

RECEIVED
OPPT NCIC

02 JAN 22 AM 4:25

TEST PLAN FOR METHYLATED 2-IMIDAZOLIDINONESDecember 29, 2001OVERVIEW

The Synthetic Urea Resins Group (SURG) of the Synthetic Organic Chemical Manufacturers Association. (SOCMA) hereby submits for review a test plan for 2-imidazolidinone, 4,5-dihydroxy-1,3-bis(hydroxymethyl)-, methylated (CAS No. 68411-61-4) under the Environmental Protection Agency's (EPA) High Production Volume (HPV) Chemical Challenge Program. It is the intent of the panel and its member companies to combine data on the methylated imidazolidinone with data on the well, studied non-methylated analog (2-imidazolidinone, 4,5-dihydroxy-1,3-bis(hydroxymethyl)-)(CAS No. 1854-26-8) to adequately fulfill the Screening Information Set (SIDS) endpoints. The non-methylated analog (CAS No. 1854-26-8) has already been reviewed as part of the OECD/SIDS program. Further comparison will be made between the methylated imidazolidinone and 2-imidazolidinone, 4,5-dihydroxy-1,3-bis(methoxymethyl)- (CAS No. 3001-61-4), for which various physical/chemical and environmental fate properties can be estimated by modeling.

TABLE OF CONTENTS

1.	Information about the Panel.....	2
2.	Identity of test substance and its analogs.....	2
3.	Background Information on the Test Substance and surrogate	3
4.	Test Plan.....	3
4.1	Chemical/Physical Properties	3
4.2	Environmental Fate Parameters	4
4.3	Biodegradation.....	5
4.4	Ecotoxicity	6
4.5	Human Health Data.....	7
4.5.1	Acute Toxicity	9
4.5.2	Repeated Dose Toxicity	9
4.5.3	Genetic Toxicity.....	10
4.5.4	Reproductive Toxicity	11
4.5.5	Developmental Toxicity.....	11
4.5.6	Other	11
4.5.6.1	Skin and eye irritation.....	11
4.5.6.2	Sensitization.....	12
4.5.6.3	Toxicokinetics.....	12
5.	Summary	12
6.	References.....	15
7.	Appendix 1- Robust Summaries	17

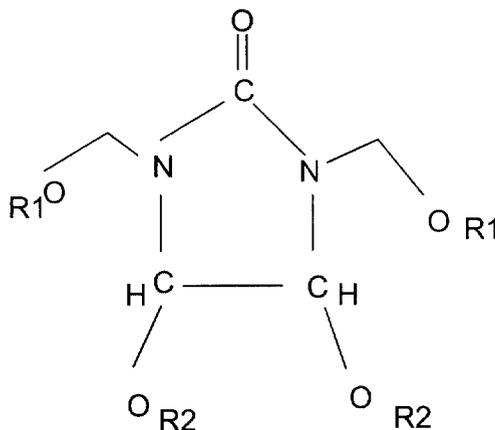
1. Information about the Panel

The Synthetic Urea Resins Group is formed under the sponsorship of the Synthetic Organic Chemical Manufacturers Association (SOCMA). The Panel consists of the following current or former manufacturers of methylated, substituted 2-imidazolidinone:

Hickson DanChem Corporation
Noveon, Inc.
OMNOVA Solutions Inc. (former manufacturer)

2. Identity of test substance and its analogs

The general molecular structure for the sponsored chemical and its analogs can be shown as follows:



Test Substance
CAS # 68411-81-4
R1, R2 = hydrogen
and/or methyl

Non-Methylated Analog
CAS # 1854-26-8
R1, R1 =hydrogen

Dimethylated Analog
CAS #3001-61-4
R1 = methyl, R2 = hydrogen

The test substance, 2-imidazolidinone, 4,5-dihydroxy-1,3-bis(hydroxymethyl)-, methylated (CAS No. 68411-81-4), is methylated to an undetermined extent. Methylation occurs primarily such that one or both R₁ groups are methyl groups instead of hydrogen atoms. It is possible that some R₂ groups are also methyl instead of hydrogen.

The primary (data rich) analog is 2-imidazolidinone, 4,5-dihydroxy-1,3-bis(hydroxymethyl)- (CAS No. 1854-26-8). All R₁s and R₂s for this substance are hydrogen atoms. This substance has already been reviewed (SIAM 10) and been assigned a low priority for further work.

A second analog is 2-imidazolidinone, 4,5-dihydroxy-1,3-bis(methoxymethyl)- (CAS No. 3001-

61-4). For this substance, the R1s are methyl groups and the R2s are hydrogen atoms. Very limited data exist for this substance. However, this analog is useful because chemical/physical and environmental fate properties can be estimated for this substance by modeling. The data obtained by modeling help to predict the effect methylation has on these properties. For simplicity in referring to the test substance and the two analogs, they will be designated,

“Methylated imidazolidinone” = CAS No. 68411-81-4 (the test substance)

“Non-methylated imidazolidinone” = CAS No. 1854-26-8 (the data-rich analog)

“Dimethylated imidazolidinone” = CAS No. 3001-61-4 (the analog useful for comparative modeling)

3. Background Information on the Test Substance and surrogate

The test substance is manufactured by the reaction of glyoxal, urea and formaldehyde followed by methylation using methanol.

