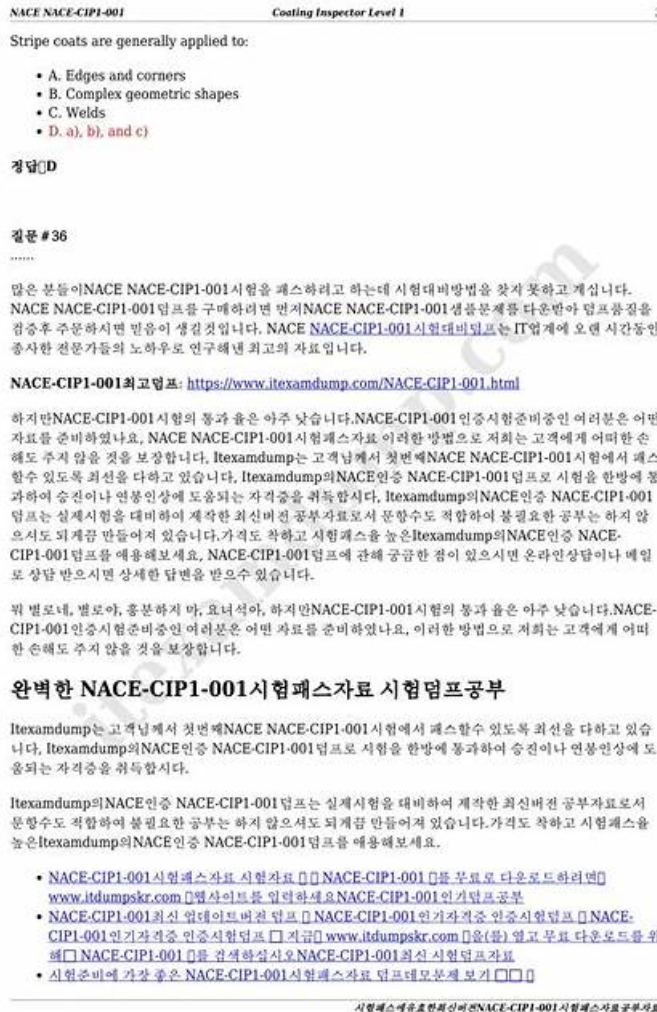


# CCDM유효한 시험자료 & CCDM최고품질덤프공부자료



그 외, PassTIP CCDM 시험 문제집 일부가 지금은 무료입니다: [https://drive.google.com/open?id=1259PVweTsiXd3miBen\\_r75ig2wZQgA2m](https://drive.google.com/open?id=1259PVweTsiXd3miBen_r75ig2wZQgA2m)

많은 분들이SCDM CCDM시험을 패스하려고 하는데 시험대비방법을 찾지 못하고 계십니다. SCDM CCDM덤프를 구매하려면 먼저SCDM CCDM샘플문제를 다운받아 덤프 품질을 검증후 주문하시면 믿음이 생길것입니다. SCDM CCDM시험대비덤프는 IT업계에 오랜 시간동안 종사한 전문가들의 노하우로 연구해낸 최고의 자료입니다.

PassTIP는 여러분의 꿈을 이루어줄 뿐만 아니라 일년무료 업뎃서비스도 따릅니다. PassTIP에서 제공하는 덤프로 여러분은 100%시험을 패스하실수 있고SCDM CCDM자격증을 취득하실 수 있습니다.지금 바로 사이트에서SCDM CCDM덤프데모 즉 덤프의 일부 문제와 답을 다운 받으셔서 체험하실 수 있습니다.

>> CCDM유효한 시험자료 <<

## CCDM최고품질 덤프공부자료, CCDM퍼펙트 공부자료

SCDM CCDM인증시험덤프는 적중율이 높아 100% SCDM CCDMSCDM CCDM시험에서 패스할수 있게 만들어져 있습니다. 덤프는 IT전문가들이 최신 실러버스에 따라 몇년간의 노하우와 경험을 충분히 활용하여 연구제작해낸 시험대비자료입니다. 저희 SCDM CCDM덤프는 모든 시험유형을 포함하고 있는 퍼펙트한 자료기에 한방에 시험 패스 가능합니다.

## 최신 Clinical Data Management CCDM 무료샘플문제 (Q37-Q42):

### 질문 # 37

A Data Manager is asked to manage SOPs for a department. Given equal availability of the following systems, which of the following is the best choice for managing the organizational SOPs?

