

RDCoberly:ccw
8/1/69

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PP# 8E0630 citrus
9F0783
BORAX

003527

- Acute Rat Oral (Boric Acid 100%) : LD₅₀ = 3.16 gm/KG

- Acute Rat Oral (Borax 10 Mol) : LD₅₀ = 5.56 gm/KG

- Acute Rat Oral (Borax) : Only summaries available
Male rat LD₅₀ = 6.08 gm/KG
Male rat LD₅₀ = 4.55 gm/KG
Female rat LD₅₀ = 4.98 gm/KG

- Acute Dog Oral (Borax) : LD₅₀ = 6.15 gm/KG only summary available.

- Acute Cavy Oral (Borax) : LD₅₀ = 5.33 gm/KG only summary available.

- Acute Rat Oral (Boric Acid) : LD₅₀ = 3.45 gm/KG only summary available.

- Acute Rat Oral (Boric Acid) : LD₅₀ = 4.08 for female rats. Only summary available.

- Acute Dog Oral (Boric Acid) : LD₅₀ = 3.9 gm/KG only summaries presented.

- Acute Mouse Oral (Boric Acid) : LD₅₀ = 4.1 gm/KG only summary presented.

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SUMMARY

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The toxicity of this formulation should not create an undue human hazard when usage as per the usage pattern.

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Acute Rat Oral (100%)

Groups of 5 male animals were tested per dosage level of 1.0, 2.15, 4.64, 6.81, and 10 gm/KG. The test material was given a 50% weight/volume suspension in distilled water. The animals were fasted 3 to 4 hours prior to dosing.

Results:

The LD₅₀ equals 3.16 gm/KG. The animals at the dosage levels of 1.0 and 2.15 showed no effects of the treatment. The animals at each of the higher dosage levels exhibited depression and shallow, rapid respiration. Each of the animals at the 4.64 and the surviving animals at the 6.81 gm/KG level showed at the 24 hour observation period depression, diarrhea, and pilo erection. The symptoms persisted until time of death.

Gross autopsies performed upon the animals that died showed markedly erythemic lungs and slightly congested adrenals. The stomachs were distended and contained approximately 2.5 ml of a clear liquid. The livers showed pale areas. Autopsies performed upon the surviving animals at termination of the study revealed no gross pathology.

Acute Rat Oral (Borax 10 Mol)

10 male animals were tested per dosage level and a dosage range of from 4.0 to 7.0 gm/KG. LD₅₀ equals 5.56 gm/KG. At 24 hours the majority of the animals at the 4.5, 5.0 and 5.5 gm/KG exhibited depression, diarrhea, ataxia, lacrimation, ptosis, and deep yellow urine stains. The animals at the higher dosage levels (6.0 to 7.0 gm/KG) exhibited ataxia, diarrhea,

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deep yellow urine stains, and ptosis.

Gross autopsies performed on the animals that died revealed congested lungs, congested adrenals, and distension of the stomachs and small intestines. Gross autopsies performed on the surviving animals revealed pale mottled kidneys, pale livers, and slightly congested adrenals.

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11/9/69

Ex. of 7/18/69

INTERDEPARTMENTAL COORDINATION
OF

ACTIVITIES RELATING TO PESTICIDES

Referral of Application for Registration under the
Federal Insecticide, Fungicide, and Rodenticide Act

2. FILE SYMBOL/REGISTRATION NO.

1624-OZ

3. DATE OF APPLICATION

003527

6/23/69

1. NAME & ADDRESS OF APPLICANT OR REGISTRANT

4. PRODUCT NAME

United States Borax & Chemical Corporation
275 Wilshire Boulevard
Los Angeles, California 90005

20 MULE TEAM BORAX For Control of Blue
and Green Mold Decay on Citrus

5. COMMENTS BY COORDINATING AGENCY

B-97
Boron Compounds

The following toxicology data on 100% Sodium Tetraborate Decahydrate ($\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$)
transmitted with label and formula for review and retention. See also referral
1624-OA:

U.S. Borax Research Corporation, Date: August 2, 1961, 10-Mol Borax, Project:
#20-0138-32 Acute Oral Administration, 7 pages.

U. S. Borax Research Corporation, Date: January 25, 1961, Boric Acid, Acute Oral
Administration, 6 pages.

Place the statement "Keep out of reach of
children" in prominent place on the front
panel

BEST AVAILABLE COPY

(NAME)	8. DATE	9. NAME OF AGENCY												
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RIN 5117-93

BORAX

TOX REVIEW 003527

Page 6 is not included in this copy.

Pages _____ through _____ are not included.

The material not included contains the following type of information:

- Identity of product inert ingredients.
- Identity of product impurities.
- Description of the product manufacturing process.
- Description of quality control procedures.
- Identity of the source of product ingredients.
- Sales or other commercial/financial information.
- A draft product label.
- The product confidential statement of formula.
- Information about a pending registration action.
- FIFRA registration data.
- The document is a duplicate of page(s) _____.
- The document is not responsive to the request.

The information not included is generally considered confidential by product registrants. If you have any questions, please contact the individual who prepared the response to your request.