Both the test substance and the surrogates are substituted 4,5-dihydroxy-1,3-hydroxymethyl-2-imidazolidinones that are used in textile manufacture as industrial intermediates to produce easy care fabrics. They are applied to textile cloth and cured to crosslink with the cellulose molecules in the cloth, so that the finished textile cloth will have “memory” to retain crease or other desired shape. The methylated test substance contains lower residual levels of formaldehyde compared to the non-methylated analog. Residual formaldehyde in these products is released during processing, therefore minimizing any potential consumer exposure.

4. Test Plan

It is the intention of the Synthetic Urea Resins Group of SOCMA, which includes the manufacturers of “methylated imidazolidinone” (CAS No. 68411-81-4) to use information on this substance, combined with available studies on the related “non-methylated imidazolidinone” (CAS No. 1854-26-8) to fulfill the screening information data needs. An IUCLID data set summarizing the available studies (with Klimisch codes) for CAS No. 1854-26-8 exists (IUCLID, 2000), as well as a SIAR. CAS No. 1854-26-8 was reviewed at SIAM 10, and assigned “low priority for further work.” The following sections discuss individual endpoints and how they are to be addressed. Study details may be reviewed in the robust summary set for CAS No. 68411-81-4.

4.1 Chemical/Physical Properties

Chemical and physical property data for the related “non-methylated imidazolidinone” (CAS No. 1854-26-8) are also shown in Table 1. Values for the two materials are comparable, demonstrating close resemblance in chemical physical properties.

Chemical/Physical Property data for CAS No. 68411-81-4 are reported in Table 1. A melting point of -39° C has been determined using OECD Guide-line 102 "Melting Point/Melting Range"

under good laboratory practices (Tognucci, 2001a)(Table 1). A boiling point of 118.5° C at 980 hPa has been determined using OECD Guide-line 103 "Boiling Point/boiling Range" employing good laboratory practices (Tognucci, 2001b). A specific gravity of 1.30-1.31 @25°C is provided in the Material Safety Data Sheet from Noveon, Inc. (2001).

Table 1. Chemical/physical properties of substituted 2-imidazolidinones

Endpoint	Methylated imidazolidinone (CAS No. 68411-81-4) ¹	Dihydroxy imidazolidinone (CAS No. 1854-26-8) ²
Melting point (° C)	-39	-35
Boiling point (° C)	118.5	106
Density	1.30- 1.31	1.36
Partition coefficient (log Kow)	-3.2	-2.2
Water solubility (g/l)	> 5000	Miscible
Vapor pressure	Similar to water	Similar to water

¹The test substance had the following composition: ca. 84% CAS No. 68411-81-4; ca. 13% H₂O, and 0.18% formaldehyde (CAS No. 50-00-0)

² Data were obtained from IUCLID data set for CAS No. 1854-26-8, dated 04-FEB-2000.

A water solubility of > 5000 g/l at 20° C was determined using OECD Guide-line 105 "Water Solubility" (Tognucci, 2001d).

The vapor pressures of the neat test product or its analogs are not known. However, these materials are commercially available as aqueous concentrates, and therefore are likely to have vapor pressures similar to water (water vapor pressure = 23.79 mm Hg or 31.71 hPa @ 25 °C).

A Log Kow of -3.2 at 20° C has been calculated from the following equation (Tognucci, 2001c):

$$\log Kow = \log (\text{n-octanol solubility}/\text{water solubility}) = (<3.25 \text{ g/l} / >5000 \text{ g/l}) = < -3.2$$

This equation used the value of >5000g/l water solubility previously determined, and the value of <3.25 g/l for the solubility in n-octanol. The n-octanol solubility of the test material was determined to be <3.25 g/l by adding 0.13-0.14 grams of test material to 40 ml n-octanol at room temperature and stirring. The result was incomplete dissolving and two phases.

The negative log Kow indicates a greater propensity for the methylated material to partition to water than organic solvents.

4.2 Environmental Fate Parameters

Modeling data discussed below suggest that the nonmethylated, methylated, and dimethylated 2-imidazolidinones will have similar environmental fate properties.

Environmental fate parameters for methylated 2-imidazolidinone (CAS No. 68411-81-4) were obtained using the EPIWIN Program. The test substance itself cannot be modeled, because it does not have a precise, defined molecular structure. Both nonmethylated 2-imidazolidinone

(CAS No.1854-26-8) and the dimethylated 2-imidazolidinone (CAS No. 3001-61-4), which have well defined molecular structures, can be modeled using EPIWIN. As discussed above, these surrogates have closely analogous molecular structures, which bracket the molecular structure of methylated 2-imidazolidinone. Therefore the modeled environmental fate parameters of the surrogates should correspond closely to estimated parameters for methylated 2-imidazolidinone.

