

TEST CHANGE

Methadone and Metabolite, Urine, Quantitative 0090362, CDCO METH

Specimen Requirements:

Patient Preparation:

Collect: Random urine.

Specimen Preparation: Transfer 1 mL with no additives or preservatives urine to an

ARUP standard transport tube. Standard Transport Tube. (Min:

Effective Date: February 20, 2024

0.5 mL)

Transport Temperature: Room temperature.

Unacceptable Conditions: Specimens exposed to repeated freeze/thaw cycles.

Remarks:

Stability: Ambient: 1 week; Refrigerated: 1 month; Frozen: 3 years

Methodology: Quantitative Liquid Chromatography-Tandem Mass

Spectrometry

Performed: Sun-Sat

Reported: 1-74 days

Note: Compare to Pain Management, Methadone, Quantitative, with

medMATCH, Urine; Pain Management, Methadone, with

Confirmation with medMATCH, Urine.

CPT Codes: 80358 (Alt code: G0480)

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Methodology: Quantitative Liquid Chromatography-Tandem Mass Spectrometry

Positive cutoff: 100 ng/mL

For medical purposes only; not valid for forensic use.

The absence of expected drug(s) and/or drug metabolite(s) may indicate noncompliance-noncompliance-noncompliance-noncompliance-noncompliance, inappropriate timing of specimen collection relative to drug administration, poor drug absorption, diluted/adulterated urine, or limitations of testing. The concentration value must be greater than or equal to the cutoff to be reported as positive. Interpretive questions should be directed to the laboratory.



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This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Reference Interval:

Orugs Covered	Cutoff

Concentrations

Methadone 100 ng/mL EDDP 100 ng/mL