Committee for Medicinal Products for Veterinary Use (CVMP)

CVMP assessment report for Versican Plus DHPPi/L4 (EMEA/V/C/003678/0000)
Common name: canine distemper, canine adenovirus, canine parvovirosis, canine parainfluenza virus (live attenuated) and canine leptospirosis (inactivated) vaccine

Assessment report as adopted by the CVMP with all information of a commercially confidential nature deleted.
**Introduction**

On 26 April 2013 the applicant Zoetis Belgium s.a. submitted an application for marketing authorisation to the European Medicines Agency (the Agency) for Versican Plus DHPPi/L4, through the centralised procedure under Article 3(2)(a) of Regulation (EC) No 726/2004 (new active substance).

The eligibility to the centralised procedure was agreed upon by the CVMP on 14 June 2012 as the product contains a new active substance. The rapporteur appointed was E. Werner and co-rapporteur T. Soós.

The dossier has been submitted in line with the requirements for submissions under Article 12(3) of Directive 2001/82/EC.

This vaccine is indicated for the immunisation of healthy puppies and dogs against canine distemper, adenovirus hepatitis, adenovirus respiratory disease, parvovirosis, parainfluenza and leptospirosis.

The route of administration is subcutaneous use.

On 13 March 2014, the CVMP adopted an opinion and CVMP assessment report.

On 7 May 2014, the European Commission adopted a Commission Decision granting a marketing authorisation for this veterinary medicinal product.

**Scientific advice**

Not applicable.

**Part 1 - Administrative particulars**

**Detailed description of the pharmacovigilance system**

The applicant has provided a detailed description of the pharmacovigilance system which fulfils the requirements of Directive 2001/82/EC. Based on the information provided the applicant has the services of a qualified person responsible for pharmacovigilance and the necessary means for the notification of any adverse reaction occurring either in the European Union (EU) or in a third country.

**Manufacturing authorisations and inspection status**

Antigen production, in-process testing, formulation, primary and secondary packaging, release testing and batch release takes place at

**BIOVETA, a.s.**
Komenského 212
683 23 Ivanovice na Hané
CZECH REPUBLIC

A valid manufacturing authorisation was presented in the dossier (dated 14 December 2011).

A valid good manufacturing practice (GMP) certificate for the Bioveta site dated 7 May 2012 was submitted.
**Overall conclusions on administrative particulars**

The detailed description of the pharmacovigilance system and the GMP certification of the manufacturing site were considered in line with legal requirements.

**Part 2 – Quality**

**Composition**

Versican Plus DHPPi/L4R contains per 1 ml dose:

Lyophilisate (live attenuated):

- Canine distemper virus, strain CDV Bio 11/A, \(10^{3.1} - 10^{5.1}\) TCID\(_{50}\)*
- Canine adenovirus type 2, strain CAV-2 Bio 13, \(10^{4.6} - 10^{5.3}\) TCID\(_{50}\)*
- Canine parvovirus type 2b, strain CPV-2b Bio 12/B, \(10^{4.3} - 10^{6.6}\) TCID\(_{50}\)*
- Canine parainfluenza type 2 virus, strain CPIV-2 Bio 15, \(10^{3.1} - 10^{5.1}\) TCID\(_{50}\)*

Solvent (inactivated):

- *Leptospira interrogans* serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain MSLB 1089, ALR** titre \(\geq 1:51\)
- *Leptospira interrogans* serogroup Canicola, serovar Canicola, strain MSLB 1090, ALR** titre \(\geq 1:51\)
- *Leptospira kirschneri* serogroup Grippotyphosa, serovar Grippotyphosa, strain MSLB 1091, ALR** titre \(\geq 1:40\)
- *Leptospira interrogans* serogroup Australis, serovar Bratislava, strain MSLB 1088, ALR** titre \(\geq 1:51\)

* Tissue culture infectious dose 50%.
** Antibody micro agglutination-lytic reaction.

Aluminium hydroxide gel is used as an adjuvant.

**Container**

Each fraction of the vaccine is filled into Type I glass vials. The vials are tested in accordance with European Pharmacopoeia (Ph. Eur.) monograph 3.2.1. The glass vials for the freeze-dried fraction are closed with 13 mm bromobutyl rubber stoppers, and the glass vials for the liquid fraction are closed with 13 mm chlorobutyl rubber stoppers. The rubber stoppers are tested in accordance with Ph. Eur. monograph 3.2.9.

The rubber stoppers are sealed with a 13 mm flip off aluminium cap. Corresponding certificates of analysis are provided. The outer packaging will be a transparent plastic box containing 25 or 50 vials of 1 dose of the lyophilisate and 25 or 50 type vials of solvent accordingly.

**Development pharmaceutics**

The Versican Plus DHPPi/L4 has been developed for the prevention of infectious diseases caused by canine distemper virus, canine parvovirus, canine adenovirus and canine parainfluenza virus (live components) and against leptospirosis (inactivated component). The product was based on the preceding vaccine Versican DHPPi/L3R. This vaccine contains canine distemper virus (CDV), canine adenovirus type 2 (CAV-2), canine parvovirus type 2 (CPV-2) and canine parainfluenza virus (CPIV) and three leptospiras: *Leptospira interrogans* serovar Canicola, *L. interrogans* serovar Icterohaemorrhagiae and *Leptospira*
kirschneri serovar Grippotyphosa. The Versican Plus DHPPi/L4 contains a new strain of canine parvovirus (2b) and one additional leptospira strain from serogroup Australis serovar Bratislava.

Method of manufacture

The manufacturing procedure is adequately described in detail to give sufficient confidence that the product will be safe, effective and stable.

The manufacturing process includes: Preparation of medium and inocula, inoculation of the cell line or growth media and growth of culture, termination of the cultivation. For the antigens for liquid fraction another procedures are described as inactivation and concentration/purification. Each leptospira antigen is purified and concentrated. The liquid fraction is blended with adjuvant (aluminium hydroxide gel).

Control of starting materials

Active substance

All 8 antigens comprised in the vaccine (i.e. CDV, CAV, CPV-2b, CPIV-2 and four leptospira antigens) are sufficiently described with regard to their origin, isolation and history. The control testing on the viral and bacterial seeds is performed in accordance with the relevant guidelines. This control testing is considered satisfactory and purity of the seed materials is sufficiently justified with regard to the risk of contamination of the materials from pathogens of the species of origin and the risk for the target species.

Excipients

Certificates of analysis of starting materials listed in Ph. Eur. monographs were provided and are satisfactory.

Up-to date the European Directorate for the Quality of Medicines and HealthCare (EDQM) certificates and/or certificates of analysis for substances of biological origin used during production were provided. Certificates of analysis of the starting materials of non-biological origin were provided and are satisfactory. Details of in-house preparation of media were provided.

Specific measures concerning the prevention of the transmission of animal spongiform encephalopathies

A detailed list of materials of animal origin included in the scope of the Note for guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products and of materials from animals other than those included in the scope of the Note for guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via human and veterinary medicinal products (EMA/410/01 rev.3) was provided in the dossier together with a transmissible spongiform encephalopathy (TSE) risk assessment. The substances comply with the TSE Note for guidance and Commission Directive 1999/104/EC.

The TSE risk assessment is provided related to the use of the bovine serum in the manufacturing process. The material is sourced only from countries classified by geographical BSE risk as either highly unlikely or unlikely but not excluded category. Therefore it can be concluded that the risk of TSE contamination from bovine serum is negligible.

The seed materials are considered in the risk assessment with regard to their origin (non-ruminant), method of preparation and recipient species of the vaccine (dogs, known as not susceptible to TSE)
The CVMP concluded that the TSE risk has been adequately addressed and is considered negligible.

**Control tests during production**

**Freeze-dried fraction:**

The control tests during production include sterility, absence of mycoplasma, virus titre, cell count, identity (CPV-2b), pH value and sterility after blending. The in-process tests are deemed to be sufficient to control all the critical steps in the manufacture.

**Liquid fraction:**

The following tests are performed for *Leptospira* components: growth, purity (during cultivation), identity, cell count (before inactivation), *Leptospira* inactivation, sterility (after inactivation), serum absence, sterility and formaldehyde (after concentration and purification).

After vaccine blending the bulk is tested for aluminium content, pH and sterility.

The in-process tests are deemed to be sufficient to control all the critical steps in the manufacture.

**Control tests on the finished product**

The description of the methods used for the control of the finished product and the specifications are provided:

**Freeze-dried fraction:**

The control tests of the finished freeze-dried fraction include appearance, sterility, extraneous agents, absence of mycoplasma, virus identity and titre, residual humidity and vacuum.

**Liquid fraction:**

The control tests of the finished liquid fraction include appearance, sterility, air tightness, extractable volume, *Leptospira* identity and potency.

**Reconstituted vaccine:**

The control tests of the reconstituted vaccine include appearance and pH determination.

A detailed overview of all in-process and finished product tests of the vaccine Versican Plus DHPPi/L4 has been submitted.

The finished product tests are considered adequate to control the quality of the finished product.

**Stability**

The proposed shelf life of the vaccine Versican Plus DHPPi/L4 is 2 years when stored at 2–8 °C.

In order to support the proposed shelf life of the finished product three batches of each final fraction of the vaccine Versican Plus DHPPi/L4R (largest combination of the Versican Plus range of vaccines) were manufactured in accordance with the proposed method. Vials from each batch were stored over a period of 27 months and tested at regular intervals at 0, 3, 6, 12, 18, 24 and 27 months in accordance with finished product specifications. In addition, at 0, 24 and 27 months of storage samples from each batch were reconstituted and tested for appearance, pH and sterility in accordance with the finished product specifications.
The proposed shelf life of 2 years can be accepted.

However, the final report on the antigen shelf life including the inactivated antigen stability data will still need to be submitted as soon as it is available. A relevant recommendation has been included in the report.

**Overall conclusions on quality**

The analytical part of the dossier is detailed and clearly states the production and control of this immunological veterinary medicinal product and demonstrates that it complies with the requirements of Directive 2001/82/EC.

All necessary information concerning qualitative and quantitative composition is submitted.

The choice of the vaccine strains and the adjuvant has been satisfactorily addressed and reference to the relevance of each strain to current epidemiological conditions is also provided.

The manufacturing process of the vaccine has been described in detail for the virus and for the *Leptospira* components. Regarding the starting materials all necessary information has been provided.

The TSE risk assessment provided by the applicant clearly demonstrates that the TSE risk of this product is negligible. Compliance with the corresponding Note for guidance is demonstrated.

Controls during manufacture and tests on the finished product are suitable to guarantee the compliance with the quality parameter mentioned. Test methods have been described and corresponding validation studies have been performed.

Batch to batch consistency has been demonstrated and a detailed overview of all in-process and finished product tests of the vaccine Versican Plus DHPPi/L4 has been provided. Based on the data provided the proposed shelf life for the finished product of 2 years can be accepted. The CVMP recommended that the final study results be provided when available in order to confirm final results for the antigen shelf life.

Based on the review of the data on quality, the manufacture and control of Versican Plus DHPPi/L4 are considered acceptable.

In addition, the applicant is recommended to provide the following information post-authorisation:

1. The antigen stability study is due to complete by the end of 2014. The final study report should be submitted during the first quarter in 2015.
2. Regarding the *Leptospira* components, for the first 10 batches produced for commercial release, the individual results of the tested rabbits including the results for all control sera (negative, hyperimmune and positive standard sera) should be submitted along with the batch release protocols.
3. The final batch release protocol of the filled batch (batch number 015820) should be provided by the end of March 2014.

**Part 3 – Safety**

Versican Plus DHPPi/L4 is a combined live virus and inactivated bacterial vaccine indicated for the immunisation of healthy puppies and dogs against canine distemper, adenovirus hepatitis, adenovirus respiratory disease, parvovirosis, parainfluenza and leptospirosis.
The applicant presented laboratory vaccination studies (safety of a single and repeated dose, overdose, clearance, shed and spread of the vaccine, increase in virulence) and two field studies to support the safety of this vaccine.

**Laboratory tests**

Methods and corresponding validations for the serological and viral isolation tests used in the clinical studies were provided.

The safety of a single and repeated dose were demonstrated using the largest combination of the Versican Plus range of vaccines, Versican Plus DHPPi/L4R, which includes an inactivated rabies component. All components of Versican Plus DHPPi/L4R are identical to those contained in Versican Plus DHPPi/L4 with only the number of component differing. This is in compliance with CVMP note for guidance: requirements for combined veterinary vaccines (CVMP/IWP/52/97) and also the CVMP Guideline on the requirements for combined vaccines and associations of immunological veterinary medicinal products (IVMPs) (EMA/CVMP/IWP/594618/2010).

**Safety of the administration of one dose**

The safety of a single dose was assessed together with the safety of repeated administration of a single dose using the fully-valent vaccine Versican Plus DHPPi/L4R. Sixteen 6 week old puppies, free of antibodies against CDV, CAV-1, CAV-2, CPV, CPIV, Leptospira and rabies virus were vaccinated subcutaneously four times at an interval of 14 days, thus the vaccination scheme applied in the study differed from the vaccination scheme in accordance with the summary of product characteristics (SPC) which recommends as basic vaccination two doses 3–4 weeks apart from 6 weeks of age. The vaccination titre per dose (1 ml) correlates with the maximum proposed titres given in the SPC. All animals were observed at defined points in time for signs of abnormal local reactions including heat, pain, erythema and swelling, and systemic reactions including rectal temperatures.

