethyloxazolidine).

D) N-Acylsulfonamides of the formula (S-IV) and their salts



15 in which

X_D is CH or N;

R_D^1 is $CO-NR_D^5R_D^6$ or $NHCO-R_D^7$;

R_D^2 is halogen, (C₁-C₄)-haloalkyl, (C₁-C₄)-haloalkoxy, nitro, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, (C₁-C₄)-alkylsulfonyl, (C₁-C₄)-alkoxycarbonyl or (C₁-C₄)-alkylcarbonyl;

20 R_D^3 is hydrogen, (C₁-C₄)-alkyl, (C₂-C₄)-alkenyl or (C₂-C₄)-alkynyl;

R_D^4 is halogen, nitro, (C₁-C₄)-alkyl, (C₁-C₄)-haloalkyl, (C₁-C₄)-haloalkoxy, (C₃-C₆)-cycloalkyl, phenyl, (C₁-C₄)-alkoxy, cyano, (C₁-C₄)-alkylthio, (C₁-C₄)-alkylsulfinyl, (C₁-C₄)-alkylsulfonyl, (C₁-C₄)-alkoxycarbonyl or (C₁-C₄)-alkylcarbonyl;

R_D^5 is hydrogen, (C₁-C₆)-alkyl, (C₃-C₆)-cycloalkyl, (C₂-C₆)-alkenyl, (C₂-C₆)-alkynyl,

25 (C₅-C₆)-cycloalkenyl, phenyl or 3- to 6-membered heterocyclyl containing v_D

heteroatoms from the group consisting of nitrogen, oxygen and sulfur, where the

seven last-mentioned radicals are substituted by v_D substituents from the group

consisting of halogen, (C₁-C₆)-alkoxy, (C₁-C₆)-haloalkoxy, (C₁-C₂)-alkylsulfinyl, (C₁-C₂)-alkylsulfonyl, (C₃-C₆)-cycloalkyl, (C₁-C₄)-alkoxycarbonyl, (C₁-C₄)-alkylcarbonyl and phenyl and, in the case of cyclic radicals, also (C₁-C₄)-alkyl and (C₁-C₄)-haloalkyl;

5

R_D^6 is hydrogen, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl or (C₂-C₆)-alkynyl, where the three last-mentioned radicals are substituted by v_D radicals from the group consisting of halogen, hydroxy, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy and (C₁-C₄)-alkylthio, or

10 R_D^5 and R_D^6 together with the nitrogen atom carrying them form a pyrrolidinyll or piperidinyll radical;

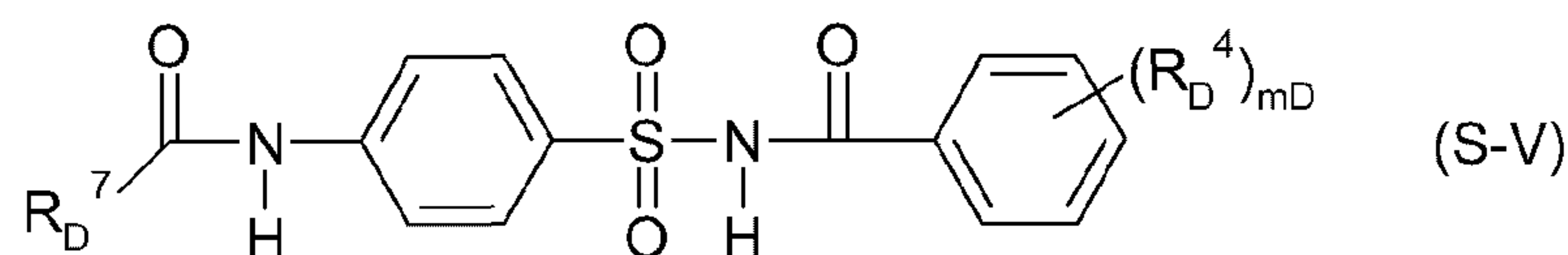
R_D^7 is hydrogen, (C₁-C₄)-alkylamino, di-(C₁-C₄)-alkylamino, (C₁-C₆)-alkyl, (C₃-C₆)-cycloalkyl, where the 2 last-mentioned radicals are substituted by v_D substituents
 15 from the group consisting of halogen, (C₁-C₄)-alkoxy, halogen-(C₁-C₆)-alkoxy and (C₁-C₄)-alkylthio and, in the case of cyclic radicals, also (C₁-C₄)-alkyl and (C₁-C₄)-haloalkyl;

n_D is 0, 1 or 2;

20 m_D is 1 or 2;

v_D is 0, 1, 2 or 3;

from among these, preference is given to compounds of the type of the N-acylsulfonamides, for example of the formula (S-V) below, which are known, for
 25 example, from WO 97/45016



in which

R_D^7 is (C₁-C₆)-alkyl, (C₃-C₆)-cycloalkyl, where the 2 last-mentioned radicals are substituted by v_D substituents from the group consisting of halogen, (C₁-C₄)-alkoxy,

65

halogen-(C₁-C₆)-alkoxy and (C₁-C₄)-alkylthio and, in the case of cyclic radicals, also (C₁-C₄)-alkyl and (C₁-C₄)-haloalkyl;

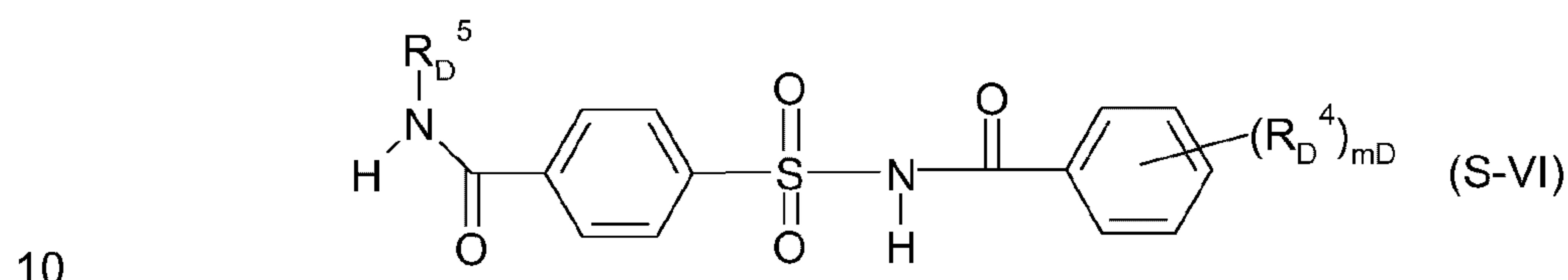
R_D⁴ is halogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, CF₃;

m_D is 1 or 2;

5 v_D is 0, 1, 2 or 3;

and also

acylsulfamoylbenzamides, for example of the formula (S-VI) below, which are known, for example, from WO 99/16744,



for example those in which

R_D⁵ = cyclopropyl and (R_D⁴) = 2-OMe ("cyprosulfamide", S3-1),

R_D⁵ = cyclopropyl and (R_D⁴) = 5-Cl-2-OMe (S3-2),

R_D⁵ = ethyl and (R_D⁴) = 2-OMe (S3-3),

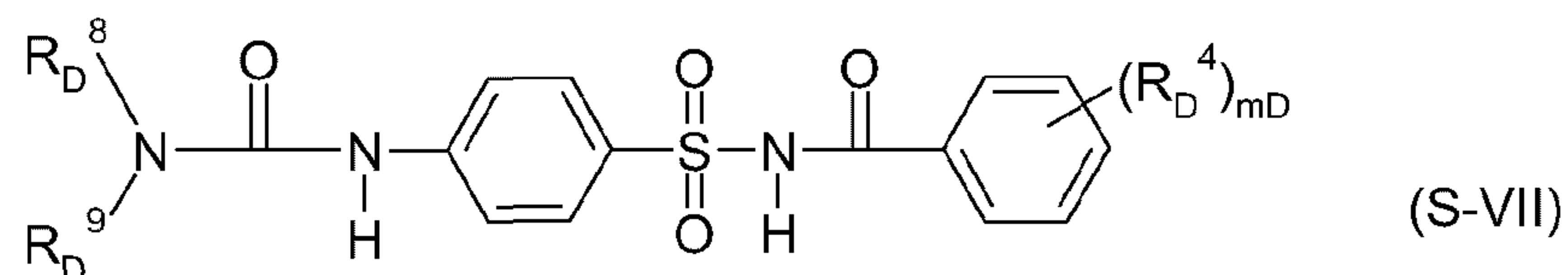
15 R_D⁵ = isopropyl and (R_D⁴) = 5-Cl-2-OMe (S3-4) and

R_D⁵ = isopropyl and (R_D⁴) = 2-OMe (S3-5);

and also

compounds of the type of the N-acylsulfamoylphenylureas of the formula (S-VII),

20 which are known, for example, from EP-A-365484



in which

R_D⁸ and R_D⁹ independently of one another are hydrogen, (C₁-C₈)-alkyl, (C₃-C₈)-cycloalkyl, (C₃-C₆)-alkenyl, (C₃-C₆)-alkynyl,

25 R_D⁴ is halogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, CF₃

m_D is 1 or 2;

from among these in particular

1-[4-(N-2-methoxybenzoylsulfamoyl)phenyl]-3-methylurea,

1-[4-(N-2-methoxybenzoylsulfamoyl)phenyl]-3,3-dimethylurea,

5 1-[4-(N-4,5-dimethylbenzoylsulfamoyl)phenyl]-3-methylurea,

1-[4-(N-naphthoylsulfamoyl)phenyl]-3,3-dimethylurea,

G) active compounds from the class of the hydroxyaromatics and aromatic-aliphatic carboxylic acid derivatives, for example

10 ethyl 3,4,5-triacetoxybenzoate, 3,5-dimethoxy-4-hydroxybenzoic acid, 3,5-dihydroxybenzoic acid, 4-hydroxysalicylic acid, 4-fluorosalicylic acid, 1,2-dihydro-2-oxo-6-trifluoromethylpyridine-3-carboxamide, 2-hydroxycinnamic acid, 2,4-dichlorocinnamic acid, as described in WO 2004084631, WO 2005015994, WO 2006007981, WO 2005016001;

15

H) active compounds from the class of the 1,2-dihydroquinoxalin-2-ones, for example

1-methyl-3-(2-thienyl)-1,2-dihydroquinoxalin-2-one, 1-methyl-3-(2-thienyl)-1,2-dihydroquinoxaline-2-thione, 1-(2-aminoethyl)-3-(2-thienyl)-1,2-dihydroquinoxalin-2-one hydrochloride, 1-(2-methylsulfonylaminoethyl)-3-(2-thienyl)-1,2-dihydroquinoxalin-2-one, as described in WO 2005112630,

20

I) active compounds which, in addition to a herbicidal action against harmful plants, also have safener action on crop plants such as rice, such as, for example, "dimepiperate" or "MY-93" (see Pestic. Man.) (=S-1-methyl-1-phenylethyl piperidine-1-thiocarboxylate), which is known as safener for rice against damage by the herbicide molinate,

25

"daimuron" or "SK 23" (see Pestic. Man.) (= 1-(1-methyl-1-phenylethyl)-3-p-tolyl-urea), which is known as safener for rice against damage by the herbicide

30

imazosulfuron,

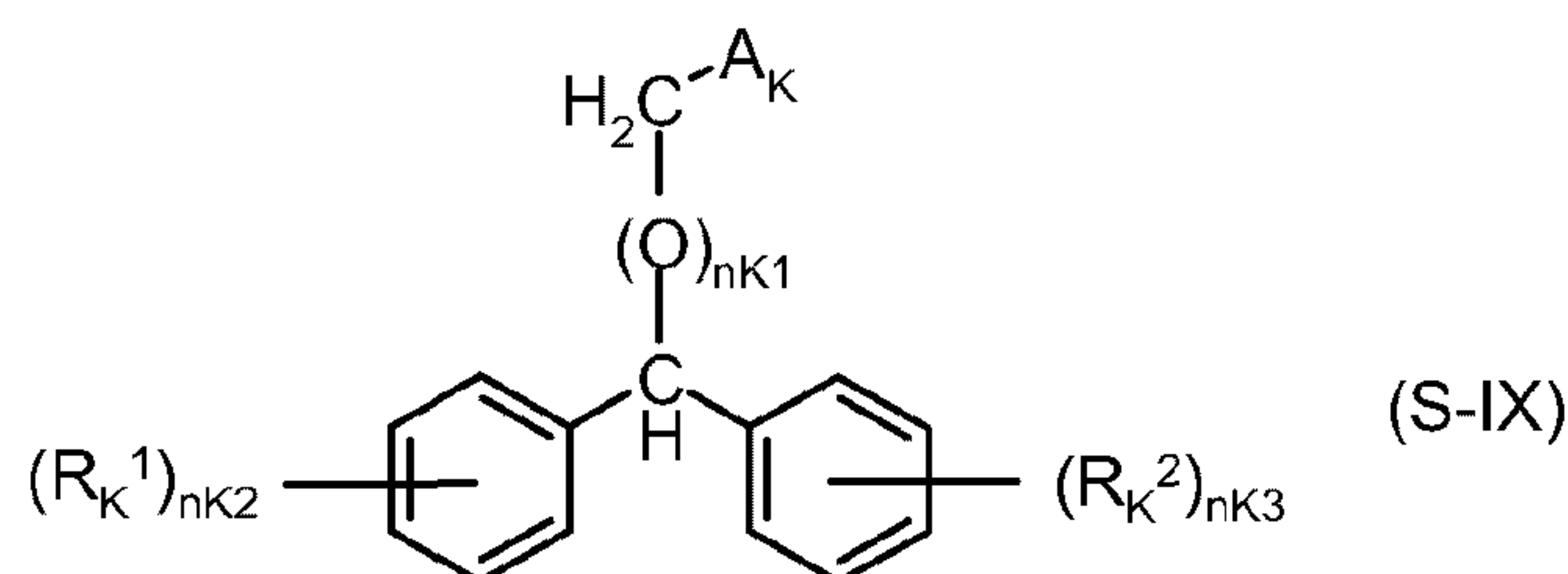
"cumyluron" = "JC-940" (= 3-(2-chlorophenylmethyl)-1-(1-methyl-1-phenyl-

ethyl)urea, see JP-A-60087254), which is known as safener for rice against damage by a number of herbicides,

"methoxyphenone" or "NK 049" (= 3,3'-dimethyl-4-methoxybenzophenone), which is known as safener for rice against damage by a number of herbicides,

- 5 "CSB" (= 1-bromo-4-(chloromethylsulfonyl)benzene) (CAS Reg. No. 54091-06-4 from Kumiai), which is known as safener against damage by a number of herbicides in rice,

- K) compounds of the formula (S-IX),
10 as described in WO-A-1998/38856



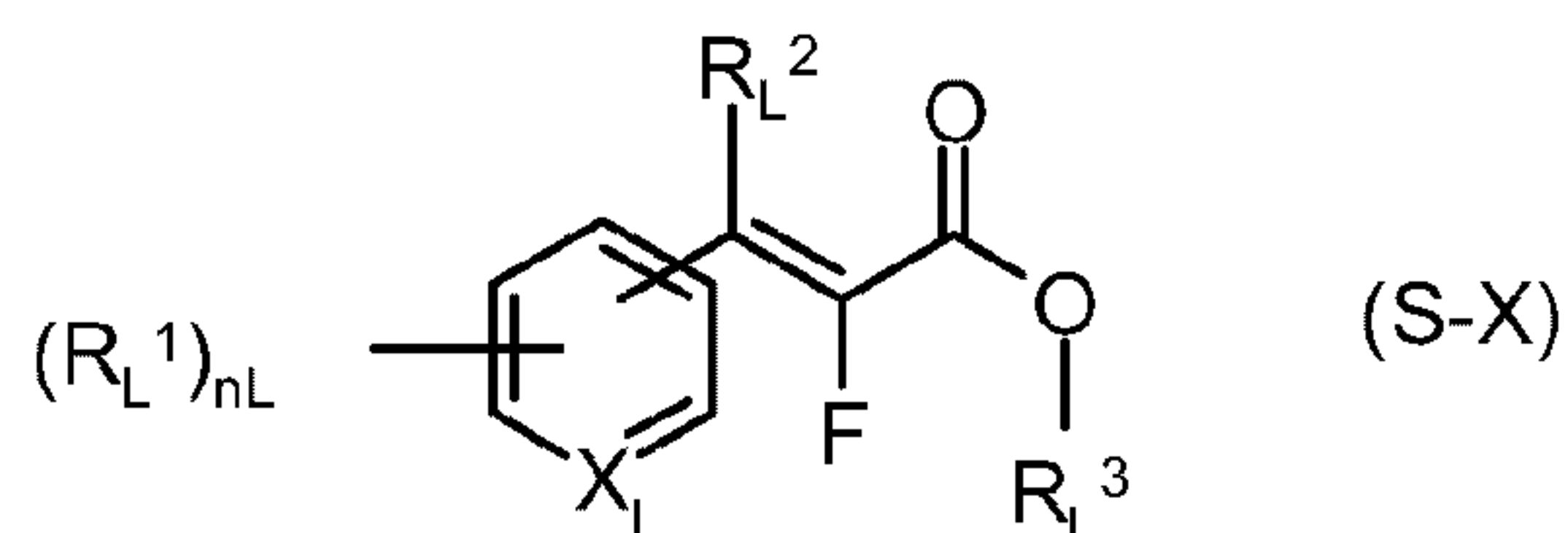
in which the symbols and indices have the following meanings:

- R_K^1, R_K^2 independently of one another are halogen, (C₁-C₄)-alkyl, (C₁-C₄)-alkoxy, (C₁-C₄)-haloalkyl, (C₁-C₄)-alkylamino, di-(C₁-C₄)-alkylamino, nitro;
15 A_K is COOR_K³ or COOR_K⁴
 R_K^3, R_K^4 independently of one another are hydrogen, (C₁-C₄)-alkyl, (C₂-C₆)-alkenyl, (C₂-C₄)-alkynyl, cyanoalkyl, (C₁-C₄)-haloalkyl, phenyl, nitrophenyl, benzyl, halobenzyl, pyridinylalkyl or alkylammonium,
20 n_K^1 is 0 or 1,
 n_K^2, n_K^3 independently of one another are 0, 1 or 2

preferably: methyl (diphenylmethoxy)acetate (CAS Reg. No.: 41858-19-9),

- 25 L) compounds of the formula (S-X),
as described in WO A-98/27049

68



in which the symbols and indices have the following meanings:

X_L is CH or N,

n_L is, in the case that $X=N$, an integer from 0 to 4 and,

5 in the case that $X=CH$, an integer from 0 to 5,

R_L^1 is halogen, (C₁-C₄)-alkyl, (C₁-C₄)-haloalkyl, (C₁-C₄)-alkoxy, (C₁-C₄)-haloalkoxy, nitro, (C₁-C₄)-alkylthio, (C₁-C₄)-alkylsulfonyl, (C₁-C₄)-alkoxycarbonyl, optionally substituted phenyl, optionally substituted phenoxy,

