

Lipase

Enzymatic, colorimetric

Diagnostic reagent for quantitative in vitro determination of lipase in human serum or plasma on photometric systems

REF	Kit Size	Configuration
D01441	5 x 50 mL	4 x 50 mL R1 + 1 x 50 mL R2
D01440	5 x 25 mL	4 x 25 mL R1 + 1 x 25 mL R2
D01443	5 x 10 mL	4 x 10 mL R1 + 1 x 10 mL R2
D44911	5 x 50 mL	4 x 50 mL R1 + 2 x 25 mL R2
D0433917	5 x 50 mL	4 x 50 mL R1 + 1 x 50 mL R2
DA0837	5 x 20 mL	4 x 20 mL R1 + 1 x 20 mL R2
DT1037	5 x 20 mL	4 x 20 mL R1 + 1 x 20 mL R2
DK0735	5 x 50 mL	4 x 50 mL R1 + 1 x 50 mL R2
DE1837	2 x 62.5 mL	2 x 50 mL R1 + 2 x 12.5 mL R2
DB20327	4 x 62.5 mL	4 x 50 mL R1 + 4 x 12.5 mL R2

Additionally available:

D98485	5 x 3 mL	Calibrator	Diacal Auto
D98485SV	1 x 3 mL	Calibrator	Diacal Auto
D98481	12 x 5 mL	Control normal	Diacon N
D14481	5 x 5 mL	Control normal	Diacon N
D98481SV	1 x 5 mL	Control normal	Diacon N
D98482	12 x 5 mL	Control abnormal	Diacon P
D14482	5 x 5 mL	Control abnormal	Diacon P
D98482SV	1 x 5 mL	Control abnormal	Diacon P

For professional in vitro diagnostic use only.

GENERAL INFORMATION

Method	Enzymatic colorimetric, kinetic, increasing reaction
Shelf life	18 months from date of production
Storage	2 – 8 °C
Wavelength	580 nm
Lightpath:	1 cm
Temperature	37 °C
Sample	Serum, heparinized plasma

INTENDED USE

Diagnostic reagent for quantitative in vitro determination of lipase in human serum or plasma on photometric systems.

DIAGNOSTIC SIGNIFICANCE [1, 2]

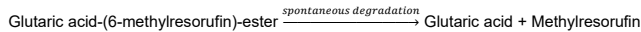
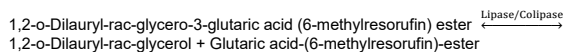
Lipases are enzymes which hydrolyze glycerol esters of long fatty acids. The enzyme and its cofactor colipase are produced in the pancreas. In small amounts, lipase is also secreted by the salivary glands as well as by gastric, pulmonary and intestinal mucosa. Bile acids and colipase form micellar complexes 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 the upper reference limit within 4 – 8 hours after the beginning of abdominal pain peaking at 24 hours and decrease within 8 to 14 days. Elevated lipase values can also be observed in chronic pancreatitis and obstruction of the pancreatic duct.

TEST PRINCIPLE

The colour substrate 1,2-o-dilauryl-rac-glycero-3-glutaric acid-(6-methylresorufin) ester is cleaved by pancreatic lipase in the presence of colipase and bile acids, and the resulting dicarboxylic acid ester is hydrolysed under alkaline test conditions to yield the chromophore methylresorufin.

The kinetic of colour formation at 580 nm is monitored and it is proportional to lipase activity in the sample.



REAGENT COMPOSITION

COMPONENTS	CONCENTRATION
Reagent 1	
Good's Buffer	pH 8.0
Colipase	≥ 2 mg/L
Desoxycholate	≥ 1.0 mmol/L
Taurodesoxycholate	≥ 1.0 mmol/L
Calcium ions	≥ 1.0 mmol/L
Detergent	
Preservative	
Reagent 2	
Tartrate Buffer	pH 4.0
Lipase Substrate	≥ 1.0 mmol/L
Stabilizer	
Preservative	

MATERIAL REQUIRED BUT NOT PROVIDED

- NaCl solution (9 g/L).
- Clinical chemistry analyser.

REAGENT PREPARATION

Reagents are ready to use.
 Avoid strong shaking!

STORAGE AND STABILITY

Conditions: Protect from direct light.
 Close immediately after use.
 Avoid contamination.
 Do not freeze the reagents.

Storage: at 2 – 8 °C
 Stability: up to the expiration date
 After first opening use preferably within 60 days 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. This is normal. It is recommended to resuspend solution before analysis, with a mild shaking.

WARNINGS AND PRECAUTIONS

1. Reagent 2: Danger.



H318: Causes serious eye damage.
 P280: Wear protective gloves/protective clothing/eye protection.
 P305+P351+P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
 P310: Immediately call a doctor.

2. Reagent 1 contains sodium azide (0.95 g/L) as preservative. Do not swallow! Avoid contact with skin and mucous membranes.
3. Many other clinical reagents contain lipase or high concentrations of detergents. Avoid contamination and carry over!
4. Special care should be taken in combination with triglycerides, HDL and LDL reagents containing microbial lipases that could stick on the surface of instrument cuvettes. Cuvettes and other glassware must be cleaned thoroughly after being used for other assays. In case of automated measurement refer to the instrument manual for special washing programs before lipase determination.
5. Please refer to the safety data sheets and take the necessary precautions for the use of laboratory reagents.
6. For diagnostic purposes, the results should always be assessed with the patient's medical history, clinical examinations and other findings.
7. For professional use only!

