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**NATIONAL INDUSTRIAL CHEMICALS NOTIFICATION AND ASSESSMENT SCHEME
(NICNAS)**

FULL PUBLIC REPORT

Complex Soap TH17

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**Director
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FULL PUBLIC REPORT

Complex Soap TH17

1. APPLICANT AND NOTIFICATION DETAILS

APPLICANT(S)

Klüber Lubrication Australia Pty Ltd (ABN 77 005 809 852) of 3 Brand Drive Thomastown VIC 3074.

NOTIFICATION CATEGORY

Standard: Chemical other than polymer (more than 1 tonne per year).

EXEMPT INFORMATION (SECTION 75 OF THE ACT)

Data items and details claimed exempt from publication: chemical name, other name, CAS number, molecular and structural formula, molecular weight, purity and spectral data.

VARIATION OF DATA REQUIREMENTS (SECTION 24 OF THE ACT)

Variation to the schedule of data requirements is claimed as follows: vapour pressure, water solubility, hydrolysis as function of pH, partition coefficient, adsorption/desorption, dissociation constant, particle size, flash point, explosive properties, reactivity, acute inhalation toxicity and chromosome damage (in vivo).

PREVIOUS NOTIFICATION IN AUSTRALIA BY APPLICANT(S)

None.

NOTIFICATION IN OTHER COUNTRIES

Austria, 2000.

2. IDENTITY OF CHEMICAL

MARKETING NAME(S)

Complex Soap TH17

Komplex Seife TH17

Isoflex NBU (formulated grease)

Isoflex Topas NB (formulated grease)

Staburags NBU (formulated grease)

Altemp Q NB (formulated grease).

3. COMPOSITION

DEGREE OF PURITY

High.

4. INTRODUCTION AND USE INFORMATION

MODE OF INTRODUCTION OF NOTIFIED CHEMICAL (100%) OVER NEXT 5 YEARS

The ready-to-use grease containing the notified chemical will be imported by sea in plastic lined 25 kg pails and 180 kg steel drums.

MAXIMUM INTRODUCTION VOLUME OF NOTIFIED CHEMICAL (100%) OVER NEXT 5 YEARS

<i>Year</i>	<i>1</i>	<i>2</i>	<i>3</i>	<i>4</i>	<i>5</i>
<i>Tonnes</i>	<5	<5	<5	<5	<5

USE

The notified chemical is a component of a ready-to-use grease (at a concentration of less than 30%) used for long term and lifelong lubrication of rolling bearings in sealed components.

5. PROCESS AND RELEASE INFORMATION**5.1. Distribution, Transport and Storage****PORT OF ENTRY**

Melbourne, Victoria

IDENTITY OF MANUFACTURER/RECIPIENTS

Various industrial sites throughout Australia.

TRANSPORTATION AND PACKAGING

The notified chemical will be imported into Melbourne in plastic lined 25 kg pails and 180 kg steel drums by sea. From the notifier's site in Victoria the grease will be transported to various industrial sites throughout Australia.

5.2. Operation Description

More than half of the notified chemical will be used in the new automotive, machinery and equipment manufacturing sites. The greases containing the notified chemical will be added via automatic metering devices, dip feed devices or centralised lubrication devices in a closed system during the assembly of automotive components or machine and equipment parts.

Maintenance fitters and other mechanics at maintenance workshops will use less than half of the notified chemical. They will apply the greases manually by brush, spatula, grease gun or grease cartridge to existing machinery.

5.3. Occupational exposure*Number and Category of Workers*

<i>Category of Worker</i>	<i>Number</i>	<i>Exposure Duration</i>	<i>Exposure Frequency</i>
Importer	8-10	2-3 hour/day	10-15 day/year
Distributor		2-3 hour/day	<50 day/year
Manufacturing worker	<35		
Maintenance worker	<230	<1 hour/time	

Exposure Details

Inhalation exposure is expected to be negligible because the product containing the notified chemical is highly viscous and therefore has reduced potential to generate aerosols. In addition, the notified chemical has a very low vapour pressure, so vapour accumulation in the workplace air is unlikely. Eye contact is possible but also unlikely due to the high viscous nature of greases. Thus, dermal contact would be the main route for occupational exposure.

Importation and distribution

The formulated greases containing <30% notified chemical will be imported from overseas and distributed to end users including car and engine manufacturers, mining sites, engineering sites and maintenance workshops. Transport, storage and distribution of the lubricants should involve little exposure to the notified chemical, except in the case of an accidental spill.

End users

At the automotive, machinery and equipment manufacturing sites, the grease will be applied automatically by automatic drip feeding systems and central automated grease delivery systems. Occupational exposure during the automatic operation is expected to be negligible. Possible occupational exposure may occur when opening the imported containers, adding the greases into storage containers, and during equipment cleaning up and maintenance. These operations generally

will last for a short period of time, and dermal contamination would be the main route of occupational exposure.

