



KWAZULU-NATAL PROVINCE
HEALTH
REPUBLIC OF SOUTH AFRICA

Quotation Advert

Opening Date: 12/09/2024
Closing Date: 16/09/2024
Closing Time: 11:00

INSTITUTION DETAILS

Institution Name: Harry Gwala Hospital
Province: KwaZulu-Natal
Department of entity: Department of Health
Division or section: Supply Chain Management
Place where goods/
service is required: Harry Gwala Regional Hospital
Date Submitted: 12/09/2024

ITEM CATEGORY AND DETAILS

Quotation number: Edn: 1593/24-25
Item Category: Goods
Item Description: Baby feeds

COMPULSORY BRIEFING SESSION / SITE VISIT

Select Type: Not applicable
Date: Not applicable
Venue: Not applicable

QUOTES CAN BE COLLECTED FROM: KZN health website

QUOTATION MUST BE DEPOSITED ON THE BLUE TENDER BOX AT HARRY GWALA REGIONAL HOSPITAL SITUATED BEHIND SECURITY GATE, BEFORE THE CLOSING DATE AND TIME OF TENDER

ENQUIRIES REGARDING ADVERT MAY BE DIRECTED TO:

Name: Miss S. Zondi
Email: Sibuyiselwe.Zondi@kznhealth.gov.za

Contact number: 033-3954247

Finance Manager Name: Ms.P.S.Hlela

Finance Manage signature:

OFFICIAL PRICE PAGE FOR QUOTATIONS COVERING R2 000.01

QUOTE NUMBER: ZNQ / EDN / 24 / 25

DESCRIPTION: Baby feeds

PREFERENCE POINTS WILL BE ALLOCATED ACCORDING TO THE IMPLEMENTATION OF SPECIFIC GOALS IN TERMS OF PPR 2022:	POINTS ALLOCATED
Race – Full/partial/combination of points allocated to companies at least 51% owned by Black People	20

ICN NUMBER	QUANTITY	UNIT OF MEASURE	DESCRIPTION	BRAND & MODEL	COUNTRY OF MANUFACTURE	PRICE	
						R	C
	3600	Unit	F100 ready to use infant formula 200ml				
	480	Unit	F75 ready to use infant formula 200ml				
			NB: Feeds need to be delivered 2 weeks after order has been placed				
			Commodity with Baby and special foods suppliers (prior to advert date)				
			Bidder to fully read and complete standard quotation and specification				
VALUE ADDED TAX @ 15% (Only if VAT Vendor)							
TOTAL QUOTATION PRICE (VALIDITY PERIOD 90 Days)							

DOES THIS OFFER COMPLY WITH THE SPECIFICATION? YES / NO
 IS THE PRICE FIRM? YES / NO
 DOES THE ARTICLE CONFORM TO THE S.A.N.S. / S.A.B.S. SPECIFICATION? YES / NO

STATE DELIVERY PERIOD (E.G. 3 DAYS, 1 WEEK) _____

NAME OF BIDDER: _____

SIGNATURE OF BIDDER: _____

[By signing this document, I hereby agree to all terms and conditions]

CAPACITY UNDER WHICH THIS QUOTE IS SIGNED: _____

DATE: _____

BIDDER'S DISCLOSURE

1 PURPOSE OF THE FORM

Any person (natural or juristic) may make an offer or offers in terms of this invitation to bid. In line with the principles of transparency, accountability, impartiality, and ethics as enshrined in the Constitution of the Republic of South Africa and further expressed in various pieces of legislation, it is required for the bidder to make this declaration in respect of the details required hereunder.

Where a person/s are listed in the Register for Tender Defaulters and / or the List of Restricted Suppliers, that person will automatically be disqualified from the bid process.

2 BIDDER'S DECLARATION

2.1. Is the bidder, or any of its directors / trustees / shareholders / members / partners or any person having a controlling interest¹ in the enterprise, employed by the state? YES / NO

2.1.1. If so, furnish particulars of the names, individual identity numbers, and, if applicable, state employee numbers of sole proprietor/ directors / trustees / shareholders / members/ partners or any person having a controlling interest in the enterprise, in table below.

FULL NAME	IDENTITY NUMBER	NAME OF STATE INSTITUTION

2.2. Do you, or any person connected with the bidder, have a relationship with any person who is employed by the procuring institution²? YES / NO

2.2.1. If so, furnish particulars: _____

2.3. Does the bidder or any of its directors / trustees / shareholders / members / partners or any person having a controlling interest in the enterprise have any interest in any other related enterprise whether or not they are bidding for this contract? YES / NO

2.3.1. If so, furnish particulars: _____

3 DECLARATION

I, the undersigned, (name) _____ in submitting the accompanying bid, do hereby make the following statements that I certify to be true and complete in every respect:

- 3.1. I have read and I understand the contents of this disclosure;
- 3.2. I understand that the accompanying bid will be disqualified if this disclosure is found not to be true and complete in every respect;
- 3.3. The bidder has arrived at the accompanying bid independently from, and without consultation, communication, agreement or arrangement with any competitor. However, communication between partners in a joint venture or consortium³ will not be construed as collusive bidding.
- 3.4. In addition, there have been no consultations, communications, agreements or arrangements with any competitor regarding the quality, quantity, specifications, prices, including methods, factors or formulas used to calculate prices, market allocation, the intention or decision to submit or not to submit the bid, bidding with the intention not to win the bid and conditions or delivery particulars of the products or services to which this bid invitation relates.
- 3.5. The terms of the accompanying bid have not been, and will not be, disclosed by the bidder, directly or indirectly, to any competitor, prior to the date and time of the official bid opening or of the awarding of the contract.
- 3.6. There have been no consultations, communications, agreements or arrangements made by the bidder with any official of the procuring institution in relation to this procurement process prior to and during the bidding process except to provide clarification on the bid submitted where so required by the institution; and the bidder was not involved in the drafting of the specifications or terms of reference for this bid.
- 3.7. I am aware that, in addition and without prejudice to any other remedy provided to combat any restrictive practices related to bids and contracts, bids that are suspicious will be reported to the Competition Commission for investigation and possible imposition of administrative penalties in terms of section 59 of the Competition Act No 89 of 1998 and or may be reported to the National Prosecuting Authority (NPA) for criminal investigation and or may be restricted from conducting business with the public sector for a period not exceeding ten (10) years in terms of the Prevention and Combating of Corrupt Activities Act No 12 of 2004 or any other applicable legislation.

I CERTIFY THAT THE INFORMATION FURNISHED IN PARAGRAPHS 1, 2 and 3 ABOVE IS CORRECT.

I ACCEPT THAT THE STATE MAY REJECT THE BID OR ACT AGAINST ME IN TERMS OF PARAGRAPH 6 OF PFMA SCM INSTRUCTION 03 OF 2021/22 ON PREVENTING AND COMBATING ABUSE IN THE SUPPLY CHAIN MANAGEMENT SYSTEM SHOULD THIS DECLARATION PROVE TO BE FALSE.

NAME OF BIDDER	SIGNATURE	POSITION	DATE
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1 The power, by one person or a group of persons holding the majority of the equity of an enterprise, alternatively, the person/s having the deciding vote or power to influence or to direct the course and decisions of the enterprise.

2 "Procuring institution" refers to all institutions under the Accounting Officer of the Department of Health.

3 Joint venture or Consortium means an association of persons for the purpose of combining their expertise, property, capital, efforts, skill and knowledge in an activity for the execution of a contract.

GENERAL CONDITIONS OF CONTRACT

NOTES

The purpose of this document is to:

- (i) Draw special attention to certain general conditions applicable to government bids, contracts and orders; and
- (ii) To ensure that clients be familiar with regard to the rights and obligations of all parties involved in doing business with government.

In this document words in the singular also mean in the plural and vice versa and words in the masculine also mean in the feminine and neuter.

- The General Conditions of Contract will form part of all bid/quotation documents and may not be amended.
- Special Conditions of Contract (SCC) relevant to a specific bid, should be compiled separately for every bid (if applicable) and will supplement the General Conditions of Contract. Whenever there is a conflict, the provisions in the SCC shall prevail.

1 Definitions

The following terms shall be interpreted as indicated:

- 1.1. "Closing time" means the date and hour specified in the bidding documents for the receipt of bids.
- 1.2. "Contract" means the written agreement entered into between the purchaser and the supplier, as recorded in the contract form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.
- 1.3. "Contract price" means the price payable to the supplier under the contract for the full and proper performance of his contractual obligations.
- 1.4. "Corrupt practice" means the offering, giving, receiving, or soliciting of any thing of value to influence the action of a public official in the procurement process or in contract execution.
- 1.5. "Countervailing duties" are imposed in cases where an enterprise abroad is subsidized by its government and encouraged to market its products internationally.
- 1.6. "Country of origin" means the place where the goods were mined, grown or produced or from which the services are supplied. Goods are produced when, through manufacturing, processing or substantial and major assembly of components, a commercially recognized new product results that is substantially different in basic characteristics or in purpose or utility from its components.
- 1.7. "Day" means calendar day.
- 1.8. "Delivery" means delivery in compliance of the conditions of the contract or order.
- 1.9. "Delivery ex stock" means immediate delivery directly from stock actually on hand.
- 1.10. "Delivery into consignees store or to his site" means delivered and unloaded in the specified store or depot or on the specified site in compliance with the conditions of the contract or order, the supplier bearing all risks and charges involved until the supplies are so delivered and a valid receipt is obtained.
- 1.11. "Dumping" occurs when a private enterprise abroad market its goods on own initiative in the RSA at lower prices than that of the country of origin and which have the potential to harm the local industries in the RSA
- 1.12. "Force majeure" means an event beyond the control of the supplier and not involving the supplier's fault or negligence and not foreseeable. Such events may include, but is not restricted to, acts of the purchaser in its sovereign capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions and freight embargoes.
- 1.13. "Fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of any bidder, and includes collusive practice among bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to deprive the bidder of the benefits of free and open competition.
- 1.14. "GCC" means the General Conditions of Contract.
- 1.15. "Goods" means all of the equipment, machinery, and/or other materials that the supplier is required to supply to the purchaser under the contract.
- 1.16. "Imported content" means that portion of the bidding price represented by the cost of components, parts or materials which have been or are still to be imported (whether by the supplier or his subcontractors) and which costs are inclusive of the costs abroad, plus freight and other direct importation costs such as landing costs, dock dues, import duty, sales duty or other similar tax or duty at the South African place of entry as well as transportation and handling charges to the factory in the Republic where the supplies covered by the bid will be manufactured.
- 1.17. "Local content" means that portion of the bidding price which is not included in the imported content provided that local manufacture does take place.
- 1.18. "Manufacture" means the production of products in a factory using labour, materials, components and machinery and includes other related value-adding activities.
- 1.19. "Order" means an official written order issued for the supply of goods or works or the rendering of a service.
- 1.20. "Project site," where applicable, means the place indicated in bidding documents.
- 1.21. "Purchaser" means the organization purchasing the goods.
- 1.22. "Republic" means the Republic of South Africa.
- 1.23. "SCC" means the Special Conditions of Contract.
- 1.24. "Services" means those functional services ancillary to the supply of the goods, such as transportation and any other incidental services, such as installation, commissioning, provision of technical assistance, training, catering, gardening, security, maintenance and other such obligations of the supplier covered under the contract.
- 1.25. "Written" or "in writing" means handwritten in ink or any form of electronic or mechanical writing.