During the study no systemic reactions were observed. At the injection sites, soft painless swellings were found. The maximum diameter after the first administrations was 25 mm. The maximum duration for until disappearance was 17 days. The rectal temperatures of all animals remained within the physiological range after each administration. In general, it can be concluded that the administration of one dose of Versican Plus DHPPi/L4 containing maximum potency of antigens by the recommended route was safe for puppies of 6 weeks of age.

**Safety of one administration of an overdose**

**Overdose safety study of the vaccine with the DHPPi component (Versican Plus DHPPi) in minimum age pups**

This study was performed in compliance with good laboratory practice (GLP) and in accordance with Annex I of Directive 2001/82/EC and International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH) GL44 on target animal safety for veterinary live and inactivated vaccines and the Ph. Eur. general monograph on safety. Sixteen 6 week old puppies free of antibodies against CDV, CAV-1, CAV-2, CPV and CPIV were vaccinated subcutaneously with a tenfold maximum dose. All animals were observed at defined points in time for signs of abnormal local reactions including heat, pain, erythema and swelling, and systemic reactions including rectal temperatures. During the study no systemic reactions were observed. Immediately after the vaccine administration, four out of eight animals showed evidence of pain at the injection site (scratching at
The signs of pain persisted for 10 seconds to 1 minute. At the injection sites, soft, painless swellings were found, which resolved within one day.

The rectal temperatures of all animals remained within the physiological range after each administration. White blood cell counts showed that no animal developed leukopenia.

In this study, only the live virus components CDV, CAV-2, CPV-2b and CPIV, diluted in water for injection instead of the actual diluent were investigated. This was justified in a sufficient manner. Further overdose testing is no longer required for inactivated vaccines.

**Master seed studies**

**Safety testing of the canine distemper virus (CDV)**

Canine distemper master seed was tested in the target species of animals according to Ph. Eur. monograph 0448 “Canine distemper vaccine (live)” – Section 2-3-1 “Safety” and based on the results, the CDV strain was considered to have met the Ph. Eur. requirements. The study proved the safety of the administration of ten-fold maximum via the subcutaneous route to 6 week old puppies.

**Irreversibility of attenuation - canine adenovirus type 2**

This study is discussed and presented in detail under "Increase in virulence – CAV-2 strain". The CAV-2 strain was generally classified as safe since no adverse effects were observed in further studies.

**Safety testing of the canine parvovirus (CPV-2b)**

Canine parvovirus master seed in first passage was tested in the target species according to Ph. Eur. monograph 0964 "Canine parvovirus vaccine (live)" – Section 2.3-1-1 "Safety" and Section 2-3-1-2 "Effect on the thymus". Based on these results the CPV-2b strain was considered to have met the requirements of the Ph. Eur. Therefore, the study proves the safety of the administration of $10^{8.5}$ TCID$_{50}$/dose via the subcutaneous route to 4 week old puppies.

**Irreversibility of attenuation of vaccine strain – canine parainfluenza virus (CPIV)**

This study is discussed and presented in detail under point “Increase in virulence – CPIV strain”. The CPIV strain was generally classified as safe since no adverse effects were observed in further studies, e.g. in the overdose study where the ten-fold dose of the maximum titre was administered.

**Safety of the repeated administration of one dose**

For details of the study design and the results see the above section on “Safety of the administration of one dose”.

The repeated administration of a single dose was presented and assessed together with the safety of a single dose. Four repeated administrations were included in the design to take account of the number of administrations for primary vaccination (2–3) and the first re-vaccination (1).

For primary vaccination two doses are recommended, but occasionally three doses may be required in areas of high risk of infection with CPV or CDV.

During the study no systemic reactions were observed. At the injection sites, soft painless swellings were found. After 4 injections a maximum of 30 mm in diameter was observed. The maximum duration for disappearance was 17 days.

The applicant chose to administer the single doses for this study four times with a 14-day interval between the administrations. While this vaccination schedule differs from the recommended SPC interval...
of 3–4 weeks, it does constitute a worse case and additionally is in line with VICH GL44 which permits shortening the intervals between administrations for repeat dose studies to at least 14 days.

Overall, it can be concluded that the administration of the repeated dose of Versican Plus DHPPi/L4 containing maximum potency of antigens by the recommended route was safe for puppies of 6 weeks of age.

**Examination of reproductive performance**

No studies have been performed on reproductive safety and a warning in this respect is included in the SPC.

The Ph. Eur. monograph 5.2.6 on requirements for examination of reproductive performance of males and non-pregnant female dogs was also considered. This requirement is applicable if the vaccine contains organisms which are known as reproductive pathogens.

CDV, CAV-2, CPV and CPiV (the live viral components of Versican Plus DHPPi/L4) are not generally associated with any pathological effects in the reproductive tract of the male or the non-pregnant female. The other components of the vaccine are inactivated viral and bacterial antigens, the aluminium hydroxide gel (adjuvant) and some inert excipients, which again are not generally associated with any pathological effects in the reproductive tract of the male or the non-pregnant female. The CVMP therefore concluded that there is no justification for requesting further examinations of reproductive performance. However, as no data regarding safety aspects of vaccine administration to pregnant or lactating dogs are available a warning is included in the SPC section 4.7: "Therefore the use is not recommended during pregnancy and lactation".

**Examination of immunological functions**

Ph. Eur. monograph 5.2.6 gives the following guidance concerning investigation of adverse effects on immunological functions: Where the product might adversely affect the immune response of the animal to which the product is administered or of its progeny, suitable tests on the immunological functions are carried out.

Except for the canine parvovirus (CPV) component, none of the components of Versican Plus DHPPi/L4 are known to have any adverse effects on the immune functions of infected animals. CDV, CPiV and CAV-2 are already known as components of Versican DHPPi/L3R. No adverse effects are known for these antigens.

Wild-type strains of CPV can cause leucopenia and thymic damage. Ph. Eur. monograph 0964 includes some specific requirements concerning the safety testing of live CPV vaccines, namely investigations of the effects of a 10-fold overdose on white blood cell count and on thymic histology. In safety study 9b data on this issue are presented and discussed. No signs of leucopenia or hypoplasia of thymus were found. The study demonstrates that this vaccine strain does not have any adverse effects on the immunological functions.

No studies have been performed on the concurrent use of any other non-Versican Plus vaccine. This fact is covered in section 4.8 of the SPC as follows:

"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 by the veterinarian."
**Special requirements for live vaccines**

**Spread of the vaccine strain**

**Specific requirements for live vaccines – canine parvovirus (CPV-2b)**

This study evaluates the dissemination, shedding and spread of the modified live CPV-2b vaccine strain of the Versican Plus DHPPi/L4R vaccine.

Six dogs were vaccinated once subcutaneously with CPV-2b vaccine strain. Six unvaccinated dogs were kept as in-contact controls and housed together with the vaccinated dogs. The six control animals were the moved and housed together with a new group of six control dogs. The details of group allocation provided were adequate.

The animals were examined for abnormal clinical observations or systemic reactions; rectal temperatures were recorded and the injection site was measured. No abnormalities or deviations were observed.

All vaccinated dogs and five animals of control group 1 had seroconverted as well as all dogs of control group 2.

Faecal samples for detection of vaccine virus by reverse transcription polymerase chain reaction (RT-PCR) were taken at defined time points. CPV could be detected in samples from vaccinated animals and animals of control group 1 and 2. The vaccine virus was found to be shed by faeces from all vaccinated and control animals. No quantitative virus titre determination was performed.

Organ samples were taken from the vaccinated animals to detect vaccine virus. The vaccine virus disseminated to thymus, spleen and mesenteric lymph nodes of the vaccinated animals. No information on or evaluation of the dissemination in the control group animals were provided.

The live attenuated vaccine virus strain CPV-2b may be shed by vaccinated animals for a number of days following vaccination and passed on to non-vaccinated dogs. The results show that the CPV-2b vaccine strain spreads from vaccinated puppies to other non-vaccinated dogs. Dogs infected through contact with vaccinated animals were able to act as a source of infection by spreading CPV-2b to a second contact group. Antibodies were detected in all animals (except for 1) of all groups. None of the dogs, which were infected with vaccine strain CPV-2b, developed any abnormal clinical signs. An appropriate statement is given in the SPC under section 4.5.

The applicant did not provide other information on the spread to other possibly susceptible non-target species. According to publications it is however known that CPV-2b can infect cats and cause parvovirus disease. Even though the vaccine strain is attenuated, it is to be expected that cats may become infected with the vaccine virus from vaccinated dogs since dogs and cats commonly have close contact. Several species of the family Canidae have been reported in publications to be susceptible as well and infection of wildlife carnivores by domestic cats may be feasible. Therefore, transmission of vaccine virus from vaccinated dogs in the wildlife population cannot definitely be excluded. Furthermore, due to the high tenacity and low infection dose of paroviruses, direct contact between individuals may not be necessary to transmit the virus to other susceptible species.

In the absence of data on the pathogenicity of the vaccine strain for domestic cats and other carnivores, a warning has been included in the SPC under section 4.5 "Special precautions for use" to reflect the need to separate vaccinated dogs from other canine and feline species after vaccination.

**Specific requirements for live vaccines – canine distemper virus (CDV)**

This study was intended to assess the safety of canine distemper master seed virus in terms of its spread and dissemination after administration by the subcutaneous route to animals of six weeks of age.
Six unvaccinated dogs were kept as in-contact controls, housed together with vaccinated dogs. The six control animals were then moved and housed together with a new group of six control dogs. All animals showed normal general health for the duration of the study. No adverse reactions were observed in the vaccinated animals. Temperatures of all animals remained within the physiological norm. Serology (serum neutralization test) to CDV was assessed. All vaccinated dogs had seroconverted, whereas all controls remained seronegative. In blood samples CDV was detected in one vaccinated animal. No virus was detected in samples from the other five vaccinated animals or any control animal. No CDV was detected in nasal swabs, samples of nasal secretions, urine, faeces and saliva by RT-PCR.

CDV was detected in lung lymph nodes of one vaccinated animal only. Based on the negative serology and virus isolation results of the first control group and the negative serology results of the second control group, secretion samples (nasal, faecal, urine and oral) of control group 2 animals were not tested for the presence of CDV. The provided studies demonstrate that no CDV is excreted. The vaccine strain Onderstepoort is known to be well attenuated. Therefore, the probability of transmission to non-target species is deemed to be negligible. No specific warning in SPC was considered necessary.

The study results show that the vaccine CDV strain, when administered by the recommended route, is not capable to spread or disseminate from vaccinated to unvaccinated puppies.

**Canine adenovirus type 2 (CAV-2)**

The objective of the study was to demonstrate potential spread of canine adenovirus type 2 vaccine strain (CAV-2) from vaccinated animals to in-contact naïve animals by evaluating the seroconversion of the control animals.

Five puppies were once vaccinated subcutaneously with CAV-2 vaccine strain. Five control puppies were kept in contact with the vaccinated animals. General health observations were carried out daily. Blood samples were collected from all puppies pre-vaccination and at defined points in time. The virus neutralizing antibodies to CAV-2 were analysed.

There was an increase in the neutralizing antibodies to CAV-2 in the 5 vaccinated animals. There were no detectable neutralizing antibodies to CAV-2 in the control animals. No secretion samples were taken to demonstrate the presence or absence of virus.

If seroconversion was not found in the co-housed control animals, it was concluded that infection by secretion must have been absent or of very low titre. The CAV-2 component of the vaccine in question corresponds to the CAV-2 strain of the product Versican DHPPi/L3R already authorised in the EU via the decentralised procedure. For the latter vaccine the SPC contains a warning with regard to potential virus excretion lasting several days after vaccination.

The study results showed that the vaccine CAV-2 strain, when administered by the recommended route and titre, is not capable to spread from vaccinated dogs to in-contact control animals and induce an active infection and therefore, section 4.5 of the SPC includes a statement reflecting the findings:

"The live virus vaccine strains CAV-2, CPIV and CPV-2b may be shed by vaccinated dogs for a number of days following vaccination. Due to the low pathogenicity of CPV-2b-strain, it is not necessary to keep vaccinated dogs separated from non-vaccinated dogs.

The vaccine virus strain CPV-2b has not been tested in domestic cats and other carnivores (except dogs) that are known to be susceptible to canine parvoviruses. Therefore vaccinated dogs should be separated from other canine and feline species after vaccination."
Canine parainfluenza virus 2 (CPiV-2)

The objective of the study was to demonstrate potential spread of CPiV-2 vaccine strain from vaccinated animals to in-contact naïve animals by evaluating the seroconversion of the control animals.

Five puppies were once vaccinated subcutaneously and five control puppies were kept in contact with the vaccinated animals.

General health observations were carried out daily until the end of the study on Day 84. No abnormal health observations were made. Blood samples were collected from all puppies at defined points in time. The virus neutralizing antibodies to CPiV-2 were analysed. No organ samples were taken. There was an increase in neutralizing antibodies to CPiV-2 in the five vaccinated animals. There were no detectable neutralizing antibodies to CPiV-2 in the control animals at any of the points in time of the test. The study results show that the vaccine CPiV-2 strain, when administered by the recommended route and titre, is not capable to spread from vaccinated dogs to in-contact control animals and induce an active infection.

Dissemination in the vaccinated animals:

Canine distemper virus

Dissemination of the CDV vaccine strain was investigated together with shedding and spread in Safety Study 4. Furthermore, the virus isolation data from the CDV reversion to virulence study Safety Study 3, discussed below, provides some information on the dissemination of the CDV vaccine strain.

