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Instructions for use

3-CAT ELISA

Enzyme Immunoassay for the quantitative determination of
adrenaline (epinephrine), noradrenaline (norepinephrine) and
dopamine in plasma and urine.

For *in-vitro* diagnostic use.

REF

IB89550



IVD

1. Introduction

1.1 Intended use and principle of the test

Enzyme Immunoassay for the quantitative determination of adrenaline (epinephrine), noradrenaline (norepinephrine) and dopamine in plasma and urine. For *in-vitro* diagnostic use.

Adrenaline (epinephrine), noradrenaline (norepinephrine), and dopamine are extracted by using a cis-diol-specific affinity gel, acylated and then converted enzymatically.

The competitive ELISA kit uses the microtiter plate format. The antigen is bound to the solid phase of the microtiter plate. The derivatized standards, controls and samples and the solid phase bound analytes compete for a fixed number of antibody binding sites. After the system is in equilibrium, free antigen and free antigen-antibody complexes are removed by washing. The antibody bound to the solid phase is detected by an anti-rabbit IgG-peroxidase conjugate using TMB as a substrate. The reaction is monitored at 450 nm.

Quantification of unknown samples is achieved by comparing their absorbance with a standard curve prepared with known standard concentrations.

1.2 Clinical application

In humans the catecholamines adrenaline (epinephrine), noradrenaline (norepinephrine) and dopamine are neurotransmitters of the sympathetic nervous system and are involved in many physiological processes. The sympathetic nervous system sets the body to a heightened state of alert, also called as the body's fight-or-flight response.

In the human body the catecholamines and their metabolites indicate the adaption of the body to acute and chronic stress.

Next to the metanephrine/normetanephrine the catecholamines are important for the diagnosis and the follow-up of tumors of the sympathoadrenal system like the pheochromocytomas. The quantitative determination of catecholamines in urine is preferred for the diagnosis of these tumors, whereas the determination of catecholamines in plasma is medically sensible for the localization of the tumor and for function testing. Values above the cut-off can provide an indication for neuroendocrine tumors.

However, in literature various diseases like hypertension, cardiovascular diseases, schizophrenia and manic depression are described with abnormal low or high levels of catecholamines.

Therapeutic consequences should never be based on laboratory results alone even if all test results are in agreement with the items as under point "Procedural cautions, guidelines and warnings". Any laboratory result is only a part of the total clinical picture of the patient.

Only in cases where the laboratory results are in an acceptable agreement with the overall clinical picture of the patient it can be used for therapeutic consequences.

The test result itself should never be the sole determinant for deriving any therapeutic consequences.

2. Procedural cautions, guidelines, warnings and limitations

2.1 Procedural cautions, guidelines and warnings

- (1) This kit is intended for professional use only. Users should have a thorough understanding of this protocol for the successful use of this kit. Only the test instruction provided with the kit is valid and has to be used to run the assay. Reliable performance will only be attained by strict and careful adherence to the instructions provided.
- (2) This assay was validated for certain types of samples as indicated in *Intended Use* (please refer to Chapter 1). Any off-label use of this kit is in the responsibility of the user and the manufacturer cannot be held liable.
- (3) The principles of Good Laboratory Practice (GLP) have to be followed.
- (4) In order to reduce exposure to potentially harmful substances, wear lab coats, disposable protective gloves and protective glasses where necessary.
- (5) All kit reagents and specimens should be brought to room temperature and mixed gently but thoroughly before use. Avoid repeated freezing and thawing of reagents and specimens.
- (6) For dilution or reconstitution purposes, use deionized, distilled, or ultra-pure water.
- (7) The microplate contains snap-off strips. Unused wells must be stored at 2 °C to 8 °C in the sealed foil pouch with desiccant and used in the frame provided.
- (8) Duplicate determination of sample is highly recommended to be able to identify potential pipetting errors.
- (9) Once the test has been started, all steps should be completed without interruption. Make sure that the required reagents, materials and devices are prepared ready at the appropriate time.
- (10) Incubation times do influence the results. All wells should be handled in the same order and time intervals.
- (11) To avoid cross-contamination of reagents, use new disposable pipette tips for dispensing each reagent, sample, standard and control.
- (12) A standard curve must be established for each run.

