

DRY-CID- dry concentrate for hemodialysis powder, for solution
Aqua Medica, S.A. de C.V.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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

Potassium

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 mEq/l

2,00 mEq/l

2,50 mEq/l

1,00 mEq/l

105,50 mEq/l

4,00 mEq/l

100,00 mg/dl

Potassium Chlorate

For use with 3-pump hemodialysis machines only.

For use with 3-pump hemodialysis machines only, using purified water (Standard 13959:2014) and in combination 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 combination 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®/100.2
DRY CONCENTRATE FOR HEMODIALYSIS

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	100,00 mEq/l
Potassium	2,00 mEq/l
Calcium	2,50 mEq/l
Magnesium	1,00 mEq/l
chlorides	105,50 mEq/l
Acetate	4,00 mEq/l
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	19,91 kg
Potassium Chloride USP	0,51 kg
Anhydrous Calcium Chloride USP	0,47 kg
Anhydrous Magnesium Chloride USP	0,16 kg
Ácido Acético USP	0,82 kg
Dextrosa anhidra USP	3,41 kg

NON STERILE PRODUCT

WARNING:
 Only for use with 3-pump hemodialysis machines, using purified water (UNE-EN ISO 23500 - 3:2019 standard) and in combination with Sodium Bicarbonate. Verify the dilution of Sodium Bicarbonate.

THE CONTENTS OF THIS BOX MAKES 20 gal (75.70 l). EMPTY THE FOUR BAGS AND THE GALLON INTO THE BLENDER. THERE SHOULD NOT BE ANY REMAINING.

PASS THE RESULTING SOLUTION THROUGH A FILTER AND STORE AT ROOM TEMPERATURE. DO NOT USE IF SEAL ON GALLON OR BAGS IS DAMAGED.

Made in Mexico by:
AQUA Médica, S.A. de C.V.
 Carr. Fed. Mex-Cuautla km 65.8 No.8,
 Col. Tetelecingo, 62757 Cuautla, Mor.

LOT No. :
 Date of Expiry:

AQUA Médica, S.A. de C.V. IS NOT RESPONSIBLE FOR THE IMPROPER DISPOSAL OF THIS CONTAINER. DISPOSE OF THE SURPLUS OF THIS PRODUCT IN ACCORDANCE WITH THE SAFETY DATA SHEET.

81943-601-01

DRY-CID			
dry concentrate for hemodialysis powder, for solution			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81943-601
Route of Administration	HEMODIALYSIS		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
POTASSIUM CHLORATE (UNII: H35KS68EE7) (CHLORATE ION - UNII:08Z8093742)	POTASSIUM CHLORATE	0.51 kg in 100 kg	
Inactive Ingredients			
Ingredient Name	Strength		
SODIUM CHLORIDE (UNII: 451W47IQ8X)	19.91 kg in 100 kg		
CALCIUM CHLORIDE ANHYDROUS (UNII: OFM21057LP)	0.47 kg in 100 kg		
MAGNESIUM CHLORIDE ANHYDROUS (UNII: 59XN63C8VM)	0.16 kg in 100 kg		
ACETIC ACID (UNII: Q40Q9N063P)	0.82 kg in 100 kg		
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)	3.41 kg in 100 kg		
WATER (UNII: 059QF0KO0R)			
Packaging			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81943-601-01	25.28 kg in 1 BOX; Type 0: Not a Combination Product	08/30/2023	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		03/30/2023		

Labeler - Aqua Medica, S.A. de C.V. (589696442)

Establishment

Name	Address	ID/FEI	Business Operations
Aqua Medica, S.A. de C.V.		589696442	manufacture(81943-601)

Revised: 11/2025

Aqua Medica, S.A. de C.V.