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## Drug Details

<b>Drug Name(s)</b>	<b>MEXATE</b>
<b>FDA Application No.</b>	<b>(ANDA) 086358</b>
<b>Active Ingredient(s)</b>	<b>METHOTREXATE SODIUM</b>
<b>Company</b>	<b>BRISTOL</b>
<b>Original Approval or Tentative Approval Date</b>	<b>September 5, 1979</b>

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

## Products on Application (ANDA) #086358

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 Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLD</a>	<a href="#">TE Code</a>
MEXATE	METHOTREXATE SODIUM	EQ 20MG BASE/VIAL	INJECTABLE;INJECTION	Discontinued	No	None
MEXATE	METHOTREXATE SODIUM	EQ 50MG BASE/VIAL	INJECTABLE;INJECTION	Discontinued	No	None
MEXATE	METHOTREXATE SODIUM	EQ 100MG BASE/VIAL	INJECTABLE;INJECTION	Discontinued	No	None
MEXATE	METHOTREXATE SODIUM	EQ 250MG BASE/VIAL	INJECTABLE;INJECTION	Discontinued	No	None

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