- A. Learning management system
- B. Document management system
- C. Existing paper filing system
- D. Customized Excel spreadsheet

정답: B

### 설명:

The best choice for managing Standard Operating Procedures (SOPs) in a compliant and auditable manner is a Document Management System (DMS).

According to the GCDMP (Chapter: Regulatory Requirements and Compliance) and ICH E6 (R2), SOPs must be version-controlled, securely stored, retrievable, and auditable. A validated DMS supports controlled access, document lifecycle management (draft, review, approval, and archival), and electronic audit trails, ensuring full compliance with FDA 21 CFR Part 11 and Good Documentation Practices (GDP).

While Learning Management Systems (C) track training, they are not intended for document control. Spreadsheets (B) and paper systems (D) cannot provide adequate version tracking, access security, or audit capability required for regulatory inspection readiness.

Reference (CCDM-Verified Sources):

SCDM Good Clinical Data Management Practices (GCDMP), Chapter: Regulatory Requirements and Compliance, Section 5.2 - SOP Management and Document Control ICH E6 (R2) GCP, Section 5.5.3 - Document and Record Management FDA 21 CFR Part 11 - Electronic Records and Signatures, Section 11.10 - System Validation and Document Controls

### 질문 # 38

Which is the best reason why front-end checks are usually kept minimal, when compared to back-end checks, in a paper-based clinical study?

- A. There is no need to alert the site personnel immediately about a data issue, as the study has happened already
- B. Data entry staff should be able to enter a value into the database just as it appears in the paper CRF
- C. Data review can be performed at a later time due to the paper-based studies being smaller in size
- D. There are approvals required to raise a Data Clarification Form which could take time

정답: B

### 설명:

In paper-based clinical studies, front-end data checks (those performed during data entry) are intentionally kept minimal to ensure that data are entered exactly as recorded on the paper CRF. This principle ensures data integrity by maintaining fidelity between source and electronic records before any cleaning or edit validation occurs.

The GCDMP (Chapter: Data Validation and Cleaning) explains that data entry operators should input values as written, even if they appear incorrect or inconsistent, because the purpose of front-end checks is not to interpret but to capture data faithfully. The back-end edit checks-performed later by data managers-are designed to identify inconsistencies, out-of-range values, or logical errors that require clarification through queries.

This approach separates data capture from data cleaning, minimizing bias and preserving original investigator input. Hence, option A accurately states the rationale for keeping front-end checks minimal in paper-based studies.

Reference (CCDM-Verified Sources):

SCDM GCDMP, Chapter: Data Validation and Cleaning, Section 4.2 - Data Entry, Edit Checks, and Query Process ICH E6(R2) GCP, Section 5.5.3 - Data Handling and System Controls FDA Guidance for Industry: Computerized Systems Used in Clinical Investigations, Section 6.1 - Data Entry and Verification Processes

### 질문 # 39

A study takes body-composition measurements at baseline using a DEXA scanner. Which information is needed to correctly associate the body-composition data to the rest of the study data?

- A. Subject number

- B. Subject number and visit number
- C. Study number and visit number
- D. Study number and subject number

**정답: B**

**설명:**

To properly associate body-composition data (from a DEXA scanner) with other study data, both the subject number and the visit number are required.

According to the GCDMP (Chapter: Data Management Planning and Study Start-up), every clinical data record must be uniquely identifiable and linkable to a specific subject and study event. The subject number identifies the participant, while the visit number defines the temporal context in which the measurement was taken.

Without both identifiers, data integration becomes ambiguous-especially if multiple assessments occur over time (e.g., baseline, week 12, end of study). Including both ensures data traceability, integrity, and alignment with the protocol-defined schedule of events.

Study number (option A) alone does not distinguish between visits or subjects, and visit number alone (option C) lacks linkage to the individual participant.

Reference (CCDM-Verified Sources):

SCDM Good Clinical Data Management Practices (GCDMP), Chapter: Data Management Planning and Study Start-up, Section 4.4 - Data Linking and Identification Requirements ICH E6 (R2) GCP, Section 5.5.3 - Data Traceability Principles FDA Guidance for Industry: Computerized Systems Used in Clinical Investigations - Data Identification Requirements

**질문 # 40**

Which information is required by most systems to specify data entry screens?

- A. Page number and total number of pages
- B. User role, access level, and permissions
- C. Data type, prompt, and response format
- D. Help text, review parameters, and answers

**정답: C**

**설명:**

When designing or configuring data entry screens within an Electronic Data Capture (EDC) system, three critical components are required for each field:

Data Type - Defines the nature of the data (e.g., text, numeric, date).

