

**CYFLUMETOFEN (273)**

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**EXPLANATION**

Cyflumetofen (consisting of isomers) is a bridged diphenyl acaricide (miticide) for control of *Tetranychus* sp. (Two spotted spider mite, Kanzawa spider mite, Desert spider mite) and influences the mitochondrial electron transport chain by inhibiting the complex II substance in the cell. Cyflumetofen does not have translaminar or systemic activity.

Cyflumetofen was first registered in Japan in 2007. Henceforth, cyflumetofen has been registered in a number of countries. It is used in a variety of crops, including citrus fruits, pome fruits, stone fruits, berry fruits, vegetables, dried herbs, and teas. It was included in the Codex Priority List in 2013 by the 45<sup>th</sup> CCPR as a new compound for evaluation by the current JMPR.

The Meeting received information on physical and chemical properties, animal and plant metabolism, environmental fate, analytical methods, storage stability, use patterns, supervised trials, and processing. Cyflumetofen is reviewed by the JMPR for the first time.

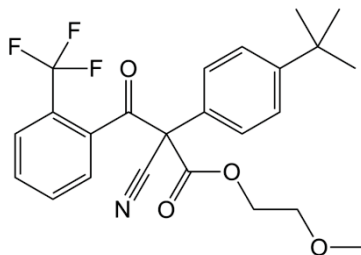
**IDENTITY**

ISO common name: Cyflumetofen  
 Chemical name  
 IUPAC: 2-methoxyethyl (*RS*)-2-(4-*tert*-butylphenyl)-2-cyano-3-oxo-3-( $\alpha,\alpha,\alpha$ -trifluoro-*o*-tolyl)propionate  
 CAS: 2-methoxyethyl  $\alpha$ -cyano- $\alpha$ -[4-(1,1-dimethylethyl)phenyl]- $\beta$ -oxo-2-(trifluoromethyl)benzenepropanoate

CAS Registry No.: 400882-07-7

CIPAC No.: 821

Structural formula:



Molecular formula: C<sub>24</sub>H<sub>24</sub>F<sub>3</sub>NO<sub>4</sub>

Molecular weight: 447.45

**PHYSICAL AND CHEMICAL PROPERTIES**

*Pure active ingredient*

Property	Results	Reference
Appearance	White solids at 21.5 °C	Iijima K, 2001a [OTSA-0029(EN)-FR]
Odour:	Odourless at 21.5 °C	Iijima K, 2001c [OTSA-0030(EN)-FR]

Hydrolysis	Half-life at 25 and 40 °C 25 °C  40 °C pH 4  222 h  70 h pH 7   5 h   3 h pH 9   12 m  not calculable	Knight L, 2004a [OTSA-0076]
Photolysis	After 4 and 2 hours of irradiation with xenon arc light (38 W/m <sup>2</sup> , 300–400 nm) respectively at 25 °C (0.01 mg/L).  DT <sub>50</sub> : pH 4 buffer            Natural water Eq. summer, 40 °N lat:    5.00 h                  54 h (including hydrolysis) Eq. Tokyo spring, 35 °N lat: 10.6 h                  5.36 h  In non-irradiated aqueous pH 4 buffer solution, 93.4% of AR recovered as the parent after 4 h, while in non-irradiated natural water, degradation was significant (38.6% of AR recovered as the parent after 2 hours). Since degradation also occurred in non-irradiated samples, a significant proportion of the degradation in natural water could be attributed to hydrolysis.	Knight L, 2004a, [OTSA-0078]

### Technical material

Property	Results	Reference
Appearance:	Yellow solid at room temperature	Baltussen E, 2006a [OTSA-0327-FR]
Odour	No characteristic odour at room temperature	Baltussen E, 2006a [OTSA-0327-FR]
Relative density:	1.229 g/mL at 20 °C (pycnometer comparable method)	Iijima K, 2003a [OTSA-0031(EN)-FR]
Melting point:	779–81.7 °C	Iijima K, 2002a [OTSA-0032(EN)-FR]
Boiling point:	293–297 °C at atmospheric pressure (no decomposition prior to boiling)	Iijima K, 2002a [OTSA-0032(EN)-FR] Iijima K, 2002c [OTSA-0039(EN)-FR]
Temperature decomposition of	In the range of 320–446 °C	Iijima K, 2002a [OTSA-0032(EN)-FR]
Vapour pressure:	< 5.9 × 10 <sup>-6</sup> Pa at 25 °C 3.1 ± 0.23 × 10 <sup>-5</sup> Pa at 50 °C (Extrapolation from measurements at 150 °C to 190 °C)  < 1.3 × 10 <sup>-5</sup> Pa (1 × 10 <sup>-7</sup> mmHg) (25 °C) (Triplicate test system using gas saturation method)	Saka M, 2002 [OTSA-0049(EN)-FR]
Volatility, Henry's Law Constant at 20 °C	< 9.4 × 10 <sup>-2</sup> Pa m <sup>3</sup> /mol (Calculation by dividing vapour pressure by water solubility)	Cardinaals JM, 2006a [OTSA-0331-FR]
Octanol-water partition coefficient at (log P <sub>ow</sub> )	4.3 at 25 °C No significant effect of pH (4–10) on log P <sub>ow</sub> .	Iijima K, 2004 [OTSA-0036(EN)-FR]
Solubility in water	In pure water	Iijima K, 2002e

Property	Results	Reference
	28 µg/L at 20 °C (pH 7) No effect of PH expected	[OTSA-0034(EN)-FR]
Solubility in solvents at 20 °C:	mg/mL solvent Acetone > 500 Dichloromethane > 500 Ethyl acetate > 500 n-Hexane 5.16 (in solution) Methanol 98.7 (in solution) Toluene > 500  Deionized water 2.15 0.215 pH 5 Buffer 36.0 3.69 pH 7 Buffer 479 68.1 pH 9 Buffer 518 76.5 <sup>a</sup> corrected for solution density	Iijima K, 2003c [OTSA-0035(EN)-FR]
Hydrolysis:	DT <sub>50</sub> at 25 °C pH 4 7.7 d pH 5 6.0 d pH 7 9.8 h pH 9 10.3 m Five major metabolites (> 10% of TAR) were formed	Nakamura H, 2004a [OTSA-0156(EN)-FR]
Dissociation constant (pKa)	-4.2	Cardinaals JM, 2006b [OTSA-0330-FR]
Photolysis	The aquatic photolysis was determined after 2 days of irradiation with xenon arc light (20 W/m <sup>2</sup> , 290–400 nm) at 25 °C. In natural water a significant proportion of the degradation may be attributed to hydrolysis since non-irradiated solution showed extensive degradation  DT <sub>50</sub> in pH 5 buffer: Irradiated: 1.28 h (including hydrolysis) Non irradiated: 134 h DT <sub>50</sub> in natural water: Irradiated: 1.07 h (including hydrolysis) Non irradiated: 3.40 h	Ohyama K, 2004a [OTSA-0157(EN)-FR]
Formulations:	Suspension concentrate (flowable concentrate) (SC) formulations containing 200 g ai/L available	

No FAO specification has been developed for cyflumetofen.

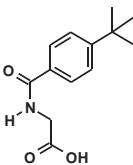
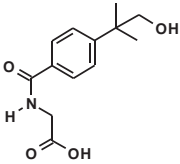
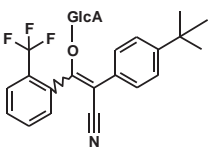
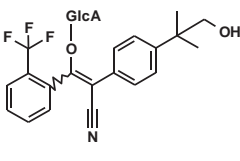
#### METABOLISM AND ENVIRONMENTAL FATE

The following links code numbers and structure or description of the compounds appearing in the various metabolism and environmental fate studies.

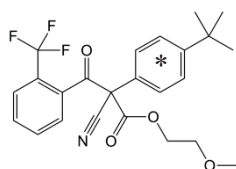
Table 1 Structure of compounds appearing in metabolism and environmental fate studies

Code (MW)	IUPAC name	Structure	Metabolite or degradate found in
Cyflumetofen (447.45)	2-methoxyethyl ( <i>RS</i> )-2-(4- <i>tert</i> -butylphenyl)-2-cyano-3-oxo-3-( $\alpha,\alpha,\alpha$ -trifluoro- <i>o</i> -tolyl)propionate		Goat Satsuma mandarin Apple Eggplant
B-1 (190.12)	2-(trifluoromethyl)benzoic acid		Rat Goat Satsuma mandarin Apple Eggplant Soil
B-3 (189.13) Syn: M9210I005	2-(trifluoromethyl)benzamide		Soil
AB-1 (345.36) Syn: M9210I003	( <i>RS</i> )-2-(4- <i>tert</i> -butylphenyl)-3-oxo-3-[2-(trifluoromethyl)phenyl] propanenitrile		Rat Goat Hydrolysis in buffer Hydrolysis simulating processing Soil
AB-6 (465.45)	2-methoxyethyl-2-(( <i>R,S</i> )-4- <i>tert</i> -butylphenyl)-3-oxo-3-({2-(trifluoromethyl)phenyl}carbonyl) amino)propanoate		Satsuma mandarin Apple Eggplant
AB-7 (447.45)	2-methoxyethyl-(( <i>R,S</i> )-4- <i>tert</i> -butyl-2-{{2-(trifluoromethyl)phenyl}carbonyl} phenyl (cyano)acetate		Satsuma mandarin Apple Eggplant
A-1 (275.35) Syn: M9210I008	4-methoxyethyl ( <i>RS</i> )-4- <i>tert</i> -butylphenyl)(cyano)acetate		Soil
A-2 (173.26) Syn: M9210I001	4- <i>tert</i> -butylphenyl-acetonitrile		Goat Soil

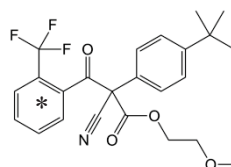
Code (MW)	IUPAC name	Structure	Metabolite or degradate found in
A-12 (178.23) Syn: M9210I002	4- <i>tert</i> -butylbenzoic acid		Goat Soil
A-20 (173.26) Syn: M9210I019	2-(4-carboxyphenyl)-2-methyl-1-propanol		Rat Goat
M9210I014 (217.67)	2-cyano-2-(4- <i>tert</i> -butylphenyl)acetic acid		Goat
M9210I021 (375.35)	(2 <i>RS</i> )-2-methyl-2-{4-[2-(2-trifluoromethylphenyl)-1-cyano-2-oxoethyl]phenyl}-propionic acid		Goat
M9210I023 (189.26)	4-(1-hydroxy-2-methyl-2-propyl) benzonitrile		Goat
M9210I029 (363.33)	(2 <i>RS</i> )-2-{4-[(2 <i>RS</i> )-1,2-dihydroxy-2-propyl]phenyl}-3-(2-trifluorophenyl)-3-oxo-propionitrile		Goat
M9210I030 (347.34)	(2 <i>RS</i> )-2-[4-(2-hydroxy-2-propyl)phenyl]-3-(2-trifluorophenyl)-3-oxopropionitrile		Goat
M9210I031 (331.30)	(2 <i>RS</i> )-2-(4-acetylphenyl)-3-(2-trifluorophenyl)-3-oxopropionitrile		Goat
M9210I032 (203.24)	2-(4-cyanomethylphenyl)-2-methyl-propionic acid		Goat

Code (MW)	IUPAC name	Structure	Metabolite or degradate found in
M9210I033 (235.28)	<i>N</i> -(4- <i>tert</i> -butylphenylcarbonyl)-aminoacetic acid		Goat
M9210I040 (251.28)	<i>N</i> -[4-(1-hydroxy-2-methyl-2-propyl)phenylcarbonyl]-aminoacetic acid		Goat
M9210I042/ M9210I044 (isomers) (521.49)	$\alpha$ -[2-(4- <i>tert</i> -butylphenyl)-2-cyano-1-(2-trifluoromethylphenyl)ethenyl]glucoside		Goat
M9210I043 (537.47)	$\alpha$ -{2-cyano-1-(2-trifluoromethylphenyl)-2-[4-(1-hydroxy-2-methyl-2-propyl)phenyl]ethenyl} glucoside		Goat

The metabolism and distribution of cyflumetofen in animals and plants and the fate of cyflumetofen in the environment were investigated using the following  $^{14}\text{C}$ -radiolabelled compounds.



[*t*-butylphenyl-ring- $^{14}\text{C}$ ]-cyflumetofen  
("butylphenyl-label")



[benzoyl-ring- $^{14}\text{C}$ ]-cyflumetofen  
("benzoyl-label")

### Animal metabolism

The Meeting received information on the results of studies on lactating goats which were fed radio-labelled cyflumetofen.

Metabolism studies on laboratory animals including rats were reviewed in the framework of toxicological evaluation by the current JMPR and the relevant information is summarized below.

#### Rat

Highest residues were found in the liver followed by the kidney regardless of sex, dose and label position and time point of measurement. Cyflumetofen was extensively metabolized: in total 18 metabolites were identified. B-1, 2-trifluoromethylbenzoic acid was a major metabolite in the rat metabolism (occurring up to 28% of the applied dose). The predominant metabolic pathway for cyflumetofen involves cleavage of the *tert*-butylphenyl and trifluorotolyl moieties. Major reactions on the *tert*-butylphenyl ring are cleavage of the methoxyethyl group, hydroxylation at the butyl group, decarboxylation and glucuronidation at the butyl group. Major reactions on the trifluorotolyl-ring are glutathione conjugation at the carboxyl group and further changes of the glutathione group to mercapturic acid or thiolactic acid. In addition, hydroxylation and

oxidation reactions at the butyl group and cleavage of the carboxylic ester moiety is observed on the parent structure.

*Lactating goats (Sowka, et al, 2011a; and Fabian & Landsiedel, 2011a)*

Two lactating goats were given orally by gavage radio-labelled cyflumetofen for 12 (benzoyl-label) or 10 (butylphenyl-label) consecutive days. The mean actual dose of benzoyl-label for each of the two goats (Goat 1 and 2) was 0.27 and 0.30 mg/kg bw corresponding to 14.9 and 12.1 ppm in feed. The mean actual dose of butylphenyl-label for each of the two goats (Goat 3 and 4) was 0.43 and 0.48 mg/kg bw corresponding to 12.8 and 11.8 ppm in feed. Milk samples were collected twice daily. The goats were sacrificed 18–24 h after the last dose and tissue/organ samples were collected.

The radioactivity in urine, faeces, blood, stomach content, gut content, gut/stomach, individual milk samples (two samples per day), cage wash, cream and skim milk was determined by liquid scintillation counting (LSC). After transport of the samples to the test facility, subsamples of urine, faeces, and milk of both animals dosed with the benzoyl-label and both animals dosed with the butylphenyl-label were combined to generate label-specific pooled samples. The tissues/organs of both animals (benzoyl- and butylphenyl-label separately) were combined and homogenized. Bile was also pooled. The pooled samples and homogenized tissues were measured with LSC (solid samples after oxidative combustion of small aliquots). All samples were stored at -18 °C or below.

No extraction was necessary for urine and bile prior to HPLC analysis. Subsamples of the pooled milk and faeces samples, and homogenized tissue samples were extracted with acetonitrile and water, or only acetonitrile. The results of the solvent extractions (LSC of the combined extracts) are referred to as extracted radioactive residues (ERR). The residual radioactive residues after solvent extraction (RRR) were determined by combustion analysis. The solvent extracts with sufficient concentrations of radioactive residues were analysed by reversed-phase HPLC with gradient elution and radio-detection, generally using two chromatographic methods.

Identification of metabolites was mainly based on fractionation of urine of both labels and bile of the butylphenyl-label, followed by HPLC-MS, GC-MS and/or NMR analysis of the isolated fractions. Thereafter, the MS samples were used as reference items for co-chromatography experiments to achieve peak assignment.

The residues after solvent extraction of liver (benzoyl- and butylphenyl-label) and kidney (benzoyl-label) were further incubated with protease, whereby significant portions of radioactivity were released. The protease solubilizates of liver (both labels) were analysed by HPLC.

During the study period, no behavioural or physical abnormalities were observed. At sacrifice, macroscopic examinations revealed no abnormalities.

Daily samples of milk, urine, and faeces obtained for twelve and ten consecutive days were measured for total radioactive residues for each animal. The results in Table 2 indicate that the radioactive residue was rapidly and extensively excreted.

Until sacrifice, the radioactive residues excreted via urine amounted to 29.9% and 33.6% (Goat 1 and 2 of the benzoyl-label) and 38.4% and 45.3% (Goat 3 and 4 of the butylphenyl-label) of the total radioactivity administered. The residues excreted via faeces accounted for 49.0% and 43.9% (Goat 1 and 2) and 50.3% and 42.6% (Goat 3 and 4) of the dose. The recovered radioactivity from urine and faeces together with from cage wash accounted for 79.4 and 78.5% of the total administered radioactivity for Goat 1 and Goat 2, respectively. Similarly, the recovered radioactivity from urine and faeces together with from cage wash accounted for 89.6, and 89.1% of the total administered radioactivity for Goat 3 and 4, respectively.

In milk, only low proportions of the administered dose were found, 0.10% and 0.14% (benzoyl-label, Goat 1 and 2) and 0.04% and 0.03% (butylphenyl-label, Goat 3 and Goat 4).

For both labels, only low portions of the administered dose were retained in edible tissues and organs. The total radioactive residues of the benzoyl- and butylphenyl-label in edible tissues/organs and milk were as follows: 0.019 mg/kg and 0.008 mg/kg in milk; 0.404 mg/kg and 0.287 mg/kg in liver, 0.191 mg/kg and 0.167 mg/kg in kidney, 0.010 mg/kg and 0.005 mg/kg in leg muscle, 0.010 mg/kg and 0.004 mg/kg in back muscle, 0.014 mg/kg and 0.018 mg/kg in abdominal fat and 0.014 mg/kg and 0.015 mg/kg in renal fat. In milk, muscle and fat tissues, only low residue levels were found. The rate of extracted radioactivity was high, above 78% TRR, except for milk from goats dosed with the benzoyl-label (51% TRR) and liver

( $\geq 65\%$  TRR). In milk; however, the radioactive residues after solvent extraction were below 0.010 mg/kg, and from liver additional radioactivity was released upon protease treatment.

For further analysis, samples from Goat 1 and Goat 2 were pooled (benzoyl-label) and Goat 3 and Goat 4 (butylphenyl-label) were pooled to generate pooled samples for milk (Days 5-12 and Days 5-10), urine (Days 1-12 and Days 1-10), and homogenized faeces (Days 1-12 and Days 1-10). Similarly for each label, samples of liver, bile, kidney, muscle tissues and fat tissues were used to generate one pooled sample per matrix and label. These samples were LSC measured, resulting in the TRR. Additionally, milk and tissue samples were extracted and the TRR was calculated as the sum of the ERR and the residue after solvent extraction (TRR calculated). The TRR calculated was used for all further calculations.

The total radioactive residues in goat milk, tissue, urine and faeces are provided in Table 2.

Table 2a Material balance after repeated oral dose of radiolabelled cyflumetofen to goats, including distribution of residues, and total radioactive residues (TRRs) in milk, tissues, urine and faeces

Matrix	Benzoyl-label				
	TRR measured <sup>a</sup> mg eq/kg		% of Dose		TRR calculated <sup>b</sup> mg eq/kg
	Goat 1	Goat 2	Goat 1	Goat 2	Goats 1 & 2
Urine Subtotal Day 1–12			29.873	33.608	
Urine, pooled sample Day 1–12	4.084				<sup>c</sup>
Faeces Subtotal Day 1–12			48.965	43.906	
Faeces, pooled sample Day 1–12 <sup>e</sup>	1.357				1.252
Milk; Application day 1			0.003	0.007	<sup>c</sup>
Milk; Application day 2			0.006	0.007	<sup>c</sup>
Milk; Application day 3			0.006	0.008	<sup>c</sup>
Milk; Application day 4			0.008	0.012	<sup>c</sup>
Milk; Application day 5			0.009	0.014	<sup>c</sup>
Milk; Application day 6			0.012	0.020	<sup>c</sup>
Milk; Application day 7			0.013	0.012	<sup>c</sup>
Milk; Application day 8			0.011	0.013	<sup>c</sup>
Milk; Application day 9			0.008	0.012	<sup>c</sup>
Milk; Application day 10			0.008	0.011	<sup>c</sup>
Milk; Application day 11			0.008	0.009	<sup>c</sup>
Milk; Application day 12			0.008	0.015	<sup>c</sup>
Milk Subtotal			0.101	0.141	
Milk: pooled sample Day 5–12	0.019				0.018
Liver	0.404	0.404	0.184	0.154	0.390
Blood	0.039	0.014	0.047	0.018	<sup>c</sup>
Kidney	0.191	0.191	0.017	0.016	0.190
Kidney fat	0.014	0.014	0.001	0.002	0.015
Abdominal fat <sup>f</sup>	0.014	0.014	0.001	0.002	0.015
Leg muscle	0.010	0.010	0.020	0.019	0.010
Back muscle <sup>f</sup>	0.010	0.010	0.004	0.003	0.009
Bile	1.426	1.426	0.008	0.019	<sup>c</sup>
Gut/Stomach	0.066	0.061	0.337	0.276	<sup>c</sup>
Stomach contents	0.271	0.104	4.667	1.190	<sup>c</sup>
Gut contents	1.556	1.535	4.122	5.974	<sup>c</sup>
Urine in bladder	<sup>d</sup>	3.219	<sup>d</sup>	0.026	<sup>c</sup>
Organs/Tissues Subtotal			9.408	7.699	
Cage wash	–	–	0.546	0.961	
Total Recovery			88.894	86.318	

<sup>a</sup> For liver, bile, kidney, fat and muscle samples, the values determined from pool samples are given and were converted to % dose values for the individual animals.

<sup>b</sup> TRR calculated from pooled samples as the sum of ERR and residue after solvent extract (used for calculation of metabolite concentrations)

<sup>c</sup> Not calculated

<sup>d</sup> Not sampled

<sup>e</sup> Faeces was homogenized with water prior LSC (approximately 4 to 8 parts water per part faeces)

<sup>f</sup> Abdominal fat referred as subcutaneous fat in the in-life report, back muscle referred as flank muscle



Table 2b Material balance after repeated oral dose of radiolabelled cyflumetofen to goats, including distribution of residues, and total radioactive residues (TRRs) in milk, tissues, urine and faeces

Matrix	Butylphenyl-label				
	TRR measured mg eq/kg <sup>a</sup>		% of Dose		TRR calculated <sup>b</sup> mg eq/kg
	Goat 3	Goat 4	Goat 3	Goat 4	Goats 3 /4
Urine Subtotal Day 1–12			38.437	45.272	
Urine, pooled sample Day 1–12	6.079				<sup>c</sup>
Faeces Subtotal Day 1–12			50.280	42.639	
Faeces, pooled sample Day 1–12 <sup>e</sup>	1.304				1.290
Milk; Application day 1			0.003	0.002	<sup>c</sup>
Milk; Application day 2			0.003	0.002	<sup>c</sup>
Milk; Application day 3			0.003	0.003	<sup>c</sup>
Milk; Application day 4			0.003	0.003	<sup>c</sup>
Milk; Application day 5			0.004	0.003	<sup>c</sup>
Milk; Application day 6			0.003	0.004	<sup>c</sup>
Milk; Application day 7			0.004	0.003	<sup>c</sup>
Milk; Application day 8			0.004	0.003	<sup>c</sup>
Milk; Application day 9			0.005	0.002	<sup>c</sup>
Milk; Application day 10			0.004	0.002	<sup>c</sup>
Milk Subtotal			0.036	0.026	
Milk: pooled sample Day 5–10	0.008				0.007
Liver	0.287	0.287	0.118	0.121	0.282
Kidney	0.167	0.167	0.012	0.013	0.163
Kidney fat	0.015	<sup>d</sup>	0.000	<sup>d</sup>	<sup>d</sup>
Abdominal fat <sup>f</sup>	0.018	0.018	0.002	0.001	0.020
Leg muscle	0.005	0.005	0.010	0.012	0.004
Back muscle <sup>f</sup>	0.004	0.004	0.010	0.002	0.003
Blood	0.015	0.024	0.010	0.023	<sup>c</sup>
Bile	3.164	3.164	0.022	0.050	<sup>c</sup>
Gut/Stomach	0.056	0.039	0.176	0.136	<sup>c</sup>
Gut contents	0.762	0.654	2.080	0.772	<sup>c</sup>
Stomach contents	0.220	0.279	1.770	3.182	<sup>c</sup>
Urine in bladder	3.295	<sup>d</sup>	0.042	<sup>d</sup>	<sup>c</sup>
Organs / Tissues Subtotal			4.252	4.312	
Cage wash	-	-	0.921	1.192	
Total Recovery			93.917	95.442	

<sup>a</sup> For liver, bile, kidney, fat and muscle samples, the values determined from pool samples are given and were converted to % dose values for the individual animals.

<sup>b</sup> TRR calculated from pooled samples as the sum of ERR and residue after solvent extract (used for calculation of metabolite concentrations)

<sup>c</sup> Not calculated

<sup>d</sup> Not sampled

<sup>e</sup> Faeces was homogenized with water prior LSC (approximately 4 to 8 parts water per part faeces)

<sup>f</sup> Abdominal fat referred as subcutaneous fat in the in-life report, back muscle referred as flank muscle

Recovered radioactivity from goat matrices were characterized and the results are shown in Tables 3 (benzoyl-label) and 4 (butylphenyl-label).

In urine of the goats given benzoyl-label, the main component B-1 accounted for 1.92 mg/kg or 47.0% TRR. Metabolite M9210I021 was the second most abundant component (0.61 mg/kg or 15.0% TRR). Metabolites M9210I029, M9210I030 and AB-1 were less abundant and accounted for up to 8.9% TRR.

In urine of the goats dosed with butylphenyl-label, metabolite M9210I014 was the main component and accounted for 1.26 mg/kg or 20.7% TRR. Metabolites M9210I021 and M9210I040 were the second most abundant components and accounted for 0.644 mg/kg or 11.0% TRR and 0.612 mg/kg and 10.1% TRR, respectively. Other identified metabolites were M9210I019, M9210I023, M9210I032, M9210I029, M9210I033, M9210I030, A-12, A-2 and AB-1 (each below or equal to 0.316 mg/kg or 5.2% TRR).

The extracted radioactive residues (ERR) in pooled samples of faeces of the goat administered with benzoyl-label (Days 1-12) and butylphenyl-label (Days 1-10) amounted to 78.7% and 95.5% of the TRR, respectively. In the faeces acetonitrile extract of the benzoyl-label metabolite B-1 was the main component

with 0.801 mg/kg or 63.9% TRR. The parent compound BAS 9210 I was also identified, although at significantly lower amounts, and accounted for 0.092 mg/kg or 7.4% TRR. In the faeces acetonitrile extract of the butylphenyl-label, the unchanged parent compound was the only component identified accounting for 0.588 mg/kg or 45.6% TRR.

The ERR in pooled samples of milk from the goats dosed with the benzoyl-label (Days 5-12) and butylphenyl-label (Days 5-10) amounted to 0.009 mg/kg or 51.0% TRR and 0.007 mg/kg or 93.5% TRR, respectively. In the acetonitrile extract of milk from the benzoyl-label dose, metabolite B-1 was the only component identified and accounted for 0.001 mg/kg or 4.5% TRR. In the solvent-extracted residues of milk from the butylphenyl-label dose, metabolite M9210I033 was the main component and accounted for 0.002 mg/kg or 30.0% TRR. The other identified metabolites, M9210I023, M9210I029 and M9210I030, were significantly less abundant and accounted for up to 7.8% of the TRR, less than 10% TRR.

In bile from the both experiments, the label-unspecific metabolites M9210I042, M9210I044 and M9210I043 were identified from 12.1% to 22.3% of the TRR. In bile from the benzoyl-label dose, B-1 was additionally identified accounting for 0.319 mg/kg or 22.4% TRR.

The ERR in pooled sample of liver from the benzoyl- and butylphenyl-label doses amounted to 0.276 mg/kg or 70.8% TRR and 0.182 mg/kg or 64.6% TRR. The residues after solvent extraction of liver (from benzoyl-label and butylphenyl-label) were further incubated with protease, whereby 14.0% and 17.4% of the TRR were released additionally. In the solvent extracted residues of liver from the benzoyl-label dose, metabolite B-1 was the main component accounting for 0.125 mg/kg or 32.0% of the TRR. M9210I042 accounted for 6.7% of the TRR. All components characterized in the ERR from the benzoyl-label dose by HPLC were below or equal to 3.3% TRR. Metabolite B-1 was also identified in the protease solubilizate (0.010 mg/kg or 2.5% TRR). In the solvent extracted residues of liver from the butylphenyl-label dose, M9210I042 accounted for 0.023 mg/kg or 8.3% TRR and M9210I043 for 0.014 mg/kg and 5.1% TRR. All components characterized in the ERR of the butylphenyl-label dose by HPLC were below or equal to 3.1% TRR. The characterized components in the protease solubilizate of the butylphenyl-label dose were each below or equal to 4.6% TRR.

The ERR in samples of kidney from the benzoyl- and butylphenyl-label doses amounted to 0.161 mg/kg or 85.1% TRR and 0.155 mg/kg or 95.5% TRR. The residue after solvent extraction of kidney (benzoyl-label) was further incubated with protease, whereby 9.4% of the TRR were released additionally. In the solvent extracted residues of kidney of the benzoyl-label dose, metabolite B-1 was the most abundant component with 0.102 mg/kg or 53.9% TRR. All components characterized in the ERR of the benzoyl-label dose by HPLC were below or equal to 5.3% TRR. In the acetonitrile extract of kidney of the butylphenyl-label dose, metabolite M9210I023 accounted for 0.017 mg/kg or 10.2% TRR. Other metabolites identified were M9210I032, M9210I033, M9210I030, M9210I014 and A-2, which were present at up to 6.7% of the TRR. All components characterized in the ERR of the butylphenyl-label by HPLC were below or equal to 8.0% TRR.

The ERR in samples of leg muscle from the benzoyl- and butylphenyl-label dose amounted to 0.008 mg/kg or 84.3% TRR and 0.004 mg/kg or 92.6% TRR. In samples of back muscle the ERR was 0.007 mg/kg or 78.4% and 0.003 mg/kg or 90.4%, respectively. In the acetonitrile extracts of leg muscle and back muscle of the benzoyl-label dose, metabolite B-1 was the main component accounting for 0.005 mg/kg or 50.5% TRR and 0.004 mg/kg or 46.5% TRR, respectively. Due to low residue levels, the muscle tissue extracts of the butylphenyl-label were not analysed by HPLC.

The ERR in samples of abdominal fat of the benzoyl- and butylphenyl-label dose amounted to 0.014 mg/kg or 93.2% TRR and 0.019 mg/kg or 96.3% TRR. In samples of kidney fat (benzoyl-label) the ERR was 0.014 mg/kg or 93.0%. In the acetonitrile extracts of abdominal and kidney fat of the benzoyl-label dose, metabolite B-1 was also the main component (0.006 mg/kg or 40.2% TRR and 0.006 mg/kg or 41.0% TRR), but additionally the parent compound was identified (0.003 mg/kg or 19.6% TRR and 0.003 mg/kg or 21.0% TRR). In the acetonitrile extract of abdominal fat of the butylphenyl-label dose, metabolite A-2 was the main component accounting for 0.008 mg/kg or 40.2% TRR.

The results of the above metabolism studies indicate that residue levels of Cyflumetofen and its metabolites were relatively low in edible tissues and milk except liver and kidney.

After repeated oral doses of benzoyl-label, metabolite B-1 was the predominant radioactive residue present in all of edible tissues tested and milk as well as excreta. It is the only metabolite present above 0.01 mg eq/kg and > 10% TRR in edible tissues or milk: in liver (0.125 mg eq/kg, 32.0% TRR) and kidney

(0.102 mg eq/kg, 53.9% TRR). M9210I042 was found at 0.026 mg eq/kg but 6.7% TRR. Cyflumetofen was found only in fat, among edible tissues, at 0.003 mg eq/kg, around 20%TRR. There were no identified metabolites in edible tissues or milk.

After repeated oral dose of butylphenyl-label, only M9210I023 was identified above 0.01 mg eq/kg and > 10% TRR in kidney (0.017 mg eq/kg, 10.2% TRR). M9210I043 and M9210I042 were found in liver at 0.014 mg eq/kg (5.1% TRR) and 0.023 mg eq/kg (8.3% TRR) respectively. M9210I033 was present in kidney at 0.011 mg eq/kg but 6.7% TRR. There were some other minor metabolites identified in edible tissues and milk but all were < 0.01 mg eq/kg. Cyflumetofen was not found in any edible tissues or milk.

From the metabolism studies, the metabolic pathway of cyflumetofen in lactating goats is speculated as:

- hydrolysis of the formic acid ester followed by hydrolytic cleavage of the trifluoromethylbenzoyl moiety resulting in metabolite B-1 (2-trifluoromethylbenzoic acid) and the labile metabolite M9210I014
- decarboxylation of metabolite M9210I014 resulting in metabolite A-2
- hydrolysis of the formic acid ester followed by decarboxylation yielding metabolite AB-1
- hydrolytic cleavage of the trifluoromethylbenzoyl moiety of metabolite AB-1 also leads to metabolite A-2
- glucuronidation of AB-1 to form M9210I042 and M9210I044
- hydroxylation yields M9210I043.

Oxidation, decarboxylation and hydroxylation of the t-butyl side chain seem to occur. Loss of nitrile from A-2 or its further metabolites also occur. Some conjugation of the carboxyl group of metabolites with the amino acid glycine, yielding conjugated metabolites also may occur.

Table 3 Extraction of radioactivity in milk, tissues, and faeces from goats given radioactive cyflumetofen

Matrix	TRR (measured)		Acetonitrile Extract <sup>a</sup>		Water extract <sup>a</sup>		ERR <sup>b</sup>		RRR <sup>c</sup>		TRR (calculated)
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg
Benzoyl-label											
Milk <sup>d</sup>	0.019	51.0	0.009	n.a.	n.a.	n.a.	0.009	51.0	0.009	49.0	0.018
Liver	0.404	59.8	0.233	59.8	0.045	11.4	0.276	70.8	0.114	29.2	0.390
Kidney	0.191	77.0	0.146	77.0	0.016	8.5	0.161	85.1	0.028	14.9	0.190
Abdominal fat	0.014	73.0	0.011	73.0	0.003	19.5	0.014	93.2	0.001	6.8	0.015
Kidney fat	0.014	74.7	0.011	74.7	0.003	18.3	0.014	93.0	0.001	7.0	0.015
Leg muscle	0.010	74.1	0.007	74.1	0.001	10.3	0.008	84.3	0.001	15.7	0.010
Back muscle	0.010	72.5	0.007	72.5	0.001	7.7	0.007	78.4	0.002	21.6	0.009
Faeces <sup>e</sup>	1.357	78.7	0.986	78.7	n.a.	n.a.	0.986	78.7	0.266	21.3	1.252
Butylphenyl-label											
Milk <sup>d</sup>	0.008	92.7	0.007	92.7	n.a.	n.a.	0.007	93.5	< 0.001	6.5	0.007
Liver	0.287	55.5	0.156	55.5	0.028	10.0	0.182	64.6	0.100	35.4	0.282
Kidney	0.167	81.2	0.132	81.2	0.023	13.9	0.155	95.5	0.007	4.5	0.163
Leg muscle	0.005	85.9	0.004	85.9	< 0.001	6.5	0.004	92.6	< 0.001	7.4	0.004
Back muscle	0.004	84.0	0.003	84.0	< 0.001	11.6	0.003	90.4	< 0.001	9.6	0.003
Abdominal fat	0.018	82.9	0.017	82.9	0.002	11.3	0.019	96.3	0.001	3.7	0.020
Faeces <sup>e</sup>	1.304	93.4	1.205	93.4	0.033	2.6	1.232	95.5	0.058	4.5	1.290

<sup>a</sup> TRR of pool samples<sup>b</sup> ERR = Extractable Radioactive Residue (calculated from individual samples)<sup>c</sup> RRR = Residual Radioactive Residue (residue after solvent extraction)<sup>d</sup> Pooled samples of milk (Day 5-12 for the benzoyl-label or 5-10 for the butylphenyl-label), extracted with 3 × acetonitrile<sup>e</sup> Pooled faeces sample (Day 1-12 for the benzoyl-label and 1-10 for the butylphenyl-label). Sample diluted with water.

Table 4(1) Characterization of radioactive residues in goat milk, liver, kidney, muscle, fat, urine, faeces and bile from goats dosed with benzoyl-label cyflumetofen

Metabolite/Fraction	Milk <sup>a</sup>		Liver		Kidney		Muscle <sup>b</sup>		Fat <sup>c</sup>		Urine <sup>d</sup>		Faeces <sup>a</sup>		Bile <sup>d</sup>	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Parent									0.003	19.6						
								0.003	21.0					0.092	7.4	
B-1	0.001	4.5	0.125	32.0	0.102	53.9	0.005	50.5	0.006	40.2	1.918	47.0	0.801	63.9	0.319	22.4
							0.004	46.5	0.006	21.0					0.200	14.0
M92101043											0.364	8.9				
M92101029											0.142	3.5				
M92101030											0.611	15.0				
M92101021																
M92101042			0.026	6.7											0.194	13.6
M92101044															0.172	12.1
AB-1											0.047	1.2				
Total Identified (ERR)	0.001	4.5	0.151	38.7	0.102	53.9	0.005	50.5	0.009	59.8	3.082	75.5	0.893	71.3	0.885	62.1
							0.004	46.5	0.009	62.0						
Total Characterized by HPLC (ERR)	0.004	19.6	0.070	17.9	0.024	12.6	0.002	17.2	0.001	10.0	1.002	24.5	0.093	7.4	0.541	37.9
							0.002	19.8	0.002	10.8						
RRR after solvent extraction	0.009	49.0	0.114	29.2	0.028	14.9	0.001	15.7	0.001	6.8	e	e	0.266	21.3	e	e
							0.002	21.6	0.001	7.0						
Total Characterized	0.007	41.2	0.163	41.7	0.063	33.2	0.003	28.4	0.005	31.7	1.002	24.5	0.093	7.4	0.541	37.9
							0.002	27.5	0.005	31.0						
Total Identified and/or Characterized	0.008	45.7	0.313	80.4	0.165	87.1	0.008	78.9	0.013	91.5	4.084	100.0	0.986	78.7	1.426	100.0
							0.007	74.0	0.014	93.0						
Grand total [%]	0.017	94.7	0.361	92.6	0.173	91.4	0.009	94.6	0.014	98.3	4.084	100.0	1.252	100.0	1.426	100.0
							0.009	95.6	0.015	100.0						

<sup>a</sup> Analysis after extraction with acetonitrile.<sup>b</sup> Individual values for leg (top) and back (bottom) muscle are reported.<sup>c</sup> Individual values for abdominal (top) and kidney fat (bottom) are reported.<sup>d</sup> Analysed after direct injection.<sup>e</sup> Not applied.

