HOUSE OF KOLOR EXEMPT CATALYST KU-150

ChemWatch Material Safety Data Sheet CHEMWATCH 5090-78 Date of Issue: Mon 11-Aug-2003

STATEMENT OF HAZARDOUS NATURE

HAZARDOUS ACCORDING TO WORKSAFE AUSTRALIA CRITERIA.

SUPPLIER

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SYNONYMS

SHIPPING NAME

PAINT None

Product Name: House of Kolor Exempt Catalyst KU-150 Other Names: Product Code: KU-150 06/03

CAS RN No(s):	None	None
UN Number:	1263	
Packing Group:	III	
Dangerous Goods Class:	3	
Subsidiary Risk:	None,	None
Hazchem Code:	3[Y]	
Poisons Schedule Number:	None	

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USE

Used according to manufacturers directions.

The use of a quantity of material in an unventilated or confined space may result in increased exposure and an irritating atmosphere developing Before starting consider control of exposure by mechanical ventilation For further information refer to the House of Kolor Technical Manual

PHYSICAL DESCRIPTION/PROPERTIES

APPEARANCE

Opaque to translucent flammable liquid with an organic solvent odour; does not mix with water. Will react with hot water to release carbon dioxide.

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Boiling Point (°C):
                            139
Melting Point (°C):
                            Not Available
Vapour Pressure (kPa):
                            Not Available
Specific Gravity:
                            1.28
Flash Point (°C):
                            42.8
                                 (TAGCC)
Lower Explosive Limit (%):
                            0.9
Upper Explosive Limit (%):
                            10.5
Solubility in Water (g/L):
                            Immiscible
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INGREDIENTS

NAME	CAS RN	90 10
4-chlorobenzotrifluoride	98-56-6	30-60
hexamethylene diisocyanate polymer	28182-81-2	30-60
hexamethylene diisocyanate	822-06-0	<1

HEALTH HAZARD

ACUTE HEALTH EFFECTS

SWALLOWED

The material is not thought to produce adverse health effects following

ingestion (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least one other route and good hygiene practice requires that exposure be kept to a minimum.

EYE

The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis.

SKIN

Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. The material may produce severe skin irritation after prolonged or repeated exposure, and may produce a contact dermatitis (nonallergic). This form of dermatitis is often characterised by skin redness (erythema) thickening of the epidermis.

Histologically there may be intercellular oedema of the spongy layer (spongiosis) and intracellular oedema of the epidermis. Prolonged contact is unlikely, given the severity of response, but repeated exposures may produce severe ulceration.

INHALED

Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful.

The material may produce respiratory tract irritation. Symptoms of pulmonary irritation may include coughing, wheezing, laryngitis, shortness of breath, headache, nausea, and a burning sensation.

Unlike most organs, the lung can respond to a chemical insult or a chemical agent, by first removing or neutralising the irritant and then repairing the damage (inflammation of the lungs may be a consequence).

The repair process (which initially developed to protect mammalian lungs from foreign matter and antigens) may, however, cause further damage to the lungs (fibrosis for example) when activated by hazardous chemicals. Often, this results in an impairment of gas exchange, the primary function of the lungs. Therefore prolonged exposure to respiratory irritants may cause sustained breathing difficulties.

Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by narcosis, drowsiness, reduced alertness, loss of reflexes, lack of coordination and vertigo.

Inhalation of quantities of liquid mist may be extremely hazardous, even lethal due to spasm, extreme irritation of larynx and bronchi, chemical pneumonitis and

pulmonary oedema.

Acute effects from inhalation of high concentrations of vapour are pulmonary irritation, including coughing, with nausea; central nervous system depression - characterised by headache and dizziness, increased reaction time, fatigue and loss of co-ordination.