2 Application

- 2.1. These general conditions are applicable to all bids, contracts and orders including bids for functional and professional services, sales, hiring, letting and the granting or acquiring of rights, but excluding immovable property, unless otherwise indicated in the bidding documents.
- 2.2. Where applicable, special conditions of contract are also laid down to cover specific supplies, services or works.
- 2.3. Where such special conditions of contract are in conflict with these general conditions, the special conditions shall apply.

3 General

- 3.1. Unless otherwise indicated in the bidding documents, the purchaser shall not be liable for any expense incurred in the preparation and submission of a bid. Where applicable a non-refundable fee for documents may be charged.
- 3.2. With certain exceptions, invitations to bid are only published in the Government Tender Bulletin. The Government Tender Bulletin may be obtained directly from the Government Printer, Private Bag X85, Pretoria 0001, or accessed electronically from www.treasury.gov.za



4 Standards

- 4.1. The goods supplied shall conform to the standards mentioned in the bidding documents and specifications.

5 Use of contract documents and information; inspection.

- 5.1. The supplier shall not, without the purchaser's prior written consent, disclose the contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the purchaser in connection therewith, to any person other than a person employed by the supplier in the performance of the contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.
- 5.2. The supplier shall not, without the purchaser's prior written consent, make use of any document or information mentioned in GCC clause 5.1 except for purposes of performing the contract.
- 5.3. Any document, other than the contract itself mentioned in GCC clause 5.1 shall remain the property of the purchaser and shall be returned (all copies) to the purchaser on completion of the supplier's performance under the contract if so required by the purchaser.
- 5.4. The supplier shall permit the purchaser to inspect the supplier's records relating to the performance of the supplier and to have them audited by auditors appointed by the purchaser, if so required by the purchaser.

6 Patent rights

- 6.1. The supplier shall indemnify the purchaser against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the goods or any part thereof by the purchaser.

7 Performance security

- 7.1. Within thirty (30) days of receipt of the notification of contract award, the successful bidder shall furnish to the purchaser the performance security of the amount specified in SCC.
- 7.2. The proceeds of the performance security shall be payable to the purchaser as compensation for any loss resulting from the supplier's failure to complete his obligations under the contract.
- 7.3. The performance security shall be denominated in the currency of the contract, or in a freely convertible currency acceptable to the purchaser and shall be in one of the following forms:
(a) a bank guarantee or an irrevocable letter of credit issued by a reputable bank located in the purchaser's country or abroad, acceptable to the purchaser, in the form provided in the bidding documents or another form acceptable to the purchaser; or
(b) a cashier's or certified cheque
- 7.4. The performance security will be discharged by the purchaser and returned to the supplier not later than thirty (30) days following the date of completion of the supplier's performance obligations under the contract, including any warranty obligations, unless otherwise specified in SCC.

8 Inspections, tests and analyses

- 8.1. All pre-bidding testing will be for the account of the bidder.
- 8.2. If it is a bid condition that supplies to be produced or services to be rendered should at any stage during production or execution or on completion be subject to inspection, the premises of the bidder or contractor shall be open, at all reasonable hours, for inspection by a representative of the Department or an organization acting on behalf of the Department.
- 8.3. If there are no inspection requirements indicated in the bidding documents and no mention is made in the contract, but during the contract period it is decided that inspections shall be carried out, the purchaser shall itself make the necessary arrangements, including payment arrangements with the testing authority concerned.
- 8.4. If the inspections, tests and analyses referred to in clauses 8.2 and 8.3 show the supplies to be in accordance with the contract requirements, the cost of the inspections, tests and analyses shall be defrayed by the purchaser.
- 8.5. Where the supplies or services referred to in clauses 8.2 and 8.3 do not comply with the contract requirements, irrespective of whether such supplies or services are accepted or not, the cost in connection with these inspections, tests or analyses shall be defrayed by the supplier.
- 8.6. Supplies and services which are referred to in clauses 8.2 and 8.3 and which do not comply with the contract requirements may be rejected.
- 8.7. Any contract supplies may on or after delivery be inspected, tested or analyzed and may be rejected if found not to comply with the requirements of the contract. Such rejected supplies shall be held at the cost and risk of the supplier who shall, when called upon, remove them immediately at his own cost and forthwith substitute them with supplies which do comply with the requirements of the contract. Failing such removal the rejected supplies shall be returned at the suppliers cost and risk. Should the supplier fail to provide the substitute supplies forthwith, the purchaser may, without giving the supplier further opportunity to substitute the rejected supplies, purchase such supplies as may be necessary at the expense of the supplier.
- 8.8. The provisions of clauses 8.4 to 8.7 shall not prejudice the right of the purchaser to cancel the contract on account of a breach of the conditions thereof, or to act in terms of Clause 23 of GCC.

9 Packing

- 9.1. The supplier shall provide such packing of the goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packing, case size and weights shall take into consideration, where appropriate, the remoteness of the goods' final destination and the absence of heavy handling facilities at all points in transit.
- 9.2. The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the contract, including additional requirements, if any, specified in SCC, and in any subsequent instructions ordered by the purchaser.

10 Delivery and documents

- 10.1. Delivery of the goods shall be made by the supplier in accordance with the terms specified in the contract. The details of shipping and/or other documents to be furnished by the supplier are specified in SCC.
- 10.2. Documents to be submitted by the supplier are specified in SCC.

11 Insurance

- 11.1. The goods supplied under the contract shall be fully insured in a freely convertible currency against loss or damage incidental to manufacture or acquisition, transportation, storage and delivery in the manner specified in the SCC.

12 Transportation

- 12.1. Should a price other than an all-inclusive delivered price be required, this shall be specified in the SCC.

13 Incidental services

- 13.1. The supplier may be required to provide any or all of the following services, including additional services, if any, specified in SCC:
- (a) performance or supervision of on-site assembly and/or commissioning of the supplied goods;
 - (b) furnishing of tools required for assembly and/or maintenance of the supplied goods;
 - (c) furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied goods;
 - (d) performance or supervision or maintenance and/or repair of the supplied goods, for a period of time agreed by the parties, provided that this service shall not relieve the supplier of any warranty obligations under this contract; and
 - (e) training of the purchaser's personnel, at the supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the
- 13.2. Prices charged by the supplier for incidental services, if not included in the contract price for the goods, shall be agreed upon in advance by the parties and shall not exceed the prevailing rates charged to other parties by the supplier for similar services.

14 Spare parts

- As specified in SCC, the supplier may be required to provide any or all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the supplier:
- 14.1.
- (a) such spare parts as the purchaser may elect to purchase from the supplier, provided that this election shall not relieve the supplier of any warranty obligations under the contract; and
 - (b) in the event of termination of production of the spare parts:
 - (i) Advance notification to the purchaser of the pending termination, in sufficient time to permit the purchaser to procure needed requirements; and
 - (ii) following such termination, furnishing at no cost to the purchaser, the blueprints, drawings, and specifications of the spare parts, if requested.

15 Warranty

- 15.1. The supplier warrants that the goods supplied under the contract are new, unused, of the most recent or current models, and that they incorporate all recent improvements in design and materials unless provided otherwise in the contract. The supplier further warrants that all goods supplied under this contract shall have no defect, arising from design, materials, or workmanship (except when the design and/or material is required by the purchaser's specifications) or from any act or omission of the supplier, that may develop under normal use of the supplied goods in the conditions prevailing in the country of final destination.
- 15.2. This warranty shall remain valid for twelve (12) months after the goods, or any portion thereof as the case may be, have been delivered to and accepted at the final destination indicated in the contract, or for eighteen (18) months after the date of shipment from the port or place of loading in the source country, whichever period concludes earlier, unless specified otherwise in SCC.
- 15.3. The purchaser shall promptly notify the supplier in writing of any claims arising under this warranty.
- 15.4. Upon receipt of such notice, the supplier shall, within the period specified in SCC and with all reasonable speed, repair or replace the defective goods or parts thereof, without costs to the purchaser.
- 15.5. If the supplier, having been notified, fails to remedy the defect(s) within the period specified in SCC, the purchaser may proceed to take such remedial action as may be necessary, at the supplier's risk and expense and without prejudice to any other rights which the purchaser may have against the supplier under the contract.

16 Payment

- 16.1. The method and conditions of payment to be made to the supplier under this contract shall be specified in SCC.
- 16.2. The supplier shall furnish the purchaser with an invoice accompanied by a copy of the delivery note and upon fulfillment of other obligations stipulated in the contract.
- 16.3. Payments shall be made promptly by the purchaser, but in no case later than thirty (30) days after submission of an invoice or claim by the supplier.
- 16.4. Payment will be made in Rand unless otherwise stipulated in SCC.

