

Quotation Advert

Opening Date: 2020-06-12
Closing Date: 2020-06-22
Closing Time: 11:00

INSTITUTION DETAILS

Institution Name: King Cetshwayo district office
Province: KwaZulu-Natal
Department or Entity: Department of Health
Division or section: Central Supply Chain Management
Place where goods / services is required King Cetshwayo Health District Office
Date Submitted 2020-06-11

ITEM CATEGORY AND DETAILS

Quotation Number: ZNQ:
101/DC28/2021
Item Category: Goods
Item Description: SYRINGES 10ML (BOX OF 100 UNITS) X 05 BOXES
NEEDLES 18G (BOX OF 100 UNITS) X 05 BOXES
BODY BAGS XXL (BAG OF 25 UNITS) X 60 BAGS
BODY BAGS XL (BAG OF 25 UNITS) X 60 BAGS
TOXICOLOGY KIT X 25 UNITS
HEAVY DUTY POST MORTEM HAND SAW X 04 UNITS

Quantity (if supplies)

COMPULSORY BRIEFING SESSION / SITE VISIT

Select Type: Not Applicable
Date :
Time:
Venue:

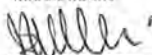
QUOTES CAN BE COLLECTED FROM: KING CETSHWAYO HEALTH DISTRICT OFFICE 2ND LOOD AVENUE
EMPANGENI RAIL 3880

QUOTES SHOULD BE DELIVERED TO: KING CETSHWAYO HEALTH DISTRICT OFFICE 2ND LOOD AVENUE
EMPANGENI RAIL 3880

ENQUIRIES REGARDING THE ADVERT MAY BE DIRECTED TO:

Name: MR N.N NDLOVU
Email: nozipho.myaka@kznhealth.gov.za
Contact Number: 035 787 6287
Finance Manager Name: Mrs S.L Msane

Finance Manager Signature:



No late quotes will be considered

DECLARATION OF INTEREST

1. Any legal person, including persons employed by the state¹, or persons having a kinship with persons employed by the state, including a blood relationship, may make an offer or offers in terms of this invitation to quote (includes a price quotation, advertised competitive quote, limited quote or proposal). In view of possible allegations of favouritism, should the resulting quote, or part thereof, be awarded to persons employed by the state, or to persons connected with or related to them, it is required that the bidder or his/her authorised representative declare his/her position in relation to the evaluating/adjudicating authority where-
 - the bidder is employed by the state; and/or
 - the legal person on whose behalf the bidding document is signed, has a relationship with persons/a person who are/is involved in the evaluation and or adjudication of the quote(s), or where it is known that such a relationship exists between the person or persons for or on whose behalf the declarant acts and persons who are involved with the evaluation and or adjudication of the quote.
2. In order to give effect to the above, the following questionnaire must be completed and submitted with the quote.

- | | |
|--|---|
| 2.1. Full Name of bidder/representative..... | 2.4. Company Registration Number: |
| 2.2. Identity Number: | 2.5. Tax Reference Number: |
| 2.3. Position occupied in the Company (director, trustee, shareholder ²):..... | 2.6. VAT Registration Number: |

2.7. The names of all directors / trustees / shareholders / members, their individual identity numbers, tax reference numbers and, if applicable, employee / persal numbers must be indicated in paragraph 3 below. [TICK APPLICABLE]

2.8. Are you or any person connected with the bidder presently employed by the state? YES NO

2.8.1. If so, furnish the following particulars:
 Name of person / director / trustee / shareholder/ member:
 Name of state institution at which you or the person connected to the bidder is employed:.....
 Position occupied in the state institution: Any other particulars:.....

2.8.2. If you are presently employed by the state, did you obtain the appropriate authority to undertake remunerative work outside employment in the public sector? YES NO

2.8.2.1. If yes, did you attach proof of such authority to the quote document?

(Note: Failure to submit proof of such authority, where applicable, may result in the disqualification of the quote.)

2.8.2.2. If no, furnish reasons for non-submission of such proof:

2.9. Did you or your spouse, or any of the company's directors / trustees / shareholders / members or their spouses conduct business with the state in the previous twelve months? YES NO

2.9.1. If so, furnish particulars:.....

2.10. Do you, or any person connected with the bidder, have any relationship (family, friend, other) with a person employed by the state and who may be involved with the evaluation and or adjudication of this quote? YES NO

2.10.1. If so, furnish particulars:.....

2.11. Are you, or any person connected with the bidder, aware of any relationship (family, friend, other) between any other bidder and any person employed by the state who may be involved with the evaluation and or adjudication of this quote? YES NO

2.11.1. If so, furnish particulars:.....

2.12. Do you or any of the directors / trustees / shareholders / members of the company have any interest in any other related companies whether or not they are bidding for this contract? YES NO

2.12.1. If so, furnish particulars:.....

3. Full details of directors / trustees / members / shareholders.

NB: The Department Of Health will validate details of directors / trustees / members / shareholders on CSD. It is the suppliers' responsibility to ensure that their details are up-to-date and verified on CSD. If the Department cannot validate the information on CSD, the quote will not be considered and passed over as non-compliant according to National Treasury Instruction Note 4 (a) 2016/17.