The vapour/mist may be highly irritating to the upper respiratory tract and lungs; the response may be severe enough to produce bronchitis and pulmonary oedema. Possible neurological symptoms arising from isocyanate exposure include headache, insomnia, euphoria, ataxia, anxiety neurosis, depression and paranoia. Gastrointestinal disturbances are characterised by nausea and vomiting. Pulmonary sensitisation may produce asthmatic reactions ranging from minor breathing difficulties to severe allergic attacks; this may occur following a single acute exposure or may develop without warning for several hours after exposure. Sensitized people can react to very low doses, and should not be allowed to work in situations allowing exposure to this material. Continued exposure of sensitised persons may lead to possible long term respiratory impairment.

Inhalation hazard is increased at higher temperatures.

CHRONIC HEALTH EFFECTS

Persons with a history of asthma or other respiratory problems or are known to be sensitised, should not be engaged in any work involving the handling of isocyanates. [CCTRADE-Bayer, APMF].

A 90-day inhalation study in rats with polymeric MDI (6 hours/day, 5 days/week) produced moderate to severe hyperplastic inflammatory lesions in the nasal cavities and lungs at levels of 8 mg/m3 or greater.

Practical evidence shows that inhalation of the material is capable of inducing a sensitisation reaction in a substantial number of individuals at a greater frequency than would be expected from the response of a normal population. Pulmonary sensitisation, resulting in hyperactive airway dysfunction and pulmonary allergy may be accompanied by fatigue, malaise and aching. Significant symptoms of exposure may persist for extended periods, even after exposure ceases. Symptoms can be activated by a variety of nonspecific environmental stimuli such as automobile exhaust, perfumes and passive smoking. Practical experience shows that skin contact with the material is capable either of inducing a sensitisation reaction in a substantial number of individuals, and/or of producing a positive response in experimental animals.

Respiratory sensitisation may result in allergic/asthma like responses; from coughing and minor breathing difficulties to bronchitis with wheezing, gasping. Sensitisation may give severe responses to very low levels of exposure, in situations where exposure may occur.

CONTAINS free organic isocyanate. Mixing and application requires special precautions and use of personal protective gear [APMF]

FIRST AID

SWALLOWED

· Immediately give a glass of water.

• First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

EYE

If this product comes in contact with the eyes:

· Wash out immediately with fresh running water.

• Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids.

· If pain persists or recurs seek medical attention.

• Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.

SKIN

If skin contact occurs:

- · Immediately remove all contaminated clothing, including footwear
- Flush skin and hair with running water (and soap if available).
- · Seek medical attention in event of irritation.

INHALED

· If fumes or combustion products are inhaled remove from contaminated area.

· Lay patient down. Keep warm and rested.

• Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures.

• Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary.

• Transport to hospital, or doctor.

ADVICE TO DOCTOR

Treat symptomatically.

For sub-chronic and chronic exposures to isocyanates:

• This material may be a potent pulmonary sensitiser which causes bronchospasm even in patients without prior airway hyperreactivity.

• Clinical symptoms of exposure involve mucosal irritation of respiratory and gastrointestinal tracts.

• Conjunctival irritation, skin inflammation (erythema, pain vesiculation) and gastrointestinal disturbances occur soon after exposure.

- \cdot Pulmonary symptoms include cough, burning, substernal pain and dyspnoea.
- Some cross-sensitivity occurs between different isocyanates.

• Noncardiogenic pulmonary edema and bronchospasm are the most serious consequences of exposure. Markedly symptomatic patients should receive oxygen,

ventilatory support and an intravenous line.
Treatment for asthma includes inhaled sympathomimetics (epinephrine [adrenalin], terbutaline) and steroids.
Activated charcoal (1 g/kg) and a cathartic (sorbitol, magnesium citrate) may be useful for ingestion.
Mydriatics, systemic analgesics and topical antibiotics (Sulamyd) may be used for corneal abrasions.
There is no effective therapy for sensitised workers.
[Ellenhorn and Barceloux; Medical Toxicology]
NOTE: Isocyanates cause airway restriction in naive individuals with the degree of response dependant on the concentration and duration of exposure. They induce smooth muscle contraction which leads to bronchoconstrictive episodes. Acute changes in lung function, such as decreased FEV1, may not represent sensitivity.
[Karol & Jin, Frontiers in Molecular Toxicology, pp 56-61, 1992]

PRECAUTIONS FOR USE

EXPOSURE STANDARDS

No data for House of Kolor Exempt Catalyst KU-150.

