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
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 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 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.

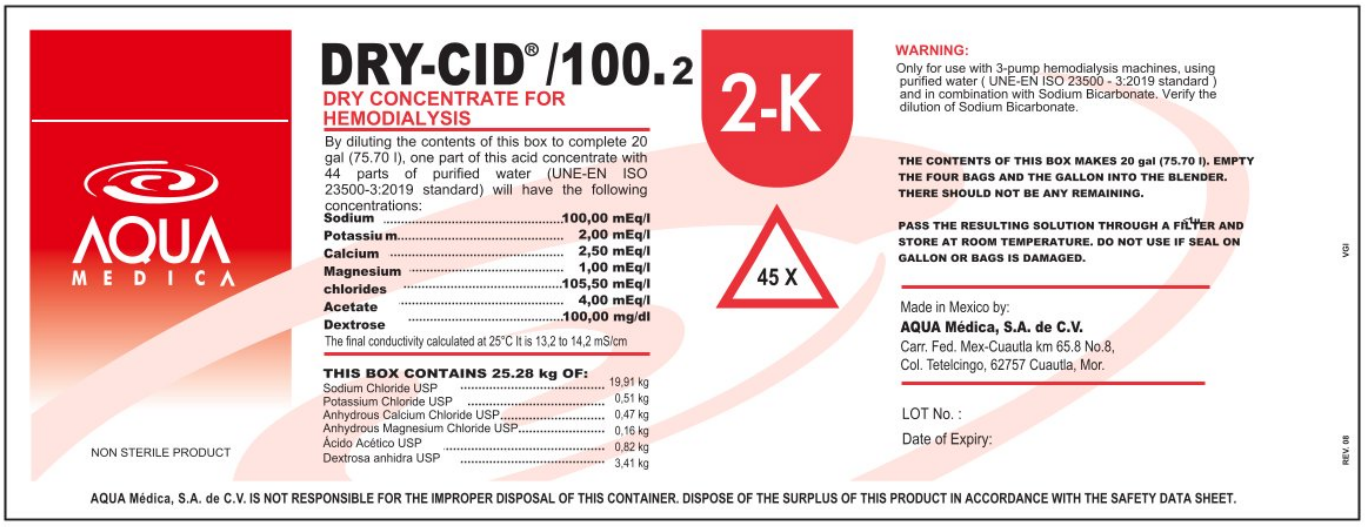
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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



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.