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Drug Details

Drug Name(s)	DOXERCALCIFEROL
FDA Application No.	(ANDA) 203929
Active Ingredient(s)	DOXERCALCIFEROL
Company	AKORN INC
Original Approval or Tentative Approval Date	May 7, 2015

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- **Labels are not available**

Products on Application (ANDA) #203929

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
DOXERCALCIFEROL	DOXERCALCIFEROL	4MCG/2ML (2MCG/ML)	INJECTABLE;INJECTION	Prescription	No AP

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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
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