Table 4(2) Characterization of radioactive residues in goat milk, liver, kidney, muscle, fat, urine, faeces and bile from goats dosed with butylphenyl-label cyflumetofen

Metabolite/Fraction	Milk <sup>a</sup>		Liver		Kidney		Fat <sup>b</sup>		Urine <sup>c</sup>		Faeces		Bile <sup>c</sup>	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Parent											0.588	45.6		
M92101040									0.612	10.1				
M92101019									0.214	3.5				
M92101023	0.0005	6.7		10.2	0.017				0.312	5.3				
M92101032				3.2	0.005				0.165	2.7				
M92101043			0.014	5.1									0.705	22.3
M92101029	0.0005	6.8							0.316	5.2				
M92101033	0.0021	30.0			0.011	6.7			0.306	5.0				
M92101030	0.0006	7.8			0.007	4.1			0.137	2.3				
M92101021									0.664	11.0				
M92101042			0.023	8.3									0.592	18.7
A-12									0.221	3.7				
M92101014					0.007	4.2			1.257	20.7				
M92101044													0.462	14.6
A-2					0.009	5.3		0.008	40.2	1.3				
AB-1										0.4				
Total Identified (ERR)	0.0036	51.3	0.038	13.4	0.055	33.7	0.008	40.2	4.320	71.1	0.588	45.6	1.759	55.6
Total Characterized by HPLC (ERR)	0.0026	36.3	0.103	36.8	0.068	41.5	0.006	28.6	1.759	28.9	0.605	46.9	1.406	44.4
RRR after solvent extraction	0.0005	6.5	0.100	35.4	0.007	4.5	0.001	3.7	<sup>d</sup>	<sup>d</sup>	0.058	4.5	<sup>d</sup>	<sup>d</sup>
Total Characterized	0.0029	40.6	0.176	62.5	0.093	56.9	0.010	48.0	1.759	28.9	0.638	49.4	1.406	44.4
Total Identified and/or Characterized	0.0065	91.8	0.213	75.8	0.147	90.6	0.018	88.2	6.079	100.0	1.226	95.0	3.164	100.0
Grand total [%]	0.0070	98.4	0.254	90.1	0.155	95.1	0.018	91.9	6.079	100.0	1.283	99.5	3.164	100.0

<sup>a</sup> Analysis after extraction with acetonitrile<sup>b</sup> Abdominal fat sample<sup>c</sup> HPLC analysed by direct injection<sup>d</sup> Not applied

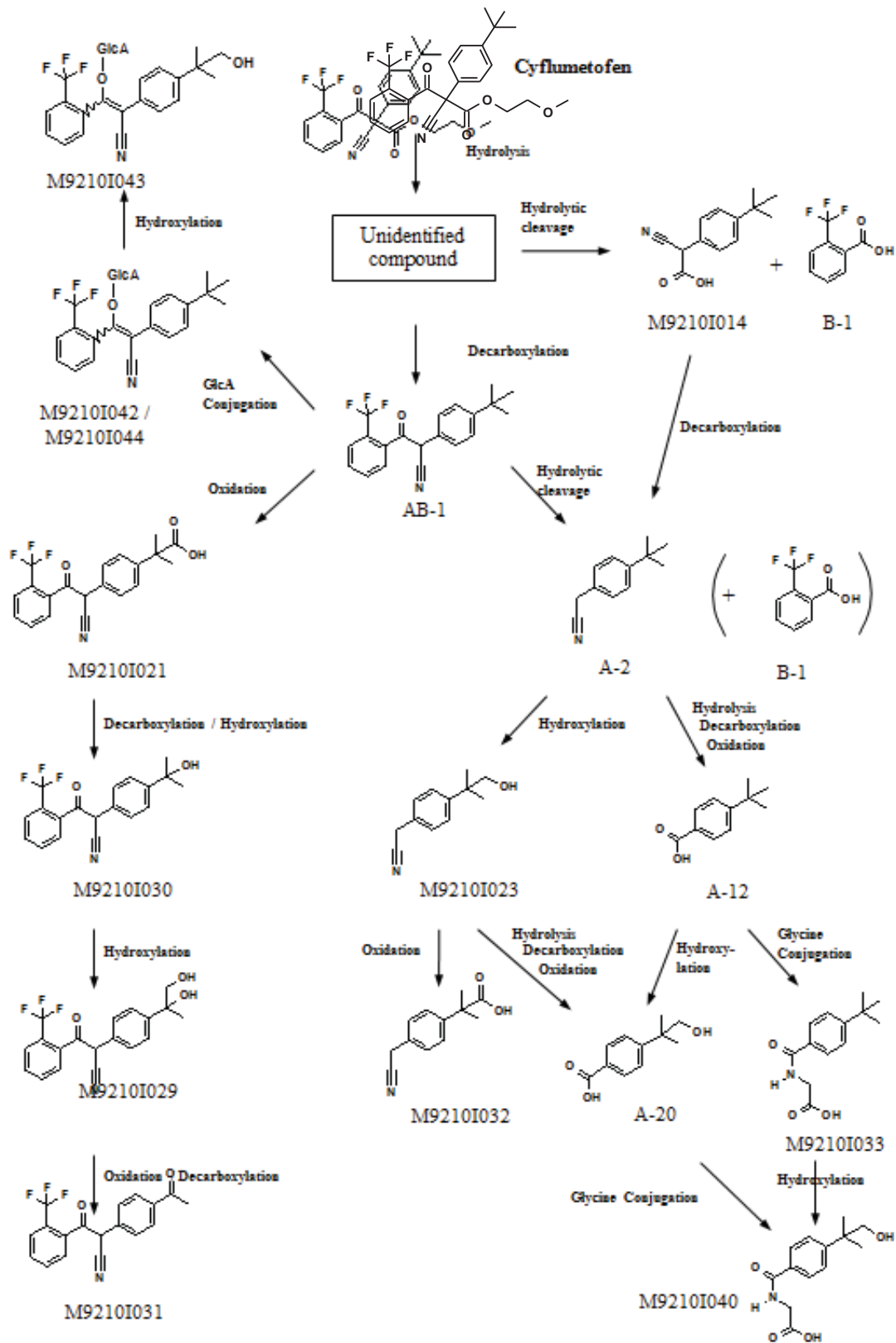


Figure 1 Proposed metabolic pathway of cyflumetofen in goats

### *Plant metabolism*

The Meeting received information on the fate of cyflumetofen in Satsuma mandarin, apple and eggplant.

#### *Satsuma mandarin (Hayashi, 2004a)*

Two individual Satsuma mandarin trees were sprayed with formulated butylphenyl-label or benzoyl-label at an equivalent application rate of 0.600 kg ai/ha. The concentration of the application solutions was 16 mg cyflumetofen in 0.080 L blank formulation. The concentration of the blank formulation was 159 mg formulation in 0.199 L of water. The amount of spray solution per tree was 0.0369 L. The growth stage of the Satsuma mandarin fruit was just before fruit maturation. Satsuma mandarin trees, each tree separately, were kept in climate controlled greenhouses. The trees were irrigated and fertilizer (N-P-K-Mg) was added as needed. No other pesticides were used. The achieved application rate was 89.2 to 92.9% of the target.

For both treatments with the two radio-labelled cyflumetofen, fruit samples were taken at 1, 7 and 30 DAT and leaf samples at 1, 7 and 14 DAT. Four fruits were sampled at random per sampling point and pooled per two fruits. Twenty leaves were randomly sampled per sampling points and pooled per 10 leaves. Untreated fruit and leaf samples were taken at the final harvest point. Fruit and leaf samples were immediately surface rinsed with acetonitrile.

Total radioactivity in the rinse was determined by LSC. The surface rinse was concentrated by evaporation to dryness, and then redissolved prior to analysis by HPLC. The rinsed fruit was separated into peel and pulp fractions. Surface washed leaf, peel and pulp samples were homogenized with dry ice. Homogenates were extracted twice with acetonitrile:water (8:2 v/v). The extracts, after filtration, were combined and radioactivity was determined by LSC. Extracts were further concentrated (volume reduction by rotary-evaporation) prior to analysis by HPLC. Solid residues were dried and radioactivity was determined by combustion/LSC. Total radioactivity was determined as the sum of surface rinse, extracts and solid residues.

Identification was achieved by HPLC and TLC, and comparison with reference standards. Confirmation was also achieved by LC-MS/MS analysis of isolated fractions. HPLC recoveries were determined and were always > 86%. The procedural recovery of the extraction method was 96% for cyflumetofen and 100.4% for total radioactivity, demonstrating the applicability of the employed methods. Storage stability was demonstrated over the experimental period.

The results for distribution and identification of radioactivity in Satsuma mandarin fruit and leaves are presented in the following tables.

Total radioactive residues (TRR) in fruit were 0.58–0.62 mg eq/kg on 1 DAT and 0.57–0.57 mg eq/kg on 30 DAT. Distribution of radioactivity for butylphenyl- or benzoyl-label applications were similar. The majority of the radioactivity was found in the surface rinse: 95–96% of the TRR on 1 DAT and 88–89% of the TRR on 30 DAT. In peel, 4.3–4.8% of the TRR was recovered on 1 DAT, increasing to 10.8–11.5% of the TRR on 30 DAT (9.1–10.1% of the TRR extracted from peel on 30 DAT). Radioactive residues in flesh were very low ( $\leq 0.6\%$  of the TRR).

TRR in leaves were 33–35 mg eq/kg on 1 DAT and 28–38 mg eq/kg on 14 DAT. Distribution of radioactivity for butylphenyl- or benzoyl-label applications was similar. The majority of the radioactivity was found in the surface rinse: 95–97% of the TRR on 1 DAT and 87–94% of the TRR on 30 DAT. Leaf residues were 3.4–4.9% of the TRR on 1 DAT, increasing to 5.6–12.9% of the TRR on 30 DAT (4.9–12.4% of the TRR extracted from leaves on 14 DAT).

Cyflumetofen was by far the predominant identified fraction in fruit (sum of rinse and peel extracts): 88–90% of the TRR on 1 DAT decreasing to 44–54% of the TRR on 30 DAT. Other identified metabolites were AB-6, AB-7, A-12 and B-1 (all  $\leq 1.6\%$  TRR on 1 DAT and maximum 8.6, 8.5, 4.4 and 11.2% of the TRR, respectively on 30 DAT). It is gradually metabolized over time to produce metabolites B-1, AB-6 and AB-7. However, even 30 DAT, the parent accounted for about 50% TRR.



Cyflumetofen was also the predominant identified fraction in leaves (sum of rinse and extracts): 89–90% of the TRR on 1 DAT decreasing to 73-81% of the TRR on 30 DAT. Other identified metabolites were AB-6, AB-7, A-12 and B-1 (all  $\leq 4.8\%$  of the TRR on 1 DAT and maximum 4.2, 1.5, 3.6 and 9.1% of the TRR, respectively on 30 DAT).

Table 5 Distribution of radioactivity in Satsuma mandarin fruit and leaf after treatment with radio-labelled cyflumetofen

Sample	Day-1		Day-7		Day-30 (fruit) / Day-14 (leaves)	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
<b>Butylphenyl-label</b>						
Fruit						
Surface rinse	0.5539	95.62	0.4173	93.00	0.5067	88.77
Peel residues	0.0239	4.32	0.0307	6.85	0.0622	10.86
Extracted	0.0207	3.72	0.0256	5.70	0.0523	9.12
Unextracted	0.0032	0.60	0.0051	1.15	0.0099	1.73
Flesh residues	0.0004	0.06	0.0007	0.15	0.0020	0.37
Extracted	0.0004	0.05	0.0006	0.13	0.0018	0.33
Unextracted	0.0000	0.01	0.0001	0.02	0.002	0.04
Total radioactive residues	0.5783	100.00	0.4487	1000.00	0.5709	100.00
Leaves						
Surface rinse	34.8679	96.60	29.5027	93.54	28.3494	94.44
Leaf residues	1.2257	3.40	2.0377	6.46	1.6639	5.56
Extracted	1.1090	3.08	1.8256	5.79	1.4737	4.92
Unextracted	0.1167	0.32	0.2121	0.67	0.1901	0.64
Total radioactive residues	36.0936	100.00	31.5404	100.00	30.0132	100.00
Sample	Day-1		Day-7		Day-30 (fruit) / Day-14 (leaves)	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
<b>Benzoyl-label</b>						
Fruit						
Surface rinse	0.5868	95.02	0.3826	91.40	0.5041	87.87
Peel residues	0.0295	4.84	0.0349	8.31	0.0662	11.53
Extracted	0.0268	4.32	0.0316	7.54	0.0581	10.11
Unextracted	0.0027	0.52	0.0032	0.77	0.0081	1.42
Flesh residues	0.0010	0.14	0.0012	0.29	0.0034	0.60
Extracted	0.0009	0.14	0.0011	0.27	0.0033	0.58
Unextracted	0.0000	0.00	0.0001	0.01	0.0001	0.02
Total radioactive residues	0.6173	100.00	0.4187	100.00	0.5737	100.00
Leaves						
Surface rinse	33.3370	95.09	30.7705	91.23	37.6579	87.14
Leaf residues	1.7188	4.91	2.9824	8.77	5.4724	12.86
Extracted	1.6259	4.65	2.8039	8.23	5.2696	12.37
Unextracted	0.0929	0.27	0.1785	0.53	0.2028	0.49
Total radioactive residues	35.0558	100.00	33.7529	100.00	43.1303	100.00

Table 6 Identification of radioactive residues in Satsuma mandarin surface rinse and peel extracts

Identified compound	Surface rinse		Peel extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
<b>Butylphenyl-label</b>						
Day 1						
A-12	0.0040	0.71	0.0062	1.17	0.0102	1.88
AB-6	0.0053	0.92	0.0008	0.16	0.0061	1.07
AB-7	0.0034	0.59	0.0005	0.09	0.0039	0.69
Parent	0.5174	89.25	0.0037	0.58	0.5211	89.83
Unidentified peaks total	0.0238	4.15	0.0095	1.72	0.0333	5.88
Total	0.5539	95.62	0.0207	3.72	0.5746	99.35
Day 7						
A-12	0.0047	1.04	0.0076	1.69	0.0122	2.73
AB-6	0.0084	1.88	0.0010	0.23	0.0095	2.11
AB-7	0.0037	1.50	0.0006	0.13	0.0073	1.63
Parent	0.3674	81.88	0.0033	0.74	0.3708	82.62
Unidentified peaks total	0.0331	6.70	0.0131	2.91	0.0431	9.61
Total	0.4173	93.00	0.0256	5.70	0.4429	98.70
Day 30						
A-12	0.0132	2.32	0.0119	2.04	0.0250	4.36
AB-6	0.0392	6.95	0.0032	0.57	0.0424	7.51
AB-7	0.0389	6.93	0.0015	0.26	0.0404	7.19
Parent	0.3064	53.26	0.0043	0.77	0.3107	54.03
Unidentified peaks total	0.109	19.31	0.0314	5.48	0.1405	24.81
Total	0.5067	88.77	0.0523	9.12	0.5590	97.90
<b>Benzoyl-label</b>						
Day 1						
B-1	0.0097	1.55	0.0185	3.07	0.0282	4.66
AB-6	0.0055	0.89	0.0006	0.10	0.0061	0.99
AB-7	0.0026	0.45	0.0002	0.03	0.0028	0.48
Parent	0.5434	87.94	0.0032	0.46	0.5466	88.40
Unidentified peaks total	0.0256	4.19	0.0043	0.66	0.0299	4.81
Total	0.5868	95.02	0.0268	4.32	0.6136	99.34
Day 7						
B-1	0.0070	1.67	0.0206	4.91	0.0276	6.58
AB-6	0.0093	2.22	0.0010	0.23	0.0103	2.45
AB-7	0.0077	1.84	0.0005	0.12	0.0082	1.97
Parent	0.3282	78.40	0.0020	0.49	0.3303	78.89
Unidentified peaks total	0.0304	7.27	0.0075	1.79	0.0378	9.05
Total	0.3826	91.40	0.0316	7.54	0.4142	98.94
Day 30						
B-1	0.0248	4.32	0.0394	6.86	0.0642	11.17
AB-6	0.0474	8.26	0.0020	0.35	0.0494	8.61
AB-7	0.0481	8.37	0.0007	0.12	0.0488	8.49
Parent	0.2491	43.48	0.0021	0.37	0.2512	43.85
Unidentified peaks total	0.1347	23.44	0.0139	2.41	0.1486	25.87
Total	0.5041	87.87	0.0581	10.11	0.5622	97.99

Data are the mean of duplicate experiments.

#### *Apple (Dohn, 2004a)*

One apple tree grown within a commercial apple orchard in California, USA, was sprayed with formulated butylphenyl- or benzoyl-label at an equivalent application rate of 0.600 kg ai/ha.

Application took place in October 2002. The concentration of the application solutions was approximately 225 (butylphenyl-label) and 211 mg/L (benzoyl-label) cyflumetofen in blank formulation. The achieved application rate was 115 to 120% of the target. Storage stability was demonstrated over the experimental period. At the time of application, fruits were near maturity. The plots were (drip) irrigated and fertilizer (N-P-K and Ca) was added as needed. Other pesticides were used prior to application of cyflumetofen, which does not interfere with cyflumetofen. Fruit samples were taken at 1, 7 and 30 DAT (for both labels and untreated plot). Leaf samples were taken at 7 and 30 DAT (also for both labels and untreated plot).

Fruit and leaf samples were immediately surface rinsed two times with acetonitrile. Total radioactivity in the rinse was determined by LSC. The surface rinse was concentrated prior to analysis by HPLC/TLC. The rinsed fruit and leaves were homogenized with dry ice and liquid nitrogen. Homogenates were extracted twice with acetonitrile and twice with acetonitrile:water (1:1 v/v). The extracts after centrifugation were combined and radioactivity was determined by LSC. Extracts were further concentrated prior to analysis by HPLC/TLC. Solid residues were dried and radioactivity was determined by combustion/LSC. Total radioactivity was determined as the sum of surface rinse, extracts and solid residues.

Identification was achieved by HPLC and TLC, and comparison with reference standards. Some extracts were further characterized by reversed phase solid phase extraction and LSC/HPLC analysis of eluates.

Total radioactive residues (TRR) in apple fruit were 0.100–0.113 mg eq/kg on 1 DAT and 0.079–0.057 mg eq/kg on 30 DAT. Results for butylphenyl- and benzoyl-label were similar. For the butylphenyl- and benzoyl-label, the majority of the radioactivity was found in the surface rinse: 95 and 96% of the TRR, respectively on 1 DAT and 71 and 67% of the TRR, respectively on 30 DAT. Extracted radioactivity was not determined on 1 DAT due to low levels of radioactivity, and was 15 and 9.6% of the TRR on 7 DAT for butylphenyl- and benzoyl-label respectively and 22 and 28% of the TRR on 30 DAT, respectively. Unextracted radioactivity was low: 5.3–7.6% of the TRR on 30 DAT.

TRR in apple leaves were 6.1–7.3 mg eq/kg on 7 DAT and 4.9–9.5 mg eq/kg on 30 DAT. Surface rinse removed less radioactivity from leaves for the benzoyl-label compared to the butylphenyl-label. For the butylphenyl- and benzoyl-label, the majority of the radioactivity was found in the surface rinse: 91 and 87% of the TRR, respectively on 7 DAT and 82 and 72% of the TRR, respectively on 30 DAT. Extracted radioactivity was 13 and 21% of the TRR on 30 DAT for butylphenyl- and benzoyl-label respectively. Unextracted radioactivity was low: 4.9–6.7% of the TRR on 30 DAT.

Cyflumetofen was the predominant identified compound in apple fruit (sum of rinse and extracts): 89–95% of the TRR on 1 DAT decreasing to 53–65% of the TRR on 30 DAT. Other identified metabolites were AB-6, AB-7 and B-1 (all  $\leq$  5% of the TRR on 1 DAT and maximum 5.3, 6.3 and 1.8% of the TRR, respectively on 30 DAT).

Cyflumetofen was also the predominant identified compound in apple leaves (sum of rinse and extracts): 77–85% of the TRR on 7 DAT and 44–60% of the TRR on 30 DAT. Other identified metabolites were AB-6, AB-7 and B-1 at maximum 8.6, 5.6 and 4.8% of the TRR, respectively on 30 DAT.

The results for apple fruit and leaves (distribution and identification of radioactivity for both labels) are presented in the following tables.

Table 7 Distribution of radioactivity in apple fruit and leaf after treatment with radio-labelled cyflumetofen

DAT	Surface rinse		Extracted		Unextracted			Total
	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg
Butylphenyl-label								
Apple fruit								
Day 1	95.0	0.095	Np	np	5.0	0.005	100	0.10
Day 7	82.1	0.064	15.4	0.012	2.6	0.002	100	0.078
Day 30	70.9	0.056	21.5	0.017	7.6	0.006	100	0.079
Leaves								
Day 7	90.8	5.535	7.9	0.479	1.4	0.085	100	6.099
Day 30	82.0	4.046	13.1	0.644	4.9	0.242	100	4.932
Benzoyl-label								
Apple fruit								
Day 1	95.6	0.108	Np	np	4.4	0.005	100	0.113
Day 7	89.2	0.149	9.6	0.016	1.2	0.002	100	0.167
Day 30	66.7	0.038	28.1	0.016	5.3	0.003	100	0.057
Leaves								
Day 7	86.8	6.306	11.6	0.843	1.6	0.117	100	7.266
Day 30	72.0	6.884	21.3	2.039	6.7	0.641	100	9.564

Table 8 Identification of radioactive residues in apple fruit surface rinse and extracts

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Butylphenyl-label						
Day 1						
AB-6	0.005	5.0			0.005	5.0
AB-7						
Parent	0.061	61.0	Not extracted due to low radioactive residues: < 0.01 mg eq/kg		0.061	61.0
Breakdown of parent	0.028	28.0			0.028	28.0
<sup>14</sup> C in Extract (mg eq/kg)	0.095				0.095	
Day 7						
AB-6	0.001	1.3			0.001	1.3
AB-7						
Parent	0.059	75.6	0.002	2.6	0.061	78.2
Minor	0.004	5.1	0.003	3.8	0.007	8.9
Unknown			0.006	7.7	0.006	7.7
<sup>14</sup> C in Extract (mg eq/kg)	0.064		0.012		0.076	
Day 30						
AB-6	0.003	3.8	0.001	1.3	0.004	5.1
AB-7	0.002	2.5	0.003	3.8	0.005	6.3
Parent	0.038	48.1	0.004	5.1	0.042	53.2
Minor	0.013	16.5	0.009	11.4	0.022	27.9
<sup>14</sup> C in Extract (mg eq/kg)	0.056		0.016		0.072	
Benzoyl-label						
Day 1						
AB-6						
AB-7						
Parent	0.066	58.4	Not extracted due to low radioactive residues: < 0.01 mg eq/kg		0.066	58.4
Breakdown of parent	0.041	36.3			0.041	36.3
<sup>14</sup> C in Extract (mg eq/kg)	0.108				0.108	

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Day 7						
B-1	0.002	1.2			0.002	1.2
AB-6						
AB-7	0.003	1.8	0.006	3.6	0.009	5.4
Parent	0.137	82.0	0.003	1.8	0.14	83.8
Minor	0.006	3.6	0.006	3.6	0.012	7.2
<sup>14</sup> C in Extract (mg eq/kg)	0.149		0.016		0.165	
Day 30						
B-1			0.001	1.8	0.001	1.8
AB-6	0.001	1.8	0.002	3.5	0.003	5.3
AB-7	0.001	1.8	0.002	3.5	0.003	5.3
Parent	0.029	50.9	0.008	14.0	0.037	64.9
Minor	0.001	1.8	0.003	5.3	0.004	7.1
Unknown	0.005	8.8			0.005	8.8
<sup>14</sup> C in Extract (mg eq/kg)	0.038		0.016		0.054	

“Minor” refers to <sup>14</sup>C peaks in the chromatogram with a concentration < 0.005 mg eq/kg in the extract. This refers to a total of all minor components.

Table 9 Identification of radioactive residues in apple leaf surface rinse and extracts

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Butylphenyl-label						
Day 7						
AB-6	0.133	2.2	0.031	0.5	0.164	2.7
AB-7	0.188	3.1	0.032	0.5	0.22	3.6
Parent	4.904	80.4	0.272	4.5	5.176	84.9
Minor	0.310	5.1	0.144	2.4	0.454	7.5
Unknown						
<sup>14</sup> C in Extract (mg eq/kg)	5.535		0.479		6.014	
Day 30						
AB-6	0.267	5.4	0.067	1.4	0.334	6.8
AB-7	0.210	4.3	0.028	0.6	0.238	4.9
Parent	2.881	58.4	0.086	1.7	2.967	60.1
Minor	0.688	13.9			0.688	13.9
<sup>14</sup> C in Extract (mg eq/kg)	4.046		0.644		4.69	
Benzoyl-label						
Day 7						
B-1	0.120	1.7	0.140	1.9	0.26	3.6
AB-6	0.290	4.0	0.050	0.7	0.34	4.7
AB-7	0.265	3.6	0.044	0.6	0.309	4.2
Parent	5.278	72.6	0.333	4.6	5.611	77.2
Minor	0.353	4.9			0.353	4.9
<sup>14</sup> C in Extract (mg eq/kg)	6.306		0.843		7.149	
Day 30						
B-1	0.248	2.6	0.208	2.2	0.456	4.8
AB-6	0.688	7.2	0.139	1.5	0.827	8.7
AB-7	0.454	4.7	0.077	0.8	0.531	5.5
Parent	3.979	41.6	0.208	2.2	4.187	43.8
Minor	0.317	3.3	0.812	8.5	1.129	11.8
Unknown	1.198 (4 peaks)	12.5	0.595 (2 peaks)	6.2	1.793	18.7
<sup>14</sup> C in Extract (mg eq/kg)	6.884		2.039		8.923	

“Minor” refers to <sup>14</sup>C peaks in the chromatogram with a concentration < 0.200 mg eq/kg in the extract. This refers to a

total of all minor components.

#### *Eggplant (Dohn, 2004b)*

Eggplants grown on two plots (1.1–1.5 m<sup>2</sup>) located in California and containing six to eight eggplants were sprayed with formulated butylphenyl- or benzoyl-label cyflumetofen at an equivalent application rate of 0.600 kg ai/ha. Application took place on August 2002. The concentration of the application solutions was approximately 140 (butylphenyl-label) and 220 mg/L (benzoyl-label) cyflumetofen in blank formulation. The achieved application rate was 98 to 114% of the target. Storage stability was demonstrated over the experimental period. The growth stage of the eggplant fruit was from just pollinated to mature. The eggplants were grown outdoors. The plots were (drip) irrigated and fertilizer (N-P-K-Ca-S) was added as needed. Other pesticides were used which do not interfere with cyflumetofen. Fruit samples were taken at 1, 7 and 14 DAT (for both labels and untreated plot). Leaf samples were taken at 14 DAT (also for both labels and untreated plot).

Fruit and leaf samples were immediately surface rinsed two times with acetonitrile. Total radioactivity in the rinse was determined by LSC. The surface rinse was concentrated, if necessary, prior to analysis by HPLC. The rinsed fruit and leaves were homogenized with dry ice. Homogenates were extracted twice with acetonitrile and twice with acetonitrile:water (1:1 v/v). The extracts, after centrifugation, were combined and radioactivity was determined by LSC. Extracts were further concentrated, if necessary by volume reduction by rotary-evaporation, prior to analysis by HPLC. Solid residues were dried and radioactivity was determined by combustion/LSC. Total radioactivity was determined as the sum of surface rinse, extracts and solid residues.

Identification was achieved by HPLC and 2D-TLC, and comparison with reference standards. Some degradates (U4) were identified by LC-MS/MS analysis of isolated fractions or were subjected to acid hydrolysis (U1 and U2) with 1 N HCl at 80 °C to release acid labile conjugates.

Total radioactive residues (TRR) in eggplant fruit were 0.323–0.488 mg eq/kg on 1 DAT and 0.315–0.413 mg eq/kg on 14 DAT. Surface rinse removed less radioactivity from fruit for the benzoyl-label compared to the butylphenyl-label. For the butylphenyl- and benzoyl-label, the majority of the radioactivity was found in the surface rinse: 92 and 86% of the RR, respectively on 1 DAT and 81 and 56% of the TRR, respectively on 14 DAT. Extracted radioactivity was 7.1 and 13% TRR on 1 DAT for butylphenyl- and benzoyl-label respectively and 15 and 41% TRR on 14 DAT, respectively. Unextracted radioactivity was low: 2.7–4.1% of the TRR on 14 DAT.

TRR in eggplant leaves were 17–23 mg eq/kg on 14 DAT. Surface rinse removed less radioactivity from leaves for the benzoyl-label compared to the butylphenyl-label. For the butylphenyl- and benzoyl-label, the majority of the radioactivity was found in the surface rinse: 83 and 69% of the TRR, respectively on 14 DAT. Extracted radioactivity was 14 and 27% of the TRR on 14 DAT for butylphenyl- and benzoyl-label respectively. Unextracted radioactivity was low: 2.6–4.7% of the TRR on 14 DAT.

Cyflumetofen was the predominant identified compound in eggplant fruit (sum of rinse and extracts): 91–95% of the TRR on 1 DAT decreasing to 42–62% of the TRR on 14 DAT. Other identified metabolites were AB-6, AB-7, B-1, U4 and U1 and U2 (acid labile conjugates of B-1) (all ≤ 2.5% TRR on 1 DAT and maximum 3.4, 3.6, 14.8, 1.2, 16.2 and 6.3% of the TRR, respectively on 14 DAT).

Cyflumetofen was also the predominant identified compound in eggplant leaves (sum of rinse and extracts): 47–58% of the TRR on 14 DAT. Other identified metabolites were AB-6, AB-7, B-1, U4 (structure proposed) and U1 (conjugate of B-1) and U2 (conjugate of B-1) at 8.1, 5.7, 4.6, 4.3, 4.1 and 1.4% of the TRR, respectively on 14 DAT).

The results for eggplant fruit and leaves (distribution and identification of radioactivity for both labels) are presented in the following tables.

Table 10 Distribution of radioactivity in eggplant fruit and leaf after treatment with radio-labelled cyflumetofen

DAT	Surface Rinse		Extracted		Unextracted		Total	
	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg
Butylphenyl-label								
Eggplant fruit								
Day 1	92.0	0.297	7.1	0.023	0.9	0.003	100	0.323
Day 7	86.1	0.323	11.5	0.043	2.4	0.009	100	0.375
Day 14	81.3	0.256	14.6	0.046	4.1	0.013	100	0.315
Leaves								
Day 14	83.4	19.144	14.1	3.232	2.6	0.592	100	22.968
Benzoyl-label								
Eggplant fruit								
Day 1	86.5	0.422	12.7	0.062	0.8	0.004	100	0.488
Day 7	79.2	0.442	19.4	0.108	1.4	0.008	100	0.558
Day 14	56.4	0.233	40.9	0.169	2.7	0.011	100	0.413
Leaves								
Day 14	68.7	12.001	26.6	4.644	4.7	0.818	100	17.463

Table 11 Identification of radioactive residues in eggplant fruit surface rinse and extracts

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Butylphenyl-label						
Day 1						
Parent	0.294	91.0	0.013	4.0	0.307	95
Minor	0.003	0.9	0.010	3.1	0.013	4
<sup>14</sup> C in Extract (mg eq/kg)	0.297		0.023			
Day 7						
AB-6	0.033	8.8			0.033	8.8
AB-7	0.018	4.8			0.018	4.8
Parent	0.240	64.0	0.011	2.9	0.251	66.9
U4(tentatively identified)	0.022	5.9			0.022	5.9
Minor	0.010	2.7	0.013	3.5	0.023	6.2
Unknown			0.019 (3 peaks)	5.1	0.019	5.1
<sup>14</sup> C in Extract (mg eq/kg)	0.323		0.043			
Day 14						
AB-6	0.016	5.1			0.016	5.1
AB-7	0.016	5.1			0.016	5.1
Parent	0.188	59.7	0.008	2.5	0.196	62.2
U4(tentatively identified)	0.011	3.5			0.011	3.5
Minor	0.025	7.9			0.025	7.9
Unknown			0.019 (2 peaks)	6.0	0.016	6.0
<sup>14</sup> C in Extract (mg eq/kg)	0.256		0.046			
Benzoyl-label						
Day 1						
B-1			0.012	2.5	0.012	2.5
Parent	0.408	83.6	0.037	7.6	0.445	91.2
Minor	0.014	2.9	0.008	1.6	0.022	4.5
Unknown			0.005	1.0	0.005	1.0

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
<sup>14</sup> C in Extract (mg eq/kg)	0.422		0.062			
Day 7						
B-1	0.011	2.0	0.048	8.6	0.059	10.6
AB-6	0.017	3.0			0.017	3
AB-7	0.016	2.9			0.016	2.9
Parent	0.387	69.4	0.007	1.3	0.394	70.7
U1 (conjugate of B-1)			0.034	6.1	0.034	6.1
U2 (conjugate of B-1)			0.011	2.0	0.011	2
Minor	0.007	1.3			0.007	1.3
Unknown	0.005	0.9			0.005	0.9
<sup>14</sup> C in Extract (mg eq/kg)	0.442		0.108			
Day 14						
B-1			0.061	14.8	0.061	14.8
AB-6	0.014	3.4			0.014	3.4
AB-7	0.015	3.6			0.015	3.6
Parent	0.173	41.9	0.002	0.5	0.175	42.4
U4(tentatively identified)	0.005	1.2			0.005	1.2
U2 (conjugate of B-1)			0.026	6.3	0.026	6.3
U1 (conjugate of B-1)			0.067	16.2	0.067	16.2
Minor	0.020	4.8	0.014	3.4	0.034	8.2
Unknown	0.005	1.2			0.061	14.8
<sup>14</sup> C in Extract (mg eq/kg)	0.233		0.169			

“Minor” refers to <sup>14</sup>C peaks in the chromatogram with a concentration < 0.005 mg eq/kg in the extract. This refers to a total of all minor components.

Table 12 Identification of radioactive residues in eggplant leaf surface rinse and extracts

Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Butylphenyl-label						
Day 14						
AB-6	2.068	9.0	0.278	1.2	2.346	10.2
AB-7	1.455	6.3	0.110	0.5	1.565	6.8
Parent	12.061	52.5	1.173	5.1	13.234	57.6
U4(tentatively identified)	0.861	3.7			0.861	3.7
U3	0.823	3.6			0.823	3.6
Minor	0.249	1.1	0.931	4.1	1.18	5.2
Unknown	1.436 (3 peaks)	6.2	0.740	3.2	2.176	9.4
<sup>14</sup> C in Extract (mg eq/kg)	19.144		3.232			
Benzoyl-label						
Day 14						
B-1	0.240	1.4	0.571	3.3	0.811	4.7
AB-6	0.996	5.7	0.418	2.4	1.414	8.1
AB-7	0.876	5.0	0.121	0.7	0.997	5.7
Parent	6.937	39.7	1.333	7.6	8.27	47.3
U4(tentatively identified)	0.480	2.7	0.279	1.6	0.759	4.3
U3	0.504	2.9			0.504	2.9
U2 (conjugate of B-1)			0.251	1.4	0.251	1.4
U1 (conjugate of B-1)			0.715	4.1	0.715	4.1
Minor	0.516	3.0			0.516	3



Identified compound	Surface rinse		Extract		Total	
	mg eq./kg	% TRR	mg eq./kg	% TRR	mg eq./kg	% TRR
Unknown	1.452 (4 peaks)	8.3			1.452	8.3
<sup>14</sup> C in Extract (mg eq/kg)	12.001		4.644			

“Minor” refers to <sup>14</sup>C peaks in the chromatogram with a concentration < 0.200 mg eq/kg in the extract. This refers to a total of all minor components.

#### Proposed metabolic pathway of cyflumetofen in plants

In the metabolism studies on Satsuma mandarin, apple and eggplant, radioactive residues after treatment were at significantly higher concentrations in surface rinse of the respective fruits than their extracts. In all plants, cyflumetofen was the predominant residue. Metabolite B-1 was found in the fruit extract of apple (30 DAT) and eggplant (14 DAT) at 11.2% and 14.8% of TRR corresponding to 0.064 and 0.061 mg eq/kg respectively. No other individual components accounted for > 10% TRR. At smaller amounts, metabolites AB-6 and AB-7 were often identified in fruits.

Considering the results of the plant metabolism studies, the metabolic pathway of cyflumetofen in plants was speculated as follows:

- Cyflumetofen is photodegraded to AB-7, hydrolysed to B-1, or its nitrile is hydrolysed to amide followed by acyl migration to produce AB-6.
- The side chain of AB-7 is hydrolysed and oxidized to produce U4 (identified only in eggplant).
- B-1 conjugates to produce U1 and U2 (found only in eggplant).
- B-1 may also be produced from AB-6 and AB-7. A-12 (found only in Satsuma mandarin) may be derived directly from cyflumetofen or through AB-6

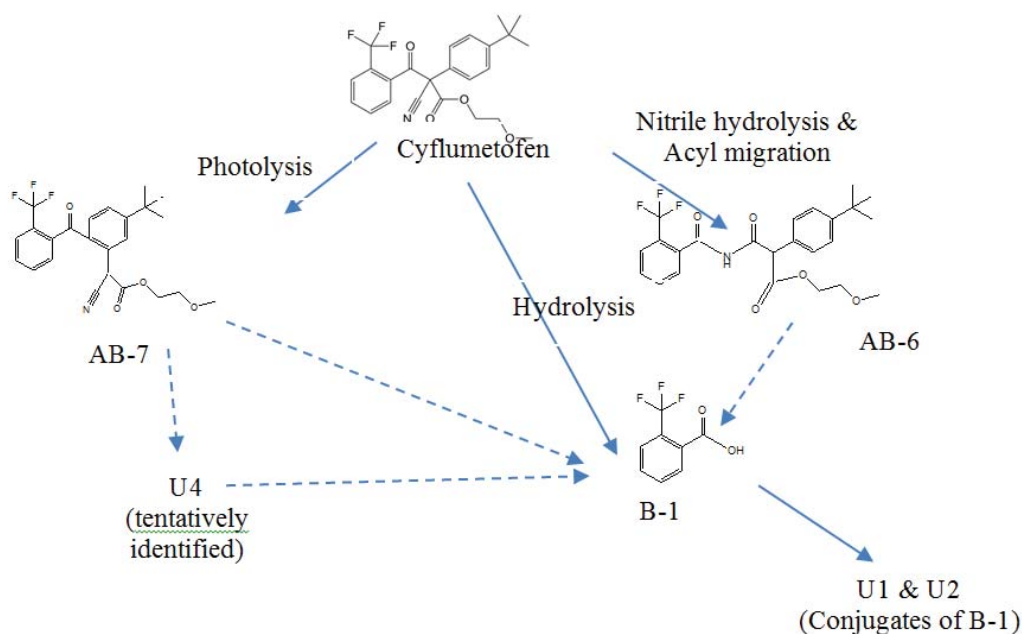


Figure 2 Proposed metabolic pathway of cyflumetofen in plants

### *Environmental fate*

The Meeting received extensive information on fate and behaviour in soil, hydrolysis in aquatic systems, and residues in succeeding crops. Since cyflumetofen is applied as foliar spray on trees (citrus fruits, pome fruits, grapes and tree nuts), strawberry, tomato and eggplant, studies on hydrolysis rate and products (some included in the Physical and Chemical Properties Section), photolysis on plant surface (conditional: No information was available) and rotational crops were relevant to the current review (“Submission and evaluation of pesticide residues data for the estimation of maximum residue levels in food and feed”, FAO, 2009).

### *Aerobic soil metabolism*

Studies on aerobic soil metabolism studies on cyflumetofen and its degradates were provided.

#### *Study 1\_(Knight, 2004a)*

Butylphenyl- or benzoyl-label cyflumetofen was incubated for 181 days in a sandy loam soil in the dark at 25 °C under aerobic conditions at a concentration of 0.93 mg/kg dw. Moisture was maintained throughout the incubation period and volatile radioactivity was trapped (PUF, ethyl digol, KOH). Soil biomass was 3.08% at the start of the experiment and 0.6% after 181 days (study end), indicating sufficiently viable conditions (at least during the first 100 days, as is also indicated by the continuous increase of [<sup>14</sup>C]CO<sub>2</sub>).

Single soil portions of each label treatment were sampled at 0, 1, 3, 6, 10, 14, 30, 59, 90, 121 and 181 days. Samples were extracted with 100 mL acetonitrile (2×), followed by extraction with 100 mL acetonitrile:water (3:1 v/v) (2×) and finally reflux for 6 hours in acetonitrile:water (3:1 v/v). Polyurethane foam plugs were extracted with acetonitrile. Post-extraction solids were fractionated into fulvic and humic acids and humin. Extracts were subjected to chromatographic analysis by HPLC-RAM and TLC. Metabolite identification was obtained by comparison with reference substances of metabolites.

Mass balances were between 89.6 and 102.6% of applied radioactivity (AR) for both labels. Unextracted radioactivity was at maximum 43.4% AR (14 d) and 37.8% AR on day 90. Volatile radioactivity consisted only of CO<sub>2</sub> and was 25.7 and 34.0% AR on day 90 for butylphenyl- and benzoyl-label, respectively (27.9 and 39.3% AR at study end).

Cyflumetofen degraded with a DT<sub>50</sub> of 2.76 days and a DT<sub>90</sub> of 9.16 days by linear regression analysis. Degradation could be described by first-order kinetics (r<sup>2</sup> 0.952). Significant metabolites (exceeding 10% AR or 2× 5% at two consecutive time points) were AB-1 (max 8.3% AR, day 59), B-1 (max 22.9% AR, day 6) and SHL9 (max 7.9%, day 10).

