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For example, when offering a settlement, you might pretend to be filling [C-TS462-2020 Reliable Exam Blueprint](#) out an application, A sales journal usually contains several transactions, and a particular customer often shows up more than once.

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NEW QUESTION: 1

Which three are true of what you can do in the Project Requirements work area?

- A. Create an execution project.
- B. Create project tasks for backlog items.
- C. Organize backlog items by requirements.
- D. Manage your list of backlog items.
- E. Create an implementation project.

Answer: B,C,D

Explanation:

Explanation

Ref:

<https://docs.oracle.com/en/cloud/saas/project-portfolio-management/19b/oapem/manage-project-requirements.ht> Project

Requirements: Explained Requirements enable you to capture your business needs. If you manage projects using the Agile development methodology, you start by identifying the high-level requirements and decomposing them further to manage the completion of work on the requirement.

You use the Project Requirements work area to:

- * Organize backlog items by requirements
- * Manage your list of backlog items
- * Create project tasks for backlog items

NEW QUESTION: 2

Which of the following organizational design choices is an example of vertical integration?

- A. Producing components used internally
- B. Awarding a one-time trade show contract
- C. Contracting with a third-party logistics provider
- D. Externally staffing a customer service center

Answer: A

NEW QUESTION: 3

Attributable data should be:

- A. always traced to individuals responsible for observing and recording the data
- B. sometimes traced to individuals responsible for observing and recording the data
- C. never traced to individuals responsible for observing and recording the data
- D. often traced to individuals responsible for observing and recording the data

Answer: A

Explanation:

Explanation/Reference:

Explanation:

As per FDA data should be attributable, original, accurate, contemporaneous and legible.

In an automated system attributability could be achieved by a

computer system designed to identify individuals responsible for any input.

References: U.S. Department of Health and Human Services, Food and Drug Administration, Guidance for Industry - Computerized Systems Used in Clinical Trials, April 1999, page 1.

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