

Product information



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TSH canine ELISA

VET

REF

DEV9955



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1 INTRODUCTION

1.1 INTENDED USE

The Demeditec TSH canine ELISA is an enzyme immunoassay for the quantitative measurement of canine TSH (thyrotropin) in serum and EDTA-plasma.

For manual processing! The usage of laboratory automats is the user's sole responsibility.

The kit is intended for single use only.

1.2 SUMMARY AND EXPLANATION

Thyroid stimulating hormone (TSH, thyrotropin) in dogs is similar in function to TSH found in other mammalian species, including humans. It is a glycoprotein produced by the anterior pituitary gland. Through its action on the thyroid gland, it plays a major role in maintaining normal circulating levels of the iodothyronines, T4 and T3. The production and secretion of TSH is controlled by negative feedback from circulating T4 and T3, and by the hypothalamic hormone TRH (thyrotropin releasing hormone). The TSH molecule is composed of two nonidentical subunits, α and β , that are bound together in a noncovalent manner. Within a species, the TSH α subunit is structurally identical to the α subunits of the related glycoprotein hormones (LH, FSH and chorionic gonadotropin). The β subunit of TSH and the β subunits of the related hormones are structurally hormone-specific, and account for their unique biological activities.

Hypothyroidism is considered to be a common endocrine disorder in dogs, whereas hyperthyroidism in this species is nearly unknown. Dogs mostly suffer from primary hypothyroidism, involving impaired production of the thyroid hormones, T4 and T3. In this condition, elevated TSH levels are expected. Secondary or tertiary hypothyroidism, where thyroid hormone production is low as a consequence of hypothalamic or pituitary disease, is believed to account for less than 5% of canine hypothyroidism cases. In the latter conditions, decreased levels of TSH would be expected. Usually, hypothyroidism in dogs is suspected on the basis of clinical history and the presence of decreased levels of thyroid hormones. However, suppressed thyroid hormone levels are nonspecific indicators of the disease, since they are often observed in nonthyroid illnesses. The evaluation of thyroid function and the diagnosis of hypothyroidism in dogs can be greatly improved through the use of the valid assay for the determination of canine TSH.

2 PRINCIPLE

The test kit is a solid phase enzyme immunometric assay (ELISA) in the microplate format, designed for the quantitative measurement of canine TSH. The microplate is coated with an antibody specific for canine TSH.

Calibrators and samples are pipetted into the antibody coated microplate. Afterwards, a horseradish peroxidase-labeled antibody is added. During a two hour incubation sandwich complexes consisting of the two antibodies and the canine TSH is formed. Non-reactive components are removed by a washing step.

A chromogenic substrate, TMB (3,3',5,5'-Tetra-Methyl-Benzidine), is added to all wells. During a 30 minutes incubation, the substrate is converted to a colored end product (blue) by the bound enzyme. Enzyme reaction is stopped by dispensing hydrochloric acid as stop solution (change from blue to yellow). The color intensity is direct proportional to the concentration of canine TSH present in the sample. The optical density of the color solution is measured with a microplate reader at 450 nm.

