

## PYRUVATE (Quantitative Enzymatic UV-Test)

REF: 335 001      100 test

R1 : 2 x 50 ml  
R2 : 1 x 5 ml  
R3 : 1 x 5 ml  
St : 1 x 20 ml

### Intended Use

Spectrum Diagnostics liquizyme Pyruvate reagent is intended for the in-vitro quantitative, diagnostic determination of pyruvate in human blood.

### Method

Enzymatic UV - Test.

### Assay Principle

In the presence of an excess of NADH pyruvate is converted to lactate. The reduction of the absorbance =  $\Delta A$ , at 340 nm, due to the oxidation of NADH to NAD<sup>+</sup>, is a measure of the amount of pyruvate originally present :



### Reagents

<b>Standard (St.) Reagents:</b>	<b>4.0 mg/dl</b>
<b>R1 Buffer Reagent :</b> (Tris buffer, pH 7.20)	1.50 mol/L
<b>R2 Coenzyme</b> NADH	10.0 mmol/L
<b>R3 Start Reagent</b> LDH	1.50 kU/mL

**Additional Reagent** ( not provided with the kit )  
**Perchloric Acid 0.6m** for deproteinization

### Precautions and Warnings

Do not ingest or inhale. In case of contact with eyes or skin; rinse immediately with plenty of soap and water. In case of severe injuries; seek medical advice immediately.

### Reagent Preparation

Spectrum Pyruvate reagents are supplied ready-to-use.

### Reagent Storage and Stability

All reagents are stable until expiration date stated on label when protected from light stored refrigerated at 2 - 8 °C.










### Specimen Preparation

Pipet 2,0 mL of **freshly drawn blood** into a centrifugation tube containing 4 mL of cold **0.6 m perchloric acid**. Vortex for about 30 seconds.

Keep the blood precipitate mixture for about 5 min in the cold to assure complete protein precipitation. Centrifuge 10 min at approximately 1500 x g. The protein free supernatant is ready for use.

The **Standard solution** has to be diluted with perchloric acid, in the same ratio as the sample.

### SYMBOLS IN PRODUCT LABELLING

	Authorised Representative		Use by/Expiration Date
	For in-vitro diagnostic use		CAUTION. Consult instructions for use
	Batch Code/Lot number		Manufactured by
	Catalogue Number		
	Consult instructions for use		
	Temperature Limitation		

### System Parameters

Wavelength	340 nm ( 334 - 365 )
Optical path	1 cm
Assay type	End point
Temperature	30 °C or 37 °C
Zero adjustment	Against Water
Linearity	400 µg/dl (61.2 mmol/l)

### Procedure

	Sample	Standard
<b>R1</b> Buffer Reagent	1 ml	1 ml
Supernatant Sample	2 ml	.....
Diluted Standard	.....	2 ml
Mix and add		
<b>R2</b> Coenzyme	50 µl	50 µl
Mix and incubate for approximately 5 min , pour into cuvette, measure initial absorbance A1		
<b>R3</b> Start Reagent	50 µl	50 µl
mix, incubate for approximately 5 min and measure absorbance A2		

$$\Delta A = A2 - A1 \text{ For Sample and Standard}$$

### CALCULATION

with Factor :

$$\text{Pyruvate (mg/dL)} = \Delta A \times 6,37 \text{ ( at 340 nm )}$$

with Standard :

$$\text{Pyruvate (mg/dL)} = \frac{\Delta A_{\text{sample}}}{\Delta A_{\text{Standard}}} \times 4.0$$

### Expected values

0.3 – 0.7 mg/dL

**Note:** It is recommended for each laboratory to establish and maintain its own reference values. The given data are only an indication.

### QUALITY CONTROL

For quality control use adequate control materials, available from Spectrum Diagnostics.

**Waste Disposal**

This product is made to be used in professional laboratories.  
Please consult local regulations for a correct waste disposal.

**S56:** dispose of this material and its container at hazardous or special waste collection point.

**S57:** use appropriate container to avoid environmental contamination.

**S61:** avoid release in environment. refer to special instructions/safety data sheets.

*Spectrum Diagnostics does not interpret the results of a clinical laboratory procedure; interpretation of the results is considered the responsibility of qualified medical personnel. All indications of clinical significance are supported by literature references.*

**ORDERING INFORMATION**

CATALOG NO.	QUANTITY
335 001	100 Test

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