

**Overview****Useful For**

Preferred screening test for detection of arsenic, cadmium, mercury, and lead in random urine specimens

**Profile Information**

Test Id	Reporting Name	Available Separately	Always Performed
ASCU	Arsenic/Creatinine Ratio, U	Yes, (order ASUCR)	Yes
CDCU	Cadmium/Creatinine Ratio, U	Yes, (order CDUCR)	Yes
HGCU	Mercury/Creatinine Ratio, U	Yes, (order HGUCR)	Yes
PBCU	Lead/Creatinine Ratio, U	Yes, (order PBU CR)	Yes
CRETR	Creatinine, Random, U	No	Yes

**Reflex Tests**

Test Id	Reporting Name	Available Separately	Always Performed
SPAS	Arsenic Speciation, Random, U	Yes	No

**Testing Algorithm**

If the total arsenic concentration is 10 mcg/L or greater, then speciation will be performed at an additional charge.

For more information see [Porphyria \(Acute\) Testing Algorithm](#)

**Special Instructions**

- [Porphyria \(Acute\) Testing Algorithm](#)
- [Metals Analysis Specimen Collection and Transport](#)

**Method Name**

ASCU, CDCU, HGCU, PBCU: Triple-Quadrupole Inductively Coupled Plasma Mass Spectrometry (ICP-MS/MS)

CRETR: Enzymatic Colorimetric Assay

**NY State Available**

Yes

**Specimen**

**Specimen Type**

Urine

**Specimen Required**

1. For the 48-hour period prior to start of collection, patient **should not** eat seafood.
2. High concentrations of gadolinium and iodine are known to potentially interfere with most inductively coupled plasma mass spectrometry-based metal tests. If either gadolinium- or iodine-containing contrast media has been administered, **a specimen should not be collected for 96 hours.**

**Supplies:** Urine Tubes, 10 mL (T068)**Collection Container/Tube:** Clean, plastic urine container with no metal cap or glued insert**Submission Container/Tube:** Plastic, 10-mL urine tube or clean, plastic aliquot container with no metal cap or glued insert**Specimen Volume:** 6 mL**Collection Instructions:**

1. Collect a random urine specimen.
2. See [Metals Analysis Specimen Collection and Transport](#) for complete instructions.

**Specimen Minimum Volume**

3 mL

**Reject Due To**

All specimens will be evaluated at Mayo Clinic Laboratories for test suitability.

**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Urine	Refrigerated (preferred)	7 days	
	Frozen	7 days	

**Clinical & Interpretive****Clinical Information**

Arsenic (As), lead (Pb), cadmium (Cd), and mercury (Hg) are well-known toxins, and toxic exposures are characterized by increased urinary excretion of these metals.

Arsenic is a naturally occurring element that is usually found in the environment combined with other elements such as oxygen, chlorine, and sulfur. Arsenic combined with these elements is called inorganic arsenic. Arsenic combined with carbon and hydrogen is referred to as organic arsenic. The organic forms (eg, arsenobetaine and arsenocholine) are relatively nontoxic, while the inorganic forms are toxic. The toxic inorganic forms are arsenite (As[3+]/As[III]) and arsenate (As[5+]/As[V]). Inorganic As(V) is readily reduced to inorganic As(III), which is then primarily broken down to the less toxic methylated metabolites monomethylarsonic acid and subsequently dimethylarsinic acid.

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People are exposed to arsenic by eating food, drinking water, or breathing air. Of these, food is usually the largest source of arsenic. The predominant dietary source of arsenic is seafood, followed by rice/rice cereal, mushrooms, and poultry. While seafood contains the greatest amounts of arsenic, from fish and shellfish, this is mostly in an organic form of arsenic called arsenobetaine, which is much less harmful. Some seaweed may contain arsenic in the inorganic form, which is more toxic. In the United States, some areas also contain high natural levels of arsenic in rock, which can lead to elevated levels in the soil and drinking water. Occupational (eg, copper or lead smelting, wood treating, or pesticide application) exposure is another source where people may be introduced to elevated levels of arsenic. Lastly, hazardous waste sites may contain large quantities of arsenic and, if not disposed of properly, may get into the surrounding water, air, or soil.

