Overview

Useful For
An auxiliary test to fractionated plasma and urine metanephrine measurements in the diagnosis of pheochromocytoma and paraganglioma

An auxiliary test to urine vanillylmandelic acid and homovanillic acid determination in the diagnosis and follow-up of patients with neuroblastoma and related tumors

Special Instructions
- Urine Preservatives-Collection and Transportation for 24-Hour Urine Specimens

Method Name
High-Performance Liquid Chromatography (HPLC)
Includes unconjugated epinephrine, norepinephrine, and dopamine

NY State Available
Yes

Specimen

Specimen Type
Urine

Advisory Information
This assay is of greatest value when the specimen is collected during a hypertensive episode.

Unless the purpose of the measurement is drug monitoring, discontinue any epinephrine, norepinephrine, or dopamine injections or infusions for at least 12 hours before specimen draw.

Do not perform the test on patients withdrawing from legal or illegal drugs known to cause rebound catecholamine release during withdrawal (see Cautions for details)

Necessary Information
24-Hour volume is required.

Specimen Required

Patient Preparation: Discontinue drugs that release or hinder metabolism of epinephrine, norepinephrine, or dopamine for at least 1 week before obtaining the specimen (see Cautions for details). If this is not possible for medical reasons, contact the laboratory to discuss whether a shorter drug-withdrawal period may be acceptable.

Supplies: Plastic, 10-mL urine tube (T068)

Specimen Volume: 2mL

Collection Instructions:
1. Collect urine for 24 hours.
2. Add 25 mL of 50% acetic acid as preservative at start of collection. Use 15 mL of 50% acetic acid for children less than 5 years old. This preservative is intended to achieve a pH of between approximately 2 and 4.

**Additional Information:** See Urine Preservatives-Collection and Transportation for 24-Hour Urine Specimens in Special Instructions for multiple collections.

**Forms**

If not ordering electronically, complete, print, and send an Oncology Test Request (T729) with the specimen.

**Urine Preservative Collection Options**

**Note:** The addition of preservative or application of temperature controls **must occur within 4 hours of completion** of the collection.

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<th>Frozen</th>
<th>50% Acetic Acid</th>
<th>Boric Acid</th>
<th>Diazolidinyl Urea</th>
<th>6M Hydrochloric Acid</th>
<th>6M Nitric Acid</th>
<th>Sodium Carbonate</th>
<th>Thymol</th>
<th>Toluene</th>
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**Specimen Minimum Volume**

1.5 mL

**Reject Due To**

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**Specimen Stability Information**

<table>
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<tr>
<th>Specimen Type</th>
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<td>Urine</td>
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<tr>
<td></td>
<td>Frozen</td>
<td>14 days</td>
</tr>
<tr>
<td></td>
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Clinical and Interpretive

Clinical Information

The catecholamines (dopamine, epinephrine, and norepinephrine) are derived from tyrosine via a series of enzymatic conversions. All 3 catecholamines are important neurotransmitters in the central nervous system and play crucial roles in the autonomic regulation of many homeostatic functions, namely, vascular tone, intestinal and bronchial smooth muscle tone, cardiac rate and contractility, and glucose metabolism. Their actions are mediated via alpha and beta adrenergic receptors and dopamine receptors, all existing in several subforms. The 3 catecholamines overlap but also differ in their receptor activation profile and consequent biological actions.

The systemically circulating fraction of the catecholamines is derived almost exclusively from the adrenal medulla, with small contributions from sympathetic ganglia. They are normally present in the plasma in minute amounts, but levels can increase dramatically and rapidly in response to change in posture, environmental temperature, physical and emotional stress, hypovolemia, blood loss, hypotension, hypoglycemia, and exercise.

In patients with pheochromocytoma, a potentially curable tumor of catecholamine producing cells of the adrenal medulla, or less commonly of sympathetic ganglia (paraganglioma), urine catecholamine levels may be elevated. This results in episodic or sustained hypertension and often in intermittent attacks of palpitations, cardiac arrhythmias, headache, sweating, pallor, anxiety, tremor, and nausea ("spells"). Elevations of the urine levels of 1 or several of the catecholamines also may be observed in patients with neuroblastoma and related tumors (ganglioneuroblastomas and ganglioneuromas) and, very occasionally, in other neuroectodermal tumors.

At the other end of the spectrum, inherited and acquired syndromes of autonomic dysfunction/failure and autonomic neuropathies are characterized by either inadequate production of 1 or several of the catecholamines, or by insufficient release of catecholamines upon appropriate physiological stimuli (eg, change in posture from supine to standing, cold exposure, exercise, stress).

Reference Values

NOREPINEPHRINE

<1 year: <11 mcg/24 hours
1 year: 1-17 mcg/24 hours
2-3 years: 4-29 mcg/24 hours
4-6 years: 8-45 mcg/24 hours
7-9 years: 13-65 mcg/24 hours
> or =10 years: 15-80 mcg/24 hours

EPINEPHRINE

<1 year: <2.6 mcg/24 hours
1 year: <3.6 mcg/24 hours
2-3 years: <6.1 mcg/24 hours
4-9 years: 0.2-10.0 mcg/24 hours
10-15 years: 0.5-20.0 mcg/24 hours
> or =16 years: <21 mcg/24 hours

DOPAMINE
<1 year: <86 mcg/24 hours
1 year: 10-140 mcg/24 hours
2-3 years: 40-260 mcg/24 hours
> or =4 years: 65-400 mcg/24 hours

For SI unit Reference Values, see https://www.mayocliniclabs.com/order-tests/si-unit-conversion.html

Interpretation
Diagnosis of Pheochromocytoma:

This test should not be used as the first-line test for pheochromocytoma. PMET / Metanephrines, Fractionated, Free, Plasma (the most sensitive assay) and/or METAF / Metanephrines, Fractionated, 24 Hour, Urine (almost as sensitive and highly specific) are the recommended first-line laboratory tests for pheochromocytoma.

