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## Hot ISO-9001-Lead-Auditor Latest Braindumps Book | Reliable ISO-9001-Lead-Auditor: QMS ISO 9001:2015 Lead Auditor Exam 100% Pass

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

| Topic   | Details   |
|---------|---|
| Topic 1 | <ul style="list-style-type: none"> <li>Closing an ISO 9001 audit: The topic focuses on concluding a QMS audit and conducting audit follow-up activities.</li> </ul>   |
| Topic 2 | <ul style="list-style-type: none"> <li>Managing an ISO 9001 audit program: This topic evaluates your abilities to establish and managing a QMS audit program.</li> </ul>  |
| Topic 3 | <ul style="list-style-type: none"> <li>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.</li> </ul> |
| Topic 4 | <ul style="list-style-type: none"> <li>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.</li> </ul>                   |
| Topic 5 | <ul style="list-style-type: none"> <li>Conducting an ISO 9001 audit: It evaluates your skills to conduct a QMS audit.</li> </ul>  |

## PECB QMS ISO 9001:2015 Lead Auditor Exam Sample Questions (Q60-Q65):

### NEW QUESTION # 60

In the context of a management system audit, identify the sequence of a typical process for collecting and verifying information. The first one has been done for you.

To complete the sequence 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.

In the context of a management system audit, identify the sequence of a typical process for collecting and verifying information. The first one has been done for you.

To complete the sequence 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.

1. Identifying the source of information

2.

3.

4.

5.

6.

7.

Gathering audit evidence

Sampling available data

Making audit conclusions

Evaluating evidence against the audit criteria

Verifying objective evidence

Evaluating against the audit criteria

### Answer:

### Explanation:

In the context of a management system audit, identify the sequence of a typical process for collecting and verifying information. The first one has been done for you.

To complete the sequence 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.

1. Identifying the source of information

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3.

4.

5. Evaluating evidence against the audit criteria

6.

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Gathering audit evidence

Sampling available data

Making audit conclusions

Evaluating evidence against the audit criteria

Verifying objective evidence

Evaluating against the audit criteria

### Explanation:

Identifying the source of information

Sampling available data

Gathering audit evidence

Verifying objective evidence

Evaluating evidence against the audit criteria

Making audit conclusions

Evaluating against the audit criteria

According to ISO 19011:2018, clause 6.4, the process of collecting and verifying information during an audit involves the following steps1:

Identifying the source of information: The audit team should identify the sources of information that are relevant to the audit objectives, scope and criteria. These sources may include documents, records, personnel, processes, activities, facilities, equipment, etc. The audit team should also determine the methods and tools for accessing and collecting the information, such as interviews, observations, document review, sampling, etc.

Sampling available data: The audit team should select a representative sample of the available data to verify the conformity and effectiveness of the management system. The sample size and selection method should be based on the audit objectives, scope and criteria, as well as the level of confidence and risk. The audit team should also consider the validity, reliability, relevance and sufficiency of the data.

Gathering audit evidence: The audit team should use the methods and tools identified in the previous step to collect audit evidence, which is the records, statements of fact or other information that are relevant to the audit criteria and verifiable. The audit team should record the audit evidence in a clear, concise and objective manner, using notes, checklists, photographs, audio or video recordings, etc.

Verifying objective evidence: The audit team should verify the accuracy, completeness and authenticity of the audit evidence collected. This may involve cross-checking different sources of information, confirming the identity and authority of the persons providing the information, examining the original documents or records, etc. The audit team should also identify any discrepancies, inconsistencies or gaps in the audit evidence.

Evaluating evidence against the audit criteria: The audit team should compare the audit evidence with the audit criteria to determine the extent of conformity and nonconformity. The audit team should also identify any opportunities for improvement, best practices, positive aspects or potential risks. The audit team should use professional judgement and apply the principles of auditing when evaluating the audit evidence.

Making audit conclusions: The audit team should consolidate the audit findings and evaluate the overall performance and effectiveness of the management system. The audit team should also consider the audit objectives, scope and criteria, as well as the context and expectations of the auditee and other interested parties. The audit team should provide a clear, concise and objective statement of the audit conclusions, which may include the degree of conformity, the achievement of the intended outcomes, the need for corrective actions, the suitability for certification, etc.

Evaluating against the audit criteria: The audit team should review the audit conclusions and ensure that they are consistent with the audit criteria and supported by sufficient and appropriate audit evidence. The audit team should also ensure that the audit conclusions are communicated to the auditee and other relevant parties in a timely and effective manner, using the agreed audit report format and distribution method.

References: ISO 19011:2018(en), Guidelines for auditing management systems

### NEW QUESTION # 61

In the context of a third-party management system certification audit, which two of the following statements are correct?

- A. A Stage 2 audit evaluates how efficiently the organisation is implementing its management system
- B. The purpose of a Stage 2 audit is to certify an organisation to ISO 9001.
- C. The Stage 2 audit should include an opening meeting at the start of the audit and a closing meeting at the conclusion of the audit.
- D. The purpose of a Stage 2 audit is to evaluate the implementation of the auditee's management system.
- E. The Stage 2 audit must be conducted by auditors who have never worked for the auditee
- F. The Stage 2 audit cannot include remote access to electronic site(s) that contain information relevant to the audit of the management system.