EXPOSURE STANDARDS FOR MIXTURE

"Worst Case" computer-aided prediction of vapour components/concentrations:

Composite Exposure Standard for Mixture (TWA) (mg/m³): 0.034 mg/m³ If the breathing zone concentration of ANY of the components listed below is exceeded, "Worst Case" considerations deem the individual to be overexposed. Component Breathing Zone ppm Breathing Zone mg/m³ Mixture Conc: (%)

hexamethylene diisocyanate 0.01 0.034 1

INGREDIENT DATA

4-CHLOROBENZOTRIFLUORIDE:

No exposure limits set by NOHSC or ACGIH OEL-STEL (Russia): 20 mg/m³ (skin) Exposure limits with "skin" notation indicate that vapour and liquid may be absorbed through intact skin. Absorption by skin may readily exceed vapour inhalation exposure. Symptoms for skin absorption are the same as for inhalation. Contact with eyes and mucous membranes may also contribute to overall exposure and may also invalidate the exposure standard. HEXAMETHYLENE DIISOCYANATE POLYMER: No exposure limits set by NOHSC or ACGIH HEXAMETHYLENE DIISOCYANATE: TLV TWA: 0.005 ppm [ACGIH] TLV TWA: 0.005 ppm [ACGIH] isocyanates, all as NCO (Mol.Wt: 42.00) ES TWA: 0.02 mg/m³; STEL: 0.07 mg/m³ sensitiser MEL TWA: 0.02 mg/m³; STEL: 0.07 mg/m³ sensitiser Some jurisdictions require that health surveillance be conducted on occupationally exposed workers. This should emphasise: · demography, occupational and medical history and health advice • completion of a standardised respiratory questionnaire • physical examination of the respiratory system and skin · standardised respiratory function tests such as FEV1, FVC and FEV1/FVC TLV TWA: 0.005 ppm, 0.034 mg/m³ MAK value: 0.005 ppm, 0.035 mg/m³ MAK Category I Peak Limitation: For local irritants Allows excursions of twice the MAK value for 5 minutes at a time, 8 times per shift. Designated S in List of MAK values: Danger of sensitization. MAK Group IIc: Substances with MAK Values but no pregnancy risk group classification. These are substances which have been investigated but for which no information regarding possible damage to the foetus/embryo was found. Mention calls attention to the absence of adequate data. MAK values, and categories and groups are those recommended within the Federal Republic of Germany The toxicological action of HDI is similar to that of toluene diisocyanate and and the TLV-TWA is analogous. In light of reported asthmatic/ respiratory sensitisation-like responses in HDI exposed workers, individuals who may be hypersusceptible or otherwise unusually responsive may not be adequately protected at this limit.

ENGINEERING CONTROLS

For flammable liquids and flammable gases, local exhaust ventilation or a process enclosure ventilation system may be required. Ventilation equipment should be explosion-resistant.

Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

Type of Contaminant: Air Speed: solvent, vapours, degreasing etc., 0.25-0.5 m/s (50-100 f/min.) evaporating from tank (in still air). aerosols, fumes from pouring 0.5-1 m/s (100-200 f/min.) operations, intermittent container filling, low speed conveyer transfers,

welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation) direct spray, spray painting in shallow 1-2.5 m/s (200-500 f/min.) booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)

Within each range the appropriate value depends on:

Lower end of the range 1: Room air currents minimal or favourable to capture 2: Contaminants of low toxicity or of 3: Intermittent, low production. 4: Large hood or large air mass in motion Lower end of the range 1: Disturbing room air currents 2: Contaminants of high toxicity 3: High production, heavy use 4: Small hood-local control only

Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min.) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.

PERSONAL PROTECTION

EYE

Safety glasses with side shields. Chemical goggles. Contact lenses pose a special hazard; soft lenses may absorb irritants and all lenses concentrate them. DO NOT wear contact lenses.