Maintenance fitters and other mechanics at maintenance workshops will apply the grease manually by brush, spatula and grease gun. Dermal exposure may occur during these manual operations. However, the exposure is expected to be infrequent (monthly or yearly) and has a short duration (<1 hour), as these products are designed to be long term lubricants.

The notifier indicated that workers will wear impermeable gloves, protective eyewear, protective clothing, and safety boots when using greases repeatedly or for prolonged periods.

5.4. Release

RELEASE OF CHEMICAL FROM USE

The notifier expects that the majority of the grease containing the notified chemical will be collected and disposed of by incineration. The notified chemical in wastes resulting from spillage and residual lubricant in import container liners and discarded machinery will be disposed of in landfill. Therefore, eventually the entire import volume is expected to need disposal.

5.5. Disposal

The notifier indicates that the majority of the grease containing the notified chemical will be collected and disposed of by incineration. Minor spills and residual lubricant in import drum liners and discarded machinery will be disposed of to landfill.

5.6. Public exposure

Significant public exposure to the notified chemical during importation, transportation and storage is unlikely, except in the event of an accidental spill. In the event of an accidental spillage, spills should be collected and placed in suitable containers for disposal. All waste materials should be disposed of in accordance with State regulations for the disposal of oil and grease by licensed oil waste and registered waste drum companies where the product will be either recycled or incinerated. Lubricants containing the notified chemical will not be available to the public for domestic use, and public exposure through the intended industrial use is expected to be negligible.

6. PHYSICAL AND CHEMICAL PROPERTIES

Appearance at 20°C and 101.3 kPa	White powder
Melting Point	270°C (decomposition)
Remarks	The melting point of the notified chemical was determined according to EC Directive 92/69/EEC A.1 Melting Freezing Temperature and OECD TG 102, Melting point/ Melting Range.
TEST FACILITY	Institut Für Biologische Analytik und Consulting IBACON GmbH (1999a).
Density	1219 kg/m ³ at 20°C
Remarks	The melting point of the notified chemical was determined according to EC Directive 92/69/EEC A.3 Relative Density and OECD TG 109, Density of liquids and Solids.
TEST FACILITY	Institut Für Biologische Analytik und Consulting IBACON GmbH (1999b).
Vapour Pressure	< 8 × 10 ⁻⁸ Pa at 25°C.
METHOD	OECD TG 104 Vapour Pressure. EC Directive 92/69/EEC A.4 Vapour Pressure.
Remarks	The vapour pressure was estimated using the lowest boiling point component of the notified chemical mixture. A boiling point of 331°C was calculated for the free sebacic acid component utilising Meissner's method. The vapour pressure of this component was calculated to be 1 × 10 ⁻⁴ Pa. However, the corresponding metal salt exhibited a lower vapour pressure than the free acid. The vapour pressure was

	estimated to be less than 8×10^{-8} Pa based on the calculated boiling point and using the Modified Watson Correlation.
TEST FACILITY	Institut Für Biologische Analytik und Consulting IBACON GmbH (1999c).
Water Solubility	219 DOC (Dissolved Organic Carbon) mg/L at 20°C
METHOD	OECD TG 105 Water Solubility.
Remarks	The notifier indicates that the solubility of notified chemical was initially determined by visual assessment. The notified chemical (0.1 g) was added to deionised water in a 10 mL glass tube. After each addition the mixture was mixed vigorously and then checked for undissolved particles. The water solubility of the notified chemical determined by visual assessment was less than 1 g/L. The water solubility was also determined by a simplified flask method. Saturated solutions were prepared by mixing an excess of the notified chemical in deionised water for 24, 48 and 72 h at 30°C followed by incubation for a further 24 h at 20°C. The resulting suspensions were filtered and concentration of the notified chemical was determined by analysis for dissolved organic carbon. The result obtained above for the water solubility can presumably be mainly attributed to Component I of the mixture.
TEST FACILITY	Institut Für Biologische Analytik und Consulting IBACON GmbH (1999d).
Hydrolysis as a Function of pH	Not determined
REMARKS	The notifier indicates that given its relatively low water solubility it was not possible to determine the notified chemical's propensity to hydrolyse. The notified chemical does contain amide groups that will not hydrolyse in the environmental pH range of 4-9 due to low solubilities of these components.
Partition Coefficient (n-octanol/water)	log Pow = 0.9 (Component I) log Pow = 9.2 (Component II) log Pow = 18 (Component III)
Remarks	The notifier indicated that the notified chemical's solubilities in both water and octanol were so low that experimental determination of the partition coefficient was not possible. Therefore, the partition coefficient was determined by theoretical fragmentation of the molecule according to the Leo-Hansch method.
TEST FACILITY	Institut Für Biologische Analytik und Consulting IBACON GmbH (1999e).
Adsorption/Desorption	log K _{oc} = 1.9 (Component I) log K _{oc} = 6.3 (Component II) log K _{oc} = 11.2 (Component III)
Remarks	Estimated through Quantitative Structure Activity Relationships (QSAR) using the relationships $\log_{10} K_{oc} = 0.544 \log P_{ow} + 1.377$ (based on partition coefficient). The adsorption/desorption coefficient indicates that components II and III of the notified chemical will be immobile in soil. However, Component I is expected to exhibit higher mobility in soil.
TEST FACILITY	RCC Ltd, Itingen, Switzerland (2001a).
Dissociation Constant	pKa = 4.9 (Aliphatic carboxylic acid) pKa = 22.7 (Amide)
Remarks	The dissociation constant was estimated based on applying a free energy relationship based on Taft and Hammett correlations. The carboxylic acid component of the notified chemical will remain fully dissociated while the amide component is unlikely to dissociate in the environmental pH range of 4-9.
TEST FACILITY	RCC Ltd, Itingen, Switzerland (2001b).
Particle Size	Not determined.
Remarks	The notified chemical is synthesised in situ in base oils and will not be isolated

