APSOPCA/ 03.c Organic System Plan for Processor

Revision number: 01

Revision date : 07.02.2023

#### ORGANIC SYSTEM PLAN FOR PROCESSOR

S. No	Particulars Particulars	
1	Name of the Company / Organization	
1.1	Name & address of the	
1.1	Responsible Person	
1.2	Address for Communication	
1.3	Pin code	
1.4	Address of the Processing	
1.4	unit	
1.5	License number & Validity	
1.6	Telephone No.	
1.7	Email Id	
1.8	Year of establishment of	
1.0	company	
	Total No. of employees	
1.9	Name of person over seeing organic	
1.10	Number of products Processed:	
1.11	Do you have a copy of NPOP standards (Yes / No)	
1.12	Mention the activity in the production subcontracted to another organization?	

2. List the brand name and the technical name of the products being processed

S.No	Brand name	Technical name	

- 3. What is the quality management system (QMS) followed by the organization?
- 4. Give the list of all ingredients, additives and processing aids used for each product with their ratios:
- 5. Give the list of all your approved suppliers (Raw material and packing material)
- 6. Assurance of Organic Integrity:
- a. Do operator has a quality assurance programme
- b. Products testing programme

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A DCODO	APSOPCA/ 03.c Organic System Plan for Processor		Revision number: 01				
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	I.	Products tested during crop prod	luction				
	II. Ingredients tested prior to purchase						
	III.	III. Ingredients tested upon receipt					
	IV. Finished products tested						
	V.	If any other (Specify)					
7. V	What a	are the quality parameters checked	d to procure each of your raw materials				
t	that go into processing?						
	Name of the Ingredient Quality Parameter Checked						

8. Give a brief description of the production process for each of your product

9. List the equipments used in processing

Type of Equipment	Capacity	Check if equipment is cleaned prior to process	Check if cleaning process is documented

## 10. In process:

a. What are the measures taken to clean the process unit prior to the organic products load? (Is the inspection/cleaning process documented?)

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LIYES	- 11	Nα

b. Are organic products are shipped at the same time as non-organic products.
 If yes checks taken to segregate organic and non-organic

- i. Separate area in transport unit
- ii. Separate sealed container to organic product
- iii. Organic product shrink wrapped

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iv. Any other method

# 11. Pest Management:

- a. What kind of Pest Management mechanism do operator use
- b. Are records kept for pest monitoring mechanisms and pesticides applied for controlling
- c. In case of pesticides applications what are the measures taken to avoid the organic product to come into with organic material/ingredients
- 12. Give the annual estimated quantity of production for each product:
- 13. Describe the precautions taken to prevent Comingling of organic and conventional products

#### 14. Sanitation:

# Check all Cleaning Methods used:

- 1. Sweeping
- 2. Scraping
- 3. Vacuuming
- 4. Compressed Air
- 5. Manual washing
- 6. Clean in place (CIP)
- 7. Steam cleaning
- 8. Sanitizing
- 9. Other (specify)

### 15. **Storage:**

Provide information about storage areas

Use	Location	Type/Capacity	Identification Name Or Number	Is Storage Unit Dedicated Organic?	Possibilities of Contamination Or Problems
Ingredients					
storage					
Packing					
material					
In-					
processed					
storage					
Finished					
product					

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Off-site			
storage			
Other			
(specify)			

- 16. What are the storage conditions for your products?
- 17. What is the packing material used for the finished product? Eco-friendly /re used material
- 18. What are the packing quantities for each of the product manufactured?
- 19. Record Keeping
- 20. Which of the following records do operators keep for organic processing/handling?

## Incoming:

- 1. Purchase orders
- 2. Contracts
- 3. Invoices
- 4. Receipts
- 5. Bills of lading
- 6. Customs forms
- 7. Scale tickets
- 8. Quality test results
- 9. Certificates of Analysis
- 10. Transaction Certificates
- 11. Copies of Certificates of Organic Operation
- 12. Verification of non-GMO ingredients
- 13. Verification of ingredients produced not using sewage sludge
- 14. Verification of ingredients produced/handled without ionizing radiation
- 15. Documentation that organic ingredients are not commercial available, when using nonorganic ingredients in products labeled as "100% organic" and/or "organic"
- 16. Receiving records
- 17. Receiving summary log (12 mos.)
- 18. Other (specify)

### **In-Process:**

- 1. Ingredient inspection forms
- 2. Blending reports
- 3. Production reports
- 4. Equipment clean-out logs
- 5. Sanitation logs
- 6. Packaging reports
- 7. QA reports
- 8. Production summary records (12 mos.)

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9. Other (specify)

## Storage:

- 1. Ingredient inventory reports
- 2. Finished product inventory reports
- 3. Other (specify)

# Outgoing:

- 1. Shipping log
- 2. Transport unit inspection/cleaning forms
- 3. Bills of lading
- 4. Scale tickets
- 5. Purchase orders
- 6. Sales orders
- 7. Sales invoices
- 8. Phytosanitary certificates
- 9. Export declaration forms
- 10. Transaction Certificates

# 21. Describe lot numbering system

- a. Can record system tract the lot numbering system yes /No
- b. Can your record keeping system balance organic ingredients in and organic products Out (yes/No)
- c. How long the operator keep records

### **Affirmation**

The above given Information on this form is true to the best of my knowledge. I agree to provide further information as required by the APSOPCA.

Place:

Date:

Signature of the Operator

//Approved//

Evaluator,

APSOPCA, Guntur