[Version 9.1,11/2024]

ANNEX I

SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combotec 134 mg/120.6 mg spot-on solution for medium dogs

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each pipette (1.34 ml) contains :

Active substances:
Fipronil 134 mg
(S)-Methoprene 120.6 mg

Excipients:

Qualitative composition of excipients and other constituents	**Quantitative composition if that information is essential for proper administration of the veterinary medicinal product**
Butylhydroxyanisole (E320)	0.27 mg
Butylhydroxytoluene (E321)	0.13 mg
Ethanol 96%	
Polysorbate 80	
Polyvidone K17	
Diethylene glycol monoethyl ether	

Clear amber spot-on solution.

3. CLINICAL INFORMATION

3.1 Target species

Dogs.

3.2 Indications for use for each target species

- For the treatment of dogs weighing 10 to 20 kg bodyweight
- To be used against infestations with fleas, alone or in association with ticks and/or biting lice.
- Treatment of flea infestations *(Ctenocephalides* spp*.)*. Insecticidal efficacy against new infestations with adult fleas persists for 8 weeks. Prevention of the multiplication of fleas by inhibiting the development of eggs (ovicidal activity) and larvae and pupae (larvicidal activity) originating from eggs laid by adult fleas for eight weeks after application.
- Treatment of tick infestations *(Ixodes ricinus, Dermacentor variabilis, Dermacentor reticulatus, Rhipicephalus sanguineus)*. The veterinary medicinal product has a persistent acaricidal efficacy for up to 4 weeks against ticks.
- Treatment of infestations with biting lice *(Trichodectes canis)*.

The veterinary medicinal product can be used as part of a treatment strategy for the control of Flea Allergy Dermatitis (FAD).

3.3 Contraindications

In the absence of available data, the veterinary medicinal product should not be used on puppies less than 8 weeks old and/or weighing less than 2 kg.
Do not use on sick (e.g. systemic diseases, fever) or convalescent animals.
Do not use on rabbits, as adverse reactions with even mortality could occur. In absence of studies, the use of the veterinary medicinal product is not recommended in non-target species.
This veterinary medicinal product is specifically developed for dogs. Do not use in cats and ferrets, as this could lead to overdosing.
Do not use in case of hypersensitivity to the active substances or to any of the excipients.

3.4 Special warnings

Bathing/immersion in water within 2 days after application of the veterinary medicinal product and more frequent bathing than once a week should be avoided, as no study has been performed to investigate how this affects the efficacy of the veterinary medicinal product. Emollient shampoos can be used prior to treatment but reduce the duration of protection against fleas to approximately 5 weeks when used weekly after application of the veterinary medicinal product. Weekly bathing with a 2% chlorhexidine medicated shampoo did not affect efficacy against fleas during a 6 week long study.
There may be an attachment of a few ticks. For this reason a transmission of infectious diseases cannot be completely excluded if conditions are unfavorable.
Fleas from pets often infest the animal's basket, bedding and regular resting areas such as carpets and soft furnishings which should be treated, in case of massive infestation and at the beginning of the control measures, with a suitable insecticide and vacuumed regularly.

Other animals living in the same household should also be treated with a suitable product.

3.5 Special precautions for use

Special precautions for safe use in the target species:

Avoid the contact with the animal’s eyes.
It is important to make sure that the veterinary medicinal product is applied to an area where the animal cannot lick it off and to make sure that animals do not lick each other following treatment.

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

This veterinary medicinal product can cause mucous membrane, skin and eye irritation. Avoid contact of the veterinary medicinal product with mouth, skin and eyes. Wash hands after use. In case of accidental contact with the skin or eyes, rinse thoroughly with water. If irritation persists, seek medical advice.

This veterinary medicinal product may cause allergic reactions. People with known hypersensitivity to fipronil and/or (S)-methoprene should not treat their animal with this veterinary medicinal product.

This veterinary medicinal product may cause neurotoxicity and be harmful if swallowed. Avoid ingestion including hand-to-mouth contact. Do not smoke, drink or eat during application. Keep the pipettes in the original packaging until ready for use and dispose of used pipettes immediately. Wash hands after use. In the event of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

Treated animals should not be handled until the application site is dry, and children should not be allowed to play with treated animals until the application site is dry. It is therefore recommended that animals are not treated during the day, but should be treated during the early evening, and that recently treated animals are not allowed to sleep with owners, especially children

Special precautions for the protection of the environment:

Dogs should not be allowed to swim in watercourses for 2 days after application (see section 5.5).

