#### FDA Home<sup>3</sup> Drug Databases<sup>4</sup> Orange Book<sup>5</sup> Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Search results from the "OB\_Rx" table for query on "019766."

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

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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 Ingredients	SIMVASTATIN
Active Ingredient:	SIMVASIATIN
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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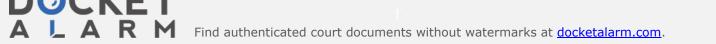
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#### FDA Home<sup>3</sup> Drug Databases<sup>4</sup> Orange Book<sup>5</sup> Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Search results from the "OB\_Rx" table for query on "019898."

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 produc	t: View

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