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CDM Test 2023-2024 Questions and Answers 100% Correct

A Certified Dietary Manager is dissatisfied with prices from current vendors. The Manager should first:

- ask vendors to lower their prices.
- ask the consultant to recommend other vendors.
- complete a comparison study of vendors.
- discontinue purchasing from the current vendors. - ANSWER-c. complete a comparison study of vendors.

The best way to prepare frozen peas is to:

- slowly cook the peas at 200°F (93.3°C) so they do not dry out.
- cook them rapidly until they reach an internal temperature of 140°F (60.0°C).
- cook them to 120°F (48.9°C) and hold them in the steam table to come up to temperature.
- cook them in batches throughout the service time. - ANSWER-d. cook them in batches throughout the service time.

Beans and legumes are essential protein substitutes for clients who are:

Choose one answer.

- lactose intolerant.
- vegan.
- ovo-facto-vegetarian.
- lacto vegetarian. - ANSWER-b. vegan.

When preparing goals for the foodservice department, a Certified Dietary Manager must show that the goals are:

Choose one answer.

- narrow.
- broad.
- listed on the bulletin board.
- transferrable to other departments. - ANSWER-b. broad.

When purchasing food, a Certified Dietary Manager must develop specifications to ensure that:

Choose one answer.

- government commodities are used when available.
- eggs are delivered in a timely manner.
- milk arrives at a temperature below 41°F (5°C).
- canned fruits are packed in water or juice. - ANSWER-c. milk arrives at a temperature below 41°F (5°C).

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SCDM Certified Clinical Data Manager Sample Questions (Q63-Q68):

NEW QUESTION # 63

What should be done if the site continues to provide inconsistent data after several re-queries?

- A. Continue to re-query until the site changes the data
- B. Gently lead the site to the correct response
- C. Escalate the issue to the appropriate site contact personnel
- D. Do nothing, the data will remain inconsistent

Answer: C

Explanation:

If a clinical site continues to provide inconsistent or illogical data after multiple queries, the correct course of action is to escalate the issue to the appropriate site contact personnel, typically the Clinical Research Associate (CRA) or Site Monitor.

According to the Good Clinical Data Management Practices (GCDMP), persistent data discrepancies often indicate a misunderstanding of the protocol, CRF instructions, or data entry procedures at the site level. Repeatedly re-querying the same data without escalation wastes time and risks introducing bias or error. By escalating through formal communication channels, the issue can be clarified through re-training, documentation review, or site monitoring visits.

The GCDMP emphasizes that escalation ensures data accuracy, site accountability, and protocol adherence, maintaining both data quality and regulatory compliance. Data managers must document the escalation process in the Data Management Plan (DMP) and ensure proper follow-up resolution is achieved.

Reference (CCDM-Verified Sources):

SCDM Good Clinical Data Management Practices (GCDMP), Chapter: Communication and Issue Escalation, Section 4.2 - Handling Persistent Data Discrepancies ICH E6 (R2) Good Clinical Practice, Section 5.18 - Monitoring and Site Communication
FDA Guidance for Industry: Oversight of Clinical Investigations - Risk-Based Monitoring, Section on Issue Escalation

NEW QUESTION # 64

The result set from the query below would be which of the following?

```
SELECT * FROM patient WHERE medical_record_number > 9000
```

- A. Wider than the patient table
- B. Narrower than the patient table
- C. Longer than the patient table
- D. Shorter or of equal length than the patient table

Answer: D

Explanation:

In Structured Query Language (SQL), the WHERE clause is used to filter records based on specified criteria. The query retrieves all columns from the patient table (SELECT *) but only those rows where the medical_record_number value is greater than 9000.

This means:

The number of columns (fields) remains the same as the original table.

The number of rows (records) will be equal to or less than the number of rows in the patient table, depending on how many patients meet the filter condition.

Hence, the result set can only be shorter or equal in length compared to the original table. It cannot be longer, wider, or narrower, since no new rows or columns are created.

Therefore, option B - "Shorter or of equal length than the patient table" - is correct.

Reference (CCDM-Verified Sources):

SCDM GCDMP, Chapter: Database Design and Build, Section 5.2 - Relational Database Queries and Filtering Logic ICH E6(R2) GCP, Section 5.5.3 - Data Retrieval, Filtering, and Storage Principles
FDA Guidance for Industry: Computerized Systems Used in Clinical Investigations, Section 6.4 - Query Logic and Record Subsetting

NEW QUESTION # 65

Which is the most important reason for why a data manager would review data before a monitor reviews it?

- A. The GCDMP recommends that data managers review data prior to a monitor's review.

