

CERTIFICATE OF ANALYSIS

Laboratory Salinen Austria AG

"PHARMASAL" – Chemically Pure Salt Sodium Chloride for pharmaceutical use, Natrii chloridum according to European Pharmacopoeia, BP, USP, JP

Lot number:

CRS231019

Retest date:

23.10.2022

Production date:

23.10.2019

26.10.2019

		Specification	Unit	Result	Unit
Idoutification	Ne	nacitive.		conforms	
Identification Identification	Na+ Cl-	positive positive		conforms conforms	-
	NaCl	99,5 - 100,5	%	99,97	%
Assay Bromides	Br-	<= 100,5		<= 100	
Iodides	I-	<= 100 <= 10	ppm	<= 100 <= 10	ppm
Todides	1-	<- 10	ppm	conforms Ph. Eur	ppm
Sulfate	SO4 2-	<= 200	nnm	<= 200	nnm
	PO4 3-	<= 200 <= 25	ppm	<= 200 <= 25	ppm
Phosphate	NO2-		ppm		ppm
Nitrite		<= 0,01	abs.	<= 0,01	abs.
Heavy metals	as Pb	<= 3	ppm	<= 3	ppm
Iron	Fe	<= 2	ppm	<= 2	ppm
Aluminium	Al	<= 0,2	ppm	<= 0,2	ppm
Arsenic	As	<= 1	ppm	<= 1	ppm
Potassium	K	<= 500	ppm	<= 500	ppm
Barium	Ва	<= 10	ppm	<= 10	ppm
				conforms Ph. Eur	
Magnesium & alkaline-earth metals	calc. as Ca	<= 100	ppm	<= 100	ppm
Ferrocyanides	[Fe(CN)6]4-	conforms	-	conforms	-
Insoluble matters		<= 50	ppm	<= 50	ppm
Loss on drying		<= 0,5	%	<= 0,5	%
Appearance of solution		clear, colourless		conforms	-
Acidity or Alkalinity		conforms		conforms	-
according to the regulations		5000 Maria 1900 Maria 1900		6000	
Residual Solvents according ICH-guideline		Impossible due to production process		conforms	-
Bacterial Endotoxins		< 5	I.U./g	< 5	I.U./g
TAMC		<= 10	CFU/g	<= 10	CFU/g
TYMC		<= 10	CFU/g	<= 10	CFU/g

This lot conforms with the current Ph. Eur, USP, BP and JP monographs. In compliance with the guidelines on good manufacturing practice for active pharmaceutical ingredients (ICH Q7). Store in a clean and dry place, nmt. 70% rel. Humidity.

It is suitable for the use in the manufacture of injectable dosage forms, peritoneal dialysis solutions, hemodialysis solutions and hemofiltration solutions.

Vigot grant

Qualified Person Dipl.-Ing.Birgit Spreitz Date:08.11.2019 Salinen Austria AG Labor Steinkogelstraße 30

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