

The Diasystem Vet Total Bile Acids diagnostic reagent for quantitative *in vitro* determination of total bile acids in serum or plasma on photometric systems.

Ordering information

Referens	Cat. Nr	Kit size
Diasystem Vet Total Bile Acids For use with Diasystem VET240	CDT-TBA	R1: 4 x 30mL R2: 4 x 10mL
Total Bile Acids For use with Thermo Fischer Indiko	1010	R1: 6 x 20mL R2: 4 x 10mL

Indication for Use

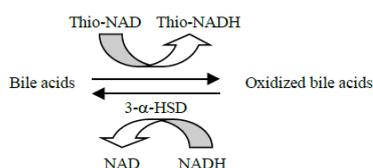
Diasystem Vet Total Bile Acids is intended for the *in vitro* quantitative determination of total bile acids (TBA) in human serum samples. Total bile acids are metabolized in the liver and serve as a marker for normal and abnormal liver function. Serum total bile acids are increased in patients with liver disease. [1,2,3].

Clinical Significance ^{1,2}

Total bile acids are metabolized in the liver and, hence, serve as a marker for normal liver function. Serum total bile acids are increased in patients with acute hepatitis, chronic hepatitis, liver sclerosis and liver cancer.

Assay Principle

The reagents of the assay kit are in a stable liquid formulation that allows for ease of use coupled with enhanced performance characteristics. In the presence of Thio-NAD, the enzyme 3- α -hydroxysteroid dehydrogenase (3- α -HSD) converts bile acids to 3-keto steroids and Thio-NADH. The reaction is reversible and 3- α -HSD can convert 3-keto steroids and Thio-NADH to bile acids and Thio-NAD. In the presence of excess NADH, the enzyme cycling occurs efficiently and the rate of formation of Thio-NADH is determined by measuring specific change of absorbance at 405 nm.



Materials Required But Not Provided

An analyzer capable of dispensing two reagents and of measuring absorbance at 405 nm with temperature control (37°C).

Calibrator for the Diasystem Vet Total Bile Acids assay (Ref 1011).

0.9% Saline is used for diluting samples and as Calibrator 0.

Controls for validating the performance of the Diasystem Vet Total Bile Acids assay (Ref DZ042A-CON).

Reagents

Components

R1: Thio-Nad >0.1nM, Buffer

R2: 3- α -HSD->2KU/L, NADH >0.1mM Buffer.

Calibrator: Conjugated cholic acids, Buffer

Reagent Preparation

Diasystem Vet Total Bile Acids Assay Reagents are ready-to-use, liquid reagents. The intrinsic yellow to yellow-brown color of the TBA reagent does not interfere with the test.

Reagent Stability and Storage

Unopened reagents are stable until the expiration date printed on the label. Reagents are light sensitive and should be stored at 2-8°C. Reagents from different lots must not be interchanged.

Specimen Collection and Handling ⁴

Use fresh patient serum samples. TBA concentration is increased after meals; hence, samples should be collected under fasting conditions*. EDTA treated plasma or Lithium heparin plasma samples are suitable for use. Serum or plasma samples are stable for a week at 4 °C, or for 3 months at -20 °C. It is the responsibility of the individual laboratory to use all available references and/or its own studies to determine specific stability criteria for its laboratory.⁵

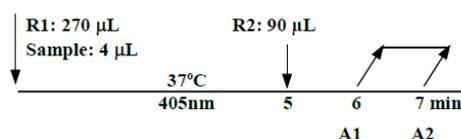
Specimens from patients, who are on Ursodeoxycholic Acid (UDCA) treatment, are not suitable for use with TBA Assay (CDT-TBA and 1010).

*This does not apply to women with intrahepatic cholestasis of pregnancy who will need peak bile acid testing and samples should therefore be taken post-prandially.

Precautions

1. This test is for *in vitro* use only and must be handled by qualified personnel.
2. Specimens and reagents containing human sourced materials should be handled as if potentially infectious, using safe laboratory procedures such as those outlined in Biosafety in Microbiological and Biomedical Laboratories (HHS Publication Number [CDC] 93-8395).
3. As with any diagnostic test procedure, results should be interpreted considering all other test results and the clinical status of the patient.
4. Avoid swallowing and contact with skin or mucous membranes.
5. Additional safety information concerning storage and handling of this product is provided within the Safety Data Sheet for this product. To obtain an SDS, please contact our customer service department at +46 36 58 59 000 or visit www.diasystemvet.se.

Assay Scheme for Chemistry Analyzers



Application sheets for use of Diasystem Vet Total Bile Acids Enzymatic Cycling Assay on automated clinical chemistry analyzers are available upon request. Please call +46 36 58 59 00 or email: service@diasystemvet.se.

Calibration

A bile acids calibrator (REF 1011), along with 0.9% saline as a zero reference, should be used as directed to calibrate the procedure. Calibration frequency may vary and is dependent

on instrument application. For more information, please call +46 36 58 59 000 or email: service@diasystemvet.se.

Calibration

We recommend that each laboratory use bile acid controls to validate the performance of bile acid reagents. A set of normal and abnormal range bile acid controls is available from Diazyme Laboratories (DZ042A-CON). If the results from the controls fall outside the acceptable limits, as determined by the manufacturer, the test should not be performed. We recommend that your quality control testing follows federal, state, and local guidelines or accreditation requirements.

Results

Bile acids concentration is expressed as $\mu\text{mol/L}$ ($\mu\text{Eq/L}$).

Reference Range ³

Serum or plasma containing 0-10 $\mu\text{mol/L}$ bile acids is considered normal range. We suggest that each laboratory establish a range of normal values for the population in their region.

Limitations

- A sample with a bile acids level exceeding the linearity limit should be diluted with 0.9% saline and reassayed incorporating the dilution factor in the calculation of the value.
- Specimens from patients, who are on Ursodeoxycholic Acid (UDCA) treatment, are not suitable for use with TBA Assay (DZ042A).

Performance Characteristics (Hitachi 717)

These performance characteristics were determined using automated procedures on a Hitachi 717.

Accuracy

The performance of this assay was compared with the performance of a similar total bile acids assay on a Hitachi 717 analyzer using serum samples.

Fifty-two (52) serum samples ranging from 0.47 – 131.25 $\mu\text{mol/L}$ gave a correlation coefficient of 0.9918. Linear regression analysis gave the following equation:
This method = 1.1536 (reference method) – 0.8567 $\mu\text{mol/L}$

A matched set of serum and lithium heparin plasma samples ranging from 0.14 – 21.18 $\mu\text{mol/L}$ gave a correlation coefficient of 0.9805. Linear regression analysis gave the following equation:

Lithium heparin = 0.9972 (serum) + 0.1178 $\mu\text{mol/L}$ information.

Precision Studies

The intra-assay precision and inter-assay precision were evaluated in samples containing two different bile acid levels (8 μM and 23 μM). The inter-assay precision was evaluated by testing these two-level specimens (low = 8 μM and high = 23 μM) in 20 runs. All tests were done using the Hitachi 717 Auto-analyzer instrument. Precision data is summarized in the table below:

Intra-Assay Precision

	Level 1 (8 μM)	Level 2 (23 μM)
Number of Replicates	20	20
Mean	7,93	23,5
SD	0,31	0,3
CV%	3,9%	1,3%

Inter-Assay Precision

	Level 1 (8 μM)	Level 2 (23 μM)
Number of Replicates	20	20
Mean	8,12	23,0
SD	0,24	0,61
CV%	2,9%	2,6%

Linearity

Linearity studies using a Hitachi 717 analyzer showed that Diasystem Vet Total Bile Acids assay has a linear range from 1 to 180 μM .

Interference

Interference for the Diasystem Vet Total Bile Acids Assay was evaluated on a Hitachi 717 analyzer. The following substances normally pre-sent in serum produced less than 10% deviation at the listed concentrations: Triglycerides at 750 mg/dL, Ascorbic acid at 50 mg/dL, Bilirubin at 50 mg/dL and Hemoglobin at 500 mg/dL.

References

1. LaRusso, N.F. et al., Dynamics of Enterohepatic Circulation of Bile Acids, *New Engl J M*, 291, 689-692, (1974).
2. Skrede S. et al: Bile acids measured in serum during fasting as a test for liver disease, *Clin Chem* 24: 1095-1099, 1978
3. Wu, Alan H. B. *Tietz Clinical Guide to Laboratory Tests*. 4th ed. St. Louis, MO: Saunders/Elsevier, 2006. 170-171.
4. Ovadia C, Seed P, Sklavounos A, et al. "Association of adverse perinatal outcomes of intrahepatic cholestasis of pregnancy with biochemical markers: results of aggregate and individual patient data meta-analyses." *The Lancet*, Elsevier Inc., 14 Feb 2019, dx.doi.org/10.1016/S0140-6736(18)31877-4.
5. CLSI, *Procedures for the Handling and Processing of Blood Specimens for Common Laboratory Tests: Approved Guideline*, H18-A4, Vol.30 No. 10.

Manufacturer

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