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Drug Details

Drug Name(s)	VENCLEXTA
FDA Application No.	(NDA) 208573
Active Ingredient(s)	VENETOCLAX
Company	ABBVIE INC
Original Approval or Tentative Approval Date	April 11, 2016
Review Classification	P Priority review drug O Orphan drug

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #208573

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
VENCLEXTA	VENETOCLAX	10MG	TABLET;ORAL	Prescription	TBD  ¹¹	None
VENCLEXTA	VENETOCLAX	50MG	TABLET;ORAL	Prescription	TBD  ¹²	None
VENCLEXTA	VENETOCLAX	100MG	TABLET;ORAL	Prescription	TBD  ¹³	None

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