The comparative modeled environmental fate parameters for nonmethylated 2-imidazolidinone and dimethylated 2-imidazolidinone are listed in Table 2. As can be seen from the table, the values for both nonmethylated and dimethylated materials in general have close similarity. It is reasonable to expect for methylated test substance that the hydroxyl rate constant will range from 73.2 to 94.5 E-12 cm³/molecule-sec. Similarly, the photolysis half-life will be between 0.11-0.15 days. There is about a 10-fold difference in the two Henry's Law Constants, but assuming a value for methylated test material between the range of 1.09E-13 to 10.6E-12 would indicate that all three compounds would display a low tendency toward volatilization from water. Fugacity Level III modeling for both the nonmethylated and dimethylated materials shows close agreement for relative concentrations in air, water, soil and sediment compartments under equilibrium conditions. It is reasonable therefore to expect that the methylated substance will preferentially partition to water and soil. Although stability of the test substance in water has not been determined, the commercial form of the product is as an aqueous concentrate. From a practical standpoint, therefore, it is likely that the product does not hydrolyze readily at neutral pHs (at ambient temperatures).

Table 2. Environmental fate parameters for substituted 2-imidazolidinones*

Environmental Fate Parameter	Nonmethylated 2-imidazolidinone (CAS No. 1854-26-8)	Dimethylated 2-imidazolidinone (CAS No. 3001-61-4)
Photolysis Hydroxyl Rate Constant (cm ³ /molecule-sec)	73.195 E-12	94.5 E-12
Photolysis half-life (days)	0.146	0.113
Stability in Water	Qualitatively stable	Qualitatively stable
Henry's Law Constant (atm-m ³ /mole)	1.06E-12	1.09E-13
Level III Fugacity: Air	0.00133 %	5.66E-6 %
Water	42.8 %	45.3 %
Soil	57.1 %	54.6 %
Sediment	0.0638 %	0.0755 %

* The values in this table for the analogs are predictive of the methylated test material of interest (CAS No. 68411-81-4), which cannot be modeled because it does not possess a precisely defined molecular structure.

4.3 Biodegradation

No biodegradation studies have been found for methylated imidazolidinone. Four biodegradation studies have been identified for the non-methylated imidazolidinone. The results shown in Table 3 indicate inherent biodegradability for the non-methylated imidazolidinone.

The EPIWIN BIOWIN Program [Version (4.0)] conducted for the non-methylated substance qualitatively predicts that it biodegrades quickly. The same program conducted for the dimethylated analog predicts slow biodegradation. This comparison suggests that methylation may retard biodegradation to some extent, and that methylated imidazolidinone will biodegrade more slowly than the unmethylated surrogate. No further biodegradation testing is recommended for methylated imidazolidinone.

Table 3. Biodegradation for substituted 2-imidazolidinones*

Study	Nonmethylated 2-imidazolidinone (CAS No. 1854-26-8)	Dimethylated 2-imidazolidinone (CAS No. 3001-61-4)
Aerobic, activated sludge, non-adapted	60-70% after 28 day, “inherently biodegradable” ¹	-
Aerobic, activated sludge, non-adapted	27% after 8 days, 28% after 58 days ²	-
Aerobic, activated sludge	38% after 28 days ³	-
Activated sludge	70% after 2 months ⁴	-
EPIWIN/BIOWIN (v4.00)	“biodegrades fast”	“Does not biodegrade fast”

* Data for the methylated test material of interest (CAS No. 68411-81-4) are not available

¹ BASF, 1996 .OECD Guideline 301. Test substance consisted of 73.9% active ingredient and 26.1% water.

² BASF, 1996 OECD Guideline 303. Test substance consisted of 73.9% active ingredient and 26.1% water.

³ BASF, 1980b. Study remark: No oxygen consumption; elimination probably not due to biodegradation.

⁴ BASF, 1974. Test substance consisted of 45 % active ingredient and 55 % water

4.4 Ecotoxicity

Results of ecotoxicity studies with the non-methylated imidazolidinone are summarized in Table 4. ECOSAR modeling predicts that both the non-methylated imidazolidinone and the dimethylated imidazolidinone exhibit low toxicity. Actual studies indicate low toxicity to fish, Daphnia and bacteria, and moderate to moderately high toxicity to algae. Although the ECOSAR predictions appear high in relation to the other studies, the modeling indicates that the presence or absence of methylation on the 1 and 3 hydroxymethyl positions does not have a significant bearing on aquatic toxicity. Therefore, it is reasonable to conclude that methylated imidazolidinone possesses a similar degree of toxicity as both non-methylated imidazolidinone and dimethylated imidazolidinone. Accordingly, no additional studies are recommended for the aquatic toxicity endpoints.