- (13) The controls should be included in each run and fall within established confidence limits. The confidence limits are listed in the QC-Report provided with the kit.
- (14) Do not mix kit components with different lot numbers within a test and do not use reagents beyond expiry date as shown on the kit labels.
- (15) Avoid contact with Stop Solution containing 0.25 M H₂SO₄. It may cause skin irritation and burns. In case of contact with eyes or skin, rinse off immediately with water.
- (16) TMB substrate has an irritant effect on skin and mucosa. In case of possible contact, wash eyes with an abundant volume of water and skin with soap and abundant water. Wash contaminated objects before reusing them.
- (17) For information on hazardous substances included in the kit please refer to Material Safety Data Sheet (MSDS). The Material Safety Data Sheet for this product is made available directly on the website of the manufacturer or upon request.
- (18) The expected reference values reported in this test instruction are only indicative. It is recommended that each laboratory establishes its own reference intervals.
- (19) The results obtained with this test kit should not be taken as the sole reason for any therapeutic consequence but have to be correlated to other diagnostic tests and clinical observations.
- (20) Kit reagents must be regarded as hazardous waste and disposed according to national regulations.

2.2 Limitations

Any inappropriate handling of samples or modification of this test might influence the results.

2.2.1 Interfering substances

Plasma

Samples containing precipitates or fibrin strands or which are haemolytic or lipemic might cause inaccurate results.

24-hour urine

Please note the *sample preparation and storage!* If the percentage of the final concentration of acid is too high, the buffer capacity of the Extraction Buffer is insufficient. As a consequence catecholamines will not be extracted adequately.

2.2.2 Drug interferences

There are no known substances (drugs) which ingestion interferes with the measurement of catecholamine level in the sample.

2.2.3 High-Dose-Hook effect

No hook effect was observed in this test.

3. Storage and stability

Store the unopened reagents at 2 - 8 °C until expiration date. Do not use components beyond the expiry date indicated on the kit labels. Once opened the reagents are stable for 1 month when stored at 2 - 8 °C. Once the resealable pouch has been opened, care should be taken to close it tightly with desiccant again.

4. Materials

4.1 Content of the kit

BA D-0090 FOILS **Adhesive Foil** - Ready to use

Content: Adhesive Foils in a resealable pouch

Volume: 3 x 4 foils

BA E-0030 WASH-CONC 50x **Wash Buffer Concentrate** - Concentrated 50x

Content: Buffer with a non-ionic detergent and physiological pH

Volume: 3 x 20 ml/vial, light purple cap

BA E-0040 CONJUGATE **Enzyme Conjugate** - Ready to use

Content: Goat anti-rabbit immunoglobulins, conjugated with peroxidase

Volume: 3 x 12 ml/vial, red cap

BA E-0055 SUBSTRATE **Substrate** - Ready to use

Content: Chromogenic substrate containing tetramethylbenzidine, substrate buffer and hydrogen peroxide

Volume: 3 x 12 ml/vial, black cap

BA E-0080	STOP-SOLN	Stop Solution - Ready to use
Content:	0.25 M sulfuric acid	
Volume:	3 x 12 ml/vial, light grey cap	
BA E-0131	ADR MN	Adrenaline Microtiter Strips - Ready to use
Content:	1 x 96 well (12x8) antigen precoated microwell plate in a resealable blue pouch with desiccant, blue coloured	
BA E-0231	NAD NMN	Noradrenaline Microtiter Strips - Ready to use
Content:	1 x 96 well (12x8) antigen precoated microwell plate in a resealable yellow pouch with desiccant, yellow coloured	
BA E-0331	DOP	Dopamine Microtiter Strips - Ready to use
Content:	1 x 96 well (12x8) antigen precoated microwell plate in a resealable green pouch with desiccant, green coloured	
BA E-6110	ADR-AS	Adrenaline Antiserum - Ready to use
Content:	Rabbit anti-adrenaline antibody, blue coloured	
Volume:	1 x 6 ml/vial, blue cap	
BA E-6210	NAD-AS	Noradrenaline Antiserum - Ready to use
Content:	Rabbit anti-noradrenaline antibody, yellow coloured	
Volume:	1 x 6 ml/vial, yellow cap	
BA E-6310	DOP-AS	Dopamine Antiserum - Ready to use
Content:	Rabbit anti-dopamine antibody, green coloured	
Volume:	1 x 6 ml/vial, dark green cap	
BA R-0050	ADJUST-BUFF	Adjustment Buffer - Ready to use
Content:	TRIS buffer	
Volume:	2 x 4 ml/vial, green cap	
BA R-6611	ACYL-BUFF	Acylation Buffer - Ready to use
Content:	Buffer with light alkaline pH for the acylation	
Volume:	1 x 20 ml/vial, white cap	
BA R-6612	ACYL-REAG	Acylation Reagent - Ready to use
Content:	Acylation reagent in DMF and DMSO	
Volume:	1 x 3 ml/vial, light red cap	
Hazards identification:	 <p>H225 Highly flammable liquid and vapour. H360 May damage fertility or the unborn child. H319 Causes serious eye irritation.</p>	
BA R-6613	ASSAY-BUFF	Assay Buffer - Ready to use
Content:	1M hydrochloric acid and a non-mercury preservative	
Volume:	1 x 6 ml/vial, light grey cap	
BA R-6614	COENZYME	Coenzyme - Ready to use
Content:	S-adenosyl-L-methionine	
Volume:	1 x 4 ml/vial, purple cap	

