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Drug Details

Drug Name(s)	ONZETRA XSAIL
FDA Application No.	(NDA) 206099
Active Ingredient(s)	SUMATRIPTAN
Company	AVANIR PHARMS
Original Approval or Tentative Approval Date	January 27, 2016
Chemical Type	3 New dosage form

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (NDA) #206099

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ONZETRA XSAIL	SUMATRIPTAN	11MG	POWDER;INHALATION	Prescription	No	None

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