

**FAO SPECIFICATIONS AND EVALUATIONS  
FOR AGRICULTURAL PESTICIDES**

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**AZIMSULFURON**

1-(4,6-dimethoxypyrimidin-2-yl)-3-[1-methyl-4-(2-methyl-*2H*-tetrazol-5-yl)pyrazol-5-ylsulfonyl]urea

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## DISCLAIMER<sup>1</sup>

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FAO specifications are developed with the basic objective of promoting, as far as practicable, the manufacture, distribution and use of pesticides that meet basic quality requirements.

Compliance with the specifications does not constitute an endorsement or warranty of the fitness of a particular pesticide for a particular purpose, including its suitability for the control of any given pest, or its suitability for use in a particular area. Owing to the complexity of the problems involved, the suitability of pesticides for a particular purpose and the content of the labelling instructions must be decided at the national or provincial level.

Furthermore, pesticides which are manufactured to comply with these specifications are not exempted from any safety regulation or other legal or administrative provision applicable to their manufacture, sale, transportation, storage, handling, preparation and/or use.

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Additionally, FAO wishes to alert users to the fact that improper storage, handling, preparation and/or use of pesticides can result in either a lowering or complete loss of safety and/or efficacy.

FAO is not responsible, and does not accept any liability, for the testing of pesticides for compliance with the specifications, nor for any methods recommended and/or used for testing compliance. As a result, FAO does not in any way warrant or represent that any pesticide claimed to comply with a FAO specification actually does so.

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<sup>1</sup> This disclaimer applies to all specifications published by FAO.



## INTRODUCTION

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FAO establishes and publishes specifications\* for technical material and related formulations of public health pesticides with the objective that these specifications may be used to provide an international point of reference against which products can be judged either for regulatory purposes or in commercial dealings.

Since 1999 the development of FAO specifications follows the **New Procedure**, described in the 5<sup>th</sup> edition of the “Manual on the development and use of FAO specifications for plant protection products” (FAO Plant Production and Protection Page No. 149). This **New Procedure** follows a formal and transparent evaluation process. It describes the minimum data package, the procedure and evaluation applied by FAO and the Experts of the FAO/WHO Joint Meeting on Pesticide Specifications (JMPS). [Note: prior to 2002, the Experts were of the FAO Panel of Experts on Pesticide Specifications, Registration Requirements, Application Standards and Prior Informed Consent, which now forms part of the JMPS, rather than the JMPS.]

FAO Specifications now only apply to products for which the technical materials have been evaluated. Consequently from the year 2000 onwards the publication of FAO specifications under the **New Procedure** has changed. Every specification consists now of two parts namely the specifications and the evaluation report(s):

**PART ONE: The Specification** of the technical material and the related formulations of the plant protection product in accordance with chapter 4, 5 and 6 of the 5<sup>th</sup> edition of the “Manual on the development and use of FAO specifications for plant protection products”.

**PART TWO: The Evaluation Report(s)** of the plant protection product reflecting the evaluation of the data package carried out by FAO and the JMPS. The data are to be provided by the manufacturer(s) according to the requirements of Appendix A, annex 1 or 2 of the “Manual on the development and use of FAO specifications for plant protection products” and supported by other information sources. The Evaluation Report includes the name(s) of the manufacturer(s) whose technical material has been evaluated. Evaluation reports on specifications developed subsequently to the original set of specifications are added in a chronological order to this report.

FAO specifications under the **New Procedure** do not necessarily apply to nominally similar products of other manufacturer(s), nor to those where the active ingredient is produced by other routes of manufacture. FAO has the possibility to extend the scope of the specifications to similar products but only when the JMPS has been satisfied that the additional products are equivalent to that which formed the basis of the reference specification.

**Specifications bear the date (month and year) of publication of the current version. Dates of publication of the earlier versions, if any, are identified in a footnote. Evaluations bear the date (year) of the meeting at which the recommendations were made by the JMPS.**

\* NOTE: PUBLICATIONS ARE AVAILABLE ON THE INTERNET AT  
(<http://www.fao.org/agriculture/crops/core-themes/theme/pests/jmps/en/> )  
OR IN HARDCOPY FROM THE PLANT PROTECTION INFORMATION OFFICER.

**PART ONE**

**SPECIFICATIONS**

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## AZIMSULFURON

### INFORMATION

*ISO common name*

Azimsulfuron (E-ISO)

*Synonyms*

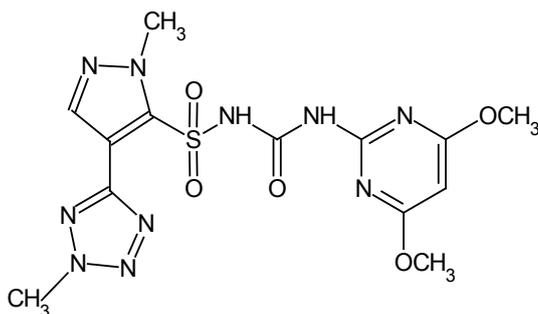
None

*Chemical names*

*IUPAC* 1-(4,6-dimethoxypyrimidin-2-yl)-3-[1-methyl-4-(2-methyl-2*H*-tetrazol-5-yl)pyrazol-5-ylsulfonyl]urea

*CA* *N*-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-1-methyl-4-(2-methyl-2*H*-tetrazol-5-yl)-1*H*-pyrazole-5-sulfonamide

*Structural formula*



*Molecular formula*

C<sub>13</sub>H<sub>16</sub>N<sub>10</sub>O<sub>5</sub>S

*Relative molecular mass*

424.40

*CAS Registry number*

120162-55-2

*CIPAC number*

584

*Identity tests*

HPLC relative retention time, IR spectrum

## AZIMSULFURON

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# TECHNICAL MATERIAL

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### FAO Specification 584/TC (February 2005\*)

*This specification, which is PART ONE of this publication, is based on an evaluation of data submitted by the manufacturer whose name is listed in the evaluation report 584/2004. It should be applicable to relevant products of these manufacturers but it is not an endorsement of those products, nor a guarantee that they comply with the specifications. The specification may not be appropriate for the products of other manufacturers. The evaluation report 584/2004, as PART TWO, forms an integral part of this publication.*

## 1 Description

The material shall consist of azimsulfuron together with related manufacturing impurities, in the form of a white crystalline solid, free from visible extraneous matter and added modifying agents.

