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cCRP VET test kit

For veterinary use only!
Veterinary test kit for quantitative in vitro determination of canine CRP in serum, EDTA or lithium heparin plasma with a solo or CUBE-VET analyser

EUROLyser

English

Order information Order number: VT 0100 Order number: VT 0101 Order number: VT 1000	Indication cCRP VET test kit cCRP VET test kit cCRP VET control kit	Kit size 10 tests/pack 6 tests/pack 1 x 0.5 ml (high level)
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Test kit preparation: Allow single test at least 10 minutes to warm up to room temperature (20 - 25 °C) by placing the test into the test kit rack. Put test kit package back into refrigerator.

Summary

This veterinary in vitro test kit is designed to measure canine C-reactive protein (cCRP) in serum and plasma. C-reactive protein (CRP) is an acute phase protein. The serum level of CRP rises in animals that suffer from different types of infection or inflammation. It correlates with degree and activity of the causative disease. Measurements of serum CRP are used to aid the evaluation of the inflammation associated with infection, tissue damage and other clinical disorders. Increased levels of CRP were described in dogs that suffered from pneumonia, pyometra, pancreatitis, parvovirus infection or traumatism. Dogs with an increased number of leukocytes had significantly higher CRP levels than dogs with noninflammatory leukograms. Elevated CRP levels were also found in dogs with arthritis, thrombophlebitis or proctitis and in dogs infected with *Bordetella bronchiseptica* or *Ehrlichia canis*.

Method

Immunoturbidimetric determination at 546 nm of the sample turbidity which is proportional to the concentration of cCRP present in the original specimen.

Measurement Range

10 - 200 mg/l (Lot dependent)

Samples with concentrations higher than the upper limit of the measurement range must be diluted 1 + 4 with physiological saline (0.9% NaCl solution), e.g. 20 µl sample + 80 µl 0.9% NaCl solution, and the result then multiplied by 5.

Values up to 800 mg/l show no prozone effect.

Sample Material

Use 5 µl serum, lithium heparin plasma or EDTA plasma. Sample stability testing showed that cCRP (in serum) was stable for two weeks when stored at 2 - 8 °C.

Test Kit

ERS cuvette filled with buffer reagent.
ERS cap filled with immunoparticles.

Stability and Storage

Stable until the expiration date stated on the label when stored in unopened vacuum package at 2 - 8 °C. Opening the vacuum package may limit the reagent stability to three months (stored at 2 - 8 °C) from the date of opening. DO NOT FREEZE!

Warnings and Precautions

DO NOT INGEST! Avoid contact with skin and eyes. Observe all necessary precautions for the use of laboratory reagents.

Waste Management

Please refer to local legal requirements.

Reference Range

Dog: < 10 mg/l (< 1 mg/dl)*

It is recommended that each laboratory establishes its own reference range.

Quality Control

For internal quality control the Eurolyser cCRP VET control kit is recommended. Order number: VT 1000

Precision

Reproducibility within-run:
Dog serum; N = 20, mean = 77 mg/l; CV = 4.7%.

Correlation

Correlation with Gentian cCRP:
42 samples splitted in serum and lithium heparin plasma:
y (solo cCRP) = 1.019 x (Gentian cCRP) - 0.76; R² = 0.98.

Interferences

The test system has been analysed for various interferences. Criterion was the recovery within 10% of initial values.

Haemoglobin	500 mg/dl
Intralipid	1000 mg/dl

References

1. HILLSTRÖM, A., HAGMAN, R., SÖDER, J., HAGGSTRÖM, J., LJUNGVALL, J., KJEL, GAARD-HANSEN, M. (2015) Validation and application of a canine-specific automated high-sensitivity C-reactive protein assay

The SOCRA CCRP certification examination is an essential component of professional development, and passing this SOCRA CCRP test can increase career options and a rise in salary. Nonetheless, getting ready for the Certified Clinical Research Professional (CCRP) (CCRP) exam may be difficult, and many working professionals have trouble locating the SOCRA CCRP practice questions they need to succeed in this endeavor.

SOCRA CCRP Exam Syllabus Topics:

Topic	Details
Topic 1	Research Study Closure: This section of the exam measures the skills of Clinical Research Coordinators and covers the activities required to properly conclude a clinical trial. It involves participating in the study closeout visit to verify documentation and account for the investigational product. The domain also includes developing and submitting final closure reports to the IRB, study sponsor, regulatory authorities, and clinicaltrials.gov. Finally, it covers the procedures for archiving study records.