3 WARNINGS AND PRECAUTIONS

1. This kit is strictly intended for veterinary use only. Use by staff, who is specially informed and trained in methods which are carried out by use of immunoassays.
2. All blood components and biological materials should be handled as potentially hazardous in use and for disposal. Follow universal precautions when handling and disposing of infectious agents.
3. Before starting the assay, read the instructions completely and carefully. Use the valid version of the package insert provided with the kit. Be sure that everything is understood.
4. The microplate contains snap-off strips. Unused wells must be stored at 2-8°C in the sealed foil pouch and used in the frame provided.
5. Pipetting of samples and reagents must be done as quickly as possible and in the same sequence for each step.
6. Use reservoirs only for single reagents. This especially applies to the substrate reservoirs. Using a reservoir for dispensing a substrate solution that had previously been used for the conjugate solution may result in colored solution. Do not pour reagents back into vials as reagent contamination may occur.
7. Mix the contents of the microplate wells thoroughly to ensure good test results. Do not reuse microwells.
8. Do not let wells dry during assay; add reagents immediately after completing the rinsing steps.
9. Allow the reagents to reach room temperature (18-25°C) before starting the test. Temperature will affect the absorbance readings of the assay.
10. Never pipet by mouth and avoid contact of reagents and samples with skin and mucous membranes.
11. Do not smoke, eat, drink or apply cosmetics in areas where samples or kit reagents are handled.
12. Wear disposable gloves when handling specimens and reagents. Microbial contamination of reagents or specimens may give false results.
13. Handling should be done in accordance with the procedures defined by an appropriate national biohazard safety guideline or regulation.
14. Do not use reagents beyond expiry date as shown on the kit labels.
15. All indicated volumes have to be performed according to the protocol. Optimal test results are only obtained when using calibrated pipettes and microtiterplate readers.
16. Do not mix or use components from kits with different lot numbers. It is advised not to exchange wells of different plates even of the same lot. The kits may have been shipped or stored under different conditions and the binding characteristics of the plates may vary slightly.
17. Avoid contact with Stop Solution. It may cause skin irritation and burns.
18. Chemicals and prepared or used reagents have to be treated as hazardous waste according to the national biohazard safety guideline or regulation.
19. For information please refer to Material Safety Data Sheets. Safety Data Sheets for this product are available upon request directly from Demeditec Diagnostics GmbH or on Demeditec homepage (www.demeditec.com).

4 REAGENTS

4.1 REAGENTS PROVIDED

1. **SORB MT Microtiter Plate**, 12 x 8 (break apart) strips with 96 wells, ready to use; wells coated with an anti-canine TSH antibody.
2. **CAL 0 – 5 LYO Calibrators**, 6 vials, lyophilized, reconstitution required; highly purified canine TSH in serum. The concentrations are 0, 0.2, 0.46, 1.05, 2.2 and 5.2 ng/ml. **For reconstitution see “Reagent preparation”.**
3. **ENZ CONJ Enzyme Conjugate**, 1 vial, 11 ml, red, ready to use; contains a horseradish peroxidase-labeled polyclonal anti canine TSH antibody, in a buffered matrix.
4. **SUB TMB TMB-Substrate Solution**, 1 vial, 22 ml, ready to use; contains tetramethylbenzidine (TMB) and hydrogen peroxide in a buffered matrix.
5. **STOP SOLN Stop Solution**, 1 vial, 7 ml, ready to use; contains 2 N Hydrochloric acid solution. Avoid contact with the stop solution. It may cause skin irritations and burns.
6. **WASH SOLN 10x Wash Solution**, 1 vial, 50 ml (10X concentrated); **see „Reagent preparation“.**

4.2 MATERIALS REQUIRED BUT NOT PROVIDED

- Microplate reader capable for endpoint measurements at 450 nm
- Vortex mixer
- Microplate mixer operating 900 rpm
- Distilled water
- Graduated cylinders for 500 ml
- Plastic containers for storage of the wash solution
- Adjustable pipette for up to 1000 µl
- Dispenser or repeatable pipet for 100 µl and 200 µl

4.3 REAGENT PREPARATION

Wash Solution:

Dilute 50 ml of 10x concentrated Wash Solution with 450 ml deionized water to a final volume of 500 ml. The diluted Wash Solution is stable for at least 12 weeks at room temperature (18-25°C). Precipitates may form when stored at 2-8°C, which should dissolve again by swirling at room temperature (18-25°C). The wash solution should only be used when the precipitates have completely dissolved.

Calibrators:

Reconstitute lyophilized Calibrators 0 through 5 with **1.0 ml dist. water** 30 minutes before use.

4.4 STORAGE CONDITIONS

When stored at 2-8°C unopened reagents will be stable until expiration date. Do not use reagents beyond this date. Opened reagents must be stored at 2-8°C. After first opening the reagents are stable for 30 days if used and stored properly. Microtiter wells must be stored at 2-8°C. Take care that the foil bag is sealed tightly.

Store Calibrators refrigerated, they will be stable at 2-8°C for 7 days after reconstitution. For longer storage aliquot and freeze at -20°C.

Protect TMB-Substrate Solution from light.

4.5 DISPOSAL OF THE KITS

The disposal of the kit must be made according to the national regulations. Special information for this product is given in the Material Safety Data Sheet.