R_L^2 is hydrogen or (C₁-C₄)-alkyl,

10 R_L^3 is hydrogen, (C₁-C₈)-alkyl, (C₂-C₄)-alkenyl, (C₂-C₄)-alkynyl or aryl, where each of the carbon-containing radicals mentioned above is unsubstituted or substituted by one or more, preferably by up to three, identical or different radicals from the group consisting of halogen and alkoxy; or salts thereof,

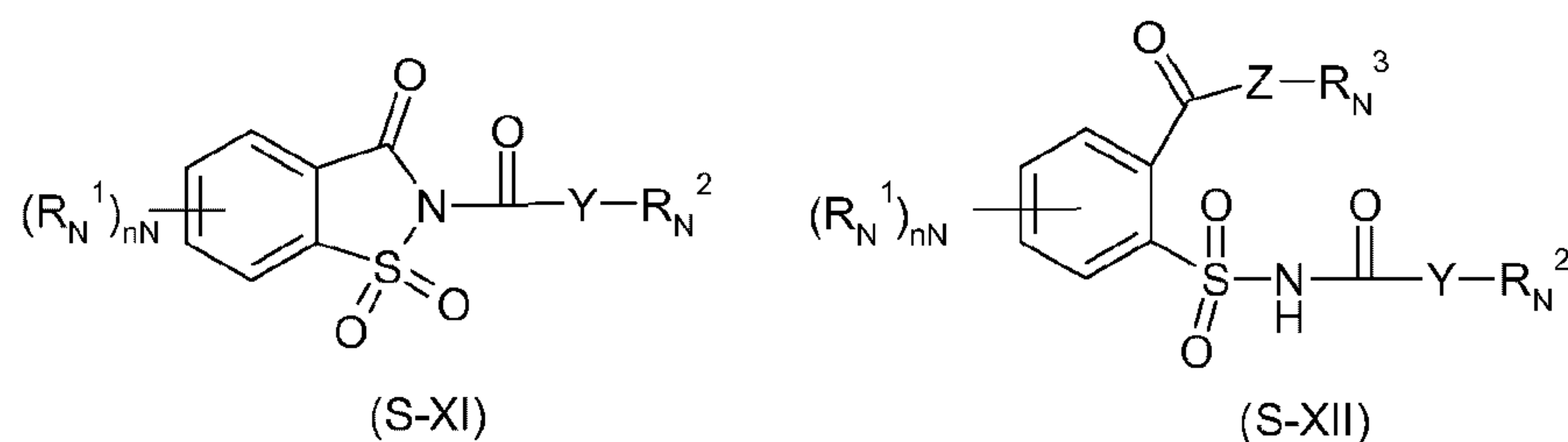
15 M) active compounds from the class of the 3-(5-tetrazolylcarbonyl)-2-quinolones, for example

1,2-dihydro-4-hydroxy-1-ethyl-3-(5-tetrazolylcarbonyl)-2-quinolone (CAS Reg. No.: 219479-18-2), 1,2-dihydro-4-hydroxy-1-methyl-3-(5-tetrazolylcarbonyl)-2-quinolone (CAS Reg. No.: 95855-00-8), as described in WO-A-1999000020,

20

N) compounds of the formula (S-XI) or (S-XII),

as described in WO-A-2007023719 and WO-A-2007023764



in which

69

R_N^1 is halogen, (C₁-C₄)-alkyl, methoxy, nitro, cyano, CF₃, OCF₃

Y, Z independently of one another are O or S,

n_N is an integer from 0 to 4,

R_N^2 is (C₁-C₁₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₆)-cycloalkyl, aryl, benzyl, halobenzyl,

5 R_N^3 is hydrogen, (C₁-C₆)-alkyl,

O) one or more compounds from the group consisting of:

1,8-naphthalic anhydride,

O,O-diethyl S-2-ethylthioethyl phosphorodithioate (disulfoton),

10 4-chlorophenyl methylcarbamate (mephenate),

O,O-diethyl O-phenyl phosphorothioate (dietholate),

4-carboxy-3,4-dihydro-2H-1-benzopyran-4-acetic acid (CL-304415, CAS Reg. No.: 31541-57-8),

2-propenyl 1-oxa-4-azaspiro[4.5]decane-4-carbodithioate (MG-838, CAS Reg. No.:

15 133993-74-5),

methyl [(3-oxo-1H-2-benzothiopyran-4(3H)-ylidene)methoxy]acetate (from

WO-A-98/13361; CAS Reg. No.: 205121-04-6),

cyanomethoxyimino(phenyl)acetonitrile (cyometrinil),

1,3-dioxolan-2-ylmethoxyimino(phenyl)acetonitrile (oxabetrinil),

20 4'-chloro-2,2,2-trifluoroacetophenone O-1,3-dioxolan-2-ylmethyloxime (fluxofenim),

4,6-dichloro-2-phenylpyrimidine (fencloirim),

benzyl 2-chloro-4-trifluoromethyl-1,3-thiazole-5-carboxylate (flurazole),

2-dichloromethyl-2-methyl-1,3-dioxolane (MG-191),

25 including the stereoisomers, and the salts customary in agriculture.

A mixture with other known active compounds, such as fungicides, insecticides, acaricides, nematocides, bird repellents, plant nutrients and soil structure improvers is likewise possible.

Some of the safeners are already known as herbicides and accordingly, in addition to the herbicidal action against harmful plants, also act by protecting the crop plants. The weight ratios of herbicide (mixture) to safener generally depend on the herbicide application rate and the effectiveness of the safener in question and may
5 vary within wide limits, for example in the range from 200:1 to 1:200, preferably from 100:1 to 1:100, in particular from 20:1 to 1:20. The safeners may be formulated analogously to the compounds of the formula (I) or their mixtures with other herbicides/pesticides and be provided and used as a finished formulation or as a tank mix with the herbicides.

10

The required application rate of the compound of the formula (I) varies depending, inter alia, on external conditions such as temperature, humidity and the type of herbicide used. It can vary within wide limits, for example between 0.001 and 10 000 g/ha or more of active substance; however, it is preferably between 0.5 and
15 5000 g/ha, particularly preferably between 0.5 and 1000 g/ha and very particularly preferably between 0.5 and 500 g/ha.

When the transgenic plant of the invention contains one or more other genes for tolerance towards other herbicides (as, for example, a gene which encodes a
20 mutated or unmutated EPSPS which confers on the plant tolerance to glyphosate herbicides or a pat or bar gene conferring tolerance to glufosinate herbicides), or when the transgenic plant is naturally resistant to another herbicide (such as sulfonylurea tolerance), the method according to the invention can comprise the simultaneous or chronologically staggered application of an HPPD inhibitor in
25 combination with the said herbicide or herbicide combination, for example glyphosate and/or glufosinate and/or sulfonylurea herbicides.

The invention also relates to the use of the chimeric gene encoding the HPPD of the invention as a marker gene during the transformation of a plant species, based on
30 the selection on the abovementioned HPPD inhibitor herbicides.

The present invention also relates to a method for obtaining a plant resistant to a triketone or a pyrazolate HPPD inhibitor, characterized in that the plant is transformed with a chimeric gene expressing in the plant an HPPD of the invention as defined herein.

5

In a particular embodiment, the invention relates to said method for obtaining a plant resistant to a triketone or a pyrazolate HPPD inhibitor, characterized in that the HPPD of the invention comprises SEQ ID No. 4 (from the amino acid position 2 to the amino acid position 368), or a synthetic DNA encoding the HPPD of the invention adapted to the codon usage of corn, rice, wheat, soya spp, sugarcane,
10 onion, Brassica species plants, or cotton.

In another particular embodiment, the invention relates to said method for obtaining a plant resistant to a triketone HPPD inhibitor selected from tembotrione,
15 mesotrione, diketonitrile, isoxaflutole, sulcotrione, tefuryltrione, and bicyclopyrone.

In another particular embodiment, the invention relates to said method for obtaining a plant resistant to a triketone or a pyrazolate HPPD inhibitor, characterized in that the plant also comprises a plant-expressible chimeric gene encoding a PDH
20 (prephenate dehydrogenase) enzyme, or an enzyme with at least PDH.

The invention also relates to a method for controlling weeds in an area or a field, which method comprises planting in this area or field transformed plants resistant to a triketone or a pyrazolate HPPD inhibitor which has been obtained according to
25 the method described above, or transformed seeds which originates from them, and in applying a dose which is toxic for the weeds of said triketone or pyrazolate HPPD inhibitor without significantly affecting the said transformed seeds or the said transformed plants.

30 The invention also relates to a method for obtaining oil or meal comprising growing a transformed plant resistant to a triketone or a pyrazolate HPPD inhibitor which

has been obtained according to the method described above, or a transformed seed which originates from such plant, optionally treating such plant or seed with a triketone or a pyrazolate HPPD inhibitor, harvesting the grains and milling the grains to make meal and extract the oil.

5

The invention also relates to the use of an HPPD of the invention as described above, characterized in that the HPPD inhibitor is a triketone HPPD inhibitor selected from tembotrione, mesotrione, topramezone, bicyclopyrone, tefuryltrione and sulcotrione.

10

The present invention also relates to a host organism, in particular plant cells or plants, which contain a chimeric gene comprising a sequence encoding an HPPD according to the invention, and which also contain a gene functional in this host organism allowing overexpression of a prephenate dehydrogenase (abbreviated
15 herein as PDH) enzyme.

The term "PDH enzyme", as used herein, refers to any natural or mutated PDH enzyme exhibiting the PDH activity of conversion of prephenate to HPP. In particular, said PDH enzyme can originate from any type of organism. An enzyme
20 with PDH activity can be identified by any method that makes it possible either to measure the decrease in the amount of prephenate substrate, or to measure the accumulation of a product derived from the enzymatic reaction, i.e. HPP or one of the cofactors NADH or NADPH.

25 Many genes encoding PDH enzymes are described in the literature, and their sequences can be identified on the website <http://www.ncbi.nlm.nih.gov/entrez/>. Particularly known is the gene encoding the PDH enzyme of the yeast *Saccharomyces cerevisiae* (Accession No. S46037) as described in Mannhaupt et al. (1989) Gene 85, 303-311, of a bacterium of the *Bacillus* genus, in particular of
30 the species *B. subtilis* (Accession No. P20692) as described in Henner et al. (1986) Gene 49 (1) 147-152, of a bacterium of the *Escherichia* genus, in particular of the

species *E. coli* (Accession No. KMECTD) as described in Hudson et al. (1984) J. Mol. Biol. 180(4), 1023-1051, or of a bacterium of the *Erwinia* genus, in particular of the species *E. herbicola* (Accession No. S29934) as described in Xia et al. (1992) J. Gen. Microbiol. 138(7), 1309-1316.

- 5 The invention further relates to a method for obtaining a host organism, particularly a plant cell or a plant, resistant to an HPPD inhibitor by integrating in such organism at least one nucleic acid sequence or one chimeric gene as defined above, and by further transforming it, simultaneously or successively, with a gene functional in this host organism allowing expression of a PDH (prephenate dehydrogenase) enzyme.
- 10 In a particular embodiment, the invention relates to a method for obtaining a host organism, particularly a plant cell or a plant, resistant to a triketone or pyrazolate HPPD inhibitor, particularly tembotrione, mesotrione topramezone, bicyclopyrone, isoxaflutole, pyrasulfotole, tefuryltrione, or sulcotrione.

Means and methods which could be used for obtaining a host organism, particularly a

15 plant cell or a plant, transformed both with a gene allowing overexpression of an HPPD enzyme, and with a gene allowing overexpression of a PDH enzyme are extensively described in WO 04/024928.

The present invention as claimed relates to:

- a chimeric gene comprising a coding sequence operably-linked to a plant
- 20 expressible promoter, wherein the coding sequence comprises a nucleic acid sequence which encodes a hydroxyphenylpyruvate dioxygenase (HPPD) protein comprising the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to amino acid position 368; and

- a method for obtaining a plant cell tolerant to a hydroxyphenylpyruvate
- 25 dioxygenase (HPPD) inhibitor herbicide, comprising introducing into said plant cell a chimeric gene of the invention.

The reference in this specification to any prior publication (or information derived from it), or to any matter which is known, is not, and should not be taken as an acknowledgement or admission or any form of suggestion that such prior publication (or information) or known matter forms part of the common general knowledge in the
5 field of this invention.

FIGURES

FIG.1 Map of the plasmid pSE420::FMP29e

FIG.2 Map of the T-DNA inserted into the tobacco plants

FIG.3 Map of the T-DNA inserted in the different plants according to

Examples 5 to 11; Abbreviations having the following meanings
 A, B, C and G, tobacco plants, D, E and F, Zea mays plants, H,
 soybean plants, I, rice plants, and J, cotton plants. 35S:
 CaMV35S promoter, KanR: gene conferring resistance to the
 5 antibiotic kanamycin, nos: nopaline synthase promoter, Ter:
 terminator, H6: sequence coding for an His TAG, OTP:
 optimized transit peptide, BAR (Bialaphos resistant, WO
 8705629) and PAT (phosphinothricin N-Acetyltransferase, EP
 257542):genes conferring tolerance to Bialaphos,
 10 phosphinothricin or glufosinate, 2mEPSPS: gene coding for the
 double mutant (Thr102Ile and Pro106Ser) EPSPS (5-
 enolpyruvylshikimate synthase) from Zea mays
 (US 20030027312), 2mAHAS: gene coding for the double
 mutant ALS (acetolactate synthase) from Arabidopsis
 15 (Pro197Ala and Trp574Leu; US 5378824, HA: histone promoter
 from Arabidopsis gene, TEV: tobacco etch virus, FMP29e: gene
 coding for FMP29 optimized for the expression in E coli with an
 sequence coding for an His TAG at its 5' extremity, FMP29t:
 gene coding for FMP29 optimized for the expression in
 20 dicotyledoneous plants with an sequence coding for an His TAG
 at its 5' extremity, FMP29t-h, gene coding for FMP29 optimized
 for the expression in dicotyledoneous plants, FMP29m, gene
 coding for FMP29 optimized for the expression in Zea mays
 plants, LB, left border, RB, right border.

25

SEQUENCES LISTING

SEQ ID No. 1: Nucleic acid sequence encoding *Picrophlus torridus* HPPD

- SEQ ID No. 2: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *E. coli*, plus containing at the 5' end a nucleic acid encoding an alanine and 6 histidine amino acids.
- 5 SEQ ID No. 3: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Nicotiana tabaccum* plus containing at the 5' end a nucleic acid sequence encoding an optimized transit peptide and an HIS Tag.
- SEQ ID No. 4: *Picrophilus torridus* HPPD amino acid sequence derived from SEQ ID No. 1
- 10 SEQ ID No. 5: Protein encoded by SEQ ID No. 2
- SEQ ID No. 6: *Picrophilus torridus* HPPD amino acid sequence (SEQ ID No. 4) fused with OTP (optimized transit peptide (WO 2009/144079))
- SEQ ID No. 7: Protein encoded by SEQ ID No. 3
- SEQ ID No. 8: Nucleic acid sequence encoding *Arabidopsis thaliana* HPPD
- 15 SEQ ID No. 9: *Arabidopsis thaliana* HPPD amino acid sequence
- SEQ ID No. 10: Protein encoded by SEQ ID No. 8 plus an additional alanine directly downstream of the initial amino acid methionine followed by 6 histidine amino acids
- SEQ ID No. 11: Protein of SEQ ID No. 9 plus the OTP sequence located at the N-terminal extremity of the protein.
- 20 SEQ ID No. 12: Protein of SEQ ID No. 10 plus the OTP sequence directly located at the N-terminal extremity of the protein.
- SEQ ID No. 13: Primer sequence Xho-OTP-for
- SEQ ID No. 14: Primer sequence NcoI-OTP-rev
- 25 SEQ ID No. 15: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for dicotyledoneous plants
- SEQ ID No. 16: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Zea mays* plants
- SEQ ID No. 17: Nucleic acid sequence encoding *Picrophilus torridus* HPPD
- 30 optimized for *Brassica napus* plants

- SEQ ID No. 18: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Beta vulgaris* plants
- SEQ ID No. 19: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Gossypium hirsutum* plants
- 5 SEQ ID No. 20: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Glycine max* plants
- SEQ ID No. 21: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Hordeum vulgare* plants
- 10 SEQ ID No. 22: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Oryza sativa* plants
- SEQ ID No. 23: Nucleic acid sequence encoding *Picrophilus torridus* HPPD optimized for *Triticum aestivum* plants

15 EXAMPLES

The various aspects of the invention will be better understood with the aid of the experimental examples which follow. All the methods or operations which are described below in these examples are given by way of example and correspond to a choice which is made from among the different methods which are available for

20 arriving at the same or similar result. This choice has no effect on the quality of the result and, as a consequence, any suitable method can be used by the skilled person to arrive at the same or similar result. The majority of the methods for manipulating DNA fragments are described in "Current Protocols in Molecular Biology" Volumes 1 and 2, Ausubel F.M. et al., published by Greene Publishing

25 Associates and Wiley Interscience (1989) or in Molecular cloning, T. Maniatis, E.F. Fritsch, J. Sambrook, 1982, or in Sambrook J. and Russell D., 2001, Molecular Cloning: a laboratory manual (Third edition)

Example 1

Preparation of *Picrophlus torridus* HPPD (named FMP29e) of SEQ ID No. 5 and of the *Arabidopsis thaliana* HPPD identified by SEQ ID No. 10.

The *Arabidopsis thaliana* AtHPPD coding sequence (1335 bp; Genbank AF047834; 5 WO 96/38567) was initially cloned into the expression vector pQE-30 (QIAGEN, Hilden, Germany) in between the restriction sites of BamHI and HindIII. The obtained vector was called "pQE30-AtHPPD".

The original *Picrophlus torridus* HPPD sequence (1107 bp) coding for the protein 10 listed under the accession number Q6K798 at UniProtKB/TrEMBL was modified and synthesized using an *Escherichia coli* K12 optimized codon usage (Eurofins MWG operon (Ebersberg, Germany), GENEius software) and cloned in a modified pBluescript vector (Eurofins MWG operon, Ebersberg, Germany). In this vector, the sequence corresponding to the MCS (multiple cloning site) was partially removed 15 that only the sequences corresponding to the recognition of the restriction enzyme HindIII remained on the both side of the insert.