A wide range of signs and symptoms may be seen in acute arsenic poisoning including headache, nausea, vomiting, diarrhea, abdominal pain, hypotension, fever, hemolysis, seizures, and mental status changes. Symptoms of chronic poisoning, also called arseniasis, are mostly insidious and nonspecific. The gastrointestinal tract, skin, and central nervous system are usually involved. Nausea, epigastric pain, colic abdominal pain, diarrhea, and paresthesias of the hands and feet can also occur.

Since arsenic is excreted predominantly by glomerular filtration, measurement of arsenic in urine is the most reliable means of detecting arsenic exposures within the last several days.

Arsenic toxicity affects a number of organ systems.

Lead toxicity primarily affects the gastrointestinal, neurologic, and hematopoietic systems.

Chronic exposure to cadmium causes accumulated kidney damage.

The correlation between the levels of mercury excretion in the urine and the clinical symptoms is considered poor.

**Reference Values****ARSENIC/CREATININE:**

0-17 years: Not established

> or =18 years: <24 mcg/g creatinine

**CADMIUM/CREATININE:**

0-17 years: Not established

> or =18 years: <0.6 mcg/g creatinine

**MERCURY/CREATININE:**

0-17 years: Not established

> or =18 years: <2 mcg/g creatinine

**LEAD/CREATININE:**

0-17 years: Not established

> or =18 years: <2 mcg/g creatinine

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**CREATININE:**

> or =18 years: 16-326 mg/dL

Reference values have not been established for patients who are younger than 18 years.

**Interpretation****Arsenic:**

Physiologically, arsenic exists in a number of toxic and nontoxic forms. The total arsenic concentration reflects all the arsenic present in the sample regardless of species (eg, inorganic vs. methylated vs. organic arsenic). The measurement of urinary total arsenic levels is generally accepted as the most reliable indicator of recent arsenic exposure. However, if the total urine arsenic concentration is elevated, arsenic speciation must be performed to identify if it is a toxic form (eg, inorganic and methylated forms) or a relatively nontoxic organic form (eg, arsenobetaine and arsenocholine).

The inorganic toxic forms of arsenic (eg, As[III] and As[V]) are found in the urine shortly after ingestion, whereas the less toxic methylated forms, monomethylarsonic acid (MMA) and dimethylarsinic acid (DMA), are the species that predominate longer than 24 hours after ingestion. In general, urinary As(III) and As(V) concentrations peak in the urine at approximately 10 hours and return to normal 20 to 30 hours after ingestion. Urinary MMA and DMA concentrations normally peak at approximately 40 to 60 hours and return to baseline 6 to 20 days after ingestion.

This test can determine if a patient has been exposed to above-average levels of arsenic. It cannot predict whether the arsenic levels in their body will affect their health.

**Cadmium:**

Urine cadmium levels primarily reflect total body burden of cadmium. Cadmium excretion above 3.0 mcg/g creatinine indicates significant exposure to cadmium.

For occupational testing, OSHA cadmium standard is below 3.0 mcg/g creatinine, and the biological exposure index is 5 mcg/g creatinine.

**Mercury:**

The correlation between the levels of mercury (Hg) excretion in the urine and the clinical symptoms is considered poor.