## 2 Active ingredient

### 2.1 Identity tests (CIPAC 584/TC/M/2, Note 1)

The active ingredient shall comply with an identity test and, where the identity remains in doubt, shall comply with at least one additional test.

### 2.2 Azimsulfuron (CIPAC 584/TC/M/3, Note 1)

The azimsulfuron content shall be declared (not less than 980 g/kg) and when determined, the average measured content obtained shall not be lower than the declared minimum content.

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Note 1 Methods for the identification and determination of azimsulfuron content were adopted by CIPAC in 2004 but are not yet published in a Handbook. Prior to publication of the Handbook, copies of the methods may be obtained through the CIPAC website, <http://www.cipac.org>

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\* Specifications may be revised and/or additional evaluations may be undertaken. Ensure the use of current versions by checking at: <http://www.fao.org/agriculture/crops/core-themes/theme/pests/jmps/en/>

## AZIMSULFURON

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# WATER DISPERSIBLE GRANULES

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### FAO Specification 584/WG (February 2005\*)

*This specification, which is PART ONE of this publication, is based on an evaluation of data submitted by the manufacturer whose name is listed in the evaluation report 584/2004. It should be applicable to relevant products of these manufacturers but it is not an endorsement of those products, nor a guarantee that they comply with the specifications. The specification may not be appropriate for the products of other manufacturers. The evaluation report 584/2004, as PART TWO, forms an integral part of this publication.*

## 1 Description

The material shall consist of an homogeneous mixture of technical azimsulfuron, complying with the requirements of FAO specification 584/TC (February 2005), together with carriers and any other necessary formulants. It shall be in the form of granules for application after disintegration and dispersion in water. The formulation shall be dry, free-flowing, essentially non-dusty and free from visible extraneous matter and hard lumps.

## 2 Active Ingredient

### 2.1 Identity tests (CIPAC 584/TC/M/2, Note 1)

The active ingredient shall comply with an identity test and, where the identity remains in doubt, shall comply with at least one additional test.

### 2.2 Azimsulfuron content (CIPAC 584/TC/M/3, Note 1)

The azimsulfuron content shall be declared (g/kg) and, when determined, the average content measured shall not differ from that declared by more than the appropriate tolerance:

Declared content, g/kg	Permitted tolerance
Above 500 g/kg	± 25 g/kg

## 3 Physical Properties

### 3.1 Wettability (MT 53.3.1)

The formulation shall be completely wetted in 10 seconds, without swirling.

### 3.2 Wet sieve test (MT 185)

Maximum: 2% retained on a 75 µm test sieve.

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\* Specifications may be revised and/or additional evaluations may be undertaken. Ensure the use of current versions by checking at: <http://www.fao.org/agriculture/crops/core-themes/theme/pests/jmps/en/>

**3.3 Degree of dispersion (MT 174)**

Dispersibility : minimum 75% after 1 minute of stirring.

**3.4 Suspensibility (MT 168) (Notes 2 & 3)**

A minimum of 60% of the azimsulfuron content found under 2.2. shall be in suspension after 30 minutes in CIPAC Standard Water D at  $30 \pm 2^\circ\text{C}$ .

**3.5 Persistent foam (MT 47.2) (Note 4)**

Maximum: 60 ml after 1 minute.

**3.6 Dustiness (MT 171) (Note 5)**

Essentially non-dusty.

**3.7 Flowability (MT 172)**

At least 99% of the formulation (after the test of storage at elevated temperature) shall pass through a 5 mm test sieve after 20 drops of the sieve.

**4 Storage Stability**

**4.1 Stability at elevated temperature (MT 46.3) (Note 5)**

After storage at  $54 \pm 2^\circ\text{C}$  for 14 days, the determined average active ingredient content must not be lower than 97% relative to the determined average content found before storage and the formulation shall continue to comply with the clauses for:

- wet sieve test (4.2);
- degree of dispersion (4.3);
- suspensibility (4.4);
- dustiness (4.6).

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**Note 1** Methods for the identification and determination of azimsulfuron content were adopted by CIPAC in 2004 but are not yet published in a Handbook. Prior to publication of the Handbook, copies of the methods may be obtained through the CIPAC website, <http://www.cipac.org> or from the Secretary, Dr László Bura, Central Service for Plant Protection and Soil Conservation, Budaörsi út 141-145, 1118 Budapest, Hungary.

**Note 2** The product should be tested at the highest and lowest rates of use recommended by the supplier, provided this does not exceed the conditions given in method MT 168.

**Note 3** Chemical assay is the only fully reliable method to measure the mass of active ingredient still in suspension. However, simpler methods such as gravimetric determination or solvent extraction determination may be used on a routine basis provided, that these methods have been shown to give equal results to those of the chemical assay method. In case of dispute, the chemical method shall be the "referee method".

**Note 4** The mass of the sample to be used in the test should be specified at the highest rate of use recommended by the supplier.

**Note 5** Measurement of dustiness must be carried out on the sample "as received" and, where practicable, the sample should be taken from a newly opened container, because changes in the water content of samples may influence dustiness significantly. The optical method, MT 171, usually shows good correlation with the gravimetric method and can, therefore, be used as an alternative where the equipment is available. Where the correlation is in doubt, it must

be checked with the formulation to be tested. In case of dispute, the gravimetric method shall be used.

Note 6 Samples of the product taken before and after the storage stability test should be analysed together after the test to reduce the analytical error.

## PART TWO

### EVALUATION REPORT(S)

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#### AZIMSULFURON

<b>2004</b>	EVALUATION REPORT based on submission of data from Du Pont de Nemours & Co. (TC, WG)	<b>10</b>
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## AZIMSULFURON

# EVALUATION REPORT 584/2004

### Explanation

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The data for azimsulfuron were evaluated in support of new FAO specifications.