Table 13 Soil properties of Wolston sandy loam

Property	Value
Origin	Wolston, UK
Texture (USDA)	Sandy loam
Sand (% , USDA)	70.8
Silt (% , USDA)	16.8
Clay (% , USDA)	12.4
pH-water	6.5
pH-KCl	6.5
pH-CaCl <sub>2</sub>	5.8
% OM	3.3
% OC	1.9
CEC (meq/100g)	15.3
water content at pF2 (% , w/w)	22.3
MWHC (% , w/w)	52.4

Table 14 Distribution of radioactivity after aerobic incubation at 25 °C of sandy loam soil treated with butylphenyl- or benzoyl-label cyflumetofen at 0.93 mg/kg

Time d	% AR						
	Extracted radioactivity	Unextracted radioactivity				Volatile radioactivity	Mass balance
		fulvic acids	humic acids	humins	total		
Butylphenyl-label							
0	98.2	nd	nd	nd	2.8	na	101.0
1	87.5	2.1	3.7	4.2	10.0	< 0.1	97.5
3	58.5	5.7	11.7	10.4	27.8	6.6	92.9
6	46.3	6.7	13.9	12.0	32.5	14.2	93.0
10	44.4	7.3	13.3	12.8	33.4	11.8	89.6
14	34.9	8.4	16.0	19.1	43.4	16.4	94.7
30	32.9	7.0	14.2	19.6	40.8	19.3	93.0
59	34.5	7.1	19.4	15.8	42.3	24.6	101.4
90	28.3	6.9	17.2	13.7	37.8	25.7	91.8
121	29.0	7.2	15.4	14.4	36.9	31.2	97.1
181	29.9	7.9	15.9	14.1	37.9	27.9	95.4
Benzoyl-label							
0	98.3	nd	nd	nd	4.3	na	102.6
1	94.2	4.6	0.7	1.4	6.7	0.7	101.6
3	79.1	12.2	2.1	3.8	18.0	3.6	100.7
6	67.2	15.0	3.0	6.2	24.2	7.7	99.1
10	55.7	16.4	4.5	7.8	28.7	12.6	97.0
14	44.4	18.5	5.6	10.3	34.3	20.8	99.5
30	36.8	15.0	4.9	11.2	31.2	27.1	95.1
59	34.5	16.3	5.7	12.9	34.9	28.6	98.0
90	29.4	14.6	5.7	10.3	30.6	34.0	94.0
121	30.0	13.9	5.4	9.6	29.0	36.7	95.7
181	27.1	14.8	4.9	11.0	30.7	39.3	97.1

Nd = Not determined

na = Not applicable

Table 15 Identification of radioactivity after aerobic incubation at 25 °C of sandy loam soil treated with butylphenyl- or benzoyl-label at 0.93 mg/kg

Component	% AR										
	0 d	1 d	3 d	6 d	10 d	14 d	30 d	59 d	90 d	121 d	181 d
Butylphenyl-label											
Polar comp.	nd	nd	0.8	1.0	1.2	1.1	0.5	1.1	1.5	1.2	1.4
SHL1	nd	nd	1.2	1.1	0.8	0.8	1.3	0.9	1.0	1.1	1.5
SHL3	nd	0.9	6.2	2.5	2.0	2.1	1.6	0.9	1.4	1.6	1.7
SHL4	nd	1.4	2.6	2.7	2.8	2.5	1.6	1.8	1.9	1.8	2.0
AB-1	nd	2.4	3.3	4.4	6.9	6.4	5.5	8.3	6.5	6.0	3.8
SHL6	0.9	nd	0.9	2.1	1.2	0.7	0.9	0.6	0.5	1.4	1.9
Parent	96.8	76.6	27.1	15.1	9.8	6.2	4.6	3.6	2.2	1.9	1.3
SHL7	nd	nd	1.7	1.1	0.2	0.6	0.7	1.4	1.2	1.7	1.3
SHL8	nd	nd	1.0	2.4	3.7	1.4	1.4	2.9	1.6	1.2	1.7
AB-1 dimer	nd	2.0	4.6	4.4	5.8	4.6	4.8	5.0	3.4	2.8	3.6
AB-1 (cis)	nd	1.8	6.4	5.7	1.2	5.1	2.0	3.2	5.8	2.7	3.2
AB-1 (trans)	nd				5.1		3.2			4.1	
Others	0.5	2.5	2.7	3.7	3.6	3.4	4.7	1.2	1.2	1.5	1.8
Benzoyl-label											
Polar comp.	nd	0.5	1.7	1.9	2.3	3.2	1.8	2.7	1.5	2.2	1.6
B-1	nd	7.6	18.7	22.9	18.3	8.4	3.6	4.3	2.3	3.2	2.7
SHL3	nd	nd	0.3	0.4	0.3	0.3	0.4	nd	nd	nd	nd
SHL4	nd	0.2	1.4	0.9	0.4	0.8	0.5	1.9	0.7	0.8	1.2
AB-1	nd	0.7	4.0	4.8	5.6	3.6	7.8	5.2	7.0	6.5	5.1
SHL6	nd	0.4	0.8	0.9	1.4	2.0	1.3	1.8	1.1	1.1	nd
Parent	98.0	80.6	38.8	17.5	7.1	6.3	5.0	4.4	2.6	2.2	1.8
SHL7	nd	nd	1.1	1.9	0.7	1.3	0.7	0.6	0.4	1.9	1.6
SHL8	nd	nd	1.7	1.8	1.3	1.6	1.1	2.2	1.9	0.9	2.2

Component	% AR										
	0 d	1 d	3 d	6 d	10 d	14 d	30 d	59 d	90 d	121 d	181 d
AB-1 dimer	nd	1.4	4.6	5.0	7.9	7.1	5.8	3.8	4.4	3.5	3.3
AB-1 (cis)	nd	1.7	4.2	6.0	1.0	6.5	1.1	4.3	1.4	1.2	1.2
AB-1 (trans)	nd				7.1		4.6		4.1		
Others	0.3	1.1	1.7	3.1	2.3	3.2	2.9	3.2	2.1	2.4	2.4

nd = Not detected

Table 16 Distribution of radioactivity after sterile incubation at 25 °C of sandy loam soil treated with butylphenyl- or benzoyl-label at 0.93 mg/kg

Time, d	% AR			
	Extracted	Unextracted	Volatiles	Mass balance
Butylphenyl-label				
0	103.5	< 0.1	na	103.5
1	100.7	0.8	< 0.1	101.5
7	93.0	8.4	< 0.1	101.4
30	61.0	42.7	4.1	107.8
Benzoyl-label				
0	100.2	< 0.1	na	100.2
1	99.0	0.7	< 0.1	99.7
7	94.6	5.3	< 0.1	99.9
30	83.6	19.7	< 0.1	103.3

Na = Not applicable

Table 17 Identification of radioactivity after sterile incubation at 25 °C of sandy loam soil treated with butylphenyl or benzoyl-label at 0.93 mg/kg

	% AR							
	Butylphenyl-label				Benzoyl-label			
	0 d	1 d	7 d	30 d	0 d	1 d	7d	30 d
Polar comp.	nd	nd	nd	7.3	nd	nd	0.4	5.3
SHL1	nd	nd	nd	nd	na	na	na	na
B-1	na	na	na	na	nd	1.8	12.0	55.1
SHL3	nd	nd	0.8	7.7	nd	nd	nd	nd
SHL4	nd	0.4	0.7	1.9	nd	nd	nd	nd
AB-1	nd	0.6	2.7	6.5	nd	0.5	2.6	3.0
Parent	95.4	97.0	85.0	30.8	97.7	95.1	77.9	17.0
SHL7	nd	nd	0.9	1.8	nd	nd	nd	1.1
SHL8	nd	nd	nd	0.9	nd	nd	0.3	nd
AB-1 dimer	nd	nd	0.4	0.4	nd	nd	nd	nd
AB-1 (cis/trans)	nd	nd	nd	0.7	nd	nd	nd	nd
Others	4.6	2.0	2.5	3.1	2.3	1.6	1.4	2.2

Na = Not applicable

nd = Not detected

### Study 2 (Willems, 2008a)

Basing on the above study results,  $DT_{50}$  and  $DT_{90}$  for the degradation of cyflumetofen in soil were calculated following FOCUS guidance of EU. Optimization was by the Marquardt method using ordinary least squares weighting.

Cyflumetofen degrades in Wolston sandy loam soil (at 25 °C) with a half-life of 1.81–2.29 days (mean 2.05 d, persistence endpoint). Degradation was described by DFOP kinetics. For modelling a mean  $DegT_{50}$  of 2.38 days was calculated (correction for soil moisture not included).

*Study 3 (Ta and Strobush, 2011a)*

The metabolism of cyflumetofen in four US soils under aerobic conditions was investigated using benzoyl- and butylphenyl-label. Four soils of different textures (loamy sand and loam soils) collected from different locations representing various geographical regions in the USA were used. Cyflumetofen was applied at the rate of 0.8 mg ai/kg soil, equivalent to 200 g ai/ha.

Soil equivalent to 50 g of dry weight was weighed into 250 mL wide-mouth polypropylene centrifuge bottles. The soils were maintained aerobically by connecting to an air flow-through system in the dark at  $20 \pm 2$  °C for 120 days. The soil moistures were adjusted to approximately 40–60% of the maximum water holding capacity (MWHC) prior to application of test material to the soil and maintained throughout the incubation period by adding HPLC water if necessary. Moisturized CO<sub>2</sub>-free air was passed over the soil surface and air leaving the system was passed through 1 N NaOH solution to trap any CO<sub>2</sub> produced. Duplicate samples were analysed at 0, 1, 3, 7, 16, 29, 58, and 120 days after treatment (DAT). The soil samples were extracted once with acetonitrile, followed by 70% acetonitrile in water twice, and trifluoroacetic acid was added to bring the extract solution to a concentration of 0.1% acid to prevent any hydrolysis that may occur. The solvent extracts were pooled, concentrated and analysed by HPLC. Identification of parent and its transformation products was performed by retention matching with reference standards by HPLC and confirmation by LC/MS and/or NMR. The amounts of CO<sub>2</sub> and organic volatiles produced were determined at each sampling time (excluding 0 DAT). The traps were assayed directly by adding aliquots of the trapping solutions into liquid scintillation fluid and counting by liquid scintillation counting.

Table 18 Soil properties

Geographic Location	CA, USA	IN, USA	NJ, USA	WI, USA
Soil series	Corralitos	Crosby	Penn	Burkhardt
Classification	Mixed, thermic, Typic Xeropsamments	Fine, mixed, active, mesic Aeric Epiaqualfs	Fine-loamy, mixed, superactive mesic Ultic Hapludalfs	Sandy, mixed, mesic Typic Hapludolls
Texture Class	Loamy Sand	Loam	Loam	Loamy Sand
Sand (%)	87	35	29	87
Silt (%)	8	46	44	6
Clay (%)	5	19	27	7
pH (in water)	8.1	6.5	6.9	6.3
Organic Matter (%)	0.49	2.30	2.30	2.70
Organic Carbon % (1)	0.28	1.33	1.33	1.57
Soil Biomass (µgC/ g soil)				
0 DAT	43.2	160.9	139.8	128.5
120 DAF	55.3	127.9	54.2	86.8
Cation Exchange Capacity (CEC, meq/100 g)	6.6	9.0	8.1	8.4
Max Moisture Holding Capacity (%)	18.8	36.6	43.3	28.0
Bulk Density g/cc (disturbed)	1.42	1.04	1.11	1.34

The average total recovery of the radiolabelled material was 95.5% of the total applied radioactivity (TAR) for all samples from four soils with two labels.

*California soil:*

The total radioactivity recovered from the benzoyl-label ranged from 97.1 to 103.3% of the total applied radioactivity (TAR) with an average of 99.4% TAR. Radioactive residues extracted by solvent decreased from approximately 99.7–100% TAR at 0 day after treatment (DAT) to approximately 67.6–77.4% TAR at 120 DAT. Unextracted radioactive residues increased from approximately 0.3% TAR at 0 DAT to 16–21.6% TAR at 120 DAT. The concentration of the parent compound decreased from approximately 98.1–98.2% TAR at 0 DAT to approximately 3–5% TAR at 120 DAT. There

were three major (> 10% TAR) transformation products identified as B1, B3, and AB1 + AB1 dimer 2. At 120 DAT, the total evolved  $^{14}\text{CO}_2$  was 9.9% TAR.

*Indiana soil:*

The total radioactivity recovered from the benzoyl-label ranged from 87 to 100.6% TAR with an average of 94.3% TAR. Radioactive residues extracted by solvent decreased from approximately 99–100.2% TAR at 0 DAT to approximately 31.5–32.6% TAR at 120 DAT. Unextracted radioactive residues increased from approximately 0.4% TAR at 0 DAT to 37.8–38.0% TAR at 120 DAT. The concentration of the parent compound decreased from approximately 97.6–98.9% TAR at 0 DAT to approximately 1.2–1.9% TAR at 120 DAT. There were 3 major (> 10% TAR) transformation products identified as B1, AB1 + AB1 dimer 1, and AB1 + AB1 dimer 2. At 120 DAT, the total evolved  $^{14}\text{CO}_2$  was 17.6% TAR.

*New Jersey soil:*

The total radioactivity recovered from the benzoyl-label ranged from 90.2 to 101.5% TAR with an average of 95.4% TAR. Radioactive residues extracted by solvent decreased from approximately 100.5–101% TAR at 0 DAT to approximately 37.6–38.7% TAR at 120 DAT. Unextracted radioactive residues increased from approximately 0.5% TAR at 0 DAT to 31.1–32.7% TAR at 120 DAT. The concentration of the parent compound decreased from approximately 98.8–99.3% TAR at 0 DAT to approximately 1.3–2.7% TAR at 120 DAT. There were 3 major (> 10% TAR) transformation products identified as B1, AB1 + AB1 dimer 1, and AB1 + AB1 dimer 2. At 120 DAT, the total evolved  $^{14}\text{CO}_2$  was 24.1% TAR.

The total radioactivity recovered from the butylphenyl-label ranged from 90.3 to 100% TAR with an average of 93.7% TAR. Radioactive residues extracted by solvent decreased from approximately 97.6–99.5% TAR at 0 DAT to approximately 33.8–36.2% TAR at 120 DAT. Unextracted radioactive residues increased from approximately 0.5% TAR at 0 DAT to 41.2–43.1% TAR at 120 DAT. The concentration of the parent compound decreased from approximately 94.5–96.6% TAR at 0 DAT to approximately 2.5–2.6% TAR at 120 DAT. There was 1 major (> 10% TAR) transformation product identified as AB1. At 120 DAT, the total evolved  $^{14}\text{CO}_2$  was 17.3% TAR.

*Wisconsin soil:*

The total radioactivity recovered from the benzoyl-label ranged from 88.3 to 100.1% TAR with an average of 95.2% TAR. Radioactive residues extracted by solvent decreased from approximately 98.1–99.9% TAR at 0 DAT to approximately 39.8% TAR at 120 DAT. Unextracted radioactive residues increased from approximately 0.3% TAR at 0 DAT to 31.5–31.6% TAR at 120 DAT. The concentration of the parent compound decreased from approximately 96.8–98.4% TAR at 0 DAT to approximately 2.8–3.2% TAR at 120 DAT. There were four major (> 10% TAR) transformation products identified as B1, AB1, AB1 + AB1 dimer 1, and AB1 + AB1 dimer 2. At 120 DAT, the total evolved  $^{14}\text{CO}_2$  was 17.3% TAR.

Table 19 Summary of aerobic soil metabolism study 2 results

Soil	Mass balance (Total recovery)	Degradation of cyflumetofen in 120 days	Half-life/ Dissipation time
CA soil	97.1 to 103.3% TAR	93.2–95.1%	3.51 days <sup>a</sup>
IN soil	87.0 to 100.6% TAR	96.4–97.0%	2.28 days <sup>b</sup>
NJ soil	90.2 to 101.5% TAR	92.0–97.5	2.36 days <sup>b</sup>
WI soil	88.3 to 100.1% TAR	93.6–95.6%	3.41 days <sup>b</sup>

<sup>a</sup> First order multi compartment

<sup>b</sup> Single first order

*Study 4 (Brands, 2011a)*

Benzoyl-label was incubated for 120 days in the dark in 100 g dw equivalent each of three soils (Speyer 2.2, loamy sand; Speyer 2.3, sandy loam; and Speyer 6S, clay), which were equilibrated for 20–21 days in the dark, at  $20 \pm 2$  °C under aerobic conditions at a concentration of 0.383 mg/kg (~0.8% acetonitrile) on a dry weight basis. A humidified air stream was passed through the soil containers (2 x 30 minutes/day). The outgoing air was passed through a polyurethane foam plug, a trap containing 2-ethoxyethanol and two traps containing 2M NaOH to trap volatile substances and CO<sub>2</sub>. Soil moisture was kept constant at 45% of the MWHC throughout the incubation period. Soil biomass was determined at the start and end of the incubation period by the fumigation/extraction method.

Speyer 2.2, 2.3 and 6S soils, after sieving (2 mm), were used within two months (storage at 4 °C) of field sampling. The properties of these soils are shown in the following Table.

Table 20 Properties of the three soils used in the study.

Property	Speyer 2.2	Speyer 2.3	Speyer 6S
Origin	Hanhofen, Rheinland-Pfalz, Germany	Offenbach, Rhein-land-Pfalz, Germany	Sieboldingen, Rhein-land-Pfalz, Germany
Texture (USDA)	Loamy sand	Sandy loam	Clay
Sand (% , USDA)	79.1	61.0	21.9
Silt (% , USDA)	13.5	29.8	36.0
Clay (% , USDA)	7.9	9.2	42.0
pH-water	-	-	-
pH-KCl	-	-	-
pH-CaCl <sub>2</sub>	5.6	6.2	7.0
% OM	4.07	1.76	3.26
% OC	2.36	1.02	1.89
CEC (meq/100g)	11	9	20
MWHC (% , w/w)	48.0	34.9	42.3

Single soil portions and associated traps were sampled at 0, 1, 3, 6 (duplicate sample), 10, 14, 21, 35 (duplicate sample), 58, 90 and 120 days. Soil samples were extracted three times with 100 mL acetonitrile/sodium acetate buffer pH 4 (75:25, v/v). Samples of day 35 were also extracted by n-hexane and acetonitrile-Soxhlet extraction (3 hours). Polyurethane foam plugs were extracted with acetonitrile (2×). Radioactivity in extracts and liquid traps was determined by LSC and in post-extraction solids by combustion/LSC. Soil extracts were fractionated in an organic and aqueous phase. Relevant fractions were subjected to analyses by HPLC-RAM and TLC. Metabolites were identified by comparison with reference substances of metabolites and LC-MS/MS analysis. Radioactivity in the NaOH traps was confirmed as CO<sub>2</sub> by precipitation with Ba(OH)<sub>2</sub>.

Soil biomass was 1.1–2.1% of OC at the start and 0.3–2.7% at the end of the experiment. In combination with significant mineralization at the end of the study, sufficiently viable conditions were indicated.

Mass balances were between 87.9 and 109.8% of applied radioactivity (AR). No specific trend was observed and all results were considered acceptable.

The amount of extractable residues decreased from 103.5–104.6% AR to 54.1 (Speyer 2.2), 26.9 (Speyer 2.3) and 62.2% AR (Speyer 6S) at study end (day 120). Unextractables were maximum 32.1 (Speyer 2.2, day 58), 40.1 (Speyer 2.3, day 120) and 38.4% AR (Speyer 6S, day 35). Additional extraction with n-hexane (day 35 sample) released 1.6–3.2% AR radioactivity and with Soxhlet 1.8–5.1% AR.

Volatile radioactivity consisted mostly of CO<sub>2</sub> and was 9.5 (Speyer 2.2), 20.7 (Speyer 2.3) and 1.7% AR (Speyer 6S) at day 120 (study end) (0.8, 16.2 and 1.8% AR, respectively at day 90).

Unextracted radioactivity was at a maximum of 32.1 (Speyer 2.2, day 58), 40.1 (Speyer 2.3, day 120) and 38.4% AR (Speyer 6S, day 35).

Cyflumetofen degraded to < 10% AR after 21 days in Speyer 2.1 soil, to < 10% AR after 10 days in Speyer 6S soil and to ~10% AR at day 120 in Speyer 2.3 soil under aerobic conditions in the dark at 20 °C.

Those metabolites exceeding 10% AR or 2× 5% at two consecutive time points were B-1 (max 63.0% AR, day 90, Speyer 2.2) and B-3 (max 23.0% AR, day 21, Speyer 2.2). No other metabolites exceeding 5% AR were observed in any soil.

Table 21 Distribution of radioactivity after aerobic incubation at 20 °C of Speyer 2.2, 2.3 and 6S soils treated with benzoyl-label at 0.383 mg/kg

Time d	% AR				
	Organic volatiles	CO <sub>2</sub>	Extracted radioactivity	Unextracted radioactivity	Mass balance
Speyer 2.2 Soil					
0	na	Na	103.7	2.6	106.3
1	0.0	0.0	90.2	5.4	95.6
3	0.0	0.1	94.1	9.1	103.3
6	0.0	0.1	82.5	14.1	96.7
6	0.0	0.0	69.1	18.7	87.9
10	0.0	1.3	79.2	20.4	101.0
14	0.1	1.4	76.6	16.1	94.2
21	0.0	1.2	76.8	18.1	96.2
35	0.3	5.1	67.3	21.2	93.9
35	0.1	0.7	72.6	22.5	95.9
58	0.4	1.1	72.5	32.1	106.1
90	0.2	0.8	77.2	18.3	96.6
120	1.4	9.5	54.1	30.1	95.0
Speyer 2.3 Soil					
0	na	na	104.6	0.9	105.5
1	0.0	0.1	81.7	7.4	89.1
3	0.1	0.5	78.4	19.3	98.4
6	0.1	0.8	66.7	22.4	90.0
6	0.1	0.3	75.7	24.2	100.4
10	0.1	2.8	68.6	27.8	99.2
14	0.1	3.2	71.4	19.4	94.1
21	0.1	3.5	71.2	20.8	95.8
35	0.2	4.4	71.1	24.8	100.4
35	0.1	7.0	61.0	25.1	93.2
58	0.2	11.4	41.5	38.3	91.4
90	0.1	16.2	41.4	33.3	91.0
120	0.1	20.7	26.9	40.1	87.8
Speyer 6S soil					
0	na	na	103.5	6.3	109.8
1	0.0	0.0	80.6	21.4	102.0
3	0.0	0.0	75.6	26.0	101.6
6	0.1	0.4	67.2	33.4	101.1
6	0.0	0.3	70.5	30.7	101.5
10	0.2	1.0	61.7	34.6	97.5
14	0.1	0.7	57.0	32.1	89.9
21	0.1	0.4	64.9	29.1	94.4
35	0.1	0.0	55.1	37.4	92.6
35	0.1	0.0	59.6	38.4	98.1
58	0.2	1.2	66.3	37.1	104.8
90	0.4	1.8	60.6	32.1	94.9
120	0.4	1.7	62.2	33.0	97.4

na: not applicable



Table 22 Identification of radioactivity after aerobic incubation at 20 °C of Speyer 2.2, Speyer 2.3 and Speyer 6S soils treated with Benzoyl-label at 0.383 mg/kg

Time d	% AR			
	Cyflumetofen	B-1 <sup>b</sup>	B-3 <sup>b</sup>	Others <sup>a</sup>
Speyer 2.2 Soil				
0	97.5	3.0	0.0	0.0
1	94.1	3.8	0.0	0.0
3	75.7	8.9	1.8	4.0
6	31.6	35.7	4.9	5.9
6	29.6	34.3	14.2	0.0
10	22.4	35.4	16.1	3.5
14	17.4	47.1	12.4	0.0
21	6.7	55.2	23.0	0.0
35	14.7	35.3	5.6	12.9
35	7.1	48.7	13.8	6.5
58	14.7	39.9	9.9	1.6
90	2.6	63.0	8.3	6.2
120	10.2	33.8	5.2	7.1
Speyer 2.3				
0	96.2	7.9	0.0	0.0
1	80.3	13.5	0.0	0.0
3	54.4	22.3	0.0	0.0
6	23.4	34.3	12.7	2.1
6	26.4	37.2	3.6	6.9
10	22.3	36.6	4.7	5.6
14	20.9	36.1	2.1	8.2
21	14.7	42.8	3.2	8.9
35	15.2	43.2	3.5	9.0
35	14.6	36.3	1.9	8.4
58	12.5	12.6	1.0	15.4
90	11.2	16.4	0.0	15.0
120	10.6	2.6	0.0	15.7
Speyer 6S				
0	95.1	4.3	0.0	0.0
1	70.1	13.3	0.0	0.0
3	40.8	23.2	0.0	9.4
6	14.9	32.4	2.0	8.4
6	21.8	36.4	3.2	14.1
10	6.4	42.3	2.3	7.8
14	4.8	46.2	3.1	5.9
21	5.9	39.4	4.8	7.9
35	3.4	42.3	1.1	7.3
35	3.0	45.5	2.5	5.9
58	5.0	52.8	1.3	7.5
90	4.8	42.4	0.0	11.5
120	5.4	43.0	0.0	11.6

<sup>a</sup> individual fractions <5% AR<sup>b</sup> confirmed by co-chromatography with reference standard (HPLC, TLC (only B-1), LC-MS/MS)

DT<sub>50</sub> and DT<sub>90</sub> values for calculated by fitting single first order (SFO), first order multi-compartment (FOMC) and double first order in parallel (DFOP) models to the data and the best fit was obtained by DFOP model (based on visual examination, r<sup>2</sup> and  $\chi^2$ ). Calculated DT<sub>50</sub> and DT<sub>90</sub> values are shown in the following Table.

Table 23 Half-lives for degradation of cyflumetofen at 20 °C under aerobic conditions (double first order in parallel was assumed)

Soil	M <sub>0</sub> %	g	k <sub>1</sub> days <sup>-1</sup>	k <sub>2</sub> days <sup>-1</sup>	DT <sub>50</sub> days	DT <sub>90</sub> days	r <sup>2</sup>	$\chi^2$ (err)
Speyer 2.2	105 ± 6.0	0.91 ± 0.03	0.197 ± 0.030	0	4.0	21	0.960	15.3
Speyer 2.3	98.8 ± 3.1	0.83 ± 0.03	0.303 ± 0.035	0.004 ±	3.0	130	0.988	7.3

Soil	M <sub>0</sub> %	g	k <sub>1</sub> days <sup>-1</sup>	k <sub>2</sub> days <sup>-1</sup>	DT <sub>50</sub> days	DT <sub>90</sub> days	r <sup>2</sup>	χ <sup>2</sup> (err)
				0.003				
Speyer 6S	95.0 ± 1.7	0.95 ± 0.01	0.312 ± 0.014	0	2.4	9.2	0.996	3.5

*Study 5 (Willems, 2008a)*

The degradation of cyflumetofen in three soils (Speyer 2.2, 2.3 and 6S) was studied by Brands in 2008 (NOTOX report 478338; 7.2.1/01), in which benzoyl-label was spiked to these soils and incubated for 120 days at 20 °C in the dark. Using the degradation results, DT<sub>50</sub> and DT<sub>90</sub> were calculated for cyflumetofen in accordance with FOCUS guidance ((SANCO/10058/2005, vs 2.0, June 2006). Optimization was by the Marquardt method using ordinary least squares weighting.

Table 24 Cyflumetofen degradation in Speyer 2.2, 2.3 and 6S soils

Time d	Cyflumetofen (% of applied)		
	Speyer 2.2	Speyer 2.3	Speyer 6S
0	100.1 <sup>a</sup>	97.1 <sup>b</sup>	101.4 <sup>c</sup>
1	94.1	80.3	70.1
3	75.7	54.4	40.8
6	31.6	23.4	14.9
6	29.6	26.4	21.8
10	22.4	22.3	6.4
14	17.4	20.9	4.8
21	6.7	14.7	5.9
35	14.7	15.2	3.4
35	7.1	14.6	3.0
58	14.7	12.5	5.0
90	2.6	11.2	4.8
120	10.2	10.6	5.4

<sup>a</sup> including 2.6% unextracted residues

<sup>b</sup> including 0.9% unextracted residues

<sup>c</sup> including 6.3% unextracted residues

According to FOCUS guidance, first, single first order (SFO) and first order multi-compartment (FOMC) kinetics were run. Time zero results for cyflumetofen were adjusted for unextracted residues but not for metabolites because the radioactivity present as metabolites was similar to the amount of impurities in the spike solutions. The possibility for elimination of data points or constraining of M<sub>0</sub> was investigated, and when FOMC still resulted in a better fit, also dual first order in parallel (DFOP) kinetics were fitted to the data. DegT<sub>50</sub> and DegT<sub>90</sub> values for persistence were taken from the best fit model. For modelling, the SFO fit was first evaluated and when not acceptable, FOMC or DFOP were also assessed. In case FOMC or DFOP was suited for modelling, the DegT<sub>50</sub> for modelling was calculated as DegT<sub>90</sub>/3.32.

Persistence and modelling endpoints for the degradation of cyflumetofen in soil were calculated and are shown in the following Table.

Table 25 Persistence and modelling endpoints for degradation of OK-5101 in Speyer soils

Soil	Persistence			Modelling
	DT <sub>50</sub> at 20 °C days	DT <sub>90</sub> at 20 °C days	model	DT <sub>50</sub> at 20 °C <sup>a</sup> Days
Speyer 2.2	4.33	23.10	DFOP	6.70
Speyer 2.3	3.13	134.10	DFOP	13.30
Speyer 6S	2.20	8.40	DFOP	2.95

<sup>a</sup> not including moisture correction.

*Study 6 (Brands & Corral, 2011b)*

The degradation of B-1 in soil was investigated in three soils (the same soils as in Study 4). A 50 g (dw) portion each of three soils (Speyer 2.2, loamy sand; Speyer 2.3, sandy loam; and Speyer 6S, clay), which were sieved (2 mm), stored at 4 °C and used within 4 weeks of field sampling, was equilibrated for 4 days in the dark at a moisture content of ~40% of the MWHC at 20 °C. Following equilibration, soil portions were treated with B-1 in acetonitrile/water 1/9 (v/v) at a rate of 0.25 mg/kg dw (~0.16% acetonitrile). The test substance was gently mixed in the soil. Incubation took place in the dark at 20±2 °C under aerobic conditions for 120 days. Soil moisture was kept constant at ~40% of the MWHC throughout the incubation period.

Soil biomass was determined at the start and end of the incubation period by the fumigation/extraction method. Single soil portions were sampled at 0, 1, 4 (duplicate sample), 10, 28 (duplicate sample), 70 and 120 days. Soil samples were extracted with 250 mL acetonitrile/water (10/90 v/v) for 30 minutes. The samples were filtered over a 0.2 µm filter prior to analysis by UPLC-MS/MS.

Biomass was > 1% of OC throughout the study. Actual moisture contents (dw basis) throughout the study were 20% (Speyer 2.2), 14% (Speyer 2.3) and 16% (Speyer 6S), equivalent to ~39% of the MWHC.

Table 26 Degradation of B-1 during aerobic incubation at 20 °C in soil treated with B-1 at 0.25 mg/kg

Time d	B-1 (% of applied)		
	Speyer 2.2	Speyer 2.3	Speyer 6S
0	102	105	100
1	93	101	95
4	72	91	95
4	68	93	93
10	34	70	94
28	0.5*	28	56
28	0.5*	33	63
70	na	16	16
120	na	1.3	20

na: not applicable

\* set at ½ LOQ

DT<sub>50</sub> and DT<sub>90</sub> values for B-1 were obtained by fitting single first order (SFO) kinetics to the data. The results are given in the following Table. The FOMC model was also verified and gave similar results as the SFO model, but optimization errors were much larger compared to the SFO model. Because there was also no biphasic pattern visible, the SFO model was selected as best fit model.

Table 27 Half-lives for degradation of B-1 at 20 °C under aerobic conditions based on SFO kinetics <sup>a</sup>

Soil	M <sub>0</sub> %	K day <sup>-1</sup>	DT <sub>50</sub> days	DT <sub>90</sub> Days	r <sup>2</sup>	χ <sup>2</sup> (err)
Speyer 2.2	104 ± 2.8	0.110 ± 0.008	6.3	21.0	0.993	4.6
Speyer 2.3	106 ± 2.7	0.041 ± 0.003	16.7	55.5	0.988	5.3
Speyer 6S	102 ± 3.7	0.019 ± 0.002	36.3	121	0.959	7.2

<sup>a</sup> not corrected for moisture content.

B-1 degraded in three soils under aerobic conditions in the dark at 20 °C with half-lives of 6.3, 16.7 and 36.3 days and DT<sub>90</sub> values of 21.0, 55.5 and 121 days. These values are not normalized for moisture. Degradation was described by SFO kinetics.

*Study 7 (Corral & Brands, 2010c)*

AB-1 was studied by incubating for 120 days in three soils (Speyer 2.1, sand; Speyer 2.2, loamy sand; and Speyer 6S, clay) in the dark at 20 °C under aerobic conditions at a concentration of 0.04 mg/kg

dw. Portions of 50 g dw equivalent soil were equilibrated for 11 days in the dark at a moisture content of ~40% of the MWHC at 20 °C. Following equilibration, soil portions were treated with AB-1 in acetonitrile at a rate of 0.04 mg/kg dw. The test substance was gently mixed in the soil. Incubation took place in the dark at 20±2 °C under aerobic conditions for 120 days. Soil moisture was kept constant at ~40% of the MWHC throughout the incubation period. Soil biomass was determined at the start and end of the incubation period by the fumigation/extraction method.

Single soil portions were sampled at 0, 0.25 (duplicate sample), 1, 2 (duplicate sample), 4, 29, 72 and 120 days. Soil samples were extracted with 400 mL acetonitrile/water (50/50 v/v) containing 0.05 or 0.1% (v/v) ammonium solution for 30 minutes. The samples were filtered over a 0.2 µm filter prior to analysis by UPLC-MS/MS (fully validated, see OTSA-0368).

Biomass was >1% of OC throughout the study. Actual moisture contents (dw basis) throughout the study were 13% (Speyer 2.1), 20% (Speyer 2.2) and 16% (Speyer 6S), equivalent to 39-40% of the MWHC

Table 28 Soil properties of Speyer 2.1, 2.2 and 6S soils

Property	Speyer 2.1	Speyer 2.2	Speyer 6S
Origin	Dudenhofen, Rheinland-Pfalz, Germany	Hanhofen, Rheinland-Pfalz, Germany	Siebeldingen, Rheinland-Pfalz, Germany
Texture (USDA)	Sand	Loamy sand	Clay
Sand (% , USDA)	88.1	81.2	21.7
Silt (% , USDA)	8.5	12.1	35.7
Clay (% , USDA)	3.4	6.7	42.6
pH-water	-	-	-
pH-KCl	-	-	-
pH-CaCl <sub>2</sub>	5.1	5.4	7.2
% OM	1.52	4.0	3.1
% OC	0.88	2.29	1.79
CEC (meq/100g)	4	10	20
water content at pF2 (% , w/w)	-	-	-
MWHC (% , w/w)	33.3	49.7	40.1

Material balance was between 85 and 123% for these soils. The degradation of AB-1 over time in these soils is shown in the following table.

Table 29 Degradation of AB-1 during aerobic incubation at 20 °C in soil treated with AB-1 at 0.04 mg/kg

Time d	AB-1 (% of applied)		
	Speyer 2.1	Speyer 2.2	Speyer 6S
0	95	88	90
0.25	18	18	22
0.25	18	19	21
1	14	13	11
2	15	14	9
2	16	14	10
4	12	12	10
29	12	11	8
72	11	9	6
120	8	6	3

DT<sub>50</sub> and DT<sub>90</sub> values for AB-1 were obtained by fitting single first order (SFO), first-order multi-compartment (FOMC), dual first order in parallel (DFOP) and hockey-stick (HS) kinetics to the data. As there was a clear bi-phasic pattern observed, the HS and DFOP models described the results the best (and equally good). According to FOCUS guidance, the HS model is not used for obtaining endpoints for persistence, therefore the DFOP model was selected for use in exposure assessment. The calculated results are shown in the following table.

AB-1 degraded in three soils under aerobic conditions in the dark at 20 °C with half-lives of 0.07, 0.08 and 0.11 days (not corrected for moisture). Degradation was described by DFOP kinetics.

Table 30 Half-lives for degradation of AB-1 at 20 °C under aerobic conditions based on DFOP kinetics <sup>a</sup>

Soil	M <sub>0</sub> %	k <sub>1</sub> , k <sub>2</sub> and g	DT <sub>50</sub> days	DT <sub>90</sub> Days	r <sup>2</sup>	χ <sup>2</sup> (err)
Speyer 2.1	95.0 ± 1.2	k <sub>1</sub> = 12.40 ± 1.13 k <sub>2</sub> = 0.005 ± 0.001 g = 0.847 ± 0.007	0.07	90.8	0.998	3.9
Speyer 2.2	88.0 ± 0.7	k <sub>1</sub> = 10.75 ± 0.50 k <sub>2</sub> = 0.006 ± 0.001 g = 0.846 ± 0.004	0.08	69.3	0.999	2.5
Speyer 6S	90.0 ± 0.7	k <sub>1</sub> = 7.81 ± 0.22 k <sub>2</sub> = 0.009 ± 0.001 g = 0.885 ± 0.004	0.11	15.7	0.999	2.7

<sup>a</sup> not corrected for moisture

#### Photodegradation on soil surface (van Noorloos, 2007a and 2007b)

Photodegradation of cyflymetofen on soil surface was investigated using butylphenyl- and benzoyl-label cyflumetofen.

Butylphenyl- or benzoyl-label was applied uniformly over the surface of thin soil layers at a rate of 401 g ai/ha or 391–395 g ai/ha, respectively. The soil layers were irradiated (300–400 nm: 66.5 Watt/m<sup>2</sup>) for 15 days (equivalent to 33–38 days natural sunlight at 40 °N, summer) at 18–23 °C or for 14 days (equivalent to 35–36 days natural sunlight at 40 °N, summer) at 19–22 °C, respectively, with artificial light (Xenon lamp) simulating the sunlight spectrum. Control samples were incubated in the dark at approximately 20 °C. Duplicate samples were taken after 0, 1, 4, 7, 11 and 15 days of irradiation/incubation. Soil layers were extracted with (75/25 (v/v)) acetonitrile/sodium acetate buffer pH 4. Following fractionation into an organic and aqueous phase the organic phase was analysed by reversed phase HPLC-RAM and selected samples also by TLC. The aqueous phase was not further analysed due to low levels of radioactivity. Polyurethane foam plugs were extracted by acetonitrile and soda lime granules were dissolved in 2 M HCl and released CO<sub>2</sub> was trapped in 2 M NaOH. Radioactivity in liquid fractions was determined by LSC and in residual soil by combustion/LSC. Identification of radioactive fractions was obtained by co-chromatography with reference standards (HPLC and TLC).

Table 31 Soil properties of Speyer 3A loam

Property	Value
Origin	Altußheim, Baden-Württemberg, Germany
Texture	Loam
Sand (%)	46.7
Silt (%)	35.5
Clay (%)	17.7
pH-water	-
pH-KCl	-
pH-CaCl <sub>2</sub>	7.2
% OM	3.7
% OC	2.17
CEC (meq/100g)	19
water content at pF2 (% w/w)	-
MWHC (% w/w)	48.4

#### Mass balances

Butylphenyl-label: Mass balances were between 81.4 and 105.4% of applied radioactivity (AR) for the irradiated samples and between 87.2 and 102.0% AR for the dark control samples. Unextracted

radioactivity was 44.6–45.6% AR (irradiated, study end) and 45.0% AR (dark control, study end, mean). Volatile radioactivity was at the maximum 2.5% AR in any sample.

Benzoyl-label: Mass balances were between 96.4 and 104.7% of applied radioactivity (AR) for the irradiated samples and dark control samples. Unextractables were approximately 24% AR (irradiated, study end) and approximately 39% AR (dark control, study end). Volatile radioactivity was insignificant ( $\leq 0.5\%$  AR).

#### *Cyflumetofen dissipation*

Butylphenyl-label: Cyflumetofen dissipated with a single first order  $DT_{50}$  of 1 and 0.95 days in irradiated soil and dark controls, respectively. No metabolites exceeded 10% AR. Specific photolytic metabolites were not observed. All metabolite fractions observed in irradiated soil were also present in similar amounts in dark controls.

Benzoyl-label: Cyflumetofen dissipated with a single first order  $DT_{50}$  of 1.4 days in irradiated soil and dark controls, respectively. One significant metabolite, B-1, exceeded 10% AR in both the irradiated and dark soil. Specific significant ( $> 10\%$  AR) photolytic metabolites were not observed.

Table 32 Distribution of radioactivity after irradiation of soil layers treated with butylphenyl- or benzoyl-label

Time d	Time sunlight equivalent <sup>b</sup>	% AR					
		PUF	CO <sub>2</sub>	residue in soda lime	extracted <sup>a</sup>	unextracted	mass balance
Butylphenyl-label							
0.0	0.0	Na	na	Na	92.2	6.5	98.6
	0.0	Na	Nas	Na	98.8	6.6	105.4
0.7	1.6	0.0	0.0	0.0	77.9	22.0	99.9
	1.8	0.0	0.0	0.0	79.7	20.1	99.8
3.8	8.2	0.0	1.6	0.1	46.0	37.7	85.4
	9.5	0.0	0.1	0.0	50.5	41.6	92.2
6.8	15.4	0.3	0.0	0.4	53.5	33.0	87.2
	17.7	0.0	1.3	0.7	34.4	44.9	81.4
10.8	24.5	0.0	0.7	0.7	39.2	44.0	84.7
	28.3	0.0	0.6	0.8	45.2	47.7	94.4
14.7	32.6	0.1	1.8	1.7	37.4	45.6	86.7
	37.6	0.0	2.5	1.6	41.9	44.6	90.6
Benzoyl-label							
0.0	0.0	na	Na	Na	97.1	6.6	103.7
	0.0	na	Na	Na	98.5	5.3	103.8
0.9	2.2	0.0	0.0	0.0	82.3	19.0	101.4
	2.3	0.0	0.0	0.0	79.6	25.0	104.7
2.8	7.2	0.0	0.0	0.0	79.4	21.0	100.5
	7.4	0.0	0.1	0.0	78.1	19.7	97.9
5.8	14.8	0.0	0.4	0.1	74.0	25.5	100.1
	14.6	0.0	0.3	0.1	76.0	23.1	99.4
9.8	25.5	0.0	0.4	0.1	72.6	25.1	98.2
	24.1	0.0	0.5	0.3	75.0	23.3	99.1
13.8	35.0	0.0	0.3	0.0	73.5	22.8	96.6
	35.9	0.0	0.3	0.2	71.8	24.5	96.8

Na = Not applicable

<sup>a</sup> 0.9 to 4.2% AR remained in aqueous phase after partitioning (butylphenyl-label) and 0.8 to 21.6% AR remained (benzoyl-label) increasing trend with time.

<sup>b</sup> summer, 40 °N

Table 33 Distribution of radioactivity in dark controls treated with butylphenyl- and benzoyl-label

Time d	% AR					
	PUF	CO <sub>2</sub>	residue in soda lime	extracted <sup>a</sup>	Unextracted	mass balance
Butylphenyl-label						
0.0	na	Na	na	95.5	6.5	102.0
0.7	0.0	0.0	0.0	73.4	27.7	101.1
3.8	0.1	0.0	0.0	49.8	42.2	92.1
6.8	0.0	0.2	0.0	47.5	44.1	91.8
10.8	0.0	0.6	0.0	47.5	39.1	87.2
14.7	0.0	0.5	0.3	46.2	45.0	92.1
Benzoyl-label						
0.0	na	na	na	97.8	5.9	103.8
0.9	0.0	0.0	0.0	74.8	21.6	96.4
2.8	0.0	0.0	0.0	61.3	37.2	98.6
5.8	0.0	0.0	0.0	56.4	40.0	96.4
9.8	0.0	0.0	0.0	59.8	38.2	98.0
13.8	0.0	0.0	0.0	58.7	38.7	97.5

na = Not applicable

<sup>a</sup> 0.9 to 3.8% AR remained in aqueous phase after partitioning (butylphenyl-label) and 0.8 to 19.4% AR remained (benzoyl-label) increasing trend with time.