from the grease.

Flash Point

Not determined.

Remarks

The notified chemical is of a very low vapour pressure.

Flammability Limits

Not highly flammable.

METHOD

EC Directive 92/69/EEC A.10 Flammability (Solids).

Remarks

In contact with the ignition source, the notified chemical glowed and coloured black. However, the test substance could not sustain a burning reaction.

TEST FACILITY

Institut Für Biologische Analytik und Consulting IBACON GmbH (1999f).

Autoignition Temperature

315°C

METHOD

92/69/EEC A.16 Relative Self-Ignition Temperature for Solids.

Remarks

The notified chemical shows two exothermic reactions. The first one started at about 226°C with maximum temperature of about 377°C. The second exothermic heat effect started at about 301°C with a maximum temperature of about 440°C. At the end of the run, the notified chemical showed a loss of mass of about 62% and coloured grey.

TEST FACILITY

Institut Für Biologische Analytik und Consulting IBACON GmbH (1999g).

Explosive Properties

Not determined.

Remarks

The notified chemical does not contain any chemical unstable or highly energetic groups that might lead to an explosion.

Reactivity

Not determined.

Remarks

The notified chemical does not contain any chemical groups that indicate it might act as oxidising agent.

Surface Tension

70 mN/m at 20°C

METHOD

OECD TG 115 Surface Tension of Aqueous Solutions.

Remarks

EC Directive 92/69/EEC A.5 Surface Tension.

The surface tension of the notified chemical was determined on a 90% saturation solution using a tensiometer. The notified chemical is not surface active.

TEST FACILITY

Institut Für Biologische Analytik und Consulting IBACON GmbH (1999h).

Physical Properties of the grease products containing the notified chemical

The density of the formulated grease products containing <30% notified chemical is in the range of 800 to 990 kg/m³. The base oil viscosity at 40°C may range from 20 to 130 mm²/s, and the drop points are between 170 and >240°C.

7. TOXICOLOGICAL INVESTIGATIONS

<i>Endpoint and Result</i>	<i>Assessment Conclusion</i>
Rat, acute oral LD50 > 2 000 mg/kg bw	low toxicity
Rat, acute dermal LD50 > 2 000 mg/kg bw	low toxicity
Rat, acute inhalation	No toxicity data were submitted
Rabbit, skin irritation	non-irritating
Rabbit, eye irritation	slightly irritating
Guinea pig, skin sensitisation - adjuvant test.	no evidence of sensitisation.
Rat, oral repeat dose toxicity - 28 days.	NOAEL = 150 mg/kg/day
Genotoxicity - bacterial reverse mutation	non mutagenic
Genotoxicity - in vitro chromosome aberration	non genotoxic
Genotoxicity - in vivo	No toxicity data were submitted

7.1. Acute toxicity – oral

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 423 Acute Oral Toxicity – Acute Toxic Class Method.
Species/Strain	Rat/Wistar
Vehicle	1% carboxymethylcellulose
Remarks - Method	GLP & QA.

RESULTS

<i>Group</i>	<i>Number and Sex of Animals</i>	<i>Dose mg/kg bw</i>	<i>Mortality</i>
1	3 males	2 000	0
2	3 females	2 000	0

LD50	> 2 000 mg/kg bw
Signs of Toxicity	None.
Effects in Organs	None.
Remarks - Results	None.

CONCLUSION The notified chemical is of low toxicity via the oral route.

TEST FACILITY Bioservice Scientific Laboratories GmbH (1999a).

7.2. Acute toxicity - dermal

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 402 Acute Dermal Toxicity – Limit Test. EC Directive 92/69/EEC B.3 Acute Toxicity (Dermal) – Limit Test.
Species/Strain	Rat/Wistar
Vehicle	1% carboxymethylcellulose
Type of dressing	Semi-occlusive.
Remarks - Method	GLP & QA.

RESULTS

<i>Group</i>	<i>Number and Sex of Animals</i>	<i>Dose mg/kg bw</i>	<i>Mortality</i>
1	5/sex	2 000	0

LD50	> 2 000 mg/kg bw
Signs of Toxicity - Local	None.
Signs of Toxicity - Systemic	None.
Effects in Organs	None.
Remarks - Results	No skin irritation observed.