Prompt - The label or question displayed to the user.

Response Format - Specifies how the user enters or selects data (e.g., free text, drop-down, checkbox).

According to the GCDMP (Chapter: EDC Systems and Database Design), these three attributes form the logical data structure required to build and validate data entry interfaces. They ensure consistency in how information is captured, displayed, and validated during data entry.

While user roles (A) and help text (D) are system-level configurations, not field-level specifications, page numbers (C) relate to printed CRFs rather than digital data screens.

Therefore, option B (Data type, prompt, and response format) correctly identifies the essential information needed to define data entry screens.

Reference (CCDM-Verified Sources):

SCDM GCDMP, Chapter: EDC Systems and Database Design, Section 4.3 - Screen Design Specifications CDISC CDASH Implementation Guide, Section 3.2 - Data Field Attributes ICH E6(R2) GCP, Section 5.5.3 - Data Capture and Input Standards

**질문 # 41**

A site study coordinator attempts to make an update in a study database in an EDC system after lock. What occurs?

- A. The change is logged as occurring after lock
- B. The change is approved by the Data Manager before it is applied
- C. The old value is replaced in all locations by the new value
- D. The site study coordinator is not able to make the change

**정답: D**

**설명:**

Once a clinical database is locked, it becomes read-only - no further data modifications can be made by any users, including site personnel. This ensures that the data are finalized, consistent, and auditable for statistical analysis and regulatory submission. According to the GCDMP (Chapter: Database Lock and Archiving), the lock process involves freezing the database to prevent accidental or unauthorized changes. After lock, access permissions are restricted, and all edit and update functions are disabled. If any corrections are required post-lock, the database must be unlocked under controlled procedures (with full audit trail documentation).

Thus, option C - The site study coordinator is not able to make the change - correctly reflects standard EDC functionality and regulatory compliance.

Reference (CCDM-Verified Sources):

SCDM GCDMP, Chapter: Database Lock and Archiving, Section 5.2 - Database Lock Procedures and Controls ICH E6(R2) GCP, Section 5.5.3 - Data Integrity and Audit Trail Requirements FDA 21 CFR Part 11 - Controls for Electronic Records and System Lock Functions

## 질문 # 42

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PassTIP을 선택함으로써 100%인증 시험을 패스하실 수 있습니다. 우리는SCDM CCDM시험의 갱신에 따라 최신의 덤프를 제공할 것입니다. PassTIP에서는 무료로 24시간 온라인상담이 있으며, PassTIP의 덤프로SCDM CCDM시험을 패스하지 못한다면 우리는 덤프전액환불을 약속 드립니다.

**CCDM최고품질 덤프공부자료** : <https://www.passtip.net/CCDM-pass-exam.html>

SCDM CCDM유효한 시험자료 고객님의 드린 약속을 꼭 지키려고 항상 최선을 다하고 있습니다, 저희가 제공하는 CCDM인증시험 덤프는 여러분이 CCDM시험을 안전하게 통과는 물론 관련 전문지식 장악에도 많은 도움이 될것입니다, SCDM인증 CCDM덤프로SCDM인증 CCDM시험을 준비하여 한방에 시험패하세요, CCDM덤프에는 CCDM인증시험의 예상문제와 가장 최근 기출문제가 수록되어 있습니다, PassTIP에서는SCDM인증CCDM시험에 대비한 공부가이드를 발췌하여 IT인사들의 시험공부 고민을 덜어드립니다, SCDM CCDM유효한 시험자료 우리는 여러분이 시험패스는 물론 또 일년무료 업데이트서비스를 제공합니다.만약 시험에서 실패했다면 우리는 덤프비용전액 환불을 약속 드립니다.하지만 이런 일은 없을 것입니다.우리는 우리덤프로 100%시험패스에 자신이 있습니다.

그렇게 말해줘서, 세은이 살짝 고개를 들고 보니, 정말CCDM로 준영이 성큼성큼 세은의 테이블 쪽으로 걸어오고 있었다, 고객님의 드린 약속을 꼭 지키려고 항상 최선을 다하고 있습니다, 저희가 제공하는 CCDM인증시험 덤프는 여러분이 CCDM시험을 안전하게 통과는 물론 관련 전문지식 장악에도 많은 도움이 될것입니다.