17 Prices

- 17.1. Prices charged by the supplier for goods delivered and services performed under the contract shall not vary from the prices quoted by the supplier in his bid, with the exception of any price adjustments authorized in SCC or in the purchaser's request for bid validity extension, as the case may be.

18 Contract amendments

- 18.1. No variation in or modification of the terms of the contract shall be made except by written amendment signed by the parties concerned.

19 Assignment

- 19.1. The supplier shall not assign, in whole or in part, its obligations to perform under the contract, except with the purchaser's prior written consent.

20 Subcontracts

- 20.1. The supplier shall notify the purchaser in writing of all subcontracts awarded under this contracts if not already specified in the bid. Such notification, in the original bid or later, shall not relieve the supplier from any liability or obligation under the contract.

21 Delays in the supplier's performance

- 21.1. Delivery of the goods and performance of services shall be made by the supplier in accordance with the time schedule prescribed by the purchaser in the contract.
- 21.2. If at any time during performance of the contract, the supplier or its subcontractor(s) should encounter conditions impeding timely delivery of the goods and performance of services, the supplier shall promptly notify the purchaser in writing of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the supplier's notice, the purchaser shall evaluate the situation and may at his discretion extend the supplier's time for performance, with or without the imposition of penalties, in which case the extension shall be ratified by the parties by amendment of contract.
- 21.3. No provision in a contract shall be deemed to prohibit the obtaining of supplies or services from a national department, provincial department, or a local authority.
- 21.4. The right is reserved to procure outside of the contract small quantities or to have minor essential services executed if an emergency arises, the supplier's point of supply is not situated at or near the place where the supplies are required, or the supplier's services are not readily available.

- 21.5. Except as provided under GCC Clause 25, a delay by the supplier in the performance of its delivery obligations shall render the supplier liable to the imposition of penalties, pursuant to GCC Clause 22, unless an extension of time is agreed upon pursuant to GCC Clause 21.2 without the application of penalties.
- 21.6. Upon any delay beyond the delivery period in the case of a supplies contract, the purchaser shall, without canceling the contract, be entitled to purchase supplies of a similar quality and up to the same quantity in substitution of the goods not supplied in conformity with the contract and to return any goods delivered later at the supplier's expense and risk, or to cancel the contract and buy such goods as may be required to complete the contract and without prejudice to his other rights, be entitled to claim damages from the supplier.
- 22 Penalties**
- 22.1. Subject to GCC Clause 25, if the supplier fails to deliver any or all of the goods or to perform the services within the period(s) specified in the contract, the purchaser shall, without prejudice to its other remedies under the contract, deduct from the contract price, as a penalty, a sum calculated on the delivered price of the delayed goods or unperformed services using the current prime interest rate calculated for each day of the delay until actual delivery or performance. The purchaser may also consider termination of the contract pursuant to GCC Clause 23.
- 23 Termination for default**
- 23.1. The purchaser, without prejudice to any other remedy for breach of contract, by written notice of default sent to the supplier, may terminate this contract in whole or in part:
- (a) if the supplier fails to deliver any or all of the goods within the period(s) specified in the contract, or within any extension thereof granted by the purchaser pursuant to GCC Clause 21.2;
- (b) if the Supplier fails to perform any other obligation(s) under the contract; or
- (c) if the supplier, in the judgment of the purchaser, has engaged in corrupt or fraudulent practices in competing for or in executing the contract.
- 23.2. In the event the purchaser terminates the contract in whole or in part, the purchaser may procure, upon such terms and in such manner as it deems appropriate, goods, works or services similar to those undelivered, and the supplier shall be liable to the purchaser for any excess costs for such similar goods, works or services. However, the supplier shall continue performance of the contract to the extent not terminated.
- 23.3. Where the purchaser terminates the contract in whole or in part, the purchaser may decide to impose a restriction penalty on the supplier by prohibiting such supplier from doing business with the public sector for a period not exceeding 10 years.
- 23.4. If a purchaser intends imposing a restriction on a supplier or any person associated with the supplier, the supplier will be allowed a time period of not more than fourteen (14) days to provide reasons why the envisaged restriction should not be imposed. Should the supplier fail to respond within the stipulated fourteen (14) days the purchaser may regard the intended penalty as not objected against and may impose it on the supplier.
- 23.5. Any restriction imposed on any person by the Accounting Officer / Authority will, at the discretion of the Accounting Officer / Authority, also be applicable to any other enterprise or any partner, manager, director or other person who wholly or partly exercises or exercised or may exercise control over the enterprise of the first-mentioned person, and with which enterprise or person the first-mentioned person, is or was in the opinion of the Accounting Officer / Authority actively associated.
- 23.6. If a restriction is imposed, the purchaser must, within five (5) working days of such imposition, furnish the National Treasury, with the following information:
- (i) the name and address of the supplier and / or person restricted by the purchaser;
- (ii) the date of commencement of the restriction
- (iii) the period of restriction; and
- (iv) the reasons for the restriction.
- These details will be loaded in the National Treasury's central database of suppliers or persons prohibited from doing business with the public sector.
- 23.7. If a court of law convicts a person of an offence as contemplated in sections 12 or 13 of the Prevention and Combating of Corrupt Activities Act, No. 12 of 2004, the court may also rule that such person's name be endorsed on the Register for Tender Defaulters. When a person's name has been endorsed on the Register, the person will be prohibited from doing business with the public sector for a period not less than five years and not more than 10 years. The National Treasury is empowered to determine the period of restriction and each case will be dealt with on its own merits. According to section 32 of the Act the Register must be open to the public. The Register can be perused on the National Treasury website.
- 24 Anti-dumping and countervailing duties and rights**
- 24.1. When, after the date of bid, provisional payments are required, or antidumping or countervailing duties are imposed, or the amount of a provisional payment or anti-dumping or countervailing right is increased in respect of any dumped or subsidized import, the State is not liable for any amount so required or imposed, or for the amount of any such increase. When, after the said date, such a provisional payment is no longer required or any such anti-dumping or countervailing right is abolished, or where the amount of such provisional payment or any such right is reduced, any such favourable difference shall on demand be paid forthwith by the contractor to the State or the State may deduct such amounts from moneys (if any) which may otherwise be due to the contractor in regard to supplies or services which he delivered or rendered, or is to deliver or render in terms of the contract or any other contract or any other amount which may be due to him.
- 25 Force Majeure**
- 25.1. Notwithstanding the provisions of GCC Clauses 22 and 23, the supplier shall not be liable for forfeiture of its performance security, damages, or termination for default if and to the extent that his delay in performance or other failure to perform his obligations under the contract is the result of an event of force majeure.
- 25.2. If a force majeure situation arises, the supplier shall promptly notify the purchaser in writing of such condition and the cause thereof. Unless otherwise directed by the purchaser in writing, the supplier shall continue to perform its obligations under the contract as far as is reasonably practical, and shall seek all reasonable alternative means for performance not prevented by the force majeure event.
- 26 Termination for insolvency**
- 26.1. The purchaser may at any time terminate the contract by giving written notice to the supplier if the supplier becomes bankrupt or otherwise insolvent. In this event, termination will be without compensation to the supplier, provided that such termination will not prejudice or affect any right of action or remedy which has accrued or will accrue thereafter to the purchaser.
- 27 Settlement of Disputes**
- 27.1. If any dispute or difference of any kind whatsoever arises between the purchaser and the supplier in connection with or arising out of the contract, the parties shall make every effort to resolve amicably such dispute or difference by mutual consultation.



- 27.2. If, after thirty (30) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the purchaser or the supplier may give notice to the other party of his intention to commence with mediation. No mediation in respect of this matter may be commenced unless such notice is given to the other party.
- 27.3. Should it not be possible to settle a dispute by means of mediation, it may be settled in a South African court of law.
- 27.4. Mediation proceedings shall be conducted in accordance with the rules of procedure specified in the SCC.
- 27.5. Notwithstanding any reference to mediation and/or court proceedings herein,
(a) the parties shall continue to perform their respective obligations under the contract unless they otherwise agree; and
(b) the purchaser shall pay the supplier any monies due the supplier.
- 28 Limitation of liability**
- 28.1. Except in cases of criminal negligence or willful misconduct, and in the case of infringement pursuant to Clause 6;
(a) the supplier shall not be liable to the purchaser, whether in contract, tort, or otherwise, for any indirect or consequential loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the supplier to pay penalties and/or damages to the purchaser; and
(b) the aggregate liability of the supplier to the purchaser, whether under the contract, in tort or otherwise, shall not exceed the total contract price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment.
- 29 Governing language**
- 29.1. The contract shall be written in English. All correspondence and other documents pertaining to the contract that is exchanged by the parties shall also be written in English.
- 30 Applicable law**
- 30.1. The contract shall be interpreted in accordance with South African laws, unless otherwise specified in SCC.
- 31 Notices**
- 31.1. Every written acceptance of a bid shall be posted to the supplier concerned by registered or certified mail and any other notice to him shall be posted by ordinary mail to the address furnished in his bid or to the address notified later by him in writing and such posting shall be deemed to be proper service of such notice
- 31.2. The time mentioned in the contract documents for performing any act after such aforesaid notice has been given, shall be reckoned from the date of posting of such notice.
- 32 Taxes and duties**
- 32.1. A foreign supplier shall be entirely responsible for all taxes, stamp duties, license fees, and other such levies imposed outside the purchaser's country.
- 32.2. A local supplier shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted goods to the purchaser.
- 32.3. No contract shall be concluded with any bidder whose tax matters are not in order. Prior to the award of a bid the Department must be in possession of a tax clearance certificate, submitted by the bidder. This certificate must be an original issued by the South African Revenue Services.
- 33 National Industrial Participation (NIP) Programme**
- 33.1. The NIP Programme administered by the Department of Trade and Industry shall be applicable to all contracts that are subject to the NIP obligation.
- 34 Prohibition of Restrictive practices**
- 34.1. In terms of section 4 (1) (b) (iii) of the Competition Act No. 89 of 1998, as amended, an agreement between, or concerted practice by, firms, or a decision by an association of firms, is prohibited if it is between parties in a horizontal relationship and if a bidder (s) is / are or a contractor(s) was / were involved in collusive bidding (or bid rigging).
- 34.2. If a bidder(s) or contractor(s), based on reasonable grounds or evidence obtained by the purchaser, has / have engaged in the restrictive practice referred to above, the purchaser may refer the matter to the Competition Commission for investigation and possible imposition of administrative penalties as contemplated in the Competition Act No. 89 of 1998.
- 34.3. If a bidder(s) or contractor(s), has / have been found guilty by the Competition Commission of the restrictive practice referred to above, the purchaser may, in addition and without prejudice to any other remedy provided for, invalidate the bid(s) for such item(s) offered, and / or terminate the contract in whole or part, and / or restrict the bidder(s) or contractor(s) from conducting business with the public sector for a period not exceeding ten (10) years and / or claim damages from the bidder(s) or contractor(s) concerned.