At the 5' end, directly downstream to the ATG was inserted a nucleic acid sequence coding for an alanine amino acid and a nucleic acid sequence encoding a N-terminal HIS6-Tag (6x HIS, encoded by: cat cac cat cat cat cac). Upstream to the 20 ATG, two additional cytosine base pairs were added in order to obtain a sequence corresponding to the recognition site of the restriction enzyme NcoI and downstream to the stop codon the sequences corresponding to the recognition site of the restriction enzyme XbaI were added. The resulting vector "pBluescript-FMP29e" was digested with the restriction enzymes NcoI and XbaI, the band migrating not to the 25 length of the size of the vector approximately 3000 bp corresponding to the DNA was separated on an agarose gel per electrophoresis. Then the DNA coding for the HPPD was purified using the MinElute™ Gel Extraction Kit (Qiagen, Hilden, Germany) and cloned into the pSE420(RI)NX vector (see below) previously cut with the same restriction enzymes.

30 The cloning and expression vector pSE420(RI)NX (5261 bp) is based on the plasmid pSE420 by Invitrogen (Karlsruhe, Germany). Modifications of this vector

include the addition of a *nptII* gene (neomycin phosphotransferase; Sambrook and Russell, 2001, Molecular Cloning: a laboratory manual (Third edition)) conferring tolerance to the antibiotic kanamycin and is missing the majority of the superlinker region (multiple cloning site).

- 5 The plasmid possesses the *trp-lac* (*trc*) promoter and the *lacI^q* gene that provides the *lac* repressor in every *E. coli* host strain. The *lac* repressor binds to the *lac* operator (*lacO*) and restricts expression of the target gene; this inhibition can be alleviated by induction with Isopropyl β -D-1-thiogalactopyranoside (IPTG).
- 10 The resulting vector was called "pSE420(RI)NX-FMP29e" (see Figure1) and it was used to transform *Escherichia coli* BL21 cells (Merck, Darmstadt, Germany). For the AtHPPD (*Arabidopsis thaliana* HPPD) that was used as reference see WO 2009/144079.
- 15 Expression of HPPD was carried out in *E. coli* K-12 BL21 containing pQE30-*AtHPPD* or pSE420(RI)NX-FMP29e. Cells were allowed to grow until OD reached 0.5, then expression was initiated from the *trp-lac* (*trc*) promoter by induction with 1 mM IPTG which binds to the *lac* repressor and causes its dissociation from the *lac* operon. Expression was carried out over 15 h at 28 °C.
- 20 To prepare the pre-starter culture, 2 mL of TB medium (100 $\mu\text{g}\cdot\text{mL}^{-1}$ carbenicillin) were inoculated with 50 μL of an *E. coli* K-12 BL21 glycerol stock. The pre-starter culture was incubated at 37 °C with shaking at 140 rpm for 15 h. 200 μL of the pre-starter culture was used to initiate the starter culture (5mL TB supplement with 100 $\mu\text{g}\cdot\text{L}^{-1}$), which was incubated 3 h at 37°C.
- 25 To prepare the main culture, 400 mL of TB medium (100 $\mu\text{g}\cdot\text{mL}^{-1}$ carbenicillin) were inoculated with 4 mL of the starter culture. This starter culture was incubated at 37 °C with shaking at 140 rpm until OD₆₀₀ 0.5 was reached. Then recombinant protein expression was induced with 400 μL of 1M IPTG solution. The cells were allowed to grow for an additional hour under these conditions, then the temperature
- 30 was lowered to 28°C and the culture was shaken at 140 rpm for 15 h. Cells were

harvested by centrifugation at 6000 x g for 15 min at 4 °C. Then cell pellets were stored at -80 °C.

5 Isolation and purification of His₆-AtHPPD and His₆-FMP29e in native form

Lysis of cells

Cells were lysed using Lysozyme, an enzyme that cleaves the 1,4-β-linkages between N-acetylmuramic acid and N-acetyl-D-glucosamine residues in
10 peptidoglycan which forms the bacterial cell wall. Cell membranes were then disrupted by the internal pressure of the bacterial cell. In addition, the lysis buffer contained Benzonase[®] Nuclease, an endonuclease that hydrolyzes all forms of DNA and RNA without damaging proteins and thereby largely reduces viscosity of the cell lysate. Lysis under native conditions was carried out on ice.

15 For purification of His₆-tagged proteins the QIAexpress[®] Ni-NTA Fast Start Kit was used following the user manual instruction.

Purification of His₆-tagged proteins by immobilized metal ion affinity chromatography (IMAC)

20

The cleared cell lysate (10 mL) obtained after centrifugation of the lysis reaction was loaded onto a Ni-NTA Fast Start Column from the QIAexpress[®] Ni-NTA Fast Start Kit (Qiagen, Hilden, Germany) and purification was carried out according to the instruction manual. The His₆-tagged protein was eluted with 2.5 mL of elution buffer.

25

Desalting of HPPD solutions by gel filtration

HPPD solutions eluted from a Ni-NTA Fast Start Column with 2.5 mL of elution
30 buffer were applied to a Sephadex G-25 PD-10 column (GE Healthcare, Freiburg,

Germany) following the user manual instruction. After the whole sample had entered the gel bed, elution was performed with 3.5 mL of storage buffer.

The HPPD solutions eluted from the desalting column were frozen at -80 °C in 1 mL aliquots.

5

Determination of HPPD protein concentration using the Bradford protein assay
Protein concentration was determined using the standard Bradford assay (Bradford, (1976), Anal Biochem 72: 248-254).

10

Determination of purity of HPPD solutions using SDS-PAGE

15

The integrity of the eluted protein was checked by SDS-PAGE protein gel electrophoresis using the gel NuPAGE[®] Novex 4-12 % Bis-Tris Gels (Invitrogen, Karlsruhe, Germany), approximately 10 µg of protein were loaded. 10 µL of Laemmli Sample Buffer was added to 1-10 µL of protein solution and the mixture was
20 incubated at 90 °C for 10 min. After short centrifugation step, the whole mixture was loaded into a slot of an SDS gel previously fixed in a XCell SureLock[™] Novex Mini-Cell gel chamber filled with NuPAGE[®] MOPS SDS Running Buffer (diluted from the 20 x-solution with ddH₂O). A voltage of 150 was then applied to the gel chamber for 1 h. For staining of protein bands, the gel was immersed in Coomassie Brilliant Blue
25 R-250 Staining Solution. For destaining of the polyacrylamide gel, it was immersed in Coomassie Brilliant Blue R-250 Destaining Solution until protein bands appear blue on a white gel.

30 Example 2

Kinetic characterization and evaluation of tolerance to HPPD inhibitors of HPPD enzymes "SEQ ID No. 5" and "SEQ ID No. 10".

The HPPD activity was checked by the standard spectrophotometric assay (method
5 extensively described in WO 2009/144079)

Determination of HPPD in vitro kinetic properties

K_m , V_{max} , and k_{cat} values for different HPPD enzyme preparations and K_i , $K_1=K_{on}$, and
10 $K_{-1}=K_{off}$ for different HPPD inhibitors were determined using a HPLC assay for
measurements of HPPD activity. The assay mixtures contained in a volume of 1 ml
150 mM Tris-HCl buffer at pH 7.8, 10 mM sodium ascorbate, 650 units of bovine
catalase (Sigma C30 (Sigma-Aldrich, Munich, Germany), 34 mg protein/ml, 23,000
units/mg), and appropriate amounts of HPP, purified HPPD enzyme and HPPD
15 inhibitors. For K_m , V_{max} , and k_{cat} value determination HPP concentrations in the
assay mixture were varied between 10 and 400 μ M. For K_i , $K_1=K_{on}$, and $K_{-1}=K_{off}$ value
determination 2 mM HPP was used. All assays were started by the addition of HPPD
enzyme to the assay mixture and stopped at a series of times between 0 and 240 s
by addition of 200 μ l of the reaction mixture to reaction assay tubes containing 20 μ l
20 10% perchloric acid. Precipitated protein was pelleted by a 5 minute centrifugation
at 10,000 g. 100 μ l of the supernatant were loaded onto a 250 x 4mm Knauer
(Berlin, Germany) Eurospher 100-5 C18-column equilibrated with 10% methanol,
0.1% trifluoroacetic acid (buffer A). The column was eluted, also at 1.5 ml/min, using
a 4 minute wash with buffer A, followed by a 3 min wash with 95% methanol and by
25 a further 2 minute wash with buffer A. The elution of HGA (homogentisic acid) and
HPP (hydroxyphenylpyruvate) was monitored at 292 nm. HGA elutes at around 5
minutes and HPP elutes later. A standard set of concentrations of HGA were used to
provide a standard curve in order to calibrate the 292 nm absorbance of the HGA
peak versus HGA concentration.
30 For K_m and V_{max} value determinations the initial rates of the HPPD reaction at
different substrate concentrations were determined from plots of HGA formed versus

time and fitted to the Michaelis-Menten equation for unireactant enzymes using the ID Business Solutions Ltd. (www.idbs.com) XLfit software suite. For the determination of K_i , $K_1=K_{on}$, and $K_{-1}=K_{off}$ values the time-courses of the HPPD reaction at different inhibitor concentrations were fitted to the equations for

- 5 Mechanism A, competitive inhibition, for tight-binding inhibitors (Cha, S. (1975) Tight-binding inhibitors – I. Kinetic behaviour. Biochemical Pharmacology 24, 2177-2185) using the ID Business Solutions Ltd. XLfit software suite

- 10 Determination of HPPD activity in presence of several HPPD inhibitors

In this content, pl_{50} -value means the log value of the concentration of inhibitor necessary to inhibit 50% of the enzyme activity in molar concentration.

- 15 pl_{50} -values for HPPD inhibitors were determined from dose-response plots of HPPD activity versus inhibitor concentration using the assay extensively described in WO 2009/144079 at 2 mM fixed HPP concentration and 3 minutes fixed incubation time using the ID Business Solutions Ltd. XLfit software suite.

- 20 Table 1: Determination of pl_{50} HPPD enzymes (*Arabidopsis thaliana* "SEQ ID No. 10" and *Picrophilus torridus* "SEQ ID No. 5") and their respective tolerance to the several listed below HPPD inhibitors tembotrione, diketonitrile, mesotrione, bicyclopyrone, pyrasulfotole, sulcotrione, pyrazolate, tefuryltrione, and benzofenap. The symbol ">>" means that the value was far higher than the one indicated but
25 could not be precisely calculated within in the range of concentration of inhibitor tested (2.5×10^{-6} , 5.0×10^{-6} , 1.0×10^{-5} , 2.5×10^{-5} , 6.3×10^{-5} , and $2.5 \times 10^{-4} M$).

	Tembotrione	Diketoneitrile	Mesotrione	Bicyclopyrone
SEQ ID				
No. 10	>>5.6	>>5.6	>>5.6	5.2

83

SEQ ID

No. 5	5.5	4.9	>>5.6	3.4
-------	-----	-----	-------	-----

	Pyrasulfotole	Sulcotrione	Pyrazolate	Tefuryltrione	Benzofenap
--	---------------	-------------	------------	---------------	------------

SEQ ID

No. 10	5.4	>>5.6	5.4	>>5.6	>>5.6
--------	-----	-------	-----	-------	-------

SEQ ID

No. 5	4.0	>5.6	4.2	5.5.	4.8
-------	-----	------	-----	------	-----

Table2: Determination of percentage of inhibition in presence of 5.0×10^{-6} M inhibitors compared to the activity measured in absence of the inhibitor for the HPPD

- 5 originated from *Arabidopsis thaliana* (SEQ ID No. 10) and from *Picrophilus torridus* (SEQ ID No. 5).

Bicyclopyron

	Tembotrione	Diketonitrile	Mesotrione	e
--	-------------	---------------	------------	---

SEQ ID

No. 10	92	87	86	29
--------	----	----	----	----

SEQ ID

No. 5	58	26	84	24
-------	----	----	----	----

	Pyrasulfotole	Sulcotrione	Pyrazolate	Tefuryltrione	Benzofenap
--	---------------	-------------	------------	---------------	------------

SEQ ID

No. 10	69	74	61	100.	90
--------	----	----	----	------	----

SEQ ID

No. 5	7	92	16	87.	44
-------	---	----	----	-----	----

- On the above Tables 1 and 2, it can be clearly seen, that the Euryarchaeota HPPD
- 10 "SEQ ID No. 5" showed superior level of tolerance to all tested HPPD inhibitors than

the plant at all tested HPPD inhibitor concentrations than observed by employing the HPPD "SEQ ID No. 10" under identical experimental conditions.

- 5 Example 3: Construction of chimeric genes for the evaluation HPPD inhibitor herbicide tolerance in tobacco plants.

A) Construction of the chimeric genes

10

The vector pRP-RD224 (extensively described in WO 2009/144079) containing the sequence coding for the OTP was used for PCR-mediated attachment upstream of the nucleic acid sequence corresponding to the recognition site of the restriction enzyme XhoI and downstream of the nucleic acid sequence corresponding to the
15 recognition site of the restriction enzyme NcoI. The obtained PCR product was cloned in the vector pCR®-Blunt II-TOPO® (Invitrogen, Karlsruhe, Germany) following the user manual instruction. The resulting vector was called "pCR-TOPO-OTP". The insertion of the correct sequence was confirmed per standard DNA sequencing. The DNA corresponding to the OTP was digested with the restriction
20 enzymes NcoI and XhoI, separated per appropriate gel electrophoresis and cloned into the plasmid pRT100 (Toepfer, (1987), Nucleic Acids Res 15:5890) previously and correspondingly digested with Nco I and XhoI restriction enzymes. The plasmid pRT100 is containing the CaMV35S promoter and CaMV35S terminator. The resulting vector was subsequently digested with the restriction enzymes NcoI and
25 XbaI. The vector pSE420(RI)NX-FMP29e (see Figure1) was subjected to the restriction enzymes NcoI and XbaI in order to obtain the DNA fragment corresponding to the "SEQ ID No. 2". The resulting vector was digested by employing the restriction enzyme HindIII to subclone the CaMV35S::OTP::FMP29e::CaMV35-term cassette (see Figure2) into the binary
30 vector pBin19 (Bevan (1984), Nucleic Acids Res. 12:8711-8721.) previously

digested with the same enzyme and dephosphorylated. The resulting vector was called "FMP29ebv".

The vectors pQE-30-AtHPPD was used for PCR-mediated attachment of an NcoI
5 restriction site and of a sequence encoding an N-terminal His₆-Tag to the 5' ends and a XbaI restriction site to the 3' ends of AtHPPD.

The PCR product of the AtHPPD gene was isolated from an agarose gel, cut with
the restriction enzymes NcoI and XbaI, purified with the MinElute™ PCR Purification
10 Kit (Qiagen, Hilden, Germany) and cloned into the pSE420(RI)NX vector cut with the
same restriction enzymes.

The generated vector was called "pSE420(RI)NX-AtHPPD" and was digested with
the restriction enzymes NcoI and XbaI and cloned into the previously opened vector
pRT100 (Toepfer et al., (1987), Nucleic Acids Res 15:5890) containing the
15 CaMV35S promoter and CaMV35S terminator. The generated vector was called
"pRT100-AtHPPD".

The vector pCR-TOPO-OTP was digested with the restriction enzymes NcoI and
XhoI, and the DNA band corresponding to the OTP was cloned in the previously
opened vector pRT100-AtHPPD with the above mentioned restriction enzymes. The
20 resulting vector was subsequently digested with restriction enzyme HindIII and the
expression cassette of interest was cloned into the previously opened and
dephosphorylated binary vector pBin19. The resulting vector was called
"AtHPPDbv".

25 The binary vectors FMP29ebv and AtHPPDbv were used to transform
Agrobacterium tumefaciens (ATHV derived from EHA101) competent cells selected
on YEB media supplemented with the antibiotics kanamycin and rifampicin
(extensively described in the patent application US005925808A).

30 These *Agrobacterium* strains containing the binary vectors of interest (FMP29ebv or
AtHPPDbv) were used to transform leaf discs from tobacco *Nicotiana tabacum* L. cv

Samsun NN plants, having approximately a size of 5x5mm² as extensively described in Horsch et al., (1985), Science 227 ; 1229-1231.

The leaf disks were co-cultivated for 2 days with *Agrobacterium tumefaciens* cells containing either the binary vector FMP29ebv or AtHPPDbv. Then the leaf disks

- 5 were transferred to a media allowing the regeneration of shoots for 6 weeks on MS (Musharige and Skoog, (1962), Physiol Plant 15(3): 473-497) media supplemented with BAP (1 mg/mL; Benzylaminopurine), carbenicillin (250 mg/mL), cefotaxime (250 mg/mL), kanamycin (75 mg/mL) and tembotrione (10⁻⁶ M)

- 10 Regenerated calli were transferred on media to induce the development of roots for 6 to 12 weeks: MS (1/2), supplemented with carbenicillin (250 mg/mL), cefotaxime (250 mg/mL), kanamycin (75 mg/mL), and tembotrione (10⁻⁶ M).

After 6 weeks on this media, the shoots transformed with *Agrobacterium tumefaciens* cells containing the binary vector AtHPPDbv, were transferred on the

- 15 same media depleted of HPPD inhibitor tembotrione.

The results are summarized on Table 5, below.

During the entire experiment, the plates containing the leaf disk were located in a growth chamber under controlled conditions (light 16 h, night 8 h, 25°C).

20

Rooting of calli

Regenerated shoot calli from a cell transformed with a nucleic acid sequence

- 25 encoding an HPPD comprising SEQ ID No. 11 (*Arabidopsis thaliana*) or SEQ ID No. 7 (*Picrophilus torridus*) were transferred to a media inducing root growth which media was further supplemented with the HPPD inhibitor tembotrione for 6 to 12 weeks. On none of the events containing the HPPD defined by SEQ ID No. 11 (*Arabidopsis thaliana*) or none transformed calli, root growth was observed under the
- 30 above given conditions. Contrary to this, under the identical conditions, the calli containing the HPPD defined by SEQ ID No. 7 clearly developed numerous and

healthy roots (see Table 3, below).

5

Table 3

Calli containing:	Events selected for molecular analysis	% Elongation & rooting on 10^{-6} M tembotrione	Numbers of events rooted on media without tembotrione
SEQ ID No. 11	21	0	5
SEQ ID No. 7	23	65	15

Leaf disk regeneration

10

Leaf disks were cut from plants containing HPPD SEQ ID No. 11 (*Arabidopsis thaliana*) or SEQ ID No. 7 (*Picrophilus torridus*), followed by regeneration for 6 weeks under standard culture conditions on MS media supplemented with BAP (1 mg/mL; Benzylaminopurine), carbenicillin (250 mg/mL), cefotaxime (250 mg/mL) and further comprising one of the following listed HPPD inhibitors at the mentioned concentration (tembotrione (10^{-6} M), diketonitrile ($5 \cdot 10^{-6}$ M), Mesotrione (10^{-6} M) and bicyclopyrone (10^{-6} M)) with a media containing none HPPD inhibitors as the positive control. At the end of the experiments the level of regeneration was evaluated as followed:

15

20 “-” means that the leaf disks looked the same as leaf disk from wild type tobacco plants on media supplemented with the inhibitor mentioned above.

