ANNEX I

SUMMARY OF PRODUCT CHARACTERISTICS
1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CYTOPOINT 10 mg solution for injection for dogs
CYTOPOINT 20 mg solution for injection for dogs
CYTOPOINT 30 mg solution for injection for dogs
CYTOPOINT 40 mg solution for injection for dogs

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Active substance:

Each vial of 1 ml contains:

CYTOPOINT 10 mg:
  Lokivetmab*  10 mg
CYTOPOINT 20 mg:
  Lokivetmab*  20 mg
CYTOPOINT 30 mg:
  Lokivetmab*  30 mg
CYTOPOINT 40 mg:
  Lokivetmab*  40 mg

*Lokivetmab is a caninised monoclonal antibody expressed through recombinant techniques in Chinese hamster ovary (CHO) cells

Excipients:

For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Solution for injection.
The product should appear clear to opalescent without any visible particle.

4. CLINICAL PARTICULARS

4.1 Target species

Dogs

4.2 Indications for use, specifying the target species

Treatment of clinical manifestations of atopic dermatitis in dogs.

4.3 Contraindications

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.
Do not use in dogs less than 3 kg bodyweight.

4.4 Special warnings for each target species

Lokivetmab may induce transient or persistent anti-drug antibodies. The induction of such antibodies is uncommon and may have no effect (transient anti-drug antibodies) or may result in a noticeable
decrease in efficacy (persistent anti-drug antibodies) in animals that responded to treatment previously.

4.5 Special precautions for use

Special precautions for use in animals

In cases of atopic dermatitis, it is recommended to investigate and treat complicating factors, such as bacterial, fungal or parasitic infections/infestations (e.g. flea and mange).

It is recommended to monitor dogs for bacterial infections associated with atopic dermatitis, especially during the first weeks of treatment.

If no or limited response is observed within one month after initial dosing, an improvement in response may be observed after administration of a second dose one month later. However, if the animal does not show a better response after the second dose, the veterinary surgeon should consider alternative treatments.

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

Hypersensitivity reactions, including anaphylaxis, could potentially occur in the case of accidental self-injection.

Accidental self-injection may result in an immune response to lokivetmab. It is not expected for this to cause any adverse effects, however, repeated self-administration may increase the risk of hypersensitivity reactions.

In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.

4.6 Adverse reactions (frequency and seriousness)

Hypersensitivity reactions (anaphylaxis, facial oedema, urticaria) may occur in rare cases. In such cases appropriate treatment should be administered immediately.

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

The safety of the veterinary medicinal product has not been established during pregnancy and lactation; therefore its use is not recommended during pregnancy, lactation or in breeding animals.

4.8 Interaction with other medicinal products and other forms of interaction

No drug interactions were observed in field studies where lokivetmab was administered concomitantly with veterinary medicinal products such as endo- and ectoparasiticides, antimicrobials, anti-inflammatories and vaccines.

If a vaccine(s) is to be administered at the same time as treatment with lokivetmab, the vaccine(s) should be administered at a different site to that of lokivetmab administration.
4.9 Amounts to be administered and administration route

Subcutaneous use.

Avoid excessive shaking or foaming of the solution. Administer the entire contents (1 ml) of the vial.

Dose according to the dosing chart below. For dogs above 40 kg, the contents of more than one vial are required to administer a single dose. In those cases, withdraw the appropriate content from each required vial into the same syringe. To allow for mixing of the solution, gently invert the syringe three or four times before administering.

Dosage and treatment schedule:

The recommended minimum dose is 1 mg/kg bodyweight, once a month.

Dose according to the dosing chart below:

<table>
<thead>
<tr>
<th>Bodyweight (kg) of dog</th>
<th>CYTOPOINT strength (mg) to be administered</th>
</tr>
</thead>
<tbody>
<tr>
<td>3.0-10.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>10.1-20.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>20.1-30.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>30.1-40.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>40.1-50.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>50.1-60.0</td>
<td>2 vials</td>
</tr>
<tr>
<td>60.1-70.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>70.1-80.0</td>
<td>2 vials</td>
</tr>
</tbody>
</table>

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

No adverse reactions other than those mentioned in section 4.6 were observed in laboratory overdose studies.