CONCLUSION The notified chemical is of low toxicity via the dermal route.

TEST FACILITY Bioservice Scientific Laboratories GmbH (1999b).

7.3. Acute toxicity - inhalation

No toxicity data were submitted.

7.4. Irritation – skin

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 404 Acute Dermal Irritation/Corrosion. EC Directive 92/69/EEC B.4 Acute Toxicity (Skin Irritation).
Species/Strain	Rabbit/New Zealand White
Number of Animals	3
Vehicle	1% carboxymethylcellulose
Observation Period	72 hours
Type of Dressing	Semi-occlusive.
Remarks - Method	GLP & QA.
RESULTS	The Draize scores for erythema/eschar and oedema were zero for all animals throughout the observation period of 72 hours after treatment.
Remarks - Results	No clinical signs of systemic toxicity were found.
CONCLUSION	The notified chemical is non-irritating to skin.
TEST FACILITY	Bioservice Scientific Laboratories GmbH (1999c).

7.5. Irritation - eye

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 405 Acute Eye Irritation/Corrosion. EC Directive 92/69/EEC B.5 Acute Toxicity (Eye Irritation).
Species/Strain	Rabbit/New Zealand White
Number of Animals	3
Observation Period	72 hours
Remarks - Method	GLP & QA.

RESULTS

<i>Lesion</i>	<i>Mean Score* Animal No.</i>			<i>Maximum Value</i>	<i>Maximum Duration of Any Effect</i>	<i>Maximum Value at End of Observation Period</i>
	1	2	3			
<i>Conjunctiva: redness</i>	0	0	0	2	1 hour	0
<i>Conjunctiva: chemosis</i>	0	0	0	0	-	0
<i>Conjunctiva: discharge</i>						
<i>Corneal opacity</i>	0	0	0	0	-	0
<i>Iridial inflammation</i>	0	0	0	0	-	0

*Calculated on the basis of the scores at 24, 48, and 72 hours for EACH animal.

Remarks - Results	Conjunctiva discharge was not observed. At 1 hour, the Draize scores of conjunctival redness for the three animals were 1, 2 and 1, respectively.
CONCLUSION	The notified chemical is slightly irritating to the eye.
TEST FACILITY	Bioservice Scientific Laboratories GmbH (1999d).

7.6. Skin sensitisation

TEST SUBSTANCE	Komplexseife TH17
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METHOD	OECD TG 406 Skin Sensitisation - Adjuvant test. EC Directive 96/54/EC B.6 Skin Sensitization - Adjuvant test.		
Species/Strain	Guinea pig/DH		
PRELIMINARY STUDY	Maximum Non-irritating Concentration: intradermal: not stated. topical: 100%		
MAIN STUDY			
Number of Animals	Test Group: 10	Control Group: 5	
induction phase	Induction Concentration: intradermal injection topical application	25% (highest applicable concentration). 100%	
Signs of Irritation	None.		
CHALLENGE PHASE			
1 st challenge	topical application: 100%		
Remarks - Method	GLP & QA.		

RESULTS

<i>Animal</i>	<i>Challenge Concentration</i>	<i>Number of Animals Showing Skin Reactions after: 1st challenge</i>		
		<i>24 h</i>	<i>48 h</i>	<i>72 h</i>
<i>Test Group</i>	100%	0/10	0/10	0/10
<i>Control Group</i>	100%	0/5	0/5	0/5

Remarks - Results	Historic data of positive controls were provided in the report.
CONCLUSION	There was no evidence of reactions indicative of skin sensitisation to the notified chemical under the conditions of the test.
TEST FACILITY	Bioservice Scientific Laboratories GmbH (1999e).

7.7. Repeat dose toxicity

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 407 Repeated Dose 28-day Oral Toxicity Study in Rodents. EC Directive 96/54/EC B.7 Repeated Dose (28 Days) Toxicity (Oral).
Species/Strain	Rat/Wistar
Route of Administration	Oral – gavage.
Exposure Information	Total exposure days: 28 days; Dose regimen: 7 days per week;
Vehicle	1% carboxymethylcellulose
Remarks - Method	GLP & QA.

RESULTS

<i>Group</i>	<i>Number and Sex of Animals</i>	<i>Dose mg/kg bw/day</i>	<i>Mortality</i>
I (control)	5/sex	0	0
II (low dose)	5/sex	30	0
III (mid dose)	5/sex	150	0
IV (high dose)	5/sex	750	0

Mortality and Time to Death
None.

Clinical Observations

A dose-dependent decrease in food consumption and bodyweight gains was observed in treated male rats. The reduced food consumption and bodyweight gain reached statistical significance in the high-dose males.

One male in the group III had diarrhoea for one day.

Laboratory Findings – Clinical Chemistry, Haematology, Urinalysis

There were no treatment-related changes in clinical chemistry, haematology and urinalysis tests.

