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## Drug Details

<b>Drug Name(s)</b>	<b>NITROFURANTOIN</b>
<b>FDA Application No.</b>	<b>(ANDA) 201722</b>
<b>Active Ingredient(s)</b>	<b>NITROFURANTOIN, MACROCRYSTALLINE</b>
<b>Company</b>	<b>SUN PHARM INDS</b>
<b>Original Approval or Tentative Approval Date</b>	<b>February 16, 2016</b>

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- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

## Products on Application (ANDA) #201722

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

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage</a>	<a href="#">Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLD</a>	<a href="#">TE Code</a>
NITROFURANTOIN	NITROFURANTOIN, MACROCRYSTALLINE	25MG		CAPSULE;ORAL	Prescription	No	AB
NITROFURANTOIN	NITROFURANTOIN, MACROCRYSTALLINE	50MG		CAPSULE;ORAL	Prescription	No	AB
NITROFURANTOIN	NITROFURANTOIN, MACROCRYSTALLINE	100MG		CAPSULE;ORAL	Prescription	No	AB

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