SPECIAL CONDITIONS OF CONTRACT

1. AMENDMENT OF CONTRACT

- 1.1. Any amendment to or renunciation of the provisions of the contract shall at all times be done in writing and shall be signed by both parties.

2. CHANGE OF ADDRESS

- 2.1. Bidders must advise the Department of Health (institution where the offer was submitted) should their address (domicilium citandi et executandi) details change from the time of bidding to the expiry of the contract.

3. GENERAL CONDITIONS ATTACHED TO THIS QUOTATION

- 3.1. The Department is under no obligation to accept the lowest or any quote.
- 3.2. The Department reserves the right to communicate in writing with vendors in cases where information is incomplete or where there are obscurities regarding technical aspects of the offer, to obtain confirmation of prices or preference claims in cases where it is evident that a typing, written, transfer or unit error has been made, to investigate the vendor's standing and ability to complete the supply/service satisfactorily.
- 3.3. **ALL DECISIONS TAKEN BY THE DEPARTMENT ARE FINAL, INCLUDING THE AWARD OR CANCELLATION OF THIS QUOTATION.**
- 3.4. The price quoted must include VAT (if VAT vendor).
- 3.5. Should a bidder become a VAT vendor after award or during the implementation of a contract, they may not request the VAT percentage from the Department as the service provider made an offer during the period they were not registered as a VAT vendor. The Department is only liable for any VAT from registered VAT vendors as originally stated on the quotation document.
- 3.6. The bidder must ensure the correctness & validity of the quotation:
(i) that the price(s), rate(s) & preference quoted cover all for the work/item (s) & accept that any mistakes regarding the price (s) & calculations will be at the bidder's risk;
(ii) it is the responsibility of the bidder to confirm receipt of their quotation and to keep proof thereof.
- 3.7. The bidder must accept full responsibility for the proper execution & fulfilment of all obligations conditions devolving on under this agreement, as the Principal (s) liable for the due fulfilment of this contract.
- 3.8. This quotation will be evaluated based on the 80/20 points system, specification, correctness of information and/or functionality criteria. All required documentation must be completed in full and submitted.
- 3.9. Offers must comply strictly with the specification.
- 3.10. Only offers that meet or are greater than the specification will be considered.
- 3.11. Late offers will not be considered.
- 3.12. Expired product/s will not be accepted. All products supplied must be valid for a minimum period of six months.
- 3.13. Used/ second-hand products will not be accepted.
- 3.14. A bidder not registered on the Central Suppliers Database or whose verification has failed will not be considered.
- 3.15. All delivery costs must be included in the quoted price for delivery at the prescribed destination.
- 3.16. Only firm prices will be accepted. Such prices must remain firm for the contract period. Non-firm prices (including rates of exchange variations) will not be considered.
- 3.17. In cases where different delivery points influence the pricing, a separate pricing schedule must be submitted for each delivery point.
- 3.18. In the event of a bidder having multiple quotes, only the cheapest according to specification will be considered.
- 3.19. Verification will be conducted to identify if bidders have multiple companies and are cover-quoting for this bid.
- 3.20. In such instances, the Department reserves the right to immediately disqualify such bidders as cover-quoting is an offence that represents both corruption and acquisition fraud.

4. SPECIAL INSTRUCTIONS AND NOTICES TO BIDDERS REGARDING THE COMPLETION OF THIS QUOTATION.

- 4.1. Unless inconsistent with or expressly indicated otherwise by the context, the singular shall include the plural and vice versa and with words importing the masculine gender shall include the feminine and the neuter.
- 4.2. Under no circumstances whatsoever may the quotation/bid forms be retyped or redrafted. Photocopies of the original bid documentation may be used, but an original signature must appear on such photocopies.
- 4.3. The bidder is advised to check the number of pages and to satisfy himself that none are missing or duplicated.
- 4.4. Quotations submitted must be complete in all respects. However, where it is identified that information in a bidder's response, which does not affect the preference points or price, is incomplete in any respect, the said supplier meets all specification requirements and scores the highest points in terms of preference points and price, the Department reserves the right to request the bidder to complete/ submit such information.
- 4.5. Any alteration made by the bidder must be initialled; failure to do so may render the response invalid.
- 4.6. Use of correcting fluid is prohibited and may render the response invalid.
- 4.7. Quotations will be opened in public as soon as practicable after the closing time of quotation.
- 4.8. Where practical, prices are made public at the time of opening quotations.
- 4.9. If it is desired to make more than one offer against any individual item, such offers should be given on a photocopy of the page in question. Clear indication thereof must be stated on the schedules attached.
- 4.10. The Department is under no obligation to pay suppliers in part for work done if the supplier can no longer for fulfil their obligation.

5. SPECIAL INSTRUCTIONS REGARDING HAND DELIVERED QUOTATIONS

- 5.1. Quotation shall be lodged at the address indicated not later than the closing time specified for their receipt, and in accordance with the directives in the quotation documents.
- 5.2. Each quotation shall be addressed in accordance with the directives in the quotation documents and shall be lodged in a separate sealed envelope, with the name and address of the bidder, the quotation number and closing date indicated on the envelope. The envelope shall not contain documents relating to any quotation other than that shown on the envelope. If this provision is not complied with, such quotations/bids may be rejected as being invalid.
- 5.3. All quotations received in sealed envelopes with the relevant quotation numbers on the envelopes are kept unopened in safe custody until the closing time of the quotation/bids. Where, however, a quotation is received open, it shall be sealed. If it is received without a quotation/bid number on the envelope, it shall be opened, the quotation number ascertained, the envelope sealed and the quotation number written on the envelope.
- 5.4. A specific box is provided for the receipt of quotations, and no quotation found in any other box or elsewhere subsequent to the closing date and time of quotation will be considered.



14. TERMINATION FOR DEFAULT

- 14.1. The purchaser, without prejudice to any other remedy for breach of contract, by written notice of default sent to the supplier, may terminate this contract in whole or in part:
- (i) if the supplier fails to deliver any or all of the goods within the period(s) specified in the contract,
 - (ii) if the supplier fails to perform any other obligation(s) under the contract; or
 - (iii) if the supplier, in the judgment of the purchaser, has engaged in corrupt or fraudulent practices in competing for or in executing the contract.
- 14.2. In the event the purchaser terminates the contract in whole or in part, the purchaser may procure, upon such terms and in such manner as it deems appropriate, goods, works or services similar to those undelivered, and the supplier shall be liable to the purchaser for any excess costs for such similar goods, works or services.
- 14.3. Where the purchaser terminates the contract in whole or in part, the purchaser may decide to impose a restriction penalty on the supplier by prohibiting such supplier from doing business with the public sector for a period not exceeding 10 years.
15. **THE DEPARTMENT RESERVES THE RIGHT TO PASS OVER ANY QUOTATION WHICH FAILS TO COMPLY WITH THE ABOVE.**

PREFERENCE POINTS CLAIM FORM IN TERMS OF THE PREFERENTIAL PROCUREMENT REGULATIONS 2022

This preference form must form part of all tenders invited. It contains general information and serves as a claim form for preference points for specific goals.

NB: BEFORE COMPLETING THIS FORM, TENDERERS MUST STUDY THE GENERAL CONDITIONS, DEFINITIONS AND DIRECTIVES APPLICABLE IN RESPECT OF THE TENDER AND PREFERENTIAL PROCUREMENT REGULATIONS, 2022

1. GENERAL CONDITIONS

- 1.1. The following preference point systems are applicable to invitations to tender:
- the 80/20 system for requirements with a Rand value of up to R50 000 000 (all applicable taxes included); and
 - the 90/10 system for requirements with a Rand value above R50 000 000 (all applicable taxes included).

1.2. The applicable preference point system for this tender is the 80/20 preference point system.

- 1.3. Points for this tender (even in the case of a tender for income-generating contracts) shall be awarded for:
- (a) Price; and
 - (b) Specific Goals.

1.4. The maximum points for this tender are allocated as follows:

	POINTS
PRICE	80
SPECIFIC GOALS	20
Total points for Price and Specific Goals	100

1.5. Failure on the part of a tenderer to submit proof or documentation required in terms of this tender to claim points for specific goals with the tender, will be interpreted to mean that preference points for specific goals are not claimed.

1.6. The organ of state reserves the right to require of a tenderer, either before a tender is adjudicated or at any time subsequently, to substantiate any claim in regard to preferences, in any manner required by the organ of state.

2. DEFINITIONS

- (a) "tender" means a written offer in the form determined by an organ of state in response to an invitation to provide goods or services through price quotations, competitive tendering process or any other method envisaged in legislation;
- (b) "price" means an amount of money tendered for goods or services, and includes all applicable taxes less all unconditional discounts;
- (c) "rand value" means the total estimated value of a contract in Rand, calculated at the time of bid invitation, and includes all applicable taxes;
- (d) "tender for income-generating contracts" means a written offer in the form determined by an organ of state in response to an invitation for the origination of income-generating contracts through any method envisaged in legislation that will result in a legal agreement between the organ of state and a third party that produces revenue for the organ of state, and includes, but is not limited to, leasing and disposal of assets and concession contracts, excluding direct sales and disposal of assets through public auctions; and
- (e) "the Act" means the Preferential Procurement Policy Framework Act, 2000 (Act No. 5 of 2000).

