1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT
Marketing authorisation holder:
Elanco Europe Ltd., Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, UK.
Tel: 01256 353131
Manufacturer responsible for batch release:
Surepharm Services Ltd., Bretby Business Park, Bretby, Burton Upon Trent, DE15 0YZ

2. NAME OF THE VETERINARY MEDICINAL PRODUCT
ACP Tablets 10 mg

3. STATEMENT OF THE ACTIVE SUBSTANCE(S) AND OTHER INGREDIENT(S)
10 mg Acepromazine (as acepromazine maleate 13.54 mg). Tablets are pale yellow in colour.

4. INDICATION(S)
The tablets are intended for use only in cats and dogs.

Anaesthetic Premedication: Following acepromazine administration, the amount of anaesthetic necessary to induce anaesthesia is considerably reduced. This reduction is approximately one-third of a suitable induction agent.

Tranquillisation: Acepromazine tranquillisation (ataraxy) involves a modification of temperament which is not associated with hypnosis, narcosis or marked sedation. This is achieved with low doses of acepromazine.

Sedation: At higher dose rates acepromazine is a sedative.

Travel sickness: A dose of 1 mg per kg given orally a quarter to half an hour before a light meal is effective in the prevention of travel sickness. Idiopathic vomiting may be controlled by acepromazine. Acepromazine possesses anti-emetic, anti-convulsant, hypothermic, hypotensive and anti-spasmodic properties and shows a marked potentiating effect on barbiturate anaesthesia.

5. CONTRAINDICATIONS
Do not use in pregnant animals.
Do not use on a long term basis in individual animals.

6. ADVERSE REACTIONS

7. TARGET SPECIES
Dog and cat.
For animal treatment only.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION
0.25 - 3 mg per kg bodyweight by oral administration. Onset of effects will be observed after 10-15 minutes. Normally single doses of acepromazine are administered.

9. ADVICE ON CORRECT ADMINISTRATION

10. WITHDRAWAL PERIOD(S)

11. SPECIAL STORAGE PRECAUTIONS
Do not store above 25°C.
Protect from light.
Keep out of the sight and reach of children.

12. SPECIAL WARNING(S)

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY
Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED
September 2020

15. OTHER INFORMATION
POM-V To be supplied only on veterinary prescription.
Vm 00879/4011
Containers of 500 x 10 mg tablets.