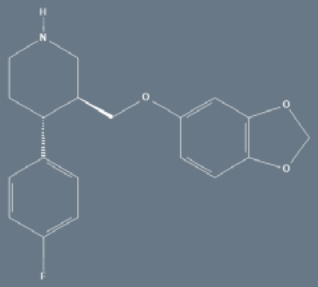
Small Molecules

Paroxetine

Quality Solutions

Category: Psychiatrics



Authorized Distributor

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

Official and Proposed Documentary Standards and Associated Physical Materials

MONOGRAPH 1

PAROXETINE HYDROCHLORIDE Official as of 01-Nov-2021	
USP Paroxetine Hydrochloride RS	USP-1500218
USP Paroxetine Related Compound B RS	USP-1500230
USP Paroxetine Related Compound C RS	USP-1500240
USP Paroxetine Related Compound E RS	USP-1500207
USP Paroxetine Related Compound F RS	USP-1500273
USP Paroxetine Related Compound G RS	USP-1500284
USP Paroxetine System Suitability Mixture A RS	USP-1500353

MONOGRAPH 2

PAROXETINE TABLETS Official as of 01-Aug-2023	
USP Paroxetine Hydrochloride RS	USP-1500218

MONOGRAPH 3

PAROXETINE EXTENDED-RELEASE TABLETS Official as of 01-Aug-2022	
USP Paroxetine Hydrochloride RS	USP-1500218
USP Paroxetine Related Compound B RS	USP-1500230
USP Paroxetine Related Compound F RS	USP-1500273
USP Paroxetine System Suitability Mixture A RS	USP-1500353
PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*	
Paroxetine Methoxy Analog | *NEW*	USP-1A03050

EXCIPIENTS INCLUDED WTIH THIS MOLECULE

MANNITOL Official as of 01-May-2020	
USP Mannitol RS	USP-1375105

MICROCRYSTALLINE CELLULOSE Official as of 01-Dec-2019	
USP Microcrystalline Cellulose RS	USP-1098388

COPOVIDONE Official as of 01-May-2021	
USP Copovidone RS	USP-1148500

SILICON DIOXIDE Official as of 01-Jun-2023

MAGNESIUM STEARATE Official as of 01-Aug-2016	
USP Palmitic Acid RS	USP-1492007
USP Stearic Acid RS	USP-1621008

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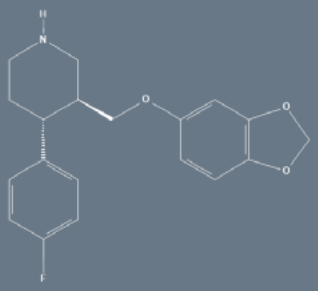

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SODIUM STARCH GLYCOLATE Official as of 01-May-2020	
USP Sodium Starch Glycolate Type B RS	USP-1614670
USP Sodium Starch Glycolate Type A RS	USP-1614669

HYPROMELLOSE Official as of 01-May-2019

TALC Official as of 01-May-2019

TITANIUM DIOXIDE Official as of 01-Jun-2023

[#] *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.*