Azimsulfuron is under patent in Taiwan until 2004; Canada until 2005; Australia, Brazil, France, Greece, Israel, Italy, New Zealand, Portugal and Turkey until 2006; Spain, South Korea and Thailand until 2007; and Japan until 2009.

Azimsulfuron has not been evaluated by the FAO/WHO JMPR or IPCS. It was reviewed by the European Commission in 1998 and has achieved Annex 1 listing.

The draft specification and the supporting data were provided by E.I. du Pont de Nemours and Company in October 2003.

### Uses

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Azimsulfuron is a herbicide which affects sensitive weeds through inhibition of the enzyme acetolactate synthase (ALS). Inhibition of ALS leads to the cessation of cell division and subsequent growth processes in plants. Azimsulfuron is taken up mainly by leaves and shoots and, to a lesser extent, roots. Once taken up, it is translocated via both xylem and phloem.

It is used post-emergence in rice fields against a variety of annual weeds.

### Identity of the active ingredient

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#### *ISO common name*

Azimsulfuron (E-ISO)

#### *Synonyms*

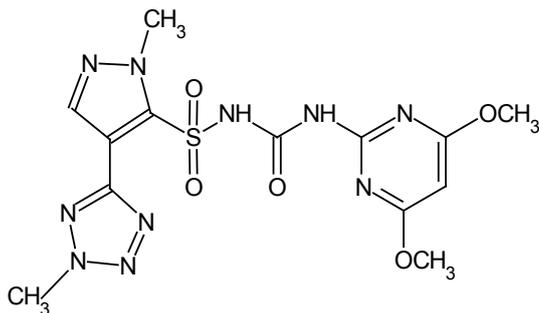
None

#### *Chemical names*

*IUPAC* 1-(4,6-dimethoxypyrimidin-2-yl)-3-[1-methyl-4-(2-methyl-2*H*-tetrazol-5-yl)pyrazol-5-ylsulfonyl]urea

*CA* *N*-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-1-methyl-4-(2-methyl-2*H*-tetrazol-5-yl)-1*H*-pyrazole-5-sulfonamide

*Structural formula*



*Molecular formula*



*Relative molecular mass*

424.40

*CAS Registry number*

120162-55-2

*CIPAC number*

584

*Identity tests*

HPLC relative retention time, IR spectrum

**Physico-chemical properties of pure azimsulfuron**

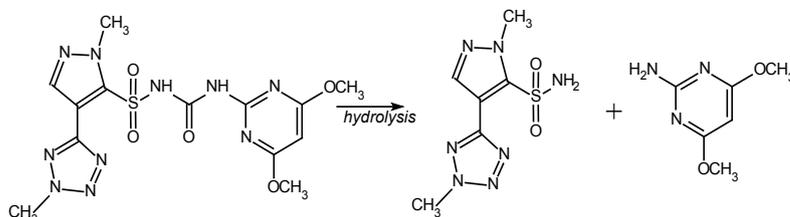
**Table 1. Physico-chemical properties of pure azimsulfuron**

Parameter	Value(s) and conditions	Purity %	Method reference	Study reference
Vapour pressure	$4 \times 10^{-9}$ Pa at 25°C (extrapolated) measurements from 86°C to 124°C	99.62%	EEC 2.3.1, Knudsen gas effusion method	Cobranchi and Schmuckler, 1994
Melting point, boiling point and/or temperature of decomposition	Melting point: 170°C Boiling point: not applicable Decomp. temp.: 180 - 250°C	99.62%	EEC 2.1, capillary, oil bath method EEC 2.12 DSC, TGA	Moore, 1994a Lesieur, 1994
Relative density	1.12 at 20°C	99.62%	EEC 2.2, OECD 109, displaced mineral oil - pycnometer method	Moore, 1994b
Solubility in water	72.3 mg/l at 20°C at pH 5 1050 mg/l at 20°C at pH 7 6536 mg/l at 20°C at pH 9	>97% <sup>a</sup>	CIPAC MT 157 Part 2, Shake flask method	Schmuckler, 1991

Parameter	Value(s) and conditions	Purity %	Method reference	Study reference
Octanol/water partition coefficient	Log K <sub>OW</sub> = 0.646 at 25°C at pH 5 Log K <sub>OW</sub> = -1.37 at 25°C at pH 7 Log K <sub>OW</sub> = -2.08 at 25°C at pH 9	96.95% <sup>a</sup>	Shake flask method, OTS Guideline CG1400	Smyser, 1991
Volatility	Henry's Law Constants (calculated) (Pa x m <sup>3</sup> x mol <sup>-1</sup> ) 8 x 10 <sup>-9</sup> at 20°C at pH5 5 x 10 <sup>-10</sup> at 20°C at pH7 9 x 10 <sup>-11</sup> at 20°C at pH9	Not applicable	Calculated	Schmuckler, 1995
Hydrolysis characteristics <sup>b</sup>	Half-life = 89 days (mean of 83 and 95) at 25°C at pH 5 Half-life = 124 days (mean of 117 and 132) at 25°C at pH 7 Half-life = 132 days (mean of 126 and 139) at 25°C at pH 9	Radiochem purity 97.4% (pyrazole label); 97.3% (pyrimidine label)	U.S. EPA Pesticide Assessment Guidelines (Hitch, 1982a)	Hausmann, 1991a
Photolysis characteristics <sup>c</sup>	Simulated sunlight: Half-life = 103 days at pH 5 Half-life = 164 days at pH 7 Half-life = 225 days at pH 9	see Hausmann, 1991b	US EPA model GCSOLAR (EPA, 1985)	Barefoot, 1998
Dissociation characteristics	pKa = 3.6	96.95% <sup>a</sup>	OECD 112, spectrophotometric method	Cooke, 1993

<sup>a</sup> Tests were conducted between 1989 and 1992. Process was optimized in 1994 to yield technical of ≥98% purity.

<sup>b</sup> Hydrolysis test in dark and sterile conditions at a concentration of 25 mg/l for 30 days. Hydrolysis cleaved the sulfonylurea bridge.



