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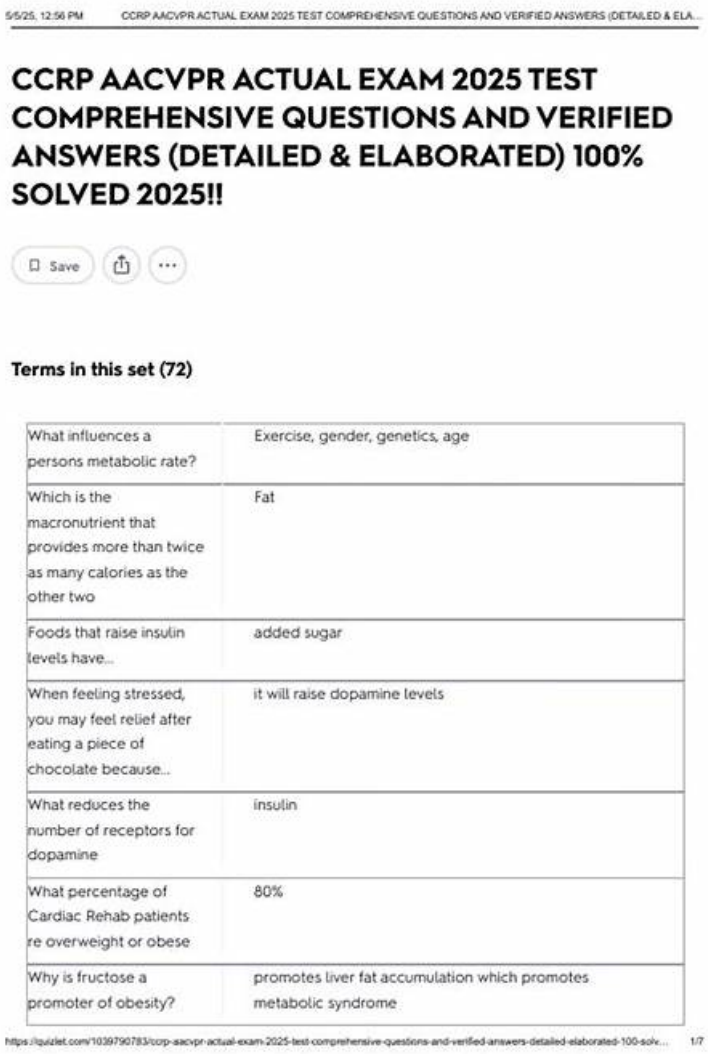

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CCRP AACVPR ACTUAL EXAM 2025 TEST COMPREHENSIVE QUESTIONS AND VERIFIED ANSWERS (DETAILED & ELABORATED) 100% SOLVED 2025!!

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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 dumps are important because they show you where you stand. After learning everything related to the Certified Clinical Research Professional (CCRP) (CCRP)certification, it is the right time to take a self-test and check whether you can clear the Certified Clinical Research Professional (CCRP) (CCRP) certification exam or not. People who score well on the Certified Clinical Research Professional (CCRP) (CCRP) practice questions are ready to give the final Certified Clinical Research Professional (CCRP) (CCRP) exam.

SOCRA CCRP Exam Syllabus Topics:

Topic	Details

Topic 1	• 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	• 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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SOCRA Certified Clinical Research Professional (CCRP) Sample Questions (Q36-Q41):

NEW QUESTION # 36
According to 21 CFR Part 11, each electronic signature must be unique and:

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

Answer: D

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 # 37
In accordance with the Belmont Report, obtaining voluntary informed consent from subjects prior to enrolling them in a clinical trial is an example of which of the following ethical principles?

- A. Respect for persons

- B. Do no harm
- C. Beneficence
- D. Justice

Answer: A

Explanation:
TheBelmont Report (1979)establishedthree key ethical principles:
* Respect for Persons:Requires informed consent, recognition of autonomy, and protection of vulnerable individuals.
* Beneficence:Obligation to maximize benefits and minimize harm.
* Justice:Ensuring fairness in subject selection and distribution of research burdens/benefits.
Voluntary informed consent embodiesRespect for Persons, as subjects are given adequate information and freedom of choice. "Do no harm" (A) is a Hippocratic principle but not Belmont terminology.
Thus, the correct answer isB (Respect for persons).
References:
The Belmont Report (1979), Part B: Basic Ethical Principles.