- B. Data managers have access to programming tools to identify discrepancies.
- **C. Data can be viewed and discrepancies highlighted prior to a monitor's review.**
- D. Data managers write the Data Management Plan that specifies the data cleaning workflow.

Answer: C

Explanation:

The primary reason data managers review data before a monitor's review is to identify and flag discrepancies or inconsistencies so that site monitors can focus their efforts more efficiently during on-site or remote source data verification (SDV).

According to the Good Clinical Data Management Practices (GCDMP, Chapter on Data Validation and Cleaning), proactive data review by data management staff ensures data completeness and accuracy by identifying missing, inconsistent, or out-of-range values. This pre-review helps streamline the monitoring process, reduces the volume of open queries, and enhances data quality.

Option A is true but not the main reason for pre-monitor review. Option C highlights a capability rather than a rationale. Option D is partially correct, but the GCDMP emphasizes process purpose, not prescriptive order. Thus, option B correctly captures the practical and process-oriented reason for early data review-to ensure data are ready and accurate for the monitor's review phase.

Reference (CCDM-Verified Sources):

SCDM GCDMP, Chapter: Data Validation and Cleaning, Section 5.3 - Data Review Timing and Purpose ICH E6(R2) GCP, Section 5.18 - Monitoring and Data Verification Requirements

NEW QUESTION # 66

On a dose escalation study, the Data Manager notices one site has a much higher number of queries than other sites and most are older than 30 days. The Data Safety Monitoring Board will meet in three weeks. What should the Data Manager providing CRO oversight do?

- A. Notify the CRO's Clinical Leader about the concerns
- **B. Consult the CRO's Lead Data Manager and the CRO's Project Leader**
- C. Ignore it for now and check back next week
- D. Call the site directly and ask the study coordinator about the concerns

Answer: B

Explanation:

The correct action is to consult the CRO's Lead Data Manager and CRO's Project Leader (Option C) to ensure the issue is addressed through the appropriate oversight and escalation process.

According to the GCDMP (Chapter: Project Management and Communication), when a sponsor Data Manager identifies significant data management issues under CRO oversight - such as aging queries or site performance disparities - communication must follow the established governance and escalation pathway defined in the Scope of Work (SOW) and Data Management Plan (DMP).

Directly contacting the site (Option B) bypasses the CRO's chain of command and violates communication protocols. Notifying only the Clinical Leader (Option A) is insufficient, and ignoring the issue (Option D) jeopardizes the Data Safety Monitoring Board (DSMB) review timeline.

Therefore, Option C ensures a documented, collaborative approach to problem resolution within the contractual oversight structure.

Reference (CCDM-Verified Sources):

SCDM Good Clinical Data Management Practices (GCDMP), Chapter: Project Management and Communication, Section 7.1 - Oversight of CRO Data Management Activities ICH E6 (R2) GCP, Section 5.2 - Contract Research Organization Responsibilities
FDA Guidance for Industry: Oversight of Clinical Investigations - Sponsor and CRO Roles and Communication Pathways

NEW QUESTION # 67

Which metrics report listed below would best help identify trends in the clinical data?

- A. Percent of data/visits cleaned
- **B. Query frequency counts per data element**
- C. Last patient/last visit date to data lock date
- D. Number of subjects screened/enrolled

Answer: B

Explanation:

The Query frequency counts per data element (Option B) is the best metric for identifying data trends and potential systemic data issues in clinical trials.

According to the Good Clinical Data Management Practices (GCDMP, Chapter: Data Quality Assurance and Control), trend analysis involves identifying recurring data issues across subjects, sites, or variables to detect training gaps, protocol misinterpretation, or CRF design flaws. A high number of queries generated for specific fields (e.g., visit date, lab values, or dosing information) may indicate systemic problems such as unclear CRF instructions or site-level misunderstandings.

While metrics such as percent of data cleaned (A) and time to database lock (B) reflect overall progress and efficiency, they do not identify specific data pattern issues. The number of subjects screened/enrolled (C) pertains to recruitment rather than data quality. Therefore, query frequency per data element provides actionable insights for quality improvement, process refinement, and early identification of potential risks.

Reference (CCDM-Verified Sources):

SCDM Good Clinical Data Management Practices (GCDMP), Chapter: Data Quality Assurance and Control, Section 6.3 - Metrics and Trend Analysis ICH E6 (R2) Good Clinical Practice, Section 5.18.4 - Risk-Based Quality Review and Data Trends FDA Guidance for Industry: Oversight of Clinical Investigations - Risk-Based Monitoring, Section 6 - Data Metrics and Trend Evaluation

NEW QUESTION # 68

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