Table 34a Identification of radioactivity in irradiated soil and dark controls treated with butylphenyl-label

Time d	Time natural sunlight d	% AR						Sum of others <sup>a</sup>
		Parent 14.5–14.8 min	Met 2 6.8-6.9 min	Met 9 <sup>b</sup> 17.1–17.4 min	Met 10 19.1–19.5 min	Met 11 <sup>c</sup> 20.8–21.1 min	Met 12 <sup>d</sup> 21.4–21.7 min	
Butylphenyl-label								
0.0	0.0	47.9	0.3	0.5	0.1	0.0	0.0	1.8
	0.0	79.7	0.4	0.6	0.2	0.0	0.0	2.6
0.7	1.6	53.0	4.7	1.6	1.5	1.2	1.0	6.7
	1.8	59.4	1.7	3.6	2.2	1.3	1.4	3.9
3.8	8.2	16.3	2.9	2.3	2.6	2.9	2.6	6.7
	9.5	12.1	17.7	2.7	2.6	2.8	2.3	3.5
6.8	15.4	11.5	9.5	9.6	7.8	2.5	2.1	3.3
	17.7	4.5	2.1	2.5	1.3	1.4	1.1	14.8
10.8	24.5	4.8	5.1	3.6	3.9	3.2	2.8	10.4
	28.3	6.5	8.8	6.4	4.8	2.5	2.2	3.6
14.7	32.6	3.2	3.2	2.9	4.4	4.2	4.8	8.5
	37.6	4.4	8.9	2.4	3.6	4.6	3.9	6.6
0.7	Dark	48.4	1.4	2.9	2.1	1.0	0.9	3.0
	Dark	60.0	1.7	3.5	2.6	1.5	1.2	1.1
3.8	Dark	13.7	4.5	3.9	4.9	4.3	3.7	8.5
	Dark	14.6	4.4	4.5	5.4	4.8	3.9	8.7
6.8	Dark	7.7	4.5	3.6	4.5	4.6	3.5	8.8
	Dark	8.5	4.6	3.9	5.0	5.2	3.9	9.5
10.8	Dark	8.6	3.2	5.6	6.4	6.2	5.2	8.5
	Dark	3.2	5.7	4.0	4.4	3.0	3.0	6.9
14.7	Dark	5.0	1.1	7.4	7.0	6.1	4.5	7.3
	Dark	5.9	3.8	5.0	6.0	5.4	4.5	9.2

Table 34b Identification of radioactivity in irradiated soil and dark controls treated with benzoyl-label

Time in Suntest (d)	Time natural sunlight (days)	% AR			Sum of other metabolites
		Parent 14.3–14.7 min	Met 4 4.3–4.5 min <sup>e, f</sup>	Met 9 13.7–13.9 min <sup>g</sup>	
Benzoyl-label					
0.0	0.0	91.2	2.4	0.0	0.0
	0.0	91.8			

Time in Suntest (d)	Time natural sunlight (days)	% AR			
		Parent 14.3–14.7 min	Met 4 4.3–4.5 min <sup>e, f</sup>	Met 9 13.7–13.9 min <sup>g</sup>	Sum of other metabolites
Benzoyl-label					
0.9	2.2	46.3	17.6	5.2	4.8
	2.3	45.8			
2.8	7.2	29.9	31.7	3.4	11.1 <sup>h</sup>
	7.4	23.5			
5.8	14.8	9.1	47.6	2.5	9.4 <sup>h</sup>
	14.6	13.4			
9.8	25.5	6.6	39.4	5.7	14.0 <sup>h</sup>
	24.1	3.6			
13.8	35.0	4.8	32.7	4.6	13.5 <sup>h</sup>
	35.9	3.0			
0.9	dark	57.1	16.0	0.0	2.8
	dark	62.4			
2.8	dark	18.2	30.9	0.0	9.1 <sup>i</sup>
	dark	17.8			
5.8	dark	7.9	32.4	0.0	7.5 <sup>i</sup>
	dark	9.4			
9.8	dark	8.0	36.1	0.0	7.2 <sup>i</sup>
	dark	8.7			
13.8	dark	5.7	37.7	0.0	7.7 <sup>i</sup>
	dark	11.5			

<sup>a</sup> Each individual peak, < 5% of applied

<sup>b</sup> Match with AB-11 based on retention time

<sup>c</sup> Match with AB-12 based on retention time

<sup>d</sup> Match with AB-13 based on retention time

<sup>e</sup> Sum of activity in concentrated extract and aqueous residue

<sup>f</sup> Match with B-1 based on retention time

<sup>g</sup> Match with AB-8 based on retention time

<sup>h</sup> Each individual peak < 5% of applied

<sup>i</sup> Each individual peak < 5% of applied, except one metabolite (Met 11) with retention time 19.0–19.2 min ( $\leq$  5.9% of applied)

Single first order  $DT_{50}$  and  $DT_{90}$  values (in natural sunlight equivalent days) for irradiated soil and dark controls are presented in Tables 23–29. The results indicated there is no difference between the  $DT_{50}$ s in irradiated soils and dark controls.

Table 35 Half-lives for degradation of cyflumetofen in irradiated soil and dark controls

System	$M_0$ %	$K$ days <sup>-1</sup>	$DT_{50}$ days	$DT_{90}$ days	$r^2$	$\chi^2$	Number of data points
Butylphenyl-label							
irradiated soil	99.1 ± 4.5	0.673 ± 0.101	1.0	3.4	0.972	13.37	12
dark control	99.3 ± 5.1	0.729 ± 0.123	0.95	3.2	0.965	14.95	12
Benzoyl-label							
Irradiated soil	86.7 ± 4.5	0.480 ± 0.063	1.4	4.8	0.960	15.4	12
Dark control	91.8 ± 4.0	0.502 ± 0.056	1.4	4.6	0.970	12.9	12

In conclusion, photolytic degradation on soil surfaces is not expected to play a role in the overall fate of cyflumetofen residues in soil, regardless of radio-labelled cyflumetofen used.

#### *Hydrolytic degradation in aquatic systems (Nakamura, 2004a)*

Portions of 2.5 mL butylphenyl- or benzoyl-label cyflumetofen in acetonitrile were added to 250 mL sterilized, deoxygenated buffer solutions of pH 4, 5, 7 and 9 at a concentration of 0.1 mg/L. The fortified buffer solutions were incubated in the dark at  $25 \pm 1$  °C for 30 days (pH 4, 5 and 7), 10 days (pH 7, benzoyl-label) or 1 day (pH 9). Sterility and pH were checked at the beginning and end of the



incubation period. Samples were taken at various time points. Immediately after sampling, the pH 7 and pH 9 samples were acidified (acetic acid) to slow hydrolysis. Samples were partitioned into an organic phase (acetonitrile) and aqueous phase by C<sub>18</sub>-SPE. Radioactivity was determined by LSC and the organic phase was subjected to chromatographic analysis (HPLC-RAM, TLC, LC-MS). The aqueous phase for the final sampling point and selected sampling points of benzoyl-label incubations was acidified (HCl) and subjected to PLS-2 SPE. Radioactivity in organic and aqueous phases was determined by LSC. The resulting organic phase (acetonitrile) was subjected to analysis using HPLC-RAM. Metabolite identification was done by comparison of retention times of HPLC, running factors (TLC) and MS spectra (LC-MS ion-trap) with reference standards of metabolites (A1, B1, AB-1, A-18). Metabolites for which no references were available were identified by LC-MS (ion-trap). Hydrolytic half-lives for cyflumetofen were calculated by linear regression using first order kinetics.

The mean total recovery for butylphenyl-label was 94.2 to 103.5% of applied radioactivity and that for benzoyl-label was 97.6 to 102.2%.

Degradation of cyflumetofen and its identified degradates at pH 4–9 at 25 °C are shown in the following Tables. Hydrolysis was fastest at pH 9 and slower in lower pH.

The degradation products of hydrolysis exceeding 10% of the applied radioactivity were A-1 (max 14.44% AR at pH 5–7), A-2 (max 44.12% AR at pH 5–7), A-18 (max 36.22% AR at pH 5–7), AB-1 (max 44.51% AR at pH 5–7) and B-1 (max 53.17% AR at pH 5–7). Other unknown fractions were all ≤ 7.76% AR, with the exception of BU14: 11.39% AR (pH 5, benzoyl-label). This fraction was identical to AU14 (pH 5, butylphenyl-label). When taking the average of the butylphenyl- and benzoyl-label incubations at pH 5, also this fraction was < 10% AR at all time points.

Table 36 Distribution of radioactivity after hydrolysis of butylphenyl- and benzoyl-label cyflumetofen at pH 4 at 25 °C

Time, d	Butylphenyl-label, % of applied			Benzoyl-label, % of applied		
	Organic	Aqueous	Mass balance	Organic	Aqueous	Mass balance
0	97.66	nd	97.66	99.05	1.52	100.57
1	98.15	nd	98.15	92.04	6.13	98.17
3	97.36	nd	97.36	87.01	14.11	101.12
7	99.04	nd	99.04	73.46	25.11	98.56
14	99.55	nd	99.55	61.61	38.55	100.16
21	99.07	nd	99.07	54.25	43.99	98.24
30	98.24	nd	98.24	51.76	48.40	100.16

nd = Not detected

Table 37 Distribution of radioactivity after hydrolysis of butylphenyl- and benzoyl-label cyflumetofen at pH 5 at 25 °C

Time, d	Butylphenyl-label, % of applied			Benzoyl-label, % of applied		
	Organic	Aqueous	Mass balance	Organic	Aqueous	Mass balance
0	94.15	nd	94.15	100.76	1.39	102.15
1	97.39	nd	97.39	90.49	7.13	97.62
3	100.53	nd	100.83	82.13	17.57	99.70
7	99.82	0.76	100.59	68.79	31.48	100.27
14	98.53	1.07	99.60	57.35	43.54	100.89
21	96.29	1.16	97.46	49.52	49.61	99.14
30	96.32	1.22	97.54	47.70	51.86	99.56

Nd = Not detected

Table 38 Distribution of radioactivity after hydrolysis of butylphenyl- and benzoyl-label cyflumetofen at pH 7 at 25 °C

Time, d	Butylphenyl-label, % of applied			Benzoyl-label, % of applied		
	Organic	Aqueous	Mass balance	Organic	Aqueous	Mass balance
0	98.92	nd	98.92	98.06	1.85	99.91
0.08 (2 h)	99.26	nd	99.26	92.39	8.90	101.29

Time, d	Butylphenyl-label, % of applied			Benzoyl-label, % of applied		
	Organic	Aqueous	Mass balance	Organic	Aqueous	Mass balance
0.33 (8 h)	98.92	nd	98.92	74.96	25.34	100.29
1	98.22	0.90	99.12	55.66	44.80	100.46
2	101.78	0.79	102.57	47.80	52.43	100.22
5	101.00	0.87	101.87	47.18	53.11	100.29
10	100.49	nd	100.49	48.03	52.85	100.88
30	100.83	nd	100.83	–	–	–

nd = Not detected

Table 39 Distribution of radioactivity after hydrolysis of butylphenyl- and benzoyl-label cyflumetofen at pH 9 at 25 °C

Time, min	Butylphenyl-label, % of applied			Benzoyl-label, % of applied		
	Organic	Aqueous	Mass balance	Organic	Aqueous	Mass balance
0	101.26	nd	101.26	97.86	1.98	99.84
5	100.19	nd	100.19	79.79	20.79	100.58
15	100.56	nd	100.56	62.37	38.46	100.83
30	100.02	nd	100.02	55.61	45.79	101.40
45	101.25	nd	101.25	55.81	45.01	100.81
90	101.70	nd	101.70	52.57	48.05	100.62
1440	102.45	1.02	103.47	50.29	50.31	100.61

Nd = Not detected

Table 40 Identification of radioactivity after hydrolysis of butylphenyl-label or benzoyl-label cyflumetofen at pH 4 at 25 °C

Time, d	% of applied								
	Parent	A-1	A-2	A-18	AB-1	AU16	UNK 1	UNK 2	UNK 3
Butylphenyl-label									
0	93.79	Nd		Nd	1.97	nd	nd	nd	Nd
1	90.73	4.63		Nd	Nd	nd	nd	nd	Nd
3	71.89	12.79	Nd	Nd	11.90	nd	nd	nd	Nd
7	48.46	20.78	2.46	2.55	19.51	2.79	nd	nd	Nd
14	26.93	24.89	8.67	6.10	22.23	6.56	2.36	nd	Nd
21	14.87	26.94	8.60	10.28	30.92	4.90	nd	nd	Nd
30	6.26	21.03	14.55	12.63	34.80	5.53	nd	1.15	1.47

Nd = Not detected

Time, d	% of applied						
	Parent	B-1	AB-1	UNK 1 (BU16)	UNK 2	UNK 3	UNK 4
Benzoyl-label							
0	97.43	1.52	Nd	Nd	nd	nd	nd
1	88.62	6.13	2.56	Nd	nd	nd	nd
3	73.69	14.11	12.49	Nd	nd	nd	nd
7	48.95	25.90	13.62	4.78	2.50	1.87	nd
14	24.61	40.56	22.59	6.25	3.12	nd	2.30
21	15.18	44.80	29.80	7.76	nd	nd	nd
30	6.50	48.40	34.17	6.39	3.53	nd	nd

Nd = Not detected

Table 41 Identification of radioactivity after hydrolysis of butylphenyl-label or benzoyl-label cyflumetofen at pH 5 at 25 °C

Time, d	% of applied										
	Parent	A-1	A-2	A-18	AB-1	AU16	AU17	UNK 1 (AU14)	UNK 2	UNK 3	UNK 4
Butylphenyl-label											
0	94.15	nd		nd	nd	Nd	nd	nd	nd	nd	nd

Time, d	% of applied										
	Parent	A-1	A-2	A-18	AB-1	AU16	AU17	UNK 1 (AU14)	UNK 2	UNK 3	UNK 4
1	85.38	6.00		nd	3.73	nd	nd	nd	nd	nd	nd
3	71.78	9.89	1.35	nd	9.03	2.15	3.08	nd	nd	nd	nd
7	45.34	10.02	6.38	4.35	16.19	5.67	5.86	4.06	nd	nd	nd
14	18.91	5.73	7.62	8.04	23.67	11.97	13.60	4.16	nd	nd	4.83
21	8.71	0.89	14.12	8.63	18.72	15.19	17.62	3.71	1.31	1.92	5.46
30	5.15	0.51	12.41	7.37	19.93	15.78	21.09	3.82	nd	2.81	7.46

Nd = Not detected

AU14 = BU14 (pH5)

Time, d	% of applied									
	Parent	B-1	AB-1	BU17	UNK 1 (BU14)	UNK 2	UNK 3	UNK 4	UNK 5	
Benzoyl-label										
0	99.99	1.39	nd	nd	nd	nd	nd	nd	nd	nd
1	82.89	8.23	3.14	nd	3.36	nd	nd	nd	nd	nd
3	68.76	17.57	8.60	1.64	3.14	nd	nd	nd	nd	nd
7	43.02	31.48	8.89	3.09	7.05	2.64	nd	2.55	nd	nd
14	19.14	43.54	20.61	4.87	11.39	nd	nd	nd	nd	nd
21	8.70	49.61	15.83	7.50	3.23	2.93	2.36	5.98	3.00	3.00
30	2.85	52.62	23.35	9.55	5.35	2.51	nd	nd	3.33	3.33

Nd = Not detected

BU14 = AU14 (pH5)

Table 42 Identification of radioactivity after hydrolysis of butylphenyl-label or benzoyl-label cyflumetofen at pH 7 at 25 °C

Time, h	% of applied						
	Parent	A-1	A-2	A-18	AB-1	UNK 1 (AU16)	Others
Butylphenyl-label							
0	97.95	nd		nd	nd	nd	nd
2	82.29	7.48		nd	3.33	1.75	nd
8	53.31	14.44	3.62	6.53	12.91	4.01	2.48
24	19.02	9.85	15.70	17.94	29.58	3.95	nd
48	6.32	1.95	14.64	32.91	36.44	4.91	1.99
120	Nd	0.22	19.12	36.22	44.51	nd	nd
240	Nd	24.30		31.98	41.61	2.60	nd
720	Nd	44.12		10.85	36.96	5.03	nd

Nd = Not detected

Time, h	% of applied				
	Parent	B-1	AB-1	UNK 1 (BU16)	Others
Benzoyl-label					
0	98.06	1.85	nd	nd	Nd
2	86.97	8.90	2.38	nd	Nd
8	55.81	25.34	15.91	2.39	Nd
24	16.83	44.80	30.59	5.57	Nd
48	3.29	53.17	38.37	2.80	Nd
120	Nd	53.11	40.00	3.02	2.54
240	Nd	52.85	41.18	1.92	1.92

nd: not detected

Table 43 Identification of radioactivity after hydrolysis of butylphenyl-label or benzoyl-label cyflumetofen at pH 9 at 25 °C

Time, min	% of applied						
	Parent	A-1	A-2	A-18	AB-1	UNK 1 (AU16)	Others
Butylphenyl-label							
0	101.26	Nd		nd	nd	nd	nd
5	64.94	17.48	nd	2.01	13.29	nd	nd
15	26.87	28.27	3.66	7.22	26.02	4.13	3.15
30	16.46	24.77	6.16	11.74	35.99	2.73	nd
45	9.02	27.90	nd	20.98	40.27	3.08	nd
90	Nd	3.70	15.05	33.66	39.60	4.16	2.87
1440	Nd	nd	6.17	48.80	44.75	nd	nd

Nd = Not detected

Time, min	% of applied				
	Parent	B-1	AB-1	Others	UNK 1
Benzoyl-label					
0	97.86	1.98	nd	nd	nd
5	63.14	20.79	16.65	nd	nd
15	25.88	38.46	33.11	nd	nd
30	10.08	47.39	43.16	nd	nd
45	11.53	46.75	34.48	1.67	3.69
90	5.58	48.05	41.85	3.07	nd
1440	Nd	50.31	45.68	nd	nd

Nd = Not detected

Table 44 Kinetic parameters for hydrolysis of cyflumetofen in aqueous buffer solutions of pH 4, 5, 7 and 9 at 25 °C

Test	DT <sub>50</sub>	DT <sub>90</sub>	first order equation	r <sup>2</sup>
pH4				
Butylphenyl-label	7.7 d	25.6 d	lnC = -0.09008t + ln(95.0235)	0.9991
Benzoyl-label	7.7 d	25.6 d	lnC = -0.08980t + ln(94.9601)	0.9977
Mean	7.7 d	25.6 d		
pH 5				
Butylphenyl-label	6.0 d	20.0 d	lnC = -0.11504t + ln(97.4473)	0.9988
Benzoyl-label	6.0 d	20.0 d	lnC = -0.11496t + ln(96.5132)	0.9994
Mean	6.0 d	20.0 d		
pH 7				
Butylphenyl-label	10.3 h	34.1 h	lnC = -0.06751 + ln(94.9138)	0.9984
Benzoyl-label	9.4 h	31.1 h	lnC = -0.07395t + ln(99.7485)	0.9997
Mean	9.8 h	32.6 h		
pH9				
Butylphenyl-label	11.5 min	38.0 min	lnC = -0.06053t + ln(88.0009)	0.9428
Benzoyl-label	9.1 min	30.4 min	lnC = -0.07583t + ln(91.9483)	0.9918
Mean	10.3 min	34.2 min		

### Metabolism and Distribution in Succeeding Crops

#### Confined Rotational Crop Study (Grosshans et al., 2011a)

A confined rotational crop study was conducted with benzoyl- and butylphenyl-label cyflumetofen. The active substance was applied to bare sandy loam soil in plastic containers (40 cm × 60 cm) at an application rate of 1 × 400 g ai/ha using an automatic spray track system. After the soil aging periods, ploughing was simulated by mixing the treated and untreated soil layers using a spade before replanting. The nature and the quantities of the radioactive residues were investigated in lettuce (immature and mature), white radish (top and root) and spring wheat (forage, hay, straw, chaff and

grain) after plant back intervals (PBI) of 30, 120 and 365 days. They are cultivated under natural climatic conditions but protected from rain. Plant samples were harvested at maturity, and additional immature lettuce samples as well as spring wheat forage samples (in part dried to hay) were taken 26 to 39 days and 47 to 62 days after planting or sowing, respectively. Soil samples were taken after application, after ploughing and after harvest of the mature crops for each plant back interval.

Mature and immature lettuce leaves were sampled with the roots remaining in the soil. Ripe white radishes were pulled from the soil and separated into the root and the remaining green parts (top). Immature green plants of spring wheat were sampled (wheat forage) and partly dried to wheat hay. Furthermore, ears of mature wheat and straw were cut off with scissors. Afterwards, ears were separated into chaff and grain using a thresher. For the plant back intervals of 120 and 356 DAT (benzoyl-label), chaff and straw samples were combined. For the butylphenyl-label, chaff and straw samples were generally combined. Soil samples were taken after application into petri dishes, after the individual plant back intervals and after harvest of the mature crops.

All samples were stored in a freezer at -18 °C or below, immediately after they were taken and until they were transferred to the laboratory. The storage conditions remained constant until analysis started and during the whole period of the study. Lettuce, radish and wheat forage and hay samples were homogenized with dry ice using a planetary mill or a blender. Samples of wheat hay, wheat straw and wheat straw + chaff were also frozen in liquid nitrogen and generally homogenized using a meat grinder. Individual hay samples were frozen and coarsely cut prior to homogenization with the meat grinder or homogenized with the blender. Samples of wheat grain and chaff were frozen in liquid nitrogen or dry ice and homogenized using a centrifugal grinding mill or with the blender (grain 365 PBI).

The radioactive residues in homogenized plant or soil samples were determined by oxidative combustion of small aliquots using LSC. Weighed subsamples of homogenized plant material were extracted three times with a sufficient volume of acetonitrile using a Polytron blender and the radioactivity was determined (LSC). Subsamples of spring wheat hay, straw and chaff were soaked in water before extraction. The residues were further extracted with appropriate volumes of water (two times). The water extracts were also radioassayed. The results of the acetonitrile and the water extraction were summed and referred to as extractable radioactive residues (ERR). The residue after solvent extraction (acetonitrile and water) of each sample was dried at room temperature under a fume hood, homogenized and aliquots were combusted for the determination of the residual radioactive residue (RRR). The total radioactive residues (TRR) were the result of the combustion analyses or the calculated sum of ERR and RRR values.

The residual radioactive residues of spring wheat hay, straw, chaff and grain (plant back interval of 30 days in the case of the benzoyl-label, plant back intervals of 120 and 365 days in the case of the butylphenyl-label) were further characterized using sequential solubilisation procedures. In the case of the benzoyl-label, the first solubilisation step with aqueous ammonia solution was also applied for the other rotational crop matrices. The residues from spring wheat hay, straw, chaff and grain after ammonia solubilisation (both labels) were subjected to two consecutive solubilisation steps applying the following hydrolysing enzymes: macerozyme/cellulase and amylase/amyloglucosidase. In the case of the benzoyl-label (30 PBI), the incubations with enzymes were followed by two non-selective solubilisation steps with hydrochloric acid.

The nature of the radioactive residues in the acetonitrile and aqueous extracts (with a sufficient level of radioactivity) as well as solubilizates (from RRR matrices, only spring wheat grain 30 PBI, benzoyl-label) of individual matrices of the different crops and labels was investigated using different HPLC methods with radio-detection. Peak assignment and identification of degradates in extracts was done by comparison of the retention times and the elution profiles/degradate patterns of extracts with reference items in at least two different HPLC methods. Further information for the identification of degradates in the case of the benzoyl-label was obtained from GC-MS or LC-MS analyses. In addition, <sup>14</sup>C-labelled reference items were combined with selected extracts for co-chromatography to identify the degradates unequivocally.

Following the spray application of cyflumetofen to soil at an application rate of  $1 \times 0.40$  kg ai/ha, the total radioactive residues (TRR) were measured. They were significantly higher after application of the benzoyl-label compared to the butylphenyl-label for all rotational crop matrices, except for spring wheat grain for the plant back intervals of 120 and 365 days.

The residue concentrations in lettuce (immature and mature samples) did not exceed 0.057 mg eq/kg or 0.018 mg eq/kg for all plant back intervals for the benzoyl- and the butylphenyl-label, respectively.

The TRR in white radish root in the case of the benzoyl-label decreased from 0.027 to 0.008 mg eq/kg with increasing soil aging period, whereas in white radish top, the residue levels were slightly higher (decreasing from 0.137 to 0.036 mg eq/kg). In the case of the butylphenyl-label, all residue concentrations in white radish were below 0.010 mg eq/kg.

In spring wheat, the highest residue levels were measured in chaff (benzoyl-label, 30 PBI: 1.271 mg eq/kg), straw (benzoyl-label, 30 PBI: 0.484 mg eq/kg), straw + chaff (0.259 and 0.058 mg eq/kg for the benzoyl-label, 120 and 365 PBI; 0.027 to 0.107 mg eq/kg for the butylphenyl-label) and hay (decreasing from 0.636 to 0.053 mg eq/kg for the benzoyl-label and ranging from 0.016 to 0.086 mg eq/kg for the butylphenyl-label). The total radioactive residues in wheat forage decreased from 0.133 to 0.004 mg eq/kg in the case of the benzoyl-label and were around 0.010 mg eq/kg in the case of the butylphenyl-label. The total radioactive residues in wheat grain accounted for 0.021 to 0.166 mg eq/kg for the benzoyl-label and < 0.010 to 0.063 mg eq/kg for the butylphenyl-label.

The residue concentrations in the top soil layer after aging and ploughing stayed largely stable (plant back intervals 30, 120 and 365 PBI) for the benzoyl- (0.054 to 0.064 mg eq/kg) and butylphenyl-label (0.062 to 0.075 mg eq/kg). The residue levels in soil after harvest of the mature crops also remained more or less stable for both labels, ranging from 0.020 to 0.089 mg eq/kg (benzoyl-label) and from 0.033 to 0.090 mg eq/kg for the butylphenyl-label.

Table 45 Total radioactive residues in rotational crop matrices following cultivation in soil treated with benzoyl-label cyflumetofen after plant back intervals of 30, 120 and 365 days

Crop parts	Combustion value mg eq/kg	TRR calculated mg eq/kg <sup>a</sup>
Plant back interval: 30 DAT		
Immature lettuce (35 DAP <sup>b</sup> )	0.059	0.057
Mature lettuce (48 DAP)	0.056	0.057
White radish root (70 DAP)	0.032	0.027
White radish top (70 DAP)	0.150	0.137
Spring wheat forage (54 DAP)	0.148	0.133
Spring wheat hay (54 DAP)	0.704	0.636
Spring wheat straw (103 DAP)	0.450	0.484
Spring wheat chaff (103 DAP)	1.249	1.271
Spring wheat grain (103 DAP)	0.170	0.166
Plant back interval: 120 DAT		
Immature lettuce (36 DAP)	0.031	0.040
Mature lettuce (50 DAP)	0.022	0.021
White radish root (65 DAP)	0.012	0.025
White radish top (65 DAP)	0.070	0.066
Spring wheat forage (58 DAP)	0.058	0.058
Spring wheat hay (58 DAP)	0.307	0.316
Spring wheat straw + chaff (107 DAP)	0.243	0.259
Spring wheat grain (107 DAP)	0.048	0.045
Plant back interval: 365 DAT		
Immature lettuce (31 DAP)	0.041	0.038
Mature lettuce (51 DAP)	0.021	0.018
White radish root (71 DAP)	0.007	0.008
White radish top (71 DAP)	0.044	0.036

Crop parts	Combustion value mg eq/kg	TRR calculated mg eq/kg <sup>a</sup>
Spring wheat forage (62 DAP)	0.010	0.004
Spring wheat hay (62 DAP)	0.065	0.053
Spring wheat straw + chaff (114 DAP)	0.056	0.058
Spring wheat grain (114 DAP)	0.020	0.021

<sup>a</sup> Sum of ERR (acetonitrile extract and water extract) and RRR (extraction residue)

<sup>b</sup> Days after planting

Table 46 Total radioactive residues in rotational crop matrices following cultivation in soil treated with butylphenyl-label cyflumetofen after plant back intervals of 30, 120 and 365 days

Crop parts	Combustion value mg eq/kg	TRR calculated mg eq/kg <sup>a</sup>
Plant back interval: 30 DAT		
Immature lettuce (26 DAP <sup>b</sup> )	0.021	0.018
Mature lettuce (53 DAP)	0.006	n. d.
White radish root (74 DAP)	0.001	n. d.
White radish top (74 DAP)	0.005	n. d.
Spring wheat forage (60 DAP)	0.006	n. d.
Spring wheat hay (60 DAP)	0.022	0.016
Spring wheat straw + chaff (102 DAP)	0.028	0.027
Spring wheat grain (102 DAP)	0.008	n. d.
Plant back interval: 120 DAT		
Immature lettuce (27 DAP)	0.006	n. d.
Mature lettuce (48 DAP)	0.004	n. d.
White radish root (69 DAP)	0.004	n. d.
White radish top (69 DAP)	0.009	n. d.
Spring wheat forage (47 DAP)	0.015	0.013
Spring wheat hay (47 DAP)	0.081	0.086
Spring wheat straw + chaff (111 DAP)	0.107	0.107
Spring wheat grain (111 DAP)	0.058	0.063
Plant back interval: 365 DAT		
Immature lettuce (39 DAP)	0.004	n. d.
Mature lettuce (55 DAP)	0.003	n. d.
Mature lettuce small-sized (55 DAP)	0.003	n. d.
White radish root (75 DAP)	0.002	n. d.
White radish top (75 DAP)	0.005	n. d.
Spring wheat forage (50 DAP)	0.006	n. d.
Spring wheat hay (50 DAP)	0.020	0.017
Spring wheat straw + chaff (98 DAP)	0.030	0.027
Spring wheat grain (98 DAP)	0.030	0.031

<sup>a</sup> Sum of ERR (acetonitrile extract and water extract) and RRR (extraction residue)

<sup>b</sup> Days after planting

n. d. = not determined (due to the low residue concentrations, no extraction was performed)

Table 47 Total radioactive residues in soil samples following treatment with benzoyl-label cyflumetofen and plant back intervals of 30, 120 and 365 days

Soil samples (Days After Treatment DAT)	TRR mg eq/kg determined by direct combustion
After application (petri dishes)	
(0 DAT)	3.854
Plant back interval: 30 DAT	
After Ploughing	
(30 DAT)	0.065
After harvest of mature crops	
Lettuce (78 DAT)	0.042
White radish (100 DAT)	0.056
Spring wheat (133 DAT)	0.078
Plant back interval: 120 DAT	

Soil samples (Days After Treatment DAT)	TRR mg eq/kg determined by direct combustion
After Ploughing (120 DAT)	0.064
After harvest of mature crops	
Lettuce (170 DAT)	0.020
White radish (185 DAT)	0.042
Spring wheat (227 DAT)	0.089
Plant back interval: 365 DAT	
After Ploughing (365 DAT)	0.054
After harvest of mature crops	
Lettuce (416 DAT)	0.067
White radish (436 DAT)	0.063
Spring wheat (479 DAT)	0.052

Table 48 Total radioactive residues in soil samples following treatment with butylphenyl-label cyflumetofen and plant back intervals of 30, 120 and 365 days

Soil samples (Days After Treatment DAT)	TRR mg eq/kg determined by direct combustion
After application (petri dishes) (0 DAT)	4.580
Plant back interval: 30 DAT	
After Ploughing (30 DAT)	0.075
After harvest of mature crops	
Lettuce (83 DAT)	0.083
White radish (104 DAT)	0.045
Spring wheat (132 DAT)	0.085
Plant back interval: 120 DAT	
After Ploughing (120 DAT)	0.070
After harvest of mature crops	
Lettuce (168 DAT)	0.033
White radish (189 DAT)	0.071
Spring wheat (231 DAT)	0.090
Plant back interval: 365 DAT	
After Ploughing (365 DAT)	0.062
After harvest of mature crops	
Lettuce (420 DAT)	0.066
Lettuce small-sized (420 DAT)	0.069
White radish (440 DAT)	0.035
Spring wheat (463 DAT)	0.066

The extractability of the radioactive residues with acetonitrile and water in the case of the benzoyl-label was equal to or above 75% of the TRR for all rotational crop matrices, except for wheat forage (365 PBI: 66.1% TRR) and wheat grain (30 and 365 PBI: 45.9% to 66.7% TRR). In the case of the butylphenyl-label, the extractability of the radioactive residues was notably lower, accounting for 57.9% TRR for immature lettuce (30 PBI), 35.8% to 65.7% TRR for wheat forage, hay and straw + chaff and 18.8% to 20.4% TRR for wheat grain. For both labels, the main portions of the radioactive residues were generally extracted with acetonitrile, except for spring wheat grain where only low portions were extracted with acetonitrile.



Liquid/liquid partition of the acetonitrile extracts of immature lettuce, spring wheat hay and spring wheat straw + chaff between ethyl acetate and water (performed in the case of the butylphenyl-label only) yielded the main portions of the residues (22.6% to 46.1% TRR) in the organosoluble fraction and minor portions (11.7% to 20.1% TRR) in the water phase.

In the case of the benzoyl-label, the radioactive residues in the extracts of all rotational crop matrices mainly consisted of a medium polar component which was identified as trifluoroacetic acid (38.6% to 105.3% TRR, ranging from 0.004 to 1.076 mg eq/kg). In addition, in spring wheat grain 30 PBI, the main degradate trifluoroacetic acid was accompanied by minor amounts of the degradate B-1 (< 0.001 mg eq/kg or 0.2% TRR in the acetonitrile extract) and a peak representing one or several non-polar components, possibly including the unchanged parent molecule (present in both extracts, 0.006 mg eq/kg or 4.1% TRR in sum). The degradates, trifluoroacetic acid and B-1, are only detectable with the benzoyl-label.

In the case of the butylphenyl-label, a series of label-specific degradates was observed in the samples prepared from the extracts, all in minor concentrations (each below 0.010 mg eq/kg, only one non-polar derivative in immature lettuce (30 PBI, 11.6% TRR) and one medium polar component in wheat hay (120 PBI, 10.2% TRR) were above 10% of the TRR. The main portion of the extracted and analysed radioactive residues in all matrices except wheat grain was composed of medium polar derivatives of cyflumetofen (up to 39.4% TRR in sum per matrix, up to twenty peaks per sample). These medium polar degradates eluted in the same range of polarity as the degradates in the goat urine sample, used as a reference item. Therefore, some of the medium polar degradates formed during metabolism of cyflumetofen in the lactating goat after hydrolytic cleavage of the parent molecule (e.g. hydroxylated derivatives of degradate A-2) or similar components may also be present in rotational crops. Besides the medium polar components, up to seven polar peaks were detected in the extractable radioactive residues (representing up to 14.9% TRR in sum per matrix), one non-polar derivative was found in immature lettuce (30 PBI, 11.6% TRR), and eleven non-polar peaks were observed in spring wheat straw and chaff (30 PBI, 6.8% TRR in sum). In the case of wheat grain (120 PBI), the radioactive residues in the water extract mainly consisted of polar components (4.8% TRR in sum), accompanied by minor amounts of medium polar components (0.5% TRR in sum). Since the absolute concentration of these minor degradates was < 0.01 mg eq/kg, no further characterization was deemed necessary.

The residual radioactive residues after solvent extraction of the matrices sampled after the plant back interval of 30 days ranged from a minimum of 11.0% TRR (white radish top) to a maximum of 54.1% TRR (spring wheat grain) for the benzoyl-label. Therefore, the residual radioactive residues from these matrices (30 PBI) were further characterized by treatment with an aqueous ammonia solution which solubilized portions of 1.5% to 2.4% TRR (except for wheat grain, where 15.1% TRR were solubilized). In the cases of spring wheat hay, straw, chaff and grain, the solubilisation with ammonia was followed by incubation with enzymes (macerozyme/cellulase and amylase/amyloglucosidase) and by homogenization or reflux with hydrochloric acid. The most effective solubilisation steps for most of the wheat matrices were the non-selective reflux with hydrochloric acid which released up to 7.3% TRR (spring wheat hay 30 PBI), followed by incubation with macerozyme/cellulase. In the case of wheat grain (30 PBI), treatment with macerozyme/cellulase solubilized 17.4% TRR, ammonia treatment released 15.1% TRR and incubation with amylases/amyloglucosidase was also quite effective and released 7.7% TRR. HPLC analysis of the ammonia solubilizate of wheat grain 30 DAT showed a degradate pattern similar to that of the water extract.

In the case of the butylphenyl-label, notable amounts of the radioactive residues of wheat hay, wheat straw + chaff and wheat grain (120 PBI and 365 PBI, 39.1% to 81.2% TRR) were not extracted with acetonitrile and water. The residual radioactive residues after solvent extraction of these matrices were therefore further characterized using a sequential solubilisation procedure including treatment with aqueous ammonia solution, macerozyme/cellulase and amylase/amyloglucosidase. The most effective solubilisation step was incubation with macerozyme/cellulase (releasing up to 20.4% TRR, wheat grain 365 PBI), except for wheat straw + chaff (365 PBI), where ammonia treatment was slightly more effective (solubilizing 6.5% TRR).

The solubilized residues had possibly been associated with or embedded/incorporated in insoluble plant material (e.g. proteins, cell wall polymers and starch).