“++++” means that the leaf disks looked like the leaf disks from the wild type tobacco plants on media without inhibitor.

“+”, “++”, and “+++” indicate regenerated leaf disks were heavily (+), medium (++)

25 and less (+++) affected by the presence of the inhibitors.

The results of the experiments are summarized in Table 4.

Table 4: Effects of various HPPD inhibitors the regeneration of leaf disk originating from transgenic plants comprising either a gene coding for an HPPD obtained from
5 *Arabidopsis* (SEQ ID No. 11) or from *Picrophilus torridus* SEQ ID No. 7.

Leaf disks containing	Control	Tembotrione	Diketonitrinile	Mesotrione	Bicyclopyrone
SEQ ID NO 11	++++	-	-	-	-
SEQ ID NO7	++++	+++	++	++	++

10 Whereas in case of plants containing HPPD defined by SEQ ID No. 7 (*Picrophilus torridus*) showed the same or only slightly reduced regeneration compared to this un-treated control, the corresponding plants containing HPPD defined by SEQ ID
15 No. 11 (*Arabidopsis thaliana*) did not show any regeneration but developed clearly visible bleaching phenotype compared to the untreated control in the presence of all tested HPPD inhibitors.

Example 4: Glasshouse trials to evaluate tolerance to HPPD inhibitor herbicides of transgenic tobacco plants expressing a gene coding for tolerant HPPD protein

20

Preparation of transgenic plant lines expressing either *Arabidopsis* or FMP29 HPPD enzymes. Glasshouse testing for herbicide tolerance.

Response to tembotrione, isoxaflutole and bicyclopyrone

T0 Tobacco plants containing either the gene from *Arabidopsis* coding for HPPD or the gene FMP29e from *Picrophilus torridus* coding for FMP29 HPPD, mentioned above (Example3), were transferred to the glasshouse (28/20°C), to develop further and produce seeds. Those seeds were harvested and put on soil (ED73 mixed with sand and osmocote Pro) to germinate in the glasshouse (28/20°C). Three to four weeks later, plantlets were transferred to single pots containing the soil mentioned above. Two weeks later, plants of a size 4-6 cm diameter were sprayed with either

- tembotrione at 100gAI/ha prepared from a WP20 (wetable powder 20%) formulation supplemented with ammonium sulfate and methyl ester raps oil, or
- 10 - isoxaflutole at 100gAI/ha prepared from a WP20 formulation supplemented with ammonium sulfate and methyl ester raps oil, or
- bicyclopyrone at 100gAI/ha prepared from a WP20 formulation supplemented with ammonium sulfate and methyl ester raps oil, or
- "blind formulation" made from a WP20 formulation without active ingredient
- 15 (AI) supplemented with ammonium sulfate and methyl ester raps oil, and were then transferred to a growth chamber with adequate light conditions (20000Lux).

Seven days after the application (DAT) of the different herbicides, the symptoms in tranformed plants were evaluated in comparison to the response observed on the wild type tobacco plants sprayed at the same time and under the same conditions as the tobacco plants containing the transgenes (100% means the plants displayed the same bleaching phenotype as the wild type plants, 0% means that the plants looked like the wild type plants treated with the "blind formulation", and intermediate percentage represent the degree of observed symptoms).

25

Table 5: Wild type tobacco plants (A) and T1 populations of tobacco events containing alternatively, the expression cassettes described above having the promoter CaMV 35S, the sequence coding for OTP and the sequence coding for *Arabidopsis* HPPD (B) or the promoter CaMV35S, the sequence encoding OTP, and the sequence FMP29e coding for the HPPD FMP29 (C). Assessments of herbicidal damage at 7 days after application (DAT) per spray with 100 g AI / ha of tembotrione

30

or isoxaflutole supplemented with ammonium sulfate and methyl ester rap seed oil. It is clear that plants containing FMP29e gene were far more tolerant to tembotrione and isoxaflutole. Plants belonging to categories (B) and (C) have not been selected for the presence of the respective transgene prior to the herbicide application.

5

A % injury, 7DAT, 100gAI/ha

Wild Type	Line	Tembotrione	Isoxaflutole
	WT 1	100	100
	WT 2	100	100
	WT 3	100	100
	WT 4	100	98
	WT 5	100	99
	WT 6	100	99
	WT 7	100	100
	WT 8	100	n.d.
	WT 9	100	n.d.
	WT 10	100	n.d.
	WT 11	100	n.d.
	WT 12	100	n.d.
	WT 13	100	n.d.
	WT 14	100	n.d.

B % injury, 7DAT, 100gAI/ha

Arabidopsis HPPD	Line	Tembotrione	Isoxaflutole
	258 1	100	100
	258 2	100	100
	258 3	100	100

B % injury, 7DAT, 100gAI/ha

Arabidopsis HPPD	Line	Tembotrione	Isoxaflutole
	258 4	100	100
	258 5	100	100
	258 6	30	100
	252 1	30	30
	252 2	40	70
	252 3	40	95
	252 4	40	98
	252 5	50	98
	252 6	60	99
	252 7	60	99
	252 8	70	99
	252 9	70	99
	252 12	75	100
	252 13	75	100
	252 14	75	100
	252 15	80	100
	327 1	10	10
	327 2	20	20
	327 3	20	60
	327 4	40	60
	327 5	50	70
	327 6	50	80
	327 7	70	95
	327 8	70	98
	327 9	70	99

92

B % injury, 7DAT, 100gAI/ha

Arabidopsis HPPD	Line		Tembotrione	Isoxaflutole
	327	10	70	100
	327	11	70	100
	327	12	80	100
	327	13	80	100
	327	14	80	100
	327	15	80	100

C % injury, 7DAT, 100gAI/ha

FMP29e	Line		Tembotrione	Isoxaflutole
	115	1	30	0
	115	2	40	2
	115	3	50	5
	115	4	50	10
	115	5	50	15
	115	6	n.d.	30
	115	7	n.d.	30
	115	8	n.d.	40
	292	1	20	20
	292	2	20	30
	292	3	30	40
	292	4	40	n.d.
	292	5	40	n.d.

Response to Bicyclopyrone.

Seeds of wild type tobacco plants and T1 tobacco plants carrying the gene from

5 *Picrophilus torridus* FMP29e coding for HPPD were sown on MS media (Murashige

and Skoog 1964) supplemented with 50g/L kanamycin. After 4 weeks, rooted green plantlets were transferred to soil and grown for 3 weeks in the glasshouse as described above then sprayed with a mixture containing bicyclopyrone (100g AI/ha), ammonium sulfate and methyl ester rap seed oil. The plants were classified in two categories based on the phenotype developed in response to the herbicide seven days after the treatment. Class I was defined as plants displayed no injuries to light injuries in response to the herbicide treatment (injury: 0-30%), Class II was defined as plants displaying strong injuries to similar injuries as seen with wild type plants submitted to the same treatment (injury: 31-100%). In this case, only plants containing at least one T-DNA were exposed to the herbicidal treatment. In general, it can be seen that even the plants containing only one T-DNA insert already showed up a significant and sufficient level of tolerance to an exposure a field dose of the HPPD inhibitor herbicide bicyclopyrone.

15 Table 6:

Bicyclopyrone, 100g AI /ha				
7 DAT				
Transgene	Line	Class I	Class II	% of tolerant plant
-	WT	0	12	0
FMP29e	121	38	11	2

The plants containing the HPPD FMP29 displayed tolerance to the HPPD inhibitor herbicide bicyclopyrone.

It can be summarized from the above presented data, that the plants expressing the gene FMP29e from *Picrophilus torridus* coding for the FMP29 HPPD obtained from several independent transgenic events are highly tolerant to several HPPD inhibitor herbicides at doses applied under standard agronomic conditions.

Example 5: Construction of binary vectors to express several dicotyledoneous optimized variants in plants and glasshouse trial to evaluate tolerance of tobacco plants containing such variants

5 Cloning into pBin19 of FMP29t (SEQ ID No. 3), FMP29t-h (SEQ ID No. 15)

A gene with codon usage optimized for the expression in dicotyledoneous plants coding for the HPPD protein FMP29 were designed, and named FMP29t-h (SEQ ID No. 15) and the same gene with an additional sequences coding for an
10 OTP and for an HIS TAG at its 5' extremity called FMP29t (SEQ ID No. 3). The sequence corresponding to FMP29t-h gene was cloned using the restriction enzymes NcoI and XbaI in the previously described vector pRT100-OTP, containing a CaMV35S promoter and terminator. The resulting vector was called pRT100-OTP-FMP29t-h. The sequence corresponding to FMP29t was cloned in the previously
15 described vector pRT100 using the restriction enzymes XhoI and XbaI, and the resulting vector was called pRT100-OTP-FMP29t. The fragments corresponding to PromCaMV35S-OTP-FMP29t-h-TerCaMV35S and PromCaMV35S-OTP-HIS6-FMP29t-TerCaMV35S were subclone in the pBIN19 vector (described above) using the restriction enzyme SbfI. The binary vectors were respectively called pBin19-
20 FMP29t-h (Fig.3C) and pBin19-FMP29t (Fig.3B) and can be used for example to transform dicotyledenous plants, such as tobacco plants as described above. Sufficiently grown transformant plants are then tested for their tolerance to HPPD inhibitor herbicides, such as tembotrione. The development of the observed symptoms in response to the herbicidal treatment is evaluated and compared to the
25 response of wild type plants under the same conditions.

30 Example 6 : Cloning of gene FMP29e, FMP29t and FMP29m coding for FMP29 HPPD in a vector to transform Zea mays plants

FMP29e (SEQ ID No. 2), FMP29t (SEQ ID No. 3), FMP29m-h (SEQ ID No. 16)

a- FMP29e in pHoe6/Ac: Gene with a codon usage optimized for *E. coli*, plus at its 5' extremity a sequence coding for OTP and sequence coding for an His TAG.

5

The vector pRT100-FMP29e containing the gene coding for the HPPD FMP29, optimized for the expression in *E. coli* under the control of the CaMV35S promoter, was digested with the restriction enzyme HindIII. The

CaMV35S::OTP::FMP29e::CaMV35S-term cassette was further cloned into the
10 binary vector pHoe6/Ac (US 6,316,694) previously digested with the same restriction enzyme and dephosphorylated. The resulting vector was called pHoe6/Ac/FMP29e.

b- FMP29t in pHoe6/Ac (SEQ ID No. 3): Gene with a codon usage optimized for dicotyledoneous plants, plus at its 5' extremity a sequence coding for OTP and
15 sequence coding for an His TAG.

FMP29t in pRT100. A version of the gene coding for the protein FMP29 optimized for the expression in *Nicotiana tabaccum* plus containing at the 5' end a nucleic acid sequence encoding an optimized transit peptide and an HIS tag was ordered and called FMP29t. Upstream to this sequence was added the recognition sequence for
20 the restriction enzyme XhoI and downstream the recognition sequence for the restriction enzyme XbaI. The DNA corresponding to the OTP and FMP29t were digested with the restriction enzymes XhoI and XbaI, separated per appropriate gel electrophoresis and cloned into the vector pRT100 (Toepfer, (1987), Nucleic Acid Res 15:5890) previously digested with XhoI and NcoI restriction enzymes. The
25 plasmid pRT100 contains the CaMV35S promoter and CaMV35S terminator. The resulting vector was called pRT100-FMP29t, and digested with the restriction enzyme HindIII to separate the DNA corresponding to
CaMV35S::OTP::FMP29t::CaMV35S-term cassette from the rest of the vector, in order to clone it into the previously restricted vector pHoe6/Ac (US 6.316.694). The
30 resulting vector was called pHoe6/Ac/FMP29t (Fig.3).

c- FMP29m in pHoe6/Ac (SEQ ID No. 16): Gene with a codon usage optimized for monocotyledoneous plants plus at its 5' extremity a sequence coding for OTP.

FMP29m in pRT100-OTP (NcoI-XbaI) then HindIII

- 5 The variant of the gene optimized for the expression in monocotyledon plants coding for FMP29, called FMP29m was ordered, and upstream of the start codon was added a NcoI restriction site while downstream of the stop codon was added the recognition sequence for the restriction enzyme XbaI. The DNA sequence corresponding to FMP29m was digested with the restriction enzymes NcoI and XbaI, then separated per gel electrophoresis, finally isolated from the gel. The isolated DNA fragment was mixed with the vector pRT100-OTP (mentioned above) previously also digested with the same restriction enzymes. The resulting vector was called pRT100-OTP-FMP29m, containing the expression cassette CaMV35S::OTP::FMP29m::CaMV35Sterm, which was isolated using the restriction enzyme HindIII then further cloned into the previously opened and dephosphorylated vector pHOE6/Ac containing the gene coding for the PAT (Phosphinothricin acetyl transferase) enzyme, conferring resistance to the herbicide glufosinate (US 6,316,694). The resulting plasmid was called pHoe/Ac/FMP29m (Fig.3F)

20

Maize transformation:

The plasmids pHoe6/Ac (US 6,316,694), pHoe6/Ac/FMP29e, pHoe6/Ac/FMP29t and pHoe6/Ac/FMP29m were used to transform maize culture.

- 25 The maize culture, protoplast isolation, transformation and regeneration of fertile transgenic maize plants have been performed according to the US Patent 6284945, "Zea mays (L.) with capability of long term, highly efficient plant regeneration including fertile transgenic maize having a heterologous gene, and their preparation".

- 30 Transformed calli were selected on media containing phosphinothricin. Regenerated rooted plants were then transferred to soil, and allowed to grow and produce seeds in the glasshouse under standard conditions (28/20°C). Adult plants were grown

until seed production and seeds were collected for further sowing, sufficiently developed plants will be treated with the respective HPPD inhibitor herbicides.

5 Example 7: Construction of binary soybean transformation vectors.

A binary vector for soybean transformation is, for example, constructed with the CaMV35 promoter driving the expression of the gene FMP29t-h (SEQ ID No. 15), with a codon usage optimized for the expression in dicotyledoneous plants and at its 5'extremity was added a sequence coding for an OTP, and further upstream a
10 sequence TEV (Tobacco etch virus) to improve the stability of the mRNA in plants followed by the CaMV35S terminator. The nucleotide sequence of the gene FMP27t-h is given in SEQ ID No. 15. Additionally, the transformation vector also contains a PAT gene cassette in which the gene is driven by a CaVM35S promoter and followed by a CaMV35S terminator for glufosinate based selection during the
15 transformation process and a 2mEPSPS gene cassette in which the gene is driven by an histone promoter from Arabidopsis to confer tolerance to the herbicide glyphosate to the transformed plants (see Fig.3 H). The binary vector was called pFCO115.

20 Example 8: Soybean T0 plant establishment and selection.

Soybean transformation is achieved using methods well known in the art, such as the one described using the *Agrobacterium tumefaciens* mediated transformation soybean half-seed explants described by Paz et al. (2006, Plant cell Rep. 25:206).
25 Transformants were identified using Isoxaflutole as selection marker. The appearance of green shoots was observed, and documented as an indicator of tolerance to the herbicide isoxaflutole.
Tolerant green shoots were transferred to rooting media or grafted. Rooted plantlets were transferred to the glasshouse after an acclimation period.
30 Plants containing the transgene are then sprayed with HPPD inhibitor herbicides, as for example with tembotrione at a rate of 100g AI/ha. Ten days after the application

the symptoms due to the application of the herbicide will be evaluated and compared to the symptoms observed on a wild type plants under the same conditions.

5 Example 9: Construction of binary cotton transformation vectors.

A binary vector for cotton transformation is, for example, constructed with the CaMV35 promoter driving the expression of the gene FMP29t-h (SEQ ID No. 15), with a codon usage optimized for the expression in dicotyledoneous plants and at its
10 5'extremity was added a sequence coding for an OTP, and further upstream a sequence TEV (Tobacco etch virus) to improve the stability of the mRNA in plants followed by the CaMV35S terminator. The nucleotide sequence of the gene FMP27t-h is given in SEQ ID No. 15. Additionally, the transformation vector also contains a PAT gene cassette in which the gene is driven by a CaVM35S promoter and
15 followed by a CaMV35S terminator for glufosinate based selection during the transformation process and a 2mEPSPS gene cassette in which the gene is driven by an histone promoter from Arabidopsis to confer tolerance to the herbicide glyphosate to the transformed plants (see Fig.3 J).

Example 10: Cotton T0 plant establishment and selection.

20

Cotton transformation is achieved using methods well known in the art, especially preferred method in the one described in the PCT patent WO 00/71733.

Regenerated plants are transferred to the glasshouse. Following an acclimation
25 period, sufficiently grown plants are sprayed with HPPD inhibitor herbicides as for example tembotrione 100 gAl/ha supplemented with ammonium sulfate and methyl ester raps oil. Seven days after the spray application, the symptoms due to the treatment with the herbicide are evaluated and compared to the symptoms observed on wild type cotton plants subjected to the same treatment under the same
30 conditions.

Example 11: Construction of binary transformation vectors to generate plants tolerant to four herbicides with distinct modes of action.

5

A binary vector for dicotyledoneous plant transformation is, for example, constructed with the CaMV35 promoter driving the expression of the gene FMP29t-h (SEQ ID No. 15), with a codon usage optimized for the expression in dicotyledoneous plants and at its 5'-extremity was added a sequence coding for an OTP followed by the CaMV35S terminator. The nucleotide sequence of the gene FMP29t-h is given in SEQ ID No. 15. Additionally, the transformation vector also contains a PAT gene cassette in which the gene is driven by a CaMV35S promoter and followed by a CaMV35S terminator to confer tolerance to glufosinate to the plant expressing the gene, a 2mEPSPS gene cassette coding for the double mutant (Thr102Ile and Pro106Ser) EPSPS in which the gene is driven by an histone promoter from Arabidopsis to confer tolerance to the herbicide glyphosate to the transformed plants, and an Arabidopsis thaliana 2mAHAS gene cassette encoding a tolerant ALS enzyme (Acetolactate, Pro197Ala, Trp574Leu) driven by a CaMV35S promoter to confer tolerance to herbicides from the sulfonylurea or imidazolinone classes to the plant expressing this gene (see Fig.3 G).

The gene cassettes is finally cloned into the vector pHoe6/Ac (US 6,316,694), and the final vector is called pHoe6/FMP29t-h/PAT/EPSPS/AHAS, and is used to transform dicotyledoneous plants via *Agrobacterium tumefaciens* mediated state of the art methods. T0 plants are transferred to soil, and after an acclimation period, sufficiently grown plants are sprayed successively with an herbicide from the HPPD inhibitor class, then with glyphosate, then with glufosinate and finally with an herbicide from the sulfonylurea class for example.