3. FORMULAE FOR PROCUREMENT OF GOODS AND SERVICES

3.1. POINTS AWARDED FOR PRICE

3.1.1. THE 80/20 OR 90/10 PREFERENCE POINT SYSTEMS

A maximum of 80 or 90 points is allocated for price on the following basis:

$$\begin{array}{ccc}
 \frac{80}{20} & & \frac{90}{10} \\
 Ps = 80 \left(1 - \frac{Pt - Pmin}{Pmin} \right) & \text{OR} & Ps = 90 \left(1 - \frac{Pt - Pmin}{Pmin} \right)
 \end{array}$$

Where

- Ps = Points scored for price of tender under consideration
- Pt = Price of tender under consideration
- Pmin = Price of lowest acceptable tender

3.2. FORMULAE FOR DISPOSAL OR LEASING OF STATE ASSETS AND INCOME GENERATING PROCUREMENT

3.2.1. POINTS AWARDED FOR PRICE

A maximum of 80 or 90 points is allocated for price on the following basis:

$$\begin{array}{ccc}
 \frac{80}{20} & & \frac{90}{10} \\
 Ps = 80 \left(1 + \frac{Pt - Pmax}{Pmax} \right) & \text{OR} & Ps = 90 \left(1 + \frac{Pt - Pmax}{Pmax} \right)
 \end{array}$$

Where

- Ps = Points scored for price of tender under consideration
- Pt = Price of tender under consideration
- Pmax = Price of highest acceptable tender



4. POINTS AWARDED FOR SPECIFIC GOALS

- 4.1. In terms of Regulation 4(2); 5(2); 6(2) and 7(2) of the Preferential Procurement Regulations, preference points must be awarded for specific goals stated in the tender. For the purposes of this tender the tenderer will be allocated points based on the goals stated in table 1 below as may be supported by proof/ documentation stated in the conditions of this tender:
- 4.2. In cases where organs of state intend to use Regulation 3(2) of the Regulations, which states that, if it is unclear whether the 80/20 or 90/10 preference point system applies, an organ of state must, in the tender documents, stipulate in the case of—
- (a) an invitation for tender for income-generating contracts, that either the 80/20 or 90/10 preference point system will apply and that the highest acceptable tender will be used to determine the applicable preference point system; or
 - (b) any other invitation for tender, that either the 80/20 or 90/10 preference point system will apply and that the lowest acceptable tender will be used to determine the applicable preference point system,
- then the organ of state must indicate the points allocated for specific goals for both the 90/10 and 80/20 preference point system.

Table 1: Specific goals for the tender and points claimed are indicated per the table below.

Note to tenderers: The tenderer must indicate how they claim points for each preference point system.

The specific goal/s allocated points in terms of this tender	Number of points allocated (80/20 system)	Number of points claimed (80/20 system)
Race – Full/partial/combination of points allocated to companies at least 51% owned by Black People	20	

DECLARATION WITH REGARD TO COMPANY/FIRM

- 4.3. Name of company/firm: _____
- 4.4. Company registration number: _____
- 4.5. TYPE OF COMPANY/ FIRM [tick applicable box]
- Partnership/Joint Venture / Consortium
 - One-person business/sole propriety
 - Close corporation
 - Public Company
 - Personal Liability Company
 - (Pty) Limited
 - Non-Profit Company
 - State Owned Company

- 4.6. I, the undersigned, who is duly authorised to do so on behalf of the company/firm, certify that the points claimed, based on the specific goals as advised in the tender, qualifies the company/ firm for the preference(s) shown and I acknowledge that:
- i) The information furnished is true and correct;
 - ii) The preference points claimed are in accordance with the General Conditions as indicated in paragraph 1 of this form;
 - iii) In the event of a contract being awarded as a result of points claimed as shown in paragraphs 1.4 and 4.2, the contractor may be required to furnish documentary proof to the satisfaction of the organ of state that the claims are correct;
 - iv) If the specific goals have been claimed or obtained on a fraudulent basis or any of the conditions of contract have not been fulfilled, the organ of state may, in addition to any other remedy it may have –
 - (a) disqualify the person from the tendering process;
 - (b) recover costs, losses or damages it has incurred or suffered as a result of that person's conduct;
 - (c) cancel the contract and claim any damages which it has suffered as a result of having to make less favourable arrangements due to such cancellation;
 - (d) recommend that the tenderer or contractor, its shareholders and directors, or only the shareholders and directors who acted on a fraudulent basis, be restricted from obtaining business from any organ of state for a period not exceeding 10 years, after the audi alteram partem (hear the other side) rule has been applied; and
 - (e) forward the matter for criminal prosecution, if deemed necessary.

_____ SIGNATURE(S) OF TENDERER(S)
SURNAME AND NAME: _____
DATE: _____
ADDRESS: _____ _____ _____

6.2 F-100 THERAPEUTIC FEED

Description: F-100 is a therapeutic feed with added vegetable fat, carbohydrates, vitamins and minerals intended for the rehabilitation phase of treatment of children diagnosed with Severe Acute Malnutrition with an energy density of 100 kcal/100ml. F-100 is a food for special medical purposes, which is administered under medical supervision.

Indication for Use: Children aged 6 months and over diagnosed with Severe Acute Malnutrition, in phase 2, or nutritional rehabilitation phase of their treatment.

Packaging:

Item no:	42231802-00410	In 300-500g
Item no:	42231802-00217	In 100-150 ml RTU container
Item no:	42231802-00218	In 200-250ml RTU container
Item no:	42231802-00219	In 400-600ml RTU container
Item no:	42231802-00220	In 1000ml RTU container

TECHNICAL SPECIFICATIONS

Quality and Safety

Milk based white or pale yellowish fine powder or liquid; free from impurities, coloured particles, caking or lumps.

Ingredients:

- milk powder
- refined vegetable oil
- Sugar
- Maltodextrin
- milk derivate
- emulsifier (e.g., lecithin)
- vitamin and mineral (optionally premix can be used).

All ingredients, including optional ingredients, shall be clean, of good quality, safe and with minimal fibre removed when necessary. All ingredients and food additives shall be gluten free. F-100 shall be free from objectionable matter. It shall not contain any toxic substances originating from microorganisms or any other poisonous or deleterious substances, including anti-nutritional factors, heavy metals or pesticide residues in amounts that may represent a hazard to health. It shall not contain detectable levels of residues of antibiotics or other veterinary drugs used in animal husbandry.

Formulation and Starting materials F100 shall be manufactured referencing the formula described in the WHO document: Management of severe malnutrition: a manual for physicians and other senior health workers World health organization, 1999 (refer to Table 7, Table 8 and Appendix 4). <http://www.who.int/nutrition/publications/severemalnutrition/9241545119/en/>

The product must provide at least 50% of protein in the form of dairy protein. After reconstitution of powdered products according to the manufacturer's preparation instruction and ready to use liquid products, the product shall be a homogenous liquid that does not separate into oil/liquid phases or leave a **solid sediment** upon standing. Frothing of the F-100 powdered product after preparation should be minimal to enable accurate dosage measurements of the feed to each individual recipient. **The product should have a characteristic milk taste and smell. It shall be white as cream, shall not have rancid, pungent or unpleasant taste or smell.**

All ingredients, including optional ingredients, shall be clean, of good quality, safe and with minimal fibre removed when necessary. All ingredients and food additives shall be gluten free. Applicable local regulations and codex references for ingredients must be adhered to.

Milk: Full cream milk powder or Skimmed milk powder and/or Whey powder (NB may produce bitter taste) may be used in the formulation of F-100. **The applicable standards reference are Codex Standard for Milk Powders and Cream Powder (CXS 207-1999) and Codex Standard for Whey Powders (CXS 289- 1995).**

Carbohydrates: Carbohydrates used shall be gluten free and readily soluble in water. Glucose polymers and lactose may be used. **Applicable standards reference is Codex Standard for Sugars (CXS 212 – 1999).**

Oil: Edible refined vegetable oil. The manufacturer shall choose the type of oil and establish specifications for oil to ensure that the specifications in finished product are met (with particular attention to requirements for omega 3 and omega 6 fatty acids. Hydrogenated vegetable oils are not to be used. Applicable standards reference: Codex Standard for Named Vegetable Oils (CXS 210 - 1999).

Protein: F-100 approved for the rehabilitation phase of treatment of children with severe acute malnutrition must provide at least 50% of protein in the form of dairy protein. For the purposes of conversion from unit weight to unit energy, protein and carbohydrate are assumed to contribute 4 kcal/g and lipids 9 kcal/g.

Additives and Flavours: The use of artificial flavourings is not permitted; only natural flavourings may be used. Natural flavourings are defined in Codex Guidelines on General Requirements for Natural Flavourings (CXG 29-1987). The use of artificial antioxidants is not permitted, only natural antioxidants such as ascorbyl palmitate or mixed tocopherols may be used.

Other additives: Essential L-amino acids, choline, taurine, carnitine, inositol, carotene and other semi-essential or biologically valuable nutrients may be added to meet the specification at levels considered safe for children with severe malnutrition.

Shelf life: The product shall retain the above-mentioned specifications for at least 18-24 months for powder and for the liquid 9-12 months from date of manufacture when stored in dry conditions at a temperature of 30°C, supported by real time shelf-life data. Shelf-life studies shall be conducted in accordance with the Requirements for stability study as per WHO standards.