Specific requirements for live vaccines – Canine distemper virus (CDV)

This study was already discussed and the design presented under point “Spread of the vaccine strain”.

Six unvaccinated dogs were kept as in-contact controls, housed together with vaccinated dogs, the six control animals were then moved and housed together with a new group of six unvaccinated control dogs. Nasal swab samples were collected from all animals. Samples of faeces, urine, oral and nasal secretions were taken from all vaccinated animals. Vaccinated animals were euthanized on study Day 14 and samples of brain, nasal mucosa, tonsils, thymus, spleen and the lungs and their local lymph nodes were collected.

In this study the CDV was found to disseminate to blood and mediastinal lymph nodes in a minority of vaccinated animals when administered to seronegative dogs. No dissemination to the brain, nasal mucosa, tonsils, thymus, spleen or lungs was detected. No CDV was detected in nasal swabs, samples of nasal secretions, urine, faeces and saliva.

The study results show that the vaccine CDV strain, when administered by the recommended route, is not capable to spread or disseminate from vaccinated to unvaccinated puppies.

Safety testing of the Canine distemper virus (CDV) – Lack of increase in virulence

This study is discussed and the design presented under point “Reversion to virulence of attenuated vaccines”.

The virus was passaged in two series. CDV could be detected by polymerase chain reaction (PCR) in pooled samples from nasal mucosa, tonsils, thymus, spleen, lungs and mediastinal lymph nodes of animals 5–8 days after inoculation with CDV. As virus could not be recovered from the 1st passage animals, the first set of passages was terminated and a second set of passages started. No vaccine virus was detected in any samples by cultivation in cell lines.
Canine adenovirus type 2

Dissemination of canine adenovirus type 2

The objective of the study was to investigate the presence of the canine adenovirus 2 (CAV-2) vaccine strain in the organs of vaccinated animals by isolation of the vaccine virus on cell cultures.

The dissemination of CAV-2 was evaluated following administration of the CAV-2 vaccine strain. Four puppies were once vaccinated subcutaneously.

CAV-2 was isolated from the mediastinal, cervical, mesenteric and axillary lymph nodes. No vaccine virus was found in nasal mucous membrane, larynx mucous membrane, tonsils, lungs, spleen, liver and kidney. Only 4 animals were used for this study. This seems to be acceptable because the CAV-2 Reversion to Virulence Study – Irreversibility of Attenuation - canine adenovirus type 2 provides some more information on dissemination in tissues. These data show that after administration of a high titre of the CAV-2 to 6/7-week old dogs vaccine virus was isolated by cell culture from the larynx mucosa, tonsils, lungs, and cervical and axillary lymph nodes, but not from the nasal mucosa, spleen, liver, kidney, mesenteric or mediastinal lymph nodes.

No abnormal health observations were made.

No secretion samples were taken to demonstrate the presence or absence of vaccine virus as required by Ph. Eur. monograph 5.2.6, Section 1-8-2 "Dissemination in vaccinated animals". The SPC section 4.5 was therefore updated as explained above:

"The live virus vaccine strains CAV-2, CPiV and CPV-2b may be shed by vaccinated dogs for a number of days following vaccination. Due to the low pathogenicity of CPV-2b-strain, it is not necessary to keep vaccinated dogs separated from non-vaccinated dogs.

The vaccine virus strain CPV-2b has not been tested in domestic cats and other carnivores (except dogs) that are known to be susceptible to canine parvoviruses. Therefore vaccinated dogs should be separated from other canine and feline species after vaccination."

Canine parvovirus

Data relevant to dissemination of the CPV-2b strain were investigated in the Shed & Spread study. The virus isolation data from the CPV-2b reversion to virulence study Safety Study, discussed in part “Increase in virulence” provide some information on the dissemination and excretion of the CPV-2b vaccine strain.

Specific requirements for live vaccines – canine parvovirus (CPV-2b)

This study is discussed and the design presented under point "Spread of the vaccine strain".

Six dogs were vaccinated with the CPV-2b. Six unvaccinated dogs were kept as in-contact controls, housed together with the vaccinated dogs. The six control animals were then housed together with a new group of control dogs.

All vaccinated and control animals excreted vaccine virus in faeces.

As demonstrated by examination of the organs the vaccine virus disseminated to thymus, spleen and mesenteric lymph nodes of the vaccinated animals.

Please see evaluation under "Spread of the vaccine strain".

Safety testing of the canine parvovirus – Irreversibility of Attenuation

This study is discussed and the design presented under point "Increase in virulence – CPV-2b strain".
In a first step, 2 dogs were once vaccinated subcutaneously with the CPV-2b strain. Faecal samples were collected and vaccine virus was isolated. These virus samples were administered oronasally to another two dogs. In total the vaccine virus was passaged through 5 pairs of animals.

In a second step, 8 dogs were inoculated oronasally with passaged virus (derived from the final passage of the first step). Eight dogs were vaccinated with un-passaged virus.

During the passages excretion of vaccine virus in faeces could be shown. No further secretion samples, e.g. oral or nasal secretion were taken and tested.

Please see evaluation under "Increase for virulence”.

**Canine parainfluenza virus**

As stated by the applicant a separate study to investigate the dissemination of the CPIV vaccine strain has not been undertaken. CPIV is not a recognised zoonosis, hence the applicant believes that under the terms of Ph. Eur. monograph 5.2.6 it is not necessary to undertake a study of dissemination of this vaccine strain in the body. Furthermore, according to the publications CPIV is known to affect only the surface epithelium of the respiratory tract and not to cause a systemic infection.

While the applicant has not performed a separate study to investigate the dissemination of the CPIV vaccine strain reference is made to study “Irreversibility of Attenuation of Vaccine Strain – Canine parainfluenza virus” and Safety Study "CPIV strain: Spread of the vaccine strain”.

In study on irreversibility of attenuation nasal swabs were taken after vaccination and isolated vaccine virus was passaged 5 times. Data show that this virus disseminates to the nasal mucosa and is excreted between approximately from 6 to 9 days after vaccination. No further secretion samples, e.g. faeces, were taken and tested. Spread of virus cannot be excluded. The CVMP therefore concluded that virus dissemination has been shown and shedding is likely to take place. Nevertheless, due to the low pathogenicity of these strains, it is not necessary to keep vaccinated dogs separated from non-vaccinated dogs.

In study on the potential spread of the vaccine virus strain CPIV-2 from vaccinated animals to in-contact animals by evaluating the seroconversion of the control animals to CPIV-2, an increase of neutralizing antibodies to CPIV-2 in the vaccinated animals could be observed, but there was no increase in neutralizing antibodies in the control animals.

Please see assessment under the corresponding studies.

**Reversion to virulence of attenuated vaccines**

**Canine distemper virus**

The increase in virulence of canine distemper master seed virus has to be evaluated according to Ph. Eur. monograph 0448 on canine distemper vaccine (live) – section 2-3-2 Increase in virulence.

In a first series of passages 2 dogs were vaccinated subcutaneously with a high dose of the CDV strain. As vaccine virus could not be recovered from the first passage animals the first set of passages was terminated and a second set of passages was started. For a second passage again two dogs were inoculated subcutaneously.

The study design was changed from those for the first set of passages to increase the probability of re-isolating vaccine virus. Apart from that the study procedure was the same.

There were no abnormal clinical observations or systemic reactions. No increase in body temperature was observed after the administrations of the tested material and/or organ suspension. No local reactions
were recorded after CDV vaccine strain administration. No vaccine virus was isolated from the organ cultures when tested in cell culture. However, the semi-quantitative RT-PCR assay confirmed presence of virus in organ suspensions from both animals administrated the maximum vaccine titre.

The amount of CDV administered corresponds to approximately ten-times the assumed maximum dose for the commercial vaccine. According to the SPC the maximum titre for CDV is $10^{5.1} \text{ TCID}_{50}$. Ph. Eur. monograph 0448 does not require tenfold of the maximum titre for the reversion to virulence study.

The results show that the virus could not be recovered at the first passage level in two series of passages. However, the Ph. Eur. monograph requires administering a quantity of the vaccine virus that will allow recovery of the virus however the strain is attenuated such that this was not possible.

**Canine adenovirus type 2**

This study was performed in 1996 and was part of the application for the vaccine Versican DHPPi/L3R to confirm the irreversibility of attenuation of the CAV-2 vaccine strain. The strain is identical with the CAV-2 strain of the new Versican Plus DHPPi/L4 vaccine.

**Step 1: Passage procedure:**
The CAV-2 vaccine strain was passaged sequentially in twelve 6 to 7 week old puppies free from antibodies against CAV-2. The first two puppies were administered subcutaneously CAV-2. Five to six days post-inoculation the 2 dogs were euthanized and tissue samples were collected and examined for the presence of virus. Positive samples were pooled and 1 ml of the suspension was administered intranasally to two other puppies for the first passage. The above operation was carried out four additional times and the presence of virus was verified at each passage.

**Step 2: Subsequent safety evaluation:**
CAV-2 positive passage material from the last two puppies (terminal passage) was administered intranasally to five puppies (Group 2). In addition, CAV-2 was directly vaccinated subcutaneously to another five puppies (Group 1).

All ten animals were blood sampled at defined points in time for determination of CAV-2 specific antibodies. General health observations were made and rectal temperatures were measured. In the subsequent safety evaluation (Step 2), no neutralizing antibodies to CAV-2 were detected before inoculation in any of the animals. Fourteen days after inoculation, titres in all animals were increased at least fourfold. These results confirm an immune response to CAV-2 antigen following inoculation. No canine adenovirus could be isolated from the tested tissues 14 days after inoculation. This study shows no indication of increased virulence of the vaccine CAV-2 and is in compliance with Ph. Eur. monograph 1951.

**Canine parvovirus**

The increase in virulence of canine parvovirus master seed virus has to be evaluated according to Ph. Eur. monograph 0964 Canine parvovirus vaccine (live) – section 2-3-2 Increase in virulence.

**Step 1: Passage procedure:**
Five MSV passages were performed in this study. In the first passage, two dogs were vaccinated subcutaneously with CPV-2b. The administered vaccine virus titre was at least ten times the maximum virus titre that can be expected in a dose of commercial vaccine.

Samples of faeces were collected at defined points in time. The sample with the highest virus titre was used for the inoculation of the next two animals. Samples of thymus from both animals inoculated with test material were collected after euthanasia.
Two new dogs were inoculated oronasally with 1 ml of the suspension of canine parvovirus recovered from the faecal sample from the first couple of animals. Samples of faeces were collected. Samples positive for virus presence were pooled and used for inoculation of further two animals, which were used for the third virus passage. 1ml of the pooled faecal sample from the previous two animals was administered oronasally. Two further passages, fourth and fifth virus passage, were performed on a couple of animals respectively. Samples of the fifth passage positive for virus presence were pooled and used for the inoculation of eight animals for a sixth passage and a final comparison of the maximal attenuated virus passage with the initial material.

Step 2: Subsequent safety evaluation:
Sixteen animals were used for a final safety evaluation. One ml of the pooled faecal suspension from the last passage (fifth) was administered oronasally to each of eight animals and the first passage from MSV was administered subcutaneously to each of another eight animals.

The results show that animals administered with un-passaged and first passage suspension showed seroconversion, other groups remained seronegative. Leucopenia was not observed after MSV administration.

During passage procedure excretion of virus in faeces was shown by haemagglutination (HA) test and isolation in sensitive cell cultures in animals inoculated with first passage only. In all other groups of animals the excretion of virus was shown by a PCR method.

During the final safety evaluation the excretion of virus in faeces was shown in animals administered CPV-2b by HA and PCR method. No excreted virus was detected in animals inoculated with high passage CPV, neither by HA test nor PCR. No further secretion samples, e.g. oral or nasal secretion were taken and tested. Excretion of CPV vaccine strains was only demonstrated in faeces as this is known to be the main route of excretion. A special warning is already implemented in the SPC.

The study results implicate that the master seed of canine parvovirus, when administered by the recommended route, does not increase in virulence when passaged in puppies.

**Canine parainfluenza virus**

The safety evaluation was performed in 2001 for the Versican DHPPi/L3R vaccine. The strain is identical with the CPiV strain of the new Versican Plus DHPPi/L4 vaccine.

Step 1: Passage procedure:
Irreversibility of attenuation of the CPiV vaccine strain was evaluated by sequential passaging of the CPiV in twelve puppies. Nasal swabs were collected and tested for presence of the virus. Nasal swabs with the maximum amount of virus were selected and 1 ml of the suspension was administered intranasally to two other puppies for the first passage. The above operation was carried out five additional times and the presence of virus was verified at each passage.

Step 2: Subsequent safety evaluation:
CPiV positive passage material from the last two puppies (terminal passage) was administered intranasally to five puppies (Group 2). In addition, the un-passaged CPiV was directly vaccinated subcutaneously to another five puppies (Group 1).

No local or systemic adverse reactions were observed in any of the study animals during the whole study. The rectal temperatures recorded during the safety study remained below 39 °C in all animals. The presence of CPiV was verified in all passages. CPiV was isolated from Day 6 to Day 9 after inoculation. In the subsequent safety evaluation, no neutralizing antibodies to CPiV were detected before inoculation. Twenty-one days after inoculation, the titres in all animals were increased.
This study shows no indication of increased virulence of the vaccine CPIV and is in compliance with Ph. Eur. monograph 1955.

**Biological properties of the vaccine strain**

The intrinsic biological properties of the live virus vaccine strains have been adequately characterised by the information provided and the safety and efficacy data supplied. A summary regarding the characterisation of the live virus strains and the stability of the attenuation was provided, which emphasizes the stability of the attenuation of each strain. No increase of virulence was observed.