- BA R-6615** ENZYME **Enzyme** - Lyophilized
 Content: Catechol-O-methyltransferase
 Volume: 6 vials, pink cap
- BA R-6617** EXTRACT-BUFF **Extraction Buffer** - Ready to use
 Content: Buffer containing carbonate
 Volume: 1 x 6 ml/vial, brown cap
- BA R-6618** EXTRACT-PLATE 48 **Extraction Plate** - Ready to use
 Content: 2 x 48 well plates coated with boronate affinity gel in a resealable pouch
- BA R-6619** HCL **Hydrochloric Acid** - Ready to use
 Content: 0.025 M Hydrochloric Acid, yellow coloured
 Volume: 1 x 20 ml/vial, dark green cap

Standards and Controls - Ready to use

Cat. no.	Component	Colour/ Cap	Concentration ng/ml			Concentration nmol/l			Volume/ Vial
			ADR	NAD	DOP	ADR	NAD	DOP	
BA R-6601	STANDARD A	white	0	0	0	0	0	0	4 ml
BA R-6602	STANDARD B	light yellow	1	5	10	5.5	30	65	4 ml
BA R-6603	STANDARD C	orange	4	20	40	22	118	261	4 ml
BA R-6604	STANDARD D	dark blue	15	75	150	82	443	980	4 ml
BA R-6605	STANDARD E	light grey	50	250	500	273	1 478	3 265	4 ml
BA R-6606	STANDARD F	black	200	1 000	2 000	1 092	5 910	13 060	4 ml
BA R-6609	STANDARD A/B	light purple	-	-	4.5	-	-	29	4 ml
BA R-6651	CONTROL 1	light green	Refer to QC-Report for expected value and acceptable range!						4 ml
BA R-6652	CONTROL 2	dark red	Refer to QC-Report for expected value and acceptable range!						4 ml

Conversion: Adrenaline (ng/ml) x 5.46 = Adrenaline (nmol/l)
 Noradrenaline (ng/ml) x 5.91 = Noradrenaline (nmol/l)
 Dopamine (ng/ml) x 6.53 = Dopamine (nmol/l)

Content: Acidic buffer with non-mercury stabilizer, spiked with defined quantity of adrenaline, noradrenaline, and dopamine

 **for the determination of dopamine in plasma the additional **Standard A/B** is mandatory!*

4.2 Additional materials and equipment required but not provided in the kit

- Calibrated precision pipettes to dispense volumes between 10 – 700 µl; 1 ml
- Microtiter plate washing device (manual, semi-automated or automated)
- ELISA reader capable of reading absorbance at 450 nm and if possible 620 - 650 nm
- Microtiter plate shaker (shaking amplitude 3 mm; approx. 600 rpm)
- Absorbent material (paper towel)
- Water (deionized, distilled, or ultra-pure)
- Vortex mixer

5. Sample collection and storage
Plasma

Whole blood should be collected into centrifuge tubes containing EDTA as anti-coagulant (Monovette™ or Vacuette™ for plasma) and centrifuged according to manufacturer's instructions immediately after collection.

