



SAMPLES REPOSITORY OFFICE

SAMPLE SUBMISSION FORM

Part A: Customer Details

1.0 Customer Name: _____

2.0 Customer Address: _____

3.0 Contact Person: _____

4.0 Designation: _____

5.0 Contact: Mobile No. _____ Telephone No. _____

6.0 E-mail: _____

Part B: Product Details

7.0 Product Generic Name: _____

7.1 Brand Name: _____

7.2 Pharmacological classification (*e.g. anti-malarial*): _____

8.0 Manufacturer: _____

8.1 Manufacturer's address: _____

8.2 Date of Manufacture: _____

8.3 Date of Expiry: _____

8.4 Batch Number: _____

8.5 Country of Origin: _____

8.6 Zimbabwe Registration Number (*if any*): _____

9.0 Sampling Site: _____

9.1 State/Province/City: _____

9.2 Storage conditions: Temperature _____ Humidity _____

9.3 Sample size: _____

10.0 Transportation conditions (*e.g. cold chain samples*): _____

Part C: Analysis Required**11.0 CHEMICAL ANALYSIS**

YES

NO

CUSTOMER REF NO:					
Details of Analysis Required <i>(Please insert (✓) or (X) in the applicable box)</i>					
	Test	Yes(✓) or No (X)		Test	Yes(✓) or No (X)
a.	Appearance		b.	pH	
c.	Identity		d.	Assay	
e.	Uniformity of Mass		f.	Dissolution	
g.	Disintegration		h.	Uniformity of Content	
i.	Friability		j.	Heavy Metals	
k.	Hardness		l.	Moisture Content	
m.	Decomposition Products		n.	Related Substances	
o.	Other test <i>(please specify)</i> :				

Please note: Tests performed, including quality performance tests (minimum test parameters), may be different from those indicated on the quotation (or on this form) depending on whether the products are USP, BP or Int'l Ph.

12.0 Any Other Comments:

13.0 MICROBIOLOGICAL ANALYSIS:

YES	NO
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CUSTOMER REF NO:							
Details of Analysis Required <i>(Please insert (✓ or X) in the applicable box)</i>							
	Test	Yes (✓)	No (X)		Test	Yes (✓)	No (X)
a.	Appearance			b.	Pyrogen (LAL/BET)		
c.	Microbial Limit (Microbiological Examination of Non-Sterile Products)			d.	Microbial Assay		
e.	Sensitivity Test			f.	Sterility Test		
g.	Preservative Efficacy			h.	Other Test <i>(please specify)</i>		
i.	Other test <i>(please specify)</i> :						

14.0 Any Other Comments:

PART D: TERMS AND CONDITIONS

1 Privacy and Confidentiality

- 1.1 For purposes of these terms and conditions, “Representatives” means MCAZ Laboratory’s officers, employees, partners, agents or other representatives (including, without limitation, its attorneys, accountants, auditors, assessors, consultants and financial advisers).
- 1.2 The MCAZ laboratory recognizes the importance of privacy and shall be responsible for the management of all information obtained or created during the performance of laboratory activities.
- 1.3 The MCAZ laboratory shall hold the information disclosed to it in strict confidence and shall use the confidential information only for testing activities.
- 1.4 The MCAZ Laboratory shall not, without your prior written consent, disclose your Confidential Information to any person, party or entity except to such of its representatives who need such information to perform duties connected with the testing activities, and who are bound by an obligation of confidentiality consistent with these terms and conditions.
- 1.5 The MCAZ laboratory shall exercise reasonable care to protect your information including your test data and results from unauthorized use and disclosure.
- 1.6 Exceptions: MCAZ Laboratory shall not be liable for disclosure or use of Confidential Information that:
 - 1.6.1 at the time of disclosure was in the public domain, or subsequently became part of the public domain otherwise than by breach of these terms and conditions;
 - 1.6.2 you authorized in writing to be disclosed; or
 - 1.6.3 is being disclosed in compliance with a Court Order or an order of any other competent body, or as required by the Medicines and Allied Substances Control Act (Chapter 15:03) or any other law, provided that the MCAZ laboratory shall first give you written notice of such disclosure and further provide that the MCAZ laboratory limits the disclosure to the least that is legally required.

2 Process requirements

- 2.1 In the event that you request a statement of conformity to a specification or standard for the test (e.g. pass/fail, complies/does not comply) the MCAZ Laboratory shall use the defined specification as stated in the method used to conduct the test. However, where a non-standard method is used, the decision rule will apply. MCAZ laboratory shall communicate to you the decision rule selected upon submission of the test results.
- 2.2 For samples submitted for testing, the Certificate of Analysis (Test report) shall be according to MCAZ standard format which includes customer details, MCAZ details, product/sample details, method of analysis, tests performed, specifications, opinions and interpretations, deviations, results and comment. If you require additional information, it shall be communicated on submission of the sample.

By signing below you agree to the terms and conditions stated above.

Signature

Printed name

Date

FOR MCAZ USE ONLY

Sample Received by: **Name** _____ **Date** _____

MCAZ Ref. No: _____