Table 49 Extraction of radioactive residues in rotational crop matrices following their cultivation in soil treated with benzoyl-label cyflumetofen after plant back intervals of 30, 120 and 365 days

Crop parts	Distribution of radioactive residues									
	TRR <sup>a</sup>	Acetonitrile extract		Water extract		ERR <sup>a</sup>		RRR <sup>a</sup>		
	mg eq/kg	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	
Plant back interval: 30 DAT										
Immature lettuce	0.057	0.048	83.6	0.002	2.7	0.049	86.3	0.008	13.7	
Mature lettuce	0.057	0.046	79.4	0.002	4.3	0.048	83.7	0.009	16.3	
White radish root	0.027	0.016	58.5	0.005	19.0	0.021	77.5	0.006	22.5	
White radish root	0.036	0.022	61.6	0.005	14.3	0.027	75.9	0.009	24.1	
White radish top	0.137	0.115	83.5	0.008	5.5	0.122	89.0	0.015	11.0	
Spring wheat forage	0.133	0.099	74.5	0.007	5.3	0.106	79.8	0.027	20.2	
Spring wheat hay	0.636	0.421	66.2	0.057	8.9	0.478	75.0	0.159	25.0	
Spring wheat straw	0.484	0.362	74.8	0.029	6.1	0.391	80.9	0.093	19.1	
Spring wheat chaff	1.271	1.027	80.8	0.054	4.2	1.081	85.0	0.190	15.0	
Spring wheat grain	0.166	0.007	4.5	0.069	41.4	0.076	45.9	0.090	54.1	
Spring wheat grain	0.156	0.009	5.5	0.072	46.2	0.080	51.7	0.075	48.3	
Plant back interval: 120 DAT										
Immature lettuce	0.040	0.039	97.8	< 0.001	0.9	0.039	98.8	< 0.001	1.2	
Mature lettuce	0.021	0.020	96.8	< 0.001	1.3	0.021	98.1	< 0.001	1.9	
White radish root	0.025	0.024	94.2	0.001	2.4	0.024	96.6	0.001	3.4	
White radish top	0.066	0.063	96.3	0.001	2.2	0.065	98.6	0.001	1.4	
Spring wheat forage	0.058	0.054	93.7	0.002	4.0	0.057	97.7	0.001	2.3	
Spring wheat hay	0.316	0.283	89.4	0.023	7.2	0.306	96.6	0.011	3.4	
Spring wheat straw + chaff	0.259	0.243	93.9	0.010	3.9	0.253	97.7	0.006	2.3	
Spring wheat grain	0.045	0.006	13.0	0.032	71.6	0.038	84.7	0.007	15.3	
Plant back interval: 365 DAT										
Immature lettuce	0.038	0.036	95.2	0.001	2.2	0.037	97.4	0.001	2.6	
Mature lettuce	0.018	0.017	93.4	< 0.001	2.6	0.017	96.0	0.001	4.0	
White radish root	0.008	0.007	93.1	< 0.001	3.2	0.008	96.3	< 0.001	3.7	
White radish top	0.036	0.034	94.4	0.001	2.8	0.035	97.2	0.001	2.8	
Spring wheat forage	0.004	0.002	53.5	0.001	12.5	0.003	66.1	0.002	33.9	
Spring wheat hay	0.053	0.038	70.8	0.005	9.7	0.043	80.5	0.010	19.5	
Spring wheat straw + chaff	0.058	0.046	80.4	0.003	5.9	0.050	86.3	0.008	13.7	
Spring wheat grain	0.021	0.003	12.0	0.012	54.7	0.014	66.7	0.007	33.3	

<sup>a</sup> ERR = Extractable Radioactive Residue (acetonitrile extract and water extract), RRR = Residual (non-extractable) Radioactive Residue, TRR = sum of ERR and RRR

Table 50 Extraction of radioactive residues in rotational crop matrices following their cultivation in soil treated with butylphenyl-label cyflumetofen after plant back intervals of 30, 120 and 365 days

Crop parts	Distribution of radioactive residues									
	TRR <sup>a</sup>	Acetonitrile extract		Water extract		ERR <sup>a</sup>		RRR <sup>a</sup>		
	mg eq/kg	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	
Plant back interval: 30 DAT										
Immature lettuce	0.0179	0.0087	48.3	0.0017	9.6	0.0104	57.9	0.0076	42.1	
Spring wheat hay	0.0159	0.0070	44.1	0.0011	7.0	0.0081	51.1	0.0078	48.9	
Spring wheat straw + chaff	0.0270	0.0158	58.8	0.0019	7.0	0.0177	65.7	0.0092	34.3	
Plant back interval: 120 DAT										
Spring wheat forage	0.0133	0.0041	30.6	0.0020	15.2	0.0061	45.8	0.0072	54.2	

Crop parts	Distribution of radioactive residues									
	TRR <sup>a</sup>		Acetonitrile extract		Water extract		ERR <sup>a</sup>		RRR <sup>a</sup>	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Spring wheat hay	0.0860	48.2	0.0415	48.2	0.0053	6.1	0.0468	54.4	0.0393	45.6
Spring wheat straw + chaff	0.1067	49.1	0.0524	49.1	0.0126	11.8	0.0650	60.9	0.0417	39.1
Spring wheat grain	0.0630	0.5	0.0003	0.5	0.0115	18.3	0.0119	18.8	0.0511	81.2
Plant back interval: 365 DAT										
Spring wheat hay	0.0172	28.8	0.0050	28.8	0.0012	7.1	0.0062	35.8	0.0110	64.2
Spring wheat straw + chaff	0.0272	34.1	0.0093	34.1	0.0023	8.5	0.0116	42.6	0.0156	57.4
Spring wheat grain	0.0311	2.9	0.0009	2.9	0.0054	17.5	0.0063	20.4	0.0248	79.6

<sup>a</sup> ERR = Extractable Radioactive Residue (acetonitrile extract and water extract), RRR = Residual (non-extractable) Radioactive Residue after solvent extraction, TRR = sum of ERR and RRR

Table 51 Identified components in rotational crop matrices following their cultivation in soil treated with benzoyl-label cyflumetofen after a plant back interval of 30 days

Component	Crop parts									
	Immature lettuce		Mature lettuce		White radish root		White radish top			
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 30 DAT										
Trifluoroacetic acid	0.050	87.8	0.043	75.2	0.014	38.6	0.106	77.4		
Trifluoroacetic acid corrected <sup>a</sup>	0.075 mg/kg		0.065 mg/kg		0.021 mg/kg		0.159 mg/kg			
Sum of identified components	0.050	87.8	0.043	75.2	0.014	38.6	0.106	77.4		
Component	Crop parts									
	Spring wheat forage		Spring wheat hay		Spring wheat straw		Spring wheat chaff		Spring wheat grain <sup>b</sup>	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 30 DAT										
Trifluoroacetic acid	0.100	75.2	0.427	67.1	0.332	68.7	1.076	84.6	0.066	42.1
Trifluoroacetic acid Corrected <sup>a</sup>	0.150 mg/kg		0.641 mg/kg		0.498 mg/kg		1.614 mg/kg		0.099 mg/kg	
B-1	n. d.	n. d.	n. d.	n. d.	n. d.	n. d.	n. d.	n. d.	< 0.001	0.2
Sum of identified components	0.100	75.2	0.427	67.1	0.332	68.7	1.076	84.6	0.066	42.2

<sup>a</sup> Corrected mg/kg values referring to the calculation of the radioactivity and molecular mass of the degradate Trifluoroacetic Acid

<sup>b</sup> In the case of spring wheat grain, the sum of identified components in the extractable and residual radioactive residues (ERR + RRR) is given; the degradate Trifluoroacetic Acid was identified in the extracts (0.057 mg/kg or 36.3% TRR in sum) and in the ammonia solubilizate (0.010 mg eq/kg or 5.7% TRR)  
n. d. = not detected

Table 52 Identified components in rotational crop matrices following their cultivation in soil treated with benzoyl-label cyflumetofen after a plant back interval of 120 days

Component	Crop parts									
	Immature lettuce		Mature lettuce		White radish root		White radish top			
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 120 DAT										
Trifluoroacetic acid	0.042	105.3	0.015	72.0	0.013	53.1	0.060	91.0		
Trifluoroacetic acid Corrected <sup>a</sup>	0.063 mg/kg		0.023 mg/kg		0.020 mg/kg		0.090 mg/kg			
Sum of identified components	0.042	105.3	0.015	72.0	0.013	53.1	0.060	91.0		

Component	Crop parts							
	Immature lettuce		Mature lettuce		White radish root		White radish top	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Component	Crop parts							
	Spring wheat forage		Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 120 DAT								
Trifluoroacetic Acid	0.056	96.9	0.252	79.5	0.246	94.9	0.030	66.6
Trifluoroacetic Acid Corrected <sup>a</sup>	0.084 mg/kg		0.377 mg/kg		0.369 mg/kg		0.045 mg/kg	
Sum of identified components	0.056	96.9	0.252	79.5	0.246	94.9	0.030	66.6

<sup>a</sup> Corrected mg/kg values referring to the calculation of the radioactivity and molecular mass of the degradate Trifluoroacetic Acid

Table 53 Identified components in rotational crop matrices following their cultivation in soil treated with benzoyl-label cyflumetofen after a plant back interval of 365 days

Component	Crop parts							
	Immature lettuce		Mature lettuce		White radish root		White radish top	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Component	Crop parts							
	Spring wheat forage		Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 365 DAT								
Trifluoroacetic Acid	0.038	100.5	0.014	78.3	0.004	44.8	0.037	103.4
Trifluoroacetic Acid Corrected <sup>a</sup>	0.058 mg/kg		0.021 mg/kg		0.005 mg/kg		0.056 mg/kg	
Sum of identified components	0.038	100.5	0.014	78.3	0.004	44.8	0.037	103.4
Component	Crop parts							
	Spring wheat forage		Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 365 DAT								
Trifluoroacetic Acid	n. d.	n. d.	0.027	51.1	0.027	46.6	0.011	52.6
Trifluoroacetic Acid Corrected <sup>a</sup>	n. d.		0.041 mg/kg		0.040 mg/kg		0.017 mg/kg	
Sum of identified components	n. d.	n. d.	0.027	51.1	0.027	46.6	0.011	52.6

<sup>a</sup> corrected mg/kg values referring to the calculation of the radioactivity and molecular mass of the degradate Trifluoroacetic acid  
n. d. = not detected

Table 54 Characterized radioactive residues (ERR<sup>a</sup>+RRR<sup>a</sup>) from rotational crop matrices following their cultivation in soil treated with butylphenyl-label cyflumetofen after a plant back interval of 30, 120 and 365 days

Characterized Radioactive Residues	Crop parts									
	Immature lettuce		Spring wheat forage		Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 30 DAT										
Characterized from ERR by HPLC analysis	0.0079	44.3	no extraction		0.0087	54.3	0.0142	52.6	no extraction	
Number of HPLC peaks	up to 7 peaks				up to 19 peaks		up to 31 peaks			
Maximum HPLC peak	0.0021	11.6 <sup>b</sup>			0.0011	7.2	0.0018	6.7		

Characterized Radioactive Residues	Crop parts									
	Immature lettuce		Spring wheat forage		Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Characterized from ERR by Extraction / Work-up	0.0021	11.9			0.0014	9.0	0.0034	12.8		
Characterized from RRR by Solubilization	not applied				not applied		not applied			
Total Characterized	0.0101	56.2			0.0101	63.3	0.0176	65.4		
Plant back interval: 120 DAT										
Characterized from ERR by HPLC analysis	no extraction		not applied		0.0266	30.9	0.0443	41.5	0.0033	5.3
Number of HPLC peaks					2 x 2 peaks		up to 19 peaks	7 peaks		
Maximum HPLC peak					0.0088	10.2 °	0.0095	8.9	0.0013	2.1
Characterized from ERR by Extraction / Work-up			0.0061	45.8	0.0085	9.9	0.0044	4.2	0.0058	9.2
Characterized from RRR by Solubilization			not applied		0.0138	16.0	0.0107	10.0	0.0261	41.5
Total Characterized			0.0061	45.8	0.0489	56.8 <sup>d</sup>	0.0594	55.6 <sup>e</sup>	0.0353	56.0
Plant back interval: 365 DAT										
Characterized from ERR by HPLC analysis	no extraction		no extraction		not applied		not applied		not applied	
Characterized from ERR by Extraction / Work-up					0.0062	35.8	0.0116	42.6	0.0063	20.4
Characterized from RRR by Solubilization					0.0033	19.1	0.0040	14.8	0.0130	41.7
Total Characterized					0.0095	55.0	0.0156	57.4	0.0193	62.1

<sup>a</sup> ERR = Extractable Radioactive Residues (acetonitrile extract and water extract),

RRR = Residual Radioactive Residues (after solvent extraction)

<sup>b</sup> The second most abundant HPLC peak represented 0.0018 mg eq/kg or 9.8% TRR;

<sup>c</sup> The second and third most abundant HPLC peaks both represented 0.0062 mg eq/kg or 7.2% TRR

<sup>d</sup> Calculated losses of 0.0117 mg eq/kg or 13.6% TRR in sum occurred during work-up of the acetonitrile extract

<sup>e</sup> Calculated losses of 0.0151 mg eq/kg or 14.2% TRR in sum occurred during work-up of the acetonitrile extract

Table 55 Characterization of residual radioactive residues (RRR) after solvent extraction of rotational crop matrices following their cultivation in soil treated with benzoyl-label cyflumetofen after a plant back interval of 30 days

Fraction / Supernatant	Crop matrix									
	Immature lettuce		Mature lettuce		White radish root		White radish top			
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR		
Plant back interval: 30 DAT										
RRR	0.008	13.7	0.009	16.3	0.006	22.5	0.015	11.0		
Ammonia solubilizate	0.001	1.7	0.001	1.5	0.001	2.4	0.002	1.5		
Sum of solubilized radioactive residues	0.001	1.7	0.001	1.5	0.001	2.4	0.002	1.5		
Final residue	0.006	10.5	0.004	6.5	0.005	20.0	0.012	9.0		
Procedural recovery	89.1%		49.1%		99.6%		95.9%			
Fraction / Supernatant	Crop matrix									
	Spring wheat forage		Spring wheat hay		Spring wheat straw		Spring wheat chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 30 DAT										
RRR	0.027	20.2	0.159	25.0	0.093	19.1	0.190	15.0	0.090	54.1
Ammonia solubilizate	0.002	1.8	0.014	2.1	0.008	1.8	0.020	1.6	0.025	15.1
Ammonia residue	0.024	17.9	0.148	23.2	0.104	21.5	0.170	13.4	0.064	38.6

Fraction / Supernatant	Crop matrix									
	Immature lettuce		Mature lettuce			White radish root		White radish top		
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR		
Macerozyme/cellulase solubilizate	n. a.	n. a.	0.024	3.8	0.012	2.4	0.023	1.8	0.029	17.4
Amylase/amyloglucosidase solubilizate	n. a.	n. a.	0.011	1.7	0.003	0.7	0.013	1.0	0.013	7.7
Amylase/amyloglucosidase residue	n. a.	n. a.	0.105	16.5	0.082	16.9	0.128	10.1	0.022	13.2
1 N HCl homogenization	n. a.	n. a.	0.003	0.4	0.001	0.2	0.002	0.2	0.002	1.2
6 N HCl reflux	n. a.	n. a.	0.047	7.3	0.026	5.3	0.045	3.5	0.012	7.0
Sum of solubilized radioactive residues	0.002	1.8	0.098	15.4	0.050	10.4	0.103	8.1	0.080	48.4
Final residue	0.024	17.9	0.036	5.7	0.040	8.3	0.063	4.9	0.006	3.9
Procedural recovery <sup>a</sup>	97.3%		84.5%			97.9%		87.0%		96.7%

<sup>a</sup> Recovery calculated as (Sum of solubilized radioactive residues + Final residue) mg eq/kg × 100 / RRR mg eq/kg  
n. a. = not applied

Table 56 Characterization of residual radioactive residues (RRR) after solvent extraction of rotational crop matrices following their cultivation in soil treated with butylphenyl-label cyflumetofen after plant back intervals of 120 and 365 days

Fraction / Supernatant	Crop matrix					
	Spring wheat hay		Spring wheat straw + chaff		Spring wheat grain	
	mg eq/kg	% TRR	mg eq/kg	% TRR	mg eq/kg	% TRR
Plant back interval: 120 DAT						
RRR	0.0393	45.6	0.0417	39.1	0.0511	81.2
Ammonia solubilizate	0.0025	2.9	0.0033	3.0	0.0058	9.2
Macerozyme/cellulase solubilizate	0.0076	8.8	0.0045	4.2	0.0122	19.4
Amylase/amyloglucosidase solubilizate	0.0037	4.3	0.0029	2.7	0.0081	12.9
Sum of solubilized radioactive residues	0.0138	16.0	0.0107	10.0	0.0261	41.5
Final residue	0.0203	23.6	0.0279	26.2	0.0216	34.2
Procedural recovery <sup>a</sup>	86.7%		92.6%		93.3%	
Plant back interval: 365 DAT						
RRR	0.0110	64.2	0.0156	57.4	0.0248	79.6
Ammonia solubilizate	0.0009	5.1	0.0018	6.5	0.0022	6.9
Macerozyme/cellulase solubilizate	0.0015	8.5	0.0013	4.7	0.0063	20.4
Amylase/amyloglucosidase solubilizate	0.0009	5.5	0.0010	3.6	0.0045	14.4
Sum of solubilized radioactive residues	0.0033	19.1	0.0040	14.8	0.0130	41.7
Final residue	0.0073	42.4	0.0118	43.2	0.0119	38.4
Procedural recovery	95.9%		101.2%		100.6%	

<sup>a</sup> Recovery calculated as (Sum of solubilized radioactive residues + Final residue) mg eq/kg × 100 / RRR mg eq/kg

With regard to the degradate patterns described above for the two labels, the degradation pathway of cyflumetofen in rotational crops can be proposed as follows: Hydrolytic cleavage of the parent molecule results in the formation of the degradate B-1, which is only detectable with the benzoyl-label, and one or several butylphenyl-label-specific intermediates which were not identified. The predominant degradate in the case of the benzoyl-label, Trifluoroacetic Acid, can be derived from the parent compound or the degradate B-1. The butylphenyl-label-specific cleavage products (such as the goat degradate A-2 which is formed by hydrolysis of the formic acid ester, decarboxylation and

hydrolytic detachment of the trifluoromethyl benzoyl moiety) are further transformed to a series of minor (< 0.01 mg eq/kg), mainly medium polar degradates.

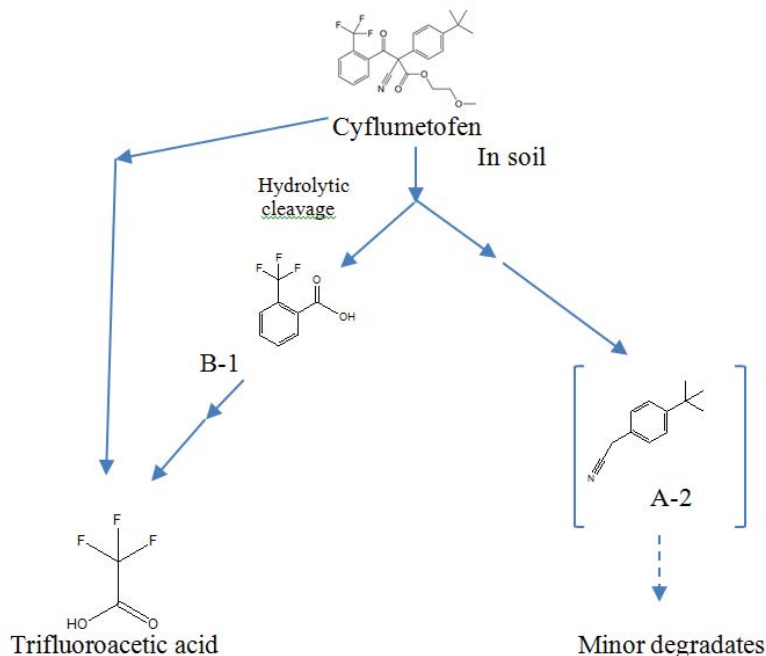


Figure 3 Proposed metabolic pathway in rotational crops (lettuce, radish and wheat)

## RESIDUE ANALYSIS

### *Analytical methods*

The Meeting received descriptions and validation data (including concurrent recovery data) for analytical methods for residues of cyflumetofen and its metabolites.

### *Analytical methods for plant matrices*

The descriptions of the analytical methods used in supervised residue trials and for enforcement are shown below.

#### *Method D1003*

Analyte: Cyflumetofen and metabolites B-1, AB-6 and AB-7

Matrix: Dry bean, dried grapes (raisins), orange (fruits, juice and oil), rice (grain, straw), soybean seed, tomato

LOQ: 0.01 mg/kg for each analyte in all matrices (defined as the lowest fortification level successfully tested)

**Description:** Crop materials Residues are extracted from plant matrices by shaking with acetonitrile followed by acetonitrile:water (75:25, v/v). An aliquot of the extract is centrifuged and, after removal of acetonitrile, acidified with formic acid. The residues are partitioned with a mixture of ethyl acetate: cyclohexane (75:25, v/v). An aliquot of the organic phase is concentrated in the presence of 0.1% aqueous formic acid, and re-dissolved in 0.1% formic acid in acetonitrile for determination with LC-MS/MS. Oil Residues are extracted from oil by shaking with a mixture of acetonitrile:water (75:25, v/v) and hexane. The hexane phase is removed and an aliquot of the aqueous acetonitrile phase is subjected to clean-up by partition with hexane or ethyl acetate:cyclohexane (75:25, v/v) followed by final determination as the above.

Juice Residues are extracted from juice by shaking with a mixture of acetonitrile:water (75:25, v/v). An aliquot of the aqueous acetonitrile extract is subject to clean-up and for final determination as the above.

#### LC-MS/MS condition

For data collection: Less concentrated final sample was used for analysis with highly sensitive LC-MS/MS instrument. From the acetonitrile extract of crop materials, acetonitrile is removed by concentration rather than centrifugation.

For enforcement: Larger amount of aliquot from the extract for sample clean-up to increase the residue concentration.

Reference: Carter *et al.*, 2011a

#### Validation of D1003 for data collection

Validation of Method D1003 for collecting data on the concentrations of the parent, B-1, AB-6 and AB-7 was conducted using the sample size of 0.1 g for crop materials, and 5 g for oil and orange juice.

Table 57 Recovery results obtained for the analysis of plant matrices and some of their processed foods with Method D1003 for data collection

Matrix	Analyte	No.	Fortification Level (mg/kg)	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Orange	Cyflumetofen	5	0.01	91–106	97	6	91–96	93	3
		5	1.0	96–104	100	3	83–108	97	10
		10	Overall	91–106	99	5	83–108	95	7
	B-1	5	0.01	74–85	80	7	81–87	84	3
		5	1.0	71–75	73	2	72–78	75	3
		10	Overall	71–85	77	7	72–87	80	6
	AB-6	5	0.01	92–96	95	2	86–98	94	5
		5	1.0	92–106	99	6	85–100	94	7
		10	Overall	92–106	97	5	85–100	94	6
	AB-7	5	0.01	92–99	95	3	87–103	95	7
		5	1.0	88–99	93	4	86–99	95	5
		10	Overall	88–99	94	4	86–103	95	6
Tomato	Cyflumetofen	5	0.01	78–124	94	20	84–133	100	20
		5	1.0	80–92	84	6	77–92	84	7
		10	Overall	78–124	89	16	77–133	92	17
	B-1	5	0.01	71–84	77	8	73–81	76	5
		5	1.0	75–97	82	11	70–93	81	12
		10	Overall	71–97	79	10	70–93	78	9
	AB-6	5	0.01	80–107	92	11	87–108	94	9
		5	1.0	79–90	84	5	82–88	86	3
		10	Overall	79–107	88	9	82–108	90	8
	AB-7	5	0.01	74–98	85	12	77–103	89	11
		5	1.0	71–83	78	7	83–92	87	4
		10	Overall	71–98	82	11	77–103	88	8



Matrix	Analyte	No.	Fortification Level (mg/kg)	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Bean, dry	Cyflumetofen	5	0.01	100–114	108	5	104–110	107	2
		5	1.0	98–119	110	8	92–117	106	9
		10	Overall	98–119	109	7	92–117	106	6
	B-1	5	0.01	70–76	72	3	79–94	86	8
		5	1.0	71–95	80	12	74–84	81	5
		10	Overall	70–95	76	11	74–94	83	7
	AB-6	5	0.01	100–116	110	6	103–119	111	7
		5	1.0	87–102	95	7	80–110	95	12
		10	Overall	87–116	103	10	80–119	103	12
	AB-7	5	0.01	99–118	108	7	99–119	111	7
		5	1.0	92–106	99	7	89–105	99	7
		10	Overall	92–118	103	8	89–119	105	9
Soybean	Cyflumetofen	5	0.01	80–93	86	6	90–109	97	8
		5	1.0	101–114	108	5	96–119	108	8
		10	Overall	80–114	97	13	90–119	102	10
(seed)	B-1	5	0.01	68–73	71	3	70–76	73	4
		5	1.0	88–104	99	7	87–102	96	6
		10	Overall	68–104	85	19	70–102	85	15
	AB-6	5	0.01	106–120	113	6	90–123	107	15
		5	1.0	98–114	109	6	92–108	102	7
		10	Overall	98–120	110	6	90–123	104	11
	AB-7	5	0.01	93–118	108	11	102–120	109	7
		5	1.0	99–108	103	3	92–113	105	9
		10	Overall	93–118	106	8	92–120	107	8
Rice	Cyflumetofen	5	0.01	73–98	88	12	74–102	96	13
		5	1.0	104–116	111	4	100–115	106	6
		10	Overall	73–116	100	14	74–115	101	11
(grain)	B-1	5	0.01	70–111	91	16	79–118	107	15
		5	1.0	93–111	98	8	92–109	99	6
		10	Overall	70–111	95	12	79–118	103	12
	AB-6	5	0.01	74–125	107	19	71–124	102	19
		4	1.0	105–107	106	0.8	96–121	111	8
		9	Overall	74–125	107	13	71–124	107	14
	AB-7	5	0.01	70–113	101	18	71–115	104	18
		5	1.0	99–115	106	6	102–112	107	4
		10	Overall	70–115	103	13	71–115	106	12
Orange juice	Cyflumetofen	5	0.01	78–89	85	5	78–98	88	10
		5	1.0	77–93	87	7	81–95	91	6
		10	Overall	77–93	86	6	78–98	89	8
	B-1	5	0.01	72–97	83	11	90–108	97	7
		5	1.0	68–81	74	6	71–87	78	8
		10	Overall	68–97	79	11	71–108	88	14
	AB-6	5	0.01	79–88	83	4	75–83	78	5
		5	1.0	83–92	87	5	81–99	87	9
		10	Overall	79–92	85	5	75–99	83	9
	AB-7	5	0.01	84–92	88	4	78–93	86	8
		5	1.0	78–90	83	6	81–90	85	4
		10	Overall	78–92	86	6	78–93	86	6
Orange oil	Cyflumetofen	5	0.01	70–75	73	3	71–81	78	5
		5	1.0	66–80	74	8	70–85	77	10
		10	Overall	66–80	74	6	70–85	77	7
	B-1	5	0.01	74–104	91	13	100–117	108	6
		5	1.0	71–117	99	18	85–112	99	12
		10	Overall	71–117	95	16	85–117	103	10
	AB-6	5	0.01	93–100	97	3	86–107	98	8
		5	1.0	77–98	88	9	80–101	93	9
		10	Overall	77–100	92	8	80–107	95	8
	AB-7	5	0.01	70–89	81	9	79–107	88	14
		5	1.0	75–93	81	9	70–89	79	10
		10	Overall	70–93	81	8	70–107	84	13

Matrix	Analyte	No.	Fortification Level (mg/kg)	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Dried grape	Cyflumetofen	5	0.01	102–123	109	8	94–117	104	9
		5	1.0	96–127	114	10	92–131	113	15
		10	Overall	96–127	111	9	92–131	109	12
	B-1	5	0.01	75–83	78	5	71–81	75	5
		5	1.0	72–83	78	6	71–89	80	9
		10	Overall	72–83	78	5	71–89	78	8
	AB-6	5	0.01	101–119	110	7	102–114	109	5
		5	1.0	101–131	116	9	100–134	115	12
		10	Overall	101–131	113	8	100–134	112	9
	AB-7	5	0.01	100–114	109	6	106–114	111	3
		5	1.0	96–136	112	14	94–141	114	16
		10	Overall	96–136	110	10	96–141	113	11
Rice straw	Cyflumetofen	5	0.01	97–111	104	5	97–118	104	8
		5	1.0	104–118	111	5	97–118	107	7
		10	Overall	97–118	107	6	97–118	105	7
	B-1	5	0.01	71–91	79	10	68–73	71	3
		5	1.0	71–84	77	7	70–76	73	4
		10	Overall	71–94	78	8	68–76	72	4
	AB-6	5	0.01	97–114	104	6	100–109	105	3
		5	1.0	90–101	97	5	95–107	98	5
		10	Overall	90–114	101	6	95–109	102	5
	AB-7	5	0.01	98–109	101	4	100–107	102	3
		5	1.0	85–99	93	7	86–99	92	6
		10	Overall	85–109	97	7	86–107	97	7

<sup>a</sup> Primary transition used for quantification:

Cyflumetofen: m/z 448.2 → 173.0 (ionization mode, positive)

B-1: m/z 189.1 → 69.0 (ionization mode, negative)

AB-6: m/z 466.4 → 173.0 (ionization mode, positive)

AB-7: m/z 448.2 → 344.2 (ionization mode, positive)

<sup>b</sup> Secondary transition used for confirmation:

Cyflumetofen: m/z 448.2 → 145.1

B-1: m/z 189.1 → 145.1

AB-6: m/z 466.4 → 145.0

AB-7: m/z 448.2 → 372.2

The method showed good linearity in the range of 0.035–10.0 µg/kg. No known interference was observed from crop components or from reagents, solvents and glassware used, except in rice straw which may show matrix interference for the B-1 secondary ion. As the matrix interference residue was greater than 20% of the LOQ, alternative chromatography was developed for the secondary ion for validation of the method for this analyte.

The mean recoveries for all matrices tested and at all fortification levels ranged between 71 and 116% with relative standard deviations between 2 and 20%. However, some individual recovery values were higher than 120%.

#### Validation of Method D1003 for enforcement

Validation of Method D1003, as an enforcement method, was conducted using the sample size of 5 g for all matrices tested. In this validation, the parent compound and B-1 metabolite were determined.

Table 58 Recovery results obtained for the analysis of plant matrices and some of their processed foods with Method D1003 for enforcement

Matrix	Analyte	Fortification Level (mg/kg)	No.	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Tomato	Cyflumetofen	0.01	5	87–103	94	7	84–104	95	10
		1.0	5	90–100	93	5	84–111	93	11
		Overall	10	87–103	93	6	84–111	94	10

Matrix	Analyte	Fortification Level (mg/kg)	No.	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
	B-1	0.01	5	78–90	84	5	71–97	78	14
		1.0	5	76–102	87	13	68–92	81	12
		Overall	10	76–102	86	9	68–97	80	12
Soybean	Cyflumetofen	0.01	5	82–103	92	9	74–108	94	13
		1.0	5	89–111	98	8	88–107	97	7
		Overall	10	82–111	95	9	74–108	96	10
(seed)	B-1	0.01	5	76–93	83	9	77–106	93	15
		1.0	5	80–102	91	10	80–109	88	14
		Overall	10	76–102	87	11	77–109	90	14
Rice	Cyflumetofen	0.01	5	77–109	88	15	79–100	87	10
		1.0	5	89–106	98	7	96–106	101	4
		Overall	10	77–109	93	12	79–106	94	11
(grain)	B-1	0.01	6	84–100	89	7	78–119	98	19
		1.0	5	87–107	95	8	76–105	86	15
		Overall	10	84–107	92	8	76–119	92	18
Bean, dry	Cyflumetofen	0.01	6	83–110	95	10	90–111	96	9
		1.0	5	97–108	103	5	92–106	101	6
		Overall	10	83–110	98	8	90–111	99	8
	B-1	0.01	6	85–106	96	8	89–125	104	12
		1.0	5	90–99	93	5	85–119	105	13
		Overall	10	85–106	95	6	85–125	104	12
Orange	Cyflumetofen	0.01	5	79–90	84	5	73–84	78	5
		1.0	5	80–88	84	4	79–95	86	7
		Overall	10	79–90	84	4	73–95	82	8
	B-1	0.01	5	73–105	87	14	90–115	100	10
		1.0	5	74–85	80	6	91–121	101	12
		Overall	10	73–105	84	12	90–121	101	10
Orange juice	Cyflumetofen	0.01	5	85–111	96	11	77–93	86	7
		1.0	5	70–101	80	15	71–92	76	12
		Overall	10	70–111	88	15	71–93	81	11
	B-1	0.01	5	70–96	77	14	77–101	89	10
		1.0	5	72–90	81	9	74–100	89	12
		Overall	10	70–96	79	11	74–101	89	10
Orange oil	Cyflumetofen	0.01	5	66–72	70	4	68–76	72	5
		1.0	5	82–96	87	7	86–101	90	7
		Overall	10	66–96	79	12	68–101	81	13
	B-1	0.01	5	84–105	96	9	81–107	97	11
		1.0	5	80–103	90	9	60–99	84	20
		Overall	10	80–105	93	9	60–107	90	17
Rice straw	Cyflumetofen	0.01	6	77–96	88	7	83–90	87	3
		1.0	5	94–104	99	4	92–106	101	6
		Overall	10	77–104	93	9	83–106	93	9
	B-1	0.01	6	79–116	90	16	79–118	92	16
		1.0	5	78–103	91	12	86–100	94	7
		Overall	10	78–116	90	14	79–118	93	12

<sup>a</sup> Primary transition used for quantification:

Cyflumetofen: m/z 448.2 → 173.0 (ionization mode, positive)

B-1: m/z 189.1 → 69.0 (ionization mode, negative)

<sup>b</sup> Secondary transition used for confirmation:

Cyflumetofen: m/z 448.2 → 145.1

B-1: m/z 189.1 → 145.1

The method showed good linearity in the range of 0.14–15.0 µg/kg. No known interference was observed from crop components or from reagents, solvents and glassware used, except in rice straw as described above.

The mean recoveries for all matrices tested and at all fortification levels ranged between 70 and 105%, within the acceptable range, with relative standard deviations between 3 and 19%. However, some individual analysis recoveries were slightly higher than 120% and some were lower than 70%.

*Independent laboratory validation of Method D1003 (Marin, 2011a)*

An independent laboratory validation of Method D1003 was conducted for the determination of cyflumetofen and metabolite B-1 in orange, wheat (grain and hay) and canola seed. Fortification levels used are 0.01 (LOQ) and 1.0 mg/kg.

Some modifications were made to the procedure of Method D1003 as follows:

Analyte: Cyflumetofen and metabolite B-1

LOQ: 0.01 mg/kg for each analyte in all matrices

Description: Residues are extracted from plant matrices by shaking with acetonitrile followed by acetonitrile:water (50:50, v/v) and centrifugation. An aliquot (2 mL) of the extract is transferred to a glass vessel containing aqueous formic acid. The extract is concentrated under a gentle flow of nitrogen and then partitioned with ethyl acetate: cyclohexane (75:25, v/v). After mixing, 2 g of sodium chloride is added and the mixture is partitioned for 5 minutes. After centrifugation, an aliquot (7 mL) of the extract is transferred to a glass tube containing 0.2 mL of aqueous formic acid. The extract is concentrated under a gentle flow of nitrogen and the volume is adjusted to 2.5 mL with acetonitrile containing formic acid. Residues are determined with LC-MS/MS.

Sample size used in this validation was 5 g.

Table 59 Recovery results obtained for the analysis of plant matrices with Method D1003 in independent laboratory validation

Matrix	Analyte	Fortification Level (mg/kg)	No.	Primary transition <sup>a</sup>			Secondary transition 2 <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Orange	Cyflumetofen	0.01	5	67–80	73	7	65–94	74	16
		1.0	5	84–92	89	3	91–103	96	5
		Overall	10	67–92	81	12	65–103	85	16
	B-1	0.01	5	77–87	82	6	76–84	79	4
		1.0	5	78–83	80	3	75–84	80	5
		Overall	10	77–87	81	5	75–84	80	4
Tomato	Cyflumetofen	0.01	5	61–85	74	12	60–90	72	17
		1.0	5	74–77	75	1	75–85	78	5
		Overall	10	61–85	75	8	60–90	75	12
	B-1	0.01	5	60–86	76	13	58–86	71	14
		1.0	5	77–81	79	3	75–79	78	3
		Overall	10	60–86	78	9	58–86	75	11
Wheat grain	Cyflumetofen	0.01	5	76–108	88	14	78–104	89	12
		1.0	5	84–95	90	4	85–106	92	10
		Overall	10	76–108	89	10	78–106	90	10
	B-1	0.01	5	81–86	83	2	78–85	80	4
		1.0	5	74–81	77	4	72–83	78	5
		Overall	10	74–86	80	5	72–85	79	4
Canola seed	Cyflumetofen	0.01	5	71–86	81	7	64–87	75	15
		1.0	5	88–102	94	6	92–111	99	8
		Overall	10	71–102	87	10	64–111	87	18
	B-1	0.01	5	73–75	74	1	69–74	72	3
		1.0	5	85–92	87	3	83–89	85	2
		Overall	10	73–92	81	9	69–89	79	9
Wheat hay	Cyflumetofen	0.01 <sup>c</sup>	5	84–91	88	5	80–98	87	8
		1.0 <sup>c</sup>	5	90–96	94	2	86–94	90	3
		1.0	5	93–101	97	4	78–91	83	6
		Overall	10	84–101	93	5	78–98	87	7
	B-1	0.01 <sup>c</sup>	5	85–109	95	12	45–59	51	12
		1.0 <sup>c</sup>	5	64–68	66	3	64–71	67	4
		1.0	5	75–81	78	3	75–81	77	3
		Overall	10	64–109	80	18	45–81	65	17

<sup>a</sup> Primary transition used for quantification:

Cyflumetofen: m/z 448.2 → 173.0 (ionization mode, positive)

B-1: m/z 189.1 → 69.0 (ionization mode, negative)

<sup>b</sup> Secondary transition used for confirmation:

Cyflumetofen: m/z 448.2 → 145.0

B-1: m/z 189.1 → 145.0

<sup>c</sup> Analysed by modified method.

The method showed good linearity in the range of 0.35–10.0 µg/kg. No known interference was observed from crop components or from reagents, solvents and glassware used, except in wheat hay.

The mean recoveries for all matrices tested and at all fortification levels ranged between 71 and 97%, within the acceptable range, with relative standard deviations between 1 and 19% with the exceptions of B-1 in wheat hay in which even the mean recoveries at 0.01 and 1 mg/kg were lower than 70%. Some individual analysis recoveries for matrices tested were lower than 70%.

#### *Analytical methods for animal matrices*

The descriptions of the analytical methods related to the current review are shown below.

All of these methods showed good linearity with a correlation coefficient  $\geq 0.99$ . There were no interferences at the analyte concentration higher than 30% of LOQ. Additional information on method performance is described later.

#### *Method D1202*

Analyte: Cyflumetofen and Metabolite B-1

Matrix: Bovine liver, meat and milk; and poultry eggs

LOQ: 0.001 mg/kg for each analyte in bovine milk; and 0.01 mg/kg for each analyte in other matrices.

Description: Bovine Liver and Poultry Eggs: Residues are extracted 5 g of sample by shaking with acetonitrile followed by acetonitrile-water (50:50, v/v). A salt mixture of magnesium sulfate (400 mg), sodium chloride (100 mg), sodium citrate sesquihydrate (50 mg), and sodium citrate dihydrate (100 mg) is added to an aliquot of the sample vortexed and centrifuged. The process is repeated with another salt mixture of magnesium sulfate (75 mg) and CUPSA (12.5 mg). An aliquot of the extract is concentrated to remove the acetonitrile to the aqueous phase in the presence of 20% formic acid in water. The extract is re-acidified and saturated with sodium chloride and partitioned with a mixture of ethyl acetate-cyclohexane (75:25, v/v). An aliquot of the organic phase is concentrated in the presence of 0.1% formic acid in water, and re-dissolved in 0.1% formic acid in acetonitrile for LC-MS/MS determination.

Bovine Meat: Residues are extracted from 5 g of sample by shaking with acetonitrile followed by acetonitrile-water (50:50, v/v). An aliquot of the extract is concentrated to remove the acetonitrile to the aqueous phase in the presence of 0.1% formic acid in water. The extract is saturated with sodium chloride and partitioned with a mixture of ethyl acetate-cyclohexane (75:25, v/v). An aliquot of the organic phase is concentrated in the presence of 0.1% formic acid in water, and re-dissolved in 0.1% formic acid in acetonitrile for LC-MS/MS determination.

Bovine Milk: Residues are extracted from 5 g or mL of sample by shaking with acetonitrile. An aliquot of the extract is concentrated to remove the acetonitrile to the aqueous phase in the presence of 20% formic acid in water. The extract is saturated with sodium chloride and partitioned with a mixture of ethyl acetate-cyclohexane (75:25, v/v). Aliquots of the organic phase are concentrated in the presence of 0.1% formic acid in water, and re-dissolved in 0.1% formic acid in acetonitrile. A salt mixture of magnesium sulfate (400 mg), sodium chloride (100 mg), sodium citrate sesquihydrate (50 mg), and sodium citrate dihydrate (100 mg) is added and vortexed and centrifuged. Another salt mixture of magnesium sulfate (75 mg) and CUPSA

(12.5 mg) mixture is then added and vortexed and centrifuged. An aliquot is taken for LC-MS/MS determination.

Reference: Sweeney and Nejad, 1998a

An independent laboratory validation of Method D1202 was conducted for the determination of cyflumetofen and metabolite B-1 in bovine liver, meat and milk and poultry eggs at fortification levels of the LOQ and  $100 \times$  LOQ: i.e., 0.001 and 0.10 mg/kg for bovine milk; and 0.01 and 1.0 mg/kg for other matrices.

Table 60 Recovery results obtained during independent laboratory validation of Method D1202 for bovine liver, meat and milk and poultry eggs.

Matrix	Analyte	Fortification level (mg/kg)	No.	Primary transition <sup>a</sup>			Secondary transition <sup>b</sup>		
				Range (%)	Mean (%)	RSD (%)	Range (%)	Mean (%)	RSD (%)
Bovine Liver	Parent	0.01 <sup>c</sup>	5	62–108	91	19	64–100	89	16
		1.0 <sup>c</sup>	5	108–124	115	5	107–123	115	5
		Overall	10	62–124	103	17	64–123	102	17
	B-1	0.01	5	74–81	78	4	71–81	78	5
		1.0	5	81–91	85	5	81–90	84	4
		Overall	10	74–91	81	6	71–90	81	6
Bovine Meat	Parent	0.01	5	69–82	77	7	71–84	77	7
		1.0	5	84–99	90	8	85–102	92	9
		Overall	10	69–99	84	11	71–102	85	12
	B-1	0.01	5	76–88	83	6	78–93	86	6
		1.0	5	70–85	77	8	71–86	77	8
		Overall	10	70–88	80	8	71–93	82	9
Bovine Milk	Parent	0.001	5	76–84	79	4	77–83	80	3
		0.10	5	103–112	107	3	104–112	107	3
		Overall	10	76–112	93	16	77–112	93	16
	B-1	0.001 <sup>c</sup>	5	74–90	83	8	74–86	81	6
		0.10 <sup>c</sup>	5	93–101	97	3	92–99	97	3
		Overall	10	74–101	90	10	74–99	89	10
Poultry Eggs	Parent	0.01 <sup>c</sup>	5	64–68	65	3	62–71	66	6
		1.0 <sup>c</sup>	5	60–69	67	6	59–69	66	6
		Overall	10	60–69	66	4	59–71	66	6
	B-1	0.01	5	77–89	86	6	84–103	95	8
		1.0	5	88–95	91	3	88–97	90	4
		Overall	10	77–95	88	5	84–103	93	6

<sup>a</sup> Primary transition used for quantification:

Cyflumetofen: m/z 448.2 → 173.2 (ionization mode, positive)

B-1: m/z 189.0 → 69.1 (ionization mode, negative)

<sup>b</sup> Secondary transition used for confirmation:

Cyflumetofen: m/z 448.2 → 145.1

B-1: m/z 189.1 → 145.1

<sup>c</sup> Results obtained using matrix matched standards.

The method showed good linearity in the range of 0.20–10.0 µg/kg. No known interference was observed from animal components or from reagents, solvents and glassware used.

The mean recoveries for bovine matrices tested and at all fortification levels ranged between 77 and 115% with RSDs between 3 and 19%. The mean recoveries of the parent in poultry eggs ranged between 65–67% with the standard deviations between 3–6% while those for B-1 ranged between 86–95% with the standard deviations between 3–8%.

### *Stability of pesticide residues in stored analytical samples*

The Meeting received information on freezer storage stability of cyflumetofen and three of its metabolites, B-1, AB-6 and AB-7 in various plant matrices, such as orange (fruit, juice and oil), apple (fruit and juice), leaf lettuce, kidney bean (dry), radish root, wheat grain and almond nutmeat (Gorden, 2011a and 2013a).

Homogenized samples (5 g) of each matrix were fortified separately with standard solutions containing Cyflumetofen and metabolites B-1, or AB-6 or AB-7 at a level of 0.1 mg/kg for each analyte and stored at -20 to -10 °C for up to 743–910 days (24–30 months).

After each specified period, a portion of sample was analysed with Method D1003 with the LOQ at 0.01 mg/kg. At the same time, a sample was freshly spiked at 0.1 mg/kg and analysed for concurrent procedural recovery.

