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Drug Details

Drug Name(s)	EXTRANEAL
FDA Application No.	(NDA) 021321
Active Ingredient(s)	ICODEXTRIN
Company	BAXTER HLTHCARE
Original Approval or Tentative Approval Date	December 20, 2002
Chemical Type	1 New molecular entity (NME)
Review Classification	S Standard review drug O Orphan drug

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #021321
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
EXTRANEAL	ICODEXTRIN	7.5GM/100ML	SOLUTION;INTRAPERITONEAL	Prescription	Yes	None

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