Pathology

No abnormal findings were observed.

Effects in Organs

The mean relative spleen and adrenal values in females of the group IV were significantly lower than corresponding control group. There were no other treatment-related differences in relative and absolute organ weight for both sexes and any of the groups.

Histopathology

Histopathological examination showed that no differences in incidence or severity between control and treatment groups were considered to be of toxicological significance.

Remarks – Results

The following changes were considered to be treatment-related in the study:

- Decreases in food consumption and bodyweight gain in high-dose male rats.
- Decreases of the mean relative spleen and adrenal weights in high-dose female rats.

CONCLUSION

The No Observed Adverse Effect Level (NOAEL) was established as 150 mg/kg bw/day in this study, based on the decreases of bodyweight gain and food consumption in high-dose male rats and the mean relative spleen and adrenal values in high-dose female rats.

TEST FACILITY

Bioservice Scientific Laboratories GmbH (1999f).

7.8. Genotoxicity - bacteria

TEST SUBSTANCE

Komplexseife TH17

METHOD

OECD TG 471 Bacterial Reverse Mutation Test.

Plate incorporation procedure & Pre incubation procedure

Species/Strain

S. typhimurium: TA1535, TA1537, TA98, TA100, TA102

Metabolic Activation System

S9-mix

Concentration Range in

a) With metabolic activation: 0 - 5 000 µg/plate.

Main Test

b) Without metabolic activation: 0 - 5 000 µg/plate.

Vehicle

DMSO

Remarks - Method

GLP & QA.

Experiment I was an incorporation test, and experiment II was a pre-incubation test.

RESULTS

<i>Metabolic Activation</i>	<i>Test Substance Concentration (µg/plate) Resulting in:</i>			
	<i>Cytotoxicity in Preliminary Test</i>	<i>Cytotoxicity in Main Test</i>	<i>Precipitation</i>	<i>Genotoxic Effect</i>
<i>Absent</i>	>5 000		≥1 000	
Test 1		Not seen	≥1 000	Not seen
Test 2		Not seen	≥1 000	Not seen

<i>Present</i>	>5 000		≥1 000	
Test 1		Not seen	≥1 000	Not seen
Test 2		Not seen	≥1 000	Not seen

Remarks - Results	<p>No significant increases in revertant colony numbers of any tested strains were observed following treatment with the notified chemical either in the presence or absence of metabolic activation in both incorporation test and pre-incubation test.</p> <p>The positive controls induced a distinct increase of induced revertant colonies.</p>
CONCLUSION	The notified chemical was not mutagenic to bacteria under the conditions of the test.
TEST FACILITY	Bioservice Scientific Laboratories GmbH (1999g).

7.9. Genotoxicity – in vitro

TEST SUBSTANCE	Komplexseife TH17
METHOD	OECD TG 473 In vitro Mammalian Chromosomal Aberration Test.
Cell Type/Cell Line	Chinese hamster V79 cells
Metabolic Activation System	S-9 mix
Vehicle	DMSO
Remarks - Method	GLP & QA.

<i>Metabolic Activation</i>	<i>Test Substance Concentration (µg/mL)</i>	<i>Exposure Period</i>	<i>Harvest Time</i>
<i>Absent</i>			
Test 1	0, 5*, 50* and 100*	4 h	20 h
Test 2	0, 5*, 50* and 100*	20 h	20 h
<i>Present</i>			
Test 1	0, 5*, 50* and 100*	4 h	20 h

*Cultures selected for metaphase analysis.

RESULTS

<i>Metabolic Activation</i>	<i>Test Substance Concentration (µg/mL) Resulting in:</i>			
	<i>Cytotoxicity in Preliminary Test</i>	<i>Cytotoxicity in Main Test</i>	<i>Precipitation</i>	<i>Genotoxic Effect</i>
<i>Absent</i>				
Test 1		Not seen	≥5	Not seen
Test 2		Not seen	≥5	Not seen
<i>Present</i>				
Test 1		Not seen	≥5	Not seen

Remarks - Results	<p>The notified chemical did not increase the frequency of aberration in Chinese V79 cells in the presence and absence of metabolic activation.</p> <p>The positive controls induced a significant increase of cells with structural chromosome aberrations above test laboratory's historic control level.</p> <p>The study was hampered by the low solubility of the test substance in DMSO and the culture medium.</p>
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CONCLUSION The notified chemical was not clastogenic to Chinese hamster V79 treated in vitro under the conditions of the test.

TEST FACILITY Bioservice Scientific Laboratories GmbH (1999h).

7.10. Genotoxicity – in vivo

No toxicity data were submitted.

8. ENVIRONMENT

8.1. Environmental fate

8.1.1. Ready biodegradability

TEST SUBSTANCE Complex Soap TH17

METHOD OECD TG 301 F Ready Biodegradability: Manometric Respirometry Test.

Exposure Period 28 days

Remarks - Method Activated sludge was mixed with the test substance to give final concentration of 20 mg/L and with the standard material, aniline at a concentration of 25.3 mg/L. The study was carried out in darkness at 22°C.