The CVMP therefore concluded that adequate information was provided on the biological properties of the vaccine strain.

**Recombination or genomic re-assortment of the strains**

The genetic stability of the each of the vaccine strains of Versican Plus DHPPi/L4 has been demonstrated. Each of the vaccine strains has been demonstrated to be consistent and stable by *in vitro* passaging. Further the strains are attenuated and of low virulence.

The potential for recombination or genomic re-assortment for each of the vaccine viruses is as follows:

CDV: Canine distemper virus is a paramyxovirus. Being a single stranded RNA virus, it is therefore not readily amenable to either recombination or re-assortment.

CAV-2: Canine adenovirus is a double stranded DNA virus and is certainly potentially capable of recombination with field strains. In practice, however, this is unlikely to occur in the host.

CPV: Canine parvovirus is a single-stranded DNA virus that replicates in the nucleus of the host cell. Both recombination and re-assortment are therefore theoretically possible.

CPiV: Canine parainfluenza virus is a paramyxovirus. Being a single stranded RNA virus, it is therefore not readily amenable to either recombination or re-assortment.

**Study of residues**

Versican Plus DHPPi/L4 is a vaccine which is indicated solely for use in dogs. Therefore, the consideration of residues is not applicable.

**Interactions**

No studies have been performed to test the effect of the vaccine on concurrent use of any other vaccine. Therefore section 4.8 of the SPC includes the common statement to this effect.

**Field studies**

One multinational study was performed to evaluate safety and efficacy of the vaccine in comparison to a positive control group. Three cohorts were analysed after vaccination. No abnormal general physical conditions were recorded. Regarding rectal temperatures no abnormalities could be identified. Examinations of the injection site show swellings with a maximum diameter of 35 mm. The maximum duration of swellings following treatment lasted 18 days.

After vaccination signs of lethargy (reduced liveliness), vomiting, diarrhoea and anorexia were observed in some dogs. These observations were reflected in the SPC.
A further field study was performed with a batch without the rabies component. Examinations of the injection site show swellings with a maximum diameter of 38 mm. The maximum duration of swelling following treatment lasted 20 days. No abnormal general physical conditions were recorded. Regarding rectal temperatures no abnormalities could be identified. These findings further supported the acceptable safety profile for this particular vaccine as well as the whole vaccine range.

**User safety**

The applicant provided a user risk assessment compliant with the CVMP Guideline on user safety for immunological veterinary medicinal products (EMEA/CVMP/IWP/54533/2006).

The following possible risks were discussed: Self-administration, skin contamination, breaking of a glass vial and toxic or infectious ingredients. Versican Plus DHPPi/L4 can be considered as presenting no particular risk to humans as many of its components are not harmful for humans and the other components have been shown to be not infectious for humans. Additionally the product must be administered by competent end users, i.e. a skilled veterinarian or a trained person under the supervision of a veterinarian.

In the SPC appropriate warnings are included concerning handling of the vaccine or in case of accidental self-injection.

The CVMP therefore concluded that the user safety for this product is acceptable when used as recommended in the SPC.

**Environmental risk assessment (ERA)**

An environmental risk assessment in compliance with the CVMP Note for guidance on environmental risk assessment of immunological veterinary medicinal products (EMEA/CVMP/074/95) was provided.

Phase I assessment

1. Hazard identification
   
   - The vaccine is composed of 8 well characterised antigens and a well-known adjuvant.
   - The stability of the attenuation of each strain was demonstrated.
   - Spread of virus cannot be excluded, but since an appropriate warning is included in the SPC the risk of virus spreading to other susceptible species can be regarded as negligible.

2. Exposure to hazard

The product is manufactured in tightly closed vials. A small volume is parentally and individually administered to dogs (subcutaneously) by a qualified person. The potential exposure to a hazard is therefore considered adequately controlled and negligible.

Based on the data provided the ERA can stop at Phase I. The product is not expected to pose a risk for the environment when used according to the SPC.

**Overall conclusions on the safety documentation**

**Laboratory studies**
The administration of one dose of Versican Plus DHPPi/L4(R) containing maximum potency of antigens by the recommended route was found to be safe for puppies of 6 weeks of age.

An overdose study with the live virus components (lyophilisate) showed an acceptable safety profile for these components. One study of a repeated dose has been performed. Overall, it can be concluded that the administration of repeated doses of Versican Plus DHPPi/L4(R) containing maximum potency of antigens by the recommended route was found to be safe for puppies of 6 weeks of age.

No studies have been performed on reproductive safety and a warning sentence that the use is not recommended during pregnancy and lactation is included in the SPC.

No studies have been performed to test the effect of the vaccine on the immune system. Except for the CPV component, none of the components of Versican Plus DHPPi/L4 is known to have any adverse effects on the immune functions of infected animals. CDV, CPiV and CAV-2 are already known as components of Versican DHPPi/L3R. No adverse effects are known for these antigens.

No studies have been performed on the concurrent use of any other vaccine not being part of the Versican Plus range. This fact is addressed in section 4.8 of the SPC.

**Spread of the vaccine strain**

**CPV-2b strain:** The virus was found to be shed by faeces in all animals, vaccinated and controls. The virus disseminated to thymus, spleen and mesenteric lymph nodes of the vaccinated animals. No evaluation of the spread to other possibly susceptible non-target species was provided.

**CDV strain:** There was an increase in the neutralizing antibodies to CDV in the vaccinated animals. There were no detectable neutralizing antibodies to CDV in the control animals. CDV was detected in the lung lymph nodes of one vaccinated animal by PCR. The control animals were negative.

The study results show that the vaccine CDV strain is not capable to spread from vaccinated to unvaccinated puppies.

**CAV-2 strain:** There was an increase in the neutralizing antibodies to CAV-2 in the vaccinated animals. There were no detectable neutralizing antibodies to CAV-2 in the control animals at any of the testing time points. No secretion samples were taken to demonstrate the presence or absence of virus.

**CPiV-2 strain:** There was an increase in neutralizing antibodies to CPiV-2 in the vaccinated animals. There were no detectable neutralizing antibodies to CPiV-2 in the control animals at any of the testing time points. No organ samples were taken.

**Dissemination in the vaccinated animal**

**CPV-2b strain:** Dissemination of the vaccine strain was investigated together in the shedding and spreading study and in the reversion to virulence study. Both studies demonstrate that all vaccinated and control animals excreted virus in faeces. No further secretion samples, e.g. oral or nasal secretion were taken and tested.

**CDV strain:** Dissemination of the CDV vaccine strain was investigated together in the shedding and spreading study. CDV was found to disseminate to blood and mediastinal lymph nodes in a minority of vaccinated animals. No dissemination to the brain, nasal mucosa, tonsils, thymus, spleen or lungs was detected. No CDV was detected in nasal swabs, samples of nasal secretions, urine, faeces and saliva.

**CAV-2 strain:** Tissue samples of vaccinated puppies were collected and prepared for passaging on cell culture. In this study CAV-2 was isolated from the mediastinal, cervical, mesenteric and axillary lymph nodes. No virus was found nasal mucous membrane, larynx mucous membrane, tonsils, lungs, spleen, liver and kidney. No secretion samples were taken to demonstrate the presence or absence of virus.
CPIV-2 strain: No investigation on the dissemination of the CPIV vaccine strain has been undertaken. Reference is made to the Spread study and the Irreversibility of Attenuation of Vaccine study. Data show that this virus disseminates to the nasal mucosa and is excreted after vaccination. Furthermore, neutralizing antibodies to CPIV-2 in the vaccinated animals could be observed, but not in the control animals.

Reversion to virulence of attenuated vaccines

CPV-2b strain: In a first passage procedure the virus was passaged several times in dogs. In a second evaluation step, animals were inoculated with this passaged virus oronasally. No excretion of virus in faeces was detected in animals inoculated with the last passage of virus, neither by HA test nor PCR. No seroconversion could be shown in this group. The study results implicate that the master seed of canine parvovirus, when administered by the recommended route, does not increase in virulence when passaged in puppies.

CDV strain: As vaccine virus could not be recovered from the 1st passage animals (two animals) the first set of passages was terminated and second set of passages started with two animals. This is acceptable because further studies are available which demonstrate a negligible risk of reversion to virulence.

CAV-2 strain: In a first passage procedure the virus was passaged several times in dogs. In a second evaluation step animals, which were inoculated with this passaged virus intranasally, show an increase of antibody titres. No canine adenovirus could be isolated from the tested tissues from these animals. This study shows no indication of increased virulence of the vaccine CAV-2.

Field studies

One multi-centre study was performed to evaluate safety and efficacy of the vaccine in comparison to a positive control group. Three cohorts were analysed after vaccination. No abnormal general physical conditions were recorded. Regarding rectal temperatures no abnormalities could be identified. Examinations of the injection site show swellings with maximum diameter of 35 mm. The maximum duration of swelling following treatment lasted 18 days.

After vaccination signs of lethargy (reduced liveliness), vomiting, diarrhoea and anorexia were observed in some dogs and these observations are reflected in the SPC.

User safety

In the SPC appropriate warnings are included concerning handling of the vaccine or in case of accidental self-injection. Furthermore it is stated that the vaccine contains no ingredients that are toxic or infectious to humans. The user safety for this product is acceptable when used as recommended in the SPC.

Environmental risk assessment

Based on the data provided the ERA can stop at Phase I. The product is not expected to pose a risk for the environment when used according to the SPC.
Part 4 – Efficacy

Introduction and general requirements

Versican Plus DHPPi/L4 is a multivalent live and inactivated virus and bacterial vaccine which is indicated for the immunisation of healthy puppies from six weeks of age and dogs against canine distemper, adenovirus hepatitis, adenovirus respiratory disease, parvovirus, parainfluenza and leptospirosis. The live virus components of the vaccine (canine distemper virus (CDV), canine adenovirus type 2 (CAV-2), canine parvovirus type 2b (CPV-2b), canine parainfluenza virus (CPiV)) are presented in freeze-dried form in a vial to be reconstituted with a vial of the inactivated components (Leptospira Bratislava, Leptospira Canicola, Leptospira Grippotyphosa and Leptospira Icterohaemorrhagiae) presented in liquid form. The liquid fraction also contains an adjuvant (aluminium hydroxide).

Laboratory vaccination/challenge studies (establishment of the minimum protective dose, onset of immunity, duration of immunity) and a field study were provided to support the efficacy claims.

Laboratory trials

The challenge trials were performed with batches of minimum protective doses.

Establishment of a challenge model

Efficacy of all components of Versican Plus DHPPi/L4 was assessed by challenges with heterologous challenge strains according to component-specific monographs. Certificates for the challenge strains were provided.

Determination of the vaccine dose

Minimum immunisation doses for viral components of Versican Plus DHPPi/L4

The minimum immunisation doses (MIDs) for the Versican Plus DHPPi/L4 viral components CDV, CAV-2, CPV and CPiV were selected based on previous experiences with these strains. The CAV-2 and CPiV components are also approved as components of Versican DHPPi/L3R. The CDV component of Versican Plus DHPPi/L4 is a new master seed of the CDV component approved in Versican DHPPi/L3R. The CPV-2b component of Versican Plus DHPPi/L4 is a new master seed not included in any vaccine approved in the EU. The MID for this component was selected based on previous experiences with polyvalent vaccines for dogs.

Minimum immunisation doses for Leptospira components of Versican Plus DHPPi/L4

Three possible MIDs (10⁷, 5x10⁷ and 10⁸ organisms/ml pre-inactivation) were selected for the Leptospira components L. Bratislava, L. Canicola, L. Grippotyphosa and L. Icterohaemorrhagiae based on previous data and experiences with the approved canine vaccine Versican DHPPi/L3R.

The results and conclusions of these studies confirmed that each Leptospira serovar in combination with the other vaccine components at an MID of 5x10⁷ organisms/ml protected minimum age dogs against clinical signs, systemic infection, and renal infection and excretion of Leptospira. No significant differences were found between formulations with or without adjuvant or rabies virus.

MIDs for all components were confirmed in onset of immunity studies.
Onset of immunity

Onset of immunity (OOI) has been demonstrated with challenge studies according to the relevant Ph. Eur. monographs.

CDV:

Seven 6-week old dogs (5 with 2 control dogs), tested seronegative against CDV were administered the vaccine Versican Plus DHPPi/L4R subcutaneously. They were challenged intravenously with the canine distemper challenge virus CHSV Snyder Hill at Day 21 after vaccination. After challenge the vaccinated group showed no clinical signs of distemper. There was no increase in the rectal temperature and a further increase in antibody titres while the control group showed clinical signs of distemper and was euthanized for humane reasons.

While six dogs are not a large number they are considered adequate and representative in this specific situation of the tight correlation between antibody titre and protection for this particular antigen. Based on the data provided titres of 1:2 or 1:4 were considered indicative of protection against CDV. It should be noted that this conclusion cannot be generalised because no standardised tests exist for determination of antibodies against CDV (like it is established for rabies virus) making it almost impossible to define “protective antibody titres against CDV” because the antigen titres will vary dependent on the individual laboratory.

CAV-1:

Seven 6-week old dogs (5 with 2 control dogs), tested seronegative against CAV-1 and CAV-2 were administered the vaccine Versican Plus DHPPi/L4R subcutaneously. They were challenged intravenously with the challenge strain CAV-1 Mirandola at Day 21 after vaccination. After challenge the vaccinated group showed no clinical signs of canine hepatitis but one dog developed a slight increase of rectal temperature (39.5 °C) above the upper physiological limit of 39.5 °C at Day 6 after challenge. There was a further increase in antibody titres while the control group showed clinical signs of canine hepatitis and was euthanized for humane reasons.

