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SuperUser EPA/ORD/CCTE/SCDCD

- User Settings
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- 2-(2-butoxyethoxy)ethyl acetate

**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
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Type at least 3 characters

REACH Complete  
2-(2-butoxyethoxy)ethyl acetate

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0 1 0

Administrative data Data source Materials and methods Results and discussion Overall remarks, attachments

Applicant's summary and conclusion

#### Administrative data

Endpoint  
toxicity to reproduction

Type of information  
experimental study

Adequacy of study  
supporting study

☐ Robust study summary

☐ Used for classification

☐ Used for SDS

Study period

Reliability  
1 (reliable without restriction)

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 1

Reason / purpose for cross-reference

reference to same study

Related information

Remarks

2

Reason / purpose for cross-reference

reference to other study

Related information

Remarks

## Data source

### Reference

- publication | Unnamed | 1993

### Data access

### Data protection claimed

## Materials and methods

### Test guideline

# Qualifier Guideline Version / remarks Deviations Actions 1

Qualifier

equivalent or similar to guideline

Guideline

other:

Version / remarks

Deviations

not applicable

Principles of method if other than guideline

GLP compliance

not specified

Limit test

no

Justification for study design

### Test material

Test material information

- Unnamed | Unnamed | 2-butoxyethanol | EC 203-905-0 | 111-76-2

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: drinking water

---

Type of inhalation exposure (if applicable)

---

Mass median aerodynamic diameter (MMAD)

---

Geometric standard deviation (GSD)

---

Remarks on MMAD

---

Vehicle

---

Details on exposure

---

Details on mating procedure

---

Analytical verification of doses or concentrations

---

Details on analytical verification of doses or concentrations

---

Duration of treatment / exposure

---

Frequency of treatment

---

Details on study schedule

---

**Doses / concentrations**

# Dose / conc. Remarks Actions

---

No. of animals per sex per dose

---

Control animals

---

Details on study design

---

Positive control

---

**Examinations**

---

Parental animals: Observations and examinations

---

Oestrous cyclicity (parental animals)

---

Sperm parameters (parental animals)

---

Litter observations

---

Postmortem examinations (parental animals)

---

Postmortem examinations (offspring)

---

Statistics

---

Reproductive indices

---

Offspring viability indices

---

**Any other information on materials and methods incl. tables**

---

**Results and discussion**

---

**Results: P0 (first parental generation)**

**General toxicity (P0)**

Clinical signs

Description (incidence and severity)

Dermal irritation (if dermal study)

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)

---

**Reproductive function / performance (P0)**

---

Reproductive function: oestrous cycle

---

Description (incidence and severity)

---

Reproductive function: sperm measures

---

Description (incidence and severity)

---

Reproductive performance

---

Description (incidence and severity)

---

**Details on results (P0)**

---

---

**Effect levels (P0)**

---

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

☐ Key result

Dose descriptor

NOAEL

Effect level

> 452 mg/kg bw/day (actual dose received)

Based on

Sex

male

Basis for effect level

- other:

Remarks on result

2

☐ Key result

Dose descriptor

NOAEL

Effect level

> 470 mg/kg bw/day (actual dose received)

Based on

Sex

female

Basis for effect level

- other:

Remarks on result

---

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

Results: P1 (second parental generation)

General toxicity (P1)

Clinical signs

Description (incidence and severity)

Dermal irritation (if dermal study)

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

**Reproductive function / performance (P1)**

Reproductive function: oestrous cycle

Description (incidence and severity)

Reproductive function: sperm measures

Description (incidence and severity)

Reproductive performance

Description (incidence and severity)

**Details on results (P1)**

**Effect levels (P1)**

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

**Target system / organ toxicity (P1)**

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

**Results: F1 generation**

**General toxicity (F1)**

Clinical signs

Description (incidence and severity)

Dermal irritation (if dermal study)

---

Description (incidence and severity)

---

Mortality / viability

---

Description (incidence and severity)

---

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)

---

Urinalysis findings

---

Description (incidence and severity)

---

Sexual maturation

---

Description (incidence and severity)

---

Anogenital distance (AGD)

---

Description (incidence and severity)

---

Nipple retention in male pups

---

Description (incidence and severity)

---

Organ weight findings including organ / body weight ratios

---

Description (incidence and severity)

---

Gross pathological findings

---

Description (incidence and severity)

---

Histopathological findings

---

Description (incidence and severity)

---

Other effects

---

Description (incidence and severity)

---

**Developmental neurotoxicity (F1)**

---

Behaviour (functional findings)

---

Description (incidence and severity)

---

**Developmental immunotoxicity (F1)**

---

Developmental immunotoxicity

---

Description (incidence and severity)

---

**Details on results (F1)**

---

---

**Effect levels (F1)**

---

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

---

**Target system / organ toxicity (F1)**

---

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

---

**Results: F2 generation**

---

**General toxicity (F2)**

---

Clinical signs

---

Description (incidence and severity)

---

Dermal irritation (if dermal study)

---

Description (incidence and severity)

---

Mortality / viability

---

Description (incidence and severity)

---

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)

Urinalysis findings

Description (incidence and severity)

Sexual maturation

Description (incidence and severity)

Anogenital distance (AGD)

Description (incidence and severity)

Nipple retention in male pups

Description (incidence and severity)

Organ weight findings including organ / body weight ratios

Description (incidence and severity)

Gross pathological findings

Description (incidence and severity)

Histopathological findings

Description (incidence and severity)

Other effects

Description (incidence and severity)

**Developmental neurotoxicity (F2)**

Behaviour (functional findings)

Description (incidence and severity)

**Developmental immunotoxicity (F2)**

Developmental immunotoxicity

Description (incidence and severity)

**Details on results (F2)**

**Effect levels (F2)**

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

**Target system / organ toxicity (F2)**

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

Overall reproductive toxicity

#	Key result	Reproductive effects observed	Lowest effective dose / conc.	Treatment related	Relation to other toxic effects	Dose response relationship	Relevant for humans	Actions	1
<input type="checkbox"/>	Key result	Reproductive effects observed	not specified	Lowest effective dose / conc.	Treatment related	Relation to other toxic effects	Dose response relationship	Relevant for humans	

Any other information on results incl. tables

Overall remarks, attachments

Overall remarks

Attachments

#	Type	Attached (confidential) document	Attached (sanitised) documents for publication	Remarks	Actions
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Illustration (picture/graph)

Applicant's summary and conclusion

Conclusions

Executive summary

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#### About IUCLID

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