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Drug Details

Drug Name(s)	PALONOSETRON HYDROCHLORIDE
FDA Application No.	(ANDA) 090713
Active Ingredient(s)	PALONOSETRON HYDROCHLORIDE
Company	TEVA PHARMS USA

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

Products on Application (ANDA) #090713

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
PALONOSETRON HYDROCHLORIDE	PALONOSETRON HYDROCHLORIDE	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	INJECTABLE;INTRAVENOUS	None (Tentative Approval)	No None
PALONOSETRON HYDROCHLORIDE	PALONOSETRON HYDROCHLORIDE	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	INJECTABLE;INTRAVENOUS	None (Tentative Approval)	No None

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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FDA

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