SPECIMEN COLLECTION AND STORAGE

Serum, heparinized plasma.

Stability [9]:
 In serum/plasma at 2 - 8 °C 7 days
 Discard contaminated specimens.

TEST PROCEDURE

Bring reagents and samples to room temperature.

Pipette into test tubes	Blank	Calibrator	Sample
Reagent 1	1000 µL	1000 µL	1000 µL
Sample	-		20 µL
Calibrator	-	20 µL	-
Dist. water	20 µL	-	-
Mix carefully (do not shake!), incubate 5 min. at 37 °C. Then add:			
Reagent 2	250 µl	250 µl	250 µl
Mix. Incubate 2 min. at 37°C, read absorbance against Reagent blank and start stop watch. Read absorbance again after exactly 1 and 2 minutes. Calculate: $\Delta A/\text{min} = [\Delta A/\text{min sample or calibrator}] - [\Delta A/\text{min blank}]$			

Automation

Special adaptations for automated analysers can be made on request.

INTERPRETATION OF RESULTS

Calculation

Serum/Plasma:

$$\text{Lipase [U/L]} = \frac{\Delta A/\text{min Sample}}{\Delta A/\text{min Calibrator}} \times \text{Conc. Calibrator [U/L]}$$

Unit Conversion

$$\text{Lipase [U/L]} \times 0.01667 = \text{Lipase [\mu\text{kat/L}]}$$

QUALITY CONTROL AND CALIBRATION

All control sera with Lipase values determined by this method can be used. We recommend the Dialab serum controls **Diacon N** (control serum with values in the normal range) and **Diacon P** (control serum with values in the abnormal range). Each laboratory should establish corrective action in case of deviations in control recovery.

Calibration

The assay requires the use of a Lipase Standard or Calibrator.
 We recommend the Dialab multi calibration serum **Diacal Auto**.

PERFORMANCE CHARACTERISTICS

LINEARITY, MEASURING RANGE

The assay is linear up to 300 U/L. If this value is exceeded, samples should be diluted 1 + 1 with NaCl solution (9 g/L) and the result multiplied by 2.

SENSITIVITY/LIMIT OF DETECTION

The limit detection is 1 U/L.

PRECISION (at 37 °C)

Intra-assay n = 10	Mean [U/L]	SD [U/L]	CV [%]
Sample 1	49.9	0.65	1.30
Sample 2	110.5	1.69	1.53

Inter-assay n = 20	Mean [U/L]	SD [U/L]	CV [%]
Sample 1	50.0	1.43	2.87
Sample 2	110.9	3.91	3.53

SPECIFICITY/INTERFERENCES

No interference was observed by the presence of:

- Ascorbic acid ≤ 50 mg/dL
- Hemoglobin ≤ 400 mg/dL
- Bilirubin ≤ 50 mg/dL
- Triglycerides ≤ 1000 mg/dL

For further information on interfering substances refer to Young DS [10].

METHOD COMPARISON

A comparison between Dialab Lipase (y) and a commercially available colorimetric test (x) using 94 samples gave following results:
 $y = 0.93 x + 5.50 \text{ U/L}$; $r^2 = 0.99$.

TRACEABILITY

The assigned values of Lipase in the calibrator Dialcal Auto have been made traceable to the molar extinction coefficient ϵ according to an available measurement procedure.

EXPECTED VALUES

Normal subjects [8]*	≤ 60 U/L (≤ 1.00 $\mu\text{kat/L}$)
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* Each laboratory should check if the reference ranges are transferable to its own patient population and determine own reference ranges if necessary.

LIMITATIONS

- Eventual Lipase (Enzymatic, colorimetric) carry-over to reagents Calcium (Arsenazo), Calcium (CPC), Magnesium (Xylidyl blue) and Triglycerides (GPO-PAP). The actual carry-over depends on the analyser.

WASTE MANAGEMENT

Please refer to local legal requirements.

LITERATURE

1. Lorentz K. Lipase. In: Thomas L, editor. Clinical laboratory diagnostics. 1st ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998, p. 95-7.
2. Moss DW, Henderson AR. Digestive enzymes of pancreatic origin. In: Burtis CA, Ashwood ER, editors. Tietz Textbook of Clinical Chemistry. 3rd ed. Philadelphia: W.B Saunders Company; 1999. p. 689-708.
3. Tietz N, Shuey DF. Lipase in serum – the elusive enzyme: an overview. Clin Chem 1993;39:746-56.
4. Lott J, Patel ST, Sawhney 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 1986;4:60-7.
6. Borgström B. The action of bile salts and other detergents on pancreatic lipase and the interaction with colipase. Biochimica et Biophysica Acta 1977;488:381-91.
7. Gargouri Y, Julien 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. Junge W, Abicht K, Goldman J. Evaluation of the colorimetric liquid assay for pancreatic lipase on Hitachi analyzers in 7 clinical centres in Europe. Clin Chem Lab Med 1999; 37, Special suppl: 469.
9. Rifai N., Horvath A.R., Wittwer C.T. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics - sixth edition ed. 2017 p. 421-424.
10. 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.