Haemolytic and lipemic samples should not be used for the assay.

Storage: up to 6 hours at 2 - 8 °C, for longer period (up to 6 month) at -20 °C.

Repeated freezing and thawing should be avoided.

Urine

Spontaneous urine or 24-hour urine, collected in a bottle containing 10 - 15 ml of 6 M HCl, can be used.

If 24-hour urine is used please record the total volume of the collected urine.

Storage: up to 48 hours at 2 - 8 °C, up to 24 hours at room temperature, for longer periods (up to 6 months) at -20 °C. Repeated freezing and thawing should be avoided.

Avoid exposure to direct sunlight.

6. Test procedure

Allow all reagents to reach room temperature and mix thoroughly by gentle inversion before use. Duplicate determinations are recommended.

The binding of the antiserum and the enzyme conjugate and the activity of the enzyme are temperature dependent, and the absorbance may vary if a thermostat is not used. The higher the temperature, the higher the absorbance will be. Varying incubation times will have a similar influence on the absorbance. The optimal temperature during the Enzyme Immunoassay is between 20 - 25 °C.

6.1 Preparation of reagents

Wash Buffer

Dilute the 20 ml Wash Buffer Concentrate with water (deionized, distilled, or ultra-pure) to a final volume of 1000 ml.

Storage: 1 month at 2 - 8 °C

Enzyme Solution

Reconstitute the content of the vial labelled 'Enzyme' with 1 ml water (deionized, distilled, or ultra-pure) and mix thoroughly. Add 0.3 ml of Coenzyme followed by 0.7 ml of Adjustment Buffer. The total volume of the Enzyme Solution is 2.0 ml.

 *The Enzyme Solution has to be prepared freshly prior to the assay (not longer than 10 - 15 minutes in advance). Discard after use!*

6.2 Sample preparation, extraction and acylation

 **for the determination of dopamine in plasma the additional **Standard A/B** is mandatory!*

1.	Pipette 10 µl of standards, controls, urine samples and 300 µl of plasma samples into the respective wells of the Extraction Plate .								
2.	Add 250 µl of water (deionized, distilled, or ultra-pure) to the wells with standards, controls and urine samples .								
3.	Pipette 50 µl of Assay Buffer into all wells.								
4.	Pipette 50 µl of Extraction Buffer into all wells.								
5.	Cover plate with Adhesive Foil and incubate 30 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm).								
6.	Remove the foil. Empty plate and blot dry by tapping the inverted plate on absorbent material.								
7.	Pipette 1 ml of Wash Buffer into all wells. Incubate the plate for 5 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm). Empty plate and blot dry by tapping the inverted plate on absorbent material.								
8.	Pipette another 1 ml of Wash Buffer into all wells. Incubate the plate for 5 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm). Empty plate and blot dry by tapping the inverted plate on absorbent material.								
9.	Pipette 150 µl of Acylation Buffer into all wells.								
10.	Pipette 25 µl of Acylation Reagent into all wells.								
11.	Incubate 15 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm).								
12.	Empty plate and blot dry by tapping the inverted plate on absorbent material.								
13.	Pipette 1 ml of Wash Buffer into all wells. Incubate the plate for 10 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm). Empty plate and blot dry by tapping the inverted plate on absorbent material.								
14.	Pipette 175 µl of Hydrochloric Acid into all wells.								
15.	Cover plate with Adhesive Foil . Incubate 10 min at RT (20 - 25 °C) on a shaker (approx. 600 rpm).  Remove the foil and discard. Do not decant the supernatant thereafter! The following volumes of the supernatant are needed for the subsequent ELISA:								
	<table border="1"><tr><td>Adrenaline</td><td>100 µl</td><td>Noradrenaline</td><td>20 µl</td></tr><tr><td>Dopamine (standards + urine)</td><td>25 µl</td><td>Dopamine (plasma)</td><td>50 µl</td></tr></table>	Adrenaline	100 µl	Noradrenaline	20 µl	Dopamine (standards + urine)	25 µl	Dopamine (plasma)	50 µl
Adrenaline	100 µl	Noradrenaline	20 µl						
Dopamine (standards + urine)	25 µl	Dopamine (plasma)	50 µl						

