

New ACRP-CP Test Tips - Reliable ACRP-CP Test Answers

ACRP Practice Exam Questions and Answers

A subject is issued 120 tablets and is instructed to take 2 tablets 4 times a day. He returns 88 tablets on the morning of day 9 fasting for laboratory tests. What percent compliant is he? -

ANSWER 50%

To be eligible for a trial, the subjects must have liver function tests no greater than twotimes the upper limit normal and renal function tests no greater than three times the upper limit normal. All of the following are normal ranges for the trial:

AST 5-65

ALT 5-35

BUN 4-25

Creat 0.5-1.2

Amylase 56-190

Lipase 4-24

ALK Phos 0-110 - ANSWER AST 130; ALT: 70; BUN 50; Creat 2.4

A subject presents at a site with her husband after pre-qualifying on a phone screen. She states that she is legally blind and cannot read the ICF. A Braille ICF is not available. This subject is able to sign her name if her hand is guided to the signature line. Which of the following is the BEST course of action to obtain legal consent from the subject? - **ANSWER The subject and an impartial witness can sign the ICF after it is read to them and she verbally states her understanding.**

Which of the following is MOST useful for scheduling trial procedures? - **ANSWER trialschedule of events**

A subject is participating in a clinical trial where only the pharmacist and sponsor knows the identity of the IP. The pharmacist has no contact with the trial subject and the clinical team.

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ACRP Certified Professional Exam Sample Questions (Q80-Q85):

NEW QUESTION # 80

Which of the following should be reviewed and evaluated by qualified experts to assess implications for the safety of the trial subjects?

- A. Project feasibility considerations
- **B. Emerging animal toxicological and clinical data**
- C. Sample collection storage, disposal, and shipment requirements
- D. PI roles and responsibilities

Answer: B

Explanation:

Qualified experts should evaluate emerging animal toxicological and clinical data to assess potential safety implications for trial subjects. These data are critical in identifying potential risks, adverse effects, or safety concerns before exposing human subjects to the investigational product. Early detection of safety issues through expert evaluation helps protect participant well-being. GCP guidelines stress the importance of expert assessment of preclinical and clinical data to identify risks and ensure participant safety.

"Emerging toxicological and clinical data should be carefully reviewed by qualified experts to identify safety concerns before clinical use." Objectives:

- * Ensure participant safety through expert data analysis.
- * Identify potential safety risks early in the trial process.

NEW QUESTION # 81

The CRC is prepared to consent a cognitively impaired participant to an Alzheimer's clinical trial. The CRC is required to utilize which of the following in the consent process?

- **A. A legally acceptable representative**
- B. A member of the research team
- C. A family member
- D. A witness

Answer: A

Explanation:

For cognitively impaired participants who may not fully understand the informed consent process, a legally acceptable representative (LAR) must be involved. This ensures that the participant's rights and interests are protected and that consent is obtained ethically and legally.

GCP guidelines mandate that, in cases where participants are unable to provide informed consent, an LAR must be involved to make decisions on their behalf.

"When a participant is cognitively impaired, informed consent must be obtained from a legally acceptable representative to ensure ethical participation." Objectives:

- * Safeguard the rights of vulnerable populations.
- * Maintain ethical standards in the consent process.

NEW QUESTION # 82

A new device trial is being considered. Before committing to participate in the trial, what is the MOST important item the PI needs to evaluate?

- A. Location of stored trial records
- **B. Availability of qualified staff to conduct the trial**
- C. Length of time to receive the approved trial device
- D. Information to be included in the advertising flyer

Answer: B

Explanation:

The availability of qualified staff to conduct the trial is essential for maintaining compliance with protocol requirements and ensuring patient safety. Without adequately trained and available staff, the trial's integrity and data quality are compromised.

This answer is based on GCP guidelines emphasizing the importance of having trained and qualified personnel before initiating a trial. "The PI must ensure that sufficient qualified staff is available to conduct the trial as per the protocol and regulatory requirements."

Objectives:

- * Assessing resource availability
- * Ensuring readiness to initiate a clinical trial

NEW QUESTION # 83

While reviewing reports of data completion, the sponsor notices low retention rates at many participating sites. What is an appropriate FIRST action for the sponsor to take?

- A. Require participants to provide documented reason for withdrawal.
- **B. Meet with the site staff to understand their workflows and to review retention strategies.**
- C. Submit revised ICFs to the IRB/IEC with increased compensation for participants.
- D. Interview participants who have dropped out.

Answer: B

Explanation:

Meeting with site staff to understand workflows and retention strategies is the most practical first step. By engaging with the team, the sponsor can identify potential issues affecting retention, such as site-related factors, participant burden, or protocol complexities. Addressing these issues collaboratively can improve retention without needing major protocol changes.

GCP guidelines recommend assessing and understanding site-specific challenges before making procedural changes or protocol amendments.

"Engaging with site staff to discuss retention issues helps identify root causes and develop practical solutions." Objectives:

- * Improve participant retention through collaboration.
- * Identify and address site-specific challenges.

NEW QUESTION # 84

Who ensures information regarding payments to subjects is documented in a written consent form?

- A. Sponsor
- **B. IRB/IEC**
- C. Coordinator
- D. PI

Answer: B

Explanation:

The IRB/IEC is responsible for reviewing and approving the consent form, which must include any information about payments to participants. This ensures transparency and ethical compliance regarding compensation.

The answer is verified according to GCP guidelines, which mandate IRB/IEC review of consent forms, including payment details.

"IRB/IEC must ensure that the informed consent process is thorough and includes disclosure of any compensation or payment to participants." Objectives:

- * Guaranteeing informed consent quality
- * Ensuring transparency about participant payments

NEW QUESTION # 85

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