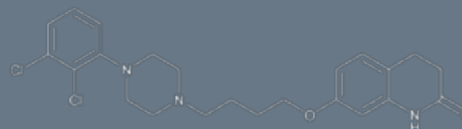


Aripiprazole

Quality Solutions

Category: Psychiatric



Authorized Distributor

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

Official and Proposed Documentary Standards and Associated Physical Materials

MONOGRAPH 1

ARIPIPAZOLE

Official as of 01-May-2020

USP Aripiprazole RS	USP-1042634
USP Aripiprazole Related Compound F RS	USP-1042689

MONOGRAPH 2

ARIPIPAZOLE TABLETS

Official as of 01-Sep-2023

USP Propylparaben RS	USP-1577008
USP Aripiprazole RS	USP-1042634
USP Aripiprazole Related Compound F RS	USP-1042689
USP Aripiprazole Related Compound G RS	USP-1042690

MONOGRAPH 3

ARIPIPAZOLE ORALLY DISINTEGRATING TABLETS

Official as of 01-May-2020

USP Aripiprazole RS	USP-1042634
USP Aripiprazole Related Compound F RS	USP-1042689
USP Aripiprazole Related Compound G RS	USP-1042690

PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*

USP Aripiprazole Acid Analog PAI	USP-1A02470
USP Aripiprazole 4,4'-dimer PAI	USP-1A02460
USP Aripiprazole Spiro Analog PAI	USP-1A02540
USP Aripiprazole Hydroxydihydroquinoline Analog PAI New	USP-1A02550

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

* 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.

01-Oct-2023. For publications related to **Aripiprazole** in PF online after this date, please visit the integrated [USP-NF/PF online](#).

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