RESULTS

<i>Test substance</i>		<i>Aniline</i>	
<i>Day</i>	<i>Mean % degradation</i>	<i>Day</i>	<i>% degradation</i>
14	18.5	14	72.0
28	16.8	28	96.0

Remarks - Results The aniline standard attained 96% biodegradation after 28 days, indicating the test conditions were valid. After 28 days, the mean biodegradation of the test substance based on ThOD_{NH4} was 16.8% and based on ThOD_{NO3} was 16.2%. Results from the toxicity control indicate that the notified chemical does not have an inhibitory effect on activated sludge micro-organisms.

CONCLUSION The notified chemical is not considered to be readily biodegradable under the conditions of OECD TG 301F.

TEST FACILITY Institut Für Biologische Analytik und Consulting IBACON GmbH (1999i).

8.1.2. Bioaccumulation

Data on the bioaccumulation potential of the notified chemical were not provided for this notification. Due to low aquatic exposure the notified chemical it is unlikely to bioaccumulate (Connell 1990).

8.2. Ecotoxicological investigations

8.2.1. Acute toxicity to fish

TEST SUBSTANCE Complex Soap TH17

METHOD OECD TG 203 Fish, Acute Toxicity Test

Species Zebra Fish (*Brachydanio rerio*)
 Exposure Period 96 h
 Water Hardness 250 mg CaCO₃/L
 Analytical Monitoring Test solutions were not measured.

RESULTS

Concentration mg/L Nominal	Number of Fish	Mortality				
		2 h	24 h	48 h	72 h	96 h
0	7	0	0	0	0	0
100	7	0	0	0	0	0

LC50 Not determined
 NOEC (or LOEC) Not determined
 Remarks – Results The definitive studies were conducted on the filtrate of a super saturated stock suspension of the notified chemical at a nominal concentration of 100 mg/L. The results of the definitive study showed that no mortalities were observed at this test substance concentration. A 96-hour EC₅₀ for the notified chemical to *Brachydanio rerio* was not determined but the 96 h NOEC is expected to be greater than the limit of its solubility.

CONCLUSION The ecotoxicity data indicates the notified chemical is not toxic to fish up to the limit of its solubility.

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8.2.2. Acute/chronic toxicity to aquatic invertebrates

TEST SUBSTANCE Complex Soap TH17
 METHOD OECD TG 202 Daphnia sp. Acute Immobilisation Test
 Species *Daphnia magna*
 Exposure Period 48 hours
 Water Hardness 250 mg CaCO₃/L
 Analytical Monitoring Test solutions were not measured.

RESULTS

Concentration mg/L Nominal	Number of <i>D. magna</i>	Number Immobilised	
		24 h	48 h
0	20	0	0
100	20	0	0

LC50 Not determined
 NOEC (or LOEC) Not determined
 Remarks - Results The definitive studies were conducted on the filtrate of a super saturated stock suspension of the notified chemical at a nominal concentration of 100 mg/L. The immobilisation tests with daphnia were performed in duplicate using 10 daphnids per flask with observations performed at 24 and 48 hours. A 48-hour EC₅₀ for the notified chemical to *Daphnia magna* was not determined but the 48 h NOEC is expected to be greater than the limit of its solubility.

CONCLUSION The ecotoxicity data indicates the notified chemical is not toxic to daphnia up to the limit of its solubility.

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8.2.3. Algal growth inhibition test

TEST SUBSTANCE	Complex Soap TH17
METHOD	OECD TG 201 Alga, Growth Inhibition Test.
Species	<i>Scenedesmus suspicatus</i>
Exposure Period	72 hours
Concentration Range	6.25, 12.5, 25, 50 and 100 mg/L
Nominal	
Water Hardness	24 mg CaCO ₃ /L
Analytical Monitoring	Test solutions were not measured

RESULTS

<i>Biomass</i>		<i>Growth</i>	
<i>E_bC50 (mg/L at 72 h)</i>	<i>NOEC (mg/L)</i>	<i>E_rC50 (mg/L at 72 h)</i>	<i>NOEC (mg/L)</i>
85.6	50	> 100	25

Remarks - Results Algae were exposed to the filtrate of a super saturated stock suspension of the notified chemical at a nominal concentration of 100 mg/L under constant illumination and shaking. After 72 h, there was no significant inhibition of algal growth and biomass at the nominal concentrations of 6.25, 12.5 and 25 mg/L. At a nominal concentration of 50 and 100 mg/L both algal growth and biomass were significantly reduced.

CONCLUSION The ecotoxicity data indicates the notified chemical shows some toxicity to algae below the limit of its solubility.