Other precautions:

The veterinary medicinal product may have adverse effects on painted, varnished or other household surfaces or furnishings.

3.6 Adverse events

Dogs

Very rare (<1 animal / 10 000 animals treated, including isolated reports):	Application site skin discolouration[1], Application site hair loss[1], Application site itching[1], Application site reddening[1], Generalised itching or hair loss Hypersalivation[2]; Vomiting Respiratory signs Increased sensitivity to stimulation[3], Central nervous system depression[3], Neurological signs (NOS)[3]

[1] Transient
[2] If licking occurs, a brief period of excessive salivation may be observed due mainly to the nature of the carrier.
[3] Reversible

Reporting adverse events is important. It allows continuous safety monitoring of a veterinary medicinal product. Reports should be sent, preferably via a veterinarian, to either the marketing authorisation holder or its local representative or the national competent authority via the national reporting system. See the package leaflet for respective contact details.

3.7 Use during pregnancy, lactation or lay

Pregnancy and lactation:

The veterinary medicinal product can be used during pregnancy and lactation.

3.8 Interaction with other medicinal products and other forms of interaction

None known.

3.9 Administration routes and dosage

Spot-on use.

Underdosing could result in ineffective use and may favour resistance development.

One pipette of 1.34 ml per dog weighing over 10 kg and up to 20 kg, corresponding to a minimum recommended dose of 6.7 mg/kg for fipronil and 6 mg/kg for (S)-methoprene.

In the absence of safety studies, the minimum treatment interval is 4 weeks.

Method of administration:
Use the easy-peel corners to remove a pipette from its blister. Do not puncture the foil with scissors, knives or other sharp instruments, as this may damage the pipette inside.

Hold the pipette upright. Tap the narrow part of the pipette to ensure the contents are within the main body of the pipette. Cut off the top of the pipette with scissors.

Part the coat between the shoulder blades until the skin is visible. Place the tip of the pipette on the skin and squeeze gently to empty its contents onto the skin.

Care should be taken to avoid excessive wetting of the hair with the veterinary medicinal product since this will cause a transient, sticky appearance of hairs at the treatment spot.

3.10 Symptoms of overdose (and where applicable, emergency procedures and antidotes)

Do not overdose.
No adverse effects were observed in target animal safety studies in 8 week-old puppies, growing dogs and dogs weighing about 2 kg treated once at five times the recommended dose. The risk of experiencing adverse effects (see section 3.6) may however increase when overdosing, so animals should always be treated with the correct pipette size according to bodyweight.

3.11 Special restrictions for use and special conditions for use, including restrictions on the use of antimicrobial and antiparasitic veterinary medicinal products in order to limit the risk of development of resistance

Not applicable.

3.12 Withdrawal periods

Not applicable.

4. PHARMACOLOGICAL INFORMATION

4.1 ATCvet code :
QP53AX65

4.2 Pharmacodynamics

The veterinary medicinal product is an insecticidal and acaricidal solution for topical use, containing an association of an adulticidal active ingredient, fipronil, in combination with an ovicidal and larvicidal active ingredient, (S)-methoprene.

Fipronil is an insecticide and acaricide belonging to the phenylpyrazole family. It acts by interacting with ligand-gated chloride channels, in particular those gated by the neurotransmitter gamma-aminobutyric acid (GABA), thereby blocking pre- and post-synaptic transfer of chloride ions across cell membranes. This results in uncontrolled activity of the central nervous system and death of insects or acarines. Fipronil kills fleas within 24 hours, ticks *(Dermacentor reticulatus, Dermacentor variabilis, Rhipicephalus sanguineus, Ixodes scapularis, Ixodes ricinus, Haemaphysalis longicornis, Haemaphysalis flava, Haemaphysalis campanulata*) and lice within 48 hours post-exposure.

(S)-Methoprene is an insect growth regulator (IGR) of the class of compounds known as juvenile hormone analogues that inhibit the development of immature stages of insects. This compound mimics the action of juvenile hormone and causes impaired development and death of the developing stages of fleas. The on-animal ovicidal activity of (S)-methoprene results from either direct penetration of the

eggshell of newly laid eggs or from absorption through the cuticle of the adult fleas. (S)-methoprene is also effective in preventing flea larvae and pupae from developing, which prevents contamination of the environment of treated animals with the immature stages of fleas.

