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Active Ingredient:	FENOFIBRATE
Dosage Form;Route:	CAPSULE;ORAL
Proprietary Name:	ANTARA (MICRONIZED)
Applicant:	LUPIN ATLANTIS
Strength:	43MG
Application Number:	N021695
Product Number:	001
Approval Date:	Nov 30, 2004
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	FENOFIBRATE
Dosage Form;Route:	CAPSULE;ORAL
Proprietary Name:	ANTARA (MICRONIZED)
Applicant:	LUPIN ATLANTIS
Strength:	130MG
Application Number:	N021695
Product Number:	003
Approval Date:	Nov 30, 2004
Reference Listed Drug	Yes
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	FENOFIBRATE
Dosage Form;Route:	CAPSULE;ORAL
Proprietary Name:	ANTARA (MICRONIZED)
Applicant:	LUPIN ATLANTIS
Strength:	30MG
Application Number:	N021695
Product Number:	004
Approval Date:	Oct 18, 2013
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	
Patent and Exclusivity Info for this product:	View

Active Ingredient:	FENOFIBRATE
Dosage Form;Route:	CAPSULE;ORAL
Proprietary Name:	ANTARA (MICRONIZED)
Applicant:	LUPIN ATLANTIS
Strength:	90MG
Application Number:	N021695
Product Number:	005
Approval Date:	Oct 18, 2013
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	

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