

# Brain CCRP Exam | Detailed CCRP Answers

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## **CCRP AACVPR EXAM LATEST** **ACTUAL EXAM TEST BANK 520+** **QUESTIONS AND CORRECT** **DETAILED ANSWERS|AGRADE**

What insulin dose adjustment type would be most effective for preventing hypoglycemia during moderate exercise - ANSWER- **Reduction of rapid acting insulin dose 2-3 hours prior to exercise**  
**-A temporary basal rate decrease 45-60 minutes prior to exercise**

Which medications place a patient at risk for hypoglycemia - ANSWER- **Insulin (Lantus, Novolog) and Sulfonylureas (Glipizide, glimepiride)**

How long can significant exercise induced enhancement of glucose uptake from muscle last - ANSWER- **24-48 hours**

Recommended calories from carbs, protein, and fats to better manage diabetes -  
ANSWER- **~40% calories from fiber-rich carbs**  
**~30% calories from lean protein**  
**~30% calories from healthy fats**

Cardiac effects of nicotine - ANSWER- **modest increase in HR, BP and myocardial contractility**

The stethoscope for smoking is.. - ANSWER- **a carbon monoxide analyzer that measures the amount of CO the patient exhales**

Carbon monoxide - ANSWER- **binds more readily than oxygen to hemoglobin resulting in relative hypoxia**

The question regarding Time to First Cigarette is primarily used to.. - ANSWER- **determine the level of dependence**

The gold standard for improving smoking quit rates is - ANSWER- **pharmacotherapy which doubles quit rates**

Smoking cessation in patients with CVD reduces the risk of death by - ANSWER- **36%**

In order to have the greatest benefit of lowering risk of MU or death, which should replace saturated fat in diet - ANSWER- **polyunsaturated fat**

What diet is recommended by AHA/ACC? - ANSWER- **DASH diet**

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## **SOCRA Certified Clinical Research Professional (CCRP) Sample Questions**

## (Q94-Q99):

### NEW QUESTION # 94

If a subject experiences a serious adverse event related to the study drug and only minimal information is available, the investigator must submit the information to the:

- A. Sponsor and IRB/IEC within five days
- B. IRB/IEC immediately, then sponsor when full details are available
- C. Sponsor and IRB/IEC within seven days
- D. **Sponsor and IRB/IEC immediately, then update later**

### Answer: D

Explanation:

\* ICH E6(R2) 4.11.1: Investigators must "immediately report all serious adverse events to the sponsor except for those the protocol identifies as not requiring immediate reporting."

\* IRB must also be informed promptly per 21 CFR 312.64(b).

\* Follow-up information is submitted later as available.

References: ICH E6(R2), §4.11.1; 21 CFR 312.64(b).

### NEW QUESTION # 95

According to the CFR, which of the following is a complete and accurate list of the signatures required on the short form consent document?

- A. The subject or else the subject's legally authorized representative; the investigator or else the investigator's designee; the witness
- B. **The subject or else the subject's legally authorized representative; the witness**
- C. The subject or else the subject's legally authorized representative
- D. The subject or else the subject's legally authorized representative; the investigator or else the investigator's designee

### Answer: B

Explanation:

The short form consent process is permitted when the subject is presented with a brief written statement that they were informed of the study, supplemented by an oral presentation.

\* 21 CFR 50.27(b)(2): Requires the short form to be signed by the subject (or legally authorized representative) and a witness.

\* The witness ensures that oral consent was properly conveyed and understood.

\* The person obtaining consent must sign a separate written summary, but not the short form itself.

Thus, the accurate answer is A: subject (or LAR) + witness.

References:

21 CFR 50.27(b)(2).

### NEW QUESTION # 96

According to 21 CFR Part 11, each electronic signature must be unique and:

- A. Identical to handwritten signature
- B. Reassignable after validation
- C. Transferable to family
- 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 # 97

A study will enroll 420 subjects over 3.5 years. What is expected average monthly accrual?

- A. 0
- B. 1
- C. 2
- D. 3

#### Answer: B

Explanation:

420 subjects  $\div$  42 months (3.5 years) = 10 subjects/month.

However, "expected average" often rounds up to next whole number, ensuring enrollment goals are met. Thus, 11/month is correct.

This calculation is important for feasibility assessments and protocol planning.

References: Standard feasibility calculations (ICH E6(R2) §5.6).

### NEW QUESTION # 98

During the closeout visit, a monitor is completing the documentation of reconciliation of investigational product. All packaging, as well as the used and unused investigational product, are being returned to the sponsor for disposition. Which of the following documents would NOT be required to be filed at the research site?

- A. A certificate of investigational product destruction
- B. Investigational product inventory forms
- C. Records of investigational product shipment
- D. Investigational product accountability forms

#### Answer: A

Explanation:

Investigators must document the receipt, use, return, or alternative disposition of investigational product (IP).

\* ICH E6(R2) 4.6.3: Requires investigators to maintain records of IP delivery, inventory, use by subjects, and return/disposition.

\* ICH E6(R2) 8.2.14-8.2.16: Essential documents include shipment records, accountability logs, and inventory records.

However, certificates of destruction are generated and retained by the sponsor (or authorized destruction facility), not required at the site unless the destruction occurred there. In this scenario, all IP was returned to the sponsor, so no destruction certificate would exist at the site.

Thus, the correct answer is D (Certificate of destruction).

References:

ICH E6(R2), §4.6.3 (Investigator product accountability).

ICH E6(R2), §8.2.14-8.2.16 (Essential documents).

### NEW QUESTION # 99

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