4.3 Pharmacokinetics

Studies of metabolism of fipronil have demonstrated that the major metabolite is the sulfone derivative of fipronil.
(S)-methoprene is extensively degraded into carbon dioxide and acetate that are subsequently incorporated into endogenous materials.

The pharmacokinetic profiles after topical application of fipronil and (S)-methoprene in combination were studied in dogs in comparison to intravenous dosing of fipronil or (S)-methoprene alone. This established absorption and other pharmacokinetic parameters. The topical application resulted in low systemic absorption of fipronil (11%) with a mean maximum concentration (C_{max}) of approximately 35 ng/ml fipronil and 55 ng/ml of fipronil sulfone in plasma.

Peak fipronil plasma concentrations are slowly attained (mean t_{max} approximately 101 h), and decline slowly (mean terminal half-life approximately 154 h, highest values are observed for males).
Fipronil is extensively metabolised to fipronil sulfone after topical administration.

Plasma concentrations of (S)-methoprene were below the limit of quantitation (20 ng/ml) in dogs after topical application.

Both (S)-methoprene and fipronil, together with its major metabolite, are well-distributed in the haircoat of a dog within one day after application. The concentrations of fipronil, fipronil sulfone and S-methoprene in the hair coat decrease with time and are detectable for at least 60 days after dosing. Parasites are killed through contact rather than systemic exposure.

No pharmacological interaction between fipronil and (S)-methoprene was noted.

5. PHARMACEUTICAL PARTICULARS

5.1 Major incompatibilities

None known.

5.2 Shelf life

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

5.3 Special precautions for storage

Store in the original package in order to protect from light.

5.4 Nature and composition of immediate packaging

Nature of primary packaging:
A blue pipette is composed of a heat-formed shell (polypropylene/cyclic olefin copolymer/ethylene-vinyl alcohol copolymer/polypropylene) and a film (polyethylene terephthalate/aluminium/polypropylene).

The blue pipette is enclosed in an aluminium blister (polyethylene/polyamide/aluminium/polyamide/polyethylene and polyamide/aluminium/polyethylene).

Package sizes:
Cardboard box containing 1 aluminium blister with 1 pipette.
Cardboard box containing 1 aluminium blister with 2 pipettes.
Cardboard box containing 1 aluminium blister with 3 pipettes.
Cardboard box containing 1 aluminium blister with 4 pipettes.
Cardboard box containing 1 aluminium blister with 6 pipettes.

Not all pack sizes may be marketed.

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

Medicines should not be disposed of via wastewater or household waste.

The veterinary medicinal product should not enter water courses as Fipronil and (S)-methoprene may be dangerous for fish and other aquatic organisms.
Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any national collection systems applicable to the veterinary medicinal product concerned.

6. NAME OF THE MARKETING AUTHORISATION HOLDER

Beaphar B.V.

7. MARKETING AUTHORISATION NUMBER(S)

8. DATE OF FIRST AUTHORISATION

<Date of first authorisation:> <{DD/MM/YYYY}> <{DD month YYYY}.>

9. DATE OF THE LAST REVISION OF THE SUMMARY OF THE PRODUCT CHARACTERISTICS

<{MM/YYYY}>
<{DD/MM/YYYY}>
<{DD month YYYY}>

10. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Veterinary medicinal product not subject to prescription.

Detailed information on this veterinary medicinal product is available in the Union Product Database (https://medicines.health.europa.eu/veterinary).

ANNEX II

LABELLING AND PACKAGE LEAFLET

A. LABELLING

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Cardboard Box

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combotec 134 mg/120.6 mg spot-on for medium dogs

2. STATEMENT OF ACTIVE SUBSTANCES

Fipronil	134 mg
(S)-Methoprene	120.6 mg

3. PACKAGE SIZE

1 x 1.34 ml
2 x 1.34 ml
3 x 1.34 ml
4 x 1.34 ml
6 x 1.34 ml

4. TARGET SPECIES

Dogs

5. INDICATIONS

For the treatment of dogs weighing 10 to 20 kg bodyweight.
Protects against infestations with fleas, alone or in combination with ticks and/or biting lice.