6.3 Adrenaline ELISA

1.	Pipette 25 µl of the Enzyme Solution (refer to 6.1) into all wells of the Adrenaline Microtiter Strips .
2.	Pipette 100 µl of the extracted standards, controls and samples into the appropriate wells.
3.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
4.	Pipette 50 µl of the respective Adrenaline Antiserum into all wells and cover plate with Adhesive Foil .
5.	Incubate for 2 h at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
6.	Remove the foil. Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
7.	Pipette 100 µl of the Enzyme Conjugate into all wells.
8.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
9.	Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
10.	Pipette 100 µl of the Substrate into all wells and incubate for 25 ± 5 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm). ⚠ Avoid exposure to direct sunlight!
11.	Add 100 µl of the Stop Solution to each well and shake the microtiter plate to ensure a homogeneous distribution of the solution.
12.	Read the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to 450 nm (if available a reference wavelength between 620 nm and 650 nm is recommended).

6.4 Noradrenaline ELISA

1.	Pipette 25 µl of the Enzyme Solution (refer to 6.1) into all wells of the Noradrenaline Microtiter Strips .
2.	Pipette 20 µl of the extracted standards, controls and samples into the appropriate wells.
3.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
4.	Pipette 50 µl of the Noradrenaline Antiserum into all wells and cover plate with Adhesive Foil .
5.	Incubate for 2 h at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
6.	Remove the foil. Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
7.	Pipette 100 µl of the Enzyme Conjugate into all wells.
8.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
9.	Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
10.	Pipette 100 µl of the Substrate into all wells and incubate for 25 ± 5 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm). ⚠ Avoid exposure to direct sunlight!
11.	Add 100 µl of the Stop Solution to each well and shake the microtiter plate to ensure a homogeneous distribution of the solution.
12.	Read the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to 450 nm (if available a reference wavelength between 620 nm and 650 nm is recommended).

6.5 Dopamine ELISA

1.	Pipette 25 µl of the Enzyme Solution (refer to 6.1) into all wells of the Dopamine Microtiter Strips .
2.	Pipette 25 µl of the extracted standards, controls, urine samples and 50 µl of the extracted plasma samples into the respective tubes.
3.	Add 25 µl of Hydrochloric Acid to the standards, controls and urine samples .
4.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
5.	Pipette 50 µl of the Dopamine Antiserum into all wells and cover plate with Adhesive Foil .
6.	Incubate for 2 h at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
7.	Remove the foil. Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
8.	Pipette 100 µl of the Enzyme Conjugate into all wells.
9.	Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
10.	Discard or aspirate the content of the wells. Wash the plate 3 x by adding 300 µl of Wash Buffer , discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
11.	Pipette 100 µl of the Substrate into all wells and incubate for 25 ± 5 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm). ⚠ Avoid exposure to direct sunlight!
12.	Add 100 µl of the Stop Solution to each well and shake the microtiter plate to ensure a homogeneous distribution of the solution.
13.	Read the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to 450 nm (if available a reference wavelength between 620 nm and 650 nm is recommended).

7. Calculation of results

Measuring range		Adrenaline	Noradrenaline	Dopamine
	Urine		0.8 - 200 ng/ml	2.5 - 1000 ng/ml
Plasma		18 - 6667 pg/ml	93 - 33 333 pg/ml	75 - 33 333 pg/ml

The standard curves are obtained by plotting the absorbance readings (calculate the mean absorbance) of the standards (linear, y-axis) against the corresponding standard concentrations (logarithmic, x-axis). Use a non-linear regression for curve fitting (e.g. spline, 4- parameter, akima).

Urine samples and controls

The concentrations of the **urine samples** and the **Controls** can be read directly from the standard curve.

Calculate the 24 h excretion for each urine sample: $\mu\text{g}/24\text{h} = \mu\text{g}/\text{l} \times \text{l}/24\text{h}$

Plasma samples

The read **Adrenaline and Noradrenaline** concentrations of the **plasma samples** have to be **divided by 30**.

The read **Dopamine** concentrations of the **plasma samples** have to be **divided by 60**.

Conversion

Adrenaline (ng/ml) \times 5.46 = Adrenaline (nmol/l)

Noradrenaline (ng/ml) \times 5.91 = Noradrenaline (nmol/l)

Dopamine (ng/ml) \times 6.53 = Dopamine (nmol/l)

Expected reference values

It is strongly recommended that each laboratory should determine its own normal and abnormal values.

