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### **INITIAL FACTORY INSPECTION REPORT**

#### NOTE

- 1. Section 1 shall be completed by the applicant and shall be returned together with the application form STA/1
- 2. Separate sheets may be included where necessary.
- 3. The provided spaces can be enlarged to capture all the details
- 4. The statements shall relate to the facilities available as of the date of completion of the form.
- 5. The information provided in this document shall be treated in Confidence.

# **SECTION 1 (TO BE COMPLETED BY THE APPLICANT)**

A.	FIRM DET	TAILS				
1.	Name of t	he Firm:				
2.	Company	Registration Number				
3.						
4.						
5.	Authorize	d Company Representative:				
6.	Company	Email Address:				
7.	Physical	Location:				
8.	Town					
9.	Total No.	of Personnel:				
10.	Compete	ncy of key personnel in charge o	f Production and Quality Control			
	S/no	Name	Qualifications Relevant (Academic/professional/experience)	to	the	Sector
	i.					
	ii.					
	iii.					

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#### B. PRODUCTION CONTROLS

#### 1. Raw Materials

Specify the type of raw materials and origin (attach their specification and testing records where applicable):

S/N O.	Product	Raw materials	Origin/Supplier	Raw materials Specifications e.g. (Standard, company specification,)	Quality checks (attach records)
1					
2					
3					
4					
5					
6					

## 2. Machinery and Plant

S/NO.	Machine and Plant	Process/ Usage	
1.			
2.			
3.			
4.			
5.			
6.			

### 3. Manufacturing process

(Attach a flowchart showing sampling/quality control points and relevant evidences where possible)

S/No.	Production process flow/ Operations	Critical Process parameters Monitored	Frequency	Process monitoring records
1.				

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2.		
3.		
4.		
5.		
6.		

4.	How do you handle nonconformities at the following manufacturing process steps:
	i. Raw materials step
	ii. In-process step
	iii.Final product step
5.	What mechanism do you have for recalling/withdrawing nonconforming products from the market?

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low do you s			
	store the following?		
Raw materials	S?		
End product?			
	·		
		testing equipment exist on your premises	
What monitor	ring, measuring and/or	testing equipment exist on your premises  Parameters Monitored /Measured/	Calibration/Verification
		Parameters Monitored /Measured/	Calibration/Verification
S/No.		Parameters Monitored /Measured/	Calibration/Verification
S/No.		Parameters Monitored /Measured/	Calibration/Verification
S/No. 1 2		Parameters Monitored /Measured/	Calibration/Verification
S/No.  1 2 3		Parameters Monitored /Measured/	Calibration/Verification
S/No.  1 2 3 4	Equipment	Parameters Monitored /Measured/ Tested	Calibration/Verification Status
S/No.  1 2 3 4	Equipment	Parameters Monitored /Measured/	Calibration/Verification Status
S/No.  1 2 3	Equipment	Parameters Monitored /Measured/ Tested	Calibration/Verification Status
S/No.  1 2 3	Equipment	Parameters Monitored /Measured/ Tested	Calibration/Verification Status
S/No.  1 2 3	Equipment	Parameters Monitored /Measured/ Tested	Calibration/Verification Status

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## 9. Marking and Labelling:

Is your product labelled and marked in compliance to the relevant standard? Specify and attach label(s) as applicable

S/No	Standard Marking/labelling requirement	Indicated/Not Indicated
1		
2		
3		
4		
5		

10.	Handling of Complaints
	How do you handle consumer complaints? Attach evidence of complaints records.
	Authorized Company Representative
	Designation
	Signature
	Date

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# SECTION 2 (FOR OFFICIAL USE ONLY)

1.

## A. INSPECTOR'S REPORT AND RECOMMENDITIONS

State	the adequacy of the following:
i.	Production facility/premises:
ii.	Plant layout:
iii.	Suitability of location:
iv.	Suitability of equipment
٧.	Cleaning and maintenance programmes

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Describe

	••••					
3.			provided with necessary wor	k wear?		
	De	scribe				
	••••					
	••••					
	••••					
	••••					
4.				statutory/regulatory requirements	(e.g. Public Health Act,	
	Oŀ	ISA requir	rements)?			
	De	scribe				
5.	Ma	rking and	Labelling:			
		S/No	Standard Marking/labelling requirements	Compliance Status (Y/N)	Remarks	
	•	1				
	•	2				
	•	3				
	•	4				

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6.	En	Environmental considerations				
	i.	Have the potential sources of contamination to the product(s) been identified? How are they managed?				
	ii.	How are wastes emanating from the process handled?				
7.	Re	commendations/Conclusions findings on processes				
	i.	Specify areas noted for Improvement i.e. nonconformities (attach separate sheet as necessary)				
	ii.	Inspector's recommendation for certification?				

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S/No.	Inspector's Name	Designation	Signature
1.			
2.			
3.			

Date	 	 
Supervisor's comments:		
Name a		
Name	 	 
Designation	 	 
Signature	 	 
Data		