

SECTION IX  
ADMINISTRATION OF VETERINARY MEDICINAL PRODUCTS

Unique life number (1)  
(See front page of passport)

Part I

Date and place of issue of this Section (1): (See front page of passport) .....

Issuing body for this Section of the identification document (1): (See front page of passport) .....

Part II

**Note: The equine animal is not intended for slaughter for human consumption.**

The equine animal may therefore undergo the administration of veterinary medicinal products authorised in accordance with Article 6(3) or those administered with Article 10(2) of Directive 2001/82/EC.

I, the undersigned Owner (2) Representative of the Owner (2) Keeper (2) declared that the equine animal described in this identification document is not intended for slaughter for human consumption.

Date and Place	Name in capitals and signature of the owner, representative of the owner or keeper of the animal	Name in capital letters and signature of the veterinarian responsible acting in accordance with Article 10(2) of Directive 2001/82/EC
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Part III

**Note: The equine animal is intended for slaughter for human consumption**

Without prejudice to Regulation (EEC) No 2377/90 and Directive 96/22/EC, the equine animal may be subject to medical treatment in accordance with Article 10(3) of Directive 2001/82/EC under the condition that animals so treated can only be slaughtered for human consumption after the end of the general withdrawal period of six months following the date of last administration of the substances listed in accordance with Article 10(3) of that Directive.

MEDICATION RECORD			
Date of last administration, as prescribed, in accordance with Article 10(°) of Directive 2001/82/EC or Date of suspension in accordance with Article 16(°) of Regulation (EC) 504/2008(°) (°) [dd/mm/yy]	– Place – country code – Postal code	Essential substance(s) incorporated in the veterinary medicinal product administered in accordance with Article 10(°) of Directive 2001/82/EC as mentioned in first column (°) (°) or in accordance with Article 16(°) of Regulation (EC) No 504/2008(°) (°)	Veterinarian responsible applying and/or prescribing administration of veterinary medicinal report
			Name: (°) ..... Signature Address: (°) ..... Postal code: (°) ..... Place: (°) ..... Telephone: (°) .....