

# Summary of Product Characteristics

## 1 NAME OF THE VETERINARY MEDICINAL PRODUCT

BOVIGEN SCOUR Emulsion for injection for cattle

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

One dose of the vaccine (3 ml) contains:

### Active substances:

Bovine rotavirus strain TM-91, serotype G6P1 (inactivated)  $\geq 6.0 \log_2$  (VNT)\*

Bovine coronavirus strain C-197 (inactivated)  $\geq 5.0 \log_2$  (HIT)\*\*

*Escherichia coli* strain EC/17 (inactivated)

expressing F5 (K99) adhesin  $\geq 44.8$  % of inhibition (ELISA)\*\*\*

\*VNT – virus neutralisation test (rabbit serology induced by 2/3 dose of vaccine).

\*\*HIT – haemagglutination inhibition test (rabbit serology induced by 2/3 dose of vaccine).

\*\*\*ELISA – Enzyme-linked immunosorbent assay (rabbit serology induced by 2/3 dose of vaccine).

### Adjuvant:

Montanide ISA 206 VG 1.6 ml

### Excipients:

Formaldehyde max. 1.5 mg

Thiomersal max. 0.36 mg.

For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Emulsion for injection.

White liquid emulsion which may form a sediment during storage.

## 4 CLINICAL PARTICULARS

### 4.1 Target Species

Cattle (pregnant cows and heifers)

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

For the active immunisation of pregnant cows and heifers to raise antibodies against *E. coli* adhesion F5 (K 99) antigen, rotavirus and coronavirus. When calves are fed colostrum from vaccinated cows during the first week of life, these antibodies have been demonstrated to reduce the severity of diarrhoea caused by bovine rotavirus, bovine coronavirus and enteropathogenic *E. coli* F5 (K99) and to reduce the shedding of virus by calves infected with bovine rotavirus or bovine coronavirus.

Onset of immunity: Passive immunity commences with colostrum feeding and is dependent on calves receiving sufficient colostrum after birth.

### 4.3 Contraindications

None.

#### 4.4 Special warnings for each target species

Vaccinate healthy animals only.

#### 4.5 Special precautions for use

##### Special precautions for use in animals

Not applicable.

##### Special precautions to be taken by the person administering the veterinary medicinal product to animals

To the user:

This veterinary medicinal product contains mineral oil. Accidental injection/self-injection may result in severe pain and swelling, particularly if injected into a joint or finger, and in rare cases could result in the loss of the affected finger if prompt medical attention is not given.

If you are accidentally injected with this veterinary medicinal product, seek prompt medical advice even if only a very small amount is injected and take the package leaflet with you.

If pain persists for more than 12 hours after medical examination, seek medical advice again.

To the physician:

This veterinary medicinal product contains mineral oil. Even if small amounts have been injected, accidental injection with this veterinary medicinal product can cause intense swelling, which may, for example, result in ischemic necrosis and even the loss of a digit. Expert, PROMPT, surgical attention is required and may necessitate early incision and irrigation of the injected area, especially where there is involvement of finger pulp or tendon.

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

A slight swelling of 5-7 cm in diameter at the site of injection is common and may in some cases be accompanied initially by increased local temperature. Typically, such swelling resolves within 15 days.

Slight, transient increases in temperature (up to 0.8 °C) may be observed within 24 hours of vaccination, resolving within 4 days after vaccination.

The frequency of adverse reactions is defined using the following convention:

- very common (more than 1 in 10 animals treated displaying adverse reaction(s))
- common (more than 1 but less than 10 animals in 100 animals treated)
- uncommon (more than 1 but less than 10 animals in 1,000 animals treated)
- rare (more than 1 but less than 10 animals in 10,000 animals treated)
- very rare (less than 1 animal in 10,000 animals treated, including isolated reports).

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

This veterinary medicinal product is intended for use in the last trimester of pregnancy.

#### 4.8 Interaction with other medicinal products and other forms of interactions

No information is available on the safety and efficacy of this vaccine when used with any other veterinary medicinal product. A decision to use this vaccine before or after any other veterinary medicinal product therefore needs to be made on a case by case basis.

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

For intramuscular use.

Common aseptic procedures should be used during vaccination.

Only sterile syringes and needles should be used.

Allow the vaccine to reach room temperature before use. Shake well before and occasionally during use to ensure that the sediment is dissolved prior to administration.

For the 90 ml and 450 ml pack sizes, it is recommended to use automated dosing equipment to protect the stopper against damage from multiple piercing.

One dose: 3 ml

One dose in the course of each pregnancy, given in the 12 – 3 week period before calving is expected.

### Feeding of colostrum

The protection of calves depends on adequate ingestion of colostrum from vaccinated cows. Measures should be taken to ensure that calves receive sufficient amounts of colostrum within the first few days of life. If calves do not get enough antibodies through the colostrum soon after they are born, they will have failure of passive transfer of antibodies. It is important that all calves receive as much colostrum as possible from the first milking within the first six hours after calving. It is recommended that at least 3 litres of colostrum are fed within the first 24 hours and this amount is equivalent to approximately 10% of the weight of a calf.

To achieve optimum results and to reduce infection pressure on the farm, a whole herd cow vaccination policy should be adopted.

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

Following the administration of an overdose, no adverse reactions other than those mentioned in section 4.6 occur.

#### **4.11 Withdrawal period(s)**

Zero days.

### **5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES**

ATCvet code: QI02AL01

Pharmacotherapeutic group: immunologicals for Bovidae, inactivated viral and inactivated bacterial vaccines for cattle.

The vaccine is designed for stimulation of the active immunity of pregnant cows against the antigenic components included in the vaccine. The antibodies are transferred to the calf via colostrum.

### **6 PHARMACEUTICAL PARTICULARS**

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

Montanide ISA 206 VG  
Formaldehyde  
Thiomersal  
Eagle's Minimum Essential Medium (MEM)  
Disodium phosphate dodecahydrate  
Sodium chloride  
Potassium chloride  
Potassium dihydrogen phosphate  
Water for injections

#### **6.2 Major incompatibilities**

Do not mix with any other veterinary medicinal product.

#### **6.3 Shelf-life**

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

Shelf life after first opening the immediate packaging: 10 hours.

#### **6.4 Special precautions for storage**

Store in a refrigerator (2 °C - 8 °C). Protect from light. Do not freeze.

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

Glass vials, type I (15 ml, 90 ml) closed with chlorobutyl rubber stoppers or glass bottle, type I (450 ml) closed with bromobutyl rubber stoppers sealed with aluminium caps.

Plastic bottles (450 ml) closed with chlorobutyl rubber stoppers and sealed with aluminium caps without outer package.

Package sizes:

Cardboard box with 1 vial of 15 ml (5 doses)

Cardboard box with 1 vial of 90 ml (30 doses)

Cardboard box with 1 glass bottle of 450 ml (150 doses)

Plastic bottle with 450 ml (150 doses)

Not all pack sizes may be marketed.

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

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**

FORTE Healthcare Ltd

Cougar Lane

Naul

Co. Dublin

Ireland

## **8 MARKETING AUTHORISATION NUMBER(S)**

VPA10806/004/001

## **9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation: 01 May 2015

Date of last renewal: 30 April 2020

## **10 DATE OF REVISION OF THE TEXT**

August 2020