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Drug Details

Drug Name(s)	VERMOX
FDA Application No.	(NDA) 208398
Active Ingredient(s)	MEBENDAZOLE
Company	JANSSEN PHARMS
Original Approval or Tentative Approval Date	October 19, 2016
Chemical Type	5 New formulation or new manufacturer

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #208398

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
VERMOX	MEBENDAZOLE	500MG	TABLET, CHEWABLE;ORAL	Prescription	TBD  ¹¹	None

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