30 Example 12: Generation of transgenic plants showing tolerance to herbicides of three different mode of action

A binary vector for tobacco transformation is, for example, constructed with the CaMV35 promoter driving the expression of the gene FMP29t-h (SEQ ID No. 15), with a codon usage optimized for the expression in dicotyledoneous plants and at its 5'extremity was added a sequence coding for an OTP, and further upstream a sequence TEV (Tobacco etch virus) to improve the stability of the mRNA in plants followed by the CaMV35S terminator. The nucleotide sequence of the gene FMP29t-h is given in SEQ ID No. 15. Additionally, the transformation vector also contains a PAT gene cassette in which the gene is driven by a CaVM35S promoter and followed by a CaMV35S terminator for glufosinate based selection during the transformation process and a 2mEPSPS gene cassette in which the gene is driven by an histone promoter from Arabidopsis to confer tolerance to the herbicide glyphosate to the transformed plants (see Fig.3 H). The binary vector was called pFCO115. The above vector was used to transform leaf discs obtained from *Nicotiana tobacum* plants, according to Example 3.

Transgenic tobacco plants were transferred to the greenhouse and treated with glyphosate at a rate of 1121 g AI/ha. Seeds were produced from such tolerant tobacco plants and harvested. These seeds were put on soil to germinate in the glasshouse. Three to four weeks later, 50 plantlets per event were transferred to single pots. Two weeks later, plants of a size 4-6 cm are sprayed either with :

- glufosinate-ammonium 1000 gAI/ha,
- glyphosate 1121 gAI/ha,
- tembotrione 100 g AI/ha, or
- tembotrione + glyphosate 100 g AI/ha + 1121 gAI/ha

After nine days, the symptoms caused by the respective heribcice applications are evaluated

SEQUENCE LISTING IN ELECTRONIC FORM

In accordance with Section 111(1) of the Patent Rules, this description contains a sequence listing in electronic form in ASCII text format (file: 30725-1375 Seq 19-JUN-12 v1.txt).

A copy of the sequence listing in electronic form is available from the Canadian Intellectual Property Office.

The sequences in the sequence listing in electronic form are reproduced in the following table.

SEQUENCE TABLE

<110> Bayer Intellectual Property GmbH

<120> Plants tolerant to HPPD inhibitor herbicides

<130> 30725-1375

<140> CA national phase of PCT/EP2010/070570

<141> 2010-12-22

<150> EP 09015987.2

<151> 2009-12-23

<150> US 61/290,589

<151> 2009-12-29

<150> EP 10190661.8

<151> 2010-11-10

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cagatcttca caaagcctgt tactgacagg ccaacgttct tctttgaggt catacagaga 1020
aaggggtgcaa ggtcattcgg caacggtaac tttaaggcac tttttgaggc gatagaaagg 1080
gagcaggcaa agagaggaaa cctatga 1107

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<210> 2

<211> 1128

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophlus torridus* HPPD optimized for *E. coli*, plus containing at the 5' end a nucleic acid encoding an alanine and 6 histidine amino acids

<220>

<221> misc_feature

<222> (4)..(6)

<223> sequence coding for Ala

<220>

<221> misc_feature

<222> (7)..(24)

<223> sequence encoding an His Tag containing 6 His

<400> 2

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atggcacatc accatcatca tcactatggc aagaacctga ttagcgaact gcgcgagaaa 60
gaaatcttca aacgcctgca tcatgtagag ttttatgtgt caagcgccaa aacctggtcg 120
tactttatga accgtggttt agggttcaaa acggtagcct atgcgggtcc agaaacgggc 180
attcgcgata aaatttccta cgttatgtcc caaggaactg ctgcatttc ttttacgagt 240
agcatgaacg atgacagcta tatctccaat catgtcaaaa aacatggaga tgggtgtgaaa 300
gacatcgctc ttgaggttga tgatctggat gaagcaaaat cgttgatcga gaagtacgg 360
accaaagtca gcaaaattaa cgaaattaaa gatgggaatg gtaaaattcg cactgccgaa 420
atcaaaacat atggcgaaac agtgcacact ctcatcgaaa ccggcgatta caatggtgtc 480
tttatgccgg ggtatgaaga gtctgaaatc aactcgaaga acaccggcat caagaaaatt 540
gaccacattg tgggtaacgt gtacgaagga gagatggatt catgggtaaa cttctatata 600
gagaaactgg gctttgaaca cctcattacg ttcgacgaca aggacattcg taccgattac 660
agcgctttac gcagtaagggt tgtcaaatac aacgatgaca tagtgtttcc gatcaatgaa 720
cccgcaaaag gcttgcgcaa aagccagatc gaggagtatc tggactacta tcgttcggaa 780
ggtgtgcagc atattgcgct tctgaccgat gacattatca aaaccgtgtc aatgatggaa 840
gaaaatggca ttgaattcct gaaaaccctt ggtagctact atgaaagtct gtcctctcgc 900
ataggggtcta ttgacgaaga tctcaacgag atagaaaagc acaatatcct ggttgatcgg 960
gatgaaaatg gctatttgct gcaaatcttt accaaaccgg ttacggatcg tccgacattc 1020
ttctttgagg tcattcagcg caaagggtgc cgtagttttg gcaatggcaa tttcaaagcg 1080
ctgtttgaag ccattgaacg ggaacaggcg aaacgtggca acttataa 1128

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<210> 3
 <211> 1503
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Nucleic acid sequence encoding *Picrophlus torridus* HPPD optimized for *Nicotiana tobaccum* plus containing at the 5' end a nucleic acid sequence encoding an optimized transit peptide and an HIS Tag

<220>
 <221> transit_peptide
 <222> (1)..(375)
 <223> Optimized transit peptide to chloroplasts

<220>
 <221> misc_feature
 <222> (376)..(378)
 <223> sequence encoding a Met

<220>
 <221> misc_feature
 <222> (379)..(381)
 <223> sequence encoding an Ala

<220>
 <221> misc_feature
 <222> (382)..(399)
 <223> sequence encoding an His Tag made of 6 His

<400> 3
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 atggtggctc cattcacagg cttgaaatcc aatgctgctt tcccaactac taagaaggct 120
 aacgatttct ctactctccc atctaattgtt ggaaggggtc agtgtatgca agtttggcca 180
 gcttacggaa ataagaagtt cgagactctt tcttaccttc caccactttc tatggctcca 240
 actgtgatga tggcttcttc tgctactgct gttgctccat tccaaggatt gaagtctact 300
 gcttctttgc cagttgctag aaggctcatc cgttctcttg gaaacgtttc taacgggtgga 360
 aggattagat gtgctatggc tcatcatcat caccatcact acggaaagaa ccttatttct 420
 gagcttagag agaaagagat cttcaagagg cttcatcacg ttgagttcta cgtttcttcc 480
 gctaagactt ggtcctactt catgaatagg ggactcggat tcaagactgt tgcttatgct 540
 ggaccagaaa ctggaatcag ggataagatc tcctacgtta tgtctcaagg tactgctagg 600
 atttctttca cttcctccat gaacgatgat tcctacattt ccaaccacgt taagaaacac 660
 ggtgatgggtg ttaaggatat cgctctcgaa gtggatgate ttgatgaggc taagtctctc 720
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 attaggactg ctgagatcaa gacttacggt gaaactgtgc acactcttat cgagactggt 840
 gattacaacg gtgttttcat gccaggatac gaagagtctg agatcaactc caagaacact 900
 ggtatcaaaa aaatcgatca cattgtggga aatgtttacg aggggtgaaat ggattcttgg 960
 gtgaacttct acattgagaa gttgggattc gagcacctta tcactttcga tgataaggat 1020
 atcaggactg attactctgc tcttaggtct aaggtggtga agtacaacga tgatatcgtg 1080
 ttccctatta acgaaccagc taagggactt aggaagtccc aaatcgaaga gtacctcgat 1140
 tattaccgtt ctgagggtgt tcaacacatt gctttgctca cagacgatat catcaagact 1200
 gtgtccatga tggaagagaa cggaattgag ttctttaaga ctccaggatc ttactacgag 1260
 tctttgtcct ctaggattgg atctatcgat gaggatctca acgaaatcga gaagcacaac 1320
 attcttgtgg atagggatga gaacggatac cttctccaga ttttactaa gccagtgact 1380
 gataggccaa cattcttctt cgaagtgatc caaagaaagg gtgctagatc ttccggaaac 1440

100d

ggaaacttca aggtctctttt cgaggctatt gagagagaac aagctaagag gggaaacctt 1500
tga 1503

<210> 4
<211> 368
<212> PRT
<213> *Picrophilus torridus*

<400> 4

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			20					25					30		
Ser	Tyr	Phe	Met	Asn	Arg	Gly	Leu	Gly	Phe	Lys	Thr	Val	Ala	Tyr	Ala
		35					40					45			
Gly	Pro	Glu	Thr	Gly	Ile	Arg	Asp	Lys	Ile	Ser	Tyr	Val	Met	Ser	Gln
	50					55					60				
Gly	Thr	Ala	Arg	Ile	Ser	Phe	Thr	Ser	Ser	Met	Asn	Asp	Asp	Ser	Tyr
65					70					75					80
Ile	Ser	Asn	His	Val	Lys	Lys	His	Gly	Asp	Gly	Val	Lys	Asp	Ile	Ala
			85						90					95	
Leu	Glu	Val	Asp	Asp	Leu	Asp	Glu	Ala	Lys	Ser	Leu	Ile	Glu	Lys	Tyr
			100					105					110		
Gly	Thr	Lys	Val	Ser	Lys	Ile	Asn	Glu	Ile	Lys	Asp	Gly	Asn	Gly	Lys
		115					120					125			
Ile	Arg	Thr	Ala	Glu	Ile	Lys	Thr	Tyr	Gly	Glu	Thr	Val	His	Thr	Leu
	130					135						140			
Ile	Glu	Thr	Gly	Asp	Tyr	Asn	Gly	Val	Phe	Met	Pro	Gly	Tyr	Glu	Glu
145					150				155						160
Ser	Glu	Ile	Asn	Ser	Lys	Asn	Thr	Gly	Ile	Lys	Lys	Ile	Asp	His	Ile
			165					170						175	
Val	Gly	Asn	Val	Tyr	Glu	Gly	Glu	Met	Asp	Ser	Trp	Val	Asn	Phe	Tyr
		180						185					190		
Ile	Glu	Lys	Leu	Gly	Phe	Glu	His	Leu	Ile	Thr	Phe	Asp	Asp	Lys	Asp
		195					200					205			
Ile	Arg	Thr	Asp	Tyr	Ser	Ala	Leu	Arg	Ser	Lys	Val	Val	Lys	Tyr	Asn
	210					215					220				
Asp	Asp	Ile	Val	Phe	Pro	Ile	Asn	Glu	Pro	Ala	Lys	Gly	Leu	Arg	Lys
225					230				235						240
Ser	Gln	Ile	Glu	Glu	Tyr	Leu	Asp	Tyr	Tyr	Arg	Ser	Glu	Gly	Val	Gln
			245					250						255	
His	Ile	Ala	Leu	Leu	Thr	Asp	Asp	Ile	Ile	Lys	Thr	Val	Ser	Met	Met
		260						265					270		
Glu	Glu	Asn	Gly	Ile	Glu	Phe	Leu	Lys	Thr	Pro	Gly	Ser	Tyr	Tyr	Glu
		275					280					285			
Ser	Leu	Ser	Ser	Arg	Ile	Gly	Ser	Ile	Asp	Glu	Asp	Leu	Asn	Glu	Ile
	290					295					300				
Glu	Lys	His	Asn	Ile	Leu	Val	Asp	Arg	Asp	Glu	Asn	Gly	Tyr	Leu	Leu
305					310					315					320
Gln	Ile	Phe	Thr	Lys	Pro	Val	Thr	Asp	Arg	Pro	Thr	Phe	Phe	Phe	Glu
			325					330						335	
Val	Ile	Gln	Arg	Lys	Gly	Ala	Arg	Ser	Phe	Gly	Asn	Gly	Asn	Phe	Lys
			340					345				350			
Ala	Leu	Phe	Glu	Ala	Ile	Glu	Arg	Glu	Gln	Ala	Lys	Arg	Gly	Asn	Leu
	355						360					365			

100e

<210> 5
 <211> 375
 <212> PRT
 <213> Artificial sequence

<220>
 <223> Protein encoded by SEQ ID No. 2

<220>
 <221> MISC_FEATURE
 <222> (2)..(2)
 <223> Ala

<220>
 <221> MISC_FEATURE
 <222> (3)..(8)
 <223> His Tag made 6 His

<400> 5
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 1 5 10 15
 Leu Arg Glu Lys Glu Ile Phe Lys Arg Leu His His Val Glu Phe Tyr
 20 25 30
 Val Ser Ser Ala Lys Thr Trp Ser Tyr Phe Met Asn Arg Gly Leu Gly
 35 40 45
 Phe Lys Thr Val Ala Tyr Ala Gly Pro Glu Thr Gly Ile Arg Asp Lys
 50 55 60
 Ile Ser Tyr Val Met Ser Gln Gly Thr Ala Arg Ile Ser Phe Thr Ser
 65 70 75 80
 Ser Met Asn Asp Asp Ser Tyr Ile Ser Asn His Val Lys Lys His Gly
 85 90 95
 Asp Gly Val Lys Asp Ile Ala Leu Glu Val Asp Asp Leu Asp Glu Ala
 100 105 110
 Lys Ser Leu Ile Glu Lys Tyr Gly Thr Lys Val Ser Lys Ile Asn Glu
 115 120 125
 Ile Lys Asp Gly Asn Gly Lys Ile Arg Thr Ala Glu Ile Lys Thr Tyr
 130 135 140
 Gly Glu Thr Val His Thr Leu Ile Glu Thr Gly Asp Tyr Asn Gly Val
 145 150 155 160
 Phe Met Pro Gly Tyr Glu Glu Ser Glu Ile Asn Ser Lys Asn Thr Gly
 165 170 175
 Ile Lys Lys Ile Asp His Ile Val Gly Asn Val Tyr Glu Gly Glu Met
 180 185 190
 Asp Ser Trp Val Asn Phe Tyr Ile Glu Lys Leu Gly Phe Glu His Leu
 195 200 205
 Ile Thr Phe Asp Asp Lys Asp Ile Arg Thr Asp Tyr Ser Ala Leu Arg
 210 215 220
 Ser Lys Val Val Lys Tyr Asn Asp Asp Ile Val Phe Pro Ile Asn Glu
 225 230 235 240
 Pro Ala Lys Gly Leu Arg Lys Ser Gln Ile Glu Glu Tyr Leu Asp Tyr
 245 250 255
 Tyr Arg Ser Glu Gly Val Gln His Ile Ala Leu Leu Thr Asp Asp Ile
 260 265 270
 Ile Lys Thr Val Ser Met Met Glu Glu Asn Gly Ile Glu Phe Leu Lys
 275 280 285


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Thr Pro Gly Ser Tyr Tyr Glu Ser Leu Ser Ser Arg Ile Gly Ser Ile
 290                               295                               300
Asp Glu Asp Leu Asn Glu Ile Glu Lys His Asn Ile Leu Val Asp Arg
305                               310                               315                               320
Asp Glu Asn Gly Tyr Leu Leu Gln Ile Phe Thr Lys Pro Val Thr Asp
                               325                               330                               335
Arg Pro Thr Phe Phe Phe Glu Val Ile Gln Arg Lys Gly Ala Arg Ser
                               340                               345                               350
Phe Gly Asn Gly Asn Phe Lys Ala Leu Phe Glu Ala Ile Glu Arg Glu
 355                               360                               365
Gln Ala Lys Arg Gly Asn Leu
 370                               375

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<210> 6

<211> 493

<212> PRT

<213> Artificial sequence

<220>

<223> Picrophlus torridus HPPD amino acid sequence (SEQ ID No. 4) fused with an optimized transit peptide (WO 2009/144079))

<220>

<221> TRANSIT

<222> (1)..(125)

<223> Optimized transit peoptide to chloroplasts

<400> 6

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Met Ala Ser Ile Ser Ser Ser Val Ala Thr Val Ser Arg Thr Ala Pro
1                               5                               10                               15
Ala Gln Ala Asn Met Val Ala Pro Phe Thr Gly Leu Lys Ser Asn Ala
                               20                               25                               30
Ala Phe Pro Thr Thr Lys Lys Ala Asn Asp Phe Ser Thr Leu Pro Ser
                               35                               40                               45
Asn Gly Gly Arg Val Gln Cys Met Gln Val Trp Pro Ala Tyr Gly Asn
 50                               55                               60
Lys Lys Phe Glu Thr Leu Ser Tyr Leu Pro Pro Leu Ser Met Ala Pro
65                               70                               75                               80
Thr Val Met Met Ala Ser Ser Ala Thr Ala Val Ala Pro Phe Gln Gly
                               85                               90                               95
Leu Lys Ser Thr Ala Ser Leu Pro Val Ala Arg Arg Ser Ser Arg Ser
                               100                              105                              110
Leu Gly Asn Val Ser Asn Gly Gly Arg Ile Arg Cys Ala Met Tyr Gly
                               115                              120                              125
Lys Asn Leu Ile Ser Glu Leu Arg Glu Lys Glu Ile Phe Lys Arg Leu
 130                              135                              140
His His Val Glu Phe Tyr Val Ser Ser Ala Lys Thr Trp Ser Tyr Phe
145                              150                              155                              160
Met Asn Arg Gly Leu Gly Phe Lys Thr Val Ala Tyr Ala Gly Pro Glu
                               165                               170                               175
Thr Gly Ile Arg Asp Lys Ile Ser Tyr Val Met Ser Gln Gly Thr Ala
                               180                               185                               190
Arg Ile Ser Phe Thr Ser Ser Met Asn Asp Asp Ser Tyr Ile Ser Asn
 195                               200                               205