Previous thought indicated urine as a more appropriate marker of inorganic mercury because organic mercury represented only a small fraction of urinary mercury. Based on possible demethylation of methylmercury within the body, urine may represent a mixture of dietary methylmercury and inorganic mercury. Seafood consumption can contribute to urinary mercury levels (up to 30%),<sup>(1)</sup> consistent with the suggestion that due to demethylation processes in the human body, a certain proportion of urinary mercury can originate from dietary consumption of fish/seafood.<sup>(2)</sup>

**Lead:**

Measurements of urinary lead levels have been used to assess lead exposure. However, like lead blood, urinary lead excretion mainly reflects recent exposure and thus shares many of the same limitations for assessing lead body burden or long-term exposure.<sup>(3,4)</sup>

Urinary lead concentration increases exponentially with blood lead and can exhibit relatively high intra-individual variability, even at similar blood lead concentrations.<sup>(5,6)</sup>

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**Cautions**

Consumption of seafood before collection of a urine specimen for arsenic testing is likely to result in a report of an elevated concentration of arsenic found in the urine, which can be clinically misleading.

**Clinical Reference**

1. Tratnik JS, Falnoga I, Mazej D, et al. Results of the first national human biomonitoring in Slovenia: Trace elements in men and lactating women, predictors of exposure and reference values. *Int J Hyg Environ Health*. 2019;222(3):563-582. doi:10.1016/j.ijheh.2019.02.008
2. Sherman LS, Blum JD, Franzblau A, Basu N. New insights into biomarkers of human mercury exposure using naturally occurring mercury stable isotopes. *Environ Sci Technol*. 2013;47(7):3403-3409. doi:10.1021/es305250z
3. Sakai T. Biomarkers of lead exposure. *Ind Health*. 2000;38(2):127-142. doi:10.2486/indhealth.38.127
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6. Skerfving S, Ahlgren L, Christoffersson JO. Metabolism of inorganic lead in man. *Nutr Res*. 1985;Suppl 1:601-607
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11. de Burbane C, Buchet JP, Leroyer A, et al. Renal and neurologic effects of cadmium, lead, mercury, and arsenic in children: evidence of early effects and multiple interactions at environmental exposure levels. *Environ Health Perspect*. 2006;114(4):584-590. doi:10.1289/ehp.8202
12. Agency for Toxic Substances and Disease Registry. Toxicological profile for arsenic. US Department of Health and Human Services; 2007. Available at [www.atsdr.cdc.gov/ToxProfiles/tp2.pdf](http://www.atsdr.cdc.gov/ToxProfiles/tp2.pdf)
13. Bernhoft RA. Mercury toxicity and treatment: a review of the literature. *J Environ Public Health*. 2012;2012:460508. doi:10.1155/2012/460508
14. Strathmann FG, Blum LM. Toxic elements. In: Rifai N, Chiu RWK, Young I, Burnham CD, Wittwer CT, eds. *Tietz Textbook of Laboratory Medicine*. 7th ed. Elsevier; 2023:chap 44

**Performance****Method Description**

The metal analytes of interest are analyzed by triple-quadrupole inductively coupled plasma mass spectrometry.(Unpublished Mayo method)

**PDF Report**

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No

**Day(s) Performed**

Monday through Friday

**Report Available**

2 to 4 days

**Specimen Retention Time**

14 days

**Performing Laboratory Location**

Mayo Clinic Laboratories - Rochester Superior Drive

**Fees & Codes****Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

**Test Classification**

This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. It has not been cleared or approved by the US Food and Drug Administration.

**CPT Code Information**

82175

82300

83825

83655

82570

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
HMUCR	Heavy Metal/Creat Ratio,w/Reflex,U	29589-9

Result ID	Test Result Name	Result LOINC® Value
CRETR	Creatinine, Random, U	2161-8
608900	Arsenic/Creatinine Ratio, U	13463-5
608901	Total Arsenic Concentration	5586-3
608902	Cadmium/Creatinine Ratio, U	13471-8
608903	Mercury/Creatinine Ratio, U	13465-0

## Test Definition: HMUCR

Heavy Metal/Creatinine Ratio, with Reflex,  
Random, Urine

608904	Lead/Creatinine Ratio, U	13466-8
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