

ACRP-CP최고덤프 - ACRP-CP최신덤프문제



참고: PassTIP에서 Google Drive로 공유하는 무료 2026 ACRP ACRP-CP 시험 문제집이 있습니다:
<https://drive.google.com/open?id=1Alam3lkIBN4wDwt3HUQIFArVetIUHngL>

만약 시험만 응시하고 싶으시다면 우리의 최신ACRP ACRP-CP자료로 시험 패스하실 수 있습니다. PassTIP 의 학습 가이드에는ACRP ACRP-CP인증시험의 예상문제, 시험문제와 답 임으로 100% 시험을 패스할 수 있습니다.우리의 ACRP ACRP-CP시험자료로 충분한 시험준비하시는것이 좋을것 같습니다. 그리고 우리는 일년무료 업데이트를 제공합니다.

PassTIP의ACRP인증ACRP-CP자료는 제일 적중률 높고 전면적인 덤프임으로 여러분은 100%한번에 응시로 패스하실 수 있습니다. 그리고 우리는 덤프를 구매 시 일년무료 업데이트를 제공합니다. 여러분은 먼저 우리 PassTIP사이트에서 제공되는ACRP인증ACRP-CP시험덤프의 일부분인 데모 즉 문제와 답을 다운받으셔서 체험해보실 수 있습니다.

>> ACRP-CP최고덤프 <<

ACRP-CP최고덤프 덤프로 ACRP Certified Professional Exam 시험을 패스하여 자격증 취득하기

PassTIP는 많은 분들이ACRP인증ACRP-CP시험을 응시하여 성공하도록 도와주는 사이트입니다PassTIP의 ACRP인증ACRP-CP 학습가이드는 시험의 예상문제로 만들어진 아주 퍼펙트한 시험자료입니다. ACRP인증ACRP-CP시험은 최근 가장 인기있는 시험으로 IT인사들의 사랑을 독차지하고 있으며 국제적으로 인정해주는 시험이라 어느 나라에서 근무하나 제한이 없습니다. PassTIP로 여러분은 소유하고 싶은 인증서를 빠른 시일내에 얻게 될것입니다.

최신 ACRP Certified Professional ACRP-CP 무료샘플문제 (Q45-Q50):

질문 # 45

A sponsor writes a protocol comparing an IP XYZ to a marketed drug ABC to determine if XYZ is more efficacious in the target population than ABC. Both drugs are prepared in identically masked IV bags and distributed according to the randomization scheme outlined in the protocol such that the study team is unaware of the treatment assignment. Which of the following is an appropriate title for this study?

- A. A randomized, open-label, comparator study comparing the efficacy of XYZ to ABC in the target population

- B. A randomized, double-blind, superiority study comparing the efficacy of XYZ to ABC in the target population
- C. A randomized, double-blind, double-dummy, superiority study comparing the efficacy of XYZ to ABC in the target population
- D. A randomized, single-blind, placebo-controlled study comparing the efficacy of XYZ to ABC in the target population

정답: B

설명:

Since both the investigator and the participant are unaware of the treatment assignment, the study is classified as double-blind. The study aims to establish the superiority of XYZ over ABC, making it a superiority study.

The use of masked IV bags confirms the double-blind design.

The answer is verified from GCP guidelines on blinding and superiority study designs.

"In double-blind studies, neither the participant nor the investigator knows the treatment assignment, which prevents bias."

Objectives:

- * Understanding blinding methods in clinical trials
- * Ensuring unbiased efficacy comparisons

질문 # 46

An impartial witness should be present during the entire informed consent discussion when:

- A. A subject has been determined to be vulnerable.
- B. A parent/guardian is consenting for a minor subject.
- C. A legally acceptable representative is unable to read.
- D. An interpreter is translating the consent form for a subject.

정답: C

설명:

An impartial witness is required when a legally acceptable representative (LAR) or the subject themselves cannot read. The witness ensures that the information is presented accurately and that the consent process is conducted ethically. The witness also signs the consent form to confirm that the subject or representative understands the study details.