Table 4. Ecotoxicity Studies for substituted 2-imidazolidinones*

Endpoint	Non-methylated imidazolidinone (CAS No. 1854-26-8)	Dimethylated imidazolidinone (CAS No. 3001-61-4)
Acute toxicity to fish (96 hr LC ₅₀ , mg/l)	2200 ^a 3.6E+9 ^b	1.9E+7 ^b
Acute toxicity to Daphnia (48 hr LC ₅₀ , mg/l)	>500 ^{c,d} 2.23E+9 ^b	1.4E+9 ^b
Chronic toxicity to Daphnia (21 day NOEC, mg/l)	≥ 100 ^c	-
Toxicity to algae (EC ₅₀ , mg/l)	36.9 ^{c,f} 28.4 ^{c,g} 8.85E+8 ^{b,g}	6.41E+6 ^{b,g}
Bacteria (mg/l)	2200 ^{c,h} > 1000 ⁱ 1995 ^j	-

* Data for the methylated test material of interest (CAS No. 68411-81-4) are not available. CAS No. 68411-81-4 cannot be modeled because it does not possess a precisely defined molecular structure.

^a BASF, 1990. Active ingredient: 40%

^b EPIWIN/ECOSAR Program (v0.99f).

^c BASF, 1988. Active ingredient: 40%

^d Directive 84/449/EEC, C.2 "Acute Toxicity for Daphnia"

^e BASF, 1999; EEC Guideline XI/681/86. Active ingredient: 70%

^f 72 hours; ^g 96 hours

^h Growth Inhibition Test, DIN 38412/8. 17 hr EC50. Active ingredient: 40%

ⁱ BASF 1996c. OECD Guideline 209 "Activated Sludge, Respiration Inhibition Test". 30 min NOEC. Active ingredient: 74%

^j BASF 1980a. Short term respiration test. Highest concentration of material tested with < 20 % inhibition in 30 min.

4.5 Human Health Data

Results of mammalian toxicity tests conducted on the non-methylated imidazolidinone are summarized in Table 5. These studies indicate that the material has a low potential for acute, repeated dose, reproductive or developmental toxicity. Based on the structural similarity of the methylated material with the non-methylated material, the methylated material is also expected to have a fairly low potential for acute, repeated dose, genetic, reproductive, or developmental toxicity.

Table 5. Mammalian Toxicity Data for substituted 2-imidazolidinones*

Endpoint	Non-methylated imidazolidinone (CAS No. 1854-26-8)	Reference
Acute oral (LD ₅₀ , mg/kg) ¹	≥ 2,880 (rat) ² ≥ 10,000 (rat) ² ≥ 10,000 (mouse) ²	BASF, 1973 IRDC, 1983a IRDC, 1983b
Acute inhalation	No mortality after 8 or 1 hr exposure to saturated atmosphere @ 20 or 150 °C (respectively)	BASF, 1973
Repeated oral dose toxicity ³ (NOAEL, mg/kg/day)	4,000 (rat, 14 day) 1,000 (rat, 90 day) ≥ 11,680 (mouse, 14 day) ² ≥ 6,000 (mouse, 90 day) ²	IRDC, 1983a IRDC, 1983a IRDC, 1983b IRDC, 1983b
In vitro genetic toxicity (Ames)	TA97, TA98, TA 1535, TA1537, and TA102 -negative TA100 - equivocal	Zeiger et al., 1987 CCR, 1992; NTP, 1984 NTP, 1984; Zeiger et al., 1987
In vivo genetic toxicity	Mouse micronucleus at 2000 mg/kg - negative Sex linked recessive lethal at 60000 ppm in Drosophila - positive Reciprocal translocation at 50000 ppm in Drosophila - negative	Biopharm, 1995 Fouremant et al., 1994 Fouremant et al., 1994
Reproductive toxicity ^{3,4} (NOAEL, mg/kg)	3,000 (rat) ≥ 6,000 (mouse) ²	IRDC, 1983a IRDC, 1983b
Developmental toxicity ¹ (NOAEL, mg/kg)	≥ 640 (rat) ²	HMR, 1998

* Data for the methylated test material of interest (CAS No. 68411-81-4) and dimethylated imidazolidinone (CAS No. 3001-61-4) are not available

¹ Values refer to 100% test material

² Highest dose used

³ Test material contained 41.4% CAS No. 1854-26-8

⁴ Examination of reproductive organs from 90-day study

4.5.1 Acute Toxicity

Acute toxicity testing has not been conducted for the methylated imidazolidinone. Two acute oral studies for the rat and one for the mouse have been performed and summarized for the nonmethylated imidazolidinone. LD₅₀ values ranged from >2,880 to >10,000 mg/kg. In mice receiving 2880 mg/kg by i.p. injection, the only symptoms observed were dyspnea and atony. Macroscopic inspection showed no pathological findings (BASF, 1973).

At ambient temperature, inhalation exposure for 8 hr to an atmosphere highly enriched in vapors from a 45 % aqueous solution (Fixapret CPN) caused no lethality, but caused dyspnea and irritation of mucous membranes (BASF, 1973). Vapors generated at 150° C produced severe irritation and dyspnea and were lethal to rats within a few hours (BASF, 1973). Spot-like hyperemia and edema of the lung were prominent, while hydrothorax was seen in isolated cases. It is assumed that decomposition products arising at temperatures greater than 40° C induced these serious effects (SIAR for CAS No. 1854-26-8; reviewed at SIAM 10).