<sup>c</sup> Photolysis test in sterile conditions at a concentration of 25 mg/l for 30 days.

## Chemical composition and properties of azimsulfuron technical material (TC)

**Table 2. Chemical composition and properties of azimsulfuron technical material (TC)**

Manufacturing process, maximum limits for impurities ≥1 g/kg, 5 batch analysis data	Confidential information supplied and held on file by FAO. Mass balances were 100.3-101.4%.
Declared minimum azimsulfuron content	980 g/kg.
Relevant impurities ≥1 g/kg and maximum limits for them	None.
Relevant impurities <1 g/kg and maximum limits for them:	None
Stabilizers or other additives and maximum limits for them:	None.
Melting temperature range of the TC	170°C (decomposition occurs at 180-250°C).

### Toxicological summaries

Notes.

- (i) The proposer confirmed that the toxicological and ecotoxicological data included in the summary below were derived from azimsulfuron having impurity profiles similar to those referred to in the table above.
- (ii) The conclusions expressed in the summary below are those of the proposer, unless otherwise specified.

**Table 3. Toxicology profile of azimsulfuron technical material, based on acute toxicity, irritation and sensitization**

Species	Test	Duration and conditions or guideline adopted	Result	Study ref
Male and female rat	Acute oral	14 days Azimsulfuron technical (97.9% purity) <sup>a,b</sup>	LD <sub>50</sub> >5000 mg/kg bw	Ueda, 1990c
Male and female rat	Acute dermal	14 days Azimsulfuron technical (97.9% purity) <sup>a,b</sup> MAFF Japan, 1985	LD <sub>50</sub> = >2000 mg/kg bw	Ueda, 1990a
Male and female rat Fischer (F344/DuCrj)	Acute inhalation	4 hours Azimsulfuron technical (99.1% purity) <sup>a</sup> OECD 403	LC <sub>50</sub> > 5.94 mg/m <sup>3</sup>	Ebino, 1993
Female rabbit (New Zealand white)	Acute skin irritation	72 hours Azimsulfuron technical (97.9% purity) <sup>a,b</sup> Draize, 1965	Non-irritant	Kosaka, 1990a
Female rabbit (New Zealand white)	Acute eye irritation	24 hours Azimsulfuron technical (97.9% purity) <sup>a,b</sup> Draize, 1965 Kay <i>et al.</i> , 1962	Non-irritant	Kosaka, 1990b
Female guinea pig (Crj:Hartley)	Acute skin sensitization	72 hours; Magnusson and Kligman method Azimsulfuron technical (97.9% purity) <sup>a,b</sup> Magnusson and Kligman, 1969	Not a sensitizer	Ueda, 1990b

<sup>a</sup> Tests were conducted between 1989 and 1992. Process was optimized in 1994 to yield technical of ≥ 98% purity.

<sup>b</sup> This study was conducted in compliance with the GLP standards of MAFF in Japan, FIFRA, EPA in USA and OECD at The Institute of Environmental Toxicology in Tokyo, Japan.

**Table 4. Toxicology profile of technical azimsulfuron based on repeated administration (sub-acute to chronic)**

Species	Test	Duration and conditions or guideline adopted	Result	Study ref
Male and female rats (Fischer F344/DuCrj)	Oral	13 weeks Azimsulfuron technical (99.1% purity) OECD Guideline 408	NOEL = 1250 ppm (75.3 and 82.4 mg/kg/day for males and females respectively)	Toshishiro, 1993
Male and female mice ICR (Crj:CD-1)	Oral	13 weeks Azimsulfuron technical (99.1% purity) <sup>b</sup>	NOEL = 300 and 3000 ppm (40.62 and 469.9 mg/kg/day) for males and females respectively)	Harada, 1992
Male and female Beagle dogs	Oral	13 weeks Azimsulfuron technical (99.3% purity) OECD Guideline 409	NOEL = 300 ppm (8.81 and 9.75 mg/kg/day for males and females respectively)	Harada, 1995b
Male and female rat (Fischer F344/DuCrj)	Oral	24 months Azimsulfuron technical (99.0% purity) OECD Guideline 453	NOAEL = 1000 ppm (34.3 and 43.8 mg/kg/day for males and females respectively)	Kosei, 1993
Male and female mice ICR (Crj/CD-1)	Oral oncogenicity	18 months Azimsulfuron technical (99.0% purity) OECD Guideline 451	NOEL = 2500 ppm and 750 ppm (247.5 and 69.9 mg/kg/day) for males and females respectively. No evidence for oncogenicity	Takahashi, 1994
Male and female dogs (Beagle)	Oral feeding	1 year Azimsulfuron technical (99.0% purity) OECD Guideline 452	NOAEL = 750 ppm (17.88 and 19.25 mg/kg/day for males and females respectively)	Harada, 1995a
Male and female rats (Sprague-Dawley)	Reproductive toxicity (2 generations)	2 generations Azimsulfuron technical (99.0% purity) OECD Guideline 416	NOEL(reproduction) = 8000 ppm (601-724 and 663-783 mg/kg/day for males and females respectively) NOEL (systemic toxicity) = 125 ppm (9.59-11.25 and 10.92–2.39 mg/kg/day for males and females, respectively).	Hojo, 1994
Female Rats Crj:CD (SD)	Teratogenicity study	15 days Azimsulfuron technical (99.0% purity) US EPA FIFRA Guideline, Subdivision F, 83-3	Non teratogenic at dose levels up to 1000 mg/kg/day Maternotoxic and fetotoxic at 1000 mg/kg/day.	Fujii, 1994
Female rabbits (Japanese White)	Teratogenicity study	13 days Azimsulfuron 99.0% purity US EPA FIFRA Guideline Subdivision F, 83-3.	NOEL for maternotoxicity 150 mg/kg/day NOEL for fetotoxicity 500 mg/kg/day	Aoyama, 1994

<sup>b</sup> This study was conducted in compliance with the GLP standards of MAFF in Japan, FIFRA, EPA in USA and OECD at The Institute of Environmental Toxicology in Tokyo, Japan.

