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Drug Details

Drug Name(s)	LURASIDONE HYDROCHLORIDE
FDA Application No.	(ANDA) 208049
Active Ingredient(s)	LURASIDONE HYDROCHLORIDE
Company	ACCORD HLTHCARE INC

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

Products on Application (ANDA) #208049

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
LURASIDONE HYDROCHLORIDE	LURASIDONE HYDROCHLORIDE	20MG	TABLET, CHEWABLE, TABLET, CAPSULE; ORAL	None (Tentative Approval)	No None
LURASIDONE HYDROCHLORIDE	LURASIDONE HYDROCHLORIDE	40MG	TABLET, CHEWABLE, TABLET, CAPSULE; ORAL	None (Tentative Approval)	No None
LURASIDONE HYDROCHLORIDE	LURASIDONE HYDROCHLORIDE	60MG	TABLET, CHEWABLE, TABLET, CAPSULE; ORAL	None (Tentative Approval)	No None
LURASIDONE HYDROCHLORIDE	LURASIDONE HYDROCHLORIDE	80MG	TABLET, CHEWABLE, TABLET, CAPSULE; ORAL	None (Tentative Approval)	No None
LURASIDONE HYDROCHLORIDE	LURASIDONE HYDROCHLORIDE	120MG	TABLET, CHEWABLE, TABLET, CAPSULE; ORAL	None (Tentative Approval)	No None

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