4.6 DAMAGED TEST KITS

In case of any severe damage of the test kit or components, Demeditec have to be informed written, latest one week after receiving the kit. Severely damaged single components should not be used for a test run. They have to be stored until a final solution has been found. After this, they should be disposed according to the official regulations.

5 SAMPLE COLLECTION AND PREPARATION

For determination of canine TSH serum and EDTA-plasma are the preferred sample matrices. The procedure calls for 100 µl sample per well. The samples may be stored refrigerated at 2-8°C for one week, or up to two months at -20°C. To avoid repeated thawing and freezing the samples should be aliquoted.

Samples expected to contain canine TSH concentrations higher than the highest calibrator 5 (CAL 5) should be diluted in the Canine TSH Zero Calibrator (CAL 0) before assay. The additional dilution step has to be taken into account for the calculation of the results.

6 ASSAY PROCEDURE

6.1 GENERAL REMARKS

- All reagents and samples must be allowed to come to room temperature (18-25°C) before use. All reagents must be mixed without foaming.
- Once the test has been started, all steps should be completed without interruption.
- Use new disposal plastic pipette tips for each standard, control or sample in order to avoid cross contamination.
- Absorbance is a function of the incubation time and temperature. Before starting the assay, it is recommended that all reagents are ready, caps removed, all needed wells secured in holder, etc. This will ensure equal elapsed time for each pipetting step without interruption.
- As a general rule the enzymatic reaction is linearly proportional to time and temperature.
- Respect the incubation times as stated in this instructions for use.
- Duplicate determination of calibrators, controls and samples is recommended in order to identify potential pipetting errors.
- A calibrator curve must be established for every run.
- For internal quality control we suggest to use **Canine Control coded DEV99CC**. For more information please contact Demeditec.

6.2 ASSAY PROCEDURE

1. Prepare a sufficient number of microplate wells to accommodate calibrators (0 through 5) and patient samples in duplicates.
2. Pipet **100 µl** of each **calibrator, control** and **sample** into the wells prepared.
3. Add **100 µl** of **Enzyme Conjugate** to all wells.
4. Rotate for **2 hours** at room temperature (18-25°C) on a plate mixer (900 rpm)
5. Discard the content of the wells and wash 4 times with 300 µl buffered wash solution. Remove as much wash solution as possible by beating the microplate carefully.
6. Add **200 µl** of **TMB-Substrate Solution** to all wells.
7. Incubate without shaking at room temperature (18-25°C) for **30 minutes** in the dark.
8. Add **50 µl** of **Stop Solution** to each well and mix carefully.
9. Determine the absorbance of each well at **450 nm** and read the wells within 15 minutes.

6.3 CALCULATION OF RESULTS

1. Calculate the average absorbance values for each set of standards, controls and samples.
2. The obtained optical densities of the standards (y-axis, linear) are plotted against their corresponding concentrations (x-axis, logarithmic) either on semi-logarithmic paper or using an automated method.
3. Using the mean absorbance value for each sample determine the corresponding concentration from the standard curve.
4. Automated method: The results in the IFU have been calculated automatically using a 4 PL (4 Parameter Logistics) curve fit. 4 Parameter Logistics is the preferred calculation method. Other data reduction functions may give slightly different results.
5. The concentration of the samples can be read directly from this standard curve. Samples with concentrations higher than that of the highest standard have to be further diluted and assayed again. For the calculation of the concentrations this dilution factor has to be taken into account.

6.3.1 EXAMPLE OF TYPICAL CALIBRATOR CURVE

The figure below shows typical results for TSH canine test kits. These data are intended for illustration only and should not be used to calculate results from another run.

Calibrator	Concentration (ng/ml)	OD (450nm)
0	0	0.071
1	0.2	0.233
2	0.46	0.423
3	1.05	0.852
4	2.2	1.569
5	5.2	3.092

7 EXPECTED NORMAL VALUES

Blood was collected from 30 apparently healthy dogs. A mean canine TSH value of 0.216 ng/ml was found, with an absolute range of 0.017 to 0.591 ng/ml.

