

## **SUMMARY OF PRODUCT CHARACTERISTICS**

### **1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Kaogel VP 19.72% w/v Oral Suspension

### **2. QUALITATIVE AND QUANTITATIVE COMPOSITION**

<i>Active substance</i>	<i>% w/v</i>
Kaolin, light	19.72

<i>Excipients</i>	
Sorbic Acid	0.10
Methyl Parahydroxybenzoate	0.20

For a full list of excipients, see Section 6.1.

### **3. PHARMACEUTICAL FORM**

Oral suspension.  
An off-white suspension

### **4. CLINICAL PARTICULARS**

#### **4.1 Target species**

Dogs and cats.

#### **4.2 Indications for use, specifying the target species**

For treatment of diarrhoea of non-specific origins in dogs and cats.

#### **4.3 Contra-indications**

Contra-indicated in cases of intestinal obstruction.

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

None.

#### **4.5 Special precautions for use**

- i) Special precautions for use in animals  
Shake the bottle before use.
- ii) Special precautions to be taken by the person administering the veterinary medicinal product to animals

Wash hands after use.

#### **4.6 Adverse reactions (frequency and seriousness)**

None.

#### **4.7 Use during pregnancy, lactation or lay**

No restriction.

#### **4.8 Interaction with other medicinal products and other forms of interaction**

May adsorb other oral medications when given concurrently therefore reducing bioavailability.

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

Shake the bottle before use. By oral administration. 0.5–1 ml/kg bodyweight as a total daily dose. May be given as a divided dose 3-4 times daily.

#### **4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary**

Overdose may lead to mild constipation.

#### **4.11 Withdrawal period**

Not applicable.

### **5. PHARMACOLOGICAL PROPERTIES**

#### **ATCVet Code: QA07BC02**

The active principal is kaolin light which is a well-established gastro-intestinal adsorbent of toxins and other substances. It also increases the bulk of the faeces to aid peristalsis.

### **6. PHARMACEUTICAL PARTICULARS**

#### **6.1 List of excipients**

Sorbic Acid

Methyl Parahydroxybenzoate

Saccharin Sodium

Citric Acid Monohydrate or Citric Acid Anhydrous

Pectin

Bentonite

Water, Purified

Citric Acid Monohydrate or Citric Acid Anhydrous or Sodium Citrate Dihydrate (for pH adjustment)

## **6.2 Incompatibilities**

None known.

## **6.3 Shelf-life**

Shelf-life of the veterinary medicinal product as packaged for sale: 5 years.

## **6.4 Special precautions for storage**

Do not store above 25°C.

## **6.5 Nature and composition of immediate packaging**

480 ml white opaque high density polyethylene bottle with white opaque high density polyethylene cap (screw fit).

## **6.6 Special precautions for the disposal of unused veterinary medicinal product or waste materials derived from the use of such products, if appropriate**

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

## **7. MARKETING AUTHORISATION HOLDER**

Zoetis UK Limited  
1<sup>st</sup> Floor, Birchwood Building  
Springfield Drive  
Leatherhead  
Surrey  
KT22 7LP

## **8. MARKETING AUTHORISATION NUMBER**

Vm 42058/4073

## **9. DATE OF THE FIRST AUTHORISATION**

31 January 1985

**10. DATE OF REVISION OF THE TEXT**

November 2023

Approved 03 November 2023

A handwritten signature in black ink, appearing to read 'M. M. M.', located below the approval date.