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100g

```

His Val Lys Lys His Gly Asp Gly Val Lys Asp Ile Ala Leu Glu Val
 210                               215                               220
Asp Asp Leu Asp Glu Ala Lys Ser Leu Ile Glu Lys Tyr Gly Thr Lys
225                               230                               235                               240
Val Ser Lys Ile Asn Glu Ile Lys Asp Gly Asn Gly Lys Ile Arg Thr
                               245                               250                               255
Ala Glu Ile Lys Thr Tyr Gly Glu Thr Val His Thr Leu Ile Glu Thr
                               260                               265                               270
Gly Asp Tyr Asn Gly Val Phe Met Pro Gly Tyr Glu Glu Ser Glu Ile
                               275                               280                               285
Asn Ser Lys Asn Thr Gly Ile Lys Lys Ile Asp His Ile Val Gly Asn
                               290                               295                               300
Val Tyr Glu Gly Glu Met Asp Ser Trp Val Asn Phe Tyr Ile Glu Lys
305                               310                               315                               320
Leu Gly Phe Glu His Leu Ile Thr Phe Asp Asp Lys Asp Ile Arg Thr
                               325                               330                               335
Asp Tyr Ser Ala Leu Arg Ser Lys Val Val Lys Tyr Asn Asp Asp Ile
                               340                               345                               350
Val Phe Pro Ile Asn Glu Pro Ala Lys Gly Leu Arg Lys Ser Gln Ile
                               355                               360                               365
Glu Glu Tyr Leu Asp Tyr Tyr Arg Ser Glu Gly Val Gln His Ile Ala
370                               375                               380
Leu Leu Thr Asp Asp Ile Ile Lys Thr Val Ser Met Met Glu Glu Asn
385                               390                               395                               400
Gly Ile Glu Phe Leu Lys Thr Pro Gly Ser Tyr Tyr Glu Ser Leu Ser
                               405                               410                               415
Ser Arg Ile Gly Ser Ile Asp Glu Asp Leu Asn Glu Ile Glu Lys His
                               420                               425                               430
Asn Ile Leu Val Asp Arg Asp Glu Asn Gly Tyr Leu Leu Gln Ile Phe
                               435                               440                               445
Thr Lys Pro Val Thr Asp Arg Pro Thr Phe Phe Phe Glu Val Ile Gln
450                               455                               460
Arg Lys Gly Ala Arg Ser Phe Gly Asn Gly Asn Phe Lys Ala Leu Phe
465                               470                               475                               480
Glu Ala Ile Glu Arg Glu Gln Ala Lys Arg Gly Asn Leu
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<210> 7
<211> 500
<212> PRT
<213> Artificial sequence

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<220>
<223> Protein encoded by SEQ ID No. 3

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<220>
<221> TRANSIT
<222> (1)..(125)
<223> Optimized transit peptide to chloroplasts

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<220>
<221> MISC_FEATURE
<222> (126)..(126)
<223> Met

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100h

<220>

<221> MISC FEATURE

<222> (127)..(127)

<223> Ala

<220>

<221> MISC FEATURE

<222> (128)..(133)

<223> His Tag made of 6 His

<400> 7

Met	Ala	Ser	Ile	Ser	Ser	Ser	Val	Ala	Thr	Val	Ser	Arg	Thr	Ala	Pro
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Ala	Gln	Ala	Asn	Met	Val	Ala	Pro	Phe	Thr	Gly	Leu	Lys	Ser	Asn	Ala
			20					25					30		
Ala	Phe	Pro	Thr	Thr	Lys	Lys	Ala	Asn	Asp	Phe	Ser	Thr	Leu	Pro	Ser
		35					40					45			
Asn	Gly	Gly	Arg	Val	Gln	Cys	Met	Gln	Val	Trp	Pro	Ala	Tyr	Gly	Asn
	50					55					60				
Lys	Lys	Phe	Glu	Thr	Leu	Ser	Tyr	Leu	Pro	Pro	Leu	Ser	Met	Ala	Pro
65					70				75					80	
Thr	Val	Met	Met	Ala	Ser	Ser	Ala	Thr	Ala	Val	Ala	Pro	Phe	Gln	Gly
				85					90					95	
Leu	Lys	Ser	Thr	Ala	Ser	Leu	Pro	Val	Ala	Arg	Arg	Ser	Ser	Arg	Ser
			100					105					110		
Leu	Gly	Asn	Val	Ser	Asn	Gly	Gly	Arg	Ile	Arg	Cys	Ala	Met	Ala	His
		115					120					125			
His	His	His	His	Tyr	Gly	Lys	Asn	Leu	Ile	Ser	Glu	Leu	Arg	Glu	
	130				135					140					
Lys	Glu	Ile	Phe	Lys	Arg	Leu	His	His	Val	Glu	Phe	Tyr	Val	Ser	Ser
145					150					155					160
Ala	Lys	Thr	Trp	Ser	Tyr	Phe	Met	Asn	Arg	Gly	Leu	Gly	Phe	Lys	Thr
				165					170					175	
Val	Ala	Tyr	Ala	Gly	Pro	Glu	Thr	Gly	Ile	Arg	Asp	Lys	Ile	Ser	Tyr
			180					185					190		
Val	Met	Ser	Gln	Gly	Thr	Ala	Arg	Ile	Ser	Phe	Thr	Ser	Ser	Met	Asn
		195					200					205			
Asp	Asp	Ser	Tyr	Ile	Ser	Asn	His	Val	Lys	Lys	His	Gly	Asp	Gly	Val
	210					215					220				
Lys	Asp	Ile	Ala	Leu	Glu	Val	Asp	Asp	Leu	Asp	Glu	Ala	Lys	Ser	Leu
225					230					235					240
Ile	Glu	Lys	Tyr	Gly	Thr	Lys	Val	Ser	Lys	Ile	Asn	Glu	Ile	Lys	Asp
				245					250					255	
Gly	Asn	Gly	Lys	Ile	Arg	Thr	Ala	Glu	Ile	Lys	Thr	Tyr	Gly	Glu	Thr
			260					265					270		
Val	His	Thr	Leu	Ile	Glu	Thr	Gly	Asp	Tyr	Asn	Gly	Val	Phe	Met	Pro
		275					280					285			
Gly	Tyr	Glu	Glu	Ser	Glu	Ile	Asn	Ser	Lys	Asn	Thr	Gly	Ile	Lys	Lys
	290					295					300				
Ile	Asp	His	Ile	Val	Gly	Asn	Val	Tyr	Glu	Gly	Glu	Met	Asp	Ser	Trp
305					310					315					320
Val	Asn	Phe	Tyr	Ile	Glu	Lys	Leu	Gly	Phe	Glu	His	Leu	Ile	Thr	Phe
				325					330					335	
Asp	Asp	Lys	Asp	Ile	Arg	Thr	Asp	Tyr	Ser	Ala	Leu	Arg	Ser	Lys	Val
			340					345					350		

Val Lys Tyr Asn Asp Asp Ile Val Phe Pro Ile Asn Glu Pro Ala Lys
 355 360 365
 Gly Leu Arg Lys Ser Gln Ile Glu Glu Tyr Leu Asp Tyr Tyr Arg Ser
 370 375 380
 Glu Gly Val Gln His Ile Ala Leu Leu Thr Asp Asp Ile Ile Lys Thr
 385 390 395 400
 Val Ser Met Met Glu Glu Asn Gly Ile Glu Phe Leu Lys Thr Pro Gly
 405 410 415
 Ser Tyr Tyr Glu Ser Leu Ser Ser Arg Ile Gly Ser Ile Asp Glu Asp
 420 425 430
 Leu Asn Glu Ile Glu Lys His Asn Ile Leu Val Asp Arg Asp Glu Asn
 435 440 445
 Gly Tyr Leu Leu Gln Ile Phe Thr Lys Pro Val Thr Asp Arg Pro Thr
 450 455 460
 Phe Phe Phe Glu Val Ile Gln Arg Lys Gly Ala Arg Ser Phe Gly Asn
 465 470 475 480
 Gly Asn Phe Lys Ala Leu Phe Glu Ala Ile Glu Arg Glu Gln Ala Lys
 485 490 495
 Arg Gly Asn Leu
 500

<210> 8
 <211> 1422
 <212> DNA
 <213> Arabidopsis thaliana

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 aaccatgatg acggcgctgc gtcgtcgccg ggattcaagc tcgtcggatt ttccaagttc 180
 gtaagaaaga atccaaagtc tgataaattc aagggttaagc gcttccatca catcgagttc 240
 tgggtgcggcg acgcaaccaa cgtcgctcgt cgcttctcct ggggtctggg gatgagattc 300
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 ggtgacctcc gattcctttt cactgctcct tactctcctg ctctctccgc cggagagatt 420
 aaaccgacaa ccacagcttc tatcccaagt ttcgatcacg gctcttgctg ttctttcttc 480
 tcttcacatg gtctcggtgt tagagccgtt gcgattgaag tagaagacgc agagtcagct 540
 ttctccatca gtgtagctaa tggcgctatt ccttcgtcgc ctcttatcgt cctcaatgaa 600
 gcagttacga tcgttgaggt taaactatac ggcgatgttg ttctccgata tgtagttac 660
 aaagcagaag ataccgaaaa atccgaattc ttgccagggt tcgagcgtgt agaggatgcg 720
 tcgtcgttcc cattggatta tggatatccg cggttgacc acgccgtggg aaacgttcct 780
 gagcttggtc cggctttaac ttatgtagcg gggttcactg gttttcacca attcgcagag 840
 ttcacagcag acgacgttgg aaccgccgag agcggtttta attcagcgtt cctggctagc 900
 aatgatgaaa tggttcttct accgattaac gagccagtgc acggaacaaa gaggaagagt 960
 cagattcaga cgtatgttga acataacgaa ggccgagggc tacaacatct ggctctgatg 1020
 agtgaagaca tattcaggac cctgagagag atgaggaaga ggagcagtat tggaggattc 1080
 gacttcatgc cttctcctcc gctacttac taccagaatc tcaagaaacg ggtcggcgac 1140
 gtgctcagcg atgatcagat caaggagtgt gaggaattag ggattcttgt agacagagat 1200
 gatcaaggga cgttgcttca aatcttcaca aaaccactag gtgacaggcc gacgatattt 1260
 atagagataa tccagagagt aggatgcatg atgaaagatg aggaaggga ggcttaccag 1320
 agtggaggat gtggtggttt tggcaaaggc aatttctctg agctcttcaa gtccattgaa 1380
 gaatacgaaa agactcttga agccaaacag ttagtgggat ga 1422

<210> 9
 <211> 445

<212> PRT

<213> Arabidopsis thaliana

<400> 9

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			20					25					30		
Val	Arg	Lys	Asn	Pro	Lys	Ser	Asp	Lys	Phe	Lys	Val	Lys	Arg	Phe	His
		35					40					45			
His	Ile	Glu	Phe	Trp	Cys	Gly	Asp	Ala	Thr	Asn	Val	Ala	Arg	Arg	Phe
	50					55					60				
Ser	Trp	Gly	Leu	Gly	Met	Arg	Phe	Ser	Ala	Lys	Ser	Asp	Leu	Ser	Thr
65					70					75					80
Gly	Asn	Met	Val	His	Ala	Ser	Tyr	Leu	Leu	Thr	Ser	Gly	Asp	Leu	Arg
			85						90					95	
Phe	Leu	Phe	Thr	Ala	Pro	Tyr	Ser	Pro	Ser	Leu	Ser	Ala	Gly	Glu	Ile
			100					105					110		
Lys	Pro	Thr	Thr	Thr	Ala	Ser	Ile	Pro	Ser	Phe	Asp	His	Gly	Ser	Cys
		115					120					125			
Arg	Ser	Phe	Phe	Ser	Ser	His	Gly	Leu	Gly	Val	Arg	Ala	Val	Ala	Ile
	130					135					140				
Glu	Val	Glu	Asp	Ala	Glu	Ser	Ala	Phe	Ser	Ile	Ser	Val	Ala	Asn	Gly
145					150					155					160
Ala	Ile	Pro	Ser	Ser	Pro	Pro	Ile	Val	Leu	Asn	Glu	Ala	Val	Thr	Ile
				165					170					175	
Ala	Glu	Val	Lys	Leu	Tyr	Gly	Asp	Val	Val	Leu	Arg	Tyr	Val	Ser	Tyr
			180					185					190		
Lys	Ala	Glu	Asp	Thr	Glu	Lys	Ser	Glu	Phe	Leu	Pro	Gly	Phe	Glu	Arg
	195						200					205			
Val	Glu	Asp	Ala	Ser	Ser	Phe	Pro	Leu	Asp	Tyr	Gly	Ile	Arg	Arg	Leu
	210					215					220				
Asp	His	Ala	Val	Gly	Asn	Val	Pro	Glu	Leu	Gly	Pro	Ala	Leu	Thr	Tyr
225					230					235					240
Val	Ala	Gly	Phe	Thr	Gly	Phe	His	Gln	Phe	Ala	Glu	Phe	Thr	Ala	Asp
				245					250					255	
Asp	Val	Gly	Thr	Ala	Glu	Ser	Gly	Leu	Asn	Ser	Ala	Val	Leu	Ala	Ser
			260					265					270		
Asn	Asp	Glu	Met	Val	Leu	Leu	Pro	Ile	Asn	Glu	Pro	Val	His	Gly	Thr
		275					280					285			
Lys	Arg	Lys	Ser	Gln	Ile	Gln	Thr	Tyr	Leu	Glu	His	Asn	Glu	Gly	Ala
	290					295						300			
Gly	Leu	Gln	His	Leu	Ala	Leu	Met	Ser	Glu	Asp	Ile	Phe	Arg	Thr	Leu
305					310					315					320
Arg	Glu	Met	Arg	Lys	Arg	Ser	Ser	Ile	Gly	Gly	Phe	Asp	Phe	Met	Pro
				325					330					335	
Ser	Pro	Pro	Pro	Thr	Tyr	Tyr	Gln	Asn	Leu	Lys	Lys	Arg	Val	Gly	Asp
				340				345					350		
Val	Leu	Ser	Asp	Asp	Gln	Ile	Lys	Glu	Cys	Glu	Glu	Leu	Gly	Ile	Leu
		355					360					365			
Val	Asp	Arg	Asp	Asp	Gln	Gly	Thr	Leu	Leu	Gln	Ile	Phe	Thr	Lys	Pro
	370					375						380			
Leu	Gly	Asp	Arg	Pro	Thr	Ile	Phe	Ile	Glu	Ile	Ile	Gln	Arg	Val	Gly
385					390					395					400
Cys	Met	Met	Lys	Asp	Glu	Glu	Gly	Lys	Ala	Tyr	Gln	Ser	Gly	Gly	Cys
				405					410					415	

100k

Gly Gly Phe Gly Lys Gly Asn Phe Ser Glu Leu Phe Lys Ser Ile Glu
 420 425 430
 Glu Tyr Glu Lys Thr Leu Glu Ala Lys Gln Leu Val Gly
 435 440 445

<210> 10
 <211> 450
 <212> PRT
 <213> Artificial sequence

<220>
 <223> Protein encoded by SEQ ID No. 8 plus an additional alanine
 directly downstream of the initial amino acid methionine followed
 by 6 histidine amino acids

<220>
 <221> MISC_FEATURE
 <222> (2)..(2)
 <223> Ala

<220>
 <221> MISC_FEATURE
 <222> (3)..(8)
 <223> His Tag made 6 His

<400> 10
 Met Ala His His His His His Gln Asn Ala Ala Val Ser Glu Asn
 1 5 10 15
 Gln Asn His Asp Asp Gly Ala Ala Ser Ser Pro Gly Phe Lys Leu Val
 20 25 30
 Gly Phe Ser Lys Phe Val Arg Lys Asn Pro Lys Ser Asp Lys Phe Lys
 35 40 45
 Val Lys Arg Phe His His Ile Glu Phe Trp Cys Gly Asp Ala Thr Asn
 50 55 60
 Val Ala Arg Arg Phe Ser Trp Gly Leu Gly Met Arg Phe Ser Ala Lys
 65 70 75 80
 Ser Asp Leu Ser Thr Gly Asn Met Val His Ala Ser Tyr Leu Leu Thr
 85 90 95
 Ser Gly Asp Leu Arg Phe Leu Phe Thr Ala Pro Tyr Ser Pro Ser Leu
 100 105 110
 Ser Ala Gly Glu Ile Lys Pro Thr Thr Thr Ala Ser Ile Pro Ser Phe
 115 120 125
 Asp His Gly Ser Cys Arg Ser Phe Phe Ser Ser His Gly Leu Gly Val
 130 135 140
 Arg Ala Val Ala Ile Glu Val Glu Asp Ala Glu Ser Ala Phe Ser Ile
 145 150 155 160
 Ser Val Ala Asn Gly Ala Ile Pro Ser Ser Pro Pro Ile Val Leu Asn
 165 170 175
 Glu Ala Val Thr Ile Ala Glu Val Lys Leu Tyr Gly Asp Val Val Leu
 180 185 190
 Arg Tyr Val Ser Tyr Lys Ala Glu Asp Thr Glu Lys Ser Glu Phe Leu
 195 200 205
 Pro Gly Phe Glu Arg Val Glu Asp Ala Ser Ser Phe Pro Leu Asp Tyr
 210 215 220

1001

Gly Ile Arg Arg Leu Asp His Ala Val Gly Asn Val Pro Glu Leu Gly
 225 230 235 240
 Pro Ala Leu Thr Tyr Val Ala Gly Phe Thr Gly Phe His Gln Phe Ala
 245 250 255
 Glu Phe Thr Ala Asp Asp Val Gly Thr Ala Glu Ser Gly Leu Asn Ser
 260 265 270
 Ala Val Leu Ala Ser Asn Asp Glu Met Val Leu Leu Pro Ile Asn Glu
 275 280 285
 Pro Val His Gly Thr Lys Arg Lys Ser Gln Ile Gln Thr Tyr Leu Glu
 290 295 300
 His Asn Glu Gly Ala Gly Leu Gln His Leu Ala Leu Met Ser Glu Asp
 305 310 315 320
 Ile Phe Arg Thr Leu Arg Glu Met Arg Lys Arg Ser Ser Ile Gly Gly
 325 330 335
 Phe Asp Phe Met Pro Ser Pro Pro Pro Thr Tyr Tyr Gln Asn Leu Lys
 340 345 350
 Lys Arg Val Gly Asp Val Leu Ser Asp Asp Gln Ile Lys Glu Cys Glu
 355 360 365
 Glu Leu Gly Ile Leu Val Asp Arg Asp Asp Gln Gly Thr Leu Leu Gln
 370 375 380
 Ile Phe Thr Lys Pro Leu Gly Asp Arg Pro Thr Ile Phe Ile Glu Ile
 385 390 395 400
 Ile Gln Arg Val Gly Cys Met Met Lys Asp Glu Glu Gly Lys Ala Tyr
 405 410 415
 Gln Ser Gly Gly Cys Gly Gly Phe Gly Lys Gly Asn Phe Ser Glu Leu
 420 425 430
 Phe Lys Ser Ile Glu Glu Tyr Glu Lys Thr Leu Glu Ala Lys Gln Leu
 435 440 445
 Val Gly
 450

<210> 11
 <211> 568
 <212> PRT
 <213> Artificial sequence

<220>
 <223> Protein of SEQ ID No. 9 plus the Optimized Transit Peptide
 sequence located at the N-terminal extremity of the protein

<220>
 <221> TRANSIT
 <222> (1)..(125)
 <223> Optimized transit peptide to chloroplasts

<400> 11
 Met Ala Ser Ile Ser Ser Ser Val Ala Thr Val Ser Arg Thr Ala Pro
 1 5 10 15
 Ala Gln Ala Asn Met Val Ala Pro Phe Thr Gly Leu Lys Ser Asn Ala
 20 25 30
 Ala Phe Pro Thr Thr Lys Lys Ala Asn Asp Phe Ser Thr Leu Pro Ser
 35 40 45
 Asn Gly Gly Arg Val Gln Cys Met Gln Val Trp Pro Ala Tyr Gly Asn
 50 55 60

100m

Lys	Lys	Phe	Glu	Thr	Leu	Ser	Tyr	Leu	Pro	Pro	Leu	Ser	Met	Ala	Pro
65					70					75					80
Thr	Val	Met	Met	Ala	Ser	Ser	Ala	Thr	Ala	Val	Ala	Pro	Phe	Gln	Gly
				85					90					95	
Leu	Lys	Ser	Thr	Ala	Ser	Leu	Pro	Val	Ala	Arg	Arg	Ser	Ser	Arg	Ser
			100					105					110		
Leu	Gly	Asn	Val	Ser	Asn	Gly	Gly	Arg	Ile	Arg	Cys	Ala	Met	Gln	Asn
		115					120					125			
Ala	Ala	Val	Ser	Glu	Asn	Gln	Asn	His	Asp	Asp	Gly	Ala	Ala	Ser	Ser
	130					135					140				
Pro	Gly	Phe	Lys	Leu	Val	Gly	Phe	Ser	Lys	Phe	Val	Arg	Lys	Asn	Pro
145					150					155					160
Lys	Ser	Asp	Lys	Phe	Lys	Val	Lys	Arg	Phe	His	His	Ile	Glu	Phe	Trp
				165					170					175	
Cys	Gly	Asp	Ala	Thr	Asn	Val	Ala	Arg	Arg	Phe	Ser	Trp	Gly	Leu	Gly
			180					185					190		
Met	Arg	Phe	Ser	Ala	Lys	Ser	Asp	Leu	Ser	Thr	Gly	Asn	Met	Val	His
		195					200					205			
Ala	Ser	Tyr	Leu	Leu	Thr	Ser	Gly	Asp	Leu	Arg	Phe	Leu	Phe	Thr	Ala
	210					215					220				
Pro	Tyr	Ser	Pro	Ser	Leu	Ser	Ala	Gly	Glu	Ile	Lys	Pro	Thr	Thr	Thr
225					230					235					240
Ala	Ser	Ile	Pro	Ser	Phe	Asp	His	Gly	Ser	Cys	Arg	Ser	Phe	Phe	Ser
				245					250					255	
Ser	His	Gly	Leu	Gly	Val	Arg	Ala	Val	Ala	Ile	Glu	Val	Glu	Asp	Ala
			260					265					270		
Glu	Ser	Ala	Phe	Ser	Ile	Ser	Val	Ala	Asn	Gly	Ala	Ile	Pro	Ser	Ser
		275					280					285			
Pro	Pro	Ile	Val	Leu	Asn	Glu	Ala	Val	Thr	Ile	Ala	Glu	Val	Lys	Leu
	290					295					300				
Tyr	Gly	Asp	Val	Val	Leu	Arg	Tyr	Val	Ser	Tyr	Lys	Ala	Glu	Asp	Thr
305					310					315					320
Glu	Lys	Ser	Glu	Phe	Leu	Pro	Gly	Phe	Glu	Arg	Val	Glu	Asp	Ala	Ser
			325					330					335		
Ser	Phe	Pro	Leu	Asp	Tyr	Gly	Ile	Arg	Arg	Leu	Asp	His	Ala	Val	Gly
			340					345					350		
Asn	Val	Pro	Glu	Leu	Gly	Pro	Ala	Leu	Thr	Tyr	Val	Ala	Gly	Phe	Thr
		355					360					365			
Gly	Phe	His	Gln	Phe	Ala	Glu	Phe	Thr	Ala	Asp	Asp	Val	Gly	Thr	Ala
	370					375					380				
Glu	Ser	Gly	Leu	Asn	Ser	Ala	Val	Leu	Ala	Ser	Asn	Asp	Glu	Met	Val
385					390					395					400
Leu	Leu	Pro	Ile	Asn	Glu	Pro	Val	His	Gly	Thr	Lys	Arg	Lys	Ser	Gln
				405					410					415	
Ile	Gln	Thr	Tyr	Leu	Glu	His	Asn	Glu	Gly	Ala	Gly	Leu	Gln	His	Leu
			420					425					430		
Ala	Leu	Met	Ser	Glu	Asp	Ile	Phe	Arg	Thr	Leu	Arg	Glu	Met	Arg	Lys
		435					440					445			
Arg	Ser	Ser	Ile	Gly	Gly	Phe	Asp	Phe	Met	Pro	Ser	Pro	Pro	Pro	Thr
	450					455					460				
Tyr	Tyr	Gln	Asn	Leu	Lys	Lys	Arg	Val	Gly	Asp	Val	Leu	Ser	Asp	Asp
465					470					475					480
Gln	Ile	Lys	Glu	Cys	Glu	Glu	Leu	Gly	Ile	Leu	Val	Asp	Arg	Asp	Asp
			485					490					495		
Gln	Gly	Thr	Leu	Leu	Gln	Ile	Phe	Thr	Lys	Pro	Leu	Gly	Asp	Arg	Pro
			500					505					510		

[illegible]

<210>	12
<211>	575
<212>	PRT
<213>	Artificial sequence

<220>
<223> Protein of SEQ ID No. 10 plus the Optimized Transit Peptide sequence directly located at the N-terminal extremity of the protein.

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<220>
<221> TRANSIT
<222> (1)..(125)
<223> Optimized transit peptide to chloroplasts
```