Laboratories should consider reference range limits *as guidelines only*. Because of differences which may exist between laboratories and locales with respect to population, laboratory technique and selection of reference groups, it is important for each laboratory to establish by similar means the appropriateness of adopting the reference range suggested here.

8 QUALITY CONTROL

Good laboratory practice requires that controls are run with each calibration curve. A statistically significant number of controls should be assayed to establish mean values and acceptable ranges to assure proper performance. It is recommended to use control samples according to state and federal regulations. The use of control samples is advised to assure the day-to-day validity of results. Use controls at both normal and pathological levels. The controls and the corresponding results of the QC-Laboratory are stated in the QC certificate included in the kit. The values and ranges stated on the QC sheet always refer to the current kit lot and should be used for direct comparison of the results.

It is also recommended to make use of national or international Quality Assessment programs in order to ensure the accuracy of the results.

Employ appropriate statistical methods for analysing control values and trends. If the results of the assay do not fit to the established acceptable ranges of control materials patient results should be considered invalid. In this case, please check the following technical areas: Pipetting and timing devices, microtiter plate reader, expiration dates of reagents, storage and incubation conditions, aspiration and washing methods. After checking the above mentioned items without finding any error contact your distributor or Demeditec directly.

9 PERFORMANCE CHARACTERISTICS

9.1 ANALYTICAL SENSITIVITY

The analytical sensitivity of the TSH canine ELISA was calculated by adding two standard deviations from the mean of twenty-two (22) replicate analyses of *Calibrator 0*. The analytical sensitivity of the assay is 0.00002 ng/ml.

9.2 REPRODUCIBILITY

9.2.1 INTRA-ASSAY

The intra-assay variation was determined by 18 replicate measurements of 3 serum samples within one run using the Demeditec ELISA. The intra-assay variability is shown below:

	Sample 1	Sample 2	Sample 3
Mean (ng/ml)	0.57	0.92	2.57
SD (ng/ml)	0.04	0.09	0.28
CV (%)	7.0	9.9	11.0
n =	18	18	18

9.2.2 INTER-ASSAY

The inter-assay variation was determined by duplicate measurements of 3 serum samples in 10 different runs using the Demeditec ELISA. The inter-assay variability is shown below:

	Sample 1	Sample 2	Sample 3
Mean (ng/ml)	0.53	0.97	3.15
SD (ng/ml)	0.05	0.08	0.15
CV (%)	10.2	8.5	4.8
n =	10	10	10

9.3 LINEARITY

In dilution experiments sera with high antibody concentrations were diluted with the zero calibrator and assayed in the Demeditec TSH canine kit.

Sample	Dilution Factor	measured Concentration	expected Concentration	Recovery
		[ng/ml]	[ng/ml]	[%]
1	-	1.81	-	-
	1:2	0.92	0.91	101
	1:4	0.45	0.45	100
	1:8	0.22	0.23	96
2	-	2.91	-	-
	1:2	1.50	1.46	103
	1:4	0.72	0.73	99
	1:8	0.37	0.36	103
3	-	3.41	-	-
	1:2	1.76	1.71	103
	1:4	0.79	0.85	93
	1:8	0.37	0.43	86

10 LIMITATIONS OF PROCEDURE

Reliable and reproducible results will be obtained when the assay procedure is performed with a complete understanding of the package insert instruction and with adherence to GLP (Good Laboratory Practice). Any improper handling of samples or modification of this test might influence the results.

10.1 INTERFERING SUBSTANCES

- Do not use any hemolytic, icteric or lipemic specimens to avoid any interferences.
- Samples containing sodium azide should not be used in the assay.
- Non-specific interferences with this in vitro immunoassay cannot be excluded. If unplausible results are suspected, they should be considered invalid and verified by further testing. For diagnostic purposes, results should always be considered only in conjunction with the clinical picture and further diagnostic tests.

10.2 DRUG INTERFERENCES

Until today no substances (drugs) are known to us, which have an influence to the measurement of TSH in a sample. Any medication should be taken into account when assessing the results.