In case of adverse clinical signs after an overdose the dog should be treated symptomatically.

4.11 Withdrawal period(s)

Not applicable.

5. IMMUNOLOGICAL PROPERTIES

Pharmacotherapeutic group: Other dermatological preparations. Agents for dermatitis, excluding corticosteroids.
Lokivetmab is a caninised monoclonal antibody (mAb) specifically targeting canine interleukin-31. The blocking of IL-31 by lokivetmab prevents IL-31 from binding to its co-receptor and thereby inhibits IL-31 mediated cell signalling, providing relief from Atopic Dermatitis-related pruritus and anti-inflammatory activity.

In a laboratory model study lokivetmab demonstrated an onset of efficacy for pruritus by the first time point at 8 hours post administration.

In field studies up to 9 months, treatment of dogs with atopic dermatitis was demonstrated to have a favourable effect on the reduction of pruritus and on the reduction of disease severity as evaluated by Canine Atopic Dermatitis Extent and Severity Index (CADESI) 03 scores. A small number of dogs showed a low or an absence of clinical response to lokivetmab. This is likely due to the highly targeted mechanism of action of lokivetmab in the context of a complex disease and heterogeneous pathogenesis. Refer also to section 4.5 of the SPC.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Histidine
Histidine hydrochloride monohydrate
Trehalose dihydrate
Disodium edetate
Methionine
Polysorbate 80
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: 2 years.
Shelf life after first opening the immediate packaging: use immediately

6.4 Special precautions for storage

Store in a refrigerator (2 °C – 8 °C).
Do not freeze.
Store in the original package.
Protect from light.

6.5 Nature and composition of immediate packaging

Primary packaging: Single dose clear glass Type I vials with fluorobutyl rubber stopper.
Secondary packaging: cardboard box.

Pack sizes:
CYTOPOINT 10 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml
CYTOPOINT 20 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

CYTOPOINT 30 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

CYTOPOINT 40 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

Not all pack sizes may be marketed.

6.6 Special precautions for the disposal of unused veterinary medicinal product 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 AUTHORITY

Zoetis Belgium SA
Rue Laid Burniat 1
1348 Louvain-la-Neuve
BELGIUM

8. MARKETING AUTHORITY NUMBER(S)

EU/2/17/205/001-008

9. DATE OF FIRST AUTHORIZATION/RENEWAL OF THE AUTHORIZATION

Date of first authorisation: 25/04/2017

10. DATE OF REVISION OF THE TEXT

Detailed information on this veterinary medicinal product is available on the website of the European Medicines Agency (http://www.ema.europa.eu/).

PROHIBITION OF SALE, SUPPLY AND/OR USE

Not applicable.
ANNEX II

A. MANUFACTURERS OF THE BIOLOGICAL ACTIVE SUBSTANCE AND MANUFACTURER RESPONSIBLE FOR BATCH RELEASE

B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE

C. STATEMENT OF THE MRLs

D. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING AUTHORIZATION
A. MANUFACTURERS OF THE BIOLOGICAL ACTIVE SUBSTANCE AND MANUFACTURER RESPONSIBLE FOR BATCH RELEASE

Name and address of the manufacturer of the biological active substance

Zoetis Inc.
601 Cornhusker Highway
68521 Lincoln, Nebraska
UNITED STATES

Name and address of the manufacturer responsible for batch release

Zoetis Belgium SA
Rue Laid Burniat 1
1348 Louvain-la-Neuve
BELGIUM

The printed package leaflet of the medicinal product must state the name and address of the manufacturer responsible for the release of the concerned batch.

B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE

Veterinary medicinal product subject to prescription.

C. STATEMENT OF THE MRLs

Not applicable.

D. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING AUTHORISATION

Endotoxin: The endotoxin results for the first 20 batches released onto the EU market should be provided and the specification revised if justified. Data to be provided within 12 months of the granting of the Marketing Authorisation.

