

#### Lactate FS\*

# Diagnostic reagent for quantitative in vitro determination of lactate in plasma on DiaSys respons®910

#### **Order Information**

Cat. No. 1 4001 99 10 921

4 twin containers for 120 tests each

#### Method

Enzymatic UV test with lactate dehydrogenase (LDH)

#### **Principle**

L-Lactate + NAD<sup>+</sup> 

◆ LDH → Pyruvate + NADH + H<sup>+</sup>

In the presence of NAD lactate is converted by the lactate dehydrogenase. This procedure releases NADH which is measured at 340 nm. The absorbance of the produced NADH is proportional to the lactate concentration in the sample.

#### Reagents

#### **Components and Concentrations**

 R1:
 Buffer LDH
 pH 9.0
 500 mmol/L

 R2:
 NAD
 ≥ 25 kU/L
 ≥ 20 mmol/L

#### Storage Instructions and Reagent Stability

The reagents are stable up to the end of the indicated month of expiry, if stored at 2 - 8 °C, protected from light and contamination is avoided. DiaSys respons containers provide protection from light. Do not freeze the reagents!

#### **Warnings and Precautions**

- The reagents contain sodium azide (0.95 g/L) as preservative.
   Do not swallow! Avoid contact with skin and mucous membranes.
- In very rare cases, samples of patients with gammopathy might give falsified results.
- Please refer to the safety data sheets and take the necessary precautions for the use of laboratory reagents. For diagnostic purposes, the results should always be assessed with the patient's medical history, clinical examinations and other findings.

#### **Waste Management**

Please refer to local legal requirements.

#### Reagent Preparation

The reagents are ready to use. The bottles are placed directly into the reagent rotor.

#### Specimen

Plasma (no serum)

As anticoagulants use glycolytic inhibitors e.g. fluoride/oxalate or fluoride/heparin.

Stability in plasma [1]:

8 hours at 20 - 25 °C

14 days at 2 - 8 °C.

Discard contaminated specimens.

#### **Calibrators and Controls**

For calibration, DiaSys TruCal U calibrator is recommended. The assigned values of the calibrator are traceable to a primary standard. For internal quality control DiaSys TruLab N and P controls should be assayed. Each laboratory should establish corrective action in case of deviations in control recovery.

	Cat. No.	Kit	t size	
TruCal U	5 9100 99 10 063	20 x	3 mL	
	5 9100 99 10 064	6 x	3 mL	
TruLab N	5 9000 99 10 062	20 x	5 mL	
	5 9000 99 10 061	6 x	5 mL	
TruLab P	5 9050 99 10 062	20 x	5 mL	
	5 9050 99 10 061	6 x	5 mL	

#### **Performance Characteristics**

Measuring range up to 120 mg/dL lactate (13.3 mmol/L) (in case of higher concentrations re-measure samples after manual dilution or use rerun function).		
Limit of detection** 1 mg/dL lactate (0.1 mmol/L)		
On-board stability 1 week		
Calibration stability 1 week		

Interferences < 10%	Lactate [mg/dL]
up to 30 mg/dL	21.5
up to 1200 mg/dL	6.31
up to 1200 mg/dL	21.8
up to 65 mg/dL	6.86
up to 65 mg/dL	21.9
up to 70 mg/dL	6.03
up to 70 mg/dL	22.1
up to 1500 mg/dL	5.85
up to 1800 mg/dL	20.9
up to 10 mg/L	21.6
up to 20 mg/L	21.3
up to 10 mg/L	21.6
up to 1200 mg/L	21.3
	< 10%  up to 30 mg/dL  up to 1200 mg/dL  up to 1200 mg/dL  up to 65 mg/dL  up to 65 mg/dL  up to 70 mg/dL  up to 70 mg/dL  up to 1500 mg/dL  up to 1500 mg/dL  up to 1800 mg/dL  up to 10 mg/L  up to 10 mg/L  up to 10 mg/L

Precision			
Within run (n=20)	Sample 1	Sample 2	Sample 3
Mean [mg/dL]	5.60	12.9	24.0
Coefficient of variation [%]	2.92	1.69	1.65
Between run (n=20)	Sample 1	Sample 2	Sample 3
Mean [mg/dL]	7.33	13.0	29.6
Coefficient of variation [%]	2.62	2.93	1.51

Method comparison (n=108)		
Test x	DiaSys Lactate FS (Hitachi 917)	
Test y	DiaSys Lactate FS (respons®910)	
Slope	0.980	
Intercept	-0.560 mg/dL	
Coefficient of correlation	0.999	

<sup>\*\*</sup> according to NCCLS document EP17-A, vol. 24, no. 34

#### **Conversion factor**

Lactate [mg/dL] x 0.1109 = Lactate [mmol/L]

#### Reference Range [3]

Plasma:

Venous 4.5 - 19.8 mg/dL (0.5 - 2.2 mmol/L) Arterial 4.5 - 14.4 mg/dL (0.5 - 1.6 mmol/L)

Each laboratory should check if the reference ranges are transferable to its own patient population and determine own reference ranges if necessary.