Microbiological criteria: Manufacturers are responsible for ensuring the compliance of finished products with the microbiological criteria as specified. With regards to limitations of the end-product testing the compliance shall be ensured through the design of an appropriate food safety control system and verification of the effectiveness of the applied control measures through appropriate auditing methods, including review of monitoring records, deviations and assuring that critical control points (CCPs) are kept under control and good hygienic practices (GHPs) are adhered to. These activities can be supplemented, as necessary, by appropriately documented microbiological sampling and analysis plans. The microbiological testing shall include, as appropriate, analysis of samples taken from starting materials, environment, production line and finished product. Environmental samples shall be taken from both: the points considered as most likely to cause product contamination and being most contaminated. Where results of monitoring the control measures and surveillance or verification indicate that the product batch, lot or consignment, or its part is unsafe (not in compliance with the set microbiological criteria), it shall be presumed that all the food in that batch, lot or consignment is also unsafe, unless following a detailed assessment there is no evidence that the rest of the batch, lot or consignment is unsafe.

Processing requirements: All processing and drying shall be carried out in a manner that minimizes loss of nutritive value, particularly protein quality and vitamin content. Products must be manufactured in accordance with the Codex Code of Hygienic Practice for Powdered Formulae for Infants and Young Children. (CXC 66 – 2008) and Recommended International Code of Practice. General Principles of Food Hygiene (CAC/RCP 1-1969, Rev. 4-2003), and other applicable codex references and GMPs (Good manufacturing practices).

Pesticides, Heavy Metals and Contaminants

Verifying that pesticide and heavy metals levels are below accepted limits is the responsibility of the manufacturer. Examples of mycotoxins, pesticides and heavy metals that must be controlled, include, but are not limited to:

- Nitrates < 200mg NO₃/kg
- Nitrites < 2mg NO₂/kg
- Aluminium < 0.6mg/kg
- Melamine < 1mg/kg

Mycotoxins (as per Codex standard when applicable for the starting materials used)

- Ochratoxin A <0.5ppb
- Aflatoxin B1 <0.1ppb
- Aflatoxin M1 <0.025ppb
- Palutin <10ppb
- Deoxynivalenol <200ppb
- Zearalenone <20ppb
- Fumonisin <200ppb

Applicable Codex standards references include: Code of Practice for Source Directed Measures to Reduce Contamination of Food with Chemicals (CXC 49-2001); General Methods of Analysis for Contaminants (CXS 228-2001); and Codex General Standard for Contaminants and Toxins in Food (CXS 193-1995).

Pesticides: Pesticide levels must be below 10 ppb. Verifying pesticide levels are below accepted limits is the responsibility of the manufacturer.

- Carbamates <10 ppb
- Organochlorines <10 ppb
- Organophosphates <10 ppb
- Pyrethroids <10 ppb

The maximum residue levels of specific pesticides or their metabolites in F100 set in below shall not be exceed:

Substance and Maximum residue level (mg/kg)

- Cadusafos: 0.006 Demeton-S-methyl/demeton-S.methyl sulfone/ oxydemeton-methyl (individually or combined, expressed as demeton-S-methyl) 0.006
- Ethoprophos 0.008
- Fipronil (sum of fipronil and fipronil-desifinyl, expressed as finpronil) 0.004
- Propineb/propylenethiourea (sum of propined and propylenethiourea) 0.006

The following pesticides shall not be used in the agricultural production intended for the production of F100:

- Disulfoton (sum of disulfoton, disulfoton sulfoxide and disulfoton sulfone, expressed as disulfoton) Fensulfotion (sum o fensulfotion, its oxygen analogue and their sulfone, expressed as fensulfotion) Fentin, expressed as triphenyltin cation

- Haloxfop (sum of haloxyfop, its salts and esters including conjugates, expressed as haloxyfop) Hexachlorobenzene Nitrofen
- Ometholate
- Terbufos (sum of terbufos, ist sulfoxide and sulfone, expressed as terbufos)
- Aldrin and dieldrin, expressed as dieldrin, Endrin

Applicable Codex standards reference is Analysis of Pesticide Residues: Recommended Methods (CXS 229-1993, REV.1-2003)

Heavy metals*: Verifying that heavy metal levels are below accepted limits is the responsibility of the manufacturer. Examples of heavy metals that must be controlled include, but are not limited to:

- Arsenic <0.052mg/kg
- Cadmium <0.112mg/kg
- Lead <0.2mg/kg
- Mercury<0.037mg/kg
- Tin <105mg/kg

*Based on 5 kg child with SAM and PTWI, General Standard for Contaminants and Toxins in Food and feed (CXS 193- 1995).

- Hydrocarbons Benzo[a]pyrene <1ppb

Radioactivity: Radioactive contamination can occur when using milk powder derived from cows that have eaten contaminated feed. This risk is managed by using certified radioactivity free milk products. The nuclear radiation level shall meet the values valid in the area of consumption. If limits are not defined, the value must not exceed 370Bq/kg (Cs 134 & Cs136). The product and its components shall not be treated by ionizing radiation.

Nutritional Composition

Table 1: Nutritional Composition per 100ml

Nutrients and nutritional values per 100ml finished product	Unit	Minimum	Maximum
Energy	kcal	95	105
Protein	g	2.3	3.1
Fat	g	4.9	6.9
Carbohydrates	g	7	12
w-3 fatty acids	0.3 – 2.5% total energy		
w-6 fatty acids	3 – 10% total energy		
Ash	Maximum 4.0%		
Moisture	Maximum 2.5%		
Lactose	G		4.4
Osmolarity	mMol/L	260	320
Vitamin A	mg RE	0.15	0.32
Thiamin	mg	0.1	
Riboflavin	mg	0.3	
Niacin	mg	1.0	
Pantothenic Acid	mg	0.6	

Pyridoxine	mg	0.1	
Biotin	mcg	11	
Folic acid	mcg	38	
Vitamin B12	mcg	0.3	
Vitamin C	mg	9.6	
Vitamin D	mcg	3.0	5.3
Vitamin E	mg	4.0	6.5
Vitamin K	mcg	3.0	
Calcium	mg	55	115
Copper	mg	0.25	0.35
Iodine	mcg	13	27
Iron	mg		0.05
Magnesium	mg	15	25
Phosphorus	mg	55	115
Potassium	mg	210	270
Selenium	mcg	3.5	7.7
Sodium	mg		55
Zinc	mg	2.0	3.0

Vitamins and minerals

- The used nutrient compounds shall comply with the criteria established in Advisory lists of Nutrient Compounds for use in foods for Special Dietary uses for Infants and Young Children (CXG 10 – 1979 (Rev. 2008 last amendment 2015).
- Another list of acceptable vitamin compounds can be found in Annex 3 of the COMMISSION DIRECTIVE 2006/141/EC of 22 December 2006 on infant formulae, follow-on formulae and amending Directive 1999/21/EC.
- If the manufacturer uses a mineral and vitamin premix(es), they must source it from an approved GMP manufacturer.
- Vitamins and minerals shall be in such forms that they are easily absorbed by patients with SAM who are often achlorhydric. The added minerals shall be water-soluble and shall not form insoluble components when mixed together. Iron salts are not to be added.
- F-100 shall have a mineral composition that will not alter the acid-base metabolism of patients with SAM. In particular, it shall have a moderate positive non- metabolisable base sufficient to eliminate the risk of metabolic acidosis.
- The non-metabolisable base can be estimated using the formula: Estimated absorbed millimoles (sodium+potassium+calcium+magnesium) minus (phosphates + chlorides) See: http://www.who.int/maternal_child_adolescent/documents/a91065/en/
- Added minerals shall be in the form of water soluble salts. Minerals used shall be in forms that are known to be bioavailable, nitrite and nitrate salts shall not be used. Recommended forms of minerals can be found in Appendix 4, Management of Severe Malnutrition a manual for senior health workers. <http://www.who.int/nutrition/publications/severemalnutrition/9241545119/en/>
http://apps.who.int/iris/bitstream/10665/44295/1/9789280641479_eng.pdf?ua=1

Packaging

Primary packaging

All packaging material being in contact with food or intended to come in contact with food, including inks and glue shall be of food-contact grade. Product shall be packed in a suitable container (refer to special conditions of contract). Packaging under inert gas (nitrogen or carbon dioxide) prolongs

products shelf-life and is recommended. Packaging must be free of damage such as, but not limited to tears, cuts, holes, and abrasions through one or more layers, leakage of any seal. The closure seal must be free of wrinkles and occluded matters.

Powdered product containers should be hermetically sealed and resistant to humid and hot climates. Seal and container integrity for F100 products shall be adequate to withstand pressure changes associated with international distribution channels such as air transport. The powdered container shall be capped with a reusable lid to adequately close the container and protect its content from external contamination and humidity during storage. The opened powdered container can be used within a minimum of 4 weeks.

The label shall contain the following information:

- Generic name: F-100 Therapeutic Feed
- Clear statement: A Food for Special Medical Purposes for the dietary management of Children > 6 months with Severe Acute Malnutrition [For the rehabilitation phase (phase 2)].
- Use under medical supervision.

Minimum requirement for release of F-100

The manufacturer must elaborate and implement an analytical plan of the finished product, starting materials and the processing environment. All analytical test procedures must be described in sufficient details, e.g., the sampling plan, acceptance/release criteria, analytical methods. SANAS accredited laboratories or ISO 17025 certified laboratories shall preferably be used.

Certificate of Analysis (CoA) may be required for every batch supplied to End-Users and shall be forwarded prior to its shipment. The manufacturer is expected to perform the necessary finish product analysis in order to prove that the batch complies with the specification. In case vitamin and mineral premix is used it may be adequate to test for tracers. In case single addition of vitamins and minerals are used, then individual analysis of each component is expected.

The principal tests listed below must be performed in order to check if the quality of F-100 meets above requirements. The validated method of analysis for the used must be indicated with the test results. The Department of Health reserves the rights to request for additional analyses in case of further quality assessment.

Table 2: List of Compulsory Tests

Nutrients and nutritional values per 100ml finished product	Unit	Minimum	Maximum
Energy	kcal	95	105
Protein	G	2.3	3.1
Fat	G	4.9	6.9
Carbohydrates	G	7	12
Ash	Maximum 4.0%		
Moisture	Maximum 2.5%		
Osmolarity	mMol/L	260	320
Vitamin A	mg RE	0.15	0.32
Vitamin C	Mg	9.6	
Iron	Mg		0.3
Potassium	Mg	210	270

Sodium	mg		55
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The manufacturer must conduct at least one complete finished product analysis annually in order to verify that the finished product is manufactured according to this specification.