Active substance stability: The applicant should complete the ongoing lokivetmab stability evaluation up to 36 months at the different batch sizes. These final reports should be provided by April 2018 and November 2019, for the ongoing 300 L and 2000 L stability studies, respectively.
A. LABELLING
PARTICULARS TO APPEAR ON THE OUTER PACKAGE

CARDBOARD BOX

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CYTOPOINT 10 mg solution for injection for dogs
CYTOPOINT 20 mg solution for injection for dogs
CYTOPOINT 30 mg solution for injection for dogs
CYTOPOINT 40 mg solution for injection for dogs
lokivetmab

2. STATEMENT OF ACTIVE SUBSTANCES

Each vial of 1 ml contains 10 mg lokivetmab.
Each vial of 1 ml contains 20 mg lokivetmab.
Each vial of 1 ml contains 30 mg lokivetmab.
Each vial of 1 ml contains 40 mg lokivetmab.

3. PHARMACEUTICAL FORM

Solution for injection

4. PACKAGE SIZE

2 x 1 ml
6 x 1 ml

5. TARGET SPECIES

Dog

6. INDICATION(S)

Read the package leaflet before use.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Subcutaneous use
Read the package leaflet before use.

8. WITHDRAWAL PERIOD(S)
9. SPECIAL WARNING(S), IF NECESSARY

Read the package leaflet before use.

10. EXPIRY DATE

EXP {month/year}

11. SPECIAL STORAGE CONDITIONS

Store in a refrigerator. Do not freeze.
Store in the original package. Protect from light.
Avoid excessive shaking.

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: read package leaflet.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, IF APPLICABLE

For animal treatment only. To be supplied only on veterinary prescription.

14. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Zoetis Belgium SA
Rue Laid Burniat 1
1348 Louvain-la-Neuve
BELGIUM

16. MARKETING AUTHORISATION NUMBER(S)

<table>
<thead>
<tr>
<th>Marketing Authorisation Number</th>
<th>Concentration (mg/ml)</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>EU/2/17/205/001</td>
<td>10</td>
<td>2 vials</td>
</tr>
<tr>
<td>EU/2/17/205/002</td>
<td>10</td>
<td>6 vials</td>
</tr>
<tr>
<td>EU/2/17/205/003</td>
<td>20</td>
<td>2 vials</td>
</tr>
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<td>EU/2/17/205/004</td>
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<tr>
<td>EU/2/17/205/005</td>
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<td>2 vials</td>
</tr>
<tr>
<td>EU/2/17/205/006</td>
<td>30</td>
<td>6 vials</td>
</tr>
<tr>
<td>EU/2/17/205/007</td>
<td>40</td>
<td>2 vials</td>
</tr>
<tr>
<td>EU/2/17/205/008</td>
<td>40</td>
<td>6 vials</td>
</tr>
</tbody>
</table>
17. MANUFACTURER’S BATCH NUMBER

Lot {number}
## MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

### VIAL – 1 ml

### 1. NAME OF THE VETERINARY MEDICINAL PRODUCT

- CYTOPOINT 10 mg solution for injection for dogs
- CYTOPOINT 20 mg solution for injection for dogs
- CYTOPOINT 30 mg solution for injection for dogs
- CYTOPOINT 40 mg solution for injection for dogs
- lokivetmab

### 2. QUANTITY OF THE ACTIVE SUBSTANCE(S)

- lokivetmab 10 mg/ml
- lokivetmab 20 mg/ml
- lokivetmab 30 mg/ml
- lokivetmab 40 mg/ml

### 3. CONTENTS BY WEIGHT, BY VOLUME OR BY NUMBER OF DOSES

1 ml

### 4. ROUTE(S) OF ADMINISTRATION

SC

### 5. WITHDRAWAL PERIOD(S)

#### 6. BATCH NUMBER

Lot

#### 7. EXPIRY DATE

EXP

#### 8. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.
B. PACKAGE LEAFLET
PACKAGE LEAFLET:
CYTOPOINT 10 mg solution for injection for dogs
CYTOPOINT 20 mg solution for injection for dogs
CYTOPOINT 30 mg solution for injection for dogs
CYTOPOINT 40 mg solution for injection for dogs