GCP guidelines require an impartial witness to be present to confirm that the consent information is correctly conveyed and understood when the subject or LAR cannot read.

"An impartial witness is required when the subject or legally acceptable representative is unable to read, ensuring the consent process is transparent and ethically sound." Objectives:

- * Protect the rights of individuals with literacy challenges.
- * Maintain ethical standards in the consent process.

질문 # 47

During a monitoring visit, a CRA notices that a piece of equipment required for the study needs to be serviced. Who is responsible for addressing this problem?

- A. PI
- B. Sponsor
- C. CRA
- D. CRC

정답: A

설명:

The Principal Investigator (PI) is responsible for ensuring that all equipment used in the clinical trial is properly maintained and serviced. If a monitor (CRA) identifies equipment that needs servicing, the PI must take immediate action to ensure the equipment is in working order to maintain the quality and integrity of the study data.

GCP guidelines emphasize the PI's responsibility to ensure that all equipment used in the study is functional, properly calibrated, and serviced as needed.

"The PI is responsible for maintaining the functionality and calibration of study-related equipment to ensure accurate data collection."

Objectives:

Maintain equipment functionality to ensure data accuracy.

Ensure proper maintenance as part of site management.

질문 # 48

While reviewing a new protocol, a PI realizes a specialized laboratory test is required that the local hospital does not perform. The PI locates a laboratory that performs the specialized test and retains their services. Responsibility for ensuring the laboratory retained is qualified for this trial-related task lies with the:

- A. PI
- B. Sponsor
- C. CRO
- D. CRC

정답: A

설명:

The Principal Investigator (PI) is responsible for ensuring that any laboratory used for trial-related testing is appropriately qualified and certified. This includes verifying the laboratory's accreditation, quality control procedures, and ability to perform the required tests accurately. The PI must document the qualification process to ensure compliance with GCP and protocol requirements. GCP guidelines state that the investigator is responsible for selecting and verifying the qualification of laboratories used in the study. "The PI must ensure that any laboratory involved in the trial is properly qualified and capable of performing the specified tests according to protocol requirements." Objectives:

- * Maintain quality control in laboratory testing.
- * Ensure the accuracy and reliability of test results.

질문 # 49

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

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

정답: B

설명:

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.

질문 # 50

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PassTIP는 여러분의 요구를 만족시켜드리는 사이트입니다. 많은 분들이 우리사이트의 i인증덤프를 사용함으로 관련시험을 안전하게 패스를 하였습니다. 이니 우리 PassTIP사이트의 단골이 되었죠. PassTIP에서는 최신의ACRP ACRP-CP자료를 제공하며 여러분의ACRP ACRP-CP인증시험에 많은 도움이 될 것입니다.

ACRP-CP최신 덤프문제 : <https://www.passtip.net/ACRP-CP-pass-exam.html>

인터넷에는ACRP인증 ACRP-CP시험대비공부자료가 헤아릴수 없을 정도로 많습니다.이렇게 많은ACRP인증 ACRP-CP공부자료중 대부분 분들께서 저희PassTIP를 선택하는 이유는 덤프 업데이트가 다른 사이트보다 빠르다는 것이 제일 큰 이유가 아닐까 싶습니다, ACRP ACRP-CP최고덤프 많은 분들이 이렇게 좋은 인증시험은 아주 어렵다고 생각합니다, ACRP인증 ACRP-CP시험덤프의 인기는 이 시험과목이 얼마나 중요한지를 증명해줍니다, ACRP인증 ACRP-CP시험을 패스하여 원하는 자격증을 취득하려면PassTIP의ACRP인증 ACRP-CP덤프를 추천해드립니다, 샘플문제는 ACRP ACRP-CP최신 덤프문제덤프의 일부분 문제로서 5~10문항이 수록되어 있습니다.

인기자격증 ACRP-CP최고덤프 시험대비 덤프문제

그리고 PassTIP ACRP-CP 시험 문제집의 전체 버전을 클라우드 저장소에서 다운로드할 수 있습니다:
<https://drive.google.com/open?id=1Alam3lkIBN4wDwt3HUQIFArVetlUHngL>