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Aftovaxpur DOE (WD) (10) O1 Manisa + A22 Iraq

Non autorisé

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated
- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

Identification du produit

Dénomination du médicament:
Aftovaxpur DOE (WD) (10) O1 Manisa + A22 Iraq

Substance(s) active(s):
Disponible uniquement en Anglais
Disponible uniquement en Anglais
Disponible uniquement en Anglais
Disponible uniquement en Anglais

Espèces cibles:
Bovins
Mouton
Porc
Bovins
Mouton

Porc

Voie d’administration:
Voie intramusculaire
Voie sous-cutanée
Voie sous-cutanée
Voie intramusculaire

Informations sur le produit

Substance(s) active(s) / Concentration:
Disponible uniquement en Anglais
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Disponible uniquement en Anglais
Presentation_strength:≥ 6 PD50 Index:11
Disponible uniquement en Anglais
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Disponible uniquement en Anglais
Presentation_strength:≥ 6 PD50 Index:11

Forme pharmaceutique:
Emulsion injectable

Temps d'attente par voie d'administration:
Voie intramusculaire:

- **Bovins**
  - Non applicable. 0 day Zero days
- **Mouton**
  - Non applicable. 0 day Zero days
- **Porc**

- Non applicable. 0 day
Zero days

Voie sous-cutanée:

- **Bovins**
  - Non applicable. 0 day
  Zero days

- **Mouton**
  - Non applicable. 0 day
  Zero days

- **Porc**
  - Non applicable. 0 day
  Zero days

Voie sous-cutanée:

- **Bovins**
  - Non applicable. 0 day
  Zero days

- **Mouton**
  - Non applicable. 0 day
  Zero days

- **Porc**
  - Non applicable. 0 day
  Zero days

Voie intramusculaire:

- **Bovins**
  - Non applicable. 0 day
  Zero days

-

Mouton

- Non applicable. 0 day Zero days

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Porc

- Non applicable. 0 day Zero days

Code anatomique, thérapeutique, chimique, vétérinaire (ATCvet):
QI02AA04

Conditions de délivrance:
Médicament vétérinaire soumis à prescription vétérinaire

Statut de l’autorisation:
Abandonnée

Autorisé en:
Autriche , Belgique , Bulgarie , Croatie , Chypre , Tchéquie , Danemark , Estonie , Finlande , France , Allemagne , Grèce , Hongrie , Islande , Irlande , Italie , Lettonie , Liechtenstein , Lituanie , Luxembourg , Malte , Pays-Bas , Norvège , Pologne , Portugal , Roumanie , Slovaquie , Slovénie , Espagne , Suède , Royaume-Uni (Irlande du Nord)

Informations complémentaires

Type d'autorisation:

Base légale de l'autorisation du produit:
Disponible uniquement en Anglais Italian

Titulaire de l’autorisation de mise sur le marché:
Boehringer Ingelheim Vetmedica GmbH

Date de l'autorisation de mise sur le marché:
15/07/2013

Site(s) de libération des lots du produit fini:
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France

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Autorité responsable:
European Commission

Numéro de l’autorisation:
Ces informations ne sont pas disponibles pour ce produit.

Date de modification du statut de l’autorisation:
15/07/2013

Pour connaître les évènements indésirables rapportés concernant les médicaments vétérinaires, consulter ce site www.adrreports.eu/vet

Documents

Fichier combinant tous les documents

français (PDF)

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