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TOLNAGRO
Rakoczi u. 142-146
H-7100 SZEKSZARD

Certificate of Analysis

Date	04.04.2018
Created at	8:19:18 PM
Customer no.	620540

Material: 90625
Description: Glucose-Lösung 40% ad us. vet. WDT
Packing unit: 10 x 500 ml

Batch: 18P4844
Expiry date: 11.2020 Manufacturing date:

Parameter	Unit	Value	Lower Limit	Upper Limit
imprint batch no. and date of expiry		complies	-	-
text ref. to print approval		complies	-	-
layout ref. to print approval		complies	-	-
print colours ref. to print approval		complies	-	-
solution appearance		complies	-	-
colour of solution		complies	-	-
visible particles		complies	-	-
primary packaging material		complies	-	-
colour of solution (CoA)		complies	-	-
extractable volume (CoA)	ml	515,00	500,00	-
pH value (CoA)		4,6	3,5	5,5
relative density (CoA)	g/ml	1,151	1,140	1,155
refractive index (CoA)		1,392	1,387	1,393
part. contamination invisible >10µm(CoA)	part./ml	0	-	25

Material: 90625
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WDT
Das Tierarztunternehmen.

Parameter	Unit	Value	Lower Limit	Upper Limit
part. contamination invisible >25µm(CoA)	part./ml	0	-	3
Parameter	**Unit**	**Value**	**Lower Limit**	**Upper Limit**
glucose (CoA)	%	102,00	95,00	105,00
Parameter	**Unit**	**Value**	**Lower Limit**	**Upper Limit**
identity glucose (CoA)		complies	-	-
Parameter	**Unit**	**Value**	**Lower Limit**	**Upper Limit**
HMF and related sub. (CoA)	%	0,000	-	0,100
Parameter	**Unit**	**Value**	**Lower Limit**	**Upper Limit**
bact. endotoxins (EP, 2.6.14) (CoA)	I.U./ml	0,05	-	0,25
Parameter	**Unit**	**Value**	**Lower Limit**	**Upper Limit**
sterility (CoA)		complies	-	-

The above mentioned batch has been manufactured and checked in accordance with the requirements of its Marketing Authorization, in accordance with the principles and guidelines of EU GMP, German Drug Law and German Drug Manufacturing Regulation. It conforms to all specifications and is released to the market hereby.

The approval was made by electronic signature of Dr. Rolf Tybussek, Sachkundige Person / Qualified Person, plant Garbsen, on Feb 17, 2018, 7:03:03 PM. This document was generated from a computer-based system validated pursuant to annex 11 of the EU GMP Guide. The data in this document, as well as the documented decision to release are legally valid without a manual signature in full.