## DRY-CID- dry concentrate for hemodialysis powder, for solution AQUA MEDICA, S.A. DE C.V.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

By diluting the contents of this box to complete 20 gal (75.70 l), one part of this acid concentrate with 44 parts of purified water (UNE-EN ISO 23500-3:2019 standard) will have the following concentrations: Sodium Potassiu m Calcium Magnesium chlorides Acetate Dextrose The final conductivity calculated at 25°C It is 13,2 to 14,2 mS/cm NON STERILE PRODUCT 100,00 mEg/l 2,00 mEg/l 2,50 mEa/l 1,00 mEg/l 105,50 mEg/l 4,00 mEg/l 100,00 mg/dl

Potassium Chlorate

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For use with 3-pump hemodialysis machines only.

For use with 3-pump hemodialysis machines only, using purified water (Standard 13959:2014) and in conbination with

For use with 3-pump hemodialysis machines only.

If the warranty seal is damaged or broken and do not allow debris to fall into the concentrate.

Using purified water (Standard 13959:2014) and in conbination with sodium bicarbonate.

If you do not use the entire contents, discard the excess.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

By diluting one part of this acid concentrate with 44 parts of purified water (ISO Standard 13959:2014).

Store at room temperature.

Keep container tightly closed when not in use.

Sodium Chloride, Anhydrous Calcium Chloride, Anhydrous Magnesium Chloride, Acetic Acid, Dextrosa

	DRY-CID <sup>®</sup> /100.2 DRY CONCENTRATE FOR HEMODIALYSIS	WARNING: Only for use with 3-pump hemodialysis machines, using purified water (UNE-KISO 23500 - 3:2019 standard) and in combination with Sodium Bicarbonate. Verify the dilution of Sodium Bicarbonate.
	By diluting the contents of this box to complete 20 gal (75.70 l), one part of this acid concentrate with 44 parts of purified water (UNE-EN ISO 23500-3:2019 standard) will have the following concentrations: 100,00 mEq/l Potassiu m	THE CONTENTS OF THIS BOX MAKES 20 gai (75.70 I). EMPTY THE FOUR BAGS AND THE GALLON INTO THE BLENDER. THERE SHOULD NOT BE ANY REMAINING. PASS THE RESULTING SOLUTION THROUGH A FRUTER AND STORE AT ROOM TEMPERATURE. DO NOT USE IF SEAL ON GALLON OR BAGS IS DAMAGED.
NON STERILE PRODUCT	Acctate     4,00 mEq/l       Acctate     100,00 mg/dl       Dextrose     100,00 mg/dl       The final conductivity calculated at 25°C It is 13.2 to 14.2 mS/cm       THIS BOX CONTAINS 25.28 kg OF:       Sodium Chloride USP       Potassium Chloride USP       Anthydrous Calcium Chloride USP       Anthydrous Magnesium Chloride USP       0.51 kg       Anthydrous Magnesium Chloride USP       0.47 kg       Antydrous Magnesium Chloride USP       0.61 kg       Acido Acético USP       0.82 kg	Made in Mexico by: AQUA Médica, S.A. de C.V. Carr. Fed. Mex-Cuautla km 65.8 No.8, Col. Tetelcingo, 62757 Cuautla, Mor. LOT No. : Date of Expiry:
	Dextrosa anhidra USP 3.41 kg	IRPLUS OF THIS PRODUCT IN ACCORDANCE WITH THE SAFETY DATA SHEET.

## 81943-601-01

DRY-CID				
dry concentrate for hemodia	lysis powder, for solutio	n		
<b>-</b>				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (S	ource)	NDC:81943-601
Route of Administration	HEMODIALYSIS			
Active Ingredient/Active	e Moiety			
Ingr	edient Name		Basis of Strength	Strength
POTASSIUM CHLORATE (UNII: H35KS68EE7) (CHLORATE ION - UNII:08Z8093742)				
	35KS68EE7) (CHLORATE ION	-	POTASSIUM CHLORATE	0.51 kg in 100 kg
	35KS68EE7) (CHLORATE ION	-		
	35KS68EE7) (CHLORATE ION			
UNII:08Z8093742)		-		
UNII:08Z8093742)	Ingredient Name		CHLORATE	in 100 kg
UNII:08Z8093742)	Ingredient Name <sup>47IQ8X)</sup>	-	CHLORATE 19.91	in 100 kg Strength
UNII:08Z 8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W	<b>Ingredient Name</b> 17IQ8X) <b>US</b> (UNII: OFM21057LP)	-	CHLORATE 19.91 0.47 I	in 100 kg <b>Strength</b> . kg in 100 kg
UNII:08Z8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W CALCIUM CHLORIDE ANHYDRO	Ingredient Name 47IQ8X) US (UNII: OFM21057LP) ROUS (UNII: 59XN63C8VM)	-	CHLORATE 19.91 0.47 I 0.16 I	in 100 kg <b>Strength</b> . kg in 100 kg kg in 100 kg
UNII:08Z 8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W CALCIUM CHLORIDE ANHYDRO MAGNESIUM CHLORIDE ANHYD	Ingredient Name 47IQ8X) US (UNII: OFM21057LP) ROUS (UNII: 59XN63C8VM)	-	CHLORATE 19.91 0.47 I 0.16 I 0.82 I	in 100 kg <b>Strength</b> . kg in 100 kg kg in 100 kg kg in 100 kg
UNII:08Z 8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W CALCIUM CHLORIDE ANHYDRO MAGNESIUM CHLORIDE ANHYDRO ACETIC ACID (UNII: Q40Q9N063F	Ingredient Name 47IQ8X) US (UNII: OFM21057LP) ROUS (UNII: 59XN63C8VM)	-	CHLORATE 19.91 0.47 I 0.16 I 0.82 I	in 100 kg <b>Strength</b> . kg in 100 kg kg in 100 kg kg in 100 kg kg in 100 kg
UNII:08Z 8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W CALCIUM CHLORIDE ANHYDRO MAGNESIUM CHLORIDE ANHYD ACETIC ACID (UNII: Q40Q9N063F ANHYDROUS DEXTROSE (UNII: 1	Ingredient Name 47IQ8X) US (UNII: OFM21057LP) ROUS (UNII: 59XN63C8VM)	-	CHLORATE 19.91 0.47 I 0.16 I 0.82 I	in 100 kg <b>Strength</b> . kg in 100 kg kg in 100 kg kg in 100 kg kg in 100 kg
UNII:08Z 8093742) Inactive Ingredients SODIUM CHLORIDE (UNII: 451W CALCIUM CHLORIDE ANHYDRO MAGNESIUM CHLORIDE ANHYD ACETIC ACID (UNII: Q40Q9N063F ANHYDROUS DEXTROSE (UNII: 1	Ingredient Name 47IQ8X) US (UNII: OFM21057LP) ROUS (UNII: 59XN63C8VM)	-	CHLORATE 19.91 0.47 I 0.16 I 0.82 I	in 100 kg <b>Strength</b> . kg in 100 kg kg in 100 kg kg in 100 kg kg in 100 kg

#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
- <b>-</b> -		25.28 kg in 1 BOX; Type 0: Not a Combination Product	08/30/2023					
		Marketing Information						
Μ	larketing l	nformation						
Μ	larketing   Marketing Category	nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
	Marketing Category	Application Number or Monograph Citation	-	-				

Labeler - AQUA MEDICA, S.A. DE C.V. (816672224)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
AQUA MEDICA, S.A. DE C.V.		816672224	manufacture(81943-601)		

Revised: 8/2023

AQUA MEDICA, S.A. DE C.V.