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Drug Details

Drug Name(s)	SUSTOL
FDA Application No.	(NDA) 022445
Active Ingredient(s)	GRANISETRON
Company	HERON THERAPEUTICS
Original Approval or Tentative Approval Date	August 9, 2016
Chemical Type	5 New formulation or new manufacturer

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #022445

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
SUSTOL	GRANISETRON	10MG/0.4ML	INJECTABLE;INJECTION, EXTENDED RELEASE	Prescription	TBD	 ¹¹ None

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