

Valid Braindumps ISO-9001-Lead-Auditor Ebook - Accurate ISO-9001-Lead-Auditor Test

Audit evidence	ISO 9001 Clause 8.3 extract
Half of all new products launched in the past 12 months were late. The NPD Manager explains he has not got enough people on his team to cope with the demand for new products.	"8.3.2 d) ... internal ... resource needs for the design and development of products ..."
The NPD Manager explains many changes are made to cosmetic formulations during product development owing to retailer feedback. Only when confirmed by the retailer is the agreed formulation documented on SWIFT.	"8.3.5 ... retain documented information ..."
The NPD Manager explains that the customer confirms their approval to proceed with a new formulation by email. These emails are kept on SWIFT.	"8.3.5 ... retain documented information ..."
The NPD Manager shows you evidence of consumer trials that are carried out for some new products prior to full-scale launch.	"8.3.4 d) ... conducted to ensure that the design and development outputs meet ..."
The NPD Manager explains that an approved external laboratory is used to perform shelf-life stability trials on some formulations during product development.	"8.3.2 e) ... external ... resource needs for the design and development of products ..."

To complete the table click on the blank section you want to complete so it is highlighted in red and then click on the ISO 9001 clause 8.3 extracts listed below. Alternatively, drag and drop each clause to the audit evidence that applies.

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PECB ISO-9001-Lead-Auditor Exam Syllabus Topics:

Topic	Details
Topic 1	<ul style="list-style-type: none"> Managing an ISO 9001 audit program: This topic evaluates your abilities to establish and managing a QMS audit program.
Topic 2	<ul style="list-style-type: none"> Quality management system (QMS) requirements: It assesses your abilities to point out and explain different requirements for a quality management system based on ISO 9001.
Topic 3	<ul style="list-style-type: none"> Preparing an ISO 9001 audit: This topic covers sub-topics related to preparing a quality management system audit.
Topic 4	<ul style="list-style-type: none"> Closing an ISO 9001 audit: The topic focuses on concluding a QMS audit and conducting audit follow-up activities.
Topic 5	<ul style="list-style-type: none"> Conducting an ISO 9001 audit: It evaluates your skills to conduct a QMS audit.
Topic 6	<ul style="list-style-type: none"> Fundamental principles and concepts of a quality management system: The main objective of this domain is to evaluate your skills of explaining and applying ISO 9001 principles and concepts.

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PECB QMS ISO 9001:2015 Lead Auditor Exam Sample Questions (Q220-Q225):

NEW QUESTION # 220

You are a member of the audit team of a second-party audit of an organisation with 625 employees. The audit procedure recommends using sampling criteria which requires the review of the documented competence for 25 personnel. The audit team leader developed an audit plan allocating one hour to audit the Human Resources department (from 11:30 am to 12:30 pm). She told you that she could not allocate any additional time. What would you do?

- A. Plan to miss lunch and review as many as possible.
- **B. Plan to review less than 25 cases.**
- C. Extend the audit until 1.00pm and ask for a quick lunch later.
- D. Plan to review as many as possible and see if you can extend the audit duration by one day.

Answer: B

Explanation:

In this scenario, the time allocated by the audit team leader for the Human Resources audit is fixed, and as an auditor, you must work within that constraint. Although the sampling criteria suggests reviewing 25 personnel files, it is acceptable to adjust the sample size based on time and resource limitations. ISO 9001:2015 emphasizes risk-based thinking and practical resource management (Clause 7.1), so it is reasonable to review a smaller sample if the time is insufficient.

Option B is a pragmatic approach, allowing you to focus on quality over quantity by reviewing as many cases as time allows without compromising the audit schedule.

Options like extending the audit (A, C, D) are impractical in a structured audit environment, especially for second-party audits where maintaining the agreed schedule is important.

NEW QUESTION # 221

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics Dispatch. The audit team includes two other internal auditors.

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics / Dispatch. The audit team includes two other internal auditors.

*If practicable

*You should not ...

You need not ...

You must ...

You must not ...

You should ...

To complete the sentences click on the blank section you want to complete so it is highlighted in red and then click on the applicable text from the options below. Alternatively, drag and drop the options to the appropriate blank section.

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Answer:

Explanation:

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics / Dispatch. The audit team includes two other internal auditors.

