

ARK™ UR-144/JWH-018 Control

This ARK Diagnostics, Inc. package insert for the ARK UR-144/JWH-018 Control must be read carefully prior to use. Package insert instructions must be followed accordingly. Reliability of the assay results cannot be guaranteed if there are any deviations from the instructions in this package insert.

Report any serious incident that has occurred in relation to the device to the manufacturer and the appropriate competent authority as applicable.

Customer Service







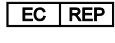





48089 Fremont Blvd
 Fremont, CA 94538 USA
 Tel: 1-877-869-2320
 Fax: 1-510-270-6298
 customersupport@ark-tdm.com
 www.ark-tdm.com
 SRN: US-MF-000023925



EC REP

Emergo Europe
 Westervoortsedijk 60
 6827 AT Arnhem
 The Netherlands

Key to Symbols Used

	Batch code	 YYYY-MM-DD	Use by/Expiration date
	Catalog Number		Manufacturer
	Authorized Representative		CE Mark with notified body number
	Consult Instructions for Use		Quality Control
	Temperature limitation		In Vitro Diagnostic Medical Device
Rx Only	For Prescription Use Only		

1 Name

ARK™ UR-144/JWH-018 Control

2 Intended Use

The ARK UR-144/JWH-018 Control is intended for use in quality control of the ARK UR-144/JWH-018 Assay.

3 Content

The ARK UR-144/JWH-018 Control is composed of a non-sterile, processed human urine matrix with the following target concentrations of UR-144 Pentanoic Acid.

REF	Product Description	Quantity/Volume
5054-0003-00	ARK UR-144/JWH-018 Control UR-144 Pentanoic Acid, human urine, stabilizer and sodium azide	Dropper Vials
	LOW / Negative (5 ng/mL)	2 X 10 mL
	HIGH / Positive (15 ng/mL)	2 X 10 mL

Traceability and Value Verification: A certified solution of UR-144 Pentanoic Acid is traceable to HPLC. Testing is performed with the ARK UR-144/JWH-018 Assay calibrated with the ARK UR-144/JWH-018 Calibrator.

Each laboratory should establish its own ranges for each new lot of controls based on its own test system and criteria.

The Low Control should be Negative and the High Control should be Positive relative to the 10 ng/mL Cutoff Calibrator.

Controls are made with non-sterile, processed human urine. Donors were non-reactive in tests for HIV 1/2, HBsAg, HCV, HIV-1 (NAT), HCV (NAT) and RPR.

4 Warnings and Precautions

- For *In Vitro* Diagnostic Use. For prescription use only.
- Harmful if swallowed.
- Contains human urine. Handle as potentially infectious.
- Do not mix controls from different lot numbers.
- Use each lot as a set.
- Product contains ≤0.09% sodium azide. As a precaution, affected plumbing including instrumentation should be flushed adequately with water to mitigate the potential accumulation of explosive metal azides.

5 Instructions For Use

- For a complete summary and explanation of the UR-144/JWH-018 Assay, refer to the package insert for the ARK UR-144/JWH-018 Assay.
- Controls are ready to use. Mix each level by gentle inversion before dispensing.
- Squeeze sufficient volume (~40µL/drop) into individual sample cups for each level. Consult instrument-specific sample volume requirements. Return caps to their original containers and keep tight.
- Store vials at 2-8°C. Once opened, use within 12 months and prior to the expiration date.

6 Limitations of Procedure

Accurate and reproducible results are dependent upon properly functioning instruments, reagents, calibrators, controls, storage of product as directed, and good laboratory technique.

7 Trademarks

ARKTM is a trademark of ARK Diagnostics, Inc.

Other brand or product names are trademarks of their respective holders.



ARK Diagnostics, Inc.
Fremont, CA 94538 USA

Revised July 2025
1600-0917-00 Rev 03