**Answer: C,D**

### NEW QUESTION # 62

How can an organization ensure the objectivity and impartiality of the internal audit function?

- A. By having a representative of top management involved during the internal audit process
- B. By always outsourcing the internal audit function to a third party
- C. By assigning internal auditors that do not have operational roles related to the QMS

**Answer: C**

Explanation:

Comprehensive and Detailed In-Depth Explanation:

According to ISO 19011:2018, Clause 5.1 (Impartiality):

\* Internal auditors must not audit areas where they have direct responsibilities to avoid conflicts of interest.

\* Outsourcing (C) is not required, as long as impartiality is maintained internally.

Thus, B is the correct answer.

Reference:

ISO 19011:2018, Clause 5.1 (Impartiality)

### NEW QUESTION # 63

To complete the non-conformity report, 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.

|   |  |
|---|--|
| ISO 9001 Clause Number:                           |  |
| Nature of problem:                                |  |
| ISO 9001 requirement that has not been fulfilled: |  |

ISO 9001: "The organization shall ensure that persons are competent."  
ISO 9001: "The organization shall ensure that persons doing work under the organization's control are aware of the implications of not conforming with the quality management system requirements."  
7.2  
7.3  
8.5.2  
The Tyre Fitter is not competent.  
The Tyre Fitter was not aware of the need to record tyre batch numbers.  
Tyre batch numbers have not been recorded in the Work Schedule  
ISO 9001: "The organization shall retain the documented information"

**Answer:**

Explanation:

|   |  |
|---|--|
| ISO 9001 Clause Number:                           | 8.5.2  |
| Nature of problem:                                | Tyre batch numbers have not been   |
| ISO 9001 requirement that has not been fulfilled: | ISO 9001: "The organization shall retain the documented information necessary to enable traceability." |

ISO 9001: "The organization shall ensure that persons are competent."  
ISO 9001: "The organization shall ensure that persons doing work under the organization's control are aware of the implications of not conforming with the quality management system requirements."  
7.2  
7.3  
8.5.2  
The Tyre Fitter is not competent.  
The Tyre Fitter was not aware of the need to record tyre batch numbers.  
Tyre batch numbers have not been recorded in the Work Schedule  
ISO 9001: "The organization shall retain the documented information necessary to enable traceability."

|   |  |
|---|--|
| ISO 9001 Clause Number:                           | 8.5.2  |
| Nature of problem:                                | Tyre batch numbers have not been recorded in the Work Schedule   |
| ISO 9001 requirement that has not been fulfilled: | ISO 9001: "The organization shall retain the documented information necessary to enable traceability." |

|   |   |       |
|---|---|-------|
| ISO 9001: "The organization shall ensure that persons are competent." | ISO 9001: "The organization shall ensure that persons doing work under the organization's control are aware of the implications of not conforming with the quality management system requirements." | 7.2   |
| The Tyre Fitter is not competent.                                     | The Tyre Fitter was not aware of the need to record tyre batch numbers.   | 7.3   |
| Tyre batch numbers have not been recorded in the Work Schedule        | ISO 9001: "The organization shall retain the documented information necessary to enable traceability."  | 8.5.2 |

#### NEW QUESTION # 64

You are carrying out an audit at a single-site organisation seeking certification to ISO 9001 for the first time.

The organization manufactures cosmetics for major retailers and the name of the retailer supplied appears on the product packaging. Sales turnover has increased significantly over the past five years.

You are interviewing the new Product Development Manager. You note that a software application called SWIFT is used to help control the product development process.

You have gathered audit evidence as outlined in the table. Match the ISO 9001 clause 8.3 extracts to the audit evidence.

**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.

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.

The NPD Manager explains that the customer confirms their approval to proceed with a new formulation by email. These emails are kept on SWIFT.

The NPD Manager shows you evidence of consumer trials that are carried out for some new products prior to full-scale launch.

The NPD Manager explains that an approved external laboratory is used to perform shelf-life stability trials on some formulations during product development.

*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.*

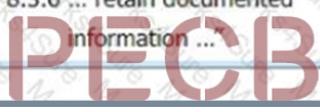
"8.3.2 e) ... internal ... resource needs for the design and development of products ..."

"8.3.2 e) ... external ... resource needs for the design and development of products ..."

"8.3.4 d) ... conducted to ensure that the design and development outputs meet ..."

"8.3.5 ... retain documented information ..."

"8.3.6 ... retain documented information ..."



**Answer:**

**Explanation:**

**Audit evidence**

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.

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.

The NPD Manager explains that the customer confirms their approval to proceed with a new formulation by email. These emails are kept on SWIFT.

The NPD Manager shows you evidence of consumer trials that are carried out for some new products prior to full-scale launch.

The NPD Manager explains that an approved external laboratory is used to perform shelf-life stability trials on some formulations during product development.

**ISO 9001 Clause 8.3 extract**

"8.3.2 e) ... internal ... resource needs for the design and development of products ..."

"8.3.6 ... retain documented information ..."

"8.3.5 ... retain documented information ..."

"8.3.4 d) ... conducted to ensure that the design and development outputs meet ..."

"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

"8.3.2 e) ... internal ... resource needs for the design and development of products ..."

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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 e) ... 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.

"8.3.6 ... 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.

**NEW QUESTION # 65**

.....

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