Based on the structural similarity of the methylated material (CAS No. 68411-81-4) with the non-methylated material (CAS No. 1854-26-8), it is likely that the methylated material would also have fairly low acute oral and inhalation toxicity.

4.5.2 Repeated Dose Toxicity

No repeated dose studies have been identified for the methylated imidazolidinone (CAS No. 68411-81-4). Repeated dose studies are available in the rat and mouse for the related non-methylated imidazolidinone (CAS No. 1854-26-8)(Table 5). Fourteen-day oral (gavage) studies in rats and mice were conducted at doses ranging from 256 to 11,600 mg/kg/day (IRDC, 1983a,b). No test-related toxicologically significant macroscopic lesions or abnormalities were observed in rats or mice treated with any dose (with the exception of a moderately inflammatory bilateral reaction in the nasal passages of rats treated with 11,600 mg/kg).

Ninety-day oral (gavage) studies have been run in both the rat and the mouse for the non-methylated analog (CAS No. 1854-26-8)(Table 5) (IRDC, 1983a,b). Dose levels were 1000, 3000 and 6000 mg/kg/day of a material containing 41.4% CAS No. 1854-26-8. Pharmacotoxic signs noted for male and female rats in the 3000 and 6000 mg/kg/day dosage level groups including discoloration of the fur, soft stool, hypoactivity, decreased grasping reflex, ataxia, and decreased temperature of extremities. No deaths were reported and no toxicologically significant organ weight changes were observed. On postmortum examination, mild to moderate mineralization was observed in the heart and testes of two male rats, and multiple yellowish linear macroscopic lesions were observed in the right testis of one male rat treated with 6000 mg/kg/day.

In the 90-day mouse study, females of all dose groups and males at 1000 and 6000 mg/kg/day dose groups showed increased weight gains compared to controls (not toxicologically significant). The 6000 mg/kg/day group and controls showed no microscopically visible changes (the animals of the

1000 and 3000 mg/kg/day doses were not examined). One death occurred in the 3000 mg/kg male dose group at week three that was not considered treatment related.

The above studies are indicative of generally low repeated dose toxicity of non-methylated imidazolidinone (CAS No. 1854-26-8), even when adjusting the effective dose downward in the 90-day studies to reflect the 41.4% concentration of active ingredient. Based on the close structural similarity of non-methylated and methylated imidazolidinones, it is reasonable to conclude that the repeated dose toxicity of the methylated material (CAS No. 68411-81-4) would not differ significantly from that of the non-methylated material.

4.5.3 Genetic Toxicity

No genotoxicity studies have been identified for methylated imidazolidinone (CAS No. 68411-81-4). Three Ames tests are reported and summarized for non-methylated imidazolidinone (CAS No. 1854-26-8) (Zeiger et al., 1987; CCR, 1992; NTP, 1984). The tests show that non-methylated analog was negative in *Salmonella* strains TA1535, TA1537, and TA102 in the presence and absence of metabolic activation. However, results of the study by Zeiger et al., 1984 (which was conducted in two different laboratories) are equivocal for strains TA98 and TA100. In the presence of S-9, approximately 50% of tests in strain TA100 were questionable in one laboratory (with test material in DMSO) and all tests were positive in the other laboratory (with test material in water). Weakly positive or questionable results were found in strain TA98 in the presence or absence of S-9 in the same laboratory that found positive results in strain TA100 (material was in water). One out of five tests in the other laboratory with strain TA98 in the presence of S-9 (and test material in DMSO) showed a weak response. Because the tests with the two solvents were performed in different laboratories, it is difficult to discern whether the variable results were due to the tests being conducted in different laboratories or the use of different solvents.

Three *in vivo* genetic toxicity studies have been conducted in *Drosophila melanogaster* [two sex linked recessive lethal (SLR) and one reciprocal translocation] with non-methylated imidazolidinone (Fouremant et al., 1994). At a very high concentration of 60000 ppm (either orally or by i.p. injection), the test material induced a four-fold increase in sex-linked recessive lethal events. However, oral administration of a similar concentration (50000 ppm) did not lead to an increase in reciprocal translocations. Because the only concentration used in the sex-linked *Drosophila* study was very high, it is not known whether these effects occur at lower, more realistic exposure concentrations. Furthermore, due to study deficiencies, the sex-linked *Drosophila* study was assigned a reliability rating of 4. The Ames study is considered to be a more reliable test for assessing genotoxic potential of CAS No. 1854-26-8.

A mouse micronucleus study conducted on CAS No. 1854-26-8 under OECD Guideline 474, (using good laboratory practices) indicated that the test material did not increase the frequency of micronuclei at 2000 mg/kg (Biopharm, 1995).