4 DECLARATION

I, THE UNDERSIGNED (NAME).....CERTIFY THAT THE INFORMATION FURNISHED IN PARAGRAPHS 2.

I ACCEPT THAT THE STATE MAY REJECT THE QUOTE OR ACT AGAINST ME SHOULD THIS DECLARATION PROVE TO BE FALSE.

.....
Name of bidder	Signature	Position	Date

¹"State" means –

- | | |
|---|---|
| a) any national or provincial department, national or provincial public entity or constitutional institution within the meaning of the Public Finance Management Act, 1999 (Act No. 1 of 1999); | c) provincial legislature; |
| b) any municipality or municipal entity; | d) national Assembly or the national Council of provinces; or |
| | e) Parliament. |

²"Shareholder" means a person who owns shares in the company and is actively involved in the management of the enterprise or business and exercises control over the enterprise.

SPECIAL CONTRACT CONDITIONS OF QUOTATIONS

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 institution is under no obligation to accept the lowest or any quote.
- 3.2. The price quoted must include VAT (if VAT vendor). However, it must be noted that the department reserves the right to evaluate all quotations excluding VAT as some bidders may not be VAT vendors.
- 3.3. The bidder must ensure the correctness & validity of quote:
- (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*
- 3.4. 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.
- 3.5. This quotation will be evaluated based on the 80/20 points system, specification & correctness of information. All required documentation must be completed in full and submitted.
- 3.6. Offers must comply strictly with the specification.
- 3.7. Only offers that meet or are greater than the specification will be considered.
- 3.8. Late quotes will not be considered.
- 3.9. Expired product/s will not be accepted. All products supplied must be valid for a minimum period of six months.
- 3.10. A bidder not registered on the Central Suppliers Database or verification has failed will not be considered.
- 3.11. All delivery costs must be included in the quote price, for delivery at the prescribed destination.
- 3.12. 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.
- 3.13. In cases where different delivery points influence the pricing, a separate pricing schedule must be submitted for each delivery point.
- 3.14. In the event of a bidder having multiple quotes, only the cheapest according to specification will be considered. Furthermore a verification will be done to identify if bidders have multiple companies and are quoting (cover-quoting) for this bid. In such instances only the cheapest bid according to specification will be considered.

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. Quotation submitted must be complete in all respects.
- 4.5. Any alteration made by the bidder must be initialled.
- 4.6. Use of correcting fluid is prohibited
- 4.7. Quotation 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.

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.

- 5.5. No quotation/bid sent through the post will be considered if it is received after the closing date and time stipulated in the quotation documentation, and proof of posting will not be accepted as proof of delivery.
- 5.6. Quotation documents must not be included in packages containing samples. Such quotations may be rejected as being invalid.

6. SAMPLES

- 6.1. In the case of the quote document stipulating that samples are required, the supplier will be informed in due course when samples should be provided to the institution. (This decreases the time of safety and storage risk that may be incurred by the respective institution). The bidders sample will be retained if such bidder wins the contract.
 - (i) If a company/s who has not won the quote requires their samples, they must advise the institution in writing of such.
 - (ii) If samples are not collected within three months of close of quote the institution reserves the right to dispose of them at their discretion.
- 6.2. Samples must be made available when requested in writing or if stipulated on the document.
 - (i) If a Bidder fails to provide a sample of their product on offer for scrutiny against the set specification when requested, their offer will be rejected. All testing will be for the account of the bidder.

7. COMPULSORY SITE INSPECTION / BRIEFING SESSION

7.1. Bidders who fail to attend the compulsory meeting will be disqualified from the evaluation process.

- (i) The institution has determined that a compulsory site meeting take place
- (ii) Date ____/____/____ Time ____:____ Place _____

Institution Stamp:	Institution Site Inspection / briefing session Official Full Name: Signature: Date:
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8. STATEMENT OF SUPPLIES AND SERVICES

8.1. The contractor shall, when requested to do so, furnish particulars of supplies delivered or services executed. If he/she fails to do so, the Department may, without prejudice to any other rights which it may have, institute inquiries at the expense of the contractor to obtain the required particulars.