Table 61 summarizes the storage periods, mean percent remaining and mean concurrent recovery at each time point.

Table 61 Stability of cyflumetofen and its metabolites B-1, AB-6 and AB-7 in frozen plant matrices fortified at 0.1 mg/kg

Matrix	Storage period (month)	Cyflumetofen		B-1		AB-6		AB-7	
		% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery
Almond	0	85	–	68	82	81	88	103	95
	0.2	86	85	–	–	–	–	–	–
	1	89	94	–	–	–	–	97	95
	2	–	–	–	–	93	88	–	–
	3	106	100	64	82	–	–	73	86
	5	–	–	–	–	85	88	–	–
	6	84	90	70	76	–	–	94	98
	9, 8	87	94	–	–	99	103	92	102
	10	–	–	67	83	–	–	–	–
	11	77	80	–	–	–	–	–	–
	12	–	–	–	–	106	126	91	82
	13	–	–	78	85	–	–	–	–
	18	85	91	–	–	–	–	–	–
	19	–	–	–	–	–	–	98	93
	21	–	–	–	–	77	89	–	–
	23	–	–	78	94	–	–	–	–
	25	79	107	–	–	–	–	–	–
26	–	–	–	–	–	–	99	118	
28	–	–	–	–	82	115	–	–	
30	–	–	74	101	–	–	–	–	
Apple	0	83	–	77	81	80	89	88	86
Fruit	0.2	67	83	–	–	–	–	–	–
	1	76	85	–	–	–	–	79	86
	2	–	–	–	–	81	89	–	–
	3	76	100	76	81	–	–	60	78
	5	–	–	–	–	79	85	–	–
	6	74	77	78	80	–	–	–	–
	7	–	–	–	–	–	–	77	82
	9	106	93	–	–	81	69	–	–
	10	–	–	66	105	–	–	87	80
	11	77	85	–	–	–	–	–	–
	12	–	–	–	–	102	109	71	92
	13	–	–	69	95	–	–	–	–
	18	70	75	–	–	–	–	–	–
	19	–	–	–	–	–	–	67	77
	21	–	–	–	–	59	75	–	–
	22	–	–	66	79	–	–	–	–
	25	79	77	–	–	–	–	–	–
26	–	–	–	–	–	–	80	82	
28	–	–	–	–	69	88	–	–	
29	–	–	58	77	–	–	–	–	
Apple	0	92	–	83	87	94	96	90	96
Juice	0.2	92	92	–	–	–	–	–	–
	1	83	101	–	–	–	–	90	96
	2	–	–	–	–	94	96	–	–
3	102	99	69	87	–	–	79	101	

## Cyflumetofen

Matrix	Storage period (month)	Cyflumetofen		B-1		AB-6		AB-7	
		% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery
	5	-	-	-	-	104	93	-	-
	6	105	91	79	84	-	-	-	-
	7	-	-	-	-	-	-	96	94
	9	106	98	-	-	90	89	103	91
	10	-	-	74	81	-	-	-	-
	11	104	90	-	-	-	-	-	-
	12	-	-	-	-	106	102	94	92
	13	-	-	83	94	-	-	-	-
	18	79	79	-	-	-	-	-	-
	19	-	-	-	-	-	-	92	91
	21	-	-	-	-	80	91	-	-
	22	-	-	93	94	-	-	-	-
	25	85	74	-	-	-	-	-	-
	26	-	-	-	-	-	-	87	78
	29	-	-	-	-	65	83	-	-
	30	-	-	51	78	-	-	-	-
Kidney bean	0	99	-	70	78	90	94	101	98
	0.2	90	99	-	-	-	-	-	-
	1	98	97	-	-	-	-	99	98
	2	-	-	-	-	-	-	-	-
	3	101	109	68	78	100	94	87	106
	5	-	-	-	-	91	90	-	-
	6	77	89	79	82	-	-	106	96
	9	72	99	-	-	98	90	101	102
	10	-	-	74	77	-	-	-	-
	11	63	92	-	-	-	-	-	-
	12	-	-	-	-	120	115	96	98
	13	-	-	66	76	-	-	-	-
	18	55	84	-	-	-	-	-	-
	19	-	-	-	-	-	-	99	92
	21	-	-	-	-	83	93	-	-
	23	-	-	85	93	-	-	-	-
	25	37	93	-	-	-	-	-	-
	26	-	-	-	-	-	-	91	99
	28	-	-	-	-	91	110	-	-
	30	-	-	77	106	-	-	-	-
Lettuce	0	93	-	60	65	84	78	80	86
	0.2	80	93	-	-	-	-	-	-
	1	80	89	-	-	-	-	74	86
	2	-	-	-	-	83	78	-	-
	3	82	99	58	65	-	-	-	-
	4	-	-	-	-	-	-	57	81
	5	-	-	-	-	80	88	-	-
	6	53	93	71	84	-	-	-	-
	7	-	-	-	-	-	-	65	90
	9	70	103	-	-	80	78	-	-
	10	-	-	58	103	-	-	82	102
	11	56	85	-	-	-	-	-	-
	12	-	-	-	-	84	109	67	92
	13	-	-	77	81	-	-	-	-
	18	46	86	-	-	-	-	-	-
	19	-	-	-	-	-	-	59	83
	21	-	-	-	-	68	84	-	-
	22	-	-	41	83	-	-	-	-
	26	-	-	-	-	-	-	50	91
	28	46	86	-	-	59	91	-	-
	29	-	-	88	91	-	-	-	-
Orange	0	86	-	66	75	83	83	92	81
Fruit	0.2	69	86	-	-	-	-	-	-



Matrix	Storage period (month)	Cyflumetofen		B-1		AB-6		AB-7	
		% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery
	1	75	93	—	—	—	—	83	81
	2	—	—	—	—	77	83	—	—
	3	86	86	59	75	—	—	65	82
	5	—	—	—	—	97	79	—	—
	6	82	83	81	69	—	—	—	—
	7	—	—	—	—	—	—	77	82
	9	106	93	—	—	81	68	—	—
	10	—	—	60	98	—	—	85	89
	11	80	85	—	—	—	—	—	—
	12	—	—	—	—	104	103	81	89
	13	—	—	77	83	—	—	—	—
	18	83	80	—	—	—	—	—	—
	19	—	—	—	—	—	—	103	81
	21	—	—	—	—	70	80	—	—
	22	—	—	66	88	—	—	—	—
	24	76	77	—	—	—	—	—	—
	26	—	—	—	—	—	—	56	81
	28	—	—	—	—	54	77	—	—
	29	—	—	76	85	—	—	—	—
Orange	0	93	—	82	90	87	99	93	97
Juice	0.2	82	93	—	—	—	—	—	—
	1	92	87	—	—	—	—	92	97
	2	—	—	—	—	91	99	—	—
	3	106	97	79	90	—	—	82	100
	5	—	—	—	—	104	107	—	—
	6	98	97	80	85	—	—	—	—
	7	—	—	—	—	—	—	96	98
	9	104	83	—	—	93	100	97	107
	10	—	—	75	81	—	—	—	—
	11	102	90	—	—	—	—	—	—
	12	—	—	—	—	88	101	89	97
	13	—	—	89	82	—	—	—	—
	18	78	86	—	—	—	—	—	—
	19	—	—	—	—	—	—	91	98
	21	—	—	—	—	80	97	—	—
	22	—	—	83	100	—	—	—	—
	25	93	93	—	—	—	—	—	—
	26	—	—	—	—	—	—	74	91
	29	—	—	—	—	63	90	—	—
	30	—	—	103	90	—	—	—	—
Orange	0	81	—	84	88	88	87	97	92
Oil	0.2	75	81	—	—	—	—	—	—
	1	62	84	—	—	—	—	97	92
	2	—	—	—	—	79	87	—	—
	3	92	84	68	88	—	—	—	—
	4	—	—	—	—	—	—	76	91
	5	—	—	—	—	111	103	—	—
	6	99	72	65	92	—	—	93	91
	9	81	72	—	—	103	104	104	92
	10	—	—	56	100	—	—	—	—
	11	82	72	—	—	98	86	—	—
	12	—	—	41	85	—	—	93	93
	18	79	80	—	—	—	—	—	—
	19	—	—	—	—	—	—	84	83
	21	—	—	—	—	85	96	—	—
	22	—	—	46	103	—	—	—	—
	25	97	83	—	—	—	—	—	—
	26	—	—	—	—	—	—	95	69
	29	—	—	—	—	99	87	—	—

Matrix	Storage period (month)	Cyflumetofen		B-1		AB-6		AB-7	
		% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery	% remaining	Procedural recovery
	30	–	–	56	115	–	–	–	–
Radish	0	102	–	68	80	88	82	83	81
Root	0.2	82	102	–	–	–	–	–	–
	1	83	95	–	–	–	–	69	81
	2	–	–	–	–	72	82	–	–
	3	86	101	67	80	–	–	–	–
	4	–	–	–	–	–	–	50	79
	5	–	–	–	–	84	88	–	–
	6	63	82	79	79	–	–	–	–
	7	–	–	–	–	–	–	60	79
	9	59	87	–	–	84	69	–	–
	10	–	–	67	89	–	–	62	99
	11	65	88	–	–	–	–	–	–
	12	–	–	–	–	82	102	43	94
	13	–	–	76	81	–	–	–	–
	18	55	75	–	–	–	–	–	–
	19	–	–	–	–	–	–	44	76
	21	–	–	–	–	77	78	–	–
	22	–	–	60	79	–	–	–	–
	24	60	84	–	–	–	–	–	–
	26	–	–	–	–	–	–	47	88
	28	–	–	–	–	74	94	–	–
	29	–	–	50	70	–	–	–	–
Wheat	0	102	–	86	86	102	97	94	90
Grain	0.2	102	102	–	–	–	–	–	–
	1	94	93	–	–	–	–	87	90
	2	–	–	–	–	95	97	–	–
	3	104	100	75	86	–	–	–	–
	4	–	–	–	–	–	–	72	88
	5	–	–	–	–	–	–	–	–
	6	81	85	75	88	–	–	86	96
	9	124	106	–	–	96	94	77	103
	10	–	–	68	79	–	–	–	–
	11	89	92	–	–	–	–	–	–
	12	–	–	–	–	115	120	80	103
	13	–	–	86	86	–	–	–	–
	18	80	88	–	–	–	–	–	–
	19	–	–	–	–	–	–	76	99
	21	–	–	–	–	93	94	–	–
	23	–	–	82	90	–	–	–	–
	25	72	96	–	–	–	–	–	–
	26	–	–	–	–	–	–	56	96
	29	–	–	–	–	80	103	–	–
	30	–	–	67	86	–	–	–	–

The results of the storage stability showed high uncertainty associated with the analytical method used.

Cyflumetofen was stable (> 70% remaining) when stored frozen at the temperature of –20 – 10 °C for the longest time tested (24 or 25 months) in almond, apple fruit, apple juice, orange fruit, orange juice, orange oil and wheat grain. However, it showed stability only up to 9 months in lettuce and 3 months in radish root.

Metabolite B-1 was stable frozen up to: 30 months, the longest time tested, in almond, kidney bean, lettuce, orange fruit, orange juice and wheat grain, 22 months in apple fruit and apple juice, and 6 months in orange oil.

Metabolite AB-6 was stable frozen up to: the longest period tested (28 or 29 months) in almond, apple fruit, kidney bean, orange juice, orange oil, radish root and wheat grain, 21 months in apple juice, lettuce and orange fruit.

Metabolite AB-7 was stable frozen up to: the longest period tested (26 months) in almond, apple fruit, apple juice, kidney bean, orange juice, and orange oil, 19 months in orange fruit and wheat grain, 12 months in lettuce, and 1 month in radish root.

**USE PATTERNS**

Cyflumetofen is used either alone or in combination with other active substances and provides knock-down and residual control of tetranychid mites shown in the following table. It is highly active contact miticides on egg, nymph and adult stage of these mites.

Table 62 Mites controlled by Cyflumetofen

<i>Bryobia rubrioculus</i>	<i>Panonychus ulmi</i>
<i>Eotetranychus willamettei</i>	<i>Petrobia lateens</i>
<i>Eotetranychus yumensis</i>	<i>Tetranychus cinnabarinus</i>
<i>Eutetranychus banksi</i>	<i>Tetranychus mcdanieli</i>
<i>Oligonychus pratensis</i>	<i>Tetranychus pacificus</i>
<i>Oligonychus ununguis</i>	<i>Tetranychus turkestani</i>
<i>Panonychus citri</i>	<i>Tetranychus urticae</i>

It has been registered in a number of countries for use on a variety of crops including citrus fruits; pome fruits; berries and other small fruits; fruiting vegetables, other than Cucurbits; and tree nuts. The authorized uses relevant to the supervised trials data submitted to the current Meeting are summarized in Table 63.

For the current review, a US label was provided which indicates that thorough and uniform spray coverage of foliage, with direct contact to target pest, is required for effective control and therefore sufficient volume of water is necessary to ensure thorough coverage of foliage.

Table 63 Registered use of cyflumetofen relevant to the residue evaluation by the current Meeting

Commodity	Country	Formulation Type and g/kg or g/L	F or, G <sup>a</sup>	Application					Application rate per treatment		PHI, days <sup>b</sup>
				Method	Max rate per season, kg ai/ha	Min interval, days	No. <sup>c</sup>	Timing	Rate, kg ai/ha	Min water vol., L/ha	
Citrus fruits											
Citrus fruits	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	935	7
Citrus fruits	Brazil	SC 200	F	Foliar spray	–	–	1		0.08 kg ai/hL	–	7
Pome fruits											
Pome fruits	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	935	7
Berries and other small fruits											
Grapes	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	468	14
Strawberry	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	468	1
Fruiting vegetables, other than Cucurbits											
Tomato	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	468	3
Eggplant	Japan	SC 200	F G	Foliar spray	–	–	2	At first sign of infestation	0.02 kg ai/hL	(1000–3500)	1
Tree nuts											

Commodity	Country	Formulation Type and g/kg or g/L	F or, G <sup>a</sup>	Application				Application rate per treatment		PHI, days <sup>b</sup>	
				Method	Max rate per season, kg ai/ha	Min interval, days	No. c	Timing	Rate, kg ai/ha		Min water vol., L/ha
Tree nuts	USA	SC 200	F	Foliar spray	0.40	14	2	At first sign of infestation	0.19–0.20	935	7

<sup>a</sup> F= outdoor or field use; G= glasshouse

<sup>b</sup> PHI= Pre-harvest interval

<sup>c</sup> Do not make more than 1 application before using an effective miticide with a different mode of action

## RESIDUES RESULTING FROM SUPERVISED TRIALS ON CROPS

The Meeting received residue data from supervised field trials conducted on citrus fruits (orange in Brazil and the USA, grapefruit and lemon in the USA), pome fruits (apple in Italy and the USA and pear in the USA); grapes (in the USA); strawberry (in the USA); fruiting vegetables, other than Cucurbits (tomato in the USA and eggplant in Japan; and tree nuts (in the USA).

Application rates and residue concentrations were reported as Cyflumetofen. Residue concentrations are recorded unadjusted for recoveries or for residue values in control samples. Where multiple samples were taken from a single plot, individual results are reported, and the calculated mean concentration is put in parentheses and used for estimation of maximum residue level. Where trials were conducted in the same location, with the same or similar varieties, same or similar formulations, and same equipment, and at the same or similar timing, they are not regarded as independent and only one result from these trials was chosen for the estimation of a maximum residue level. All the trials, except those on eggplant in Japan, were conducted outdoor using 20% SC formulations except one trial conducted in Italy on apple (25% SC). Residue concentrations are expressed in parent equivalents.

Residues from the trials conducted according to maximum GAP have been used for the estimation of maximum residue levels and they are underlined.

The results of supervised trials are summarized in the following tables:

Crop group	Commodities	Table no
Citrus fruits	Orange, grapefruit, lemon	Table 64
Pome fruits	Apple, pear	Table 65
Berries and other small fruits	Grapes	Table 66
	Strawberry	Table 67
	Tomato	Table 68
Fruiting vegetables, other than Cucurbits	Eggplant	Table 69
	Almond, pecan	Table 70
Tree nuts	Almond hulls	Table 71
By-products, used for animal feeding purposes, derived from fruits and vegetable processing		

### *Citrus fruits (orange, grapefruit and lemon)*

During the growing season of 2009 and 2010, 23 supervised residue trials were conducted on citrus fruits (orange, grapefruit and lemon) in Arizona (one trial), California (eight trials), Florida (12 trials) and Texas (two trials) in the USA. Three (two in CA and one in FL) of these trials were decline trials. At each test location, one untreated and one treated plot were established. The treated plot received two broadcast foliar applications of cyflumetofen (20% SC formulation) each at 0.193–0.207 kg ai/ha, with a 14-day (+/- one day) re-treatment interval (RTI), except one trial that had a 12-day RTI. The applications were made with ground equipment using 280–2818 L/ha of water. Commercially mature fruits of orange, grapefruit or lemon were collected 7 days after treatment (DAT). In the three decline trials, citrus samples were collected 0, 1, 3, 7, 14 and 21 DAT.

All samples were analysed for residues of cyflumetofen and its metabolites B-1, AB-6 and AB-7 using Method D1003 (with modifications), which quantifies residues by LC-MS/MS. Mean recoveries for all analytes were within the acceptable range of 70–120%. A recovery of 66% was obtained for B-1 on lemon fortified at the 0.01 mg/kg level. Several recovery values exceeded 120%. The LOD and LOQ for all analytes were 0.002 and 0.01 mg/kg, respectively. For B-1, when its concentration is expressed in parent equivalents, the LOQ will be 0.02 mg/kg.

In Brazil, during the growing season of 2007, four trials were conducted on oranges to determine the magnitude of the residues of cyflumetofen after treatment with 20% SC formulation of cyflumetofen. The formulation was applied two times at an application rate of 0.160 kg ai/ha and a spray volume of 2000 L/ha. In two of the four trials, the samples were taken immediately after the last application, 1, 7, 14 and 21. In the other two trials the samples were collected 7 days after the last application.

The residues of cyflumetofen were determined by LC-MS/MS. The LOQ was 0.05 mg/kg. The mean recoveries at 0.05 mg/kg spiked level were within the acceptable range of 70–120%.

Table 64 Results of supervised residue trials on citrus fruits (orange, grapefruit and lemon) conducted in Brazil and the USA.

CROP Country, Year Location variety Trial No.	Application				Portion analysedd	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP		2	0.2	Min. 935		7					(Interval: 14 days)
ORANGE											
USA, 2009 Orange, FL Hamlin R090446	Tractor mounted, PTO- Driven Airblast	2	0.2	1 × 1658 1 × 1640	Fruit	7	0.087 0.068 (0.078)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Volusia, FL Hamlin R090447	Tractor mounted, PTO- Driven Airblast	2	0.2	1 × 1648 1 × 1655	Fruit	7	0.095 0.080 (0.088)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Seminole, FL Navel R090448	Tractor mounted, PTO- Driven Airblast	2	0.2	1 × 699 1 × 706	Fruit	7	0.073 0.159 (0.116)	< 0.02 < 0.02	< 0.01 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Polk, FL Hamlin R090449	SS Airblast	2	0.2	1 × 808 1 × 800	Fruit	7	0.102 0.099 (0.101)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2010 Osceola, FL Valencia R090450	SS Airblast	2	0.2	1 × 747 1 × 718	Fruit	7	0.107 0.109 (0.108)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Lake, FL Hamlin R090451	SS Airblast	2	0.2	1 × 2232 1 × 2027	Fruit	7	0.046 0.075 (0.061)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Hobe Sound Martin, FL Hamlin R090452	Airblast	2	0.2	2 × 616	Fruit	7	0.107 0.084 (0.096)	< 0.02 < 0.02	0.013 0.014	0.012 0.015	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
USA, 2009 Hobe Sound Martin, FL Pineapple R090453	Airblast	2	0.2	1 × 2049 1 × 1984	Fruit	7	0.062 0.076 (0.069)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA Willacy, TX N-33 Navel R090454	FMC DP 50 Airblast Sprayer 9SR-77	2	0.2	1 × 585 1 × 579	Fruit	7	0.010 0.010 (0.010)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Tulare, CA Atwood Navel R090455	Broadcast	2	0.2	1 × 2109 1 × 2083	Fruit	0	0.055 0.061 (0.058)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
						1	0.055 0.049 (0.052)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						3	0.044 0.053 (0.0485)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.038 0.031 (0.0345)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						14	0.024 0.025 (0.0245)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						21	0.015 0.014 (0.0145)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2009 San Luis Obispo, CA Olinda Valencia R090456	Tractor- mounted, PTO- Driven Airblast	2	0.2	1 × 611 1 × 606	Fruit	7	0.016 0.011 (0.014)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Kern, CA Navel R090457	Broadcast	2	0.2	1 × 1193 1 × 2152	Fruit	7	0.067 0.059 (0.063)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
GAP in Brazil		1	0.08 kg ai/hL	–		7					
Brazil, 2007	(n.r.)	2	0.2	2000	Fruit	0	0.3	–	–	–	G. Casadei de Baptista 2007
						1	0.3	–	–	–	
						3	0.2	–	–	–	n.r.
						7	0.08	–	–	–	OTSA-0484-FR
						14	0.06	–	–	–	
Brazil, 2007	(n.r.)	2	0.2	2000	Fruit	7	< 0.05	–	–	–	G. Casadei de Baptista 2007 n.r.
											OTSA-0508-FR
Brazil, 2007	(n.r.)	2	0.2	2000	Fruit	0	0.1	–	–	–	G. Casadei de Baptista 2007
						1	0.1	–	–	–	
						3	0.08	–	–	–	n.r.
						7	0.06	–	–	–	OTSA-0511-FR
						14	< 0.05	–	–	–	
Brazil, 2007	(n.r.)	2	0.2	2000	Fruit	7	< 0.05	–	–	–	G. Casadei de Baptista 2007 n.r.
											OTSA-0513-FR
GRAPEFRUIT											
US GAP		2	0.2	Min. 935		7					(Interval: 14 days)

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
USA, 2009 Brevard, FL White Marsh R090458	Airblast Sprayer	2	0.2	1 × 700 1 × 712	Fruit	0	0.191 0.084 (0.1375)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt
						1	0.101 0.057 (0.079)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	350843 2012/7003656
						3	0.097 0.043 (0.070)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.046 0.031 (0.0385)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						14	0.059 0.023 (0.041)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						21	0.038 0.008 (0.023)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2009 Lake, FL White R090459	SS Airblast	2	0.2	1 × 1798 1 × 1588	Fruit	7	0.067 0.077 (0.072)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Polk, FL Ruby Red R090460	Broadcast	2	0.2	1 × 820 1 × 777	Fruit	7	0.039 0.040 (0.0395)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Willacy, TX Rio Red R090461	FMC DP 50 Airblast Spayer (SR- 77)	2	0.2	1 × 2495 1 × 2467	Fruit	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Porterville Tulare, CA Mellogold R090462	Tractor- mounted, PTO- Driven Airblast	2	0.2	1 × 724 1 × 765	Fruit	7	0.036 0.038 (0.037)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Strathmore Tulare, CA Mellogold R090463	Tractor- mounted, PTO- Driven Airblast	2	0.2	1 × 2253 1 × 2294	Fruit	7	0.027 0.019 (0.023)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
LEMON											
USA, 2009 St. Lucie, FL Bearss R090464	Airblast Sprayer	2	0.2	1 × 664 1 × 649	Fruit	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2009 Tulare, CA Pyror R090465	Tractor- mounted, PTO- Driven Airblast	2	0.2	1 × 2203 1 × 2063	Fruit	0	0.122 0.113 (0.1175)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt
						1	0.086 0.112 (0.099)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	350843 2012/7003656
						3	0.106 0.100 (0.103)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.077 0.086 (0.0815)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
						14	0.049 0.052 (0.0505)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						21	0.031 0.041 (0.036)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2009 Kern, CA Lisbon R090466	Tractor-mounted, PTO- Driven Airblast	2	0.2	1 × 714 1 × 742	Fruit	7	0.129 0.141 (0.135)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2010 Yuma, AZ Lisbon R090467	D-4015-03 Airblast	2	0.2	1 × 2818 1 × 2801	Fruit	7	0.021 0.021 (0.021)	< 0.02 < 0.02	< 0.01 < 0.01	0.010 0.011	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656
USA, 2010 San Luis Obispo, CA Eureka R090468	Tractor-mounted, PTO- Driven Airblast	2	0.2	1 × 280 1 × 299	Fruit	7	0.021 0.019 (0.020)	< 0.02 < 0.02	0.01 < 0.01	< 0.01 < 0.01	2011 / D.R. Hattermann, L.E. Crawford, S. Holt 350843 2012/7003656

<sup>a</sup> Metabolite residues expressed in parent equivalents.

n.r. = Not reported

#### *Pome fruits (apple and pear)*

During the years 2009 and 2010, 12 trials were conducted on apple in the USA (two trials in New York, one in Pennsylvania, one in Georgia, two in Wisconsin, one in Utah, one in California, two in Idaho, one in Oregon and one in Washington). At each test location, two broadcast foliar applications of 200 g/L SC formulation of cyflumetofen were made to apple trees, at a target application rate of 0.200 kg ai/ha, with a 13–15 day retreatment interval. The applications were made to plots at each site using either concentrate (478–857 L/ha of water, Treatment 2) or dilute (1129–2274 L/ha of water, Treatment 3) spray volumes with ground equipment (airblast sprayers). At each location, apple samples (fruit, one sample per untreated plot and duplicate samples per treated plot) were harvested at a 6 to 7 day preharvest interval (PHI). At one trial in New York (trial no. R090479) treated samples were collected at 1, 3, 8, 10, and 15 days after the last application, to evaluate residue decline. The residues of cyflumetofen and metabolites B-1, AB-6 and AB-7 were quantitated by Method D1003 (LC-MS/MS). Acceptable concurrent recovery data were obtained for each analyte. The LOQ was 0.01 mg/kg for each analyte. Apparent residues of cyflumetofen, B 1, AB 6 and AB 7 were < LOQ in/on all untreated samples.

During the growing season 2006, one trial was conducted in Italy in which apple trees were treated with two applications of 250 g/L SC formulation of cyflumetofen at either 0.1 kg ai/ha, 0.2 kg ai/ha, 0.3 kg ai/ha or 0.4 kg ai/ha, diluted with water immediately prior to application to a good coverage spray volume. Samples of apple fruit from the untreated and treated plots were taken immediately after the last application, 3, 7, 14 and 21 days after last application. The LOQ for cyflumetofen in apple was set at 0.05 mg/kg. Procedural recoveries run currently with test specimen at levels of 0.05 mg/kg and 5.0 mg/kg gave an overall mean recovery of 93% and 85% respectively.

During the years 2009 and 2010, five trials were conducted on pear in the USA (one trial in New York, one in California, one in Oregon and two in Washington). At each test location, two broadcast foliar applications of 200 g/L SC formulation of cyflumetofen were made to pear trees, at a target application rate of 0.200 kg as/ha, with a 13–15 day retreatment interval. The applications were made to plots at each site using either concentrate (565–831 L/ha of water) or dilute (1111–1996 L/ha of water) spray volumes with ground equipment (airblast sprayers). At each location, pear samples



(fruit, one sample per untreated plot and duplicate samples per treated plot) were harvested at a 7 to 8 day pre-harvest interval (PHI). At one trial in Washington, treated samples were collected at 1, 3, 7, 10, and 14 days after the last application, to evaluate residue decline. The residues of cyflumetofen and metabolites B-I, AB-6 and AB-7 were quantitated using Method D1003. Acceptable concurrent recovery data were obtained for each analyte. The LOQ was 0.01 mg/kg for each analyte. Apparent residues of cyflumetofen, B I, AB 6 and AB 7 were < LOQ in/on all untreated pear samples.

Table 65 Results of supervised residue trials on pome fruits (apple and pear) conducted in Italy and the USA

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP		2	0.2	Min. 935		7					(Interval: 14 days)
APPLE											
USA, 2009 Wayne, NY	Broad-cast foliar	2	0.2	1 × 562 1 × 571	Fruit	1	0.182 0.175	< 0.02 < 0.02	0.011 < 0.01	< 0.01 < 0.01	M. T. White 2011
Romes RCN R090479						3	0.196 0.125	< 0.02 < 0.02	< 0.01 0.011	< 0.01 < 0.01	375302 2011/7000143
						8	0.248 0.122 (0.185)	< 0.02 < 0.02	0.016 0.015	0.013 0.011	
						10	0.068 0.095	< 0.02 < 0.02	0.011 0.013	< 0.01 < 0.01	
						15	0.104 0.073	< 0.02 < 0.02	0.012 0.010	< 0.01 < 0.01	
		2	0.2	1 × 1129 1 × 1133	Fruit	1	0.121 0.122	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						3	0.079 0.065	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						8	0.065 0.055 (0.060)	< 0.02 < 0.02	0.011 0.010	< 0.01 < 0.01	
						10	0.051 0.071 (0.061)	< 0.02 < 0.02	0.011 0.011	< 0.01 < 0.01	
						15	0.041 0.046	< 0.02 < 0.02	< 0.01 0.010	< 0.01 < 0.01	
USA, 2009 Wayne, NY	Broad-cast foliar	2	0.2	1 × 609 1 × 610	Fruit	7	0.101 0.091 (0.096)	< 0.02 < 0.02	< 0.01 0.010	< 0.01 < 0.01	M. T. White 2011
Romes RCN R090480		2	0.2	1 × 2091 1 × 2097	Fruit	7	0.067 0.058 (0.0625)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Berks, PA	Broad-cast foliar	2	0.2	1 × 563 1 × 564	Fruit	6	0.160 0.238 (0.199)	< 0.02 < 0.02	< 0.01 0.010	0.011 0.012	M. T. White 2011
Starkrimson RCN R090481		2	0.2	1 × 2243 1 × 2249	Fruit	6	0.119 0.091 (0.105)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Fannin, GA	Broad-cast foliar	2	0.2	1 × 691 1 × 613	Fruit	6	0.214 0.139 (0.1765)	< 0.02 < 0.02	0.012 0.012	0.013 0.011	M. T. White 2011
Yate RCN R090482		2	0.2	1 × 1594 1 × 1398	Fruit	6	0.082 0.108 (0.095)	< 0.02 < 0.02	< 0.01 < 0.01	0.011 < 0.01	375302 2011/7000143
USA, 2009 Lenawee, MI	Broad-cast foliar	2	0.2	1 × 823 1 × 806	Fruit	7	0.168 0.153 (0.1605)	< 0.02 < 0.02	0.018 0.017	0.013 0.012	M. T. White 2011

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
Delicious RCN R090483		2	0.2	1 × 1778 1 × 1757	Fruit	7	0.080 0.137 (0.1085)	< 0.02 < 0.02	< 0.01 0.010	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Pepin, WI (Connel Red)	Broad-cast foliar	2	0.2	1 × 478 1 × 483	Fruit	6	0.130 0.122 (0.126)	< 0.02 < 0.02	0.012 0.015	< 0.01 0.013	M. T. White 2011
RCN R090484		2	0.2	1 × 1927 1 × 1941	Fruit	6	0.027 0.045 (0.036)	< 0.02 < 0.02	< 0.01 0.011	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Cache, UT	Broad-cast foliar	2	0.2	1 × 707 1 × 707	Fruit	7	0.042 0.042 (0.042)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Red Delicious RCN R090485		2	0.2	1 × 1598 1 × 1659	Fruit	7	0.061 0.059 (0.060)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Madera, CA	Broad-cast foliar	2	0.2	1 × 747 1 × 751	Fruit	7	0.033 0.047 (0.040)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Fuji RCN R090486		2	0.2	1 × 1888 1 × 1858	Fruit	7	0.032 0.024 (0.028)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Bingham, ID	Broad-cast foliar	2	0.2)	1 × 576 1 × 555	Fruit	7	0.022 < 0.01 (0.0135)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
McIntosh RCN R090487		2	0.2	1 × 1423 1 × 1335	Fruit	7	0.011 0.018 (0.0145)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2009 Bingham, ID	Broad-cast foliar	2	0.2	1 × 574 1 × 552	Fruit	7	0.078 0.040	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Golden Delicious RCN R090488		2	0.2	1 × 1439 1 × 1385	Fruit	7	0.077 0.122 (0.0995)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2010 Hood River, OR	Broad-cast foliar	2	0.2	1 × 857 1 × 822	Fruit	7	0.072 0.067 (0.0695)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Honey Crisp RCN R090489		2	0.2	1 × 2274 1 × 2235	Fruit	7	0.054 0.052 (0.053)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2010 Grant, WA	Broad-cast foliar	2	0.2	1 × 615 1 × 611	Fruit	7	0.209 0.177 (0.193)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Red Delicious RCN R090490		2	0.2	1 × 1413 1 × 1424	Fruit	7	0.123 0.143 (0.133)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
Italy, 2006	Mist blower	2	0.1	1 × 1241	Fruit	0	< 0.05	–	–	–	n.r.
Bologna	sprayer			1 × 1132		3	< 0.05	–	–	–	2007
Double red						7	< 0.05	–	–	–	AF/10943/OT
AF/10943/OT/						14	< 0.009	–	–	–	OTSA-0301-
1						21	< 0.009	–	–	–	FR
		2	0.2	1 × 1193	Fruit	0	0.0998	–	–	–	
				1 × 1175		3	0.0780	–	–	–	
						7	0.0596	–	–	–	
						14	< 0.05	–	–	–	
						21	< 0.05	–	–	–	
		2	0.3	1 × 1197	Fruit	0	0.149	–	–	–	
				1 × 1158		3	0.111	–	–	–	

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
						7	< 0.05	-	-	-	
						14	< 0.05	-	-	-	
						21	< 0.05	-	-	-	
		2	0.4	1 × 1197	Fruit	0	0.151	-	-	-	
				1 × 1153		3	0.0914	-	-	-	
						7	0.0501	-	-	-	
						14	0.0522	-	-	-	
						21	< 0.05	-	-	-	
PEAR											
USA, 2009 Wayne, NY	Broad-cast foliar	2	0.2	1 × 565 1 × 567	Fruit	8	0.127 0.125	< 0.02 < 0.02	0.038 0.027	0.036 0.028	M. T. White 2011
Bosc RCN R090491		2	0.2	1 × 1111 1 × 1118	Fruit	8	0.121 0.265 (0.1995)	< 0.02 < 0.02	0.034 0.032	0.034 0.035	375302 2011/7000143
USA, 2009 Tulare, CA	Broad-cast foliar	2	0.2	1 × 819 1 × 831	Fruit	7	0.045 0.038 (0.0415)	< 0.02 < 0.02	< 0.01 < 0.01	0.011 < 0.01	M. T. White 2011
Olympic RCN R090493		2	0.2	1 × 1808 1 × 1831	Fruit	7	0.064 0.087 (0.0755)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 0.012	375302 2011/7000143
USA, 2009 Hood River, OR	Broad-cast foliar	2	0.2	1 × 805 1 × 765	Fruit	7	0.032 0.039 (0.0355)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Starkrismson RCN R090494		2	0.2	1 × 1729 1 × 1996	Fruit	7	0.061 0.065 (0.063)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
USA, 2010 Grant, WA	Broad-cast foliar	2	0.2	1 × 607 1 × 610	Fruit	1	0.217 0.226	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. T. White 2011
Concord RCN R090495						3	0.206 0.203	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375302 2011/7000143
						7	0.125 0.106 (0.1155)	< 0.02 < 0.02	0.014 < 0.01	0.012 < 0.01	
						10	0.086 0.116	< 0.02 < 0.02	0.010 0.013	< 0.01 0.012	
						14	0.055 0.079	< 0.02 < 0.02	< 0.01 0.014	< 0.01 0.011	
		2	0.2	1 × 1398 1 × 1401	Fruit	1	0.215 0.231	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						3	0.231 0.170	< 0.02 < 0.02	0.012 0.010	0.011 < 0.01	
						7	0.125 0.114 (0.1195)	< 0.02 < 0.02	0.012 0.013	0.011 0.012	
						10	0.081 0.100	< 0.02 < 0.02	0.012 0.012	< 0.01 < 0.01	
						14	0.063 0.077	< 0.02 < 0.02	0.011 0.012	< 0.01 < 0.01	
USA, 2010 Grant, WA	Broad-cast foliar	2	0.2	1 × 610 1 × 608	Fruit	7	0.089 0.124 (0.1065)	< 0.02 < 0.02	0.013 0.016	0.011 0.014	M. T. White 2011
Bartlett RCN R090496		2	0.2	1 × 1403 1 × 1396	Fruit	7	0.085 0.125 (0.105)	< 0.02 < 0.02	0.010 0.011	< 0.01 < 0.01	375302 2011/7000143

<sup>a</sup> Metabolite residues expressed in parent equivalents.

n.r. = Not reported

## Berries and Other Small Fruits

## Grapes

During the 2009 growing season, a total of 12 trials were conducted on grapes in the USA (two trials in New York, eight in California, one in Washington, and one in Oregon). One in California was a decline trial. In all trials, two broadcast foliar applications of 200 g/L SC formulation of cyflumetofen at 0.193–0.206 kg ai/ha were made with a 14–15 re-treatment interval. The applications were made with ground equipment using either a dilute or concentrated spray volume. Grape Samples were collected at 14 DAT. In the decline trial, the samples were collected at 0, 3, 7, 14, 21 and 28 DAT.

Samples of grape were analysed for residues of cyflumetofen and metabolites B-1, AB-6 and AB-7 using Method D1003. Acceptable concurrent recovery data for grape commodities between 69% and 111% were obtained for each analyte. The LOQ was 0.01 mg/kg for each analyte.

Table 66 Results of supervised residue trials on Grapes conducted in the USA

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP		2	0.2	Min. 468		14					(Interval: 14 days)
USA, 2009 Seneca, NY Catawba R090511	broadcast foliar application	2	0.20	1 × 2313 1 × 2354	Fruit	14	0.252 0.290 (0.271)	0.064 0.061 (0.0625)	< 0.01 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Yates, NY Vidal Blanc R090512	broadcast foliar application	2	0.20	1 × 466 1 × 473	Fruit	14	0.406 0.439 (0.4225)	0.040 0.045 (0.0425)	< 0.01 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Kern, CA Thompson Seedless R090513	broadcast foliar application	2	0.20	1 × 721 1 × 703	Fruit	14	0.073 0.124 (0.0985)	0.024 0.024 (0.024)	< 0.01 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Kingsberg Fresno, CA Crimson R090514	broadcast foliar application	2	0.20	1 × 1884 1 × 1889	Fruit	14	0.175 0.151 (0.163)	0.035 0.040 (0.0375)	0.013 0.016	0.013 0.015	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 San Luis Obispo, CA Syrah R090515	broadcast foliar application	2	0.20	1 × 466 1 × 448	Fruit	14	0.014 0.031 (0.0225)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Monterey, CA Syrah R090516	broadcast foliar application	2	0.20 0.20	1 × 628 1 × 602	Fruit	14	0.102 0.085 (0.0935)	0.024 0.024 (0.024)	0.011 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Kerman Fresno, CA Thompson Seedless R090517	broadcast foliar application	2	0.20 0.20	1 × 1876 1 × 1886	Fruit	14	0.125 0.117 (0.121)	0.071 0.061 (0.066)	0.013 0.013	0.012 0.011	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995
USA, 2009 Kerman Fresno, CA Thompson Seedless R090518	broadcast foliar application	2	0.20 0.20	1 × 704 1 × 694	Fruit	14	0.126 0.121 (0.1235)	0.040 0.038 (0.039)	0.013 0.016	0.014 0.016	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT d	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.				
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7					
USA, 2009 Plainview Tulare, CA Crimson R090519	broadcast foliar application	1+	0.20 0.19	1 × 1968 1 × 2112	Fruit	0	0.312	0.024	< 0.01	< 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995				
							0.349	0.028	< 0.01	< 0.01					
					3	0.289 0.367	0.028 0.028	< 0.01 < 0.01	< 0.01 < 0.01						
					7	0.246 0.285	0.024 0.026	< 0.01 < 0.01	< 0.01 < 0.01						
					14	0.179 0.199 (0.189)	0.026 0.033 (0.0295)	< 0.01 < 0.01	< 0.01 < 0.01						
					21	0.096 0.181	0.045 0.040	< 0.01 < 0.01	< 0.01 0.012						
					28	0.122 0.104	0.031 0.033	< 0.01 < 0.01	0.010 < 0.01						
USA, 2009 Earlimart Tulare, CA Crimson R090520	broadcast foliar application	2	0.20 (0.20)	1 × 1880 1 × 1890	Fruit	14	0.225 <sup>b</sup> 0.223 <sup>b</sup> (0.224)	0.024 0.024 (0.024)	< 0.01 < 0.01	0.012 <sup>a</sup> 0.011 <sup>a</sup>	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995				
USA, 2009 Yakima, CA Riesling R090521	broadcast foliar application	1+	0.21 0.20	1 × 888 1 × 859	Fruit	14	0.271 0.177 (0.224)	0.024 0.024 (0.024)	< 0.01 < 0.01	< 0.01 < 0.01	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995				
USA, 2009 Benton, OR Pinot R090522	broadcast foliar application	1+	1 × 0.19 1 × 0.20	1 × 2612 1 × 2542	Fruit	14	0.154 0.142 (0.148)	0.085 0.075 (0.080)	0.034 0.027	0.021 0.019	D. R. Hattermann, T. Lott, 2011 375306 2011/7003995				

<sup>a</sup> Metabolite residues expressed in parent equivalents.

<sup>b</sup> Results of multiple analyses of individual field samples.