The above studies are indicative of low potential for non-methylated imidazolidinone (CAS No. 1854-26-8) to produce genotoxicity. Only high doses of the material (generally 3333 mg/plate or higher) were shown to be mutagenic in some strains in the presence of S-9. Based on the close structural similarity of non-methylated and methylated imidazolidinones, the *in vitro* genetic

toxicity profile of the methylated material (CAS No. 68411-81-4) is not expected to differ significantly from that of the non-methylated material. Also, because the material is largely excreted unchanged in the urine upon oral administration and is not absorbed well from the skin (see Toxicokinetics below), the potential for DNA-reactive metabolites to be formed after in vivo exposure is low. Therefore, no additional in vitro testing is planned.

4.5.4 Reproductive Toxicity

No reproductive toxicity study has been identified for methylated imidazolidinone (CAS No. 68411-81-4). However, ninety-day oral (gavage) toxicity studies (that included examination of the reproductive organs of both sexes) on a material containing 41.4% of the non-methylated analog (CAS No. 1854-26-8) have been conducted in rats and mice (IRDC, 1983a,b). Microscopic inspection of these organs (including testes, epididymis, prostate, preputial gland/uterus, ovaries, clitoral gland) gave no indication of morphological abnormalities. No histopathological changes were observed up to 3000 mg/kg/day in male rats and up to 6000 mg/kg/day in female rats. No changes were seen up to 6000 mg/kg/day in both genders of mice. These studies are predictive of low reproductive toxicity for the methylated imidazolidinone. No additional studies are planned for this endpoint.

4.5.5 Developmental Toxicity

No developmental toxicity study has been identified for methylated imidazolidinone (CAS No. 68411-81-4). Teratogenicity testing has been conducted for non-methylated imidazolidinone (CAS No. 1854-26-8) in pregnant Wistar rats, following OECD Guideline 414 (HMR, 1998)(Table 5). No compound-related effects were observed at doses up to 1000 mg/kg. Since the test substance was 61.4% CAS No. 1854-26-8 in water, a NOAEL of 640 mg/kg was determined for both maternal toxicity and teratogenicity. Based on the structural similarity of the methylated imidazolidinone to the non-methylated material, this study should be predictive of developmental toxicity for methylated imidazolidinone. Therefore, no further testing is planned.

4.5.6 Other

4.5.6.1 Skin and eye irritation

No skin and eye irritation studies have been identified for methylated imidazolidinone (CAS No. 68411-81-4). Irritation studies are available for non-methylated imidazolidinone (CAS No. 1854-26-8). One skin irritation study (BASF, 1973) using the rabbit showed no irritation, and a secondary reference (Marhold [1972] in Czech) cited in RTECs indicated severe irritation. One eye irritation study, using the rabbit, showed no irritation (BASF, 1973), whereas the same secondary RTECs Czech reference noted above indicated mild irritation. The BASF study is identified as the critical, valid study for this endpoint. Therefore, it is concluded that CAS No. 1854-26-8 is not highly irritating to the skin and eye. Based on the structural similarities between the methylated and nonmethylated material, strong skin and eye irritation is not likely to be associated with the methylated imidazolidinone (CAS No. 68411-81-4).

4.5.6.2 Sensitization

Human experience data with methylated imidazolidinone (CAS No. 68411-81-4) is limited. However, none of the sponsors identified any reports of skin sensitization in people who work with this material. Several cases of skin sensitization, dermatitis or eczema have been reported in humans who have contacted resin-treated textiles (Malten, 1964; Andersen and Harman, 1982; Tegner, 1985; Fregert and Tegner, 1971; Hatch and Maibach, 1986, Scheman et al., 1998; Sommer et al., 1999; BG Chemie, 1995;). Many of the examined cases showed patch test results to both CAS No. 1854-26-8 and formaldehyde (a probable contaminant). Because the methylated imidazolidinone used for patch testing was not analyzed in any of the studies, one cannot conclude that the methylated imidazolidinone (and not contaminating formaldehyde) was the sensitizer. Because the methylated material is less likely to release formaldehyde than the nonmethylated material, the potential for sensitization to occur due to formaldehyde exposure is expected to be low.

4.5.6.3 Toxicokinetics

Results of studies in rats and monkeys indicate that non-methylated imidazolidinone (CAS No. 1854-26-8) is poorly absorbed from the skin (Jeffcoat, 1984; 1985). Hydration of the skin increases absorption. After oral or intravenous administration, the material is quickly distributed to the skin, muscle, blood, liver and kidney (Robbins et al., 1984, Robbins and Norred, 1984; Jeffcoat, 1985). Within 24-hours of oral or intravenous administration, the vast majority of the material is excreted unchanged in the urine (Jeffcoat, 1985). Based on the structural similarities between the methylated and nonmethylated material, the methylated imidazolidinone (CAS No. 68411-81-4) is likely to have a similar pharmacokinetic profile as the nonmethylated material.

5. Summary

In summary, based on the structural/physical similarities and chemical physical between methylated imidazolidinone (CAS No. 68411-81-4), dimethylated material (CAS No. 3001-61-4), and non-methylated imidazolidinone (CAS No. 1854-26-8), the data for the non-methylated and dimethylated materials will be predictive of toxicity for the methylated material. Physical/chemistry data for the actual test material, modeled environmental fate data for the dimethylated material (CAS No. 3001-61-4) and experimental ecotoxicity and mammalian toxicity data for the nonmethylated material are present to satisfy all endpoints (Tables 6 and 7). No further testing is planned.