9. SUBMISSION AND COMPLETION OF SBD 6.1

9.1. Should a bidder wish to qualify for preference points they must complete a SBD 6.1 document. Failure by a bidder to provide all relevant information required, will result in such a bidder not being considered for preference point's allocation. The preferences applicable on the closing date will be utilized. Any changes after the closing date will not be considered for that particular quote.

10. TAX COMPLIANCE REQUIREMENTS

- 10.1. In the event that the tax compliance status has failed on CSD, *it is the suppliers' responsibility to provide a SARS pin in order for the institution to validate the tax compliance status of the supplier.*
- 10.2. In the event that the institution cannot validate the suppliers' tax clearance on SARS as well as the Central Suppliers Database, *the quote will not be considered and passed over as non-compliant according to National Treasury Instruction Note 4 (a) 2016/17.*

11. TAX INVOICE

11.1. A tax invoice shall be in the currency of the Republic of South Africa and shall contain the following particulars:

- (i) the name, address and registration number of the supplier;
- (ii) the name and address of the recipient;
- (iii) an individual serialized number and the date upon which the tax invoice is issued;
- (iv) a description and quantity or volume of the goods or services supplied;
- (v) the official department order number issued to the supplier;
- (vi) the value of the supply, the amount of tax charged;
- (vii) the words tax invoice in a prominent place.

12. PATENT RIGHTS

The supplier shall indemnify the KZN Department of Health (hereafter known as 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.

13. PENALTIES

- 13.1. If at any time during the contract period, the service provider is unable to perform in a timely manner, the service provider must notify the institution in writing/email of the cause of and the duration of the delay. Upon receipt of the notification, the institution should evaluate the circumstances and, if deemed necessary, the institution may extend the service provider's time for performance.
- 13.2. In the event of delayed performance that extends beyond the delivery period, the institution is entitled to purchase commodities of a similar quantity and quality as a substitution for the outstanding commodities, without terminating the contract, as well as return commodities delivered at a later stage at the service provider's expense.
- 13.3. Alternatively, the institution may elect to terminate the contract and procure the necessary commodities in order to complete the contract. In the event that the contract is terminated the institution may claim damages from the service provider in the form of a penalty. The service provider's performance should be captured on the service provider database in order to determine whether or not the service provider should be awarded any contracts in the future.
- 13.4. 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.

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. FAILURE TO COMPLY WITH ABOVE WILL RESULT IN YOUR QUOTE BEING PASSED OVER.

9. DECLARATION WITH REGARD TO COMPANY/FIRM

9.1 Name of company/firm:.....

9.2 VAT registration number:.....

9.3 Company registration number:.....

9.4 TYPE OF COMPANY/ FIRM [TICK APPLICABLE BOX]

- Partnership/Joint Venture / Consortium
- One person business/sole propriety
- Close corporation
- Company
- (Pty) Limited

9.5 DESCRIBE PRINCIPAL BUSINESS ACTIVITIES

.....
.....

9.6 COMPANY CLASSIFICATION [TICK APPLICABLE BOX]

- Manufacturer
- Supplier
- Professional service provider
- Other service providers, e.g. transporter, etc.

9.7 Total number of years the company/firm has been in business:.....

9.8 I/we, the undersigned, who is / are duly authorised to do so on behalf of the company/firm, certify that the points claimed, based on the B-BBE status level of contributor indicated in paragraphs 1.4 and 6.1 of the foregoing certificate, qualifies the company/ firm for the preference(s) shown and I / we 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 6.1, the contractor may be required to furnish documentary proof to the satisfaction of the purchaser that the claims are correct;
- iv) If the B-BBEE status level of contributor has been claimed or obtained on a fraudulent basis or any of the conditions of contract have not been fulfilled, the purchaser may, in addition to any other remedy it may have –
 - (a) disqualify the person from the bidding 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 bidder or contractor, its shareholders and directors, or only the shareholders and directors who acted on a fraudulent basis, be restricted by the National Treasury 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.

WITNESSES	
1.
2.

