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Certificate of Analysis
Bulletin d'Analyse

MSD Santé Animale

BA 5 Edition 15 du 09/05/2014

CEPRAVIN DRY COW **3 G**

BATCH N° **4482** **CSSA**

Specification Reference: Dossier HRA/88/035

TEST	METHOD	RESULT	ACCEPTANCE LIMITS AT RELEASE
Appearance	Q-PF- CEPRAVIN	Conform	White to cream coloured suspension
Identity	Colorimetric Q-PF- CEPRAVIN	Conform	Complies with test
Assay	UV/ HPLC Q-PF- CEPRAVIN	0.250	0.243 to 0.283 g/syringe (MFW)
Sterility	BP Vet Q-PF- CEPRAVIN	Conform	Complies with the BP (Vet)
Uniformity of fill weight /extractable weight	In a random sample of 10 applicators Q-PF- CEPRAVIN	Conform	NMT 1 in 10 shall deviate from the mean fill weight by more than +/- 10% and none shall deviate by more than +/- 20%
Related substances	HPLC Q-PF- CEPRAVIN	0.7 0.4 1.4 2.0	Isonicotinamide ≤ 2% Any other individual impurity ≤ 1% Total unknown impurities ≤ 3% Total impurities ≤ 5%

MFW : Mean Fill Weight

Manufacturing Date : June 30th 2014 Expiry Date : May 2017

This batch has been manufactured and released in accordance with GMP requirements and according to the Marketing Authorisation.

Release date : November 19th 2014

Mikaël CAUCHE
Pharmacist Quality Assurance

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