**Table 5. Mutagenicity profile of technical azimsulfuron based on *in vitro* and *in vivo* tests**

Species	Test	Conditions	Result	Study ref
<i>Salmonella typhimurium</i>	Mutagenicity	Azimsulfuron technical (97.9% purity) <sup>a</sup> U.S. EPA Pesticide Assessment Guidelines, Subdivision F, 84-2	Negative	Reynolds, 1989
Chinese Hamster ovary cells	CHO/HRPT gene mutation	Azimsulfuron technical (98.9% purity) OECD Guideline 476	Negative with and without activation	Gerber, 1994b
Rat hepatocytes	<i>In vitro</i> Unscheduled DNA synthesis (UDS)	Azimsulfuron technical (98.9% purity) OECD Guideline 482	UDS was not observed	Gerber, 1994a
Chines hamster lung cells	<i>In vitro</i> cytogenetics	Azimsulfuron technical (97.9% purity) <sup>a,b</sup>	No structural nor numerical chromosome aberrations in the metabolic activation method	Sasaki, 1990

<sup>a</sup> Tests were conducted between 1989 and 1992. Process was optimized in 1994 to yield technical of  $\geq 98\%$  purity.

<sup>b</sup> This study was conducted in compliance with the GLP standards of MAFF in Japan, FIFRA, EPA in USA and OECD at The Institute of Environmental Toxicology in Tokyo, Japan.

**Table 6. Ecotoxicology profile of technical azimsulfuron**

Species	Test	Duration and conditions	Result	Study ref
<i>Lepomis macrochirus</i> (bluegill sunfish)	Acute	96 hr, static Azimsulfuron technical (98.9% purity) OECD Guideline 203	LC <sub>50</sub> = >1000 mg/l NOEC = 780 mg/l	Kreamer, 1994d
<i>Oncorhynchus mykiss</i> (rainbow trout)	Acute	96 hr, static Azimsulfuron technical (98.9% purity) OECD Guideline 203	LC <sub>50</sub> = 154 mg/l NOEC = 49 mg/l	Kreamer, 1994e
<i>Oncorhynchus mykiss</i> (rainbow trout)	Sub-chronic	28 days, flow-through Azimsulfuron technical (98.9% purity) ECD Guideline 204	NOEC = 23 mg/l	Kreamer, 1994b
<i>Daphnia magna</i> (water flea)	Acute toxicity	48 hr, static Azimsulfuron technical (98.9% purity) ECD Guideline 202	EC <sub>50</sub> = 941 mg/l NOEC = 650 mg/l	Kreamer, 1994c
<i>Daphnia magna</i> (water flea)	Chronic toxicity	21 days, static renewal Azimsulfuron technical (98.9% purity) OECD Guideline 202	NOEC = 5.4 mg/l	Kreamer, 1995

Species	Test	Duration and conditions	Result	Study ref
<i>Oncorhynchus mykiss</i> (rainbow trout)	Chronic toxicity	90 days, flow-through Azimsulfuron technical (98.9% purity) OECD Guideline 210	NOEC = 6.3 mg/l	Kreamer, 1994a
<i>Lemna gibba</i>	Growth and reproduction	14 days Azimsulfuron technical (98.9% purity) FIFRA, Subdivision J, 123-2	EC <sub>50</sub> = 0.8 µg/l NOEC < 0.46 µg/l	Thompson, 1995
<i>Selenastrum capricornutum</i> (green alga)	Growth and reproduction	120 hours Azimsulfuron technical (98.9% purity) FIFRA, Subdivision J, 123-2,122-2	EC <sub>50</sub> = 12 µg/l NOEC < 8.1 µg/l	Thompson, 1994
<i>Eisenia foetida andrei</i> earthworm	Acute toxicity	14 days Azimsulfuron technical (98.9% purity) OECD Guideline 207	LC <sub>50</sub> = >1000 ppm	Caley <i>et al.</i> 1994
<i>Apis mellifera</i> (honey bee)	Acute oral and contact toxicity	48 hours Azimsulfuron technical (98.9% purity) FIFRA Subdivision L, Series 71-1, Hazard Evaluation: Non-target Insects	LC <sub>50</sub> = >1000 ppm (oral) and > 25.0 µg/bee (contact)	Beavers & Palmer, 1994 Palmer & Beavers, 1994
<i>Colinus Virginianus</i> and <i>Anas Platyrhynchos</i> Bobwhite quail and Mallard duck	Acute oral toxicity	14 days Azimsulfuron technical (98.9% purity) Pesticide Assessment Guidelines, FIFRA Subdivision E, Hazard Evaluation: Wildlife and Aquatic Organisms	LD <sub>50</sub> = >2250 mg/kg	Beavers & Campbell, 1994c, 1994a
<i>Colinus Virginianus</i> and <i>Anas Platyrhynchos</i> Bobwhite quail and Mallard duck	Dietary toxicity	5 days Azimsulfuron technical (98.9% purity) OECD Guideline 205	LC <sub>50</sub> = >5620 ppm	Beavers & Campbell, 1994b. Jaber & Campbell, 1994

Azimsulfuron has not been evaluated by the IPCS or by the FAO/WHO JMPR.

Azimsulfuron is classified by WHO as “unlikely to present acute hazard in normal use” (WHO 2002). It does not meet the criteria established in the UN Recommendations on the Transport of Dangerous Goods (published by the United Nations Committee of Experts on the Transport of Dangerous Goods) and, therefore, is not considered as dangerous/hazardous for transportation purposes.

### Formulations and co-formulated active ingredients

The main formulation type available is water dispersible granules (WG). Azimsulfuron may be co-formulated with other herbicides including bensulfuron-methyl or metsulfuron-methyl. These formulations are registered and sold in many countries throughout the world.

## Methods of analysis and testing

The analytical method for determination of the active ingredient (including identity tests) was validated by collaborative study (Bura 2003) and adopted, with provisional status, by CIPAC in 2004 (CIPAC 2004).

