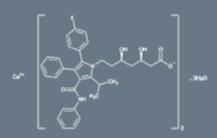
Atorvastatin Calcium

Quality Solutions

Category: Antihyperlipidemics



Authorized Distributor

USP can support your development and manufacturing activities on Atorvastatin Calcium-based medicines with these existing and upcoming standards.

Official and Proposed Documentary Standards and Associated Physical Materials

MONOGRAPH 1

ATORVASTATIN CALCIUM Official as of 01-Jun-2023	
USP Atorvastatin Calcium RS	<u>USP-1044516</u>
USP Atorvastatin Related Compound A RS	<u>USP-1044527</u>
USP Atorvastatin Related Compound B RS	<u>USP-1044538</u>
USP Atorvastatin Related Compound C RS	<u>USP-1044549</u>
USP Atorvastatin Related Compound D RS	<u>USP-1044593</u>

MONOGRAPH 2

ATORVASTATIN CALCIUM TABLETS Official as of 01-Sep-2023	
USP Atorvastatin Calcium RS	<u>USP-1044516</u>
USP Atorvastatin Related Compound B RS	<u>USP-1044538</u>
USP Atorvastatin Related Compound C RS	<u>USP-1044549</u>
USP Atorvastatin Related Compound H RS	<u>USP-1044582</u>

01-Oct-2023. For publications related to **Atorvastatin Calcium** in *PF* online after this date, please visit the integrated *USP-NF/PF* online.

MONOGRAPH 3

AMLODIPINE AND ATORVASTATIN TABLETS Official as of 01-Dec-2020	
USP Amlodipine Besylate RS	<u>USP-1029501</u>
USP Amlodipine Related Compound A RS	<u>USP-1029512</u>
USP Atorvastatin Calcium RS	<u>USP-1044516</u>
USP Atorvastatin Related Compound A RS	<u>USP-1044527</u>
USP Atorvastatin Related Compound B RS	<u>USP-1044538</u>
USP Atorvastatin Related Compound C RS	<u>USP-1044549</u>
USP Atorvastatin Related Compound H RS	<u>USP-1044582</u>

PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*		
USP Atorvastatin Sodium Pyrrolidone Analog PAI I New	<u>USP-1A00810</u>	
USP Atorvastatin Pyrrolidone Lactone PAI	<u>USP-1A00820</u>	
USP Atorvastatin Calcium Epoxy Pyrroloo- xazin 7-Hydroxy Analog PAI I <mark>New</mark>	<u>USP-1A00790</u>	
USP Atorvastatin Epoxy Tetrahydrofuran Analog PAI	<u>USP-1A00800</u>	

^{*} Documents in PF Online are not official and not suitable to demonstrate compliance. They may never become official.

Disclaimer: USP Quality Solution Sheets are provided as a convenience and for informational purposes only. Labmix24 makes reasonable efforts to provide correct information but assumes no liability for the completeness, timeliness or accuracy of the information contained in each QSS. It is the responsibility of the user to verify the information. The US Pharmacopeia remains the official source of this information.



^{*} Pharmaceutical Analytical Impurities (PAI products) are released using a process developed by USP's subject matter experts. The release process is based on internal policies, standard operating procedures, and requirements as defined by USP's Quality Management System. USP is an ISO 9001:2015 certified facility. PAI products are different from official USP Reference Standards. PAI products are not required for compendial compliance.