

# CCRP최신업데이트인증시험자료 & CCRP최신기출자료



Research C\_S4CPR\_2302 최신업데이트 시험자료 덤프샘플문제를 무료로 Google Drive에서 다운로드하세요.  
[https://drive.google.com/open?id=1Uk8kZZKp\\_1mwgdTMTACU82gATFFTSrT0](https://drive.google.com/open?id=1Uk8kZZKp_1mwgdTMTACU82gATFFTSrT0)

SAP인증 C\_S4CPR\_2302 시험자료의 문제와 답을 모두 무료로 얻을 수 있습니다. 자신의 지식과 별 남김의 부활로  
 보충해 주시기 바랍니다. 그리고 SAP인증 C\_S4CPR\_2302 시험을 준비하는 분들께  
 피아를 드립니다. 이 자료를 사용 하려하시면 SAP인증 C\_S4CPR\_2302 시험을 위한 샘플문제를 무료로  
 받으실 수 있습니다. SAP인증 C\_S4CPR\_2302 시험을 준비하고 싶으시다면 이 자료를 꼭 다운로드 받으시기 바랍니다.

**SAP C\_S4CPR\_2302 시험요강:**

주제	소개
주제 1	<ul style="list-style-type: none"> <li>Explain features of SAP Cloud for Customer</li> <li>Identify Key Solution configurations related to SAP Archa Integration scope events</li> </ul>
주제 2	<ul style="list-style-type: none"> <li>Perform Test related to Reporting (12), Provisioning of Direct Material, and other related to SAP for Customer</li> </ul>
주제 3	<ul style="list-style-type: none"> <li>Identify Key Solution configurations related to integration</li> <li>Describe data structures for integration data migration between release libraries</li> </ul>
주제 4	<ul style="list-style-type: none"> <li>Identify the implementation details and scope</li> <li>Describe the contents of SAP Fiori Provisioning and review</li> </ul>
주제 5	<ul style="list-style-type: none"> <li>Identify SAP SuccessFactors configuration for data migration</li> <li>Perform data migration related to RISE with SAP S/4HANA Cloud</li> </ul>
주제 6	<ul style="list-style-type: none"> <li>Explain SAP SuccessFactors Technology Platform Identity Authentication Service</li> <li>Scope from Level 20 implementation and Configuration for Provisioning</li> </ul>

BONUS!!! Itcertkr CCRP 시험 문제집 전체 버전을 무료로 다운로드하세요: [https://drive.google.com/open?id=1Uk8kZZKp\\_1mwgdTMTACU82gATFFTSrT0](https://drive.google.com/open?id=1Uk8kZZKp_1mwgdTMTACU82gATFFTSrT0)

Itcertkr 의 SOCRA인증 CCRP덤프는 PDF버전과 소프트웨어버전 두가지 버전으로 되어있는데 소프트웨어버전은 시뮬레이션버전입니다. 소프트웨어버전의 문제를 푸는 과정은 시험현장을 연상케하여 시험환경에 먼저 적응하여 실제시험에서 높은 점수를 받도록 도와드릴수 있습니다.

## SOCRA CCRP 시험요강:

주제	소개
주제 1	<ul style="list-style-type: none"> <li><b>Research Study Closure:</b> This section of the exam measures the skills of Clinical Research Coordinators and covers the activities required to properly conclude a clinical trial. It involves participating in the study closeout visit to verify documentation and account for the investigational product. The domain also includes developing and submitting final closure reports to the IRB, study sponsor, regulatory authorities, and clinicaltrials.gov. Finally, it covers the procedures for archiving study records.</li> </ul>

주제 2	<ul style="list-style-type: none"> <li>• <b>Research Study Start-Up:</b> This section of the exam measures the skills of Clinical Research Coordinators and covers the initial planning and setup of a clinical trial. It involves coordinating the development of the study protocol, ensuring it considers ethical guidelines and regulatory pathways like IND or IDE. It also includes creating essential study documents like informed consent forms and case report forms. The domain covers obtaining necessary approvals from stakeholders like the IRB and sponsor, selecting study sites, training staff, and ensuring the study's compliance with various laws. Additionally, it involves obtaining the research product and preparing all necessary tools and documentation for the study's commencement.</li> <li>• <b>Research Study Implementation:</b> This section of the exam measures the skills of Clinical Research Associates and covers the active management and execution of the clinical trial. It focuses on following the study protocol and standard operating procedures, managing the investigational product, and ensuring ongoing regulatory compliance. The domain includes identifying, documenting, and reporting any study anomalies such as adverse events or protocol deviations. It also involves managing subject recruitment, consent, and retention, as well as maintaining all study records and essential documents. Furthermore, it covers communicating with all study stakeholders and participating in study audits to ensure quality and adherence to regulations.</li> </ul>
------	--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