This study is considered valid because it fulfils the requirements of Ph. Eur. monograph 1951. The claims stated for this antigen are supported by this study.

CAV-2:

Twenty 6-week old dogs (10 with 10 control dogs), tested seronegative against CAV-2 were administered the vaccine Versican Plus DHPPi/L4R subcutaneously. They were challenged intranasally with the challenge strain CAV-2 Manhattan at Day 21 after vaccination. After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. There was a further increase in antibody titres against CAV-2 and on virus isolation 5/10 dogs excreted CAV-2 for 1–3 days from Day 3 to Day 6 after challenge. The duration of virus excretion was significantly lower than in the controls.

This study complies with the requirements of Ph. Eur. monograph 1951 and is considered acceptable. The claims stated for this antigen are supported by this study.

CPV-2b:

Seven 6-week old dogs (5 with 2 control dogs), tested seronegative against CPV (haemagglutination inhibition (HAI) and virus neutralisation (VN)), were administered the vaccine Versican Plus DHPPi/L4R subcutaneously. They were challenged oronasally with the challenge strain CPV-2b strain 212/98 at Day 21 after vaccination. After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. There was a further increase in antibody titres against CPV-2b while the WBC count demonstrated no leukopenia.
On virus isolation and HA 3/5 dogs excreted virus on one day between 3 and 5 days after challenge. This was less than 1/100 of the geometric mean of the maximum titres found in control animals by HA. No isolation of infectious CPV-2b was found in cell culture.

This study complies with the requirements of Ph. Eur. monograph 0964 and is considered acceptable. The claims stated for this antigen are supported by this study.

**CPIV:**

Fifteen 6-week old dogs (10 with 5 control dogs), tested seronegative against CPIV were administered the vaccine Versican Plus DHPPi/L4R subcutaneously. They were challenged intranasally with the challenge strain CPIV D008 at Day 21 after vaccination. After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. There was a further increase of antibody titres against CPIV.

On virus isolation no CPIV-excretion was found prior to challenge whereas 8/10 vaccinated dogs excreted CPIV for 1–4 days starting from Day 2 until Day 6 after challenge. The duration of virus excretion was found to be significantly lower than in the controls.

As regards immunogenicity Ph. Eur. monograph 1955 states: “The vaccine complies with the test if the scores for coughing or virus excretion for the vaccinated dogs are significantly lower than in the controls.” In this study no coughing could be observed in the control group.

This clinical sign is not induced in the challenge model used by the applicant. Signs following experimental infection with CPIV are known to be very mild and clinical signs observed in the field are mostly due to concurrent (secondary bacterial) infection. A more severe challenge model could only be achieved in dogs from the Bioveta SPF colony which is free of respiratory pathogens by introducing a second pathogen (e.g. *Bordetella bronchiseptica*), but this may interfere with the assessment of the efficacy against CPIV. As the monograph states that scores for either coughing or virus excretion need to be significantly lower than in controls the CVMP concluded that this wording implies that only one of the two options (i.e. coughing or virus excretion) needed to be fulfilled for the vaccine to comply with the test.

However, taking into consideration that the challenge was very mild, it cannot be excluded that in case of a severe challenge even the vaccinated animals would have developed mild clinical signs. Therefore more detailed information as regards the claim for CPIV, i.e. “to prevent clinical signs (nasal and ocular discharge)” has been added to the SPC.

**Leptospira**

The study design for each Leptospira component was as follows. Twelve 6-week-old dogs (6 with 6 control dogs), tested seronegative against the principle serovars of Leptospira, were subcutaneously administered the vaccine Versican Plus DHPPi/L4R. They were challenged by conjunctival and by intraperitoneal route. After challenge they were observed for clinical signs, measurement of rectal temperature, blood samples for serology, for haematology, biochemistry and isolation of the challenge organism, urine samples for isolation of the challenge organism. Twenty-eight days after challenge they were euthanized and the liver and kidneys examined macroscopically and microscopically and tested for the presence of the challenge organism.

**Results:**

*L. Bratislava*

All six vaccinated animals seroconverted. After challenge, only control animals showed typical clinical signs of canine leptospirosis such as apathy (5 out of 6 animals), mild to moderate anorexia (6/6), fever (3/6), dehydration (4/6), conjunctivitis (2/6), diarrhoea (1/6) and jaundice (2/6) starting four days after challenge and persisting in one animal until 28 days.
No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

For the post-challenge, the difference between vaccinated and control animals was significant regarding the total clinical score, the number of days that the challenge organism was detected in blood and the number of liver and kidney samples in which the organism was detected.

*L. Canicola*

All six vaccinated animals seroconverted. After challenge, only control animals showed typical clinical signs of canine leptospirosis such as apathy (5 out of 6 animals), mild to moderate anorexia (6/6), fever (4/6), dehydration (5/6), conjunctivitis (4/6), diarrhoea (2/6) and jaundice (2/6) starting two days after challenge and persisting until Day 25.

No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

For the post-challenge, the difference between vaccinated and control animals was significant regarding the total clinical score, the number of days that the challenge organism was detected in blood and the number of liver and kidney samples in which the organism was detected.

*L. Grippotyphosa*

All six vaccinated animals seroconverted. After challenge, only control animals showed typical clinical signs of canine leptospirosis such as apathy (3 out of 6 animals), mild to moderate anorexia (5/6), fever (4/6), dehydration (3/6), conjunctivitis (2/6) and jaundice (4/6) starting three days after challenge and persisting until Day 18.

No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

For the post-challenge, the difference between vaccinated and control animals was significant regarding the total clinical score, the number of days that the challenge organisms were detected in blood and the number of liver and kidney samples in which the organism was detected.

*L. Icterohaemorrhagiae*

All six vaccinated animals seroconverted. After challenge, only control animals showed typical clinical signs of canine leptospirosis such as apathy (6 out of 6 animals), mild to moderate anorexia (6/6), fever (4/6), dehydration (5/6), conjunctivitis (4/6) and jaundice (4/6) starting four days after challenge and persisting until Day 28.
Increases in creatinine (1 out of 6 control animals), AST (3/6), ALT (3/6) and ALP (2/6) and increase of ALT in two vaccinates were detected but the difference was not significant.

Increases in WBC above the upper limit of the physiological range were detected in four control animals but in none of the vaccinated dogs.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

For the post-challenge, the difference between vaccinated and control animals was significant regarding the total clinical score, the mean total haematological score, the number of days that the challenge organism was detected in blood and the number of liver and kidney samples in which the organism was detected.

Conclusion:

Twenty-five days after vaccination of naïve dogs from six weeks of age according to the vaccination scheme with batches of Versican Plus DHPPi/L4R of minimum potency, the animals were protected against challenge with heterologous *Leptospira* strains that resulted in typical signs of leptospirosis in control animals.

**Influence of maternal antibodies on the efficacy of the vaccine**

The presented studies clearly show the influence of maternally derived antibodies (MDA) regarding CDV, CPV and CAV antigens. While MDA negative dogs seroconvert after the first vaccination and are boostered after the second vaccination, MDA positive dogs show a titre decrease until Days 28–35. Seroconversion is observed from Days 35–42 onwards. Although the animals were protected against challenges with virulent CDV, CPV and CAV-1 strains, the possible interference of MDA should always be taken into consideration when vaccinating very young puppies. The studies demonstrate the importance of the second vaccination as part of the primary vaccination: animals with MDA do not seroconvert after the first vaccination but after the second one. As immunological responses to CDV, CPV and CAV-2 may be delayed due to MDA the vaccination scheme for young dogs should be planned carefully. This has been reflected in the SPC.

The data show that the interference of MDA against CPIV does not play a considerable role. Therefore, a corresponding warning in the SPC is unnecessary.

**Duration of immunity (DOI)**

Ten laboratory challenge studies in dogs were performed to demonstrate 1-year duration of immunity for the CDV, CAV-2, CPV, CPIV, *Leptospira* and rabies components of Versican Plus DHPPi/L4R.

**CDV:**

Nine 6-week-old dogs (6 with 3 control dogs), tested seronegative against CDV were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21. They were challenged intravenously with the canine distemper challenge virus CHSV Snyder Hill at 12 months after vaccination.

After challenge the vaccinated group showed no clinical signs of distemper, no increase in rectal temperature above the upper limit (39.5 °C) and a satisfactory anamnestic antibody response while the control group showed clinical signs of distemper and was euthanized for humane reasons.
CAV-1:
Nine 6-week old dogs (6 with 3 control dogs), tested seronegative against CAV-1 and CAV-2 were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21. They were challenged intravenously with the challenge strain CAV-1 Mirandola.

After challenge the vaccinated group showed no clinical signs of canine hepatitis and a satisfactory anamnestic antibody response.

This study fulfils the requirements of Ph. Eur. monograph 1951 and is considered acceptable. Duration of immunity for at least one year was demonstrated for CAV-1.

CAV-2:
Twenty 6-week old dogs (10 with 10 control dogs), tested seronegative against CAV-1 and CAV-2 were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21. They were challenged intranasally with the challenge strain CAV-2 Manhattan at 12 months after vaccination.

As controls were co-housed with vaccinated animals and therefore served as sentinels, it can be concluded that no concurrent infections with CAV-2 boosted the immunity in vaccinated animals before challenge.

After challenge the vaccinated group showed no clinical signs of canine hepatitis and a satisfactory anamnestic antibody response. The mean total score was significantly lower in vaccinated animals than in the controls.

Virus isolation: 3/10 vaccinated animals excreted CAV-2 for one day either 2 or 3 days after challenge. Duration of virus excretion: the number of days was significantly lower in vaccinated animals than in the controls.

In the control group 4/10 dogs showed mild nasal discharge and mild ocular discharge starting 2 days after challenge and persisting for up to 4 days until 6 days after challenge. The mean total score of abnormal clinical observations was significantly higher in control than in vaccinated animals. Virus isolation: 10/10 controls excreted CAV-2 for 2–4 days starting from 2 until 5 days after challenge. Duration of virus excretion: the number of days was significantly higher in the controls than in the vaccinates. The area under the curve of CAV-2 titres was significantly higher in control than in vaccinated animals.

This study fulfils the requirements of Ph. Eur. monograph 1951 and is considered acceptable. Duration of immunity for at least one year was demonstrated for CAV-2.

CPV-2b:
Seven 6-week old dogs (5 with 2 control dogs), tested seronegative against CPV (HAI and VN) were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21. They were challenged oronasally with the challenge strain CPV-2b strain 212/98 at 12 months after vaccination.

Control animals remained seronegative until challenge. As they were housed together with vaccinated animals, these results confirm that vaccine induced immunity to CPV was not further boosted through concurrent infections with field strains during the study.

After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. Virus isolation: by HA, 3/5 animals showed CPV excretion for 2 days between 5 and 10 days after challenge. This was less than 1/100 of the geometric mean of the maximum titres found in control animals by HA. By cell culture, 0/5 vaccinated animals excreted infectious CPV.

The control group showed no abnormal clinical signs, there was hyperthermia in 1/2 controls 10 and 11 days after challenge. WBC count: leukopenia - white blood cell numbers decreased more than 50%
compared to pre-challenge mean values (leukopenia) in 2/2 controls. Virus isolation: by HA, 2/2 controls showed CPV excretion starting 5 days and peaking 7 to 10 days after challenge. By cell culture, 2/2 controls started to excrete CPV 5 days and peaked 7 days post-challenge.

This study fulfils the requirements of Ph. Eur. monograph 0964 and is considered acceptable. Duration of immunity for at least one year was demonstrated for CPV-2b.

**CPiV:**

Sixteen 6–7 week old dogs (10 with 6 control dogs), tested seronegative against CPiV were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21. They were challenged intranasally with the challenge strain CPiV D008 at 12 months after vaccination.

As controls were co-housed with vaccinated animals and therefore served as sentinels, it can be concluded that no concurrent infections with CPiV boosted the immunity in vaccinated animals before challenge.

After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. Virus excretion: 70% started excreting CPiV from Day 2 until Day 4 after challenge (duration 1–5 days). The number of days of viral shedding was significantly higher in control than in vaccinated animals.

In the control group 67% showed respiratory signs (mild or moderate nasal discharge, mild or moderate ocular discharge) 3–7 days after challenge. Additionally, sneezing was seen for two days in one of the four control animals with discharge. No coughing was observed. There was no increase in rectal temperature and no virus excretion before challenge. 100% excreted virus starting from 2 to 3 days after challenge. There was a further increase of antibody titres against CPiV. On virus isolation no CPiV-excretion was found prior to challenge whereas 8/10 vaccinated dogs excreted CPiV for 1–4 days starting from 2 days until 6 days after challenge. The duration of virus excretion was found to be significantly lower than in the controls.

As regards immunogenicity Ph. Eur. monograph 1955 states: “The vaccine complies with the test if the scores for coughing or virus excretion for the vaccinated dogs are significantly lower than in the controls.” However, in this study no coughing could be observed in the control group.

This clinical sign is not induced in the challenge model used by the applicant. Signs following experimental infection with CPiV are known to be very mild and clinical signs observed in the field are mostly due to concurrent (secondary bacterial) infection. A more severe challenge model could only be achieved in dogs from the Bioveta SPF colony which is free of respiratory pathogens by introducing a second pathogen (e.g. *Bordetella bronchiseptica*), but this may interfere with the assessment of the efficacy against CPiV. As the monograph states that scores for *either* coughing or virus excretion need to be significantly lower than in controls the CVMP concluded that this wording implies that only one of the two options (i.e. coughing or virus excretion) needed to be fulfilled for the vaccine to comply with the test.

However, taking into consideration that the challenge was very mild, it cannot be excluded that in case of a severe challenge even the vaccinated animals would have developed mild clinical signs. Therefore, more detailed information as regards the claim for CPiV, i.e. “to prevent clinical signs (nasal and ocular discharge)” has been added to the SPC.

Twelve months after the second vaccination the antibody titres have declined drastically (one dog 8, one dog 16, one dog 32, one dog 2 and 6 dogs < 2). However, the animals were protected against a challenge with a virulent CPiV challenge. Based on existing knowledge it can be assumed that in this case cellular immunity plays an important role in protection.
Leptospira

The study design for each Leptospira component was as follows. Twelve 6-week-old dogs (6 with 6 control dogs), tested seronegative against the principle serovars of Leptospira, were subcutaneously administered the vaccine Versican Plus DHPPi/L4R at Day 0 and 21. They were challenged by conjunctival and by intraperitoneal route at 12 months after vaccination. After challenge they were observed for clinical signs, measurement of rectal temperature, blood samples for serology, for haematology, biochemistry and isolation of the challenge organism, urine samples for isolation of the challenge organism. Twenty-eight days after challenge they were euthanized and the liver and kidneys examined macroscopically and microscopically and tested for the presence of the challenge organism.

Results:

L. Bratislava

After challenge, 5 out of 6 control animals showed typical clinical signs of canine leptospirosis such as apathy (1 out of 6 animals), mild to moderate anorexia (2/6), dehydration (2/6) and conjunctivitis (4/6). Vaccinated animals did not show any abnormal clinical signs.

No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge Leptospira were re-isolated from the blood of all control animals (100%) and from urine, kidney and liver of 5 out of 6 control animals (83.3%). No Leptospira were detected in the blood, urine, kidney or liver of vaccinated animals.

Macroscopic and microscopic examination of liver and kidney samples showed more gross pathological and more severe histopathological changes in organs from control than from vaccinated animals.

For the post-challenge, the difference between vaccinated and control animals was significant regarding the total clinical score, the number of days that the challenge organism was detected in blood and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented clinical signs, infection and excretion caused by L. Bratislava following two administrations, in seronegative animals from six weeks of age for at least one year after completion of the basic immunisation.

L. Canicola

After challenge, 3 out of 6 control animals showed typical clinical signs of canine leptospirosis such as apathy (1 out of 6 animals), mild to moderate anorexia (2/6), fever (2/6) and dehydration (2/6).

Vaccinated animals did not show any abnormal clinical signs.

No noteworthy changes in biochemical parameters were observed. An increase in WBC count was detected in 1 vaccinated animal 5 days post-challenge. A decrease in thrombocyte numbers was seen in 4 out of 6 control and 1 vaccinated animal.

Post-challenge Leptospira were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No Leptospira were detected in the blood, urine, kidney or liver of vaccinated animals except for 1 vaccinated animal that had positive blood samples 3 and 5 days after challenge.

Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

For the post-challenge the difference between vaccinated and control animals regarding the total clinical score was not significant. For the number of days that the challenge organisms were detected in blood
and the number of liver and kidney samples in which the organism was detected the difference between vaccinated and control animals was significant.

It could be shown that Versican Plus DHPPi/L4(R) prevented excretion and clinical signs and reduced infection caused by *L. Canicola* following two administrations, in seronegative animals from six weeks of age for at least one year after completion of the basic immunisation.

**L. Grippotyphosa**

After challenge, 4 out of 6 control animals showed typical clinical signs of canine leptospirosis such as mild anorexia (1/6), fever (3/6), dehydration (1/6), conjunctivitis (2/6), diarrhoea (1/6) and jaundice (1/6). Vaccinated animals did not show any abnormal clinical signs including fever after challenge except for 1 animal with jaundice. However, *Leptospira* could not be isolated from blood, urine or organs of this animal.

There were increases in the biochemical variable AST, ALT and creatinine detected in control and vaccinated animals pre- and post-challenge; but none of the increases observed pre- and post-challenge was considered to be clinically relevant.

Decreases in thrombocyte numbers were seen in 4 control animals; the decrease was noticeable in only 1 control animal reaching >50%. The animal had developed jaundice after challenge.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). In vaccinated animals, *Leptospira* were detected in the blood of 2 animals for 1 and 2 days post-challenge, respectively. The vaccinated animal with 2 positive blood samples post-challenge also tested positive for *Leptospira* in urine for one day 14 days post-challenge, and in kidney and liver.

Macroscopic and microscopic examination of liver and kidney samples showed more gross pathological and more severe histopathological changes in organs from control than from vaccinated animals.

The difference between vaccinated and control animals was significant regarding the number of days that the challenge organisms were detected in blood and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented clinical signs and reduced excretion and infection caused by *L. Grippotyphosa* following two administrations, in seronegative animals from six weeks of age for at least one year after completion of the basic immunisation.

**L. Icterohaemorrhagiae**

After challenge, 3 out of 6 control animals showed typical clinical signs of canine leptospirosis such as apathy (2 out of 6 animals), mild to moderate anorexia (2/6) and fever (3/6).

Vaccinated animals neither showed abnormal clinical signs nor hyperthermia after challenge.

A more than 100% increase from the pre-challenge baseline value in ALP was noted in one control animal with clinical signs 3 days post-challenge. An increase in AST of > 25–50% was detected in the control animal with hyperthermia 2 days post-challenge. A decrease in thrombocyte numbers was seen in 4 out of 6 controls, including the 3 animals which showed clinical signs and hyperthermia post-challenge, and 1 vaccinated animal.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals except for two vaccinated animal that had positive blood samples 3 days after challenge.
Macroscopic and microscopic examination of liver and kidney samples showed more pathological changes of greater severity in organs from control than from vaccinated animals.

The difference between vaccinated and control animals was significant regarding the number of days that the challenge organisms were detected in blood and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented excretion and clinical signs and reduced infection caused by L. Icterohaemorrhagiae following two administrations, in seronegative animals from six weeks of age for at least one year after completion of the basic immunisation.

**Immunity after revaccination – response to booster (RTB)**

To demonstrate protective immunity of the components of Versican Plus DHPPi/L4R following re-vaccination (annual booster) with a single dose 12 months after completion of the primary vaccination course laboratory response-to-booster (RTB) studies in dogs were performed. For the CPIV and *Leptospira* components of Versican Plus DHPPi/L4R, protective immunity following an annual booster was demonstrated by challenge. For the CDV, CAV-2, CPV components, protective immunity following an annual booster was demonstrated serologically by comparing antibody titres in response to the annual booster with those after primary vaccination.

**CPIV:** Eighteen 6 weeks old dogs (12 with 6 control dogs) tested seronegative against CPIV were administered the vaccine Versican Plus DHPPi/L4R subcutaneously at Day 0 and 21 and one year after the 2nd vaccination. They were challenged intranasally with the CPIV D008 at Day 21 after the one year booster.

After challenge they were observed for clinical signs, measurement of rectal temperature, serology at Day 14 after challenge, nasal swabs from 2 to 10 days after challenge for virus isolation of the challenge organism.

As controls were co-housed with vaccinated animals and therefore served as sentinels, it can be concluded that no concurrent infections with CPIV boosted immunity in vaccinated animals before challenge.

After challenge the vaccinated group showed no clinical signs and no increase of rectal temperature. There was a further increase of antibody titres against CPIV.

The number of days of viral shedding was significantly lower in the vaccinates (42% excreted CPIV on a single day 3 days after challenge) than in the controls (83% excreted virus starting from 2 to 5 days after challenge).

As regards the CPIV component the applicant proposes the following claim: “prevent clinical signs and reduce viral excretion caused by canine parainfluenza virus”.

The challenge data clearly demonstrate that there were no clinical signs in the vaccinated group. However, taking into consideration that the challenge was very mild (no coughing, only mild nasal and ocular discharge in the control group) it cannot be excluded that in case of a severe challenge even the vaccinated animals would have developed mild clinical signs. Therefore, the claim for CPIV has been amended. More detailed information has been included: “to prevent clinical signs (nasal and ocular discharge)”.

As regards the conclusion that the minimum protective titre for CPIV is 1:32, it should be taken into consideration that it was only one animal which was protected by this titre. All other animals showed higher titres. From a statistical point of view this conclusion cannot be supported. Basing the minimum
protective titre for CPiV on the titre of one animal alone cannot be accepted. As protection also depends on other factors (e.g. general health status, individual immune system, cellular immunity) it is basically very difficult to determine fixed protective titres for the antigens.

Twelve months after the second vaccination the antibody titres have declined drastically (one dog 32, three dogs 8, three dogs 4, three dogs 2 and two dogs < 2). However, after a single booster vaccination the titres increased again until challenge (the titres were comparable to those observed three weeks after primary vaccination course) and the animals were protected against a challenge with a virulent CPiV.

As part of this study, the immunogenicity of the other viral vaccine components was assessed by serology following a booster vaccination one year after the primary immunisation of minimum age dogs.

The antibody titres against CDV of vaccinated animals 12 months after the second vaccination have declined drastically (two dogs 32, two dogs 8, four dogs 4, two dogs 2 and two dogs < 2). However, after a single booster vaccination the titres increased again and were comparable to those observed three weeks after primary vaccination course.

The antibody titres against CAV-1 and CAV-2 of vaccinated animals 12 months after the second vaccination have declined. After a single booster vaccination the titres increased again. They were comparable (even higher) to those observed three weeks after primary vaccination course. However, it should be noted that in this study the titres were lower compared to those of the DOI studies (presented in the submitted dossier).

The antibody titres against CPV of vaccinated animals 12 months after the second vaccination have declined slightly. After a single booster vaccination the titres increased again and were comparable to those observed three weeks after primary vaccination course.

**Leptospira**

The study design for each *Leptospira* component was as follows. Twelve 6-week-old dogs (6 with 6 control dogs), tested seronegative against the principle serovars of *Leptospira*, were subcutaneously administered the vaccine Versican Plus DHPPi/L4R at Day 0 and 21 and one year (the control group received vaccination only at Day 0 and 21). They were challenged by conjunctival and by intraperitoneal route at 12 months after vaccination. After challenge they were observed for clinical signs, measurement of rectal temperature, blood samples for serology, for haematology, biochemistry and isolation of the challenge organism, urine samples for isolation of the challenge organism. Twenty-eight days after challenge they were euthanized and the liver and kidneys examined macroscopically and microscopically and tested for the presence of the challenge organism.

**Results:**

*L. Bratislava*

After challenge, four out of six control animals showed typical clinical signs of canine leptospirosis, such as anorexia (1 out of 6 animals), dehydration (3/6), conjunctivitis (1/6) and jaundice (1/6). In two control animals and in one vaccinated animal the body temperatures exceeded the upper physiological limit of 39.5 °C slightly. Apart from this, no abnormal clinical signs were found in vaccinated dogs.

No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of five out of six control animals (83.3%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Findings of macroscopic and microscopic examinations of liver and kidney samples were inconclusive.
Concerning the post-challenge, the difference between vaccinated and control animals was significant regarding the number of days that the challenge organism was detected in blood and urine and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented clinical signs, infection and excretion caused by L. Bratislava following booster vaccination one year after primary vaccination to animals from six weeks of age.

**L. Canicola**

After challenge, four out of six control animals showed typical clinical signs of canine leptospirosis such as apathy (2 out of 6 animals), anorexia (4/6), conjunctivitis (3/6), jaundice (1/6) and dehydration (3/6). In two control animals the body temperatures exceeded the upper physiological limit of 39.5 °C.

In the vaccinated animals no abnormal clinical signs and temperature increases were observed.

No noteworthy changes in biochemical parameters were observed. An increase in WBC count was detected in all control animals and in four vaccinated animal after challenge. A decrease in thrombocyte numbers was seen in three out of six control animals and one vaccinated animal.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals.

Findings of macroscopic and microscopic examinations of liver and kidney samples were inconclusive.

Concerning the post-challenge, a significant difference was found between total clinical scores of male vaccinated and male control animals, but not between female animals because the two female control dogs did not show any abnormal clinical signs. The difference between vaccinated and control animals was significant as regards the number of days that the challenge organisms were detected in blood and urine and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented clinical signs, excretion and infection caused by L. Canicola following booster vaccination one year after primary vaccination to animals from six weeks of age.

**L. Grippotyphosa**

After challenge, five out of six control animals showed typical clinical signs of canine leptospirosis such as anorexia (2 out of 6 animals), dehydration (4/6), vomiting (2/6), diarrhoea (1/6) and jaundice (1/6). Body temperatures exceeding the upper physiological limit of 39.5 °C were not observed in any control animal after challenge.

Vaccinated animals did not show any abnormal clinical signs including fever after challenge.

Except for an increase of AST in three out of six control animals and one vaccinate no clinically relevant post-challenge changes in any of the biochemical variables were seen in the other control animals and in the vaccinates.

No increases in WBC counts above the physiological range and no decreases in thrombocyte numbers were detected neither in control animals nor in vaccinated animals.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of five out of six control animals (83%)

No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals after challenge.