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Marketing authorisation holder and Manufacturer responsible for batch release:
Zoetis Belgium SA
Rue Laid Burniat 1
1348 Louvain-la-Neuve
BELGIUM

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

CYTOPOINT 10 mg solution for injection for dogs
CYTOPOINT 20 mg solution for injection for dogs
CYTOPOINT 30 mg solution for injection for dogs
CYTOPOINT 40 mg solution for injection for dogs
lokivetmab

3. STATEMENT OF THE ACTIVE SUBSTANCE AND OTHER INGREDIENTS

Active substances:

Each vial of 1 ml contains:

CYTOPOINT 10 mg:
  Lokivetmab*  10 mg
CYTOPOINT 20 mg:
  Lokivetmab*  20 mg
CYTOPOINT 30 mg:
  Lokivetmab*  30 mg
CYTOPOINT 40 mg:
  Lokivetmab*  40 mg

*Lokivetmab is a caninised monoclonal antibody expressed through recombinant techniques in Chinese hamster ovary (CHO) cells

4. INDICATION(S)

Treatment of clinical manifestations of atopic dermatitis in dogs.

5. CONTRAINDICATIONS

Do not use in case of hypersensitivity to the active substance or to any of the excipients.
Do not use in dogs less than 3 kg bodyweight.
6. ADVERSE REACTIONS

Hypersensitivity reactions (anaphylaxis, facial oedema, urticaria) may occur in rare cases. In such cases appropriate treatment should be administered immediately.

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).

If you notice any side effects, even those not already listed in this package leaflet or you think that the medicine has not worked, please inform your veterinary surgeon.

7. TARGET SPECIES

Dogs.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Subcutaneous use.

Avoid excessive shaking or foaming of the solution. Administer the entire contents (1 ml) of the vial.

Dose according to the dosing chart below. For dogs above 40 kg, the contents of more than one vial are required to administer a single dose. In those cases, withdraw the appropriate content from each required vial into the same syringe. To allow for mixing of the solution, gently invert the syringe three or four times before administering.

Dosage and treatment schedule:

The recommended minimum dose is 1 mg/kg bodyweight, once a month. Dose according to the dosing chart below:
<table>
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<th>CYTOPOINT strength (mg) to be administered</th>
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<td>1 vial</td>
</tr>
<tr>
<td>20.1-30.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>30.1-40.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>40.1-50.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>50.1-60.0</td>
<td>2 vials</td>
</tr>
<tr>
<td>60.1-70.0</td>
<td>1 vial</td>
</tr>
<tr>
<td>70.1-80.0</td>
<td>2 vials</td>
</tr>
</tbody>
</table>

9. **ADVICE ON CORRECT ADMINISTRATION**

Avoid excessive shaking or foaming.

10. **WITHDRAWAL PERIOD(S)**

Not applicable.

11. **SPECIAL STORAGE PRECAUTIONS**

Store in a refrigerator (2°C - 8°C). Do not freeze.
Store in the original package. Protect from light.
Do not use this veterinary medicinal product after the expiry date which stated on the label after EXP.
Shelf life after first opening the container: use immediately.
Keep out of the sight and reach of children.

12. **SPECIAL WARNING(S)**

Special warnings for each target species

Lokivetmab may induce transient or persistent anti-drug antibodies. The induction of such antibodies is uncommon and may have no effect (transient anti-drug antibodies) or may result in a noticeable decrease in efficacy (persistent anti-drug antibodies) in animals that responded to treatment previously.
Special precautions for use in animals

In cases of atopic dermatitis, it is recommended to investigate and treat complicating factors, such as bacterial, fungal or parasitic infections/infestations (e.g. flea and mange).

It is recommended to monitor dogs for bacterial infections, especially during the first weeks of treatment.

If no or limited response is obtained within one month after initial dosing, a second dose one month later may increase effectiveness. If the animal does not show a better response after a second dose, the veterinary surgeon should consider alternative treatments.

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

Hypersensitivity reactions, including anaphylaxis, could potentially occur in the case of accidental self-injection.

