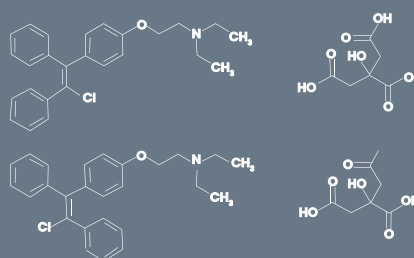


Clomiphene Citrate

Quality Solutions

Category: Steroid



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

MONOGRAPH 1

CLOMIPHENE CITRATE Official as of 1-Oct-2020	
USP Clomiphene Citrate RS	USP-1140000
USP Clomiphene Related Compound A RS	USP-1140101

MONOGRAPH 2

CLOMIPHENE CITRATE TABLETS Official as of 1-Aug-2023	
USP Clomiphene Citrate RS	USP-1140000
USP Clomiphene Related Compound A RS	USP-1140101
PHARMACEUTICAL ANALYTICAL IMPURITIES (PAI)*	
CLOMIPHENE BENZOPHENONE ANALOG - NEW	USP-1A08060
DESCHLOROCLOMIPHENE CHLOROPHENYL CITRATE ANALOG - NEW	USP-1A07420

INCLUDED GENERAL CHAPTERS

<11> USP REFERENCE STANDARDS Official as of 01-Nov-2020	
<191> IDENTIFICATION TESTS—GENERAL Official as of 01-May-2021	
<197> SPECTROSCOPIC IDENTIFICATION TESTS Official as of 01-Sep-2021	
<281> RESIDUE ON IGNITION Official as of 31-Dec-2012	
<621> CHROMATOGRAPHY - NEW Official as of 01-Dec-2024	
<711> DISSOLUTION Official as of 01-May-2023	
USP Dissolution Performance Verification Standard Prednisone RS	USP-1222818
<905> UNIFORMITY OF DOSAGE UNITS Official as of 01-Aug-2023	
<921> WATER DETERMINATION Official as of 01-May-2022	
USP Sodium Tartrate Dihydrate RS	USP-1614909

INCLUDED EXCIPIENTS

MAGNESIUM STEARATE Official as of 1-Aug-2016	
USP Palmitic Acid RS	USP-1492007
USP Stearic Acid RS	USP-1621008

LACTOSE MONOHYDRATE Official as of 01-May-2020	
USP Dextrose RS	USP-1181302
USP Fructose RS	USP-1286504
USP Lactose Monohydrate RS	USP-1356701
USP Sucrose RS	USP-1623637

MICROCRYSTALLINE CELLULOSE Official as of 1-Dec-2019	
USP Microcrystalline Cellulose RS	USP-1098388

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

CORN STARCH Official as of 1-Jun-2023	
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Receive a **15% discount** when purchasing all of the products listed below:

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- USP Clomiphene Related Compound A RS ([USP-1140101](#))
- CLOMIPHENE BENZOPHENONE ANALOG PAI ([USP-1A08060](#))
- DESCHLOROCLOMIPHENE CHLOROPHENYL CITRATE ANALOG PAI ([USP-1A07420](#))

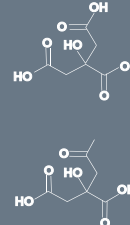
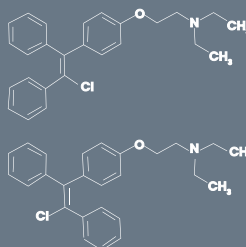
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* *Pharmaceutical Analytical Impurities (PAI products) are released using a process developed by USP's subject matter experts. The release process is based on internal policies, standard operating procedures, and requirements as defined by USP's Quality 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.*

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