HANDS/FEET

Wear chemical protective gloves, eg. PVC. Wear safety footwear or safety gumboots, eg. Rubber.

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NOTE: The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact.

OTHER

Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower.

RESPIRATOR

Respiratory protection is required when ANY "Worst Case" vapour-phase concentration is exceeded (see Computer Prediction in "Exposure Standards").

Protection	Factor	(Min)	Half-Face Respirator	Full-Face Respirator
$10 \times ES$			A-AUS	-
			A-PAPR-AUS	-
50 x ES			-	A-AUS
			-	A-PAPR-AUS
100 x ES			-	A-2
			_	A-PAPR-2

^ - Full-face

The local concentration of material, quantity and conditions of use determine the type of personal protective equipment required. For further information consult site specific CHEMWATCH data (if available), or your Occupational Health and Safety Advisor.

SAFE HANDLING

STORAGE AND TRANSPORT

SUITABLE CONTAINER

Packing as supplied by manufacturer. Plastic containers may only be used if approved for flammable liquid. Check that containers are clearly labelled and free from leaks.

· For low viscosity materials (i) : Drums and jerry cans must be of the

non-removable head type. (ii) : Where a can is to be used as an inner package, the can must have a screwed enclosure.

- · For materials with a viscosity of at least 2680 cSt. (23 deg. C)
- · For manufactured product having a viscosity of at least 250 cSt. (23 deg. C)
- Manufactured product that requires stirring before use and having a viscosity of at least 20 cSt (25 deg. C)
- (i) : Removable head packaging;
- (ii) : Cans with friction closures and
- (iii) : low pressure tubes and cartridges may be used.

• Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with inner and outer packages

• In addition, where inner packagings are glass and contain liquids of packing group I there must be sufficient inert absorbent to absorb any spillage, unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.

STORAGE INCOMPATIBILITY

Avoid reaction with water, alcohols, strong bases, alkalis, metal compounds and detergent solutions. Reacts with water, may generate a large volume of foam, carbon dioxide gas (CO2) and heat. Foaming in confined space may produce pressure. Isocyanates will attack and embrittle some plastics and rubbers. Avoid cross contamination between the two liquid parts of product (kit). If two part products are mixed or allowed to mix in proportions other than manufacturer's recommendation, polymerisation with gelation and evolution of heat (exotherm) may occur. This excess heat may generate toxic vapour.

- · Avoid contamination with water, alkalies and detergent solutions.
- \cdot Material reacts with water and generates gas, pressurises containers with even drum rupture resulting.
- · DO NOT reseal container if contamination is suspected.
- · Open all containers with care.

Avoid reaction with oxidising agents

STORAGE REQUIREMENTS

- · Store in original containers in approved flammable liquid storage area.
- \cdot DO NOT store in pits, depressions, basements or areas where vapours may be trapped.
- No smoking, naked lights, heat or ignition sources.
- · Keep containers securely sealed.
- · Store away from incompatible materials in a cool, dry, well-ventilated area.
- · Protect containers against physical damage and check regularly for leaks.
- · Observe manufacturer's storing and handling recommendations.

TRANSPORTATION

Class 3 - Flammable liquids shall not be loaded in the same vehicle or packed in

the same vehicle or packed in the same freight container with:

Class 1 - Explosives;

Class 2.1 - Flammable gases (where both flammable liquids and flammable gases are in bulk);

- Class 2.3 Poisonous gases;
- Class 4.2 Spontaneously combustible substances;
- Class 5.1 Oxidising agents;
- Class 5.2 Organic peroxides;
- Class 7 Radioactive substances.

SPILLS AND DISPOSAL

MINOR SPILLS

- Remove all ignition sources.
- · Clean up all spills immediately.
- · Avoid breathing vapours and contact with skin and eyes.
- · Control personal contact by using protective equipment.
- Contain and absorb small quantities with vermiculite or other absorbent material.
- · Wipe up.
- · Collect residues in a flammable waste container.