- Kills fleas (*Ctenocephalides* spp.) and protects against new infestations during 8 weeks.
- Kills ticks (*Ixodes ricinus, Dermacentor variabilis, Dermacentor reticulatus, Rhipicephalus sanguineus*) and protects against new infestations during 4 weeks.
- Kills biting lice (*Trichodectes canis*)
- Inhibits development of eggs, larvae and pupea of fleas (during 8 weeks in dogs) and prevents infestation of the environment of the treated animal with premature fleas during the same period.
- The veterinary medicinal product can be used as part of a strategy for the control of Flea Allergy Dermatitis (FAD) in dogs.

Fipronil kills fleas within 24 hours and ticks and lice within 48 hours after treatment.

Optional (to print on front side of the container):

- Prevents infestations in the environment of the treated animal with premature fleas
- Kills fleas, ticks and biting lice

6. ROUTES OF ADMINISTRATION

Spot-on

Use one pipet to treat one dog. External use only. The minimum treatment interval is 4 weeks.

Do not use on dogs less than 8 weeks old or weighing less than 2 kg.

The product can be used during pregnancy and lactation.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Store in the original package in order to protect from light.
Do not use after the expiry date stated on the package.

10. THE WORDS "READ THE PACKAGE LEAFLET BEFORE USE"

Read the package leaflet before use.

11. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only.

12. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Beaphar B.V.

14. MARKETING AUTHORISATION NUMBERS

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL ~~IMMEDIATE~~ PACKAGING UNITS

ALU foil (blister)

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combotec

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Fipronil 134 mg
(S)-Methoprene 120.6 mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

PP pipette

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combotec

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Fipronil 134 mg
(S)-Methoprene 120.6 mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

B. PACKAGE LEAFLET

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Combotec 134 mg/120.6 mg spot-on solution for medium dogs

2. Composition

Each pipette (1.34 ml) contains:

Fipronil	134 mg
(S)-Methoprene	120.6 mg
Butylhydroxyanisole (E320)	0.27 mg
Butylhydroxytoluene (E321)	0.13 mg

Clear amber spot-on solution

3. Target species

Dogs

4. Indications for use

For the treatment of medium dogs (10 to 20 kg bodyweight):
Protects against infestations with fleas, alone or in combination with ticks and/or biting lice.

- Kills fleas (*Ctenocephalides* spp.) and protects against new infestations during 8 weeks.
- Kills ticks (*Ixodes ricinus, Dermacentor variabilis, Dermacentor reticulatus, Rhipicephalus sanguineus*) and protects against new infestations during 4 weeks.
- Kills biting lice (*Trichodectes canis*)
- Inhibits development of eggs, larvae and pupea of fleas (during 8 weeks in dogs) and prevents infestation of the environment of the treated animal with premature fleas during the same period.
- The veterinary medicinal product can be used as part of a strategy for the control of Flea Allergy Dermatitis (FAD) in dogs.

Fipronil kills fleas within 24 hours and ticks and lice within 48 hours after treatment.

5. Contraindications

In the absence of available data, the veterinary medicinal product should not be used on puppies less than 8 weeks old and/or weighing less than 2 kg.
Do not use on sick (e.g. systemic diseases, fever) or convalescent animals.
Do not use on rabbits, as serious reactions, including death, could occur. In absence of studies, the use of the veterinary medicinal product is not recommended in non-target species.
Do not use in case of hypersensitivity to the active substances or to any of the excipients.
This veterinary medicinal product is specifically developed for dogs. Do not use in cats and ferrets, as this could lead to overdosing.

6. Special warnings

Bathing/immersion in water within 2 days after application of the veterinary medicinal product and more frequent bathing than once a week should be avoided, as no study has been performed to investigate how this affects the efficacy of the veterinary medicinal product. Emollient shampoos can be used prior to treatment, but reduce the duration of protection against fleas to approximately 5 weeks when used weekly after application of the veterinary medicinal product. Weekly bathing with a 2% chlorhexidine medicated shampoo did not affect efficacy against fleas during a 6 week long study. There may be an attachment of a few ticks. For this reason a transmission of infectious diseases cannot be completely excluded if conditions are unfavourable.
Fleas from pets often infest the animal's basket, bedding and regular resting areas such as carpets and soft furnishings which should be treated, in case of massive infestation and at the beginning of the control measures, with a suitable insecticide and vacuumed regularly.

Other animals living in the same household should also be treated with a suitable product.