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<220>
<221> MISC_FEATURE
<222> (126)..(126)
<223> Met
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<220>
<221> MISC_FEATURE
<222> (127)..(127)
<223> Ala
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<220>
<221> MISC_FEATURE
<222> (128)..(133)
<223> His Tag made of 6 His
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<400>	12															
Met	Ala	Ser	Ile	Ser	Ser	Ser	Val	Ala	Thr	Val	Ser	Arg	Thr	Ala	Pro	
1				5					10					15		
Ala	Gln	Ala	Asn	Met	Val	Ala	Pro	Phe	Thr	Gly	Leu	Lys	Ser	Asn	Ala	
			20					25						30		
Ala	Phe	Pro	Thr	Thr	Lys	Lys	Ala	Asn	Asp	Phe	Ser	Thr	Leu	Pro	Ser	
		35					40					45				
Asn	Gly	Gly	Arg	Val	Gln	Cys	Met	Gln	Val	Trp	Pro	Ala	Tyr	Gly	Asn	
	50					55					60					
Lys	Lys	Phe	Glu	Thr	Leu	Ser	Tyr	Leu	Pro	Pro	Leu	Ser	Met	Ala	Pro	
65					70					75					80	
Thr	Val	Met	Met	Ala	Ser	Ser	Ala	Thr	Ala	Val	Ala	Pro	Phe	Gln	Gly	
				85					90					95		
Leu	Lys	Ser	Thr	Ala	Ser	Leu	Pro	Val	Ala	Arg	Arg	Ser	Ser	Arg	Ser	
			100					105						110		

Leu	Gly	Asn	Val	Ser	Asn	Gly	Gly	Arg	Ile	Arg	Cys	Ala	Met	Ala	His	115	120	125
His	His	His	His	His	Gln	Asn	Ala	Ala	Val	Ser	Glu	Asn	Gln	Asn	His	130	135	140
Asp	Asp	Gly	Ala	Ala	Ser	Ser	Pro	Gly	Phe	Lys	Leu	Val	Gly	Phe	Ser	145	150	155
Lys	Phe	Val	Arg	Lys	Asn	Pro	Lys	Ser	Asp	Lys	Phe	Lys	Val	Lys	Arg	165	170	175
Phe	His	His	Ile	Glu	Phe	Trp	Cys	Gly	Asp	Ala	Thr	Asn	Val	Ala	Arg	180	185	190
Arg	Phe	Ser	Trp	Gly	Leu	Gly	Met	Arg	Phe	Ser	Ala	Lys	Ser	Asp	Leu	195	200	205
Ser	Thr	Gly	Asn	Met	Val	His	Ala	Ser	Tyr	Leu	Leu	Thr	Ser	Gly	Asp	210	215	220
Leu	Arg	Phe	Leu	Phe	Thr	Ala	Pro	Tyr	Ser	Pro	Ser	Leu	Ser	Ala	Gly	225	230	235
Glu	Ile	Lys	Pro	Thr	Thr	Thr	Ala	Ser	Ile	Pro	Ser	Phe	Asp	His	Gly	245	250	255
Ser	Cys	Arg	Ser	Phe	Phe	Ser	Ser	His	Gly	Leu	Gly	Val	Arg	Ala	Val	260	265	270
Ala	Ile	Glu	Val	Glu	Asp	Ala	Glu	Ser	Ala	Phe	Ser	Ile	Ser	Val	Ala	275	280	285
Asn	Gly	Ala	Ile	Pro	Ser	Ser	Pro	Pro	Ile	Val	Leu	Asn	Glu	Ala	Val	290	295	300
Thr	Ile	Ala	Glu	Val	Lys	Leu	Tyr	Gly	Asp	Val	Val	Leu	Arg	Tyr	Val	305	310	315
Ser	Tyr	Lys	Ala	Glu	Asp	Thr	Glu	Lys	Ser	Glu	Phe	Leu	Pro	Gly	Phe	325	330	335
Glu	Arg	Val	Glu	Asp	Ala	Ser	Ser	Phe	Pro	Leu	Asp	Tyr	Gly	Ile	Arg	340	345	350
Arg	Leu	Asp	His	Ala	Val	Gly	Asn	Val	Pro	Glu	Leu	Gly	Pro	Ala	Leu	355	360	365
Thr	Tyr	Val	Ala	Gly	Phe	Thr	Gly	Phe	His	Gln	Phe	Ala	Glu	Phe	Thr	370	375	380
Ala	Asp	Asp	Val	Gly	Thr	Ala	Glu	Ser	Gly	Leu	Asn	Ser	Ala	Val	Leu	385	390	395
Ala	Ser	Asn	Asp	Glu	Met	Val	Leu	Leu	Pro	Ile	Asn	Glu	Pro	Val	His	405	410	415
Gly	Thr	Lys	Arg	Lys	Ser	Gln	Ile	Gln	Thr	Tyr	Leu	Glu	His	Asn	Glu	420	425	430
Gly	Ala	Gly	Leu	Gln	His	Leu	Ala	Leu	Met	Ser	Glu	Asp	Ile	Phe	Arg	435	440	445
Thr	Leu	Arg	Glu	Met	Arg	Lys	Arg	Ser	Ser	Ile	Gly	Gly	Phe	Asp	Phe	450	455	460
Met	Pro	Ser	Pro	Pro	Pro	Thr	Tyr	Tyr	Gln	Asn	Leu	Lys	Lys	Arg	Val	465	470	475
Gly	Asp	Val	Leu	Ser	Asp	Asp	Gln	Ile	Lys	Glu	Cys	Glu	Glu	Leu	Gly	485	490	495
Ile	Leu	Val	Asp	Arg	Asp	Asp	Gln	Gly	Thr	Leu	Leu	Gln	Ile	Phe	Thr	500	505	510
Lys	Pro	Leu	Gly	Asp	Arg	Pro	Thr	Ile	Phe	Ile	Glu	Ile	Ile	Gln	Arg	515	520	525
Val	Gly	Cys	Met	Met	Lys	Asp	Glu	Glu	Gly	Lys	Ala	Tyr	Gln	Ser	Gly	530	535	540
Gly	Cys	Gly	Gly	Phe	Gly	Lys	Gly	Asn	Phe	Ser	Glu	Leu	Phe	Lys	Ser	545	550	555
																		560

Ile Glu Glu Tyr Glu Lys Thr Leu Glu Ala Lys Gln Leu Val Gly
 565 570 575

<210> 13
 <211> 26
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Primer Xho-OTP-for

<400> 13
 ctcgagatgg ctccgatctc ctccctc

26

<210> 14
 <211> 26
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Primer NcoI-OTP-rev

<400> 14
 cccatggcgc accggattct tccgcc

26

<210> 15
 <211> 1110
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
 optimized for dicotyledoneous plants

<400> 15
 atggccttacg gaaagaacct tatttctgag cttagagaga aagagatctt caagaggctt 60
 catcacgttg agttctacgt ttcttccgct aagacttggt cctacttcat gaatagggga 120
 ctccgattca agactgttgc ttatgctgga ccagaaactg gaatcaggga taagatctcc 180
 tacgttatgt ctcaaggtag tgctaggatt tctttcactt cctccatgaa cgatgattcc 240
 tacatttcca accacgttaa gaaacacggt gatggtgtta aggatatcgc tctcgaagtg 300
 gatgatcttg atgaggctaa gtctctcatt gagaagtacg gaactaaggt gtccaagatc 360
 aacgagatca aggatggaaa cggaaagatt aggactgctg agatcaagac ttacggtgaa 420
 actgtgcaca ctcttatcga gactggtgat tacaacggtg ttttcatgcc aggatacgaa 480
 gagtctgaga tcaactccaa gaacactggt atcaaaaaaa tcgatcacat tgtgggaaat 540
 gtttacgagg gtgaaatgga ttcttgggtg aacttctaca ttgagaagtt gggattcgag 600
 caccttatca ctttcgatga taaggatata aggactgatt actctgctct taggtctaag 660
 gtggtgaagt acaacgatga tatcgtgttc cctattaacg aaccagctaa gggacttagg 720
 aagtcaccaaa tcgaagagta cctcgattat taccgttctg aggggtgttca acacattgct 780
 ttgctcacag acgatatcat caagactgtg tccatgatgg aagagaacgg aattgagttc 840
 cttaagactc caggatctta ctacgagtct ttgtcctcta ggattggatc tatcgatgag 900
 gatctcaacg aaatcgagaa gcacaacatt cttgtggata gggatgagaa cggatacctt 960
 ctccagattt tcactaagcc agtgactgat aggccaacat tcttcttcga agtgatccaa 1020

100q

agaaaggggtg ctagatcttt cggaaacgga aacttcaagg ctcttttcga ggctattgag 1080
 agagaacaag ctaagagggg aaacctttga 1110

<210> 16
 <211> 1110
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
 optimized for *Zea mays* plants

<400> 16
 atggcatatg gcaaaaattt aatctcagaa ctaagggaaa aggagatctt taaacgatta 60
 catcacgtgg aattttacgt tagcagtgcc aaaacatggg catatttcat gaacaggggt 120
 cttggattta aaacagtggc atatgccggg ccagaaaccg ggataaggga caagatatcc 180
 tatgttatgt cccagggcac tgcaaggata tcttttacat catcaatgaa tgatgatagc 240
 tatatatcga atcatgttaa aaaacacggg gatggcgtaa aggatatagc acttgaggtc 300
 gatgatctgg acgaggcaaa aagcctgata gaaaagtatg gaacaaagggt ttcaaaaata 360
 aatgaaataa aggatggaaa tggaaagata agaactgcag agataaaaaac gtacggtgaa 420
 accgttcata cattaataga aaccggggat tacaatggcg tattcatgcc cggttatgag 480
 gaatctgaaa taaattcaaa aaacactggg ataaaaaaga tcgatcatat agttggaaat 540
 gtctatgagg gcgagatgga tagctgggtt aatttttaca tagaaaaact tggctttgag 600
 catttaataa cttttgatga taaagatata agaactgatt acagcgcatt aagatcaaag 660
 gttgtaaaat acaatgacga tatcgtattt ccaataaatg agcctgcaaa gggcttaaga 720
 aaatcacaga tagaggaata tcttgactat tacaggctct agggcggttca gcacatagca 780
 ctgttaactg atgatataat aaaaactgta tccatgatgg aggaaaacgg catagaattt 840
 ttaaaaacac caggatcata ctatgaatcc ctatcatcaa ggataggctc aatagacgag 900
 gatttaaatg aaatagagaa acataacata cttgtggatc gtgatgagaa cggataccta 960
 ttacagatct tcacaaagcc tggtactgac aggccaacgt tcttctttga ggtcatacag 1020
 agaaaggggt caaggtcatt cggcaacggg aactttaagg cactttttga ggcgatagaa 1080
 agggagcagg caaagagagg aaacctatga 1110

<210> 17
 <211> 1110
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
 optimized for *Brassica napus* plants

<400> 17
 atggcttatg ggaaaaacct catctcagag cttagggaaa aggaaatatt caaaagactc 60
 catcacgtgg agttttatgt ttctctgcc aaaacttggg cttattttat gaacagaggt 120
 ctcggtttta agactgtggc ctacgccggg cctgaaacag gaattcgaga caaaatttcc 180
 tacgtcatgt cacaaggtac agctagaatt agtttcacat cctcaatgaa tgatgactcg 240
 tacataagca accatgtcaa gaagcacgga gacggagtaa aagatatcgc gttggaggta 300
 gacgatcttg acgaagctaa gagtttgata gaaaagtacg gtacgaaagt ttccaagatc 360
 aatgagatta aagatggtaa cggtaaaata cgcactgccg agatcaagac ttacggggaa 420
 acggtgcaca ctttgattga gacaggagac tacaatggag tattcatgcc cggttacgaa 480
 gaaagcgaaa ttaacagcaa gaataccggg atcaagaaga tagatcacat cgtaggaaac 540
 gtctacgaag gagagatgga ctcatgggtc aatttctata tcgagaagtt gggtttcgaa 600
 cacttaataa catttgacga caaagatata cgaacagact attccgcatt gcgttcaaaa 660

gttgtgaaat	acaacgatga	tatagtgttc	cctatcaatg	agcccgetaa	gggcctaaga	720
aagtcgcaga	ttgaagagta	cttggactac	taccgctctg	aggggggttca	gcatattgct	780
ctgttgactg	atgatattat	taagacagtc	agtatgatgg	aggagaacgg	tattgaattc	840
ttgaaaacac	cggctcgtta	ctacgaatcc	ttgtcttcaa	gaatcggaag	tatcgatgag	900
gatcttaacg	aaatcgagaa	gcacaacatc	ctgggtggatc	gtgacgagaa	cggatatctc	960
ctgcaaattct	tcacaaagcc	agtaactgac	cgccctactt	tcttttttga	agtgattcaa	1020
cgtaagggag	ctagatcctt	tggtaatggg	aacttcaaag	cgctgttcga	agctatcgaa	1080
agagagcagg	ccaagagagg	caatctgtga				1110

<210> 18

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for *Beta vulgaris* plants

<400> 18

atggccttacg	gaaaaaattt	aatcagcgaa	ctacgtgaaa	aggagatatt	caaaaggctg	60
catcatgtag	agttttacgt	atctagtgcg	aaaacttggt	catatttcat	gaaccgagga	120
ttaggattca	agacagtagc	gtatgctgga	cctgaaacag	gaatacgtga	caagatttcg	180
tacgttatga	gtcaggggtac	ggcccggaata	tctttttacca	gttccatgaa	tgacgattcg	240
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ctccaaatct	ttactaaacc	tgtgacagat	cgtccaacat	tttttttcga	ggcatccaa	1020
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<210> 19

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for *Gossypium hirsutum* plants

<400> 19

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tacgtgatgt	ctcaggggaac	cggccgcatt	tccttcaactt	cttccatgaa	cgatgattcc	240
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catctcatta ctttcgatga caaggacatt cggaccgact attctgcttt gagatctaaa 660
gttgtaaaat acaacgacga catcgtgttt cccattaacg agccggcaaa aggcttgaga 720
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<210> 20

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for Glycine max plants

<400> 20

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<210> 21

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for Hordeum vulgare plants

<400> 21

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<210> 22

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for *Oriza sativa* plants

<400> 22

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<210> 23

<211> 1110

<212> DNA

<213> Artificial sequence

<220>

<223> Nucleic acid sequence encoding *Picrophilus torridus* HPPD
optimized for *Triticum aestivum* plants

<400> 23

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CLAIMS:

1. A chimeric gene comprising a coding sequence operably-linked to a plant expressible promoter, wherein the coding sequence comprises a nucleic acid sequence which encodes a hydroxyphenylpyruvate dioxygenase (HPPD) protein
5 comprising the amino acid sequence of SEQ ID No. 4 from amino acid position 2 to amino acid position 368.
2. The chimeric gene according to claim 1 comprising, upstream to the HPPD coding sequence, a nucleic acid sequence which encodes a transit peptide active in plants so that a transit peptide/HPPD fusion protein is encoded by said chimeric gene.
- 10 3. A vector comprising at least one chimeric gene according to any one of claims 1 to 2.
4. A plant cell comprising a chimeric gene according to claim 1 or 2.
5. The plant cell of claim 4, which also comprises a chimeric gene encoding a PDH (prephenate dehydrogenase) enzyme.
- 15 6. The plant cell of claim 4 or 5 further comprising one or more chimeric gene(s) conferring tolerance to 2,4-D herbicide, dicamba herbicide, herbicide inhibiting acetolactate synthase (ALS), herbicide inhibiting EPSP synthase (EPSPS), herbicide inhibiting glutamine synthase (GS), or any combination thereof.
- 20 7. A method for obtaining a plant cell tolerant to a hydroxyphenylpyruvate dioxygenase (HPPD) inhibitor herbicide, comprising introducing into said plant cell a chimeric gene according to one of claims 1 to 2.
8. A method for controlling weeds, the method comprising applying a hydroxyphenylpyruvate dioxygenase (HPPD) inhibitor herbicide which is toxic for said weeds to a field planted with plants or seeds obtained from the plant cell as defined in
25 any one of claims 4 to 6.