..... SIGNATURE(S) OF BIDDERS(S)
DATE:
ADDRESS.....



health

Department:
Health
PROVINCE OF KWAZULU-NATAL

DIRECTORATE:

2ND Lood Avenue, Kuleka & Chrome Crescent Corner, Empangeni Rail, Old Telkom Building
Private Bag X 20034, EMPANGENI, 3880
Tel: 035 787 0631 Fax: 035 787 0641/46
www.kznhealth.gov.za

King Cetshwayo District Office
Specification Document

DESCRIPTION	Mandatory requirements to be Completed by bidder
Syringe Hypodermic	COMPLY/ DO NOT COMPLY
- Syringe, hypodermic	
- Plastic, eccentric nozzle, luer slip, two-part syringe	
- Nominal capacity: 10ml	
- Single use, sterile, individually packed	
- To comply with relevant test of the latest issue of SANS 1124-2	
- Specification number 6063	
- 100 syringes per box	

Bidder's Comment:

.....
.....

1. The bidder is requested to put a company stamp on this specification document.
2. The bidder is requested to complete "Mandatory requirements to be completed by bidder" segment

NB: FAILURE TO COMPLY WITH THE ABOVE WILL RESULT FOR A BIDDER TO BE DISQUALIFIED.

Bidder's Official Stamp

Bidder Signature:



health

Department:
Health
PROVINCE OF KWAZULU-NATAL

DIRECTORATE:

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King Cetshwayo District Office
Specification Document

DESCRIPTION	Mandatory requirements to be Completed by bidder
Needle Hypodermic	COMPLY/ DO NOT COMPLY
- Hypodermic Needles	
- Latex free	
- Disposable	
- Regular bevel	
- Thin wall	
- Sterile, non-toxic and non-pyrogenic	
- Polypropylene hubs and boxes are colour coded for easy gauge identification	
- Each needle comes ready to use	
- Fits luer-lock, luerislip and Eccentric luer slip syringes.	
- Made of stainless steel and with a polypropylene hub	
- Size: 18g x 1.5"	
- Packaging: Box of 100 needles	

Bidder's Comment:

.....
.....

1. The bidder is requested to put a company stamp on this specification document.
2. The bidder is requested to complete "Mandatory requirements to be completed by bidder" segment

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Bidder's Official Stamp

Bidder Signature:



health

Department:
Health
PROVINCE OF KWAZULU-NATAL

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DIRECTORATE:

King Cetshwayo District Office
Specification Document

DESCRIPTION	Mandatory requirements to be Completed by bidder	
	YES	NO
Body bag		
- Micron Strength: 180 microns		
- Breaking strength: 200kg		
- Material: Poly propylene laminated, waterproof, 100% leak-proof, extremely water resistant.		
- Manufacture: double stitched, formulated folding U-shaped.		
- Colour: White		
- Size: XXL Large = 2600mm long X 1200mm wide.		
- Zip: U shaped nylon zip closure to facilitate placing the body in the bag, total leak proof.		
- Seams: Must be double stitched and heat bonded to prevent leakage of body fluids.		
- Handles: 3 handles of 50mm width on each side of bag. Webbed handles to stuck on and then stitched. Must be from one side to other to accommodate weight and maximum breaking strain of 200kg.		
- Ties: there will be 3 ties inside of the bag, at the top, middle and bottom.		
- Absorbent Pad: to be attached inside of the bag to absorb body fluids, size of pad to be from the head to the toes. Must be one complete pad.		
- Luggage Label: Transparent sleeve at the bottom right hand side, on the top surface of the bag. The label size: 12cm x 9cm.		

1. The bidder is requested to put a company stamp on this specification document.
2. The bidder is requested to complete "Mandatory requirements to be completed by bidder" segment

NB: FAILURE TO COMPLY WITH THE ABOVE WILL RESULT FOR A BIDDER TO BE DISQUALIFIED.

Bidder's Official Stamp

Bidder Signature: _____



health

Department:
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Specification Document

DESCRIPTION	Mandatory requirements to be Completed by bidder	
	YES	NO
Body Bag		
- Micron Strength: 180 microns		
- Breaking strength: 200kg		
- Material: Poly propylene laminated, waterproof, 100% leak-proof, extremely water resistant.		
- Manufacture: double stitched, formulated folding U-shaped.		
- Colour: White		
- Size: XLarge = 2400mm long X 1000mm wide.		
- Zip: U shaped nylon zip closure to facilitate placing the body in the bag, total leak proof.		
- Seams: Must be double stitched and heat bonded to prevent leakage of body fluids.		
- Handles: 3 handles of 50mm width on each side of bag. Webbed handles to stuck on and then stitched. Must be from one side to other to accommodate weight and maximum breaking strain of 200kg.		
- Ties: there will be 3 ties inside of the bag, at the top, middle and bottom		
- Absorbent Pad: to be attached inside of the bag to absorb body fluids, size of pad to be from the head to the toes. Must be one complete pad.		
- Luggage Label: Transparent sleeve at the bottom right hand side, on the top surface of the bag. The label size: 12cm x 9cm.		