Topic 2	• Research Study Start-Up: This section of the exam measures the skills of Clinical Research Coordinators and covers the initial planning and setup of a clinical trial. It involves coordinating the development of the study protocol, ensuring it considers ethical guidelines and regulatory pathways like IND or IDE. It also includes creating essential study documents like informed consent forms and case report forms. The domain covers obtaining necessary approvals from stakeholders like the IRB and sponsor, selecting study sites, training staff, and ensuring the study's compliance with various laws. Additionally, it involves obtaining the research product and preparing all necessary tools and documentation for the study's commencement. • Research Study Implementation: This section of the exam measures the skills of Clinical Research Associates and covers the active management and execution of the clinical trial. It focuses on following the study protocol and standard operating procedures, managing the investigational product, and ensuring ongoing regulatory compliance. The domain includes identifying, documenting, and reporting any study anomalies such as adverse events or protocol deviations. It also involves managing subject recruitment, consent, and retention, as well as maintaining all study records and essential documents. Furthermore, it covers communicating with all study stakeholders and participating in study audits to ensure quality and adherence to regulations.
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SOCRA Certified Clinical Research Professional (CCRP) Sample Questions (Q126-Q131):

NEW QUESTION # 126

According to 21 CFR Part 11, each electronic signature must be unique and:

- A. Cannot be reused or reassigned
- B. Transferable to family
- C. Reassignable after validation
- D. Identical to handwritten signature

Answer: A

Explanation:

* 21 CFR 11.100(a): Requires that electronic signatures be "unique to one individual and shall not be reused or reassigned to anyone else."

* This ensures accountability and audit trail integrity.

References: 21 CFR 11.100(a).

NEW QUESTION # 127

Which of the following is one of the responsibilities of an investigator?

- A. Selecting qualified monitors on the basis of training, experience, and expertise
- B. Maintaining accurate and current case histories of study subjects
- C. Participating in the IRB/IEC voting process for approval of their protocol
- D. Updating the investigator brochure with new safety information

Answer: B

Explanation:

Investigators are required to maintain accurate subject records, often referred to as case histories.

* 21 CFR 312.62(b): "An investigator shall prepare and maintain adequate and accurate case histories that record all observations

and other data pertinent to the investigation."

* ICH E6(R2) 4.9.0: Reinforces that investigators are responsible for recording, handling, and storing clinical trial data.

Incorrect options:

* B: Investigators may present protocols but cannot vote on IRB approval.

* C: Sponsor responsibility (ICH E6 §5.18).

* D: Sponsors are responsible for IB updates (ICH E6 §7.3.1).

Correct answer: A.

References:

21 CFR 312.62(b).

ICH E6(R2), §4.9.0.

NEW QUESTION # 128

Protocol increases drug dose by 20%. Baseline: 370 mg/m². New dose = ?

- A. 492 mg/m²
- **B. 444 mg/m²**
- C. 400 mg/m²
- D. 420 mg/m²

Answer: B

Explanation:

$370 \times 1.20 = 444 \text{ mg/m}^2$.

Accurate dosing calculations are critical for protocol adherence and patient safety.

References: Standard pharmacology dose adjustment principles; ICH E6(R2) §4.5.1.

NEW QUESTION # 129

An investigator received \$60,000 equity interest three years after study completion. What is the financial reporting requirement?

- **A. None**
- B. Report to FDA
- C. Report to sponsor
- D. Report to OHRP

Answer: A

Explanation:

* 21 CFR 54.4(b): Requires disclosure during the study and for 1 year after completion.

* After three years, no disclosure is required.

References: 21 CFR 54.4(b).

NEW QUESTION # 130

Before approving a research protocol, an IRB/IEC must determine compliance with which of the following requirements?

- A. The sponsor is qualified to provide oversight of the trial
- B. The investigator has adequate access to patients eligible for the trial
- **C. The selection of subjects is equitable**
- D. A plan for the publication of study results is in place

Answer: C

Explanation:

IRB/IEC review focuses on ethical protection of human subjects. Equitable subject selection is a cornerstone principle.

* 45 CFR 46.111(a)(3): "In order to approve research... the IRB shall determine that: Selection of subjects is equitable."

* ICH E6(R2) 3.1.2: "The IRB/IEC should safeguard the rights, safety, and well-being of all trial subjects... with particular attention to trials that may include vulnerable subjects." Publication plans (A) are not required by IRBs. Access to patients (C) and sponsor qualifications (D) are evaluated by sponsors, not IRBs. The IRB's role is ensuring fairness, minimizing coercion, and protecting vulnerable populations.

Thus, the correct answer is B (The selection of subjects is equitable).

References:

45 CFR 46.111(a)(3) (Equitable subject selection).

ICH E6(R2), §3.1.2 (IRB/IEC role).

NEW QUESTION # 131

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When we are not students, we have more responsibility. The time we can be dedicated to learning is less, but if you want to have a better development in the IT industry, it is very important to pass the international recognized IT certification exam such as CCRP exam. However, the IT elite our ExamCost make efforts to provide you with the quickest method to help you Pass CCRP Exam. We provide three type version of CCRP exam materials: PDF, online and software version, and each version has its unique benefit. You can combine what you like and to choose a free trial of our demo.

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