

US EPA ARCHIVE DOCUMENT

TECHNICAL SUPPORT SECTION TOXICITY REVIEW - I

Disinfectants Branch

Reviewed by	<u>James E. Wilson, Jr.</u>	IN <u>12/03/84</u>	OUT <u>01/14/85</u>
			Date <u>01/10/85</u>
EPA Reg. No. or File Symbol	<u>21164-L,A</u>		
EPA Petition or EUP No.	<u>None</u>		
Date Division Received	<u>11/29/84</u>		
Type Product(s):	<u>I, (D,) H, F, N, R, S</u>		
Data Accession No(s)	<u>255801, 255802</u>		
Product Mgr. No.	<u>32 (Castillo)</u>		
Product Name(s)	<u>AKta Klor 80, AKta Klor 25</u>		
Company Name (s)	<u>Rio Linda Chemical Company, Inc.</u>		
Submission Purpose	<u>Resubmission - Tox. Data</u>		
Chemical & Formulation	<u>80 - Powder/25 - liquid</u>		

Active Ingredient (s):

Sodium chlorite

8
80 and 25

1/3

BACKGROUND

Acute oral studies were required in a previous study since all of the test animals died at the 5.0 g/kg level where the 80% formulation was tested. Data on both the 25 and 80% formulations are submitted in this package.

RECOMMENDATION

Based on the data submitted the 80% formulations and the 25% should be placed in toxicity categories 2 and 3 respectively for oral toxicity.

LABELING

The statement "Harmful if swallowed" on the 80% label should be expanded to read "Harmful or fatal if swallowed". The changes recommended in the previous safety review still apply.

CRP STATUS

Special packaging is not required based on the area of intended use (industrial).

DATA REVIEW

Reports by Northview Pacific Laboratories, Inc., submitted to Rio Linda Chemical Company, Inc., Rio Linda, CA 95673. (Accession Nos. 255801, 255802)

Acute Oral (AKTA KLOR 80)

Report dated November 15, 1984.

Method - After range finding doses of 50-750 mg/kg were administered to rats, doses of 315 and 355 mg/kg were administered to 5 male and 5 female rats in each group. Doses of 180 and 280 mg/kg were administered to 6 and 8 male and female rats per group respectively. All animals were weighed on the day of dosing and weekly thereafter. After the 14 day observation period all animals were given a gross necropsy examination.

Results - The mortality produced appears in the chart below:

Dose (mg/kg)	Mortality (M:F)
180	1/6:1/6
280	6/8:6/8
315	3/5:3/5
355	5/5:4/5

The signs recorded were bleeding from the nose and mouth, lethargy, paleness, diarrhea and scruffy coat. Weight and appetite losses were noted. Animals that died had discolored lungs and intestines darken adrenals, kidneys and livers and stomachs filled with fluid; reddened lungs were found in survivors.

Conclusion - The calculated LD₅₀s are 225 and 270 mg/kg in male and female rats respectively.

Acute Oral (AKTA KLOR 25)

Report dated November 12, 1984.

Method - The method used in the previously was also used in this study. Males were administered doses of 750, 795, 890, 1000 and 1500 mg/kg; females received doses of 600, 650, 710, 795 and 890 mg/kg.

Results - The mortality is presented in the chart below:

Dose (mg/kg)	Mortality (M:F)
600	--:2/6
650	--:0/5
710	--:4/5
750	0/5:--
795	2/5:5/5
890	3/5:6/6
1000	5/6:--
1500	6/6:--

The signs and necropsy finding were essentially the same as those found in the preceding study.

Conclusion - The LD₅₀s are 820 and 640 mg/kg for male and female rats respectively.