CATEGORY SIX: NUTRITION PRODUCTS FOR THE MANAGEMENT OF SEVERE ACUTE MALNUTRITION FOR CHILDREN 6 MONTHS-5 YEARS OF AGE

6.1 F-75 THERAPEUTIC FEED

Description: F-75 is a therapeutic feed with added vegetable fat, carbohydrates, vitamins and minerals intended for the stabilisation phase of treatment of children diagnosed with Severe Acute Malnutrition (SAM) with an energy density of 75 kcal/100ml. F-75 is a feed for special medical purposes which is administered under medical supervision. A cautious approach is required because of the child's fragile physiological state and reduced homeostatic capacity; hence, F-75 is not designed for weight gain.

Indication for Use: Children above the age of 6 months diagnosed with Severe Acute Malnutrition, in phase 1 of their treatment.

Packaging:

Item number:	42231802-00409	In 300g-500g container
Item no:	42231802-00213	In 100ml-150ml RTU container
Item no:	42231802-00214	In 200ml-250 ml RTU container
Item no:	42231802-00215	In 400ml-600ml RTU container
Item no:	42231802-00216	In 1000ml RTU container

Quality and Safety

General quality milk based white or pale yellowish fine powder or liquid and should be free from impurities, coloured particles, caking and lumps.

Ingredients:

- milk powder
- refined vegetable oil
- Sugar
- Maltodextrin
- milk derivate
- emulsifier (e.g. lecithin)
- vitamin and mineral (optionally premix can be used).

F75 shall be free from objectionable matter. It shall not contain any toxic substances originating from microorganisms or any other poisonous or deleterious substances, including anti-nutritional factors, heavy metals or pesticide residues in amounts that may represent a hazard to health. It shall not contain detectable levels of residues of antibiotics or other veterinary drugs used in animal husbandry. Formulation and Starting materials in F75 shall be manufactured referencing the formula described in the WHO document: Management of severe malnutrition: a manual for physicians and other senior health workers World health organization, 1999 (refer to Table 7, Table 8 and Appendix 4). <http://www.who.int/nutrition/publications/severemalnutrition/9241545119/en/>

The product must provide at least 50% of protein in the form of dairy protein. After reconstitution of powdered products according to the manufacturer's preparation instruction and ready to use liquid products, the product shall be a homogenous liquid that does not separate into oil/liquid phases or leave a **solid sediment** upon standing. Frothing of the F75 powdered product after preparation should be minimal to enable accurate dosage measurements of the feed to each individual recipient. **The product should have a characteristic milk taste and smell. It shall be white as cream, shall not have rancid, pungent or unpleasant taste or smell.**

All ingredients, including optional ingredients, shall be clean, of good quality, safe and with minimal fibre removed when necessary. All ingredients and food additives shall be gluten free. Applicable local regulations and codex references for ingredients must be adhered to.

Milk: Full cream milk powder or Skimmed milk powder and/or Whey powder (NB: may produce bitter taste) may be used in the formulation of F-75. **The applicable standards reference are Codex Standard for Milk Powders and Cream Powder (CXS 207-1999) and Codex Standard for Whey Powders (CXS 289- 1995).**

Carbohydrates: Carbohydrates used shall be gluten free and readily soluble in water. Lactose shall not be added. Glucose polymers can be used. **Applicable standards reference is Codex Standard for Sugars (CXS 212 – 1999).**

Oil: Edible refined vegetable oil. The manufacturer shall choose the type of oil and establish specifications for oil to ensure that the specifications in the finished product are met (with particular attention to requirements for omega 3 and omega 6 fatty acids). Hydrogenated vegetable oils are not to be used. Applicable standards reference: Codex Standard for Named Vegetable Oils (CXS 210 - 1999).

Protein: F75 approved for the initial feeding or starting phase of treatment of children with severe acute malnutrition must provide at least 50% of protein in the form of dairy protein. For the purposes of conversion from unit weight to unit energy, protein and carbohydrate are assumed to contribute 4 kcal/g and lipids 9 kcal/g.

Additives and Flavours: The use of artificial flavourings is not permitted; only natural flavourings may be used. Natural flavourings are defined in Codex Guidelines on General Requirements for Natural Flavourings (CXG 29-1987). The use of artificial antioxidants is not permitted, only natural antioxidants such as ascorbyl palmitate or mixed tocopherols may be used.

Other additives: Essential L-amino acids, choline, taurine, carnitine, inositol, carotene and other semi-essential or biologically valuable nutrients may be added to meet the specification at levels considered safe for children with severe acute malnutrition.

Shelf life: The product shall retain the above-mentioned specifications for at least 18-24 months for powder and 9-12 months for liquids from date of manufacture when stored in dry conditions at a temperature of 30°C, supported by real time shelf life data. Shelf life studies shall be conducted in accordance with the Requirements for stability studies as per the WHO standards.

Microbiological criteria: Manufacturers are responsible for ensuring the compliance of finished products with the microbiological criteria as specified. In regard to limitations of the end-product testing the compliance shall be ensured through the design of an appropriate food safety control system and verification of the effectiveness of the applied control measures through appropriate auditing methods, including review of monitoring records, deviations and assuring that critical control points (CCPs) are kept under control and good hygienic practices (GHPs) are adhered to. These activities can be supplemented, as necessary, by appropriately documented microbiological sampling and analysis plans. The microbiological testing shall include, as appropriate, analysis of samples taken from starting materials, environment, production line and finished product. Environmental samples shall be taken from both: the points considered as most likely to cause product contamination and being most contaminated. **Where results of monitoring the control measures and surveillance or verification indicate that the product batch, lot or consignment, or its part is unsafe (not in compliance with the set microbiological criteria), it shall be presumed that all the food in that batch,**

lot or consignment is also unsafe, unless following a detailed assessment there is no evidence that the rest of the batch, lot or consignment is unsafe.

Processing requirements: All processing and drying shall be carried out in a manner that minimizes loss of nutritive value, particularly protein quality and vitamin content. Products must be manufactured in accordance with the Codex Code of Hygienic Practice for Powdered Formulae for Infants and Young Children, (CXC 66 – 2008) and Recommended International Code of Practice, General Principles of Food Hygiene (CAC/RCP 1-1969, Rev. 4-2003), and other applicable codex references and GMPs (Good manufacturing practices).

Pesticides, Heavy Metals and Contaminants

Verifying that pesticide and heavy metals levels are below accepted limits is the responsibility of the manufacturer. Examples of mycotoxins, pesticides and heavy metals that must be controlled, include, but are not limited to:

- Nitrates < 200mg NO₃/kg
- Nitrites < 2mg NO₂/kg
- Aluminium < 0.6mg/kg
- Melamine < 1mg/kg

Mycotoxins (as per Codex standard when applicable for the starting materials used)

- Ochratoxin A <0.5ppb
- Aflatoxin B1 <0.1ppb
- Aflatoxin M1 <0.025ppb
- Palutin <10ppb
- Deoxynivalenol <200ppb
- Zearalenone <20ppb
- Fumonisin <200ppb

Applicable Codex standards references include: Code of Practice for Source Directed Measures to Reduce Contamination of Food with Chemicals (CXC 49-2001); General Methods of Analysis for Contaminants (CXS 228-2001); and Codex General Standard for Contaminants and Toxins in Food (CXS 193-1995).

Pesticides: Pesticide levels must be below 10 ppb. Verifying pesticide levels are below accepted limits is the responsibility of the manufacturer.

- Carbamates <10 ppb
- Organochlorines <10 ppb
- Organophosphates <10 ppb
- Pyrethroids <10 ppb

The maximum residue levels of specific pesticides or their metabolites in F75 set in below shall not be exceeded:

Substance and Maximum residue level (mg/kg)

- Cadusafos: 0.006 Demeton-S-methyl/demeton-S.methyl sulfone/ oxydemeton-methyl (individually or combined, expressed as demeton-S-methyl) 0.006
- Ethoprophos 0.008
- Fipronil (sum of fipronil and fipronil-deslfinyl, expressed as finpronil) 0.004
- Propineb/propylenethiourea (sum of propined and propylenethiourea) 0.006

The following pesticides shall not be used in the agricultural production intended for the production of F75:

- Disulfoton (sum of disulfoton, disulfoton sulfoxide and disulfoton sulfone, expressed as disulfoton) Fensulfotion (sum o fensulfothion, its oxygen analogue and their sulfone, expressed as fensulfothion) Fentin, expressed as triphenyltin cation
- Haloxfop (sum of haloxyfop, its salts and esters including conjugates, expressed as haloxyfop) Hexachlorobenzene Nitrofen
- Ometholate
- Terbufos (sum of terbufos, ist sulfoxide and sulfone, expressed as terbufos)
- Aldrin and dieldrin, expressed as dieldrin, Endrin

Applicable Codex standards reference is Analysis of Pesticide Residues: Recommended Methods (CXS 229-1993, REV.1-2003)

Heavy metals*: Verifying that heavy metal levels are below accepted limits is the responsibility of the manufacturer. Examples of heavy metals that must be controlled include, but are not limited to:

- Arsenic <0.052mg/kg
- Cadmium <0.112mg/kg
- Lead <0.2mg/kg
- Mercury<0.037mg/kg
- Tin <105mg/kg

*Based on 5 kg child with SAM and PTWI, General Standard for Contaminants and Toxins in Food and feed (CXS 193- 1995).

- Hydrocarbons Benzo[a]pyrene <1ppb

Radioactivity: Radioactive contamination can occur when using milk powder derived from cows that have eaten contaminated feed. This risk is managed by using certified radioactivity free milk products. The nuclear radiation level shall meet the values valid in the area of consumption. If limits are not defined, the value must not exceed 370Bq/kg (Cs 134 & Cs136). The product and its components shall not be treated by ionizing radiation.