The azimsulfuron content is determined by reversed-phase HPLC using a 15 cm x 4.6 mm i.d. Zorbax<sup>®</sup> SB-C8 column, 5 µm particle size (or equivalent). The mobile phase is composed of pH 3.0 water and acetonitrile. The compound is detected using a UV detector at 240 nm and quantification is done by internal standardization with 4,4'-biphenol as the internal standard.

Within-laboratory validation data were also provided for an HPLC method (method ESB-22-93) designed for analysis of WG formulations (Blank, 1994). Azimsulfuron is determined on a reversed-phase HPLC system using a Zorbax<sup>®</sup> cyano column and a mobile phase of 30% acetonitrile and 70% water (adjusted to pH 3 with phosphoric acid), with UV detection at 240 nm. Phenyl sulphone is added as an internal standard. Validation data are summarized in Table 7.

**Table 7. Validation data for azimsulfuron content of WG formulations (Blank, 1994)**

Substrate	Test	Result
Method ESB-22-93 (HPLC, internal standard) Validation: Blank, 1994.		
WG	linearity	Sensitivity of response from 25% below to 25% above the target assay level (0.1 mg/l) is unchanged
	specificity	no co-elution of the azimsulfuron active ingredient or the internal standard, phenyl sulfone, occurred with impurities of the formulation or components of a formulation placebo.
	precision Note 1	Repeatability $S_r = 0.002$ at an azimsulfuron level of 49% (n=5). Reproducibility $S_r = 0.012$ at an azimsulfuron level of 49% (n=5), 2 different analysts on different days.
	LOQ	method as written is valid down to 5% of ai in formulations
	recovery	mean 101.1% (n=4) for spiking placebo at 48% and 51%.

Note 1:  $S_r$  is relative standard deviation.

The method for determination of impurities is based on reversed-phase gradient-elution HPLC, using UV detection at 254 nm and external standardisation (Simons, 2003).

Test methods for determination of physico-chemical properties of the technical active ingredient were OECD, CIPAC, EPA, EEC, while those for the formulations were CIPAC, as indicated in the specifications.

## Physical properties

The physical properties, the methods for testing them and the limits proposed for the WG formulations, comply with the requirements of the FAO/WHO Manual (FAO/WHO 2002). Physical testing of azimsulfuron 50% WG prepared in March 1992 was reported by Metzger and Koehler, 1995 (Table 8).

**Table 8. Physical testing of azimsulfuron 50% WG (Metzger and Koehler 1995)**

Test	Method	Result		
Appearance		Colour: tan.		
pH of 1% aq soln	CIPAC MT 75	5.7		
Bulk density (tap density)	CIPAC MT 169	0.63 g/ml		
Storage stability at 54°C	CIPAC MT 46	Test	Ambient	54°C
		Assay	49.2% ai	49.2% ai
		pH	5.7	5.6
		Wet sieve	0.6%	0.6%
		Suspensibility	66.1%	64.4%
		Dispersibility	91%	89%
Wettability	CIPAC MT 53.3.1	Wetting time: immediate		
Persistent foaming	CIPAC MT 47	Volume of foam at 1 minute: 0 ml		
Suspensibility	CIPAC MT 168	66%		
Dispersibility	CIPAC MT 174	91%		
Wet sieve test	CIPAC MT 167	0.6% retained on 75 µm sieve		
Nominal size range (dry sieve analysis)	CIPAC MT 170	≥90% retained on 500 µm sieve, ≤10% retained on 1400 µm sieve		
Dustiness	CIPAC MT 171	0.03% dust content		
Flowability	CIPAC MT 172	All of the test substance flowed freely through the sieve spontaneously (without agitation).		

### Containers and packaging

No extraordinary container or package issues need be considered.

### Expression of the active ingredient

The active ingredient is expressed as azimsulfuron.

### Appraisal

The Meeting considered the data and draft specifications in support of the development of new FAO specifications for azimsulfuron TC and WG. The data submitted were in accordance with the requirements of the FAO/WHO Manual (1<sup>st</sup> edition).

The water solubility of azimsulfuron depends on pH (72.3, 1050 and 6536 mg/l at 20°C at pH 5, 7 and 9 respectively). Azimsulfuron is acidic, with a pKa of 3.6. It is reasonably stable to hydrolysis and photolysis.

The Meeting was provided with commercially confidential information on the manufacturing process and batch analysis data on all impurities present at or above 1 g/kg. The process typically produces azimsulfuron having a minimum assay of 980 g/kg. Analyses of 5 batches of azimsulfuron produced in 2002 accounted for 100.3-101.4% of the material (azimsulfuron 99.9-100.9%, water 0.16-0.18%, total other impurities 0.23-0.32%). The Meeting was informed that several minor

impurities included in the manufacturing specification but which did not appear in the 5-batch data were included on the basis that they were only rarely detectable. The confidential data agreed with those submitted to the Ministry of Health in Italy (Desideri, 2004). The Meeting agreed that none of the impurities of azimsulfuron should be considered as relevant impurities.

The toxicological and ecotoxicological data were derived from azimsulfuron having impurity profiles similar to those described in the 5-batch analyses (manufacturing QC minimum limit 980 g/kg for azimsulfuron) or, for tests conducted between 1989 and 1992, on material of a lesser purity.

Azimsulfuron has not been evaluated by the WHO IPCS or by JMPR.

Azimsulfuron appears to be generally of low mammalian toxicity.

Ecotoxicological study summaries on fish, *Daphnia*, *Lemna gibba*, algae, birds, bees and earthworms were provided. Azimsulfuron is generally of low ecotoxicity except for the plants, *Lemna gibba* and green algae, with NOEC values of <0.46 and <8.1 µg/l respectively. In the view of WHO/PCS, as a herbicide it would be classified as very toxic to aquatic organisms.

Azimsulfuron does not meet the criteria established in the UN Recommendations on the Transport of Dangerous Goods (published by the United Nations Committee of Experts on the Transport of Dangerous Goods) and therefore, is not considered as dangerous or hazardous for transportation purposes.

The main formulation type is water dispersible granules (WG).