Findings of macroscopic and microscopic examinations of liver and kidney samples were inconclusive.
Concerning the post-challenge, a significant difference was found between total clinical scores of vaccinated and control animals. The difference between vaccinated and control animals was significant as regards the number of days that the challenge organisms were detected in blood and urine and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented clinical signs, excretion and infection caused by *L. Grippotyphosa* following booster vaccination one year after primary vaccination to animals from six weeks of age.

*L. Icterohaemorrhagiae*

After challenge, five out of six control animals showed typical clinical signs of canine leptospirosis such as apathy (1 out of 6 animals), anorexia (2/6), dehydration (3/6), conjunctivitis (2/6) and jaundice (3/6). In two control animals the body temperatures exceeded the upper physiological limit of 39.5 °C.

Vaccinated animals neither showed abnormal clinical signs nor hyperthermia after challenge.

No noteworthy changes in biochemical and haematological parameters were observed.

Post-challenge *Leptospira* were re-isolated from the blood, urine, kidney and liver of all control animals (100%). No *Leptospira* were detected in the blood, urine, kidney or liver of vaccinated animals except for two vaccinated animals which had positive blood samples for one day 2 days after challenge.

Findings of macroscopic and microscopic examinations of liver and kidney samples were inconclusive.

Concerning the post-challenge, a significant difference was found between total clinical scores of vaccinated and control animals. The difference between vaccinated and control animals was significant as regards the number of days that the challenge organisms were detected in blood and urine and the number of liver and kidney samples in which the organism was detected.

It could be shown that Versican Plus DHPPi/L4(R) prevented excretion and clinical signs and reduced infection caused by *L. Icterohaemorrhagiae* following booster vaccination one year after primary vaccination to animals from six weeks of age.

**Additional studies**

**Compatibility**

No studies on immunological compatibility of Versican Plus DHPPi/L4 with other products were undertaken. Section 4.8 of the proposed SPC for Versican Plus DHPPi/L4 contains the following text:

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

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.

In view of this text, data on the concurrent administration of Versican Plus DHPPi/L4 with other veterinary medicinal products are not required for this application. The justification and the proposed text of the SPC are acceptable.

**Field trials**

The applicant performed a multi-centre, positively controlled, randomised, blinded field study in two countries, in compliance with CVMP/VICH/595/98. The animals were either vaccinated two times at an
interval of 3 or 4 weeks (V1= Versican Plus DHPPi/L4; V2= Versican Plus DHPPi/L4R) or received one annual booster (Versican Plus DHPPi/L4R). The efficacy was assessed by determining antibody titres prior to and after vaccinations. As no unvaccinated dogs were included for ethical reasons, a competitor vaccine was used for antibody comparison. The antibody responses were categorised as follows:

No increase of antibody titres prior to and after vaccination.

**Increase 1:**
- < 2-fold for CDV, CAV-1, CAV-2, CPV, CPIV (SN)
- < 4-fold (SN) and < 0.5 IU (FAVN) for rabies virus
- < 4-fold for *Leptospira* (MAT)

**Increase 2:**
- ≥ 2-fold for CDV, CAV-1, CAV-2, CPV, CPIV (SN)
- ≥ 4-fold (SN) and > 0.5 IU (FAVN) for rabies virus
- ≥ 4-fold for *Leptospira* (MAT)

An overview of the field trial is presented below. These were carried out in 3 centres in France (FR) and 3 centres in Germany (DE). A total of 128 dogs (FR 63, DE 65) were included in the field trials. Cohort 1: 54 dogs (FR 27, DE 27) with an age range of 8 weeks to 15 years; Cohort 2: 41 dogs (FR 21, DE 20) with an age range of 1 year to 11 years; Cohort 3: 33 dogs (FR 15, DE 18) with an age range of 8 to 9 weeks. The comparator products used were Enduracell 7 and Enduracell 8 (FR) and Vanguard 7 and Vanguard R (DE).

Cohort 1: two vaccinations to puppy and adult dogs 3–4 weeks apart followed by owner observations;

Cohort 2: one annual booster vaccination to adult dogs followed by owner observations;

Cohort 3: two vaccinations 3–4 weeks apart to puppies followed by observations through trained personnel.

There were 45 mixed bred and 84 pure bred dogs of 28 breeds including toy breeds, utility/hunting breeds and large breeds (Boxer, Bulldog, Collie, Dachshund, Dalmatian, Pinscher, Pointers, Poodle, Retrievers, Shepherd Dogs, Setter, Spaniels, Terriers) and their genders were 50 females, 23 neutered females, 41 males and 14 neutered males.

Serology was performed on cohort 1 and 3 before vaccination, and 21 days after the second vaccination, i.e. V1, V2 (+3 days) and V2+21 (+3 days) and on cohort 2 before vaccination and 21 days after vaccination, i.e. V1 and V1+21 (+3 days).

Results:

The serological data were weighted on basis of the age and/or the presence or absence of antibodies and also country- and cohort-wise.

**Puppies (consisting of part of Cohort 1 [n=11; only 7 vaccinated with Versican Plus DHPPi/L4R] and Cohort 3 [n=33])**

Forty-four dogs from 8 weeks of age but younger than 6 months, without a previous history of vaccination.

Forty out of 44 puppies (91%) had MDA against at least one vaccine component at the time of the first vaccination: MDA against rabies virus, *Leptospira* and CPIV were detected only sporadically, MDA against CPV consistently persisted for the longest period of time and were still detected in two puppies at 14 weeks of age. MDA against CAV-1, CAV-2 and CDV were detected in 16–25% and were still present in one puppy at 14 weeks of age.
Puppies without MDA:
- Full response (increase 2) against the live viral components CDV, CAV-2, CPV, CPiV, and the inactivated bacteria L. Canicola and L. Icterohaemorrhagiae.
- 11% of the puppies did not respond to L. Grippotyphosa and L. Bratislava.
- 8% of the puppies did not respond to rabies virus. 19% of puppies did not reach 0.5 IU (increase 1). 27% of puppies did not reach 0.5 IU (increase 2).

Puppies with MDA:
- Full response (increase 2) to primary immunisation (two vaccinations 3-4 weeks apart) against CAV-1 (n=5), CAV-2 (n=8) and rabies virus (only 1 dog).
- 20% of puppies did not show an antibody titre increase against CDV (n=1) and CPV (n=5):
  CDV: The animal had an antibody titre of 4 before the first vaccination, was seronegative (< 2) before the second vaccination and had a titre of 4 after the second vaccination => MDA interfered with the first, but not the second vaccination.
  CPV: In three puppies MDA interfered with responses after both vaccinations leaving the puppies without any protective antibody levels (< 2) after primary immunisation.

 The other two pups had higher than average MDA titres before the first vaccination which decreased to levels that allowed them to respond to the second vaccination.
- Serological results for CPiV, L. Bratislava and L. Grippotyphosa did not allow more general conclusions as there was only one MDA positive animal. MDA titres were moderately high before the first vaccination and remained at this level (CPiV) or increased after the second vaccination (L. Bratislava and Grippotyphosa) indicating that it responded to primary immunisation.

Cohort 1
Fifty-four dogs: 43 dogs over 6 months of age and without a previous history of vaccination or with a previous history of vaccination that had lapsed by more than 14 months, i.e. adult animals due to re-start a primary vaccination course. 11 dogs ≤ 6 months of age without previous vaccinations (Versican Plus DHPPi/L4R: n=7; comparator groups: n=4).

Animals without pre-existing antibodies:
- 63–100% responded fully (increase 2) to two vaccinations.
- One dog did not respond to the CPiV component.
- One dog only showed a very small titre increase against L. Bratislava.
- Three dogs showed antibody titre increase against rabies virus that did not reach 0.5 IU. Two were puppies and are discussed above. One was an adult and reached a titre of 0.39 IU after immunisation which is considered protective.

Animals with pre-existing antibodies:
The proportions of dogs with pre-existing antibodies that showed type 2 increases in serological responses were lower than those of dogs without pre-existing antibodies against all antigens except for CDV, CAV-1 and CAV-2. Consequently the proportion of dogs with pre-existing antibodies that showed lower serological increases (increase 1) or no increases were higher than those of dogs without pre-existing antibodies.
Adult boosters (cohort 2)

Forty-one dogs > 6 months of age, with a previous history of vaccination and requiring an annual booster, i.e. adult animals for re-vaccination:

Animals without pre-existing antibodies:

- 100% responded to the annual booster vaccination with antibody titre increase against CDV, CAV-1, CAV-2, CPV, CPIV, L. Canicola and L. Icterohaemorrhagiae.

- Two adults did not respond to L. Bratislava and L. Grippotyphosa, one dog showed only a slight response to L. Bratislava (commercial vaccines against these serovars were not available before).

Animals with pre-existing antibodies:

Proportion of dogs that showed type 2 antibody titre increase was lower than those of dogs without pre-existing antibodies against all antigens. Consequently the proportion of dogs with pre-existing antibodies that showed lower serological increases (increase 1) or no increases were generally higher than those of dogs without pre-existing antibodies.

Serological responses as measure for protection

Since antibody titres from field and laboratory studies were determined using the same assay systems in the same laboratory, it was possible to directly compare field titres with minimum protective titres established in laboratory studies.

The results of the field study show that vaccination with Versican Plus DHPPi/L4R induced antibody titres that can be considered protective against CDV, CAV-1 and CAV-2 in all dogs.

For CPV 100% of the adult dogs were protected following an annual booster vaccination (Cohort 2) and all adult dogs of Cohort 1 were protected following a primary immunisation. Puppies did not all respond with protective antibody levels against CPV (Cohorts 1 and 3) because of pre-existing MDA.

For CPIV, all except for one adult dog in Cohort 1 responded with titres $\geq 16$ to primary immunisation. In Cohorts 2 and 3, 11% and 27% of dogs which were mostly seronegative before vaccination showed titres of $>2$ but $<16$ after vaccination, respectively.

For Leptospira, the majority of dogs in all cohorts responded to vaccination with Versican Plus DHPPi/L4R. Cohort 2 dogs without pre-existing antibodies required a second vaccination to achieve full protection. In Cohorts 1 and 3, there were dogs that responded to vaccination with increases albeit below 16 for L. Bratislava and 32 for the other three Leptospira. As antibody titres against Leptospira do not greatly correlate with protection, animals that responded with an antibody titre increase to vaccination may still be protected. However, it is questionable whether dogs that did not show any increase following vaccination are protected.

The applicant summarized all serological data irrespective of their antibody status pre-vaccination via descriptive statistics and compared the minimally induced antibody titre per antigen with the titre that was fixed as minimum protective titre in the challenge studies. From this, the applicant concluded the percentage of dogs that were protected in the case of an infection.

Overall summary and comments:

Evaluable serological data from 86 animals were generated.

The data of the animals were analysed following the prerequisites given below:

- Animals with/without antibodies pre-vaccination.
- Animals with/without MDA.
The dogs were vaccinated with Versican Plus DHPPi/L4 (V1) and Versican Plus DHPPi/L4R (V2) at an interval of 3 or 4 weeks, except for the booster Cohort 2 that received one vaccination with Versican Plus DHPPi/L4R.

Subsequently, the applicant compared the serological data from the field with the serological results of challenge studies by descriptive statistics and deduced the percentage of dogs that are expected to be protected in the case of an infection.

From the CVMP point of view, the following critical issues were identified:

a) None of the field trial animals was at minimum age (6 weeks). The youngest were 8 weeks that received the vaccine without the rabies component.

b) The joint evaluation of sub-groups (Cohort 1: dogs > 6 months subdivided into a) non-vaccinated, seronegative and b) vaccinated more than 14 months ago, seronegative) is not acceptable from an immunological point of view. Seronegative animals with a previous vaccination history, so-called primed animals, react immunologically differently compared to naïve seronegative animals.

Furthermore, from a statistical point of view, the CVMP could not agree with the applicant’s conclusions regarding the protective titres for CDV and CAV-1. It was only one animal in each challenge study which was protected by these low titres. As protection also depends on other factors (e.g. general health status, individual immune system, cellular immunity) it is very difficult to determine fixed protective titres for the antigens. In addition, it should be noted that there do not exist standardised tests for determination of antibodies against CDV and CAV-1 (like it is established for rabies virus). Therefore, it is almost impossible to define “protective antibody titres against CDV or CAV-1” because the antigen titres will vary dependent on the individual laboratory.

The presented field study clearly shows the influence of MDA regarding CDV, CPV and CAV antigens. While the MDA negative dogs mostly seroconvert after the first vaccination and are boostered after the second vaccination, MDA positive dogs react after the second vaccination and in general have lower titres than those without MDA. The study demonstrates again the importance of the second vaccination as part of the primary vaccination scheme. As immunological responses to CDV, CPV and CAV-2 may be delayed due to MDA, the vaccination scheme for young dogs should be planned carefully. This has been reflected in the SPC.

It should be noted that 17 out of 33 seronegative dogs did also not seroconvert against CPIV after the first vaccination and 2 dogs even remained seronegative after the second vaccination.

The data show that the interference of MDA against Leptospira does not play a considerable role. Therefore, a corresponding warning in the SPC is unnecessary.

**Overall conclusion on efficacy**

Versican Plus DHPPi/L4 is intended for use in dogs from six weeks of age.

