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**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

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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