In a study conducted with apparently normal healthy adults, using the 3-CAT ELISA ^{Fast Track} the following values are observed:

	Adrenaline	Noradrenaline	Dopamine
Urine	< 20 µg/day (110 nmol/day)	< 90 µg/day (535 nmol/day)	< 600 µg/day (3 900 nmol/day)
Plasma	< 100 pg/ml	< 600 pg/ml	< 100 pg/ml

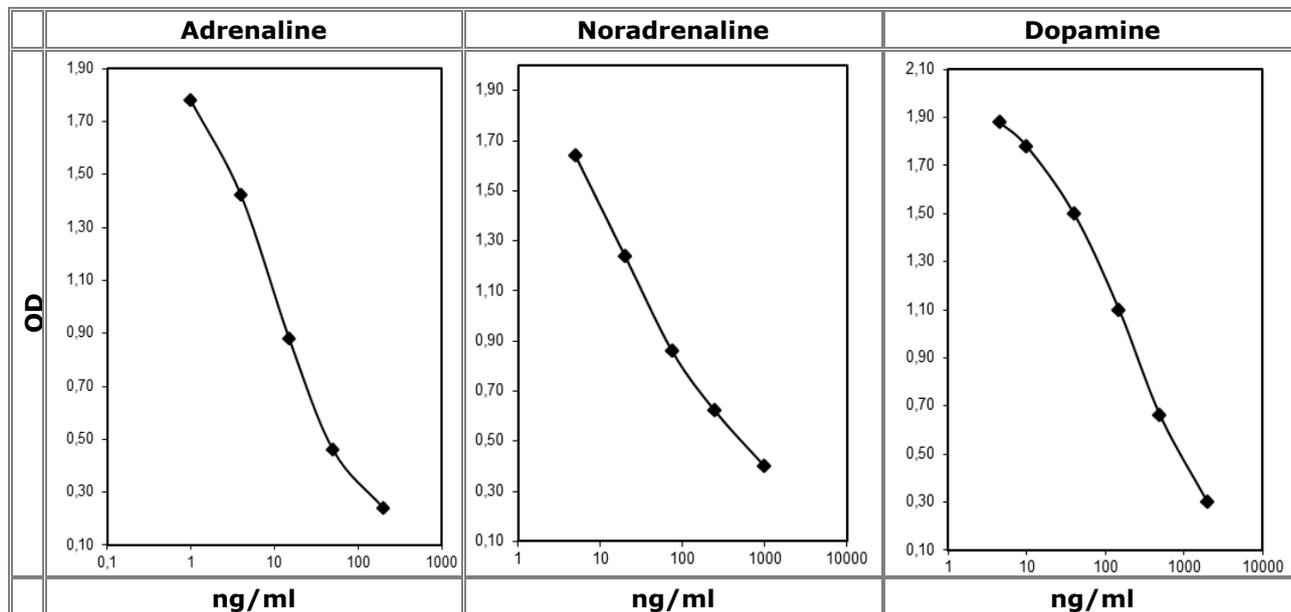
7.1 Quality control

It is recommended to use control samples according to national regulations. Use controls at both normal and pathological levels. The kit or other commercial controls should fall within established confidence limits. The confidence limits of the kit controls are printed on the QC-Report.

7.2 Typical standard curves



Examples, do not use for calculation!



8. Assay characteristics

Analytical Sensitivity	LOD		Adrenaline	Noradrenaline	Dopamine
		Urine (ng/ml)	0.9	1.7	2.5
	Plasma (pg/ml)	0.01	0.04	1.9	
	LOQ	Urine (ng/ml)	0.7	2.5	4.8
Plasma (pg/ml)	18.1	93	75		

Analytical Specificity (Cross Reactivity)	Substance	Cross Reactivity (%)		
		Adrenaline	Noradrenaline	Dopamine
	Derivatized Adrenaline	100	0.08	0.02
	Derivatized Noradrenaline	0.13	100	6.4
	Derivatized Dopamine	< 0.01	0.03	100
	Metanephrine	0.18	< 0.01	< 0.01
	Normetanephrine	< 0.01	0.16	0.01
	3-Methoxytyramine	< 0.01	< 0.01	0.49
	3-Methoxy-4-hydroxyphenylglycol	< 0.01	< 0.01	< 0.01
	Tyramine	< 0.01	< 0.01	0.18
	Phenylalanine, Caffeinic acid, L-Dopa, Homovanillic acid, Tyrosine, 3-Methoxy-4-hydroxymandelic acid	< 0.01	< 0.01	< 0.01