Nutritional Composition

Table 1: Nutritional Composition per 100ml

Nutrients and nutritional values per 100ml finished product	Unit	Minimum	Maximum
Energy	kcal	70	80
Protein	g	0.75	1.5
Fat	g	2.0	3.0
w-3 fatty acids	0.3 – 2.5% total energy		
w-6 fatty acids	3 – 10% total energy		
Ash	Maximum 4.0%		
Moisture	Maximum 4%		
Lactose	<1.4g		
Osmolarity	mMol/L	240	320
carbohydrates	g	10.5	14
Vitamin A	mg RE	0.1	0.3
Thiamin	mg	0.08	

Riboflavin	mg	0.3	
Niacin	mg	0.8	
Pantothenic Acid	mg	0.5	
Pyridoxine	mg	0.1	
Biotin	mcg	10	
Folic acid	mcg	35	
Vitamin B12	mcg	0.3	
Vitamin C	mg	10	
Vitamin D	mcg	2.5	5.0
Vitamin E	mg	3.3	6.5
Vitamin K	mcg	2.5	
Calcium	mg	50	100
Copper	mg	0.2	0.3
Iodine	mcg	12.3	24.5
Iron	mg		0.05
Magnesium	mg	8.5	11
Phosphorus	mg	50	100
Potassium	mg	122	156
Selenium	mcg	3.5	7
Sodium	mg	17	17
Zinc	mg	1.8	3.0

Vitamins and minerals

- The used nutrient compounds shall comply with the criteria established in Advisory lists of Nutrient Compounds for use in foods for Special Dietary uses for Infants and Young Children (CXG 10 – 1979 (Rev. 2008 last amendment 2015).
- Another list of acceptable vitamin compounds can be found in Annexure 3 of the COMMISSION DIRECTIVE 2006/141/EC of 22 December 2006 on infant formulae, follow-on formulae and amending Directive 1999/21/EC.
- If the manufacturer uses a mineral and vitamin premix(es), they must source it from a premix manufacturer/s who adhere to GMP standards.
- Vitamins and minerals shall be in such forms that they are easily absorbed by patients with SAM who are often achlorhydric. The added minerals shall be water-soluble and shall not form insoluble components when mixed together. **Iron salts are not to be added.**
- The F75 shall have a mineral composition that will not alter the acid-base metabolism of patients with SAM. In particular, it shall have a moderate positive non-metabolisable base sufficient to eliminate the risk of metabolic acidosis.
- The non-metabolisable base can be estimated using the formula: Estimated absorbed millimoles (sodium+potassium+calcium+magnesium) minus (phosphates + chlorides) See: http://www.who.int/maternal_child_adolescent/documents/a91065/en/
- Added minerals shall be in the form of water-soluble salts. Minerals used shall be in forms that are known to be bioavailable, nitrite and nitrate salts shall not be used. Recommended forms of minerals can be found in Appendix 4, Management of Severe Malnutrition a manual for senior health workers. <http://www.who.int/nutrition/publications/severemalnutrition/9241545119/en/>
http://apps.who.int/iris/bitstream/10665/44295/1/9789280641479_eng.pdf?ua=1

Packaging

Primary packaging

All packaging material being in contact with food or intended to come in contact with food, including inks and glue shall be of food-contact grade. Product shall be packed in a suitable container (refer to special conditions of contract). Packaging under inert gas (nitrogen or carbon dioxide) prolongs products shelf-life and is recommended. Packaging must be free of damage such as, but not limited to tears, cuts, holes, and abrasions through one or more layers, leakage of any seal. The closure seal must be free of wrinkles and occluded maters.

Powdered product containers should be hermetically sealed and resistant to humid and hot climates. Seal and container integrity for F75 products shall be adequate to withstand pressure changes associated with international distribution channels such as air transport. The powdered container shall be capped with a reusable lid to adequately close the container and protect its content form external contamination and humidity during storage. The opened powdered container can be used within a minimum of 4 weeks.

The label shall contain the following information:

- Generic name: F-75 Therapeutic Feed
- Clear statement: For the initial phase (or Phase 1) of treatment of Children > 6 months with Severe Acute Malnutrition.
- Use under medical supervision.

Minimum requirement for release of F 75

The manufacturer must elaborate and implement an analytical plan of the finished product, starting materials and the processing environment. All analytical test procedures must be described in sufficient details, e.g. the sampling plan, acceptance/release criteria, analytical methods. SANAS accredited laboratories or ISO 17025 certified laboratories shall preferably be used.

Certificate of Analysis (CoA) may be required for every batch supplied to End-Users and shall be forwarded prior to its supply. The manufacturer is expected to perform the necessary finish product analysis in order to prove that the batch complies with the specification. In case vitamin and mineral premix is used it may be adequate to test for tracers. In case single addition of vitamins and minerals are used, then individual analysis of each component is expected.

The principal tests listed below must be performed in order to check if the quality of F-75 meets above requirements. The validated method of analysis for the used must be indicated with the test results. The Department of Health reserves the rights to request for additional analyses in case of further quality assessment.

Table 2: List of Compulsory Tests per 100ml

Nutrients and nutritional values per 100ml finished product	Unit	Minimum	Maximum
Energy	kcal	70	80
Protein	g	0.75	1.5
Fat	g	2.0	3.0
Ash	Maximum 4.0%		
Moisture	Maximum 4%		
Lactose	<1.4g		

Osmolarity	mMol/L	240	320
Carbohydrates	g	10.5	14
Vitamin A	mg RE	0.1	0.3
Vitamin C	mg	10	
Vitamin D	mcg	2.5	5.0
Iron	mg		<0.3
Potassium	mg	122	156
Sodium	mg		17

The manufacturer must conduct at least one complete finished product analysis annually in order to verify that the finished product is manufactured according to this specification.



Physical Address: 110189, Selaywe Road, Pietermaritzburg
Tel: 033-3195-4000 Fax: 033-3195-4000 Email: info@kznhealth.gov.za
www.kznhealth.gov.za

QUOTATION NO.	ZNQ/EDN 1593/24-25
QUOTATION DESCRIPTION	Baby feeds
BIDDER NAME	

EVALUATION CRITERIA:

The Institution intends to evaluate valid quotations using four (4) evaluation stages. These are peremptory requirements, should the bidder/ tenderer fail to comply with any of the stages as stated below, the quotation will be regarded as non-responsive, and will not progress to the final stage of evaluation:

STAGE 1: ADMINISTRATIVE, COMPULSORY COMPLIANCE AND MANDATORY REQUIREMENTS

STAGE 2: CAPACITY TO DELIVER

STAGE 3: COMPLIANCE WITH SPECIFICATION

STAGE 4: PRICE AND PREFERENCE POINTS

STAGE 1: ADMINISTRATIVE, COMPULSORY COMPLIANCE AND MANDATORY REQUIREMENTS

NO.	REQUIREMENTS	INCLUDED IN THE PUBLISHED DOCUMENT?	TO BE RETURNED BY BIDDER/ TENDERER?
Administrative Compliance			
1.	PARTICULARS OF QUOTATION	YES	YES
2.	OFFICIAL PRICE PAGE FOR QUOTATIONS OVER R2 000.01, MUST BE CALCULATED CORRECTLY AND IN COMPLIANCE WITH SARS REQUIREMENTS	YES	YES
3.	BIDDER'S DISCLOSURE (SBD4)	YES	YES
4.	GENERAL CONDITIONS OF CONTRACT (GCC)	YES	YES
5.	SPECIAL CONDITIONS OF CONTRACT (SCC)	YES	YES
6.	PREFERENCE POINTS CLAIM FORM IN TERMS OF THE PREFERENTIAL PROCUREMENT REGULATIONS 2022 (SBD 6.1)	YES	YES
Compulsory Compliance			
7.	A B-BBEE STATUS LEVEL VERIFICATION CERTIFICATE/ SWORN AFFIDAVIT (For EMEs & QSEs)	NO	YES
8.	COPY OF CENTRAL SUPPLIER DATABASE COMPLIANCE REPORT (CSD)	NO	YES
9.	TAX COMPLIANT AS PER SARS REQUIREMENTS	NO	YES
Mandatory Requirements			
10.	NOT APPLICABLE	NO	YES

Note: This relates to administrative, compulsory and mandatory returnable documents which must be fully completed, and submitted, should you fail to submit any of the above returnable documents, your offer will be treated as non-responsive and will not proceed to the next stage of evaluation. The department reserve a right to verify validity of the documents submitted, should it be discovered that the information submitted is misrepresented the quotation will be disqualified.

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STAGE 2: CAPACITY TO DELIVER

1.	If there is valid proof that the bidder was previously issued with an order and failed to deliver without acceptable reasons, the bidder will be treated as a defaulter and will not progress to the next stage of evaluation.
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STAGE 3: COMPLIANCE WITH SPECIFICATION

Requirement	Complies With Specification Yes /No
The bidder / Tenderer to confirm that the work to be supplied comply with attached specification document, should you fail to indicate with yes, it will mean, you do not comply and your quotation will not progress to the next stage of evaluation	

STAGE 4: PRICE AND PREFERENCE POINTS

The value of this quotation is estimated not to exceed R 500 000 (inclusive of all applicable taxes), therefore the 80/20 preference point system shall be applicable. Points for this quotation will be awarded for:

CATEGORY	POINTS
PRICE	80
SPECIFIC GOALS	20
Total points for Price and must not exceed	100

The Department has identified the following specific goal:

Specific Goal	Number of Points allocated	Proof To Claim Specific Goal (Returnable Documents)
Race – Full/partial/combination of points allocated to companies at least 51% owned by Black People	20	<ol style="list-style-type: none"> 1. Ownership Certificate issued by the Companies and Intellectual Property Commission (CIPC). 2. A B-BBEE status level/ sworn affidavit. 3. The Department will download CSD to verify this information.
<p>NOTE: Should a responsive bidder fail to submit proof to claim points, as stated above this will not result in disqualification, however the bidder will not be awarded points for specific goals.</p>		

SIGNATURE: 

PS HLELA
DD: FINANCE AND SCM

DATE: 13.09.2024

Name and surname of the bidder: _____

Signature _____

Date: _____

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