

## LIPASE SYSTEM PACK

(Methyl Resorufin Method)

B Auto 200, Unicorn 230, Unicorn 120 & Bonavera Chem 200 ,  
Beaconic chem 200, Beaconic B200, Beaconic analyzer 120,  
Bonavera chem 100 (Fully Auto Biochemistry Analyzer)



Code	Product Name	Pack Size
BA226	Lipase System Pack	2x20 + 2 x 5 ml
BA226A	Lipase System Pack	2x40 + 2x 10 ml
BA226B	Lipase System Pack	1x20 + 1 x 5 ml

### INTENDED USE

Quantitative determination of lipase in serum/plasma.

### CLINICAL SIGNIFICANCE

Lipases are enzymes which hydrolyze glycerol ester of long fatty acids. The enzyme and its cofactor colipase is produced in the pancreas, lipase being also secreted in small amounts by the salivary glands as well as by gastric, pulmonary and intestinal mucosa. Bile acids and colipase form micellar complexes with the lipids and bind lipase on the substrate / water interface. Determination of lipase is used for investigation of pancreatic disorders. In acute pancreatitis the lipase concentrations rise to 2-50 fold to upper reference limit within 4-8 hours after begin of abdominal pain peaking at 24 hours and decreasing within 8 to 14 days. Elevated lipase values can also be observed in chronic pancreatitis and obstruction of the pancreatic duct.

### PRINCIPLE

Enzymatic color test.

The colorimetric substrate 1,2-o-dilauryl-rac-glycero-3-glutaric acid-(6- methylresorufin)-ester is cleaved by pancreatic lipase and the resulting dicarboxylic acid ester is hydrolysed under the alkaline test condition to yield the chromophore methylresorufi. The kinetic of color formation at 580 nm is monitored and it is proportional to lipase activity in sample.

### REAGENT COMPOSITION

#### Reagent 1: Lipase Reagent 1

Bicine Buffer	>40 mmol/l
Colipase	>0.98 mg/l
Na-Deoxycholate	>1 mmol/l
Calcium Chloride	>8 mmol/l

#### Reagent 2: Lipase Reagent 2

Buffer	>8 mmol/l
Taurodeoxy-Cholate	>8 mmol/l

### REAGENT PREPARATION

Reagents are liquid, ready to use.

### STORAGE AND STABILITY

The unopened reagents are stable till the expiry date stated on the bottle and kit label when stored at +2-+8°C. Reagent R2 is a microemulsion. Therefore, a slight

apparent precipitation could occur, showing a light red deposit on the bottom of vial. It is a normal behaviour and it is recommended to resuspend solution before analysis with a mild shaking  
On board stability: Min 30 days if refrigerated (+8-+14°C) and not contaminated.

### SPECIMEN COLLECTION & HANDLING

Use serum, Plasma (heparin, EDTA).

It is recommended to follow NCCLS procedures (or similar standardized conditions).

### STABILITY

in serum / plasma: 7 days at +4- +8°C  
Discard contaminated specimens.

### CALIBRATION

Calibration with the Beacon Multicalibrator is recommended.

### QUALITY CONTROL

Each laboratory should establish its own Quality Control scheme and corrective actions if controls do not meet the acceptable tolerances.

### UNIT CONVERSION

U/l x 0.017 =  $\mu$ kat/l

### EXPECTED VALUES

Serum  
at +37°C = Up to 60 U/L (=1.0  $\mu$ kat/l)

### PERFORMANCE CHARACTERISTICS

Data contained within this section is representative of performance on Beacon System. Data obtained in your laboratory may differ from these values.

Limit of quantification : 3 U/L

Linearity : 300 U/L

Measuring range : 3-300U/L

### Precision

Intra-assay precision Within run (n=20)	Mean (U/L)	SD (U/L)	CV (%)
Sample 1	131	2.56	1.96
Sample 2	471	0.91	1.94

Inter-assay precision Run to run (n=20)	Mean (U/L)	SD (U/L)	CV (%)
Sample 1	66	0.90	1.36

### COMPARISON

A comparison between Beacon Lipase (y) and a commercially available test (x) using 20 samples gave following results:

$$y=1.0163x-1.2407 \text{ U/L } r=0.998$$

### INTERFERENCES

Following substances do not interfere:

Hemoglobin upto 4.5 g/l, bilirubin up to 40 mg/dl, triglycerides up to 1000 mg/dl.

### NOTE:

Reagents such as Triglycerides, Cholesterol, LDL, HDL, Albumin contain high concentration of detergent and hydrolysing enzymes, cross contamination from such reagent should be avoided.

### WARNING AND PRECAUTIONS

For in vitro diagnostic use. To be handled by entitled and professionally educated person.

Reagents 1 is not classified as dangerous. It contains less than 0.1% sodium azide, which is classified as very toxic and dangerous substance for environment.

Reagent 2 of the kit contains less than 5% propan-1-ol.

### WASTE MANAGEMENT

Please refer to local legal requirements.

### NOTES

Clinical diagnosis should not be made on findings of a single test result, but should integrate both clinical and laboratory data.

Parameter For B Auto 200, Unicorn 230, Unicorn 120 & Bonavera Chem 200, Beaconic chem 200, Beaconic B200, Beaconic analyzer 120, Bonavera chem 100 (Fully Auto Biochemistry Analyzer)

Test Name	LIPASE
Full Name	LIPASE
Pri Wave	578 nm
Sec Wave	-
Assay/point	Fixed time
Start	18
End	24
Decimal	2
Unit	U/L
Linearity Range Low	3 U/L
Linearity Range High	300 U/L
Sample Volume	4 µl
Reagent 1 (R1) Volume	200 µl
Reagent 2 (R2) Volume	50 µl
Substrate Depleted	-
Linearity	300 U/L
Out Of Linearity Range	-
Calibration Type	2 Point linear
Points	2
Blank Type	Reagent
Concentration Blank	0.00
Concentration Std	Refer calibrator value sheet

### REFERENCES

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3. Tietz N, Shuey DF. Lipase in serum - the elusive enzyme; an overview. Clin Chem 1993;39:746-56.
4. Lott J, Patel ST, Sawheney AK, Kazmierczak SC, Love JE. Assays of serum lipase analytical and clinical considerations. Clin chem 1986;32:1290-1302.
5. Leybold A, Junge W. Importance of colipase for the measurement of serum lipase activity. Adv Clin enzymol 1984;4:60-7.
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7. Garguori Y, Jiilen R, Bois A, Verger R, Sarda L. Studies on the detergent inhibition of pancreatic lipase activity. J of Lipid Research 1983;24:1336-42.
8. Guder WG, Zafta B et al. The quality of Diagnostic samples. 1st ed. Darmstadt: GIT Verlag; 2001.p.36-7.
9. 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.
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### Symbols Used On Labels



Catalogue Number



Manufacturer



See Instruction for Use



Lot Number



Content



Storage Temperature



Expiry Date



In Vitro Diagnostics

BEA/24/LIP/SB/IFU Ver-03  
09/09/2025

