ACD-A- anticoagulant citrate dextrose solution formula a solution Fenwal. Inc.

Anticoagulant Citrate Dextrose Solution USP (ACD) Formula A

CAUTION —

DO NOT REMOVE UNIT FROM OVERWRAP UNTIL READY FOR USE

THE OVERWRAP IS A MOISTURE BARRIER
THE INNER BAG MAINTAINS THE STERILITY OF THE PRODUCT

RECOMMENDED STORAGE-STORE AT CONTROLLED ROOM TEMPERATURE.

CODE 4B7891X NDC 0942-0641-04

1000 mL

Fenwal™

Anticoagulant Citrate Dextrose Solution USP (ACD) Formula A

EACH 100 mL CONTAINS 2.45 g DEXTROSE (MONOHYDRATE) USP 2.2 g SODIUM CITRATE (DIHYDRATE) USP 730 mg CITRIC ACID (ANHYDROUS) USP

STERILE

NONPYROGENIC

SINGLE USE CONTAINER

DISCARD UNUSED PORTION

FOR USE WITH CYTAPHERESIS DEVICE ONLY

NOT FOR DIRECT INTRAVENOUS INFUSION

DO NOT USE UNLESS SOLUTION IS CLEAR AND NO LEAKS ARE DETECTED

AFTER REMOVING OVERWRAP CHECK FOR MINUTE LEAKS BY SQUEEZING INNER BAG FIRMLY. IF LEAKS ARE FOUND DISCARD SOLUTION AS STERILITY MAY BE IMPAIRED

Rx only

PL 146 PLASTIC

Manufactured by:

Baxter Healthcare Corporation

Deerfield, IL 60015 USA

Manufactured for:

Fenwal, Inc. Lake Zurich, IL 60047 USA

07-25-58-006 Made in USA

PACKAGE/LABEL DISPLAY PANEL

	LOT	0	EXP	0			
1 —	DO NOT REM UNTIL READY THE OVERWE THE INNER B OF THE PROI RECOMMENI ROOM TEMP	FOR USE RAP IS A MOI AG MAINTAI DUCT DED STORAG	ISTURE NS THE	BARRIER STERILIT		_	1
2-	CODE 487891					_	2
3-	NDC 0942-064			10	000 mL	-	3
4-	Antico Dextro (ACD)	se So	luti	on US		_	4
5 -	EACH 100 mL (MONOHYDRA (DIHYDRATE) USP	CONTAINS 2	.45 g DI	EXTROSE IUM CITRA		_	5
6- —	STERILE NO SINGLE USE OF FOR USE WITH NOT FOR DIR DO NOT USE	CONTAINER H CYTAPHER RECT INTRAV	DISCAL ESIS DI ENOUS UTION	EVICE ONL'	Y N	_	6
7 -	FOUND DISCA BE IMPAIRED	VING OVERW UEEZING INN ARD SOLUTIO	RAP CH	G FIRMLY. TERILITY M	IF LEAKS ARE	-	7
8-	Rx only Manufactured I Baxter Health Deerfield, IL 60 Manufactured f	care Corporat 015 USA	tion	F	PL 146 PLASTIC	-	8
9-	Fenwal, Inc. Lake Zurich, IL 07-25-58-006 LABEL ISSUE D	Made in USA					9

Product Information

Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:0942-0641

Route of Administration INTRAVENOUS

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Dextrose Monohydrate (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SL0G7R0OK)	Dextrose Monohydrate	24.5 g in 1000 mL
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR) (Anhydrous Citric Acid - UNII:XF417D3PSL)	SODIUM CITRATE, UNSPECIFIED FORM	22 g in 1000 mL
Anhydrous Citric Acid (UNII: XF417D3PSL) (Anhydrous Citric Acid - UNII: XF417D3PSL)	Anhydrous Citric Acid	7.3 g in 1000 mL

Inactive Ingredients				
Ingredient Nam	Strength			
Water (UNII: 059QF0KO0R)				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:0942-0641- 04	1000 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
NDA	BN160918	03/01/2007		

Labeler - Fenwal, Inc. (794519020)

Registrant - Fenwal, Inc. (794519020)

Establishment					
Na me	Address	ID/FEI	Business Operations		
Baxter Healthcare Corporation		059140764	MANUFACTURE(0942-0641)		

Revised: 2/2022 Fenwal, Inc.