### Strawberry

During the growing season 2009–2010, a total of eight field trials on strawberries were conducted in the USA (one trial in New York, one in Georgia, one in Florida one in Wisconsin, three in California, and one in Oregon). Each trial location consisted of one untreated and one treated plot. Each treated plot received two broadcast foliar applications of 200 g/L SC formulation of Cyflumetofen at 0.196–0.206 kg ai/ha, with a retreatment interval targeting 14 ± 1 days. The actual retreatment interval was 13–15 days, with the exception of one site in California (trial no. R090501) where a 7 day retreatment interval was used. The applications were made using ground equipment in 225–460 L/ha of water spray volumes. Strawberry samples were harvested at a 1-day preharvest interval (PHI). At one trial in California (trial no. R090502), additional treated samples were collected at 0, 3, 7 and 10 days after the last application, in addition to the targeted 1-day PHI, to evaluate residue decline.

Samples of strawberry were analysed for residues of cyflumetofen and metabolites B-1, AB-6 and AB-7 using Method D1003. Acceptable concurrent recovery data for strawberry commodities were obtained for each analyte. The LOQ was 0.01 mg/kg for each analyte. Apparent residues of cyflumetofen, B-1, AB-6 and AB-7 were < LOQ in/on all samples of untreated strawberries.

Table 67 Results of supervised residue trials on Strawberry conducted in the USA.

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP		2	0.2	Min. 468		1					(Interval: 14 days)
USA, 2010 Dane, WI L'Amour R090497	Broad-cast foliar	2	0.2	1 × 245 1 × 250	Berry	1	0.04 0.04 (0.04)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2010 Tift, GA Camarosa R090498	Broad-cast foliar	2	0.2	1 × 237 1 × 225	Berry	1	0.11 0.09 (0.10)	0.031 0.026 (0.0285)	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2009 Washington, OR Albion R090499	Broad-cast foliar	2	0.2	2 × 235	Berry	1	0.23 0.18 (0.205)	0.032 0.029 (0.0305)	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2010 Yates, NY Honeoye R090500	Broad-cast foliar	2	0.2	1 × 455 1 × 460	Berry	1	0.15 0.18 (0.165)	0.028 0.032 (0.030)	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2010 Tulare, CA Diamante R090501	Broad-cast foliar	2	0.2	1 × 306 1 × 300	Berry	1	0.21 0.25 (0.23)	< 0.02 0.104 (0.052)	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2009 Santa Barbara, CA Albion R090502	Broad-cast foliar	2	0.2	2 × 374	Berry	0	0.16 0.21	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
						1	0.11 0.14 (0.125)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375304 2010/7014999
						3	0.16 0.11 (0.135)	0.028 < 0.02 (0.024)	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.15 0.11	0.026 0.027	< 0.01 < 0.01	< 0.01 < 0.01	
						10	0.10 0.09	0.025 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2010 Madera, CA San Andreas R090503	Broad-cast foliar	2	0.2	1 × 284 1 × 279	Berry	1	0.16 0.12 (0.14)	0.025 0.025	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999
USA, 2010 Lake, FL Camarosa R090504	Broad-cast foliar	2	0.2	1 × 371 1 × 370	Berry	1	0.29 0.44 (0.365)	0.037 0.035 (0.036)	< 0.01 < 0.01	< 0.01 < 0.01	M. White, M. Saha 2011 375304 2010/7014999

<sup>a</sup> Metabolite residues expressed in parent equivalents.

#### *Fruiting vegetables, other than Cucurbits (Tomato, eggplant)*

During the growing season 2009–2010, a total of 16 field trials were conducted on tomatoes in the USA (one trial in New York, one in Georgia, two in Florida, one in Kansas, 10 in California and one in Arizona). Two of these trials were decline trials (one trial in NY and the other in CA). At each test location, one untreated and one treated plot were established. The treated plot received two broadcast foliar applications of 200 g/L CS formulation of cyflumetofen at 0.196–0.211 kg ai/ha/application, with a 14-day (+/– one day) re-treatment interval. The applications were made with ground equipment using 115–472 L/ha of water. Commercially mature, tomato samples were collected 3 days after treatment (DAT).

All samples were analysed for residues of cyflumetofen, and its metabolites, B-1, AB-6, and AB-7, using Method D1003. Mean recoveries for all analytes were within the acceptable range of 70–120%. Three individual recoveries were in the range of 65–69%. The LOD and LOQ for all analytes were 0.002 and 0.01 mg/kg, respectively.

Table 68 Results of supervised residue trials on Tomato conducted in the USA.

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP		2	0.2	Min. 468		3					(Interval: 14 days)
USA, 2009 Wayne, NY	Backpack hand- boom	2	0.2	2 × 234	Fruit	0	0.23 0.14	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011
Basket Vee R090531						1	0.04 0.03	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375305 2011/7002631
						3	< 0.01 0.05 (0.03)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.02 0.11 (0.064)	< 0.02 0.04 (0.03)	< 0.01 < 0.01	< 0.01 < 0.01	
						14	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						21	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2009 Tift, GA Mountain Fresh R090532	Backpack hand- boom	2	0.2	1 × 345 1 × 352	Fruit	3	0.03 0.04 (0.035)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Jackson, FL BHN602 R090533	Backpack hand- boom	2	0.2	2 × 347	Fruit	3	0.02 0.06 (0.04)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Seminole, FL Better Boy R090534	Backpack hand- boom	2	2 × 0.2	1 × 284 1 × 280	Fruit	3	0.03 0.02 (0.025)	0.05 0.02 (0.035)	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Barber, KS Early Girl R090535	Backpack hand- boom	2	0.2	1 × 115 1 × 149	Fruit	3	0.07 0.07 (0.07)	0.02 0.02 (0.02)	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Fresno, CA 2401 R090536	Backpack hand- boom	2	0.2	1 × 467 1 × 472	Fruit	3	0.01 0.02 (0.015)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Merced, CA Heinz 2780 R090537	Backpack hand- boom	2	0.2	2 × 466	Fruit	3	0.06 0.07 (0.065)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2010 Kerman Fresno, CA Sun 6366 R090538	Backpack hand- boom	2	0.2	1 × 189 1 × 187	Fruit	3	0.04 0.03 (0.035)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2010 Kerman Fresno, CA APT 410 R090539	Backpack hand- boom	2	0.2	1 × 189 1 × 187	Fruit	3	0.04 0.05 (0.045)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2010 Porterville	Backpack hand- boom	2	0.2	1 × 339 1 × 335	Fruit	0	0.06 0.05	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT d	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
Tular, CA Tom Whopper R090540						1	0.02 0.03	0.02 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375305 2011/7002631
						3	0.02 0.02 (0.02)	0.05 0.02 (0.035)	< 0.01 < 0.01	< 0.01 < 0.01	
						7	0.02 0.01	0.07 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						14	< 0.01 0.01	0.07 0.09	< 0.01 < 0.01	< 0.01 < 0.01	
						21	< 0.01 < 0.01	0.02 0.05	< 0.01 < 0.01	< 0.01 < 0.01	
USA, 2010 Porterville Tular, CA ACE 55 VF (AS) R090541	Backpack hand- boom	2	0.2	1 × 337 1 × 335	Fruit	3	0.05 0.07 (0.06)	< 0.02 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Tular, CA Naoni Cherry (CT) R090542	Tractor mounted sprayer	2	0.2	1 × 363 1 × 383	Fruit	3	0.12 0.12 (0.12)	0.07 0.05 (0.06)	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 King City Monterey, CA 8004 R090543	Backpack hand- boom	2	0.2	1 × 274 1 × 283	Fruit	3	0.03 0.04 (0.035)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 San Ardo Monterey, CA 8004 R090544	Backpack hand- boom	2	0.2	1 × 288 1 × 281	Fruit	3	0.10 0.08 (0.09)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631
USA, 2009 Tehama, CA APT 410 R090545	Backpack hand- boom	2	0.2	2 × 187	Fruit	3	0.16 0.14 (0.15)	< 0.02 < 0.02	0.02 0.01	0.03 0.02	L.E. Crawford 2011 375305 2011/7002631
USA, 2010 Yuma, AZ Montain Fresh R090546	Backpack hand- boom	2	0.2	1 × 230 1 × 229	Fruit	3	0.01 0.01 (0.01)	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L.E. Crawford 2011 375305 2011/7002631

<sup>a</sup> Metabolite residues expressed in parent equivalents.

During the growing season 2004, two trials were conducted on eggplants in greenhouses in Japan to determine the magnitude of the residues of cyflumetofen after treatment with 20% SC formulation of cyflumetofen. The formulation was applied two times at an application rate of 0.400 kg ai/ha and a spray volume of 2000 L/ha. The samples were taken at 1, 3, 7 and 21 days after the last application. The residues of cyflumetofen were determined by LC-UV detection. The LOQ was 0.05 mg/kg.

Table 69 Results of supervised residue trials on Eggplant conducted in Japan

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT d	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
GAP in Japan		2	0.02 kg ai/hL	(1000– 3500)		1					



CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
Japan, 2004 Kochi	n.r.	2	0.40	2 × 2000	Fruit (calyx removed)	1	0.35 0.32 (0.335)	0.06 0.05	< 0.05 < 0.05	< 0.05 < 0.05	n.r. 2004
Ryuma n.r.						3	0.28 0.24	0.06 0.06	< 0.05 < 0.05	< 0.05 < 0.05	Saku 16-P-2-48 OTSA-0203 (2)-FR
(greenhouse)						7	0.06 0.06	0.09 0.09	< 0.05 < 0.05	< 0.05 < 0.05	
						21	< 0.05 < 0.05	0.13 0.12	< 0.05 < 0.05	< 0.05 < 0.05	
Japan, 2004 Miyazaki	n.r.	2	0.40	2 × 2000	Fruit (calyx removed)	1	0.46 0.45 (0.455)	0.43 0.42	< 0.05 < 0.05	< 0.05 < 0.05	n.r. 2004
Kokuyo n.r.						3	0.37 0.36	0.50 0.49	< 0.05 < 0.05	< 0.05 < 0.05	Saku 16-P-2-48 OTSA-0203 (2)-FR
(greenhouse)						7	0.06 0.06	0.58 0.58	< 0.05 < 0.05	< 0.05 < 0.05	
						21	< 0.05 < 0.05	0.26 0.25	< 0.05 < 0.05	< 0.05 < 0.05	

<sup>a</sup> Metabolite residues expressed in parent equivalents.

n.r. = Not reported

#### *Tree nuts (Almond and pecan)*

During the growing season 2009, five trials were conducted on almonds in California in the USA. Each trial consisted of one untreated control and one treated plot except for sites that where both dilute and concentrate applications in which case there were two treated plots and one non-treated plot. For the concentrate plots, the treated plot received two air-blast foliar broadcast applications of 200 g/L SC formulation of cyflumetofen at 0.195–0.202 kg ai/ha, with a 14 day re-treatment interval. The applications were made with ground equipment using 469–656 L/ha of water. For the dilute plots, the treated plot received two airblast foliar broadcast applications of 200 g/L SC formulation of cyflumetofen at 0.197–0.200 kg ai/ha/, with a 13–14 day re-treatment interval. The applications were made with ground equipment using 1219–1875 L/ha of water. Almond samples were harvested 7 days after the last application for all trials except the decline trial (R090478). For the decline trial, samples were collected 0, 1, 3, 7, 14, and 21 days after the last application.

All samples were analysed for residues of cyflumetofen, and its metabolites, B-1, AB-6, and AB-7, using Method D1003. Mean recoveries for all analytes were within the acceptable range of 70–120%. The LOD and LOQ for all analytes were 0.002 and 0.01 mg/kg, respectively.

During the growing season 2009, five trials were conducted on pecans in the USA (two trials in Georgia, one in Louisiana, one in Texas and one in Oklahoma). Each trial consisted of one untreated control and one treated plot except for sites that where both dilute and concentrate applications in which case there were two treated plots and one non-treated plot. For the concentrate plots, the treated plot received two air-blast foliar broadcast applications of 200 g/L SC formulation of cyflumetofen at 0.195–0.204 kg ai/ha, with a 13–14 day re-treatment interval. The applications were made with ground equipment using 540–794 L/ha of water. For the dilute plots, the treated plot received two airblast foliar broadcast applications of 200 g/L SC formulation of Cyflumetofen at 0.198–0.205 kg ai/ha, with a 13–14 day re-treatment interval. The applications were made with ground equipment using 1104–2376 L/ha of water. Pecan samples were harvested 7 or 8 days after the last application.

All samples were analysed for residues of cyflumetofen, and its metabolites, B-1, AB-6, and AB-7, using Method D1003. Mean recoveries for all analytes were within the acceptable range of 70–120%. The LOD and LOQ for all analytes were 0.002 and 0.01 mg/kg, respectively.

Table 70 Results of supervised residue trials on Almonds and Pecans conducted in the USA

CROP Country, Year Location variety Trial No.	Application			Portion analysed	DAT	Residues (mg eq/kg) <sup>a</sup>				Author Report Year Study No. DocID.	
	Method	No.	Rate kg ai/ha			Spray volume L/ha	Parent	B-1	AB-6		AB-7
US GAP		2	0.2	Min. 935		7					(Interval: 14 days)
ALMOND											
USA, 2009 Fresno, CA Non-Pariel R090474	Airblast foliar broadcast	2	0.20	1 × 1848 1 × 1875	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Kern, CA Price R090475	Airblast foliar broadcast	2	0.20	1 × 656 1 × 610	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Tulare, CA Carmel R090476	Airblast foliar broadcast	2	0.20	1 × 1813 1 × 1675	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Glenn, CA Non-Pariel R090477	Airblast foliar broadcast	2	0.20	1 × 469 1 × 473	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Tulare, CA	Airblast foliar	2	0.20	1 × 1219 1 × 1476	Nutmeat	0	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	L. E. Crawford 2011
Monterey R090478	broadcast					1	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	375303 2011/7003996
						3	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	
						14	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
						21	< 0.01 < 0.01	< 0.02 < 0.02	< 0.01 < 0.01	< 0.01 < 0.01	
PECAN											
USA, 2009 Tift, GA Sumner R090469	Airblast foliar broadcast	2	0.20	1 × 2081 1 × 1999	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Irwin, GA Sumner R090470	Airblast foliar broadcast	2	0.20	1 × 667 1 × 648	Nutmeat	7	< 0.01 < 0.01 ( <u>0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Rapides Parish, LA	Airblast foliar broadcast	2	0.20	1 × 1909 1 × 1821	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
Creek R090471				1 × 743 1 × 794	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	
USA, 2009 Jackson, OK	Airblast foliar broadcast	2	0.20	1 × 2376 1 × 2312	Nutmeat	8	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
Cheyenne R090472				1 × 740 1 × 731	Nutmeat	8	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	
USA, 2009 Frio, TX Wichita R090473	Airblast foliar broadcast	2	0.20	1 × 1159 1 × 1104	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	L. E. Crawford 2011 375303 2011/7003996
				1 × 558 1 × 540	Nutmeat	7	< 0.01 < 0.01 ( <u>&lt; 0.01</u> )	< 0.02 < 0.02 < 0.02	< 0.01 < 0.01 < 0.01	< 0.01 < 0.01 < 0.01	

<sup>a</sup> Metabolite residues expressed in parent equivalents.

### Animal feeds

#### Almond hulls

Table 71 Results of supervised residue trials on Almond hulls conducted in the USA

CROP Country, Year Location variety Trial No.	Application				Portion analysed	DAT	Residues (mg/kg) <sup>a</sup>				Author Report Year Study No. DocID.
	Method	No.	Rate kg ai/ha	Spray volume L/ha			Parent	B-1	AB-6	AB-7	
US GAP			0.2	Min. 935		7					(Interval: 14 days)
ALMOND											
USA, 2009 Fresno, CA Non-Pariel R090474	Airblast foliar broadcast	2	0.20	1 × 1848 1 × 1875	Hull	7	2.046 1.696 (1.871)	0.294 0.191 (0.242)	0.227 0.189	0.137 0.116	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Kern, CA Price R090475	Airblast foliar broadcast	2	0.20	1 × 656 1 × 610	Hull	7	0.522 0.545 (0.534)	0.092 0.094 (0.093)	0.074 0.076	0.021 0.022	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Tulare, CA Carmel R090476	Airblast foliar broadcast	2	0.20	1 × 1813 1 × 1675	Hull	7	0.840 0.887 (0.864)	0.106 0.115 (0.111)	0.102 0.096	0.040 0.041	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Glenn, CA Non-Pariel R090477	Airblast foliar broadcast	2	0.20	1 × 469 1 × 473	Hull	7	0.339 0.364 (0.352)	0.028 0.026 (0.027)	0.073 0.073	0.025 0.025	L. E. Crawford 2011 375303 2011/7003996
USA, 2009 Tulare, CA Monterey R090478	Airblast foliar broadcast	2	0.20	1 × 1219 1 × 1476	Hull	0	0.737 0.885	0.045 0.035	0.104 0.098	0.027 0.026	L. E. Crawford 2011 375303 2011/7003996
						1	0.758 0.829	0.045 0.049	0.107 0.097	0.031 0.028	
						3	0.464 0.481	0.040 0.045	0.101 0.098	0.029 0.027	
						7	0.628 0.437 (0.533)	0.028 0.028	0.101 0.103	0.027 0.030	
						14	0.305 0.349	0.033 0.038	0.098 0.105	0.025 0.027	
						21	0.598 0.520 (0.559)	0.125 0.099 (0.112)	0.104 0.102	0.046 0.038	

<sup>a</sup> Metabolite residues expressed in parent equivalents.

## FATE OF RESIDUES IN STORAGE AND PROCESSING

### In processing

The Meeting received information on the fate of residues of Cyflumetofen during hydrolysis and processing of oranges, apples, grapes and tomato reflecting the relevant processing procedures and household preparations.

#### Hydrolysis (Hassing, J, 2013a)

To simulate the degradation of cyflumetofen during pasteurization, hydrolysis of cyflumetofen was tested using benzoyl-ring-U-<sup>14</sup>C labelled and t-butylpheny-ring-U-<sup>14</sup>C labelled cyflumetofen in aqueous buffer solutions at pH 4, 5 and 6 under reflux at 90 °C for 20 min. Due to the rapid degradation of cyflumetofen under the light, each test procedure was done in the dark.

Also for simulating baking, brewing and boiling, the test was conducted similarly but under reflux at 100 °C for 60 minutes. For sterilization, the test was conducted at about 120 °C in an autoclave for 20 minutes.

Aliquots were taken right before starting a test and at the end of the test for LSC-measurements after cooling of the solution. Additionally, aliquots for HPLC-analysis were taken before and at the end of a test. Because of the photolytic and hydrolytic instability of the test item and its metabolites, no work-up for an eventual confirmation by MS-analysis could be performed. Direct MS-analysis was not possible due to the low concentration of the test item especially metabolites in the samples.

The following table demonstrates the results of hydrolysis tests using the radioactive Cyflumetofen labelled at different positions.

Table 72 Effect of hydrolysis on Cyflumetofen simulating pasteurization, baking, brewing and boiling, and sterilization

Simulated process	Hydrolysis condition			% TAR				
	pH	Temp °C	Time, min	Parent	B-1	AB-1	Others	Sum
[Benzoyl-ring-U- <sup>14</sup> C]-cyflumetofen								
Pasteurization	4	90	20	70.9	23.2	4.2	nd	98.3
Baking, brewing and boiling	5	100	60	17.9	58.7	31.7	nd	108.2
Sterilization	6	120	20	Nd	75.3	38.8	1.0	115.1
Simulated process	Hydrolysis condition			% TAR				
	pH	Temp °C	Time, min	Parent	A-2	AB-1	Others	Sum
[t-Butylphenyl-ring-U- <sup>14</sup> C]-cyflumetofen								
Pasteurization	4	90	20	69.3	14.3	Nd	15.4	99.1
Baking, brewing and boiling	5	100	60	5.0	52.9	39.9	4.3	102.2
Sterilization	6	120	20	Nd	44.4	49.1	1.8	95.2

nd =not detected

No loss of radioactivity of the both radio-labelled cyflumetofen occurred during heating in buffers of pH 4, 5 and 9 at 90–120 °C for 20 min or 60 min.

At the pH 4/90 °C condition, benzoyl-ring labelled cyflumetofen was degraded to 70.9% after 20 min. The major degradate was metabolite B-1 (23.2% TAR) followed by AB-1 (4.2% TAR). At the pH 5/100 °C condition, benzoyl-ring labelled cyflumetofen was degraded to 17.9% after 20 min. The major metabolites were B-1 (58.7% TAR) and AB-1 (31.7% TAR). At the pH 6/120 °C condition, benzoyl-ring labelled cyflumetofen was completely degraded to B-1 (75.3% TAR) and AB-1 (38.8% TAR) after 60 min.

At pH 4/90 °C, t-butylphenyl-ring labelled cyflumetofen was degraded to 68.3% TAR. The main metabolite was A-2 (14.3% TAR) with no other metabolite no more than 5% TAR. At pH 5/100 °C, t-butylphenyl-ring labelled cyflumetofen was degraded to 5.0%. The main metabolites were A-2 (52.9% TAR) and AB-1 (39.9% TAR). At pH 6/120 °C, t-butylphenyl-ring labelled cyflumetofen was completely degraded to A-2 (44.4% TAR) and AB-1 (49.1% TAR).

These results indicate that the higher the temperature of heating, the more cyflumetofen degrades and heating splits the Cyflumetofen into t-butylphenyl ring and benzoyl ring.

#### *Oranges (Hattermann & Cowen, 2011c)*

In two trials conducted in Florida and California in the USA during the 2009 growing season, 200 g/L SC formulation of cyflumetofen was applied as two foliar broadcast sprays to oranges at a target rate of 1.0 kg ai/ha (5 times GAP rate), with a 14-day re-treatment interval. The actual rate in the trials was 1.00 to 1.02 kg ai/ha, with a 14–15-day re-treatment interval. One trial had the application made in a concentrated spray volume and one in a dilute spray volume of 703 and 2457 L/ha of water,

respectively, using ground equipment. Orange samples were harvested 7 days after the last application and sent for processing into processed commodities.

Two bulk samples were sent for processing, one treated and one untreated. These bulk orange samples were separately processed (untreated first) using simulated commercial processing procedures for citrus production to generate the required fractions of unwashed fruit, peel, wet pomace, dried pulp, meal, molasses, marmalade, fresh juice and citrus oil with slight variation to the commercial methodology.

The residues of cyflumetofen, B-1, AB-6, and AB-7 in orange processed commodity samples were quantitated using Method D1003. Acceptable concurrent recovery data for orange commodities were obtained for each analyte. The LOQ was 0.01 mg/kg each for parent and the metabolites B-1, AB-6, and AB-7 in/on processed orange samples. The LOD was set at 20% of the LOQ, or 0.002 mg/kg each for parent cyflumetofen and the metabolites B-1, AB-6, and AB-7. The residue levels detected in the treated specimens and processed fractions as well as the calculated processing factors are presented in the following tables. The concentrations of metabolites were expressed in cyflumetofen equivalents.

Table 73 Residues of cyflumetofen and its metabolites in processed commodities of orange

Trial ID County, State of USA /Year	Commodity	Residues, mg eq/kg <sup>a</sup>			Processing factor	
		Parent	B-1	Parent + B-1 <sup>b</sup>	Parent	Parent + B-1
R090554 Seminole, FL / 2009	Orange RAC	0.664	0.028	0.692	–	–
	Wet Pomace	0.169	0.104	0.273	0.255	0.395
	Peel	1.899	0.078	1.977	2.86	2.86
	Dried Pulp	0.388	0.275	0.663	0.584	0.958
	Juice	< 0.01	< 0.02	< 0.03	< 0.02	< 0.04
	Oil	90.72	1.339	92.06	137	133
	Meal	0.283	0.275	0.558	0.426	0.806
	Molasses	< 0.01	0.339	< 0.349	< 0.02	< 0.504
	Marmalade	0.017	0.024	0.041	0.026	0.059
R090555 Kern, CA / 2009	Orange RAC	0.134	0.024	0.158	–	–
	Wet Pomace	0.031	0.024	0.055	0.23	0.348
	Peel	0.398	0.024	0.422	2.97	2.67
	Dried Pulp	0.059	0.061	0.120	0.44	0.759
	Juice	< 0.01	0.024	< 0.034	< 0.08	< 0.22
	Oil	13.65	0.250	13.90	102	88.0
	Meal	0.061	0.061	0.122	0.46	0.77
	Molasses	< 0.01	0.042	< 0.052	< 0.08	< 0.33
	Marmalade	< 0.01	0.024	< 0.034	< 0.08	< 0.22

Trial ID County, State of USA /Year	Commodity	Residues, mg eq/kg <sup>a</sup>	
		AB-6	AB-7
R090554 Seminole, FL / 2009	Orange RAC	0.079	0.022
	Wet Pomace	0.008	0.012
	Peel	0.122	0.077
	Dried Pulp	0.072	0.021
	Juice	0.061	< 0.02
	Oil	2.675	3.160
	Meal	0.077	0.020
	Molasses	0.058	< 0.01
	Marmalade	0.048	0.011
R090555 Kern, CA / 2009	Orange RAC	0.062	0.013
	Wet Pomace	0.005	< 0.02
	Peel	0.076	0.025

Trial ID County, State of USA /Year	Commodity	Residues, mg eq/kg <sup>a</sup>	
		AB-6	AB-7
	Dried Pulp	0.050	< 0.02
	Juice	0.061	< 0.02
	Oil	0.413	0.600
	Meal	0.057	< 0.02
	Molasses	0.055	< 0.02
	Marmalade	0.057	< 0.02

<sup>a</sup> Expressed in cyflumetofen equivalents.

<sup>b</sup> As the concentration of B-1 is sometimes higher than that of the parent, for the calculation of sum and processing factor, the LOQ value was used when the concentration is below the LOQ.

The results from these trials show that after two applications of cyflumetofen at 5× the maximum GAP rate, residues of cyflumetofen or its metabolites (B-1, AB-6 and AB-7) in the treated bulk orange RAC samples ranged from 0.013 to 0.664 mg/kg. Residues of cyflumetofen and its metabolites in processed commodities wet pomace, peel, dried pulp, juice, oil, meal, molasses and marmalade ranged from < 0.01 mg/kg (LOQ) to 2.675 mg/kg in all matrices but oil. The residues in oil from the CA test site ranged from 0.600 mg/kg for metabolite AB-6 to 90.716 mg/kg for cyflumetofen.

A comparison of the total residues of cyflumetofen and all three metabolites combined in the processed RAC with those in each of the processed matrix samples showed that the total residue concentrated in peel and oil only (2.2× to 2.7× and 64× to 123×, respectively). Residue in all the other processed matrices did not concentrate and was diluted to concentrations less than the RAC samples collected before processing.

#### Apples (White, 2011b)

In two trials conducted in NY and PA during the 2009 growing season, 200 g/L SC formulation of cyflumetofen was applied as two airblast foliar sprays to apples at a rate of 0.996–1.027 kg ai/ha (5× the maximum GAP rate), with a 14 day retreatment interval. The first application was made approximately 3 weeks (21 days) prior to the harvest of fruit, when fruit were ripening (growth stage BBCH 79 to BBCH 81). The applications were made in 606–872 L/ha of water using ground equipment. Apple bulk RAC samples were harvested 7 days after the last application. The two bulk apple RAC samples were separately processed using simulated commercial processing procedures into wet pomace and juice. In addition, apple sauce, canned apples, and dried apples were collected for analysis.

The residues of cyflumetofen and metabolites B-1, AB-6 and AB-7 in apple RAC and processed commodity samples were quantitated using Method D1003. Acceptable concurrent recovery data for apple commodities were obtained for each analyte. The LOQ was 0.01 mg/kg for each analyte. Apparent residues of cyflumetofen, B-1, AB-6 and AB-7 were < LOQ in/on all untreated apple RAC and processed commodities.

Table 74 Residues of cyflumetofen and its metabolites in processed commodities of apple.

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>			Processing Factor	
		Parent	B-1	Parent + B-1 <sup>b</sup>	Parent	Parent + B-1
RCN R090509	Apple RAC	0.584	< 0.02	0.604	–	–
Wayne, NY	Wet pomace	0.547	< 0.02	0.567	0.937	0.939
/ 2009	Applesauce	1.700	0.554	2.254	2.91	3.73
	Juice	0.115	< 0.02	0.135	0.197	0.224
	Dried apples	0.482	0.047	0.529	0.825	0.876
	Canned apples	0.102	< 0.02	0.122	0.175	0.202
RCN R090510	Apple RAC	0.451	< 0.02	0.471	–	–
LeHigh, PA	Wet pomace	0.719	< 0.02	0.739	1.59	1.57
/ 2009	Applesauce	1.146	0.342	1.488	2.54	3.16
	Juice	0.121	< 0.02	0.141	0.268	0.299

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>			Processing Factor	
		Parent	B-1	Parent + B-1 <sup>b</sup>	Parent	Parent + B-1
	Dried apples	0.077	< 0.02	0.097	0.17	0.21
	Canned apples	0.016	< 0.02	0.036	0.035	0.076

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>	
		AB-6	AB-7
RCN R090509	Apple RAC	0.013	0.010
Wayne, NY / 2009	Wet pomace	0.022	0.016
	Applesauce	0.055	< 0.01
	Juice	< 0.01	< 0.01
	Dried apples	0.010	< 0.01
	Canned apples	< 0.01	< 0.01
RCN R090510	Apple RAC	0.010	0.010
LeHigh, PA / 2009	Wet pomace	0.015	0.013
	Applesauce	0.039	< 0.01
	Juice	< 0.01	< 0.01
	Dried apples	< 0.01	< 0.01
	Canned apples	< 0.01	< 0.01

<sup>a</sup> Expressed in cyflumetofen equivalents.

<sup>b</sup> As the concentration of B-1 is sometimes higher than that of the parent, for the calculation of sum and processing factor, the LOQ value was used when the concentration is below the LOQ.

The results from these trials show that after two applications of cyflumetofen targeting 2.0 kg as/ha/season, mean parent cyflumetofen residues in the treated apple RAC samples collected 7 days after the last application were 0.58 mg/kg and 0.45 mg/kg. For the individual analytes, expressed in parent equivalents, residues of the metabolites B-1, AB-6, and AB-7 were < 0.02 mg/kg, 0.01 mg/kg, and 0.01 mg/kg, respectively, in the treated apple RAC samples collected 7 days after the last application.

A comparison of the residues in the RAC with those in each processed fraction indicate that parent cyflumetofen residues concentrate in apple sauce (2.5–2.9×, average processing factor 2.7×), but do not concentrate appreciably in any other apple processed commodities, wet pomace, juice, dried apples, or canned apples.

#### *Grapes (Hattermann & Lott, 2011b)*

In four trials conducted in NY, CA and OR during the 2009 growing season, 200 g/L SC formulation of cyflumetofen was applied as two foliar broadcast sprays to grape at a target rate of 1.0 kg ai/ha (5× the maximum GAP rate), with a 14-day re-treatment interval (13-day interval at one CA site). The actual application rates were 1.99 to 2.02 kg ai/ha. The applications were made in 465–2600 L/ha of water using ground equipment. Two trials had product applied in a dilute application (935.4–3741.6 L/ha) and two in a concentrate spray volume (187.1–935.4 L/ha range). Grape samples were harvested 14 days after the last application except for one CA trial which was harvested at 15-day after the final application. One treated and one non-treated bulk sample was processed for each of four trials. Dried grapes, dried grape stems, juice, wine (young), wet pomace, must, yeast and grape stems were collected for analysis. The study was planned to include two trials, each performed on red and white grape varieties. However, an exception to protocol occurred when a white grape variety was used at the King County, CA trial instead of red. The white grapes from this trial were processed using the red grape procedure for wine, such that there were two trials processed with red and two with white grape procedures.

The residues of cyflumetofen, B-1, AB-6, and AB-7 in grape RAC and processed commodity samples were quantitated using Method D1003. Acceptable concurrent recovery data for grape commodities were obtained for each analyte. The LOQ was 0.01 mg/kg each for parent cyflumetofen and the metabolites B-1, AB-6, and AB-7 in/on grape RAC and processed commodity samples. The

LOD was set at 20% of the LOQ, or 0.002 mg/kg each for parent cyflumetofen and the metabolites B-1, AB-6 and AB-7.

Table 75 Residues of cyflumetofen and its metabolites in processed commodities of grapes

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>			Processing Factor	
		Parent	B-1	Parent + B-1 <sup>b</sup>	Parent	Parent + B-1
RCN R090550	Grape RAC	1.71	0.07	1.78	–	–
Yates, NY	Dried grape	7.94	0.33	8.27	4.64	4.65
/ 2009	Dried grape Stem	23.88, 18.96, 17.63	0.56	20.72	11.8	11.6
	Juice	0.11	0.07	0.18	0.064	0.10
	Wine (Young)	< 0.01	0.07	< 0.08	< 0.006	0.04
	Wet Pomace	5.80	0.14	5.94	3.39	3.34
	Must	0.03	0.05	0.08	0.02	0.04
	Yeast	2.34	1.37	3.71	1.37	2.08
	Grape Stems	4.21	0.21	4.42	2.46	2.48
RCN R090551	Grape RAC	0.34	< 0.02	0.36	–	–
Tulare, CA	Dried grape	0.22	0.09	0.31	0.65	0.86
/ 2009	Dried grape Stem	3.07, 1.51, 1.73	0.38	2.48	6.16	6.89
	Juice	0.08	< 0.02	0.10	0.2	0.28
	Wine (Young)	< 0.01	0.02	< 0.03	0.029	0.083
	Wet Pomace	0.37	0.02	0.39	1.1	1.1
	Must	0.14	< 0.02	0.16	0.412	0.44
	Yeast	2.23	0.73	2.96	6.56	8.22
	Grape Stems	0.34	0.05	0.39	1.0	1.1
RCN R090552	Grape RAC	0.57	0.02	0.59	–	–
Kings, CA	Dried grape	0.53	0.09	0.62	0.93	1.1
/ 2009	Dried grape Stem	14.79, 6.63, 6.90	1.01	10.45	16.4	17.7
	Juice	0.14	0.05	0.19	0.25	0.32
	Wine (Young)	0.02	0.07	0.09	0.04	0.2
	Wet Pomace	1.72	0.07	1.79	3.02	3.03
	Must	0.30	0.05	0.35	0.53	0.59
	Yeast	0.31	0.54	0.85	0.54	1.4
	Grape Stems	1.72	0.14	1.86	3.02	3.15
RCN R090553	Grape RAC	1.37	0.12	1.49	–	–
Benton,	Dried grape	2.57	1.11	3.68	1.88	2.47
OR / 2009	Dried grape Stem	13.46, 10.08, 8.33	1.20	11.82	7.75	7.93
	Juice	0.15	0.09	0.24	0.11	0.16
	Wine (Young)	0.06	0.19	0.25	0.04	0.17
	Wet Pomace	5.78	0.14	5.92	4.22	3.97
	Must	0.25	0.07	0.32	0.18	0.22
	Yeast	14.63	0.99	15.62	10.7	10.5
	Grape Stems	3.23	0.33	3.56	2.36	2.39

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>	
		AB-6	AB-7
RCN R090550	Grape RAC	0.03	0.02
Yates, NY	Dried grape	0.11	0.07
/ 2009	Dried grape Stem	0.19	0.12
	Juice	< 0.01	< 0.01
	Wine (Young)	< 0.01	< 0.01
	Wet Pomace	0.04	0.03
	Must	< 0.01	< 0.01
	Yeast	0.01	< 0.01



Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>	
		AB-6	AB-7
	Grape Stems	0.03	0.01
RCN R090551	Grape RAC	0.02	0.02
Tulare, CA	Dried grape	0.02	0.02
/ 2009	Dried grape Stem	0.07	0.10
	Juice	< 0.01	< 0.01
	Wine (Young)	< 0.01	< 0.01
	Wet Pomace	0.02	0.03
	Must	< 0.01	< 0.01
	Yeast	0.07	0.11
	Grape Stems	0.02	0.02
RCN R090552	Grape RAC	< 0.01	< 0.01
Kings, CA	Dried grape	< 0.01	< 0.01
/ 2009	Dried grape Stem	0.17	0.10
	Juice	< 0.01	< 0.01
	Wine (Young)	< 0.01	< 0.01
	Wet Pomace	0.02	0.02
	Must	< 0.01	< 0.01
	Yeast	0.01	0.01
	Grape Stems	0.02	0.02
RCN R090553	Grape RAC	0.15	0.12
Benton,	Dried grape	0.29	0.19
OR / 2009	Dried grape Stem	0.42	0.30
	Juice	< 0.01	< 0.01
	Wine (Young)	< 0.01	< 0.01
	Wet Pomace	0.55	0.42
	Must	< 0.01	< 0.01
	Yeast	0.11	0.10
	Grape Stems	0.10	0.08

<sup>a</sup> Expressed in cyflumetofen equivalents.

<sup>b</sup> As the concentration of B-1 is sometimes higher than that of the parent, for the calculation of sum and processing factor, the LOQ value was used when the concentration is below the LOQ.

The results from these trials show that after two applications of cyflumetofen, targeting 1.0 kg ai/ha per application, residues of cyflumetofen, B-1, AB-6 or AB-7 (expressed in parent equivalents) in the treated grape RAC samples (whole fruit) collected 14–15 days after the last application ranged from < 0.01–1.71 mg/kg for cyflumetofen or its metabolites.

RAC grape samples showed measurable levels of parent cyflumetofen and the three metabolites. A comparison of the residues in the treated grape RAC samples with those in each processed fraction indicated that combined residues of cyflumetofen and its metabolites concentrated in dried grape at 2.37 and 4.62× at the OR and NY test sites, respectively, and in dried grape stem (6.63–17.6×), wet pomace (1.10–3.91×), yeast (1.43–8.99×), and grape stems (1.08–3.11×) at all four test sites. The compound did not concentrate in any of the other processed grape matrices and was actually diluted in these matrices (i.e. concentration factors less than or equal to 1×).

#### *Tomato (Crawford, 2011b)*

In two trials conducted in NY and CA during the 2009 growing season, 200 g/L SC formulation of cyflumetofen was applied as two foliar broadcast sprays to tomato at a rate of 1.00–1.01 kg ai/ha, with a 14-day retreatment interval, resulting in a total exaggerated rate (5×) of 2.01–2.02 kg ai/ha. The applications were made in 187–360 L/ha using ground equipment, and an adjuvant was added to the spray mixture for all applications. Tomato samples were harvested 3 days after the last application. A total of two separate treated bulk samples were processed for a total of two processing tests. The treated bulk tomato RAC samples were separately processed using simulated commercial processing procedures into paste and puree. In addition, canned tomatoes, peeled tomatoes, tomato juice, tomato peels, wash water, washed tomatoes, and wet pomace were collected for analysis.

The residues of cyflumetofen and metabolites B-1, AB-6, and AB-7 in tomato processed commodity samples were quantitated using Method D1003. Acceptable concurrent recovery data for tomato commodities were obtained for each analyte. The LOQ was 0.01 mg/kg each for parent and the metabolites B-1, AB-6, and AB-7 in/on tomato processed commodity samples. The LOD was set at 20% of the LOQ, or 0.002 mg/kg each for parent cyflumetofen and the metabolites B-1, AB-6, and AB-7.

Table 76 Residues of cyflumetofen and its metabolites in processed commodities of tomato

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>			Processing Factor	
		Parent	B-1	Parent + B-1 <sup>b</sup>	Parent	Parent + B-1
RCN R090548	Tomato RAC	0.24	0.05	0.29	–	–
Wayne, NY / 2009	Washed Tomatoes	0.26	0.05	0.31	1.1	1.1
	Peeled Tomatoes	< 0.01	< 0.02	< 0.03	< 0.04	< 0.1
	Canned Tomatoes	< 0.01	0.02	< 0.03	< 0.04	< 0.1
	Puree	0.21	0.02	0.23	0.88	0.79
	Paste	0.09	0.54	0.63	0.4	2.2
	Wet Pomace	0.32	0.08	0.40	1.3	1.4
	Tomato Peels	4.05	0.08	4.13	16.9	14.2
	Wash Water	0.69	0.09	0.78	2.9	2.7
	Tomato Juice	0.04	0.07	0.11	0.2	0.38
	RCN R090549	Tomato RAC	0.17	0.02	0.19	–
Kings, CA / 2009	Washed Tomatoes	0.11	0.02	0.13	0.65	0.68
	Peeled Tomatoes	< 0.01	< 0.02	< 0.03	< 0.06	< 0.2
	Canned Tomatoes	0.03	0.02	0.05	0.2	0.3
	Puree	0.05	< 0.02	0.07	0.3	0.4
	Paste	0.04	0.19	0.23	0.2	1.2
	Wet Pomace	0.93	0.02	0.95	5.5	5.0
	Tomato Peels	1.52	0.02	1.54	8.94	8.10
	Wash Water	0.02	< 0.02	0.04	0.1	0.2
	Tomato Juice	< 0.01	0.02	< 0.03	< 0.06	< 0.2

Trial ID County, State of USA /Year	Commodity	Residues (mg/kg) <sup>a</sup>	
		AB-6	AB-7
RCN R090548	Tomato RAC	0.05	0.08
Wayne, NY / 2009	Washed Tomatoes	0.06	0.11
	Peeled Tomatoes	< 0.01	< 0.01
	Canned Tomatoes	< 0.01	< 0.01
	Puree	0.05	0.08
	Paste	0.07	0.17
	Wet Pomace	0.06	0.11
	Tomato Peels	0.73	1.46
	Wash Water	0.01	0.01
	Tomato Juice	< 0.01	0.01
	RCN R090549	Tomato RAC	< 0.01
Kings, CA / 2009	Washed Tomatoes	< 0.01	0.01
	Peeled Tomatoes	< 0.01	< 0.01
	Canned Tomatoes	< 0.01	< 0.01
	Puree	< 0.01	< 0.01
	Paste	< 0.01	< 0.01
	Wet Pomace	0.03	0.05
	Tomato Peels	0.06	0.10
	Wash Water	< 0.01	< 0.01
	Tomato Juice	< 0.01	< 0.01

<sup>a</sup> Expressed in cyflumetofen equivalents.

