CERTIFICATE OF ANALYSIS

Product	:	**AMOXICILLIN SODIUM/CLAVULANATE POTASSIUM 5:1 (Augmentin)**
Product number	:	A0189
Batch number	:	011232
Date of production	:	08-03-2015
Date of analysis	:	09-03-2015
Date of expiry	:	28-02-2019
Packaging and storage	:	in a well-closed container, protected from light at a temperature of 2 – 8 ^{0}C

	SPECIFICATION	RESULT	UNITS	REMARKS
Description	white or almost white powder	complies		
Identification	conform	complies		HPLC
pH	8.0 – 10.0	9.2		
Appearance of solution	clear and colour ≤ ref. sol. Y_6 or YG_6	complies		
Water	≤ 4.0	0.70	%	
Related substances:				
- any impurity	≤ 2.0	0.34	%	
- total impurities	≤ 5.0	0.68	%	
Particle matter:				
- NLT 10 µm	≤ 6000	300	pc/1.2g	
- NLT 25 µm	≤ 600	10	pc/1.2g	
Assay :				
- Amoxicillin	≥ 660	720	µg/mg	
- Clavulanic acid	≥ 132	145	µg/mg	
- Ratio	(4.65 – 5.35):1	4.97:1		

This certificate of analysis is conform the results of the manufacturer.

Conclusie: goedgekeurd
Vrijgifte door: drs. M. van Wissen, apotheker
Kwaliteitsbewaking Duchefa Farma B.V.

Datum: 07-05-2015

The buyer is responsible to test each batch to ensure the product is suitable for their specific purpose.