MAJOR SPILLS

- · Clear area of personnel and move upwind.
- · Alert Fire Brigade and tell them location and nature of hazard.
- · May be violently or explosively reactive.
- · Wear breathing apparatus plus protective gloves.
- Prevent, by any means available, spillage from entering drains or water course.
- · No smoking, naked lights or ignition sources.
- · Increase ventilation.
- · Stop leak if safe to do so.
- \cdot Water spray or fog may be used to disperse / absorb vapour.
- \cdot Contain spill with sand, earth or vermiculite.
- · Use only spark-free shovels and explosion proof equipment.
- \cdot Collect recoverable product into labelled containers for recycling.
- · Absorb remaining product with sand, earth or vermiculite.
- \cdot Collect solid residues and seal in labelled drums for disposal.
- \cdot Wash area and prevent runoff into drains.
- \cdot If contamination of drains or waterways occurs, advise emergency services.

DISPOSAL

- · Recycle wherever possible.
- Consult manufacturer for recycling options or consult local or regional waste

management authority for disposal if no suitable treatment or disposal facility can be identified.

• Dispose of by: Burial in a licenced land-fill or Incineration in a licenced apparatus (after admixture with suitable combustible material)

• Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

Puncture containers to prevent re-use and bury at an authorised landfill.

FIRE FIGHTERS' REPORT

EXTINGUISHING MEDIA

Foam. Dry chemical powder. BCF (where regulations permit). Carbon dioxide. Water spray or fog - Large fires only.

FIRE FIGHTING

· Alert Fire Brigade and tell them location and nature of hazard.

- · May be violently or explosively reactive.
- · Wear breathing apparatus plus protective gloves.
- Prevent, by any means available, spillage from entering drains or water course.
- · If safe, switch off electrical equipment until vapour fire hazard removed.
- · Use water delivered as a fine spray to control fire and cool adjacent area.
- · Avoid spraying water onto liquid pools.
- DO NOT approach containers suspected to be hot.
- \cdot Cool fire exposed containers with water spray from a protected location.
- · If safe to do so, remove containers from path of fire.
- When any large container (including road and rail tankers) is involved in a fire, consider evacuation by 500 metres in all directions.

FIRE/EXPLOSION HAZARD

- · Liquid and vapour are flammable.
- · Moderate fire hazard when exposed to heat or flame.
- · Vapour forms an explosive mixture with air.
- Moderate explosion hazard when exposed to heat or flame.
- · Vapour may travel a considerable distance to source of ignition.
- Heating may cause expansion or decomposition leading to violent rupture of containers.
- \cdot On combustion, may emit toxic fumes of carbon monoxide (CO).
- Combustion products include.
- carbon dioxide (CO2).

isocyanates.
and minor amounts of.
hydrogen cyanide.
hydrogen chloride.
phosgene.
nitrogen oxides (NOx).
other pyrolysis products typical of burning organic material.
Flooding quantities of water only.
Combustible.
Moderate fire hazard when exposed to heat or flame.
When heated to high temperatures decomposes rapidly generating vapour which pressures and may then rupture containers with release of flammable and highly toxic isocyanate vapour.
Burns with acrid black smoke and poisonous fumes.

Combustion yields traces of highly toxic hydrogen cyanide HCN, plus toxic nitrogen oxides NOx and carbon monoxide.

 Small quantities of water in contact with hot liquid may react violently with generation of a large volume of rapidly expanding hot sticky semi-solid foam.

· Presents additional hazard when fire fighting in a confined space.

· Cooling with flooding quantities of water reduces this risk.

FIRE INCOMPATIBILITY

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result

HAZCHEM

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CONTACT POINT

COMPANY CONTACT (+61 2) 9737 9422

AUSTRALIAN POISONS INFORMATION CENTRE 24 HOUR SERVICE: 13 11 26 POLICE, FIRE BRIGADE OR AMBULANCE: 000

NEW ZEALAND POISONS INFORMATION CENTRE 24 HOUR SERVICE: 0800 764 766 NZ EMERGENCY SERVICES: 111

End of Report

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