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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q332-Q337):

**NEW QUESTION # 332**
Which is the most appropriate disposal method for rejected raw materials?

- A. Return to vendor or destroy

- B. Use for training
- C. Give to another manufacturer
- D. Store indefinitely

**Answer: A**

**NEW QUESTION # 333**
A major change in a validated process requires:

- A. No action if risk is low
- B. Change control and potential re-validation
- C. Engineering notification only
- D. Approval but no documentation

**Answer: B**

**NEW QUESTION # 334**
The impact analysis for a proposed change should NOT:
Response:

- A. Ignore potential impacts on regulatory compliance
- B. Consider only the positive outcomes
- C. Exclude input from external consultants
- D. Be limited to immediate effects

**Answer: B**

**NEW QUESTION # 335**
Regulatory requirements for compressed air and gas systems include all the following EXCEPT:
Response:

- A. Storage and flow regulation
- B. Filtration and venting
- C. Purging requirements
- D. The color coordination of piping systems

**Answer: D**

**NEW QUESTION # 336**
For tablet granulation and compression, critical process parameters include:
Response:

- A. Social media trends
- B. Color of the granules
- C. Granule size and compression force
- D. Preferences of the production team

**Answer: C**

**NEW QUESTION # 337**
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