However, urine catecholamine measurements can still be useful in patients whose plasma metanephrines or urine metanephrines measurements do not completely exclude the diagnosis. In such cases, urine catecholamine specimens have an 86% diagnostic sensitivity when cut-offs of >80 mg/24 hour for norepinephrine and >20 mg/24 hour for epinephrine are employed. Unfortunately, the specificity of these cut-off levels for separating tumor patients from other patients with similar symptoms is only 88%. When more specific (98%) decision levels of >170 mg/24 hours for norepinephrine or >35 mg/24 hours for epinephrine are used, the assay’s sensitivity falls to about 77%.

Diagnosis of Neuroblastoma:

Vanillylmandelic acid, homovanillic acid, and sometimes urine catecholamine measurements on spot urine or 24-hour urine are the mainstay of biochemical diagnosis and follow-up of neuroblastoma; 1 or more of these tests may be elevated.

Cautions

Many alterations in physiologic and pathologic states can profoundly affect catecholamine concentrations.

Any environmental factors that may increase endogenous catecholamine production should be avoided. These include noise, stress, discomfort, body position, and the consumption of food, caffeinated beverages, and nicotine. Caffeine and nicotine effects are short term, a few minutes to hours only.

Other substances and drugs that may affect the results include:

Substances that result in increased release or diminished metabolism of endogenous catecholamines:

-Monamine oxidase inhibitors (MOIs): a class of anti-depressants with marked effects on catecholamine levels, particularly if the patient consumes tyrosine rich foods, such as nuts, bananas, or cheese
- Catecholamine reuptake inhibitors including cocaine and synthetic cocaine derivatives, such as many local anesthetics, which also can be antiarrhythmic drugs (eg, lidocaine)

- Some anesthetic gases, particularly halothane

- Withdrawal from sedative drugs, medical or recreational, in particular alcohol, benzodiazepines (eg, Valium), opioids, and some central acting antihypertensive drugs, particularly Clonidine, but, generally not cannabis or other hallucinogens such as lysergic acid diethylamide (LSD), mescal, or peyote

- Vasodilating drugs (eg, calcium antagonists, alpha-blockers)

- Tricyclic antidepressants usually exert a negligible effect

Substances that reduce or increase plasma volume acutely (eg, diuretics, radiographic contrast media, synthetic antidiuretic hormone [eg, desmopressin 1-deamino-8-d-arginine vasopressin: DDAVP])

Historically, a third category of potentially interfering substances was represented by molecules that are either similar in chemical structure, antibody epitopes, or chromatographic migration pattern to the catecholamines, or have metabolites that can be mistaken for the catecholamines. Our current HPLC-based assay is not subject to any significant direct interference of this kind. In most cases, the following drugs do not cause problems with the current assay that cannot be resolved: acetaminophen, allopurinol, amphetamines and its derivatives (methylphenidate, methylenedioxyamphetamine [MDMA: ecstasy]), atropine, beta blockers (atenolol, labetalol, metoprolol, sotalol), buspirone, butalbital, carbamazepine, clorazepate, chlor diazepoxide, chlorpromazine, chlorothiazide, chlorothalidone, clonidine, codeine, diazepam, digoxin, dimethindene, diphenhydramine, diphenoxylate, dobutamine, doxycycline, ephedrine and pseudoephedrine, fludrocortisone, flurazepam, guanethidine, hydralazine, hydrochlorothiazide, hydroflumethiazide, indomethacin, insulin, isoprenaline, isosorbide dinitrate, L-Dopa, methenamine mandelate (mandelic acid), methyl dopa, methylprednisolone, nitrofurantoin, nitroglycerine, oxazepam, entazocine, phenacetin, phenobarbital, phenytoin, prednisone, probenecid, progesterone, propoxyphene, propranolol, quinidine, spironolactone, tetracycline, thyroxine, and tripelennamine.

On occasion, when interference cannot be resolved, an interference comment will be reported.

The variability associated with age, gender, and renal failure is uncertain.

**Clinical Reference**


Test Definition: CATU
Catecholamine Fract, Free, U

Performance

Method Description
A 1-mL aliquot of a 24-hour urine collection preserved in acid is absorbed on aluminum oxide at an alkaline pH and eluted with acid. An aliquot of the eluate is injected onto a high-performance reverse-phase paired ion-chromatography column where the catecholamines are resolved into individual components. The catechol is oxidized electronically to an O-quinone. The current generated at the detector is converted by amplifier to a voltage signal and an XY recording is generated. (Jiang NS, Machacek D: Measurement of catecholamines in blood and urine by liquid chromatography with amperometric detection. In Progress in HPLC. Vol 2. Edited by Parvez. VNU Science Press, 1987, pp 397-426; Moyer TP, Jiang NS, Tyce GM, Sheps SG: Analysis for urinary catecholamines by liquid chromatography with amperometric detection: methodology and clinical interpretation of results. Clin Chem 1979;25:256-263)

PDF Report
No

Day(s) and Time(s) Test Performed
Monday through Saturday; 8 a.m.

Analytic Time
2 days (not reported on Sundays)

Maximum Laboratory Time
4 days

Specimen Retention Time
14 days

Performing Laboratory Location
Rochester

Fees and 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 Regional Manager. 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. This test has not been cleared or approved by the U.S. Food and Drug Administration.

CPT Code Information
82384

LOINC® Information

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<td>Catecholamine Fract, Free, U</td>
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