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**Date: Tuesday, February 17, 2026 14:52**

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**INCURIN® 1 mg tablet**

**Species:** Dogs

**Therapeutic indication: Pharmaceuticals: Hormones and therapeutically related products:** Other sex hormones

**Active ingredient:** Estriol

**Product:** INCURIN® 1 mg tablet

**Product index:** INCURIN®

**Incorporating:**

**Qualitative and quantitative composition**

**Active substance:**

Estriol 1 mg/tablet

For the full list of excipients, see section “Pharmaceutical particulars”.

**Pharmaceutical form**

White, round, single-scored tablets.

**Clinical particulars**

**Target Species**

Dogs (bitch).

**Indications for use**

The treatment of hormone-dependent urinary incontinence due to sphincter mechanism incompetence in ovariohysterectomised bitches.

**Contraindications**

Do not use in intact bitches as the efficacy has only been established in ovariohysterectomised bitches.

Do not use in animals with symptoms of polyuria-polydipsia.

Do not use in animals younger than 1 year.

### **Special warnings for each target species**

High doses of oestrogen may have a tumour-promoting effect in target organs with oestrogen receptors (mammary glands).

### **Special precautions for use**

In case of oestrogenic effects, the dose should be lowered.

### **Operator warnings**

Not applicable.

### **Adverse Reactions**

Dog (bitch):

Very common (> 1 animal / 10 animals treated): Swollen vulva<sup>1,2</sup>, mammary gland oedema<sup>1,2</sup>, attr

Rare (1 to 10 animals / 10 000 animals treated): Vaginal haemorrhage. Alopecia.

1 Observed at the highest recommended dose of 2 mg per dog.

2 These effects are reversible after lowering the dose.

Reporting adverse events is important. It allows continuous safety monitoring of a veterinary medicinal product. Reports should be sent, preferably via a veterinarian, to either the marketing authorisation holder or the national competent authority via the national reporting system. See the package leaflet for respective contact details.

### **Use during pregnancy or lactation**

The use is not recommended during pregnancy or lactation.

### **Interactions**

None known.

### **Amounts to be administered and administration route**

Oral use.

A relationship between the final effective dose and body weight has not been established and therefore the dose must be determined for each dog on an individual basis.

Recommended dosing schedule:

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Start treatment with 1 tablet (1 mg estriol) every day.

If treatment is successful, lower the dose to half a tablet per day.

If treatment is not successful, increase the dose to 2 tablets per day to be administered as a single dose.

Do not use more than 2 tablets per dog per day.

The minimum dose should not be less than half a tablet (0.5 mg estriol) per dog per day.

Ensure the dose used to achieve the therapeutic effect is as low as possible.

Some dogs do not need daily treatment. Once an effective daily dose has been established, treatment every other day may be tried.

If no response to treatment is achieved, the diagnosis should be reconsidered and other causes for the incontinence such as neurological disorders, bladder neoplasia, etc. investigated.

Animals should be re-examined every 6 months during treatment.

### **Overdose**

In case of overdose, typical oestrogen effects may occur. These effects are reversible after lowering the dose.

### **Withdrawal periods**

Not applicable.

### **Pharmacological particulars**

**ATCvet code:** QG03CA04

### **Pharmacodynamic properties**

Estriol is a short-acting natural oestrogen. In ovarioectomised female dogs it has a beneficial effect on urinary incontinence. In the target animal safety study and the clinical trials, including long-term treatment, no signs of bone marrow suppression were observed. This is probably due to the short-acting oestrogenic character of estriol.

### **Pharmacokinetic particulars**

After oral administration, estriol is nearly completely absorbed from gastrointestinal tract. Almost all of the absorbed estriol is bound to plasma albumin.

Estriol is excreted in conjugated form via the urine.

After oral administration of multiple doses, no accumulation occurs.

### **Pharmaceutical particulars**

#### **Excipients**

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Amylopectin

Potato starch

Magnesium stearate

Lactose

### **Major incompatibilities**

Not applicable.

### **Shelf life**

Shelf life of the veterinary medicinal product as packaged for sale: 3 years.

### **Special precautions for storage**

Do not store above 30°C.

### **Immediate packaging**

Blister package of clear PVC film backed by aluminium foil provided with heat seal coating (vinyl copolymer) on the side in contact with the tablets. One blister contains 30 tablets.

Pack size: carton box with 1 blister of 30 tablets.

### **Disposal**

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived therefrom in accordance with local requirements and with any national collection systems applicable to the veterinary medicinal product concerned.

### **Marketing Authorisation Holder (if different from distributor)**

UK(NI): Intervet International B.V., The Netherlands

### **Marketing Authorisation Number**

UK(GB): Vm 01708/5038

UK(NI): EU/2/00/018/001

### **Significant changes**

#### **Date of the first authorisation or date of renewal**

Date of first authorisation: 24/03/2000

Date of last renewal: 15/03/2010

#### **Date of revision of the text**

## **NOAH Compendium**

September 2025.

### **Any other information**

For animal treatment only. Keep out of the sight and reach of children.

### **Legal category**

**Legal category:**POM-V

### **GTIN**

**GTIN description:**Incurin 1mg 1x30tab:

**GTIN:**8713184156327