NEW QUESTION # 38
A sponsor's monitor is conducting a site selection visit for an interventional drug trial. In accordance with ICH GCP, which pharmacy drug storage facility information should be collected in order to determine whether the site could be selected for the trial?

- A. Available storage square footage
- B. Storage cost
- C. Storage facility temperature range
- D. Number of staff members

Answer: C

Explanation:
Drug storage conditions are essential to maintaining investigational product (IP) integrity. According to ICH:
* ICH E6(R2) 5.13.3:"The sponsor should ensure that investigational products are stored... under appropriate conditions as specified by the sponsor and in accordance with applicable regulatory requirement(s)."
* ICH E6(R2) 4.6.4:"The investigator/institution should store the investigational product(s) as specified by the sponsor (and in accordance with applicable regulatory requirement(s)), and ensure that product (s) are used only in accordance with the approved protocol." During site qualification/selection, the monitor evaluates storage conditions - particularlytemperature ranges- to ensure the site can meet the stability requirements for the IP. Factors like staff numbers, space, and cost are operational considerations but not regulatory determinants of site qualification.
Thus, the correct answer isC (Storage facility temperature range). This ensures compliance with sponsor specifications, product stability, and ultimately subject safety.
References:
ICH E6(R2), §5.13.3 (Product storage requirements).
ICH E6(R2), §4.6.4 (Investigator product storage responsibilities).

NEW QUESTION # 39
In accordance with the ICH GCP Guideline, which of the following can an Independent Data Monitoring Committee provide?

- A. The selection of qualified investigators
- B. An initial review and approval of a trial
- C. Suggestions for a new trial design
- D. Recommendations to stop a trial

Answer: D

Explanation:
AnIndependent Data Monitoring Committee (IDMC or DSMB)is a group of independent experts established to review accumulating safety and efficacy data during a trial. Their main role is to ensure subject protection and trial integrity.
* ICH E6(R2) 5.5.1:"The sponsor may consider establishing anindependent data-monitoring committee (IDMC)to assess the progress of a clinical trial, including the safety data and the critical efficacy endpoints, and to recommend to the sponsor whether to continue, modify, or stop a trial." Thus, DSMBs/IDMCsdo not perform trial approvals (A),do not design trials (C), anddo not select

investigators (D). Their authority is strictly advisory, providing recommendations to sponsors about safety and whether continuation of the study is ethically justified. The sponsor makes the final decision, but DSMB recommendations are highly influential.
Therefore, the correct answer isB (Recommendations to stop a trial).
References:
ICH E6(R2), §5.5.1 (Independent Data Monitoring Committees).

NEW QUESTION # 40
An investigator is working with a new sponsor to submit a cardiovascular trial to the IRB/IEC. In accordance with the ICH GCP Guidelines, which parties should sign the protocol to confirm agreement that the trial will be conducted as agreed?

- A. The sponsor and the IRB/IEC
- B. The sponsor and the FDA
- C. The investigator/institution and the sponsor
- D. The investigator/institution and the delegated site staff

Answer: C

Explanation:
Theprotocol signature pagedocuments agreement between the sponsor and the investigator/institution to conduct the trial in compliance with ICH GCP, the protocol, and regulatory standards.
* ICH E6(R2) 8.2.2 (Signed protocol and amendments):Requires "the sponsor and investigator /institution to sign the protocol and amendments, confirming agreement."
* ICH E6(R2) 4.5.1:"The investigator should conduct the trial in compliance with the protocol agreed to by the sponsor and, if required, approved by the regulatory authority(ies) and by the IRB/IEC." The sponsor-investigator signatures ensure shared accountability for subject protection, data integrity, and adherence to trial methodology. Neither delegated staff (B) nor IRB/IEC (C) nor FDA (D) sign protocols.
These bodies approve or oversee, but do not formally enter into execution of the protocol.
Thus, the correct answer isA (The investigator/institution and the sponsor).
References:
ICH E6(R2), §8.2.2 (Signed protocol and amendments).
ICH E6(R2), §4.5.1 (Investigator compliance with protocol).

NEW QUESTION # 41
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