The analytical method for active ingredient relies on reversed-phase HPLC with internal standard phenyl sulfone and has been adopted by CIPAC. The HPLC method provides one identity test, with IR spectrophotometry for further identification.

The physical properties of azimsulfuron WG comply with the proposed specifications. No special requirements for containers and packaging have been identified.

## Recommendations

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The Meeting recommended that the draft specifications for azimsulfuron TC and WG proposed by E.I. du Pont de Nemours and Company, as amended by agreement between the company and the Meeting, should be adopted by FAO.

## References

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|----------------|---|
| Aoyama H. 1994 | DPX-A8947: Teratogenicity study in rabbits. Study IET 90-0041. The Institute of Environmental Toxicology, Japan. Unpublished. |
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- Barefoot A.C. 1998. Photodegradation of [pyrimidine-2-<sup>14</sup>C]DPX-A8947 and [pyrazole-4-<sup>14</sup>C]DPX-A8947 in water conducted in simulated sunlight. Quantum efficiency for degradation of azimsulfuron in water (supplement 1). Report. AMR 1724-90 Sup 1. DuPont Experimental Station. Unpublished.
- Beavers B.J. and Campbell S.M. 1994a. An acute oral toxicity study with the mallard. Report. HLO 412-94. Wildlife International Ltd (USA). Unpublished.
- Beavers B.J. and Campbell S.M. 1994b. DPX-A8947-24 (technical) a dietary LC<sub>50</sub> study with the Northern Bobwhite. Report. HLO 215-94. Wildlife International Ltd (USA). Unpublished.
- Beavers B.J. and Campbell S.M. 1994c. DPX-A8947-24 (technical): an acute oral toxicity study with the Northern Bobwhite. Report. HLO 216-94. DuPont Haskell Laboratory. Unpublished.
- Beavers J.B. and Palmer S.J. 1994. DPX-A8947-24 (technical): a dietary LC<sub>50</sub> toxicity study with the honey bee. Report. HLO 213-94. Wildlife International Ltd (USA). Unpublished.
- Blank D.J. 1994. Validation of analytical method for the determination of DPX-A8947 in A8947 50DF. DuPont report AMR 2774-93. Unpublished.
- Bura L. 2003. CIPAC Information Sheet 247. 11 Nov 2003. Azimsulfuron.
- Caley C.Y., Knight B. and Boyle J. 1994. DPX-A8947 Technical determination of acute toxicity in earthworms (limit test). Report. AMR 2863-93. Inveresk Research International (IRI) Limited (Scotland). Unpublished.
- CIPAC 2004. <http://www.cipac.org>. Status of new CIPAC methods. Decisions of the 48<sup>th</sup> meeting in Brno. 584 Azimsulfuron (note: originally published as validation of the method for analysis of EC, not WG, but subsequently corrected). The method is not yet published in a CIPAC Handbook.
- Cobranchi D.P. and Schmuckler M.E. 1994. Vapor pressure determination of DPX-A8947. Report. AMR 3020-94. DuPont Chambers Works. Unpublished.
- Cooke L.A. 1993. Dissociation constant of DPX-A8947. Report. AMR 2031-91. DuPont Experimental Station. Unpublished.
- Desideri A.. 2004. Azimsulfuron specifications. Letter (22 July 2004), Ministry of Health, Rome, Italy.
- Draize J.H., 1965. Appraisal of the safety of chemicals in foods, drugs, and cosmetics – dermal toxicity, pp. 45-59, *Assoc. of Food and Drug Officials of the United States*. Topeka, Kans.
- Ebino K.. 1993. Acute inhalation toxicity study in rats. Report. IET 90-0044. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- EPA, 1985. GCSOLAR model, United States Environmental Protection Agency, Environmental Research Laboratory, Athens, Georgia.
- FAO/WHO 2002. Manual on development and use of FAO and WHO specifications for pesticides, 1<sup>st</sup> edition, 2002, Rome.
- Fujii S. 1994. DPX-A8947: Teratogenicity study in rats. Report. IET 90-0039. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Gerber K.M. 1994a. Assessment of DPX-A8947-24 (technical) in the *in vitro* unscheduled DNA synthesis assay in primary rat hepatocytes. Report. HLR 481-94. DuPont Haskell Laboratory. Unpublished.
- Gerber K.M. 1994b. Mutagenicity evaluation of DPX-A8947-24 (technical) in the CHO/HPRT assay. Report. HLR 460-94. DuPont Haskell Laboratory. Unpublished.
- Harada T. 1992. DPX-A8947: 13-week oral subchronic toxicity study in mice. Report. IET 90-0033. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Harada T. 1995a. DPX-A8947: 12-month oral chronic toxicity study in dogs. Report. IET 90-0037 VOL1 - VOL2. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Harada T. 1995b. DPX-A8947: 13-week oral subchronic toxicity study in dogs. Report. IET 90-0036. Institute of Environmental Toxicology (IET) (Japan). Unpublished.

