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Drug Details

Drug Name(s)	ACZONE
FDA Application No.	(NDA) 207154
Active Ingredient(s)	DAPSONE
Company	ALLERGAN
Original Approval or Tentative Approval Date	February 24, 2016
Chemical Type	5 New formulation or new manufacturer

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

Products on Application (NDA) #207154

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ACZONE	DAPSONE	7.5%	GEL; TOPICAL	Prescription	TBD  ¹¹	None

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