



FOOD AND DRUGS AUTHORITY

DOC. TYPE: FORM

DOC NO.: FDA/FER/RQT - 07

Ver. No.: 01

Effective Date: 01/06/2020

Page 1 of 1

CERTIFICATE OF ANALYSIS REPORTING FORMAT

Each test report shall include at least the following information:

- 1. A Title (e.g. Analytical Test Report)
- 2. Name and address of laboratory
- 3. Location of performance of laboratory activity
- 4. Name of Product
- 5. Unique identification of the Test Report
- 6. The name/ address of the client
- 7. Manufacturer
- 8. Identification of the method used
- 9. Temperature condition of testing area.(where applicable)
- 10. A description of, the condition of, and identification of the sample.
- 11. Date of receipt of sample
- 12. Date(s) on which test was conducted
- 13. Units of Measurement, where appropriate
- 14. Identification of person(s) authorizing the test report
- 15.A statement to the effect that the results relate only to the product tested where applicable
- 16. Where relevant, a statement on compliance/non-compliance with requirements and/or specifications
- 17. Where applicable or when requested by the client, a statement on the estimated uncertainty of measurement.
- 18. Where appropriate and needed, opinions and interpretations.