11 LEGAL ASPECTS

11.1 RELIABILITY OF RESULTS

The test must be performed exactly as per the manufacturer's instructions for use. Moreover the user must strictly adhere to the rules of GLP (Good Laboratory Practice) or other applicable national standards and/or laws. This is especially relevant for the use of control reagents. It is important to always include a sufficient number of controls within the test procedure for validating the accuracy and precision of the test.

The test results are only valid if all controls meet the specified ranges and all other test parameters are also within the given assay specifications. In case of any doubt or concern please contact Demeditec.

11.2 THERAPEUTIC CONSEQUENCES

Therapeutic consequences should never be based on laboratory results alone even if all test results are in agreement with the items as stated under point 11.1. Any laboratory result is only a part of the total clinical picture of a patient.

Only in cases where the laboratory results are in acceptable agreement with the overall clinical picture of the patient therapeutic consequences should be derived. The test result itself should never be the sole determinant for deriving any therapeutic consequences.

11.3 LIABILITY

Any modification of the test kit and/or exchange or mixture of any components of different lots from one test kit to another could negatively affect the intended results and validity of the overall test. Such modification and/or exchanges invalidate any claim for replacement.

Claims submitted due to customer misinterpretation of laboratory results subject to point 11.2. are also invalid. Regardless, in the event of any claim, the manufacturer's liability is not to exceed the value of the test kit. Any damage caused to the test kit during transportation is not subject to the liability of the manufacturer.














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13 SHORT INSTRUCTION(all sample sizes given in μl)

MP Well		CAL 0	CAL 1	CAL 2	CAL 3	CAL 4	CAL 5	Sample
	ng/ml	0	0.20	0.46	1.05	2.20	5.20	
Steps	Solution							
Pipet	Calibrator	100	100	100	100	100	100	-
Pipet	Sample	-	-	-	-	-	-	100
Pipet	Enzyme Conjugate	100	100	100	100	100	100	100
Incubate for 2h at RT (18-25°C) on a shaker (900 rpm)								
Decant Wash 4x with 300 μL of buffered wash solution								
Pipet	Substrate Solution	200	200	200	200	200	200	200
Incubate for 30 min at RT (18-25°C) in the dark								
Pipet	Stop Solution	50	50	50	50	50	50	50
Read at $\lambda = 450 \text{ nm}$								

SYMBOLS USED WITH DEMEDITEC ELISA

Symbol	English	Deutsch	Française	Espanol	Italiano
	European Conformity	CE-Konformitätskennzeichnung	Conforme aux normes européennes	Conformidad europea	Conformità europea
	Consult instructions for use	Gebrauchsanweisung beachten	Consulter les instructions d'utilisation	Consulte las Instrucciones	Consultare le istruzioni per l'uso
	In vitro diagnostic device	In-vitro-Diagnostikum	utilisation Diagnostic in vitro	Diagnóstico in vitro	Per uso Diagnostica in vitro
	For research use only	Nur für Forschungszwecke	Seulement dans le cadre de recherches	Sólo para uso en investigación	Solo a scopo di ricerca
	Catalogue number	Katalog-Nr.	Référence	Número de catálogo	No. di catalogo
	Lot. No. / Batch code	Chargen-Nr.	No. de lot	Número de lote	Lotto no
	Contains sufficient for <n> tests/	Ausreichend für "n" Ansätze	Contenu suffisant pour "n" tests	Contenido suficiente para <n> ensayos	Contenuto sufficiente per "n" saggi
	Note warnings and precautions	Warnhinweise und Vorsichtsmaßnahmen beachten	Avertissements et mesures de précaution font attention	Tiene en cuenta y advertencias precauciones	Annoti avvisi e le precauzioni
	Storage Temperature	Lagerungstemperatur	Température de conservation	Temperatura de conservacion	Temperatura di conservazione
	Expiration Date	Mindesthaltbarkeitsdatum	Date limite d'utilisation	Fecha de caducidad	Data di scadenza
	Legal Manufacturer	Hersteller	Fabricant	Fabricante	Fabbricante
<i>Distributed by</i>	Distributed by	Vertrieb durch	Distribution par	Distribución por	Distribuzione da parte di
	Version	Version	Version	Versión	Versione
	Single-use	Einmalverwendung	À usage unique	Uso único	Uso una volta



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