Special precautions for safe use in the target species:
It is important to make sure that the veterinary medicinal product is applied to an area where the animal cannot lick it off and to make sure that animals do not lick each other following treatment. Avoid the contact with the animal's eyes.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
This veterinary medicinal product can cause mucous membrane, skin and eye irritation. Avoid contact of the veterinary medicinal product with mouth, skin and eyes. Wash hands after use. In case of accidental contact with the skin or eyes, rinse thoroughly with water. If irritation persists, seek medical advice.

This veterinary medicinal product may cause allergic reactions. People with known hypersensitivity to fipronil and/or (S)-methoprene should not treat their animal with this veterinary medicinal product.

This veterinary medicinal product may cause neurotoxicity and may be harmful if swallowed. Avoid ingestion including hand-to-mouth contact. Do not smoke, drink or eat during application. Keep the pipettes in the original packaging until ready for use and dispose of used pipettes immediately. Wash hands after use. In the event of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

Treated animals should not be handled until the application site is dry, and children should not be allowed to play with treated animals until the application site is dry. It is therefore recommended that animals are not treated during the day, but should be treated during the early evening, and that recently treated animals are not allowed to sleep with owners, especially children.

Special precautions for the protection of the environment:
Dogs should not be allowed to swim in watercourses for 2 days after application.
See also section on special precautions for disposal.

Other precautions:
The veterinary medicinal product may have adverse effects on painted, varnished or other household surfaces or furnishings.

Pregnancy and lactation:
The veterinary medicinal product can be used during pregnancy and lactation.

Overdose:
Do not overdose.

No adverse effects were observed in target animal safety studies in 8 week-old puppies, growing dogs and dogs weighing about 2 kg treated once at five times the recommended dose. The risk of experiencing adverse effects may however increase when overdosing, so animals should always be treated with the correct pipette size according to bodyweight.

Special restrictions for use and special conditions for use:
Other animals living in the same household should also be treated with a suitable product.

7. Adverse events

Dogs:

Very rare (<1 animal / 10 000 animals treated, including isolated reports):
Application site skin discolouration[1], Application site hair loss[1], Application site itching[1], Application site reddening[1]. Generalised itching or hair loss. Hypersalivation[2]; Vomiting; Respiratory signs; Increased sensitivity to stimulation[3], Central nervous system depression[3], Neurological signs (NOS)[2]

[1] Transient
[2] If licking occurs, a brief period of excessive salivation may be observed due mainly to the nature of the carrier.
[3] Reversible

Reporting adverse events is important. It allows continuous safety monitoring of a product. 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 contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder or its local representative using the contact details at the end of this leaflet, or via your national reporting system: {national system details}.

8. Dosage for each species, routes and method of administration

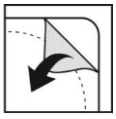
Spot-on use, by topical application to the skin.

One pipette of 1.34 ml per dog weighing over 10 and up to 20 kg, corresponding to a minimum recommended dose of 6.7 mg/kg for fipronil and 6 mg/kg for (S)-methoprene.
In the absence of safety studies, the minimum treatment interval is 4 weeks.

Underdosing could result in ineffective use and may favour resistance development.

9. Advice on correct administration

Use the easy-peel corners to remove a pipette from its blister. Do not puncture the foil with scissors, knives or other sharp instruments, as this may damage the pipette inside.



Hold the pipette upright. Tap the narrow part of the pipette to ensure the contents are within the main body of the pipette. Cut off the top of the pipette with scissors.

Part the coat between the shoulder blades until the skin is visible. Place the tip of the pipette on the skin and squeeze gently to empty its contents onto the skin.

Care should be taken to avoid excessive wetting of the hair with the product since this will cause a transient sticky appearance of hairs at the treatment spot.

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Keep out of the sight and reach of children.
Store in the original package in order to protect from light.
Do not use this veterinary medicinal product after the expiry date which is stated on the label. The expiry date refers to the last day of that month.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.

This veterinary medicinal product should not enter water courses as Fipronil and (S)-methoprene may be dangerous for fish and other aquatic organisms.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product not subject to prescription.

14. Marketing authorisation numbers and pack sizes

Cardboard box containing 1 aluminium blister with 1 pipette.
Cardboard box containing 1 aluminium blister with 2 pipettes.
Cardboard box containing 1 aluminium blister with 3 pipettes.
Cardboard box containing 1 aluminium blister with 4 pipettes.
Cardboard box containing 1 aluminium blister with 6 pipettes.