- Hausmann S.M. 1991a. Hydrolysis of [pyrimidine-2-<sup>14</sup>C] DPX-A8947 and [pyrazole-4-<sup>14</sup>C] DPX-A8947 in buffer solutions at pH 5, 7, and 9. Report. AMR 1725-90. DuPont Experimental Station. Unpublished.
- Hausmann S.M. 1991b. Photodegradation of [pyrimidine-2-<sup>14</sup>C] DPX-A8947 and [pyrazole-4-<sup>14</sup>C] DPX-A8947 in water conducted in simulated sunlight. Report. AMR 1724-90. DuPont Experimental Station. Unpublished.
- Hitch R.K, 1982a. EPA Guidelines: "Hydrolysis Studies", Pesticide Assessment Guidelines, Subdivision N, Chemistry: Environmental Fate 161-1, pp 44-46; October 1982; National Technical Service Information No. PB83-153973
- Hitch R.K, 1982b. EPA Guidelines: "Photodegradation Studies in Water", Pesticide Assessment Guidelines, Subdivision N, Chemistry: Environmental Fate 161-2, pp 46-49; October 1982; National Technical Service Information No. PB83-153973
- Hojo H. 1994. DPX-A8947: Two-generation reproduction study in rats. Report. IET 90-0043. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Jaber M.J and Campbell S.M. 1994. DPX-A8947-24 (technical): a dietary LC<sub>50</sub> study with the mallard. Report. HLO 217-94. Wildlife International Ltd (USA). Unpublished.
- Kay J.H. and Calandra J.C., 1962. Interpretation of eye irritation tests, *J. Soc. Cosm. Chem.* 13; pp. 281-289.
- Kosaka T. 1990a. DPX-A8947: Primary dermal irritation study in rabbits. Report. IET 90-0002. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Kosaka T. 1990b. DPX-A8947: Primary eye irritation study in rabbits. Report. IET 90-0001. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Kosei I. 1993. DPX-A8947: 24-month oral chronic toxicity and oncogenicity study in rats. Report. IET 90-0031 VOL1 - VOL7. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Kreamer C. 1994a. Early life-stage toxicity of DPX-A8947-24 (technical) with Rainbow Trout; *Oncorhynchus mykiss* (revision 1). Report. HLR 391-94 RV1. DuPont Haskell Laboratory. Unpublished.
- Kreamer C. 1994b. Flow-through, 28-day toxicity of DPX-A8947-24 (technical) to Rainbow Trout, *Oncorhynchus mykiss*. Report. HLR 342-94. DuPont Haskell Laboratory. Unpublished.
- Kreamer C. 1994c. Static, acute, 48-hour EC<sub>50</sub> of DPX-A8947-24 (technical) to *Daphnia magna*. Report. HLR 170-94. DuPont Haskell Laboratory. Unpublished.
- Kreamer C. 1994d. Static, acute, 96-hour LC<sub>50</sub> OF DPX-A8947-24 (technical) to Bluegill Sunfish, *Lepomis macrochirus*. Report. HLR 115-94. DuPont Haskell Laboratory. Unpublished.
- Kreamer C. 1994e. Static, acute, 96-hour LC<sub>50</sub> of DPX-A8947-24 (technical) to Rainbow Trout, *Oncorhynchus mykiss*. Report. HLR 116-94. DuPont Haskell Laboratory. Unpublished.
- Kreamer C. 1995. Chronic toxicity of DPX-A8947-24 (technical) to *Daphnia magna*. Report. HLR 459-94. DuPont Haskell Laboratory. Unpublished.
- Lesieur L.B. 1994. Thermal stability of A8947. Report. AMR 2794-93. DuPont Experimental Station. Unpublished.
- MAFF Japan, 1985. "Testing guidelines for Toxicology Studies"; Notification of Director, Plant Protection Division, Agricultural Protection Bureau, Ministry of Agriculture, Forestry and Fisheries in Japan; NohSan No. 4200, January 28, 1985
- Magnusson B. and Kligman A.M., 1969. The identification of contact allergens by animal assay. The guinea pig maximization test. *J. Invest. Dermatol.*, 52, 268-276.
- Metzger J.D. and Koehler J.A. 1995. Physical, chemical and technical properties of DPX-A8947 50% water-dispersible granular formulation. DuPont report AMR 3146-94.
- Moore L.A. 1994a. Melting point of DPX-A8947. Report. AMR 3004-94. DuPont Experimental Station. Unpublished.
- Moore L.A. 1994b. Relative density of DPX-A8947 AT 20°C. Report. AMR 2965-94. DuPont Experimental Station. Unpublished.
- Palmer S.J. and Beavers J.B. 1994. DPX-A8947-24 (technical): an acute contact toxicity study with the honey bee. Report. HLO 214-94. Wildlife International Ltd (USA). Unpublished.

- Reynolds V.L. 1989. Mutagenicity testing of INA-8947 in the *Salmonella typhimurium* plate incorporation assay. Report. HLR 450-89. DuPont Haskell Laboratory. Unpublished.
- Sasaki Y.F.X. 1990. DPX-A8947: *In vitro* cytogenetics test. Report. IET 89-0107. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Schmuckler M.E. 1991. Water solubility of DPX-A8947 using continuous sample agitation. Report. AMR 1585-90. DuPont Experimental Station. Unpublished.
- Schmuckler M.E. 1995. Volatility of DPX-A8947 AT 20°C (Calculation of Henry's law constant). Report. AMR 3079-94. DuPont Experimental Station. Unpublished.
- Simons C. 2003. Azimsulfuron (DPX-A8947): Analysis and certification of product ingredients for technical grade material produced at the Manati, Puerto Rico manufacturing facility. DuPont report 11123. Study 008-122, Exygen Research, USA. Unpublished.
- Smyser B.P. 1991. n-octanol/water partition coefficient determination of DPX-A8947 at pH 5, pH7, and pH 9. Report. AMR 2029-91. DuPont Experimental Station. Unpublished.
- Takahashi K. 1994. DPX-A8947: 18-month oral oncogenicity study in mice. Report. IET 90-0034 Vol1 - Vol6. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Thompson S.G. 1994. DPX-A8947: Influence on growth and reproduction of four select algal species. Report. AMR 3025-94. Wildlife International Ltd (USA). Unpublished.
- Thompson S.G. 1995. DPX-A8947: Influence on growth and reproduction of *Lemna gibba* G3. Report. AMR 2997-94. Wildlife International Ltd (USA). Unpublished.
- Toshishiro K. 1993. DPX-A8947: 13-week oral subchronic toxicity study in rats. Report. IET 90-0030. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Ueda H. 1990a. DPX-A8947: Acute dermal toxicity study in rats. Report. IET 89-0105. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Ueda H. 1990b. DPX-A8947: Dermal sensitization study in guinea pigs. Report. IET 90-0003. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- Ueda J. 1990c. DPX-A8947: Acute oral toxicity study in rats. Report. IET 89-0103. Institute of Environmental Toxicology (IET) (Japan). Unpublished.
- WHO 2002. The WHO recommended classification of pesticides by hazard and guidelines to classification 2000-2002, WHO, Geneva, 2002.