UTAK

Certificate of Analysis

Product Information

Product Name:	DRUGS OF ABUSE LEVEL 1
Product Number:	DAU 1 ID
Date of Manufacture:	12/22
Lot Number:	D2359
Expiration Date:	2025-06-30
Intended Use:	In vitro diagnostic
Homogeneity:	Sufficiently homogeneous for analytical testing

Precautions

Matrix:	Urine
Form:	Lyophilized
Anticoagulant:	N/A
Preservative:	Sodium Azide (0.01%)
pH:	4.8-7
Virology:	Non-reactive for HBsAg, HCV, and HIV
Microbial Growth:	<100 colony-forming units per milliliter

Storage and Stability

Storage Condition:	2-8˚C (35-46˚F)
Open Vial Stability:	25 Days
Vial Volume:	3 mL

Origin of Raw Material

UTAK Laboratories, Urine Pool T1-121922-UR1

Analyte	Supplier	Lot#	Purity
D-Amphetamine	Cerilliant Co	FE06242001	100%
D-Methamphetamine	Cerilliant Co	FE12172016	100%
(±)-3,4-Methylenedioxymethamphetamine (MDMA)	Cerilliant Co	FE06102011	100%
Secobarbital	Cerilliant Co	FE05042111	100%
Nordiazepam	Cerilliant Co	FE10012008	100%
Oxazepam	Cerilliant Co	FE02042105	100%
(l)-9-Carboxy-11-Nor-Delta-9-Thc (THC-COOH)	Cerilliant Co	FN09252110	100%
Benzoylecgonine (BE)	Cerilliant Co	FE03032102	100%
Cocaethylene	Cerilliant Co	FE09301903	100%
Lysergic Acid Diethylamide (LSD)	Cerilliant Co	FE04142117	100%
Methadone	Cerilliant Co	FE06252002	100%
Methaqualone	Lipomed INC	765.1B2.1	100%
Morphine	Cerilliant Co	FE03232010	100%

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Analyte	Supplier	Lot#	Purity
Oxycodone	Cerilliant Co	FE01082008	100%
Phencyclidine (PCP)	Cerilliant Co	FE03222003	100%
(+)-Propoxyphene	Cerilliant Co	FE12051901	100%
Nortriptyline	Cerilliant Co	FE06112121	100%

Certification and Compliance Statements

I hereby certify that the above information is authentic and accurate. This batch of product has been manufactured, including packaging/labelling and quality control at the below mentioned site in full compliance with the GMP requirements of the local Regulatory Authority and with the specifications in the Marketing Authorization of the importing country. The batch processing, packaging and analysis records were reviewed and found to be in compliance with GMP. Additional product information is available on the device labeling.

Approved by: Andrew Hartmann
Quality Assurance