Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

Search results from the "OB_Rx" table for query on "019766."

Active Ingredient:	SIMVASTATIN
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	ZOCOR
Applicant:	MERCK
Strength:	5MG
Application Number:	N019766
Product Number:	001
Approval Date:	Dec 23, 1991
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	SIMVASTATIN
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	ZOCOR
Applicant:	MERCK
Strength:	10MG
Application Number:	N019766
Product Number:	002
Approval Date:	Dec 23, 1991
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	SIMVASTATIN
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	ZOCOR
Applicant:	MERCK
Strength:	20MG
Application Number:	N019766
Product Number:	003
Approval Date:	Dec 23, 1991
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	SIMVASTATIN
Dosage Form;Route:	TABLET;ORAL

Proprietary Name:	ZOCOR
Applicant:	MERCK
Strength:	40MG
Application Number:	N019766
Product Number:	004
Approval Date:	Dec 23, 1991
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	SIMVASTATIN
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	ZOCOR
Applicant:	MERCK
Strength:	80MG
Application Number:	N019766
Product Number:	005
Approval Date:	Jul 10, 1998
Reference Listed Drug	Yes
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

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10903 New Hampshire Avenue
Silver Spring, MD 20993
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Contact FDA

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Search results from the "OB_Rx" table for query on "019898."

Active Ingredient:	PRAVASTATIN SODIUM
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	PRAVACHOL
Applicant:	BRISTOL MYERS SQUIBB
Strength:	20MG
Application Number:	N019898
Product Number:	003
Approval Date:	Oct 31, 1991
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	PRAVASTATIN SODIUM
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	PRAVACHOL
Applicant:	BRISTOL MYERS SQUIBB
Strength:	40MG
Application Number:	N019898
Product Number:	004
Approval Date:	Mar 22, 1993
Reference Listed Drug	No
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

Active Ingredient:	PRAVASTATIN SODIUM
Dosage Form;Route:	TABLET;ORAL
Proprietary Name:	PRAVACHOL
Applicant:	BRISTOL MYERS SQUIBB
Strength:	80MG
Application Number:	N019898
Product Number:	008
Approval Date:	Dec 18, 2001
Reference Listed Drug	Yes
RX/OTC/DISCN:	RX
TE Code:	**AB**
Patent and Exclusivity Info for this product:	View

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