The minimum protective dose is indicated below:

<table>
<thead>
<tr>
<th>Component</th>
<th>Minimum potency/ Antigen content</th>
</tr>
</thead>
<tbody>
<tr>
<td>Freeze-dried fraction (live attenuated)</td>
<td></td>
</tr>
<tr>
<td>Canine distemper virus, strain Bio 11/A</td>
<td>$10^{3.1}$ TCID$_{50}$</td>
</tr>
<tr>
<td>Canine adenovirus type 2, strain CAV-2-Bio 13</td>
<td>$10^{3.6}$ TCID$_{50}$</td>
</tr>
<tr>
<td>Canine parvovirus type 2b, strain CPV-Bio 12/B</td>
<td>$10^{4.3}$ TCID$_{50}$</td>
</tr>
<tr>
<td>Canine parainfluenza type 2 virus, strain CPIV-2-Bio 15</td>
<td>$10^{3.1}$ TCID$_{50}$</td>
</tr>
</tbody>
</table>
**Liquid fraction (inactivated)**

- *Leptospira interrogans* serovar Bratislava, strain MSLB 1088: GMT$^2 \geq 1:51$ ALR$^3$
- *Leptospira interrogans* serovar Canicola, strain MSLB 1090: GMT $\geq 1:51$ ALR
- *Leptospira kirschneri* serovar Grippotyphosa, strain MSLB 1091: GMT $\geq 1:40$ ALR
- *Leptospira interrogans* serovar Icterohaemorrhagiae, strain MSLB 1089: GMT $\geq 1:51$ ALR

1. Tissue culture infectious dose 50%
2. Geometric mean titre
3. Antibody agglutination-lytic reaction

The proposed minimum titres for the virus components are acceptable.

**Leptospira**

**Indication**

The proposed claims regarding the *Leptospira* components are acceptable as follows:

- to prevent clinical signs, infection and urinary excretion caused by *L. interrogans* serogroup Australis serovar Bratislava,
- to prevent clinical signs and urinary excretion and reduce infection caused by *L. interrogans* serogroup Canicola serovar Canicola and *L. interrogans* serogroup Icterohaemorrhagiae serovar Icterohaemorrhagiae,
- to prevent clinical signs and reduce infection and urinary excretion caused by *L. interrogans* serogroup Grippotyphosa serovar Grippotyphosa.

This could be shown for at least one year after completion of the basic immunisation.

**Maternally derived antibodies (MDA)**

Several studies have been performed to assess the possible influence of MDA on the antibody response to the antigens of Versican Plus DHPPi/L4R. Challenge studies have been conducted to demonstrate that the vaccine is protective against virulent challenge in the presence of MDA against CDV, CAV and CPV at levels equal to or higher than those likely to be encountered under field conditions. The presented laboratory and field studies clearly show the influence of MDA regarding CDV, CPV and CAV antigens. While the MDA negative dogs mostly seroconvert after the first vaccination and are boostered after the second vaccination, MDA positive dogs react after the second vaccination and, in general, have lower titres than those without MDA. These studies demonstrate the importance of the second vaccination as part of the primary vaccination because animals with MDA do not seroconvert after the first vaccination but after the second one. As immunological responses to CDV, CPV and CAV-2 may be delayed due to MDA the vaccination scheme for young dogs should be planned carefully. This has been reflected in section 4.4 of the SPC:

“Immunological responses to the CDV, CAV and CPV components of the vaccine may be delayed due to maternally derived antibody interference. However, the vaccine has been proven to be protective against virulent challenge in the presence of maternally derived antibodies against CDV, CAV and CPV at levels equal or higher to those likely to be encountered under field conditions. In situations where very high maternally derived antibody levels are expected, the vaccination protocol should be planned accordingly.”
As related to the Leptospira components the data show that the interference of MDA does not play a considerable role.

The following vaccination scheme is justified:

**Dosage and route of administration:**

Aseptically reconstitute the lyophilisate with the solvent. Shake well and administer immediately the entire content (1 ml) of the reconstituted product.

**Primary vaccination scheme:**

Two doses of Versican Plus DHPPi/L4 3–4 weeks apart from 6 weeks of age.

**Rabies**

If protection against rabies is required:

First dose: Versican Plus DHPPi/L4 from 8–9 weeks of age.

Second dose: Versican Plus DHPPi/L4R 3–4 weeks later but not before 12 weeks of age.

The efficacy of the rabies fraction is proven after a single dose from 12 weeks of age in laboratory studies. However, in field studies 10% of seronegative dogs did not show seroconversion (> 0.1 IU/ml) 3–4 weeks after single primary vaccination against rabies. Another 17% did not show the 0.5 IU/ml antibody titre against rabies virus required by some non-EU countries to travel in. In case of travelling to risk areas or for travel outside the EU veterinary surgeons may wish to use a two dose primary course including rabies or give an additional rabies vaccination after 12 weeks.

In case of need, dogs younger than 8 weeks can be vaccinated as safety of Versican Plus DHPPi/L4R has been demonstrated in 6 weeks old dogs.

**Re-vaccination scheme:** A single dose of Versican Plus DHPPi/L4 to be given annually.

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**Part 5 – Benefit-risk assessment**

**Introduction**

Versican Plus DHPPi/L4 is a multivalent vaccine which is indicated for the immunisation of healthy puppies and dogs against canine distemper, adenovirus hepatitis, adenovirus respiratory disease, parvovirosis, parainfluenza and leptospirosis. The live virus components of the vaccine (canine distemper virus (CDV), canine adenovirus type 2 (CAV-2), canine parvovirus type 2b (CPV-2b), canine parainfluenza virus (CPiV)) are presented in freeze-dried form in a vial to be reconstituted with a vial of the inactivated components (L. Bratislava, L. Canicola, L. Grippotyphosa and L. Icterohaemorrhagiae) which are presented in liquid form. The liquid fraction also contains the adjuvant (aluminium hydroxide).

**Benefit assessment**

**Direct therapeutic benefit**

Versican Plus DHPPi/L4 is a fixed combination containing eight active substances as detailed in the introduction. The vaccine components are directed against canine infectious diseases present and widespread in most European countries. CDV, CPV and CAV-2 are commonly included in canine vaccines to protect against distemper, parvovirosis and adenovirus type 1 hepatitis which are serious illnesses and
often fatal in young dogs. Rabies is a notifiable disease and a zoonosis which inevitably always results in death. CAV-2 and CPIV infect the canine respiratory tract and on their own usually only cause mild disease. In combination with bacterial pathogens such as *Bordetella bronchiseptica*, however, they can result in kennel cough. In rare cases, severe bronchopneumonia may occur. Kennel cough is highly contagious and can persist for many weeks, and thus of great concern.

Leptospirosis is a zoonotic disease with a variable clinical presentation. Infections can be subclinical or cause acute to chronic renal or hepatic disease which may result in death. An acute icterohaemorrhagic form is known as Weill’s disease. Subclinically infected dogs or dogs that survive acute disease may shed the organism via urine and thereby serve as reservoirs of infection for other dogs and even for humans. Historically, *L. Canicola* and *L. Icterohaemorrhagiae* were seen as the primary causative agents of canine leptospirosis and most canine *Leptospira* vaccines include only these two serovars. In recent years, however, *L. Bratislava* and *L. Grippotyphosa* have been identified with increasing frequency as pathogens of dogs in Europe. This has resulted in the inclusion of these two serovars in novel canine vaccines.

Controlled clinical trials demonstrated that the product is efficacious for the following indications:

Active immunisation of dogs from 8–9 weeks of age:

- to prevent mortality and clinical signs caused by canine distemper virus,
- to prevent mortality and clinical signs caused by canine adenovirus type 1,
- to prevent clinical signs and reduce viral excretion caused by canine adenovirus type 2,
- to prevent clinical signs, leukopenia and viral excretion caused by canine parvovirus,
- to prevent clinical signs (nasal and ocular discharge) and reduce viral excretion caused by canine parainfluenza virus,
- to prevent clinical signs, infection and urinary excretion caused by *L. interrogans* serogroup Australis serovar Bratislava,
- to prevent clinical signs and urinary excretion and reduce infection caused by *L. interrogans* serogroup Canicola serovar Canicola and *L. interrogans* serogroup Icterohaemorrhagiae serovar Icterohaemorrhagiae, and
- to prevent clinical signs and reduce infection and urinary excretion caused by *L. interrogans* serogroup Grippotyphosa serovar Grippotyphosa.

**Additional benefits**

The vaccine Versican Plus DHPPi/L4 is administered subcutaneously which is easy to apply by the veterinarian and reduces stress for the animal. Using large multivalent vaccines for dogs has advantages for the animal and the pet owner. Stress and pain for the animal are reduced as only one vaccine injection is required per visit to the veterinary practice and compliance by the pet owners is usually improved because the appointments at the veterinarian are reduced.

In addition, the use of this vaccine potentially reduces the need for antimicrobial treatment against *Leptospira* infections.

**Risk assessment**

**Main potential risks:**

**Quality:**
The choice of the vaccine strains and the adjuvant has been satisfactorily addressed and reference to the relevance of each strain to current epidemiological conditions is also provided. The manufacturing process of the vaccine has been described in detail for the virus and for the *Leptospira* components. Regarding the starting materials all necessary information has been provided. The TSE risk assessment provided by the applicant clearly demonstrates that the TSE risk of this product is negligible. Compliance with the corresponding Note for Guidance is demonstrated. Controls during manufacture and tests on the finished product should guarantee the compliance with the quality parameter mentioned. Test methods have been described and corresponding validation studies have been performed. Batch to batch consistency has been demonstrated and a detailed overview of all in-process and finished product tests of the vaccine Versican Plus DHPPi/L4 has been provided. Some data is needed for finalisation of reports to fully consolidate the risk assessment. The antigen stability study is due to complete by the end of 2014 and the final study report will be submitted in early 2015. The first 10 batches produced for commercial release will have the individual results for *Leptospira* of the tested rabbits submitted along with the batch release protocols. The final batch release protocol of the filled batch (batch number 015820) will be provided by the end of March 2014. With all these recommendations in place the quality of Versican Plus DHPPi/L4 is considered to be satisfactorily demonstrated.

**Safety for the target animal**

The administration of one dose and a repeated dose of Versican Plus DHPPi/L4 containing maximum potency of antigens by the recommended route was found to be safe for puppies of 6 weeks of age. An overdose study was performed with the live virus components (lyophilisate).

Several studies have been performed to assess the possible influence of MDA on the antibody response to the antigens of Versican Plus DHPPi/L4. The presented laboratory and field studies clearly show the influence of MDA regarding CDV, CPV and CAV antigens.

It has been demonstrated that immunological responses to the CDV, CAV and CPV components of the vaccine may be delayed due to MDA interference. However, the vaccine has been proven to be protective against virulent challenge in the presence of MDA against CDV, CAV and CPV at levels equal or higher to those likely to be encountered under field conditions. The studies demonstrate the importance of the second vaccination as part of the primary vaccination because animals with MDA do not seroconvert after the first vaccination but after the second one. Therefore, the vaccination scheme for young dogs should be planned carefully.

No studies have been performed on reproductive safety and a warning is given in the SPC. The use is not recommended during pregnancy and lactation. No studies have been performed on the concurrent use of any other non-Versican Plus vaccine. This fact is covered in section 4.8 of the SPC. The possible spread and dissemination of the vaccine strain has been correctly reflected in the SPC and is acceptable. No indication of reversion to virulence during the passages was found. In the field study dogs showed signs of lethargy, vomiting, diarrhoea and anorexia after vaccination and these observations are reflected in the SPC.

**User safety:**

The user safety for this product is acceptable when used as recommended and taking into account the safety advice in the SPC.

**Environmental risk assessment:**

The product is not expected to pose any risk to the environment when used as recommended.
Risk management or mitigation measures

Appropriate information has been included in the SPC to inform on the potential risks of this product relevant to the target animal, user and environment and so provide advice on how to prevent or reduce these risks.

Evaluation of the benefit-risk balance

The product has been shown to have a positive benefit-risk balance overall.

The formulation and manufacture of Versican Plus DHPPi/L4 is well described and specifications set will ensure that a product of consistent quality will be produced.

Versican Plus DHPPi/L4 has been demonstrated to be efficacious for the active immunisation of dogs from 8–9 weeks of age

- to prevent mortality and clinical signs caused by canine distemper virus,
- to prevent mortality and clinical signs caused by canine adenovirus type 1,
- to prevent clinical signs and reduce viral excretion caused by canine adenovirus type 2,
- to prevent clinical signs, leukopenia and viral excretion caused by canine parvovirus,
- to prevent clinical signs (nasal and ocular discharge) and reduce viral excretion caused by canine parainfluenza virus,
- to prevent clinical signs, infection and urinary excretion caused by *L. interrogans* serogroup Australis serovar Bratislava,
- to prevent clinical signs and urinary excretion and reduce infection caused by *L. interrogans* serogroup Canicola serovar Canicola and *L. interrogans* serogroup Icterohaemorrhagiae serovar Icterohaemorrhagiae,
- to prevent clinical signs and reduce infection and urinary excretion caused by *L. interrogans* serogroup Grippotyphosa serovar Grippotyphosa.

Based on the laboratory and field studies the proposed vaccination scheme is acceptable. The product is well tolerated by the target animals and presents a low risk for users and the environment. Appropriate warnings of findings in the laboratory and field trials have been included in the SPC.

Conclusion on benefit-risk balance

The overall benefit-risk evaluation for the product is deemed positive with sufficiently clear and complete product information.

Conclusion

Based on the original and complementary data presented, the CVMP concluded that the quality, safety and efficacy of Versican Plus DHPPi/L4 were considered to be in accordance with the requirements of Directive 2001/82/EC.

Based on the CVMP review of the data on quality, safety and efficacy, the CVMP recommended the granting of the marketing authorisation for Versican Plus DHPPi/L4.