*If practicable

*You should not ...

audit production

You need not ...

change the audit team

You must ...

raise audit findings if necessary

You must not ...

send the audit report to the Quality Manager

You should ...

carry out a formal opening meeting

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audit production

carry out a formal opening meeting

raise audit findings if necessary

change the audit team

send the audit report to the Quality Manager

Explanation:

Here is the correct matching of actions to the statements in the context of leading the internal audit:

* If practicable carry out a formal opening meeting

* You should not audit production (as you are a supervisor in that area, and this would compromise audit objectivity)

* You need not change the audit team (unless there is a specific reason, such as conflict of interest)

* You must raise audit findings if necessary (this is a key responsibility of an auditor when nonconformities are found)

* You must not send the audit report to the Quality Manager (the audit report must be reviewed first; it is typically part of the internal audit process to go through necessary channels before final submission)

* You should send the audit report to the Quality Manager (after appropriate reviews and approvals) This reflects key principles of conducting an internal audit according to ISO 9001:2015, ensuring objectivity, proper documentation, and clear reporting procedures.

NEW QUESTION # 222

Scenario 2:

Bell is a Canadian food manufacturing company that operates globally. Their main products include nuts, dried fruits, and confections. Bell has always prioritized product quality and has maintained a good reputation for many years. However, the company's production error rate increased significantly, leading to more customer complaints.

To increase efficiency and customer satisfaction, Bell implemented a Quality Management System (QMS) based on ISO 9001. The top management established a QMS implementation team comprising five middle managers from various departments, including Leslie, the quality manager.

Leslie was responsible for assigning responsibilities and authorities for QMS-related roles. He also suggested including a top management representative in the QMS team, but top management declined due to other priorities.

The team defined the QMS scope as:

"The scope of the QMS includes all activities related to food processing." Leslie established a quality policy and presented it to the team for review before top management approval

. Top management also proposed a new strategy for handling customer complaints, requiring biweekly customer surveys to monitor customer perceptions.

The quality policy was established by Leslie and approved by top management. Is this acceptable?

Please refer to scenario 2.

- A. No, the quality policy must be established and approved only by the quality manager.
- **B. No, the quality policy must be established and approved by top management.**
- C. Yes, as long as top management is informed, the policy can be established by any responsible employee.
- D. Yes, the quality policy can be established by the QMS implementation team and be approved by top management.

Answer: B

Explanation:

Comprehensive and Detailed In-Depth Explanation: ISO 9001:2015, Clause 5.2.1 (Establishing the Quality Policy) states that top management must establish, implement, and maintain a quality policy.

In the scenario, the quality manager (Leslie) created the policy, but top management did not establish it themselves, which violates Clause 5.2.1. While the policy can be drafted by a team, top management must take full ownership of its development and approval.

NEW QUESTION # 223

Which of the following is correct with regard to the internal audit?

- A. It may be conducted on an ongoing basis
- B. It has no advisory role within the organization for the improvement of the QMS
- C. It considers only the effectiveness of the QMS

Answer: A

Explanation:

Comprehensive and Detailed In-Depth Explanation:

According to ISO 9001:2015, Clause 9.2 (Internal Audit):

* Internal audits can be conducted on an ongoing basis as part of continual improvement.

* Audits consider both conformity and effectiveness (A is incorrect).

Thus, C is the correct answer.

Reference:

ISO 9001:2015, Clause 9.2 (Internal Audit)

NEW QUESTION # 224

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics Dispatch. The audit team includes two other internal auditors.

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics / Dispatch. The audit team includes two other internal auditors.

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You must not ...

You should ...

To complete the sentences click on the blank section you want to complete so it is highlighted in red and then click on the applicable text from the options below. Alternatively, drag and drop the options to the appropriate blank section.

Answer:

Explanation:

You are the supervisor in Production of a medium size manufacturing organisation. You are qualified as an internal auditor. The Quality Manager asks you to lead the next internal audit of Production and Logistics / Dispatch. The audit team includes two other internal auditors.

*If practicable
 *You should not ...
 You need not ...
 You must ...
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To complete the sentences click on the blank section you want to complete so it is highlighted in red and then click on the applicable text from the options below. Alternatively, drag and drop the options to the appropriate blank section.

Explanation:

Here is the correct matching of actions to the statements in the context of leading the internal audit:

- * If practicable carry out a formal opening meeting
- * You should not audit production (as you are a supervisor in that area, and this would compromise audit objectivity)
- * You need not change the audit team (unless there is a specific reason, such as conflict of interest)
- * You must raise audit findings if necessary (this is a key responsibility of an auditor when nonconformities are found)
- * You must not send the audit report to the Quality Manager (the audit report must be reviewed first; it is typically part of the internal audit process to go through necessary channels before final submission)
- * You should send the audit report to the Quality Manager (after appropriate reviews and approvals) This reflects key principles of conducting an internal audit according to ISO 9001:2015, ensuring objectivity, proper documentation, and clear reporting procedures.

NEW QUESTION # 225

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