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8.2.4. Inhibition of microbial activity

TEST SUBSTANCE	Complex Soap TH17
METHOD	OECD TG 209 Activated Sludge, Respiration Inhibition Test.
Inoculum	Activated sludge
Exposure Period	3 hours
Concentration Range	10, 32, 100, 320, 1000 mg/L
Nominal	

RESULTS The activated sludge study was conducted using sludge obtained from sewage treatment plant in Groß-Zimmern, Germany. The definitive study was conducted on nominal concentrations of 10, 32, 100, 320 and 1000 mg/L. Amounts of test material (5, 16, 50, 160 and 500 mg) were added to water (284 mL) and sewerage (16 mL) and samples were stirred continuously. The reference material used in the study was 3,5-dichlorophenol. When compared to the control, activated sludge after 3 h experienced differences in respiration of between -8.5 to -18.6%. The 3-hour EC₅₀ for the notified substance to activated sludge could not be quantified. However, the 3-hour EC₅₀ for the notified substance to activated sludge is expected to be greater than 1000 mg/L. The EC₅₀ of the reference substance was 9 mg/L, therefore confirming the suitability of the activated sludge.

EC₅₀ > 1000 mg/L
NOEC Not determined

CONCLUSION The ecotoxicity data indicates the notified chemical is not toxic to activated sludge up to 1000 mg/L in suspension.

TEST FACILITY

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(1999m).

9. RISK ASSESSMENT

9.1. Environment

9.1.1. Environment – exposure assessment

The notified chemical will be used as a component of ready-to-use grease used at industrial sites throughout Australia. As waste, the majority of the import volume will be collected and disposed of by incineration, which will result in the formation of water vapour and oxides of carbon and nitrogen and metal salts in the sludge. A small amount will be discarded in landfill through the disposal of plastic drum liners and machinery to which the grease containing the notified chemical has been applied.

Although it is not considered to be readily biodegradable, the notified chemical is expected to biodegrade to a certain extent in landfill. The high octanol-water partition coefficient calculated for Component's II and III of the notified chemical mixture and the expected low water solubilities indicate that they will partition to soil and sediment and be immobile in the environment. The third component of the mixture, Component I, is moderately soluble, has a low partition coefficient and is potentially mobile in soil. However, as a consequence of its anionic nature it is expected to associate with metal ions on the surface of soil and be immobile.

9.1.2. Environment – effects assessment

The notified chemical is not toxic to fish, daphnia and micro-organisms up to the limit of its solubility. However, it shows some toxicity to algae below this limit. Bioaccumulation is not expected due to the notified chemical's limited exposure to the aquatic compartment.

9.1.3. Environment – risk characterisation

The notified chemical will be used as a component of ready-to-use grease, and most of it is expected to eventually be collected and incinerated, which will result in the formation of water vapour and oxides of carbon and nitrogen. A small amount will be discarded in landfill primarily through the disposal of plastic drum liners. Here, given it is not readily biodegradable (< 20% over 28 days), has a relatively high partition coefficients and low water solubility, the notified chemical would associate with soil and sediment and slowly degrade over time.

The above considerations indicate minimal hazard to the environment when the notified chemical is used in the manner and levels indicated by the notifier.

9.2. Human health

9.2.1. Occupational health and safety – exposure assessment

The notified chemical will be imported in pails and drums as a component (<30%) of ready-to-use greases. The exposure for importation and distribution workers is expected to be negligible except in the event that the packaging is breached.

More than half of the notified chemical will be used at the manufacturing sites. Minimal exposure is expected because the systems for applying the greases are generally enclosed and automated. However, the possibility of exposure to drips and spills exists during the processes of preparation, cleaning and maintenance. Dermal exposure would be the predominant route of occupational exposure to workers during these activities.

Fitters or other mechanics at the maintenance sites will apply the greases manually to existing machinery by brush, spatula, grease gun or grease cartridge. Dermal exposure may occur. However, the exposure is considered to be of short duration and intermittent.

9.2.2. Public health – exposure assessment

Exposure of the general public to the notified chemical as a result of transport or through environmental release is assessed as being negligible.

9.2.3. Human health - effects assessment

Acute toxicity.

The notified chemical was of low oral and dermal toxicity in acute rat studies. No acute inhalation study data were submitted.

Irritation and Sensitisation.

In the eye irritation study, two rabbits had Draize scores of 1 and one had Draize score of 2 of conjunctival redness at 1 hour after treatment. All other scores were zero. Thus, the notified chemical is considered to be a slight eye irritant.

The notified chemical was non-irritant to rabbit skin and negative to guinea-pigs in a skin sensitisation adjuvant test.

Repeated Dose Toxicity (sub acute, sub chronic, chronic).

In the 28-day oral repeat study, decreases in food consumption and bodyweight gains in high-dose male rats, and decreases of the mean relative spleen and adrenal values in high-dose female rats were considered to be treatment-related. No significant histopathological changes were observed. The No Observed Adverse Effect Level (NOAEL) was considered to be 150 mg/kg bw/day in this study.

Genotoxicity

The notified chemical was negative in an Ames test and found not to be clastogenic in a chromosomal aberration study in Chinese hamster V79 cells. However, the latter study was hampered by poor solubility, so the concentrations tested were lower than usual for this type of test. No in vivo genotoxicity study data were submitted.

