

CCRP Top Exam Dumps, CCRP Real Questions

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CCRP AACVPR ACTUAL EXAM 2025 TEST COMPREHENSIVE QUESTIONS AND VERIFIED ANSWERS (DETAILED & ELABORATED) 100% SOLVED 2025!!



Terms in this set (72)

What influences a persons metabolic rate?	Exercise, gender, genetics, age
Which is the macronutrient that provides more than twice as many calories as the other two	Fat
Foods that raise insulin levels have...	added sugar
When feeling stressed, you may feel relief after eating a piece of chocolate because...	it will raise dopamine levels
What reduces the number of receptors for dopamine	insulin
What percentage of Cardiac Rehab patients re overweight or obese	80%
Why is fructose a promoter of obesity?	promotes liver fat accumulation which promotes metabolic syndrome

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SOCRA CCRP Exam Syllabus Topics:

Topic	Details

Topic 1	<ul style="list-style-type: none"> • 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.
Topic 2	<ul style="list-style-type: none"> • 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.

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A generally accepted view on society is only the professionals engaged in professionally work, and so on, only professional in accordance with professional standards of study materials, as our Certified Clinical Research Professional (CCRP) study questions, to bring more professional quality service for the user. Our study materials can give the user confidence and strongly rely on feeling, lets the user in the reference appendix not alone on the road, because we are to accompany the examinee on CCRP Exam, candidates need to not only learning content of teaching, but also share his arduous difficult helper, so believe us, we are so professional company.

SOCRA Certified Clinical Research Professional (CCRP) Sample Questions (Q64-Q69):

NEW QUESTION # 64

An investigator discovered a new serious unanticipated adverse device effect. Who must they notify?

- A. Research pharmacist
- **B. Sponsor**
- C. FDA
- D. OHRP

Answer: B

Explanation:

* 21 CFR 812.150(a)(1):Investigators must report unanticipated adverse device effects to the sponsor within 10 working days.

* Sponsor is then responsible for notifying FDA and all investigators.

References:21 CFR 812.150(a)(1).

NEW QUESTION # 65

In order to adequately monitor a clinical trial, the monitor must be familiar with each of the following, EXCEPT the:

- A. IRB/IEC requirements for reporting to the regulatory authority
- B. Written information to be provided to the subjects
- C. Sponsor's SOPs
- D. Requirements for storage of the investigational product

Answer: A

Explanation:

Monitors verify compliance with protocol, sponsor SOPs, GCP, and regulations.

* ICH E6(R2) 5.18.4: Outlines monitor responsibilities, including verifying informed consent, protocol compliance, investigational product accountability, and adherence to sponsor SOPs.

* Monitors must also be familiar with subject-facing documents (A) and storage requirements for investigational product (B). However, IRB/IEC requirements for reporting to regulatory authorities are outside a monitor's scope.

That responsibility lies with investigators and IRBs under 21 CFR 56.108(b).

Thus, the correct answer is D.

References:

ICH E6(R2), §5.18.4.

21 CFR 56.108(b).

NEW QUESTION # 66

An unconscious patient experiencing life-threatening cardiac arrhythmias has been admitted to an emergency room. No FDA-approved treatment is available, and no legal representative is present. The clinical investigator determined that the use of an investigational antiarrhythmic drug is required. In accordance with the CFR, who must certify the investigator's determination?

- A. The sponsor's study monitor
- B. An independent physician
- C. A sub-investigator
- D. The sponsor's medical monitor

Answer: B

Explanation:

This scenario falls under emergency use of investigational drugs without informed consent.

* 21 CFR 50.23(a): Allows waiver of informed consent if subject faces a life-threatening condition, available treatments are unproven, and immediate use is required.

* 21 CFR 50.23(a)(3): Requires that "the determination... be reviewed and concurred with by a physician who is not otherwise participating in the clinical investigation." Thus, an independent physician (not part of the trial team) must certify the necessity of emergency investigational use.

Sponsors and monitors (C, D) are not authorized by regulation to make such determinations. Sub-investigators (A) lack independence and would be conflicted.

Correct answer: B (Independent physician).

References:

21 CFR 50.23(a)(3).

NEW QUESTION # 67

In a completed multi-site Phase I drug study using remote EDC, who ensures the system complies with accuracy and reliability requirements?

- A. Institution
- B. Sponsor
- C. Investigator
- D. Regulatory authority

Answer: B

Explanation:

* ICH E6(R2) 5.5.3: Sponsors are responsible for validating computerized systems used in trials.

Investigators ensure proper data entry, but system compliance lies with sponsor.

References: ICH E6(R2), §5.5.3.