Accidental self-injection may result in an immune response to lokivetmab. It is not expected for this to cause any adverse effects, however, repeated self-administration may increase the risk of hypersensitivity reactions.

In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.

Pregnancy and lactation:

The safety of the veterinary medicinal product has not been established during pregnancy and lactation; therefore its use is not recommended during pregnancy, lactation or in breeding dogs.

Interaction with other medicinal products and other forms of interaction:

No drug interactions were observed in field studies where lokivetmab was administered concomitantly with veterinary medicinal products such as endo- and ectoparasiticides, antimicrobials anti-inflammatories and vaccines.

When administering vaccines at the same time as this veterinary medicinal product, it is advised that each injection be given at different sites.

Overdose (symptoms, emergency procedures, antidotes):

No adverse reactions other than those mentioned in section 6 were observed in laboratory overdose studies.

In case of adverse clinical signs after an overdose the dog should be treated symptomatically.

Incompatibilities:

Do not mix with any other veterinary medicinal product.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

Medicines should not be disposed of via wastewater or household waste.
Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required. These measures should help to protect the environment.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

Detailed information on this veterinary medicinal product is available on the website of the European Medicines Agency (http://www.ema.europa.eu/).

15. OTHER INFORMATION

Primary packaging: Single dose clear glass Type I vials with fluorobutyl rubber stopper.

Secondary packaging: cardboard box.

Pack sizes:
CYTOPOINT 10 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

CYTOPOINT 20 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

CYTOPOINT 30 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

CYTOPOINT 40 mg solution for injection for dogs:
Cardboard box with 2 vials of 1 ml
Cardboard box with 6 vials of 1 ml

Not all pack sizes may be marketed.

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

België/Belgique/Belgien
Zoetis Belgium SA
Tél/Tel.: +32 (0) 800 99 189

Lietuva
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Tel: +370 610 05088

Республика България
Zoetis Belgium SA
Tel: +359 2 4775791

Luxembourg/Luxemburg
Zoetis Belgium SA
Tél/Tel.: +352 8002 4026

Česká republika
Zoetis Česká republika, s.r.o.
Tel: +420 257 101 111

Magyarország
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Tel: +361 224 5222
**Danmark**
Orion Pharma Animal Health
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**Deutschland**
Zoetis Deutschland GmbH
Tel: +49 30 330063 0

**Eesti**
Oriola Vilnius UAB
Tel: +370 610 05088

**Ελλάδα**
Zoetis Hellas S.A.
Τηλ.: +30 210 6791900

**España**
Zoetis Spain, S.L.
Tel: +34 91 4191900

**France**
Zoetis France
Tél: +33 (0)810 734 937

**Hrvatska**
Zoetis B.V., Podružnica Zagreb za promidžbu
Tel: +385 1 644 1460

**Ireland**
Zoetis Belgium SA
Tel: +353 (0) 1 256 9800

**İslan**
Icepharma hf.
Sími: +354 540 80 00

**Italia**
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Tel: +39 06 3366 8133

**Κύπρος**
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**Malta**
Agrimed Limited
Tel: +356 21 465 797

**Nederland**
Zoetis B.V.
Tel: +31 (0)10 714 0900

**Norge**
Orion Pharma Animal Health
Tlf: +47 40 00 41 90

**Österreich**
Zoetis Österreich GmbH
Tel: +43 1 2701100 110

**Polska**
Zoetis Polska Sp. z o.o.
Tel: +48 22 2234800

**Portugal**
Zoetis Portugal, Lda.
Tel: +351 21 042 72 00

**România**
Zoetis România S.R.L.
Tel: +40 21 202 3083

**Slovenija**
Zoetis B.V., Podružnica Zagreb za promidžbu
Tel: +385 1 644 1460

**Slovenská republika**
Zoetis Česká republika, s.r.o.
Tel: +420 257 101 111

**Suomi/Finland**
Zoetis Finland Oy
Puh/Tel: +358 (0)9 4300 40

**Sverige**
Orion Pharma Animal Health
Tel: +46 (0)8 623 64 40

**United Kingdom**
Zoetis UK Limited
Tel: +44 (0) 845 300 8034