Hazard classification for health effects.

Based on the available toxicological data, the notified chemical would not be classified as hazardous under the NOHSC *Approved Criteria for Classifying Hazardous Substances* (NOHSC, 1999). However, the notified chemical is a soluble barium salt, which is on the NOHSC List of Designated Hazardous Substances, with the risk phrases R20/22 (Harmful by inhalation and if swallowed). Therefore, the notified chemical must be classified as a hazardous substance.

9.2.4. Occupational health and safety – risk characterisation

Dermal exposure would be the predominant route of occupational exposure to the notified chemical. Although the notified chemical is neither a skin irritant nor a skin sensitiser, repeated or prolonged skin contact with lubricant and grease products should be avoided since human experience has shown that prolonged skin contact with lubricant or grease products may cause skin irritation and/or dermatitis (oil acne or folliculitis).

The health risk for importation and distribution workers is expected to be negligible except in the event that the packaging is breached.

At sites manufacturing new machinery, minimal exposure is expected because the systems are enclosed and automated. However, gloves, eyewear and protective clothing should be worn when the possibility of exposure to drips and spills exists during the processes of preparation, cleaning and maintenance. With the application of personal protective equipment, the risk of adverse health effects at the manufacturing sites is expected to be low.

Fitters or other mechanics at maintenance sites will apply the greases manually to existing machinery. Adverse skin effects may ensue if dermal contact is repeated or prolonged. It is recommended that the workers wear gloves, eyewear and protective clothing to minimise the risk of adverse skin effects from the greases.

In the case that any oil mist or aerosol of the notified chemical will be generated at workplaces, the NOHSC Exposure Standards for oil mist (5 mg/m³, TWA) and soluble barium compounds

(0.5 mg/m³, TWA) must be applied (NOHSC, 1995).

9.2.5. Public health – risk characterisation

Given that the notified chemical will be a component in grease intended for industrial use only and it has low acute oral and dermal toxicity, the risk to public health is considered to be minimal.

10. CONCLUSIONS – ASSESSMENT LEVEL OF CONCERN FOR THE ENVIRONMENT AND HUMANS

10.1. Hazard classification

Based on the available data the notified chemical is classified as hazardous under the NOHSC *Approved Criteria for Classifying Hazardous Substances*, with the risk phrases R20/22 (Harmful by inhalation and if swallowed).

10.2. Environmental risk assessment

On the basis of the available information, the overall environmental hazard of the notified chemical is expected to be low.

10.3. Human health risk assessment

10.3.1. Occupational health and safety

There is Low Concern to occupational health and safety under the conditions of the occupational settings described.

10.3.2. Public health

There is No Significant Concern to public health when used in the proposed manner.

11. MATERIAL SAFETY DATA SHEET

11.1. Material Safety Data Sheet

The MSDS of the [products containing the chemical](#) provided by the notifier was in accordance with the NOHSC *National Code of Practice for the Preparation of Material Safety Data Sheets* (NOHSC, 1994a). It is published here as a matter of public record. The accuracy of the information on the MSDS remains the responsibility of the applicant.

11.2. Label

The labels for the [products containing the chemical](#) provided by the notifier [were](#) in accordance with the NOHSC *National Code of Practice for the Labelling of Workplace Substances* (NOHSC, 1994b). The accuracy of the information on the label remains the responsibility of the applicant.

12. RECOMMENDATIONS

CONTROL MEASURES

Occupational Health and Safety

- Employers should implement the following safe work practices to minimise occupational exposure during handling of products containing the notified chemical:
 - avoid repeated or prolonged dermal exposure.
 - avoid generation of any oil mist or aerosol of the notified chemical.
- Employers should ensure that the following personal protective equipment is used by workers to minimise occupational exposure to products containing the notified chemical:

- gloves
- safety eyewear, and
- protective clothing.

Guidance in selection of personal protective equipment can be obtained from Australian, Australian/New Zealand or other approved standards.

- A copy of the MSDS should be easily accessible to employees.
- If products and mixtures containing the notified chemical are classified as hazardous to health in accordance with the NOHSC *Approved Criteria for Classifying Hazardous Substances*, workplace practices and control procedures consistent with provisions of State and Territory hazardous substances legislation must be in operation.

Disposal

- The notified chemical should be disposed of by incineration.

Emergency procedures

- Spills/release of the notified chemical should be contained as described in the MSDS and the resulting waste disposed of by incineration.

12.1. Secondary notification

The Director of Chemicals Notification and Assessment must be notified in writing within 28 days by the notifier, other importer or manufacturer:

- (1) Under Section 64(1) of the Act; if
 - the notified chemical will be used in a manner that generation of aerosol of the notified chemical is possible.or
- (2) Under Section 64(2) of the Act:
 - if any of the circumstances listed in the subsection arise.

The Director will then decide whether secondary notification is required.

No additional secondary notification conditions are stipulated.

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