>> CCRP최신 업데이트 인증시험자료 <<

## 100% 유효한 CCRP최신 업데이트 인증시험자료 인증덤프

Itcertkr 의 IT전문가들이 자신만의 경험과 끊임없는 노력으로 최고의 SOCRA CCRP학습자료를 작성해 여러분들이 SOCRA CCRP시험에서 패스하도록 최선을 다하고 있습니다. 덤프는 최신 시험문제를 커버하고 있어 시험패스율이 높습니다. SOCRA CCRP시험을 보기로 결심한 분은 가장 안전하고 가장 최신인 적응율 100%에 달하는 SOCRA CCRP시험대비덤프를 Itcertkr에서 받을 수 있습니다.

### 최신 Clinical Research Professional CCRP 무료 샘플문제 (Q34-Q39):

#### 질문 # 34

A clinical investigator wants to publish a subject's unique results. The consent form did not mention publication. What is required?

- A. Approval from monitor
- **B. Consent from subject**
- C. IRB chair approval
- D. Nothing further

**정답: B**

#### 설명:

\* ICH E6(R2) 4.8.10(n):Consent must include explanation about confidentiality and possible publication.

\* If not included, specific subject consent must be obtained before publishing identifiable results.

Thus, subject's explicit permission is required.

References:ICH E6(R2) §4.8.10(n).

#### 질문 # 35

An investigator's responsibilities for conducting clinical trials include:

- A. Observing preclinical drug effects
- **B. Administering or overseeing investigational drug administration**
- C. Maintaining financial documentation for study staff
- D. Maintaining IRB meeting minutes

**정답: B**

#### 설명:

\* ICH E6(R2) 4.6.1:The investigator is responsible for investigational product accountability at the site.

\* 21 CFR 312.61:Investigators must administer the investigational drug only to subjects under their supervision.

The IRB maintains meeting minutes (A), preclinical studies are sponsor tasks (B), and financial interest documentation (C) is covered

under sponsor reporting. Thus, D is correct.  
References: ICH E6(R2) §4.6.1; 21 CFR 312.61.

### 질문 # 36

In accordance with ICH, which of the following is an acceptable protocol review frequency for an IRB?

- A. 36 months
- B. 12 months
- C. 24 months
- D. 6 months

정답: B

설명:

IRBs must review protocols at least annually to ensure ongoing subject protection.

\* 21 CFR 56.109(f): "An IRB shall conduct continuing review of research at intervals appropriate to the degree of risk, but not less than once per year."

\* ICH E6(R2) 3.1.4: "The IRB/IEC should conduct continuing review of each ongoing trial at intervals appropriate to the degree of risk, but at least once per year." This establishes 12 months as the minimum required interval. More frequent reviews (e.g., 6 months) may occur for higher-risk studies, but longer intervals (24-36 months) are not permitted.

Correct answer: B (12 months).

References:

21 CFR 56.109(f).

ICH E6(R2), §3.1.4.

### 질문 # 37

According to the ICH GCP Guidelines, what is the purpose of source documents?

- A. To validate insurance reimbursement
- B. To establish diverse subject enrollment
- C. To validate reports submitted to the IRB/IEC
- D. To provide a record of subjects' investigational medical treatment

정답: D

설명:

\* ICH E6(R2) 1.52: Source documents are "original documents, data, and records... necessary for the reconstruction and evaluation of the trial."

\* Their main role is to document treatment and verify CRFs.

References: ICH E6(R2), §1.52.

### 질문 # 38

In accordance with the CFR, clinical trial sponsors are required to retain records and reports after a marketing application is approved for at least:

- A. 5 years
- B. 2 years
- C. 3 years
- D. 15 years

정답: B

설명:

The FDA record retention requirement for investigational drug studies is clearly outlined in 21 CFR 312.57 (c) and 21 CFR 312.62(c).

\* 21 CFR 312.57(c): "A sponsor shall retain the records and reports... for 2 years after a marketing application is approved for the drug; or, if an application is not approved, until 2 years after shipment and delivery of the drug for investigational use is discontinued and FDA has been so notified."

\* 21 CFR 312.62(c): Investigators also must retain study-related records for 2 years following the date a marketing application is