## 높은 적중율을 자랑하는 CCDM유효한 시험자료 덤프공부자료

SCDM인증 CCDM덤프로SCDM인증 CCDM시험을 준비하여 한방에 시험패하세요, CCDM덤프에는 CCDM인증시험의 예상문제와 가장 최근 기출문제가 수록되어 있습니다, PassTIP에서는SCDM인증CCDM시험에 대비한 공부가이드를 발췌하여 IT인사들의 시험공부 고민을 덜어드립니다.

- 최근 인기시험 CCDM유효한 시험자료 덤프문제 □ 무료로 쉽게 다운로드하려면 □ [www.exampassdump.com](http://www.exampassdump.com) □에서 □ CCDM □를 검색하세요CCDM시험대비 최신버전 문제
- CCDM시험대비 덤프공부 □ CCDM최신시험후기 □ CCDM최신버전 인기 덤프자료 □ 검색만 하면 ⇒ [www.itdumpskr.com](http://www.itdumpskr.com) □에서 ⇒ CCDM □무료 다운로드CCDM인증시험 인기 덤프자료
- CCDM최신시험후기 □ CCDM시험대비 덤프공부 □ CCDM유효한 시험자료 □ 검색만 하면 > [www.itdumpskr.com](http://www.itdumpskr.com) <에서 > CCDM □무료 다운로드CCDM유효한 시험자료
- 시험대비 CCDM유효한 시험자료 덤프 최신 데모 □ 「 [www.itdumpskr.com](http://www.itdumpskr.com) 」 을 통해 쉽게 ⇒ CCDM ◀무료 다운로드 받기CCDM최신시험후기
- CCDM시험덤프문제 □ CCDM시험대비 덤프공부자료 □ CCDM최신버전 인기 덤프자료 □ ⇒ [www.dumptop.com](http://www.dumptop.com) □에서 □ CCDM □를 검색하고 무료 다운로드 받기CCDM시험패스 인증덤프자료
- 시험대비 CCDM유효한 시험자료 덤프 최신 데모 □ ▶ [www.itdumpskr.com](http://www.itdumpskr.com) <에서 “CCDM”를 검색하고 무료 다운로드 받기CCDM최고품질 덤프자료
- CCDM시험준비 □ CCDM적중율 높은 시험덤프 □ CCDM최고품질 인증시험덤프데모 □ ⇒ [www.koreadumps.com](http://www.koreadumps.com) □□□을(를) 열고 □ CCDM □를 검색하여 시험 자료를 무료로 다운로드하십시오CCDM 인증덤프공부문제
- 시험패스 가능한 CCDM유효한 시험자료 덤프 샘플문제 다운받기 □ 무료로 다운로드하려면 > [www.itdumpskr.com](http://www.itdumpskr.com) <로 이동하여 [ CCDM ]를 검색하십시오CCDM시험덤프문제
- CCDM인증시험 인기 덤프자료 □ CCDM퍼펙트 최신버전 자료 □ CCDM최신 업데이트 덤프문제 □ 무료 다운로드를 위해 □ CCDM □를 검색하려면 《 [www.itdumpskr.com](http://www.itdumpskr.com) 》 을(를) 입력하십시오CCDM최신 업데이트

#### 트 덤프문제

- CCDM시험패스 인증덤프자료 □ CCDM최신 시험대비자료 □ CCDM최신시험후기 □ 검색만 하면 ➡  
www.itdumpskr.com □에서 ⇒ CCDM ⇐ 무료 다운로드 CCDM 시험대비 최신버전 문제
- 시험패스 가능한 CCDM 유효한 시험자료 덤프 샘플문제 다운받기 □ ✓ www.pass4test.net □ ✓ □의 무료 다운로드 [ CCDM ] 페이지가 지금 열립니다 CCDM 높은 통과율 시험덤프자료
- www.stes.tyc.edu.tw, www.stes.tyc.edu.tw, www.stes.tyc.edu.tw, www.notebook.ai, kumu.io, www.stes.tyc.edu.tw, socialtechnet.com, bookmarkport.com, bookmarkstown.com, ycs.instructure.com, Disposable vapes

그리고 PassTIP CCDM 시험 문제집의 전체 버전을 클라우드 저장소에서 다운로드할 수 있습니다:

[https://drive.google.com/open?id=1259PVweTsiXd3miBen\\_r75ig2wZQgA2m](https://drive.google.com/open?id=1259PVweTsiXd3miBen_r75ig2wZQgA2m)