Table 6. Test Plan

CAS No. 68411-81-4	Information	OECD Study	Other	Estimation	GLP	Acceptable	New Testing Required
ENDPOINT	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N
PHYS/CHEM PROPERTIES							
Melting Point	Y	Y	N	N	Y	Y	N
Boiling Point	Y	Y	N	Y	Y	Y	N
Vapor Pressure	Y ¹	N	N	N	N	N	N
Partition Coefficient	Y	N	Y	N	Y	Y	N
Water Solubility	Y	Y	N	N	Y	Y	N
ENVIRONMENTAL FATE							
Photodegradation	Y*	N	Y	Y	N	Y	N
Stability in Water	Y ²	N	N	N	N	N	N
Biodegradation	Y*	Y	N	N	Y	Y	N
Transport between Environmental Compartments (Fugacity)	Y*	N	Y	Y	N	Y	N
ECOTOXICITY							
Acute Toxicity to Fish	Y*	N	Y	N	N	Y	N
Acute Toxicity to Aquatic Invertebrates	Y*	N	Y	N	N	Y	N
Toxicity to Aquatic Plants	Y*	N	Y	N	N	Y	N
TOXICOLOGICAL DATA							
Acute Toxicity	Y*	N	Y	N	N	N	N
Repeated Dose Toxicity	Y*	Y	N	N	N	Y	N
Genetic Toxicity-Mutation	Y*	N	Y	N	N	N	N
Genetic Toxicity-Chromosomal Aberrations (mouse micronucleus)	Y*	Y	N	N	Y	Y	N
Toxicity to Reproduction	Y*	N	Y	N	N	Y	N
Developmental Toxicity	Y*	Y	N	N	Y	Y	N
OTHER TOXICITY DATA							
Skin Irritation	Y*	N	Y	N	N	Y	N
Eye Irritation	Y*	N	Y	N	N	Y	N
Skin Sensitization	Y*	N	Y	N	N	Y	N
Absorption	Y*	N	Y	N	N	Y	N

*Data on surrogate chemical (CAS No. 1854-26-8) are used

¹ Actual value is not known; however is likely to be close to that of water, since the product is commercially available as an aqueous concentrate.

² Actual value is not known; however material is likely to be stable in water because it is sold in the form of an aqueous concentrate.

Table 7. Analog Matrix for 2-Imidazolidinone, 4,5-dihydroxy-1,3-bis(hydroxymethyl)-, methylated*

ENDPOINT	68411-81-4 (Methylated) (Test substance)	1854-26-8 (Non-methylated) (Analog)	3001-61-4 (Dimethylated) (Analog)
PHYSICAL CHEMISTRY			
Melting point	A	A	NR
Boiling point	A	A	NR
Density	A	A	NR
Vapor Pressure	E	E	NR
Water Solubility	A	A	NR
Kow	A	A	NR
ENVIRONMENTAL FATE			
Photodegradation	R	Calc.	Calc.
Stability in Water	E	E	NR
Biodegradation	R	A & Calc.	Calc.
Transport between Environmental Compartments (Fugacity)	R	Calc.	Calc.
ECOTOXICITY			
Acute Toxicity to Fish	R	A & Calc.	Calc.
Acute Toxicity to Aquatic Invertebrates	R	A & Calc.	Calc.
Toxicity to Aquatic Plants	R	A & Calc.	Calc.
TOXICOLOGICAL DATA			
Acute Toxicity	R	A	NR
Repeated Dose Toxicity	R	A	NR
Genetic Toxicity-Mutation	R	A	NR
Genetic Toxicity-Chromosomal Aberrations	R	A	NR
Carcinogenicity (NR)	NR	NR	NR
Toxicity to Reproduction	R	A	NR
Developmental Toxicity	R	A	NR
OTHER TOXICITY DATA			
Skin Irritation (NR)	R	A	NR
Eye Irritation (NR)	R	A	NR
Skin Sensitization (NR)	R	A	NR
Toxicokinetics (NR)	R	A	NR

* Data on analogs CAS No. 1854-26-8 and 3001-61-4 are shown for comparison.

