Australian Government
Australian Pesticides and Veterinary Medicines Authority

# PERMIT TO ALLOW MINOR USE OF A REGISTERED AGVET CHEMICAL

# PRODUCT FOR CONTROL OF LIVERFUKE INFESTATION IN RUMINANTS, CAMELIDS AND EQUINES

## PERMIT NUMBER – PER13882

This permit is issued to the Permit Holder in response to an application granted by the APVMA under section 112 of the Agvet Codes of the jurisdictions set out below. This permit allows a person, as stipulated below, to use the product in the manner specified in this permit in the designated jurisdictions. This permit also allows the Permit Holder and any person stipulated below to claim that the product can be used in the manner specified in this permit.

**THIS PERMIT IS IN FORCE FROM 01 JANUARY 2013 TO 31 JANUARY 2027**

**Permit Holder:**
DEPARTMENT OF PRIMARY INDUSTRIES & REGIONAL DEVELOPMENT
1 Nash Street
PERTH, WA, 6000, AUSTRALIA

**Persons who can use the product under this permit:**
The Products must only be used by an inspector of stock, a registered veterinary surgeon, or a person acting under the supervision of an inspector of stock or a registered veterinary surgeon for the treatment of animals for liver fluke for the purposes of transport of the animals into Western Australia.

## CONDITIONS OF USE

**Products to be used:**

FASINEX 120 FLUKICIDE FOR CATTLE AND SHEEP (APVMA No. 47675)
Containing 120 g/L TRICLABENDAZOLE as the only active constituent.

FLUKARE C FLUKICIDE FOR CATTLE AND SHEEP (APVMA No. 51308)
Containing 120 g/L TRICLABENDAZOLE as the only active constituent.

FASINEX 240 ORAL FLUKICIDE FOR CATTLE AND SHEEP (APVMA No. 51262)
Containing 240 g/L TRICLABENDAZOLE as the only active constituent.

TREMACIDE 120 FLUKICIDE FOR CATTLE AND SHEEP (APVMA No. 52463)
Containing 120 g/L TRICLABENDAZOLE as the only active constituent.

WSD LV TRICLABENDAZOLE ORAL FLUKICIDE FOR SHEEP, CATTLE AND GOATS (APVMA No. 60617)
Containing 100 g/L TRICLABENDAZOLE as the only active constituent.

EXIFLUKE 240 ORAL FLUKICIDE FOR CATTLE (APVMA No. 63770)
Containing 240 g/L TRICLABENDAZOLE as the only active constituent.

**Restraints:**

DO NOT USE in lactating or pregnant animals where milk may be used or processed for human consumption, except in cattle.

**Directions for Use:**

| Target animals | Purpose | Dose |
|---|---|---|
| Ruminants and camelids | Adult and immature Liver Flukicide (Importation treatment) | 15 mg/kg body weight triclabendazole administered as an oral treatment |
| Equines | | 12 mg/kg body weight triclabendazole administered as an oral treatment |

**Precautions:**

The safety of the products listed in this permit has not been fully established in young or pregnant animals. Use of this product in pregnant or young animals, at the recommended dose as specified in this permit, must be based on a risk-benefit analysis by a registered veterinarian.

**Critical Use Comments:**

- Target animal species covered by this permit are those identified in the Published import conditions as set out by the *Biosecurity and Agriculture Management Act 2007* and the *Biosecurity and Agriculture Management Regulations 2013.*
- The maximum number of treatments within any period of six (6) months is six (6) treatments.
- The minimum retreatment interval is 14 days.
- Resistance may develop to any drenches. Ask your local veterinarian or animal health adviser for recommended parasite management practices for your area to reduce development of resistance. It is advisable that a resistance test be conducted before any parasite treatment is used.

**Withholding periods [MEAT]:**

MEAT [Cattle, sheep, goats] following single dose: DO NOT USE less than 28 days before slaughter for human consumption.
MEAT [Cattle, sheep, goats] following multiple doses: DO NOT USE less than 42 days before slaughter for human consumption.

A meat withholding period has not been established for ruminants, other than cattle, sheep and goats. If the Products listed in this permit are to be used in other species of ruminants which will be processed for human consumption, an extended withholding period (WHP) may be required.

**(**CAMELIDS): DO NOT USE less than 49 days before slaughter for human consumption.
**(**HORSES): DO NOT USE less than 49 days before slaughter for human consumption.

**Withholding periods [MILK]:**

MILK [Cattle]: LACTATING: Where an animal is lactating at the time of treatment, milk used or intended to be used for human consumption MUST NOT BE COLLECTED from that animal until 35 days after that treatment. Calves fed this milk should be withheld from slaughter for 28 days after last exposure.

MILK [Cattle]: NON-LACTATING: DO NOT USE less than 35 days before calving in cows where milk and milk products from treated animals may be used for human consumption. Where triclabendazole is accidentally given within this period or should any cow calve earlier than 35 days after treatment, the milk will contain residues. This milk must not be used for human consumption or supplied for processing for at least 35 days following treatment. Calves fed this milk should be withheld from slaughter for 28 days after last exposure.

**TRADE ADVICE:** Export Slaughter Interval (ESI): [Cattle, sheep] DO NOT USE less than six (6) months before slaughter for export.

**Jurisdiction:**
All States and Territories.

**Additional Conditions:**
This permit allows for the use of a product in a manner specified on the permit. Persons who wish to prepare for use and/or use products for the purposes specified in this permit must read, or have read to them, the details and conditions of this permit. Unless otherwise stated, the use of the product must be in accordance with the registered product labels.

*Use*

1. Users must report any adverse events (including lack of efficacy) arising from the use of the Products to the permit Holder and to the APVMA, Adverse Experience Reporting Program. Phone: 1800 700 583; Email: aerp@apvma.gov.au and online reporting - https://portal.apvma.gov.au/aerpexternal/welcome.htm. **More information:** Find out more about how to report adverse experiences.

*Holder*

2. The holder of the permit must notify the APVMA of new information, including relevant information in accordance with section 161 of the Schedule to the *Agricultural*

*and Veterinary Chemicals Code Act 1994*, in accordance with the obligation imposed by that section.

Issued by the Australian Pesticides and Veterinary Medicines Authority

Note: 7/9/2015 – Permit expiry extended to 31 /1/2017. Permit issued as Version 2
Note: 30/01/2020 – Permit updated to remove the product Flukare S Flukicide For Sheep, Cattle and Goats (50 g/L Triclabendazole) as it is no longer registered. Permit expiry extended to 31/1/2023. Permit issued as Version 3
Note: 11/01/2023 – Permit updated to current standard. Products listed on previous permit, 58529, 64929, 58982 were removed as they are no longer registered. Permit expiry extended to 31/01/2026. Permit issued as Version 4.
Note: 23/01/2026 – Permit updated to remove products listed on previous permit, 52185, 47676, 60489, and 52865 as they are no longer registered. Restraints and precaution statements added. Critical comment added to clarify scope of target animals to be treated under this permit. Withholding period statements updated for clarity on target species. Additional conditions on use added. Permit expiry extended to 31/01/2027. Permit issued as Version 5 on 23/01/2026.