

DOC NO.: FDA/FER/FOR - 01

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Effective Date: 13/09/2024

1. DETAILS OF APPLIC	CANI			
Name of Applicant: (Company)				
Company TIN number:				
Postal Address:				
Telephone:	Email:			
Contact Person Name:				
2. DETAILS OF MANUFACTURER				
Name of Manufacturer:				
Manufacturing Facility Location Address (Please include; Plot No./				
HNo/Ghana Post Digital Address):				
Telephone:	Email:			
3. DETAILS OF THIRD-PARTY REPRESENTATION (WHERE APPLICABLE)				
Name:				
Business Address:				
Telephone:	Email:			
Contact Person Name:				



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4. INFORMATION ON PRODUCT			
Brand Name:			
Product Description:			
Variants (If any)	Yes:	No:	
Type of Packaging:			
Primary Packaging:			
Secondary Packaging:			
Recommended Storage & Handling Condition:			
Shelf-Life:		Best Before/ Use By/Expiry Date:	
Country of Origin:		Date.	
Labelling Language:			
Number of Samples Submitted:			
Type of Samples Submitted (Please tick the box that Applies)	Final Samples	Mock Samples	
Unit Size (s) Volume or Weight)			
Raw materials/ Ingredients:			
For Packaged water Source of Water: Bore-hole/ Pipe borne Other-specify			



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5. PRODUCTION PROCESS			
Briefly describe your Production Process			
6. TYPE OF APPLICATION (P	ease tick the box that applies) Renewal		
the last registration? (Please tick the box that applies, and briefly describe the modifications			
Yes, labels			
accordingly) No Changes Yes, labels Yes, location of manufacturing facility Yes, Product Formulation			
Yes, Product Formulation			
7. DECLARATION I Hereby, declare that the above information is correct to the best of my knowledge.			
Name:	Signature:		
Position:	Date:		
NB: Please attach the following documents:			
1. Copy of Business Registration Certificate			
2. Certificate of Ana	ysis (per product and variant), Sanitary or Phytosanitary		
Certificate (where a	pplicable), and Radiation Certificate (where applicable)		



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- 3. Details of any investigations carried out to determine whether or not ingredient(s) used in Manufacturing/processing the product is injurious to health
- 4. Model Labels (per product and variant)
- 5. Copy of previous Registration Letter or Certificate