Precision							
Intra-Assay Urine (n = 60)				Intra-Assay Plasma (n = 60)			
	Sample	Range (ng/ml)	CV (%)		Sample	Range (pg/ml)	CV (%)
Adrenaline	1	6.2 ± 1.1	17.4	Adrenaline	1	64.7 ± 15.9	24.7
	2	21.4 ± 2.7	12.4		2	258 ± 32.5	12.7
	3	59.4 ± 7.8	13.1		3	948 ± 105	11.0
Noradrenaline	1	26.1 ± 3.6	13.8	Noradrenaline	1	510 ± 65	12.8
	2	97 ± 12.8	13.4		2	1358 ± 194	14.3
	3	267 ± 35	13.1		3	3363 ± 374	11.1
Dopamine	1	82 ± 16.1	19.7	Dopamine	1	75 ± 22	29.8
	2	253 ± 41.1	16.3		2	353 ± 86	24.4
	3	714 ± 67	9.4		3	1186 ± 271	22.5
Inter-Assay Urine (n = 33)				Inter-Assay Plasma (n = 18)			
	Sample	Range (ng/ml)	CV (%)		Sample	Range (pg/ml)	CV (%)
Adrenaline	1	5.2 ± 0.9	17.9	Adrenaline	1	76.4 ± 11.1	14.5
	2	17.8 ± 2.1	11.7		2	247 ± 27.5	11.1
	3	54.2 ± 6.6	12.1		3	771 ± 101	13.1
Noradrenaline	1	19.5 ± 3.9	20.0	Noradrenaline	1	445 ± 40.9	9.2
	2	80.6 ± 10.6	13.2		2	1232 ± 134	10.9
	3	226 ± 39.5	17.4		3	3283 ± 302	9.2
Dopamine	1	79.3 ± 18.8	23.7	Dopamine	1	238 ± 67.0	28.2
	2	222 ± 27.0	12.1		2	1072 ± 201	18.8
	3	630 ± 69.0	11.0		3	3449 ± 491	14.2

			Serial dilution up to	Range (%)	Mean (%)
	Linearity	Adrenaline	Urine	1:512	92 - 123
Plasma			1:512	94 - 115	105
Noradrenaline		Urine	1:512	100 - 127	112
		Plasma	1:512	102 - 125	112
Dopamine		Urine	1:512	83 - 126	104
		Plasma	1:512	85 - 132	106

			Mean (%)	Range (%)	Range
	Recovery	Adrenaline	Urine	106	94 - 120
Plasma			105	88 - 117	9.1 - 4268 pg/ml
Noradrenaline		Urine	103	91 - 113	58.6 - 230 ng/ml
		Plasma	87	75 - 107	51 - 16 735 pg/ml
Dopamine		Urine	110	89 - 121	252 - 1306 ng/ml
		Plasma	89	84 - 92	57.4 - 17 438 pg/ml

9. References/Literature

- (1) Dai et al. Association of plasma epinephrine level with insulin sensitivity in metabolically healthy but obese individuals. *Autonomic Neuroscience: Basic and Clinical*, 167:66-69 (2012)
- (2) Gruber et al. Increased Dopamine Is Associated With the cGMP and Homocysteine Pathway in Female Migraineurs. *Headache* 50:109-116 (2010)
- (3) Mobine et al. Pheochromocytoma-Induced Cardiomyopathy is Modulated by the Synergistic Effects of Cell-Secreted Factors. *Circulation: Heart Failure*, 2(1):121-128 (2009)

For orders, please contact:

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 **For updated literature or any other information please contact your local supplier.**

Symbols:

	Storage temperature		Manufacturer		Contains sufficient for <n> tests
	Expiry date		Batch code		For in-vitro diagnostic use only!
	Consult instructions for use		Content		CE labelled
	Caution		Catalogue number		For research use only!