R = Required endpoint fulfilled by surrogate, SAR; Test = Testing planned to fulfill requirement; Calc. = Calculated value; E = estimated qualitatively; A = Adequate existing data; NR = Not required

6. References

- Andersen KE, Hamann K. 1982. Cost benefit of patch testing with textile finish resins. *Contact Dermatitis* 8:64-67.
- BASF AG, Department of Toxicology. 1973. Unpublished study (XXII/230), 23.01.73
- BASF AG, Analytical Department, 1974. Unpublished data (Fixapret CPN 45%), (J.Nr. 12501; 22.07.74)
- BASF AG, Laboratory of Ecology. 1980a. Unpublished data, (Fixapret CP; 28.08.80)
- BASF AG, Laboratory of Ecology. 1980b. Unpublished data, (beginning of the test: 13.08.80)
- BASF AG, Laboratory of Ecology. 1988. Unpublished data, (1200/87)
- BASF AG, Department of Toxicology. 1990. Unpublished study (89/183), 15.05.90
- BASF. 1996a. Study No. 96/0117/21/1
- BASF. 1996b. Study No. 96/0117/30/2
- BASF. 1996c. Study No. 96/0117/08/1
- BASF AG, Laboratory of Ecology. 1999. Unpublished data (98/0419/51/3), 02.02.1999
- BG Chemie. 1995. Dimethyloldihydroxyethylenharnstoff; No. 230, Toxikologische Bewertung, Berufsgenossenschaft der chemischen Industrie, Heidelberg.
- BIOPHARM. 1995. Unpublished results, report No. 022 TOX94, June 2, 1995. Sponsored by BG Chemie, Germany
- CCR (Cytotest Cell Research GmbH & CoKG). 1992. Unpublished report CCR no. 291407, ZHT Proj. No. 40MO502/919014 (sponsored by BASF AG, Germany, Nov. 10, 1992).
- Fouremant P, Mason JM, Valencia R, Zimmering S. 1994. Chemical mutagenesis testing in *Drosophila*. IX. Results of 50 coded compounds tested for the National Toxicology Program. *Environ Molec Mutagen* 23:51-63.
- Fregert S, Tegner E. 1971. *Contact Dermatitis News Lett.* 9:200.
- Hatch KL, Maibach HI. 1986. Textile chemical finish dermatitis. *Contact Dermatitis* 14:1-13.
- HMR (Hoechst Marion Roussel) Deutschland. 1998. Unpublished results, report No. 97.0590, Sept. 25, 1998. Sponsored by BG Chemie, Germany

Int. Research & Development Corp (IRDC). 1983a. Unpublished Report No. 5701-1-307, Dated July 15, 1983 (short communication).

Int. Research & Development Corp (IRDC). 1983b. Unpublished Report No. 5701-1-303, Dated July 13, 1983 (short communication).

IUCLID. 2000. Substance ID: 1854-26-8. BASF AG, 04-FEB-2000.

Jeffcoat AR. 1984. Percutaneous penetration of formaldehyde. NTIS/OTS 0512137, Doc ID. 40-8470033

Jeffcoat AR. 1985. Nat. Inst. Environ. Health, Research Triangle Institute, "Adsorption, Disposition, Metabolism and Excretion of 1,3-Dimethylol-4,5-Dihydroxy-2-imidazolidinone (DMDHEU)", Contract No. N01-ES-1-5007, December 1985

Malten KE. 1964. Textile finish contact hypersensitivity. Arch Dermatol 89:215-221.

Marhold JV (1972 in Czech), cited in RTECS (1999) NIOSH, USA.

MSDS for Freerez® MTH Conc. Document: RZMTHCNC CFLN: AUUS Effective Date: 18 October 2001. Noveon, Inc., 9911 Brecksville Rd., Cleveland, OH 44141-3247. NTP Annual Plan, Fiscal Year 1984, S. 58

Robbins JD and Norred WP. 1984. NTIS/OTS0512125, 40-8470042.

Robbins JD, Norred WP, Bathija A, Ulsamer AG. 1984. Bioavailability in rabbits of formaldehyde from durable-press textiles. J Toxicol Environ Health 14:453-463.

SIAR for CAS no. 1854-26-8. (Reviewed at SIAM 10)

Scheman AJ, Carroll PA, Brown KH, Osburn AH. 1998. Formaldehyde-related textile allergy: an update. Contact Dermatitis 38:332-336.

Sommer S, Wilkinson SM, Dodman B. 1999. Contact dermatitis due to urea-formaldehyde resin in shin pads. Contact Dermatitis 40:159-160.

Tegner E. 1985. Acta Derm Venereol. (Stockholm). 65:254-257.

Tognucci A. 2001a. "Determination of the Freezing Point/Freezing Range of Freerez® MTH Conc," Study No. 822690, RCC Ltd 4452 Itingen Switzerland.

Tognucci A. 2001b. "Determination of the Boiling Point/Boiling Range of Freerez® MTH Concentrate," Study No. 822701, RCC Ltd, 4452 Itingen, Switzerland.

Tognucci A, 2001c. "Determination of the Partition Coefficient (N-Octanol/Water) of Freerez® MTH Concentrate," Study No. 882723, RCC Ltd, 4452 Itingen, Switzerland.

Tognucci A. 2001d. "Determination of the Water Solubility of Freerez® MTH Concentrate," Study No. 822712, RCC Ltd, 4452 Itingen, Switzerland.

Zeiger E, Anderson B, Haworth S et al. 1987. Salmonella mutagenicity tests: III. Results from the testing of 255 chemicals. Environ Mutagen 9(Suppl. 9):1-110.

7. Appendix 1- Robust Summaries