CAFFEINE AND SODIUM BENZOATE 20% solutio pro injectionibus

Awtorizzat

- Sodium benzoate
- Caffeine

Product identification

Isem tal-medicina:
CAFFEINE AND SODIUM BENZOATE 20% solutio pro injectionibus
КОФЕИН НАТРИЕВ БЕНЗОАТ 20% разтвор за инжекции

Sustanza attiva:
Disponibbli biss fi English
Disponibbli biss fi English

Speci li fuqhom ser jintuża l-prodott:
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian

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Metodu ta' amministrazzjoni:
Użu għal taħt il-ġilda

Użu għal ġol-muskoli

Product details

Sustanza attiva / Qawwa:

Disponibbli biss fi English
1.00 gram(s) / 1.00 Ampoule

Disponibbli biss fi English
1.00 gram(s) / 1.00 Ampoule

Forma farmaċewtika:

Soluzzjoni għall-injezzjoni

Withdrawal period by route of administration:

Użu għal taħt il-ġilda:

- **Horse**
- **Pig**
- **Dog**

Użu għal ġol-muskoli:

- **Horse**
- **Pig**
- **Dog**

Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):

QV04CG30

Status legali tal-provvista:

Disponibbli biss fi Czech Estonian English French Italian Latvian Portuguese Slovenian Finnish Swedish Icelandic Norwegian

Status tal-awtorizzazzjoni:

Valid

Authorised in:

Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Deskrizzjoni tal-pakkett:

Disponibbli biss fi Bulgarian

Additional information

Entitlement type:
Disponibbli biss fi English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Bażi ġuridika tal-awtorizzazzjoni tal-prodott:
Disponibbli biss fi English French Italian Latvian Norwegian

Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:
Vetprom AD

Marketing authorisation date:
5/02/2007

Siti ta' manifattura b'rilaxx tal-lott:
Vetprom AD

Awtorità responsabbli:
Bulgarian Agency For Food Safety

Numru tal-awtorizzazzjoni:
701/ 06.02.2007

Data tal-bidla fl-istatus tal-awtorizzazzjoni:
5/02/2007

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Source URL: *https://medicines.health.europa.eu/veterinary/600000099374*