

## **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:

CRS171120

Retest date:

17.11.2023

Production date:

17.11.2020

22.11.2020

		Specification	Unit	Result	Unit
Talanakifi anki an	AI.			-	
Identification Identification	Na+ Cl-	positive		conforms	-
Assay	NaCl	positive	0/	conforms	- 0/
Assay Bromides	Br-	99,5 - 100,5	%	99,97	%
	I-	<= 100 <= 10	ppm	<= 100	ppm
Iodides	1-	<= 10	ppm	<= 10	ppm
Sulfate	CO4 3	. 200		conforms Ph. E	
	SO4 2-	<= 200	ppm	<= 200	ppm
Phosphate	PO4 3-	<= 25	ppm	<= 25	ppm
Nitrite	NO2-	<= 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					
Residual Solvents		Impossible due to		conforms	-
according ICH-guideline		production process		. =	T 11 /
Bacterial Endotoxins		< 5	I.U./g	< 5	I.U./g
ГАМС		<= 10	CFU/g	<= 10	CFU/g
ΓΥΜC		<= 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 alinen Austria AG solutions, hemodialysis solutions and hemofiltration solutions.

Qualified Person Dipl.-Ing.Birgit Spreitz Date:01.12.2020 Steinkogelstraße 30 4802 Ebensee Tel. +43 6132 / 200-0

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