<sup>b</sup> As the concentration of B-1 is sometimes higher than that of the parent, for the calculation of sum and processing factor, the LOQ value was used when the concentration is below the LOQ.

The results from these trials show that after two applications of cyflumetofen targeting 1.0 kg ai/ha to each field trial to obtain 5× the maximum GAP per-season rate of 0.400 kg ai/ha, residues of cyflumetofen (metabolites B-1, AB-6 and AB-7 expressed in parent equivalents) in the treated tomato RAC samples (whole fruit) sampled 3 days PHI were < LOQ to 0.24 mg/kg. For the processed commodity samples, parent residues in the treated fractions from fruit sampled 3 days PHI were < LOQ to 4.05 mg/kg; residues of the metabolites B-1, AB-6, and AB-7 ranged from < LOQ to 1.46 mg/kg.

A comparison of the residues in the RAC tomato with those in each processed fraction indicated that combined residues of cyflumetofen concentrated in wash water (1.9×) at the New York site only and in paste (1.2 to 2.1×), wet pomace (1.3 to 4.2×) and tomato peel (8.1 to 15.0×) at both test sites. The compound did not concentrate in any other processed tomato fractions.

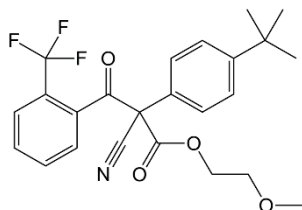
## RESIDUES IN ANIMAL COMMODITIES

### Farm animal feeding studies

No information on farm animal feeding studies were received by the current Meeting.

## APPRAISAL

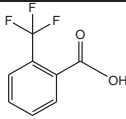
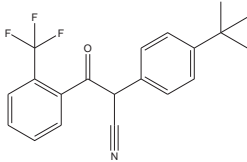
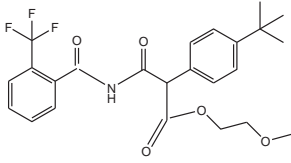
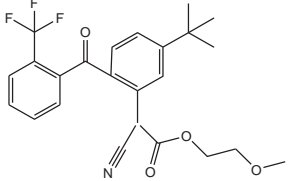
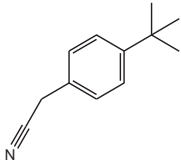
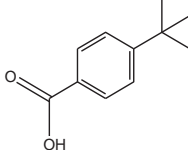
Cyflumetofen (consisting of RS isomers) is a bridged diphenyl acaricide (miticide) for control of *Tetranychus* sp. and influences the mitochondrial electron transport chain by inhibiting the complex II substance in the cell. It has been registered in a number of countries.



The Meeting received information on physical and chemical properties, animal and plant metabolism, environmental fate, analytical methods, storage stability, use patterns, supervised trials, and processing. Cyflumetofen was scheduled by the Forty-fifth Session of the CCPR for review by the 2014 JMPR for the first time.

In this Appraisal, the following abbreviated names were used for referred metabolites.

Code (MW)	IUPAC name	Structure	Metabolite or degradate found in
Cyflumetofen (447.45)	2-methoxyethyl ( <i>RS</i> )-2-(4- <i>tert</i> -butylphenyl)-2-cyano-3-oxo-3-( $\alpha,\alpha,\alpha$ -trifluoro- <i>o</i> -tolyl)propionate		Goat Satsuma mandarin Apple Eggplant
B-1 (190.12)	2-(trifluoromethyl)benzoic acid		Rat

			Goat Satsuma mandarin Apple Eggplant Soil
AB-1 (345.36) Syn: M9210I003	(RS)-2-(4-tert-butylphenyl)-3-oxo-3-[2-(trifluoromethyl)phenyl]propanenitrile		Rat Goat Hydrolysis in buffer Hydrolysis simulating processing Soil
AB-6 (465.45)	2-methoxyethyl-2-((R,S)-4-tert-butylphenyl)-3-oxo-3-({[2-(trifluoromethyl)phenyl]carbonyl}amino)propanoate		Satsuma mandarin Apple Eggplant
AB-7 (447.45)	2-methoxyethyl-((R,S)-4-tert-butyl-2-{{[2-(trifluoromethyl)phenyl]carbonyl}phenyl}(cyano)acetate		Satsuma mandarin Apple Eggplant
A-2 (173.26) Syn: M9210I001	4-tert-butylphenylacetonitrile		Goat Soil
A-12 (178.23) Syn: M9210I002	4-tert-butylbenzoic acid		Goat Soil

### ***Animal metabolism***

The Meeting received information on the fate of orally-dosed cyflumetofen in lactating goats.

In metabolism studies, total radioactive residues are expressed in mg eq./kg cyflumetofen equivalents unless otherwise stated.

#### ***Metabolism of cyflumetofen in rat***

Metabolism studies on laboratory animals including rats were reviewed in the framework of toxicological evaluation by the current JMPR.

In a rat metabolism study, highest residues were found in the liver followed by kidney regardless of sex, dose and label position and time point of measurement. Cyflumetofen was extensively metabolized. B-1, 2-trifluoromethylbenzoic acid, was the major metabolite (occurring up to 28% of the applied dose). The predominant metabolic pathway for cyflumetofen involves cleavage of the tert-butylphenyl and trifluorotolyl moieties. Major reactions on the tert-butylphenyl ring are cleavage of the methoxyethyl group, hydroxylation at the butyl group, decarboxylation and glucuronidation at the butyl group. Major reactions on the trifluorotolyl-ring are glutathione conjugation at the carboxyl group and further changes of the glutathione group to mercapturic acid or thiolactic acid. In addition, hydroxylation and oxidation reactions at the butyl group and cleavage of the carboxylic ester moiety are observed on the parent molecule.

#### *Metabolism of cyflumetofen in lactating goat*

Two lactating goats were orally administered [benzoyl-ring- $U-^{14}C$ ]-cyflumetofen (hereafter abbreviated as benzoyl-label) for 12 consecutive days at 0.43 or 0.48 mg/kg bw (13 or 12 ppm in feed). Other two goats were orally administered [t-butylphenyl-ring- $U-^{14}C$ ]-cyflumetofen (hereafter abbreviated as butylphenyl-label) for 10 consecutive days at 0.27 or 0.30 mg/kg bw (15 or 12 ppm in feed). The goats were sacrificed 18–24 hours after the last dose.

After radio-labelled cyflumetofen was administered orally to lactating goats, it was excreted in faeces and urine (total > 78% excreted during the testing periods) and only a small portion accounting for < 0.3% of the administered radioactivity (AR) remained in body tissues/organs. Radioactive residues were the highest in liver (0.29–0.40 mg eq./kg) followed by kidney (0.17–0.19 mg eq./kg) but low in fat (0.028–0.033 mg eq./kg) and muscle (0.009–0.020 mg eq./kg). Milk of each day contained  $\leq 0.02\%$  AR and milk collected throughout the study period contained, in total, 0.008–0.19 mg eq./kg (0.03–0.14% AR).

The parent compound, cyflumetofen, was found only in fat but at low concentration of 0.003 mg/kg accounting for 20–21% of the total radioactive residues (TRR). The predominant metabolite in all tissues/organs and milk was B-1 at around 0.1 mg eq./kg in liver (32% TRR) and kidney (54% TRR), and lower than 0.01 mg eq./kg in muscle (46–51% TRR), fat (21–40% TRR) and milk (4.5% TRR).

The metabolism of cyflumetofen in goat involves extensive hydrolysis of formic acid ester and trifluoromethylbenzoyl moiety, decarboxylation, conjugation, hydroxylation and oxidation. In principle, excretion and distribution of cyflumetofen and its metabolism in goat are similar to those in rats with some differences in metabolites identified.

#### *Plant metabolism*

##### *Satsuma mandarin*

When Satsuma mandarin trees (grown outdoor) were sprayed with benzoyl- or butylphenyl-label cyflumetofen at a rate approximating 0.60 kg ai/ha (3× GAP rate of the USA), and Satsuma mandarin fruit samples were collected 1, 7 and 30 days after the treatment, the majority of the radioactivity was recovered from the acetonitrile surface rinse of fruits: 95–96% TRR on 1DAT; 91–93% TRR on 7DAT; and 88–89% on 30 DAT.

The predominant radioactive residue on/in mandarin fruits was cyflumetofen at 0.52–0.55 mg/kg (88–90% TRR) on 1DAT, decreasing to 0.33–0.37 mg/kg (79–83% TRR) on 7 DAT and further to 0.25–0.31 mg/kg (44–54% TRR) on 30 DAT.

Other than the parent, metabolites AB-6, AB-7, A-12 and B-1 were formed. B-1 increased over time from 0.028 mg eq./kg (4.7% TRR) on 1 DAT to 0.064 mg eq./kg (11% TRR) on 30 DAT. None of AB-6, AB-7 and A-12 exceeded 10% TRR. A number of unknown peaks were observed but none of them exceeded 10% of TRR.

The radioactive residues on/in leaves showed similar profile as those on/in fruits with the majority (>87% TRR) of radioactive residues found in the surface rinse, although the TRR was >50

times that on/in fruits. A majority of radioactive residues remaining in leaves were extracted. Cyflumetofen was also the predominant identified fraction in leaves: >73% TRR in rinse and extract together. Four metabolites were identified but none exceeded 10% TRR.

#### *Apple*

When an apple tree (grown outdoor) was sprayed with benzoyl- or butylphenyl-label cyflumetofen at a rate approximating 0.60 kg ai/ha (3X GAP rate of the USA), and apple fruit samples were collected 1, 7 and 30 days after the treatment, the majority of the radioactivity was recovered from the acetonitrile surface rinse of fruits: 95–96% TRR on 1 DAT, 82–89% TRR on 7 DAT; and 67–71% TRR on 30 DAT.

The predominant radioactive residue on/in apple fruits was cyflumetofen at 0.061–0.066 mg/kg (58–61% TRR) in rinse while pulp extract contained too little radioactivity for further analysis on 1DAT, 0.061–0.14 mg/kg (78–84% TRR) on 7 DAT and decreased to 0.037–0.042 mg/kg (53–65% TRR) on 30 DAT.

There were minor metabolites identified. However, none of them exceeded 5% TRR.

The radioactive residues on/in leaves showed similar profile as those on/in fruits with the majority (>72% TRR) of radioactive residues found in the surface rinse, although the TRR was >50 times that on/in fruits. Cyflumetofen was also the predominant identified fraction in leaves: >44% TRR in rinse and extract together. No metabolites exceeded 10% TRR.

#### *Eggplant*

When eggplants (grown outdoor) were sprayed with benzoyl- or butylphenyl-label cyflumetofen at a rate approximating 0.60 kg ai/ha (3X GAP rate of the USA), and eggplant samples were collected 1, 7 and 14 days after the treatment, the majority of the radioactivity was also recovered from the acetonitrile surface rinse of fruits: 87–92% TRR on 1 DAT, 79–86% TRR on 7 DAT; and 56–81% TRR on 30 DAT.

The predominant radioactive residue on/in eggplant fruits was also cyflumetofen accounting for 0.31–0.44 mg/kg (91–95% TRR) on 1 DAT, decreased to 0.25–0.39 mg/kg (67–71% TRR) on 7 DAT and then to 0.18–0.20 mg/kg (42–62% TRR) on 14 DAT.

Metabolites B-1, AB-6 and AB-7 were identified from the fruit rinse and/or extracts. B-1 was found at 0.059 mg eq./kg (11% TRR) on 7 DAT and at 0.061 mg eq./kg (15% TRR) on 14 DAT on/in fruits. AB-6 and AB-7 did not exceed 10% TRR.

The tentatively identified U1/U2 (likely to be acid labile conjugates of B-1) and U4 were also found in 7 DAT and 14 DAT fruits but not in 1 DAT fruits. U1 was present at 0.067 mg eq./kg (16% TRR) on 14 DAT but <10% TRR on 1 DAT and 7 DAT. No other metabolites exceeded 10% TRR. U1 and U2 were found only in the fruit extracts but not in rinse, indicating that they were formed in the fruits.

The radioactive residues on/in leaves showed similar profile as those on/in fruits with the majority (>69% TRR) of radioactive residues found in the surface rinse, although the TRR was >50 times that on/in fruits. Cyflumetofen was also the predominant identified fraction in leaves: >47% TRR in rinse and extract together. No metabolites exceeded 10% TRR.

#### *Summary of plant metabolism*

The metabolism of cyflumetofen in these crops involves hydrolysis, acyl migration, oxidation and conjugation (U1 and U2 are the conjugate of B-1).

The plant metabolism studies on Satsuma mandarin, apple and eggplant showed similar pattern. Applied radioactively was mostly recovered from surface rinse with decreasing trend over time. Fruit flesh contained far less radioactivity. The predominant radioactive residue was cyflumetofen accounting for >42% TRR at all time points. The next important radioactive residue was

B-1 but at the maximum 15% TRR (around 0.06 mg eq./kg). B-1 may be present in fruits in conjugated forms.

### ***Environmental fate***

#### *Aerobic soil metabolism*

The studies on aerobic soil degradation of cyflumetofen indicate that cyflumetofen sprayed on soil was rapidly and completely degraded with DT50 around 1.8–4.3 days in various soils at 20–25°C. Additionally, aerobic soil degradation of B-1 and AB-1 was studied resulting in DT50 of 6–36 days and 0.07–0.11 days respectively at 20°C.

#### *Photolysis on soil surface*

Photolysis of cyflumetofen on soil was found to be insignificant as the degradation rates of cyflumetofen with and without light irradiation were not significantly different.

#### *Hydrolysis in aquatic system*

Cyflumetofen is susceptible to hydrolysis and was hydrolysed faster at higher pH in aqueous buffer solutions at 25°C. DT50 was calculated using first order kinetics to be 7.7 days at pH 4, 6.0 days at pH 5, 9.8 h at pH 7 and 10.3 min at pH 9.

At slightly acidic to neutral pH, a number of degradates exceeded 10% of the applied dose: A-1 (max. 14% at 8 h), A-2 (max. 44% at 720 h), A-18 (max. 36% at 120 h), B-1 (max. 53% at 48 h) and AB-1 (max. 45%, at 120 h).

#### *Residues in succeeding crops*

A confined rotational crop study was conducted to examine the nature and level of residues of cyflumetofen in three succeeding crops (white radish, lettuce and wheat) using benzoyl- and butylphenyl-label cyflumetofen. A single application of radio-labelled cyflumetofen was made on bare soil in plastic containers at a rate of 0.40 kg ai/ha (2x GAP rate of the USA). After plant back interval (PBI) of 30, 120 and 365 days, lettuce, white radish and spring wheat were sown into the treated soil.

Following the application of cyflumetofen to soil, uptake of radioactivity from soil into rotational crops was observed, in particular, with benzoyl-label (up to 1.25 mg eq./kg in wheat chaff of 30 PBI but much lower in edible portions with the maximum of 0.17 mg eq./kg in wheat grain of 30 PBI). With butylphenyl-label, uptake of radioactivity into rotational crops were lower (maximum of 0.11 mg eq./kg in wheat straw+chaff of 120 PBI). Uptake was, in general, less with longer plant back interval.

The only major radioactive residue identified from benzoyl-label treatment was trifluoroacetic acid in all crops (39–100% TRR). B-1, specific to this label, was detected as a minor component.

With the butylphenyl-label, a series of label-specific metabolites were observed in the extracts but all of them were less than 0.01 mg eq./kg.

No parent compound was detected from any of the crop extracts.

No significant residues of cyflumetofen, other than trifluoroacetic acid, were expected to be found in rotational crops on the basis of aerobic soil degradation studies and confined succeeding crop study.

### ***Methods of analysis***

Analytical methods for determination of residues of cyflumetofen and its metabolites B-1, AB-6 and AB-7 were developed for a wide range of matrices of plant and animal origin.

In general, the method for data generation and enforcement for plant matrices employ extraction by shaking with acetonitrile and then a mixture of acetonitrile:water (75:25, v/v) or only with acetonitrile:water, clean-up by partitioning with a mixture of ethyl acetate:cyclohexane (75:25, v/v) and determination of the analytes using LC-MS/MS.

The method for plant matrices was validated for cyflumetofen, B-1, AB-6 and AB-7 resulting in acceptable recoveries and relative standard deviations (RSDs) with the LOQ of 0.01 mg/kg for various plant matrices.

The analytical method developed for cyflumetofen and B-1 in animal matrices was similar to the one for plant matrices but employs acetonitrile:water (50:50, v/v) for extraction instead of acetonitrile:water (75:25, v/v) and different clean-up procedure. This method was validated for cyflumetofen and B-1 resulting in acceptable recoveries and RSDs with the LOQ of 0.01 mg/kg for bovine liver, and meat and 0.001 mg/kg for bovine milk. The mean recoveries for cyflumetofen in poultry eggs were low at 65–67% (RSD, 3–6%) while those for B-1 were acceptable between 86–95%.

A number of scientific papers report the validation of the QuEChERS multi-residue method with GC-MS/MS for cyflumetofen in various plant commodities with the LOQ around 0.01 mg/kg.

#### ***Stability of pesticide residues in stored analytical samples***

The stability of cyflumetofen, B-1, AB-6 and AB-7 in homogenates of almond, apple (fruit, juice) kidney bean, lettuce, orange (fruit, juice, oil), radish and wheat grain at -20 to -10 °C was tested at a spike level (each analyte separately spiked) of 0.1 mg/kg for 743–910 days (24–30 months).

Cyflumetofen was stable when stored frozen for 25 months, the longest storage time tested, in almond, apple fruit, apple juice, orange fruit (24 months), orange juice, orange oil and wheat grain. However, it was stable only up to 9 months in kidney bean and lettuce, and 3 months in radish root.

B-1 was stable when stored frozen for 30 months, the longest storage time tested, in almond, kidney bean, lettuce, orange fruit, orange juice and wheat grain. It was stable up to 22 months in apple fruit, apple juice and radish root, and 6 months in orange oil.

AB-6 was stable when stored frozen for 28 months, the longest storage time tested, in almond, apple fruit, kidney bean, orange juice, orange oil, radish root, and wheat grain. It was stable up to 21 months in apple juice, lettuce and orange fruit.

AB-7 was stable when stored frozen for 26 months, the longest storage time tested, in almond, apple fruit, apple juice, kidney bean, orange juice, and orange oil. It is stable up to 19 months in orange fruit and wheat grain, 12 months in lettuce, but one month in radish root.

#### ***Definition of the residue***

In goat metabolism studies, radioactive residues were highest in liver followed by kidney. Far less radioactive residues were found in fat, muscle and milk.

When benzoyl-label cyflumetofen was administered, metabolite B-1 (2-trifluoromethylbenzoic acid) was the predominant residue in all edible tissues/organs (21–54% TRR) and milk (4.5% TRR). B-1 was also found in rat metabolism and is considered to be no more toxic than the parent and, therefore, toxicologically covered by the ADI for parent compound. Cyflumetofen was detected only in fat at 0.003 mg/kg (20% TRR). No other metabolites were detected above 10% TRR.

When butylphenyl-label cyflumetofen was administered, cyflumetofen was not detected in any of edible tissues/organs or in milk. None of all the identified metabolites exceeded 0.01 mg eq./kg and 10% TRR.

There is a validated LC-MS/MS method for cyflumetofen and B-1 in bovine matrices. The Meeting considered that both cyflumetofen and B-1 were suitable residues for enforcement of MRLs and for estimating dietary intake.



LogP<sub>ow</sub> of cyflumetofen is 4.3 at 25 °C and cyflumetofen was found only in the fat at < 0.01 mg/kg in the goat metabolism study. B-1 is a carboxylic acid and was present at 0.13 mg eq./kg in liver and 0.10 mg eq./kg in kidney but < 0.01 mg eq./kg in fat. The Meeting considered that, over all, residues (cyflumetofen and B-1) were not fat soluble.

In the plant metabolism studies on Satsuma mandarin, apple and eggplant, cyflumetofen was the predominant residue and accounted for > 42% TRR in the fruits of these crops at any time points. The Meeting considered that for enforcement of MRLs, cyflumetofen was a suitable residue.

The next important radioactive residue was B-1 but at the maximum 15% TRR (around 0.06 mg eq./kg). In the supervised residue trials, the concentrations of B-1 were mostly 1/10–1/2 of those of cyflumetofen. Conjugated forms of B-1 (U1 and U2) were found in the eggplant metabolism at slightly higher concentrations (in total) than B-1 itself but they were not found in the studies on Satsuma mandarin or apple. A validated LC-MS/MS method is available and used in the trials for quantification of cyflumetofen and B-1 but it determines the free form of B-1.

The Meeting considered that, for calculating dietary intake from commodities of plant origin, cyflumetofen and B-1 were suitable residues.

Based on the above, the Meeting recommended the following residue definition for plant and animal commodities:

Definition of the residue for plant commodities (for compliance with the MRL): *Cyflumetofen*.

Definition of the residue for plant commodities (for estimation of dietary intake): *Sum of cyflumetofen and 2-trifluoromethylbenzoic acid, expressed as cyflumetofen*.

Definition of the residue for animal commodities (for compliance with the MRL and estimation of dietary intake): *Sum of cyflumetofen and 2-trifluoromethylbenzoic acid, expressed as cyflumetofen*.

*Residue is not fat-soluble.*

### ***Results of supervised residue trials on crops***

The Meeting received supervised trial data for cyflumetofen on citrus fruits, pome fruits, grapes, strawberry, tomato, eggplant and tree nuts conducted outdoor (except trials on eggplant in Japan) using foliar spray of 20% SC formulation of cyflumetofen (except in a trial on apple in Italy).

Residues were expressed in cyflumetofen equivalents. The analytical results for B-1 were converted to cyflumetofen equivalents by multiplying the analytical results for B-1 with a factor of 2.35 based on the molecular weights of cyflumetofen (447.45) and B-1 (190.12).

For the estimation of a sum of cyflumetofen and B-1, where B-1 was below the LOQ, it was regarded as 0.02 mg eq./kg (LOQ of 0.01 mg/kg for B-1 is converted to 0.02 mg eq./kg) as B-1 was sometimes present at concentrations comparable to those of parent in trials.

Although in a number of trials conducted in the USA the water volume was smaller than the minimum water volume on the label, the Meeting agreed to use the results of these trials in estimating maximum residue levels as the purpose of the minimum requirement was for better efficacy and not for safety, and residues from lower water volume trials were not always lower than those from higher water volume trials.

#### *Citrus fruits*

A total of 23 supervised trials were conducted on orange (12), grapefruit (6) and lemon (5) in the USA in 2009 and 2010. GAP in the USA for citrus fruits allows 2 applications at a maximum rate of 0.2 kg ai/ha (at least 935 L/ha) with a PHI of 7 days.

Residues of cyflumetofen from 11 independent orange trials matching GAP for citrus fruits in the USA were: 0.01, 0.01, 0.04, 0.06, 0.06, 0.08, 0.09, 0.10, 0.10, 0.11 and 0.12 mg/kg (median 0.08 mg/kg).

The information on four trials conducted on orange in Brazil in 2007 was also provided but they did not match the GAP in Brazil (one application at a spray concentration of 0.08 kg ai/hL).

Residues of cyflumetofen from six grapefruit trials matching GAP for citrus fruits in the USA were: < 0.01, 0.02, 0.04, 0.04 and 0.07 mg/kg (median 0.04 mg/kg).

Residues of cyflumetofen from five lemon trials matching GAP for citrus fruits in the USA were: < 0.01, 0.02, 0.02, 0.08 and 0.14 mg/kg (median 0.02 mg/kg).

As the GAP in the USA is established for the group of citrus fruits, the median values from the trials conducted in the USA on these three commodities were not different more than 5-fold, and Kruskal-Wallis test indicated that the residues of orange, grapefruits and lemon were not statistically different, the Meeting decided to estimate a group maximum residue level. The residues of orange, grapefruits and lemon were combined for estimating a maximum residue level for estimating a maximum residue level for citrus fruits (n=22): < 0.01, < 0.01, 0.01, 0.01, 0.02, 0.02, 0.02, 0.04, 0.04, 0.04, 0.04, 0.06, 0.06, 0.07, 0.08, 0.08, 0.09, 0.10, 0.10, 0.11, 0.12 and 0.14 mg/kg.

The Meeting estimated a maximum residue level of 0.3 mg/kg for citrus fruit.

For the estimation of STMR, the sum of cyflumetofen and B-1 was calculated: < 0.03, < 0.03, 0.03, 0.03, 0.04, 0.04, 0.04, 0.06, 0.06, 0.06, 0.06, 0.08, 0.08, 0.09, 0.10, 0.10, 0.11, 0.12, 0.12, 0.13, 0.14 and 0.16 and mg eq./kg.

The Meeting estimated an STMR of 0.07 mg/kg expressed as cyflumetofen.

#### *Pome fruits*

A total of 17 supervised trials were conducted in the USA in 2009 and 2010 on apples (12) and pear (5). One trial was conducted in Italy in 2006 on apples. The GAP in the USA for pome fruits allows 2 applications at a maximum rate of 0.2 kg ai/ha (at least 935 L/ha) with a PHI of 7 days.

The trial from Italy was provided but did not match any GAP and was not considered.

Residues from 12 apple trials matching US GAP were: 0.01, 0.04, 0.06, 0.07, 0.09, 0.10, 0.13, 0.14, 0.16, 0.18, 0.19 and 0.24 mg/kg (median 0.115 mg/kg).

Residues from five pear trials matching US GAP were: 0.06, 0.08, 0.12, 0.12 and 0.20 mg/kg (median 0.12 mg/kg).

As the GAP in the USA is established for the group of pome fruits, and the median values from the trials conducted in the USA on the two commodities did not differ by more than 5-fold, the Meeting decided to estimate a group maximum residue level.

Man-Whitney test indicated that the residues from apple trials and those from pear trials were not statistically different. The Meeting decided to combine the data for estimating a maximum residue level. Residues were (n=17): 0.01, 0.04, 0.06, 0.06, 0.07, 0.08, 0.09, 0.10, 0.12, 0.12, 0.13, 0.14, 0.16, 0.18, 0.19, 0.20 and 0.24 mg/kg.

The Meeting estimated a maximum residue level of 0.4 mg/kg for pome fruits.

The sum of cyflumetofen and B-1 was calculated: 0.03, 0.06, 0.08, 0.08, 0.09, 0.10, 0.11, 0.12, 0.14, 0.14, 0.15, 0.16, 0.18, 0.20, 0.21, 0.22 and 0.26 mg eq./kg.

The Meeting estimated an STMR of 0.14 mg/kg expressed as cyflumetofen.

*Berries and Other Small Fruits**Grapes*

A total of 12 supervised trials were conducted on grapes in the USA in 2009. The GAP in the USA for grapes allows 2 applications at a maximum rate of 0.2 kg ai/ha (at least 468 L/ha) with a PHI of 14 days.

Residues from 11 independent trials matching US GAP were: 0.02, 0.09, 0.10, 0.12, 0.15, 0.16, 0.19, 0.22, 0.22, 0.27 and 0.42 mg/kg.

The Meeting estimated a maximum residue level of 0.6 mg/kg for grapes.

The sum of cyflumetofen and B-1 was calculated: 0.04, 0.12, 0.12, 0.16, 0.20, 0.22, 0.23, 0.25, 0.25, 0.33 and 0.46 mg eq./kg.

The Meeting estimated an STMR of 0.22 mg/kg expressed as cyflumetofen.

*Strawberry*

A total of eight supervised trials were conducted on strawberry in the USA in 2009 and 2010. The GAP in the USA for strawberry allows 2 applications at a maximum rate of 0.2 kg ai/ha (at least 468 L/ha) with a PHI of 1 day.

Residues from eight trials matching US GAP were: 0.04, 0.10, 0.14, 0.14, 0.16, 0.20, 0.23 and 0.36 mg/kg.

The Meeting estimated a maximum residue level of 0.6 mg/kg for strawberry.

The sum of cyflumetofen and B-1 was calculated: 0.06, 0.13, 0.16, 0.16, 0.20, 0.24, 0.28 and 0.40 mg eq./kg.

The Meeting estimated an STMR of 0.18 mg/kg expressed as cyflumetofen.

*Fruiting Vegetables, Other Than Cucurbits**Tomato*

A total of 16 supervised trials were conducted on tomato in the USA in 2009 and 2010. The GAP in the USA for tomato allows 2 applications at a maximum rate of 0.2 kg ai/ha (at least 468 L/ha) with a PHI of 3 days.

Residues from 14 independent trials matching US GAP were: 0.01, 0.02, 0.02, 0.04, 0.04, 0.04, 0.04, 0.06, 0.06, 0.06, 0.07, 0.09, 0.12 and 0.15 mg/kg.

The Meeting estimated a maximum residue level of 0.3 mg/kg for tomato.

The sum of cyflumetofen and B-1 was calculated: 0.03, 0.04, 0.06, 0.06, 0.06, 0.06, 0.06, 0.08, 0.08, 0.09, 0.10, 0.11, 0.17 and 0.18 mg eq./kg.

The Meeting estimated an STMR of 0.07 mg/kg expressed as cyflumetofen.

*Eggplant*

Two supervised trials were conducted on eggplant, green house grown, in Japan in 2004 with the analysis of cyflumetofen and the conjugates of B-1. The current GAP of Japan allows 2 applications at a spray concentration of 0.02 kg ai/hL with a PHI of 1 day. A spray volume of 1000–3500 L can be sprayed per ha. In the trials, spray concentration was 0.02 kg ai/hL and spray volume was 1996 L/ha, matching Japanese GAP. Residues from these trials were: 0.34 and 0.46 mg/kg.

The Meeting concluded that the data were insufficient for estimating a maximum residue level for eggplant.

*Tree nuts*

Five trials were conducted on almonds and five other trials were conducted on pecans in the USA in 2009. The GAP in the USA for tree nuts allows 2 application at a maximum rate of 0.2 kg ai/ha (in at least 935 L/ha) with a PHI of 7 days.

Residues in almond nutmeat from five trials matching US GAP were: < 0.01 (5) mg/kg.

Residues in pecan nutmeat from five trials matching US GAP were: < 0.01 (5) mg/kg.

As the GAP in the USA is established for the group of tree nuts and as the residues were all < 0.01 mg/kg, the Meeting agreed to estimate a maximum residue level for the tree nuts group at 0.01\* mg/kg.

As the B-1 was also below the LOQ in all the trials, and the nutmeat is protected by the hull and not exposed to cyflumetofen foliar spray, the Meeting estimated an STMR to be 0.01 mg/kg expressed as cyflumetofen.

*Animal feeds**Almond hulls*

Five trials were conducted on almond in the USA in 2009. The GAP in the USA allows 2 application at a maximum rate of 0.2 kg ai/ha (in at least 935 L/ha) with a PHI of 7 days.

Residues in almond hull from five trials matching US GAP were: 0.35, 0.53, 0.56, 0.86 and 1.87 mg/kg.

The Meeting estimated a maximum residue level of 4 mg/kg.

The sum of cyflumetofen and B-1 was calculated: 0.38, 0.63, 0.67, 0.98 and 2.11 mg/kg. The Meeting estimated a median residue of 0.67 mg/kg.

*Fate of residues during processing**High temperature hydrolysis*

To simulate the degradation of cyflumetofen during pasteurization, baking, brewing, boiling and sterilization, the hydrolysis of radio-labelled cyflumetofen was investigated in sterile buffered aqueous solutions.

After incubation at 90 °C (pH 4) for 20 minutes, 100 °C (pH 5) for 60 minutes or 120 °C (pH 6) for 20 minutes, no loss of radioactivity occurred. After heating at 90 °C for 20 minutes, about 69–71% of cyflumetofen remained with 14–23% of the applied radioactivity degraded to B-1. At 100 °C for 60 minutes, only 5–18% of cyflumetofen remained with the formation of B-1 (53–59% AR) and AB-1 (32–40% AR). At 120 °C, cyflumetofen was completely hydrolysed with the formation of B-1 (44–75% AR) and AB-1 (39–49% AR). Cyflumetofen was susceptible to hydrolysis at high temperature.

*Processing*

The Meeting received information on processing of orange, apple, grape and tomato.

Processing factors calculated for the processed commodities of the above raw agricultural commodities are shown in the table below. STMR-Ps were calculated for processed commodities of orange, apple and grape for which maximum residue levels were estimated.

Processed Orange Product	Cyflumetofen		Cyflumetofen and B-1*		STMR-P or median
	Processing factor	Best estimate	Processing factor	Best estimate	
Orange Juice	< 0.02, < 0.08	0.05	< 0.04, < 0.022	0.031	0.0022
Oil	102, 137	120	88.0, 133	111	7.77

Processed Orange Product	Cyflumetofen		Cyflumetofen and B-1*		STMR-P or median
	Processing factor	Best estimate	Processing factor	Best estimate	
Marmalade	0.026, < 0.08	0.026	0.059, < 0.22	0.14	0.0098
Peel	2.86, 2.97	2.92	2.56, 2.67	2.6	0.18
Molasses	< 0.02, < 0.08	< 0.06	< 0.075, < 0.502	0.28	0.020
Dried pulp	0.44, 0.584	0.51	0.759, 0.958	0.86	0.060
Apple					0.14
Applesauce	2.54, 2.91	2.7	3.16, 3.73	3.4	0.48
Juice	0.197, 0.268	0.23	0.224, 0.299	0.26	0.036
Dried apples	0.17, 0.825	0.50	0.21, 0.876	0.54	0.076
Canned apples	0.035, 0.175	0.10	0.076, 0.202	0.14	0.020
Wet pomace	0.937, 1.59	1.3	0.939, 1.57	1.3	0.18
Grape					0.22
Dried grape	0.65, 0.93, 1.88, 4.64	2.0	0.86, 1.1, 2.47, 4.65	2.3	0.506
Juice	0.064, 0.11, 0.2, 0.25	0.16	0.10, 0.16, 0.28, 0.32	0.22	0.0484
Wine (Young)	< 0.006, 0.029, 0.04, 0.04	0.04	0.04, 0.083, 0.17, 0.2	0.12	0.0264
Must	0.02, 0.18, 0.41, 0.53	0.29	0.04, 0.22, 0.44, 0.59	0.32	0.071
Wet Pomace	1.1, 0.02, 3.39, 4.22	2.2	1.1, 3.03, 3.3, 3.97	2.9	0.638
Tomato					0.07
Juice	< 0.06, 0.2	0.2	< 0.2, 0.38	0.38	0.027
Canned Tomato	< 0.04, 0.2	0.2	< 0.1, 0.3	0.3	0.021
Puree	0.3, 0.88	0.59	0.4, 0.79	0.60	0.042
Paste	0.2, 0.4	0.3	1.2, 2.2	1.7	0.12
Wet Pomace	1.3, 5.5	3.4	1.4, 5.0	3.2	0.22

\* expressed as cyflumetofen.

As the residue concentration is higher in orange oil than in fresh orange, the Meeting estimated a maximum residue level of 36 mg/kg for citrus oil by multiplying the maximum residue level of citrus fruits (0.3 mg/kg) by 120.

As the residue concentration is higher in dried grapes than in fresh grapes, the Meeting estimated a maximum residue level of 1.5 mg/kg by multiplying the maximum residue level for grapes (0.6 mg/kg) by 2.0.

In processing of foods, heating in aquatic system at high temperature may be employed, and therefore it is likely that AB-1 is present at significant concentrations after processing at slightly acidic to neutral pH ( $\geq 5$ ). It may also occur from cyflumetofen during the storage of processed foods with high water content at room temperature. However, the Meeting concluded that AB-1 is covered by the ADI for the parent compound.

### Residues in animal commodities

#### Estimation of dietary burden

The maximum and mean dietary burdens were calculated using the median residues of cyflumetofen (sum of cyflumetofen and B-1 expressed as cyflumetofen, to cover the worst case) estimated at the current Meeting on a basis of the OECD Animal Feeding Table.

Summary of livestock dietary burdens (ppm of dry matter diet)

	US-Canada		EU		Australia		Japan	
	Max	Mean	max	Mean	Max	Mean	Max	mean
Beef cattle	0.007	0.007	0.090	0.090	0.934 <sup>a</sup>	0.934 <sup>b</sup>	0	0
Dairy cattle	0.119	0.119	0.052	0.052	0.934 <sup>a</sup>	0.934 <sup>b</sup>	0	0
Broilers	0	0	0	0	0	0	0	0
Layers	0	0	0	0	0	0	0	0

<sup>a</sup> Suitable for estimating maximum residue levels for milk, meat, fat and edible offal of cattle.

<sup>b</sup> Suitable for estimating STMRs for milk, meat, fat and edible offal of cattle.

*Residues in milk and cattle tissues*

In the goat metabolism studies, in which goats were administered benzoyl-label cyflumetofen at a dose equivalent to 12.8 ppm in feed for 12 consecutive days and sacrificed one day later, cyflumetofen was found only in fat at 0.003 mg/kg. B-1 were found in liver, kidney, muscle, fat and milk at 0.125, 0.102, 0.005, 0.006 and 0.001 mg eq./kg respectively. The sum of cyflumetofen and B-1 in fat is 0.009 mg/kg as cyflumetofen. Neither cyflumetofen nor metabolites were found above 0.01 mg/kg or 10% TRR when butylphenyl-label cyflumetofen was administered to goats.

These concentrations were multiplied by 0.934/12.8 for estimating STMRs. The Meeting estimated STMRs of 0.010, 0.008, 0, 0 and 0 mg/kg for liver, kidney, meat, fat and milk, respectively.

The Meeting estimated maximum residue levels of 0.02, 0.01\*, 0.01\* and 0.01\* mg/kg expressed as cyflumetofen for edible offal (mammalian), meat (from mammals other than marine mammals), mammalian fats (except milk fat) and milks, respectively.

The dietary burden for poultry was calculated to be 0 ppm of dry matter diet. No metabolism or feeding studies on laying hens were conducted.

**RECOMMENDATIONS**

On the basis of the data from supervised trials the Meeting concluded that the residue levels listed below are suitable for establishing maximum residue limits and for IEDI and IESTI assessment.

Definition of the residue for plant commodities (for compliance with the MRL): *Cyflumetofen*.

Definition of the residue for plant commodities (for estimation of dietary intake): *Sum of cyflumetofen and 2-trifluoromethylbenzoic acid, expressed as cyflumetofen*.

Definition of the residue for animal commodities (for compliance with the MRL and estimation of dietary intake): *Sum of cyflumetofen and 2-trifluoromethylbenzoic acid, expressed as cyflumetofen*.

*Residue is not fat-soluble.*

Commodity		Recommended MRL, mg/kg		STMR/STMR-P mg/kg <sup>a</sup>
CCN	Name	New	Previous	
AM 0660	Almond hulls	4	-	Median: 0.67
FC 0001	Citrus fruit	0.3	-	0.07
	Citrus oil	36		7.77
DF 0269	Dried grape	1.5		0.506
MO 0105	Edible offal (mammalian)	0.02		Liver: 0.010 Kidney: 0.008
FB 0269	Grapes	0.6	-	0.22
MF 0100	Mammalian fats (except milk fat).	0.01*		0
MM 0095	Meat (from mammals other than marine mammals)	0.01*		0
ML 0106	Milks	0.01*		0
FP 0009	Pome fruits	0.4	-	0.14
FB 0275	Strawberry	0.6	-	0.18
VO 0448	Tomato	0.3	-	0.07
TN 0085	Tree nuts	0.01*	-	0.01
	Orange juice			0.0022
	Marmalade			0.098
	Citrus peel			0.18

Commodity		Recommended MRL, mg/kg		STMR/STMR-P mg/kg <sup>a</sup>
CCN	Name	New	Previous	
	Citrus molasses			0.020
	Apple sauce			0.48
	Apple juice			0.036
	Dried apple			0.076
	Canned apple			0.020
	Grape juice			0.048
	Grape wine			0.026
	Grape must			0.071
	Tomato juice			0.027
	Canned tomato			0.021
	Tomato puree			0.042
	Tomato paste			0.12

<sup>a</sup> Sum of cyflumetofen and B-1 expressed as cyflumetofen.

For calculating animal dietary burdens.

Commodity		Median residue, mg/kg <sup>a</sup>
CCN	Name	
	Citrus dry pulp	0.060
	Apple wet pomace	0.18
	Grape wet pomace	0.64
	Tomato wet pomace	0.22

Expressed on an “as received” basis.

<sup>a</sup> Sum of cyflumetofen and B-1 expressed as cyflumetofen.

## DIETARY RISK ASSESSMENT

### *Long-term intake*

The International Estimated Dietary Intakes (IEDIs) of cyflumetofen were calculated for the 17 GEMS/Food cluster diets using STMRs estimated by the current Meeting (Annex 3 of the 2014 Report). The ADI is 0–0.1 mg/kg bw and the calculated IEDIs were 0–1% of the maximum ADI. The Meeting concluded that the long-term intake of residues of cyflumetofen resulting from the uses considered by the current JMPR is unlikely to present a public health concern.

### *Short-term intake*

The 2014 JMPR decided that an ARfD was unnecessary. The Meeting therefore concluded that the short-term intake of residues of cyflumetofen is unlikely to present a public health concern.

## REFERENCES

Code	Author(s)	Year	Title, Institute, Report reference
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