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Drug Details

Drug Name(s)	DAPSONE
FDA Application No.	(ANDA) 204074
Active Ingredient(s)	DAPSONE
Company	NORTH CREEK PHARMS
Original Approval or Tentative Approval Date	May 10, 2016

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

Products on Application (ANDA) #204074

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DAPSONE	DAPSONE	25MG	TABLET;ORAL	Prescription	No	AB
DAPSONE	DAPSONE	100MG	TABLET;ORAL	Prescription	No	AB

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U.S. Food and Drug Administration
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Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)
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