## **INTRODUCTION TO MS ISO/IEC 17025**

## **EXERCISE 1**

1. A testing lab shall comply with this International Standard and regulatory authorities under which it operates

Answer : Yes / No

Reference:

2. Only permanent lab can be accredited to this International Standard.

Answer: Yes / No

Reference:

3. Must the organization and management structure of the lab be properly defined?

Answer: Yes / No

Reference:

4. Management quality policy and objectives shall only cover management's commitment to comply with this International Standard.

Answer: Yes / No

Reference:

5. The laboratory is required to prepare a document called "quality manual", however named.

Answer: Yes / No

Reference:

6. The roles and responsibilities of Technical and Quality Manager shall be defined in the quality manual

Answer: Yes / No

Reference:

7. Records of review of requests, tenders and contracts shall be maintained.

Answer: Yes / No

Reference:

8. The lab shall control all type of documents in the lab

Answer: Yes / No

Reference:

9. Customers must agree in writing for sub-contract test/calibration.

Answer: Yes / No

Reference:

10. Document review and approval shall be done by the same function that performed the original review unless specifically designated otherwise.

Answer: Yes / No

Reference:

11. The laboratory shall maintain documented procedure on how they wish to provide service to the customers.

Answer: Yes / No

Reference:

12. Amendment of document by hand is not allowed

Answer: Yes / No

Reference:

13. Review of request, tenders and contract only covers method of test

Answer: Yes / No

Reference:

14. The testing laboratory shall be also responsible to review work that the laboratory subcontracts.

Answer: Yes / No

Reference:

15. The laboratory shall only evaluate suppliers who provide supplies and services which can affect the quality of test/calibration results

Answer: Yes / No

Reference: