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SuperUser EPA/ORD/CCTE/SCDCD

- User Settings
- Logout

- Dashboard
- Substances
- Ethylenediamine

Filtered aggr_1

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View Dossiers

Validate

- Export to i6z
- Extract to dataset
- Create component PDF/RTF
- Create document PDF/RTF
- Compare
- Generate report
- Dissemination preview

Type at least 3 characters

REACH Complete
Ethylenediamine

- 1 General information
15
- 2 Classification & Labelling and PBT assessment
- 3 Manufacture, use and exposure
- 4 Physical and chemical properties
27
- 5 Environmental fate and pathways
12
- 6 Ecotoxicological information
23
- 7 Toxicological information
45
 - 080e61d2-4648-4f7c-8630-d58f23efa01b
 - 7.1 Toxicokinetics, metabolism and distribution
3

- 7.2 Acute Toxicity
6
- 7.3 Irritation / corrosion
4
- 7.4 Sensitisation
4
- 7.5 Repeated dose toxicity
6
 - 7.5.1 Repeated dose toxicity: oral
3
 - 8c1a9bb5-f907-42b7-9d03-73149581dafa
 - 6aaf8e00-2e54-4768-8ec8-14496328ffff
 - 66c427b9-34c9-4e32-a8e9-035e7b879bd8
 - 7.5.2 Repeated dose toxicity: inhalation
1
 - 7.5.3 Repeated dose toxicity: dermal
2
 - 7.5.4 Repeated dose toxicity: other routes
- 7.6 Genetic toxicity
12
- 7.7 Carcinogenicity
2
- 7.8 Toxicity to reproduction
5
- 7.9 Specific investigations
- 7.10 Exposure related observations in humans
2
- 7.11 Toxic effects on livestock and pets
- 7.12 Additional toxicological information

- 8 Analytical methods
- 9 Residues in food and feedingstuffs
- 10 Effectiveness against target organisms
- 11 Guidance on safe use
- 12 Literature search
- 13 Assessment reports
- 14 Information requirements
- Inherited templates

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1

0

Administrative data

Data source

Materials and methods

Results and discussion

Overall remarks, attachments

Applicant's summary and conclusion

Administrative data

Endpoint
sub-chronic toxicity: oral

Type of information
experimental study

Adequacy of study
supporting study

☐ Robust study summary

☐ Used for classification

☐ Used for SDS

Study period

Reliability
2 (reliable with restrictions)

Rationale for reliability incl. deficiencies

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification Reason / purpose Actions

Cross-reference

Reason / purpose for cross-reference Related information Remarks Actions

Data source

Reference

- publication | Unnamed | 1982
-

Data access

Data protection claimed

Materials and methods

Test guideline

Qualifier Guideline Version / remarks Deviations Actions 1

Qualifier
no guideline followed

Guideline

Version / remarks

Deviations

Principles of method if other than guideline

GLP compliance
not specified

Limit test
no

Test material

Test material information

- Unnamed | Unnamed | ethane-1,2-diamine | EC 203-468-6 | 107-15-3

Additional test material information

Specific details on test material used for the study

Specific details on test material used for the study (confidential)

Test animals

Species
rat

Strain
Fischer 344

Details on species / strain selection

Sex
male/female

Details on test animals or test system and environmental conditions

Administration / exposure

Route of administration
oral: gavage

Details on route of administration

Vehicle

Details on oral exposure

Analytical verification of doses or concentrations

Details on analytical verification of doses or concentrations

Duration of treatment / exposure
90 days

Frequency of treatment

Doses / concentrations

Dose / conc. Remarks Actions

No. of animals per sex per dose

Control animals

Details on study design

Positive control

Examinations

Observations and examinations performed and frequency

Sacrifice and pathology

Optional endpoint(s)

Other examinations

Statistics

Any other information on materials and methods incl. tables

Results and discussion

Results of examinations

Clinical signs

Description (incidence and severity)

Mortality

Description (incidence)

Body weight and weight changes

Description (incidence and severity)

Food consumption and compound intake (if feeding study)

Description (incidence and severity)

Food efficiency

Description (incidence and severity)

Water consumption and compound intake (if drinking water study)

Description (incidence and severity)

Ophthalmological findings

Description (incidence and severity)

Haematological findings

Description (incidence and severity)

Clinical biochemistry findings

Description (incidence and severity)

Endocrine findings

Description (incidence and severity)

Urinalysis findings

Description (incidence and severity)

Behaviour (functional findings)

Description (incidence and severity)

Immunological findings

Description (incidence and severity)

Organ weight findings including organ / body weight ratios

Description (incidence and severity)

Gross pathological findings

Description (incidence and severity)

Neuropathological findings

Description (incidence and severity)

Histopathological findings: non-neoplastic

Description (incidence and severity)

Histopathological findings: neoplastic

Description (incidence and severity)

Other effects

Description (incidence and severity)

Details on results

Effect levels

Key result Dose descriptor Effect level Based on Sex Basis for effect level Remarks on result Actions 1

☐ Key result

Dose descriptor

LOAEL

Effect level

100 mg/kg bw/day (actual dose received)

Based on

test mat.

Sex

male/female

Basis for effect level

- other:

Remarks on result

Target system / organ toxicity

Key result Critical effects observed Lowest effective dose / conc. System Organ Treatment related Dose response relationship Relevant for humans Actions 1

☐ Key result

Critical effects observed

not specified

Lowest effective dose / conc.

System

Organ

Treatment related

Dose response relationship

Relevant for humans

Overall remarks, attachments

Overall remarks

Attachments

Type Attached (confidential) document Attached (sanitised) documents for publication Remarks Actions

Illustration (picture/graph)

Applicant's summary and conclusion

Conclusions

Executive summary

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Inventory manager

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User management

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