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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**
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Active Ingredient: SIMVASTATIN
 Dosage Form;Route: 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 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**
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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**
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