

Pass Guaranteed Quiz 2026 ACRP-CP: Updated ACRP Certified Professional Exam Test Review

NEW ACRP-CP UPDATED 2025-2026 EXAM WITH Q&A 100% VERIFIED ANSWERS {CLEAN VERSION}

The Belmont Report (1979) - CORRECT ANSWER-ethical principles and guidelines for the protection of human subjects of research.

respect for persons - CORRECT ANSWER-individuals should be treated as autonomous agents and persons with diminished autonomy are entitled to protection

An autonomous person - CORRECT ANSWER-A person capable of deliberation about personal goals and of acting under the direction of such deliberation

Beneficence - CORRECT ANSWER-Do not harm and maximize possible benefits and minimize possible harms

Justice - CORRECT ANSWER-1) to each person an equal share 2) to each person according to individual need 3) to each person according to individual effort 4) to each person according to societal contributions 5) to each person according to merit

Justice - CORRECT ANSWER- The selection of research subjects needs to be scrutinized in order to determine whether some classes are being systematically selected simply because of their easy availability, their compromised position, or their manipulability, rather than for reasons directly related to the problem being studied.

Justice - CORRECT ANSWER- Whenever research supported by public funds leads to the development of therapeutic devices and procedures, justice demands both that these not provide advantages only to those who can afford them and that such research should not unduly involve persons from groups unlikely to be among the beneficiaries of subsequent applications of the research.

Three elements of the consent process - CORRECT ANSWER-information, comprehension, voluntariness

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ACRP Certified Professional Exam Sample Questions (Q120-Q125):

NEW QUESTION # 120

While consenting a potential subject, a coordinator discovered the subject could not read. Under what conditions can this potential subject be enrolled?

- **A. Get an impartial witness involved in consenting this person.**
- B. There are no circumstances under which this person can be enrolled.
- C. Locate a translator and have them work with the person.
- D. Have the Principal Investigator consent the person.

Answer: A

Explanation:

When a participant cannot read, an impartial witness must be present during the consent process. The witness ensures that the information is accurately presented and that the subject's consent is truly informed. The witness must sign the consent form to attest that the subject understands the information.

GCP guidelines require the use of an impartial witness during the informed consent process for individuals who are illiterate or visually impaired.

"Informed consent must involve an impartial witness when the subject cannot read, ensuring the consent process is conducted ethically." Objectives:

- * Maintain ethical standards during consent.
- * Protect the rights of participants with literacy challenges.

NEW QUESTION # 121

Who is responsible for securing agreement from all involved parties to ensure direct access of all trial-related source documents?

- **A. Sponsor**
- B. CRC
- C. CRO
- D. Investigator

Answer: A

Explanation:

The sponsor is responsible for ensuring that agreements are in place with all involved parties (including investigators and institutions) to grant direct access to trial-related source documents. This is crucial for monitoring, auditing, and inspection purposes, ensuring transparency and compliance with regulatory requirements.

According to GCP guidelines, the sponsor must establish agreements to secure direct access to trial data for verification and compliance checks.

"The sponsor should ensure that agreements are in place to permit direct access to source data and documents for monitoring and inspection." Objectives:

- * Maintain compliance with regulatory requirements.
- * Facilitate data verification and quality assurance.

NEW QUESTION # 122

Which document confirms the PI's agreement to permit auditing at the study site?

- A. IB
- B. Delegation Log
- C. Protocol

- **D. ICF**

Answer: D

Explanation:

The Informed Consent Form (ICF) typically includes a statement indicating that the participant's records may be reviewed by monitors, auditors, and regulatory authorities. This ensures transparency and compliance with regulatory requirements, allowing for audits and inspections when necessary.

This answer follows GCP guidelines which specify that the ICF should include consent for audits and inspections to protect subject confidentiality while ensuring data integrity.

"The ICF must include a statement allowing access to trial data for monitoring, auditing, and regulatory inspection purposes."

Objectives:

- * Ensure informed consent for data access.
- * Facilitate compliance with auditing requirements.

NEW QUESTION # 123

The sponsor calls the site and informs the research team that they have decided to temporarily suspend the study. What step should the research team take FIRST?

- **A. Inform participant and assure proper care is provided.**
- B. Inform the monitor of the termination of the study.
- C. Schedule participant for early termination visit.
- D. Inform the IRB/IEC of the study closure.

Answer: A

Explanation:

When a study is temporarily suspended, the primary responsibility of the research team is to ensure the ongoing care and safety of the enrolled participants. Participants must be informed about the suspension and assured that their health and safety will continue to be monitored.

GCP guidelines prioritize the safety and well-being of participants, which must be maintained even during a temporary suspension.

"In the event of a study suspension, the first priority is to inform the participants and ensure that appropriate care and monitoring are continued." Objectives:

- * Protect participant safety during study suspension.
- * Maintain clear communication with study participants.

NEW QUESTION # 124

The IB contains which of the following sections?

- A. Participant eligibility criteria
- **B. Known pharmacokinetic effects**
- C. IP preparation instructions
- D. Participant dosing schedule

Answer: B

Explanation:

The Investigator's Brochure (IB) contains comprehensive information about the investigational product (IP), including its pharmacokinetic and pharmacodynamic properties, preclinical and clinical trial data, and any known side effects. The IB provides a scientific basis for the study protocol and helps investigators understand the potential risks and benefits.

GCP guidelines state that the IB must include relevant scientific data, including pharmacokinetic and pharmacodynamic profiles.

"The IB should provide data on the pharmacokinetics and pharmacodynamics of the investigational product to inform clinical practice." Objectives:

- * Provide comprehensive data on investigational products.
- * Support safe and informed clinical decision-making.

NEW QUESTION # 125

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