#### Literature

- Westgard JO, Lahmeyer BL, Birnbaum ML. Use of the Du Pont "Automatic Clinical Analyzer" in Direct Determination of Lactic Acid in Plasma Stabilized with Sodium Fluoride. Clin Chem 1972; 18: 1334-8.
- Young DS. Effects of Drugs on Clinical Laboratory Tests. 5th. ed. Volume 1 and 2. Washington, DC: The American Association for Clinical Chemistry Press, 2000.
- Section I General Clinical Tests In: Tietz NW, editor. Clinical Guide to Laboratory Tests. 3rd ed. Philadelphia: Saunders; 1995. p. 382-3.
- David B. Sacks, M.B., Ch.B., F.A.C.P. Carbohydrates In: Burtis CA, Ashwood ER, editors. Tietz Textbook of Clinical Chemistry. 3rd ed. Philadelphia: W.B Saunders Company; 1999. p. 787-790.
- Thomas L. Clinical Laboratory Diagnostics. 1st ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998. p. 160-166.

#### Manufacturer

IVD (€

DiaSys Diagnostic Systems GmbH Alte Strasse 9 65558 Holzheim Germany

Reagent information \* fluid stable



### Lactate

## Application for serum and plasma samples

This application was set up and evaluated by DiaSys. It is based on the standard equipment at that time and does not apply to any equipment modifications undertaken by unqualified personnel

Identification	
This method is usable for analysis:	Yes
Name:	LACT
Shortcut:	
Reagent barcode reference:	043
Host reference:	

Technic	
Type:	Endpoint
First reagent:[µL]	180
Blanc correction	Yes
Second reagent:[µL]	45
Blanc correction	Yes
Main wavelength:[nm]	340
Secondary wavelength:[nm]	800
Polychromatic factor:	1.000
1 st reading time [min:sec]	(04:24)
Last reading time [min:sec]	10:00
Reaction way:	Increasing
Linear Kinetics	
Substrate deplation: absorbance limit	
Linearity: Maximum deviation [%]	
Fixed Time Kinetics	
Substrate deplation: absorbance limit	
Endpoint	x
Stability: largest remaining slope	
Prozone Limit [%]	

Sample Diluent	NaCl
Concentration technical limits-Lower	1
	120
SERUM	120
	3
	1
rterrial andien (raeter)	6
	1
zeren nermai anatien (ideter)	3
	6
URIN	0
	3
	1
Treatment (reserve)	6
	1
\ /	3
£1 2	6
PLASMA	
Normal volume [µL]	3
Normal dilution (factor)	1
	6
Below normal dilution (factor)	1
Above normal volume [µL]	3
Above normal dilution (factor)	6
CSF	
Normal volume [µL]	3
Normal dilution (factor)	1
	6
Below normal dilution (factor)	1
Above normal volume [µL]	3
Above normal dilution (factor)	6

Results	
Decimals	2
Units	mg/dL
Correlation factor-Offset	0.000
Correlation factor-Slope	1.000

Range	
Genre	Venous
Age	
SERUM	
URINE	
PLASMA	>=4.5 <=19.8
CSF	
Genre	Arterial
Age	
SERUM	
URINE	
PLASMA	>=4.5 <=14.4
CSF	

Contaminants	
Contaminant 1	
Wash with	
Cycle	
Volume [µL]	
Contaminant 2	
Wash with	
Cycle	
Volume [µL]	

Calibrators details		
Calibrator li	st	Concentration
Cal. 1		0
Cal. 2		*
Cal. 3		*
Cal. 4		*
Cal. 5		*
Cal. 6		*
	Max delta abs.	
Cal. 1	0.015	
Cal. 2	0.015	
Cal. 3		
Cal. 4		
Cal. 5		
Cal. 6		
Drift limit [%]	0.8	
Calculations		
Model		X degree
Degree	Degree 1	
* Enter calibrator value		

Application respons®910 March 2013/3