Not all pack sizes may be marketed.

15. Date on which the package leaflet was last revised

<{MM/YYYY}>
<{DD/MM/YYYY}>
<{DD month YYYY}>

Detailed information on this veterinary medicinal product is available in the Union Product Database (https://medicines.health.europa.eu/veterinary).

16. Contact details

Marketing authorisation holder:
Beaphar B.V.
Drostenkamp 3
8101 BX, Raalte
The Netherlands

Manufacturer responsible for batch release:
Beaphar B.V.
Oude Linderteseweg 9
8102 EV, Raalte
The Netherlands

Local representatives and contact details to report suspected adverse events:

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

België/Belgique/Belgien
{Nom/Naam/Name}
{Adresse/Adres/Anschrift }
BE-0000 {Localité/Stad/Stadt}
Tél/Tel: + {N° de téléphone/Telefoonnummer/Telefonnummer}
<{E-mail}>

Република България
{Наименование}
{Адрес}
BG {Град} {Пощенски код}
Тел: + {Телефонен номер}
<{E-mail}>

Česká republika
{Název}
{Adresa}
CZ {město}
Tel: +{telefonní číslo}
<{E-mail}>

Lietuva
{pavadinimas}
{adresas}
LT {pašto indeksas} {miestas}
Tel: + {telefono numeris}
<{E-mail}>

Luxembourg/Luxemburg
{Nom}
{Adresse}
L-0000 {Localité/Stadt}
Tél/Tel: + {N° de téléphone/Telefonnummer}
<{E-mail}>

Magyarország
{Név}
{Cím}
HU-0000 {Város}
Tel.: + {Telefonszám}
<{E-mail}>

Danmark
{Navn}
{Adresse}
DK-0000 {by}
Tlf.: + {Telefonnummer}
<{E-mail}>

Deutschland
{Name}
{Anschrift}
DE-00000 {Stadt}
Tel: + {Telefonnummer}
<{E-mail}>

Eesti
(Nimi)
(Aadress)
EE - (Postiindeks) (Linn)
Tel: +(Telefoninumber)
<{E-mail}>

Ελλάδα
{Όνομα}
{Διεύθυνση}
EL-000 00 {πόλη}
Τηλ: + {Αριθμός τηλεφώνου}
<{E-mail}>

España
{Nombre}
{Dirección}
ES-00000 {Ciudad}
Tel: + {Teléfono}
<{E-mail}>

France
{Nom}
{Adresse}
FR-00000 {Localité}
Tél: + {Numéro de téléphone}
<{E-mail}>

Hrvatska
{Ime}
{Adresa}
{Poštanski broj} {grad}
Tel: + {Telefonski broj}
<{e-mail}>

Ireland
{Name}
{Address}
{Town} {Postal code} - IE
Tel: + {Telephone number}
<{E-mail}>

Malta
{Isem}
{Indirizz}
MT-0000 {Belt/Raħal}
Tel: + {Numru tat-telefon}
<{E-mail}>

Nederland
{Naam}
{Adres}
NL-0000 XX {stad}
Tel: + {Telefoonnummer}
<{E-mail}>

Norge
{Navn}
{Adresse}
N-0000 {poststed}
Tlf: + {Telefonnummer}
<{E-mail}>

Österreich
{Name}
{Anschrift}
A-00000 {Stadt}
Tel: + {Telefonnummer}
<{E-mail}>

Polska
{Nazwa/ Nazwisko:}
{Adres:}
PL – 00 000{Miasto:}
Tel.: + {Numer telefonu:}
<{E-mail}>

Portugal
{Nome}
{Morada}
PT-0000–000 {Cidade}
Tel: + {Número de telefone}
<{E-mail}>

România
{Nume}
{Adresă}
{Oraş} {Cod poştal} – RO
Tel: + {Număr de telefon}
<{E-mail}>

Slovenija
{Ime}
{Naslov}
SI-0000 {Mesto}
Tel: + {telefonska številka}
<{E-mail}>

Ísland
{Nafn}
{Heimilisfang}
IS-000 {Borg/Bær}
Sími: + {Símanúmer}
<{Netfang}>

Italia
{Nome}
{Indirizzo}
IT-00000 {Località}
Tel: + {Numero di telefono}
<{E-mail}>

Κύπρος
{Όνομα}
{Διεύθυνση}
CY-000 00 {πόλη}
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17. Other information