9. The method for controlling weeds according to claim 8, wherein the HPPD inhibitor is selected from the group of isoxaflutole, tembotrione, mesotrione, sulcotrione, pyrasulfotole, topramezone, 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-CF₃phenyl)propane-1,3-dione and 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-2,3 Cl₂phenyl)propane-1,3-dione, bicyclopyrone, benzobicyclon, tefuryltrione, diketonitrile, and pyrazoxyfen.
10. Use of a hydroxyphenylpyruvate dioxygenase (HPPD) protein comprising the amino acid sequence of SEQ ID No. 4 to render the plant cell according to claim 4, 5, or 6 and plants or seeds comprising said cell tolerant to HPPD inhibitor herbicides.
- 10 11. The use according to claim 10, wherein the HPPD inhibitor herbicides are selected from the group consisting of: isoxaflutole, tembotrione, mesotrione, sulcotrione, pyrasulfotole, topramezone, 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-CF₃-phenyl)propane-1,3-dione and 2-cyano-3-cyclopropyl-1-(2-SO₂CH₃-4-2,3 Cl₂phenyl)propane-1,3-dione, bicyclopyrone, benzobicyclon, tefuryltrione, diketonitrile, and pyrazoxyfen.
- 15

Figures:

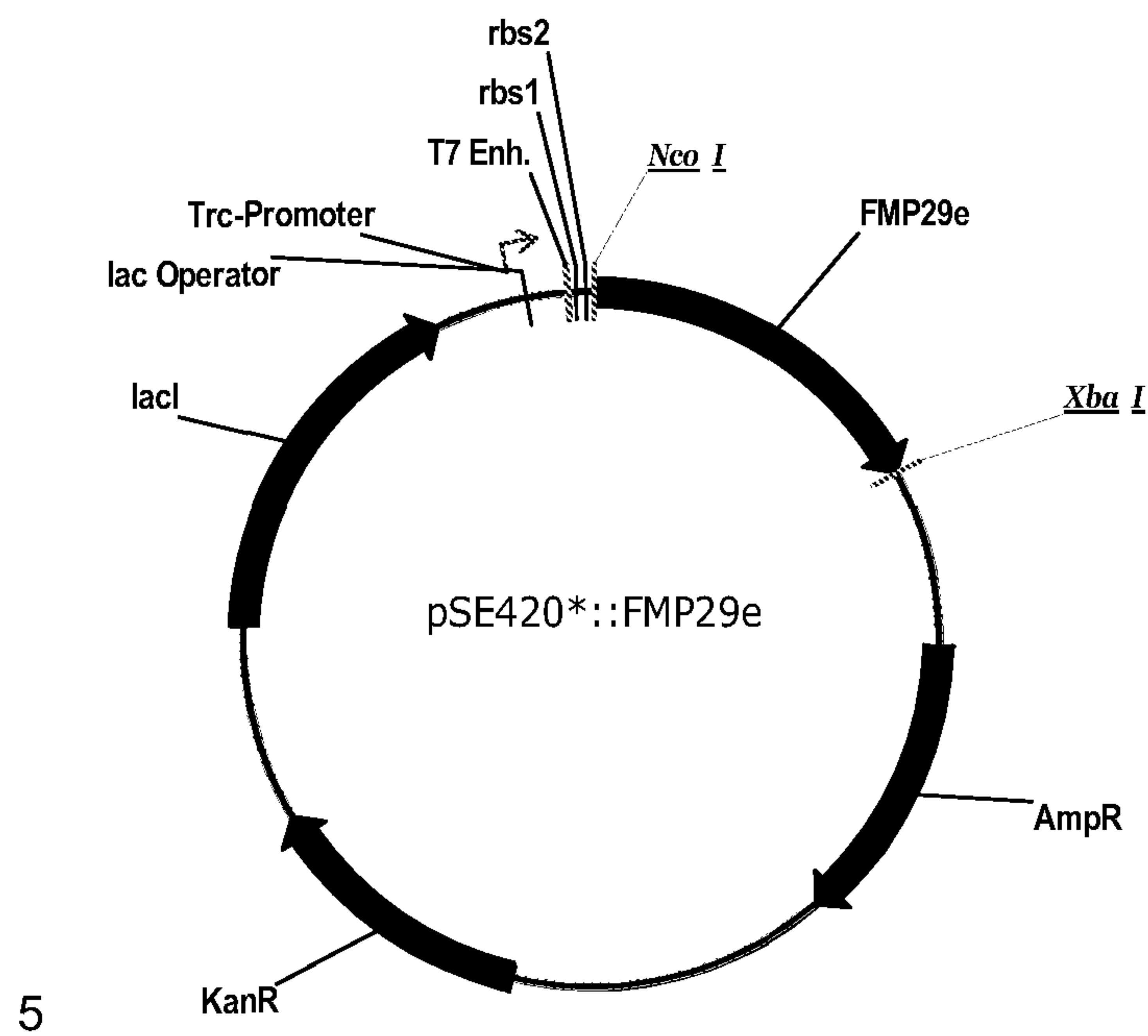
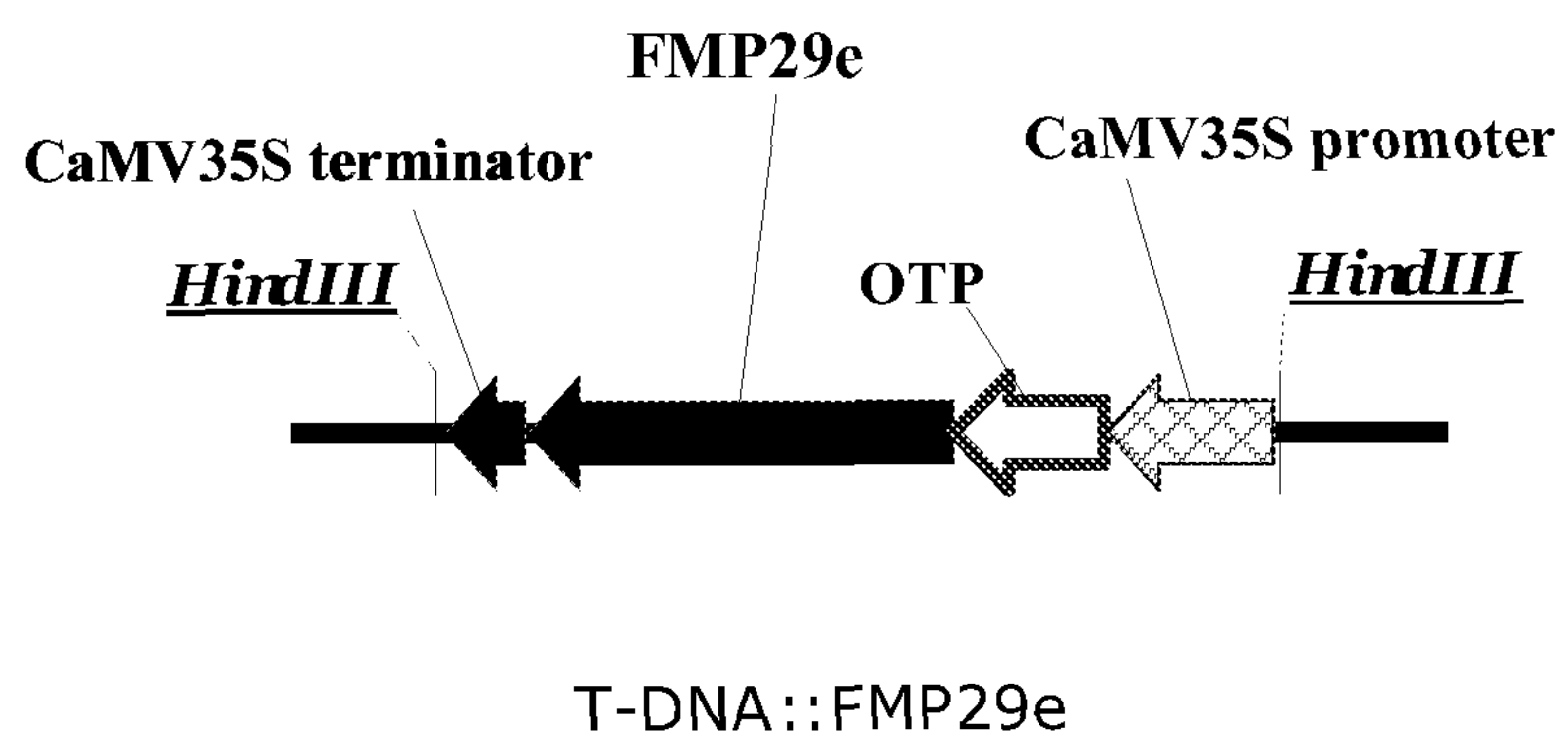


Figure1: map of the plasmid pSE420-FMP29e



10

Figure 2: map of the T-DNA inserted into the tobacco plants.

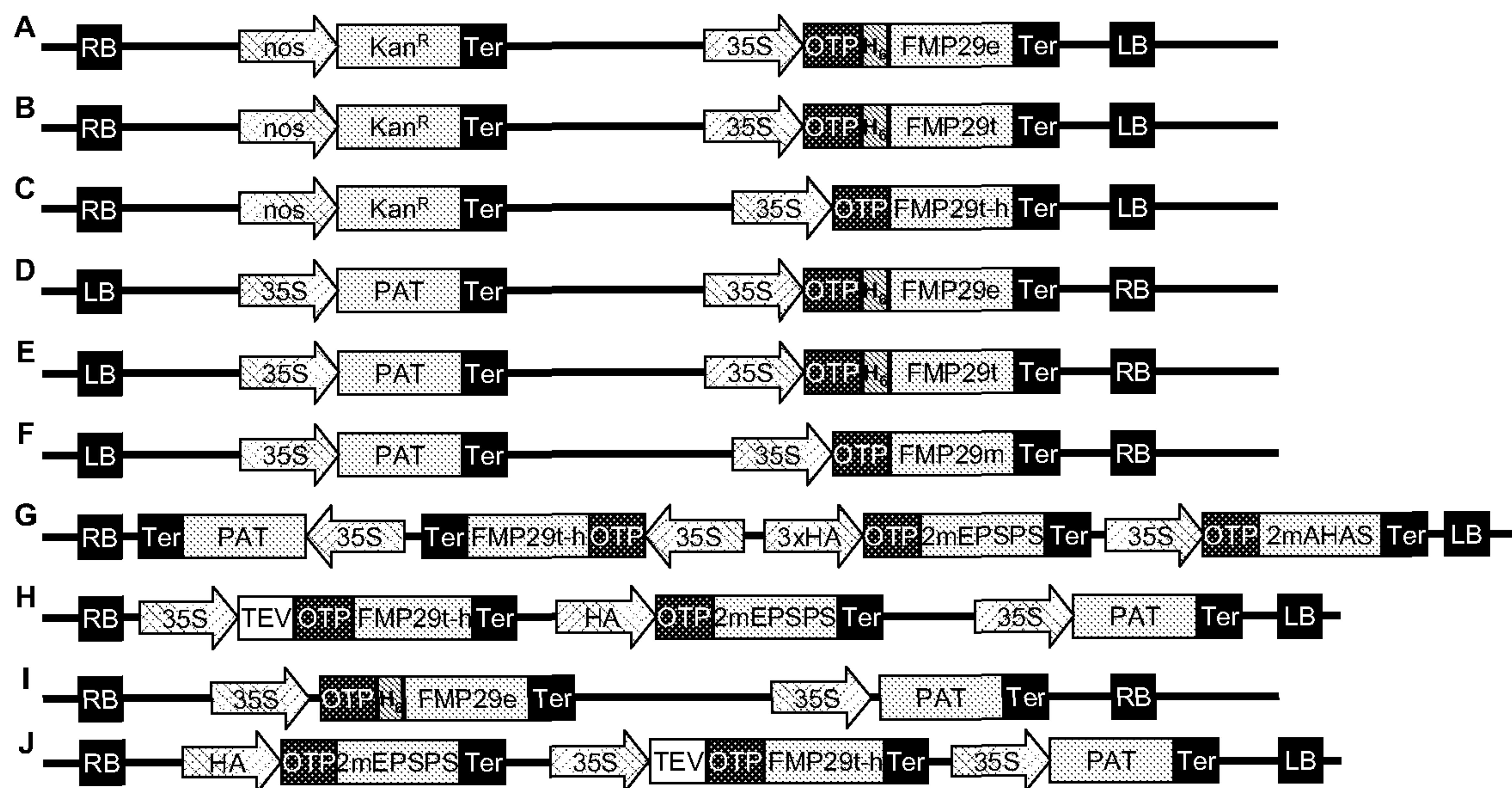
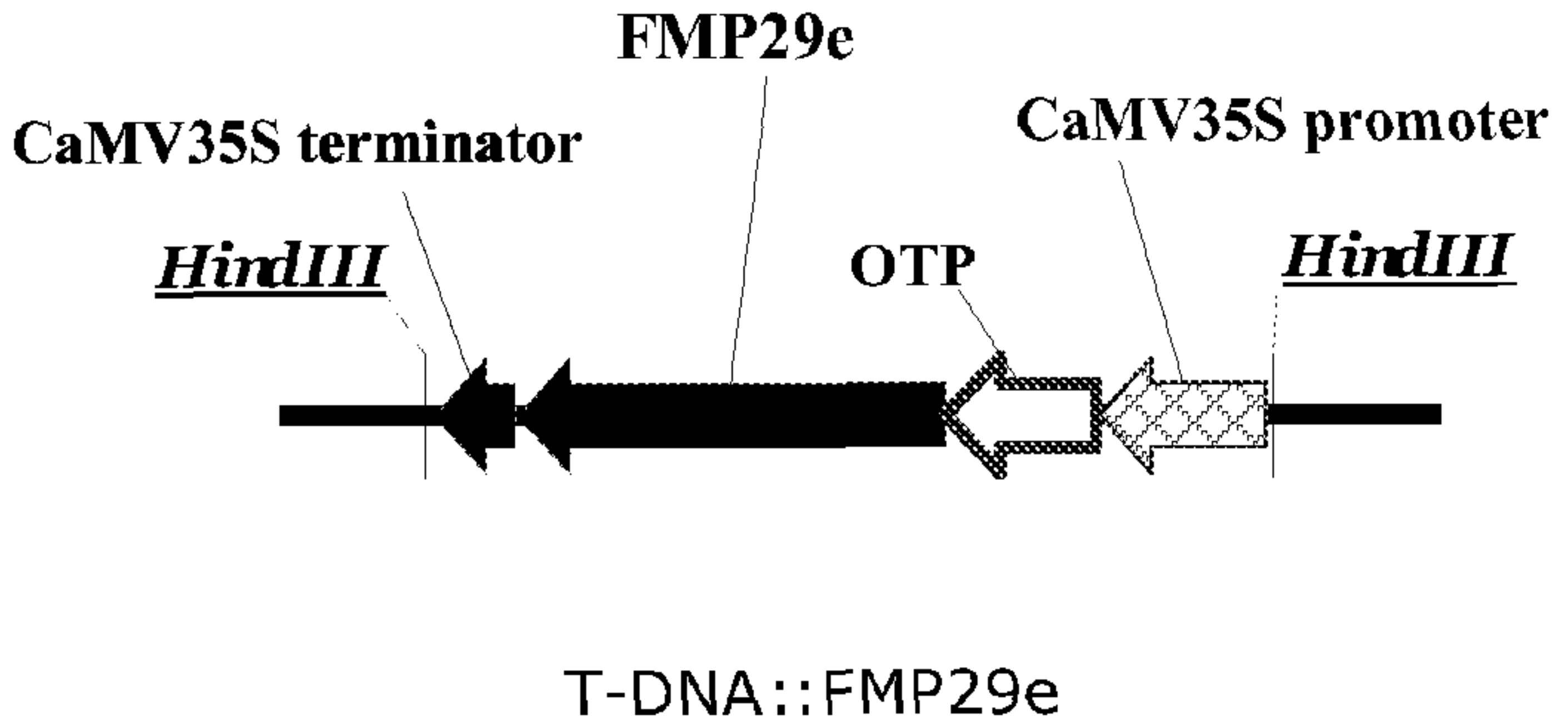


Figure 3: Map of the different T-DNA inserted into plants.



map of the T-DNA inserted into the tobacco plants.