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NB: FAILURE TO COMPLY WITH THE ABOVE WILL RESULT FOR A BIDDER TO BE DISQUALIFIED.

Bidder's Official Stamp

Bidder Signature:



health

Department:
Health

PROVINCE OF KWAZULU-NATAL

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DIRECTORATE:
UTHUNGULU DISTRICT OFFICE
SPECIFICATION DOCUMENT

DESCRIPTION	Mandatory requirements to be Completed by bidder COMPLY / DO NOT COMPLY
Toxicology Sampling Kit	
- Toxicology sampling kit for the collection and transportation of post-mortem biological fluids and human tissue for the purpose of forensic toxicological analysis, to be used in mainly in the Department of Health – Forensic Mortuaries.	
- Kit to comprise of clean suitable plastic sample containers that will not leak or break if dropped.	
- Packaged in a suitable container and labelled as specified below: Packaging, content, labelling and quality requirements are listed as follows:	
The following qualities is required:	
1. It is imperative that the kit must be prepared in a manner that would avoid the possibility of contamination.	
2. Quality control measures must be in place to prove this.	
3. The intention is for a standardized Toxicology Sample Kit to be procured at provincial level and used at mortuaries nationally for the collection of post-mortem samples from where it would be transported to the relevant Department of Health Forensic Chemistry Laboratories for the toxicological analysis of samples.	
4. The prescribed minimum specifications of the required Toxicology Sample Kit are stipulated under the following:	
THE BIDDER TO STATE:	
1. The sample containers must be of clean suitable plastic material that would not allow leaking of samples and would not break easily when dropped.	
2. The specimen containers must be suitably clean not to result in contamination of samples and to avoid interference on analytical results. Prior to the bid being awarded, samples of the containers will be requested for testing to verify the suitability thereof.	
3. Labels on the containers must be printed clearly, eg. Font size 14, in water fast ink on labels that would not come loose or get damaged when exposed to repeated freezing, thawing and wetting.	
4. The specimen containers must be supplied capped with self-sealing, leak-free lids with a watertight fit where relevant.	
5. The outer packaging must be sealed with tamper evident tape bearing a unique serial number with the prefix T e.g. T000001 (bar-coded and numerical) that must be recorded with full serial tracking as per ISO standard. It must be easy to see if a seal or the packaging has undergone tampering.	

Blood Alcohol Kit

HP m

TOXICOLOGY SAMPLE KIT: PACKAGING/BUCKET

1. A 1 to 2 litre plastic bucket, with handle and lid, that can hold the required contents, and meeting the following:
2. Must close tightly and must be simple to seal for tamper proof purposes.
3. The opening must be wide enough to allow easy access to the contents.
4. For convenient storage purposes, a shape that is easily stackable.
5. The packaging/bucket must be labelled with a standard sign (at least 40 mm x 40 mm) for biological hazards that is not affected by wetting or freezing.
6. The lid of the packaging/bucket must be labelled with a unique serial number (bar-code) mirrored by the sample containers inside.

TOXICOLOGY SAMPLE KIT: CONTENT

1. The following content is required for the Toxicology Sampling Kit:
2. One plastic specimen jar of approximately 150 ml capacity, labelled "STOMACH CONTENTS".
3. One plastic specimen jar of approximately 150 ml capacity, with a blank label.
4. One 10 ml bottle, with a cap that seals airtight and that contains an anti-leak mechanism. Included in the bottle must be the correct amount of "Potox" preservative of pharmaceutical quantity, ground to an ultra-fine powder, mixed to the correct ratios. (Information to this regard can be provided, if required).
5. One 20 ml McCartney bottle, with a cap that seals airtight and that contains an anti-leak mechanism. Included in the bottle must be the correct amount of "Potox" preservative of pharmaceutical quantity, ground to an ultra-fine powder, mixed to the correct ratios. (Information to this regard can be provided, if required).
6. One 20 ml McCartney bottle for the collection of blood, with a cap that seals airtight and that contains an anti-leak mechanism. Included in the bottle must be the correct amount of "Potox" preservative of pharmaceutical quantity, ground to an ultra-fine powder, mixed to the correct ratios. (Information to this regard can be provided, if required).
7. One 20 ml McCartney bottle, with a cap that seals airtight and that contains an anti-leak mechanism. Included in the bottle must be the correct amount of "Potox" preservative of pharmaceutical quantity, ground to an ultra-fine powder, mixed to the correct ratios. (Information to this regard can be provided, if required).
8. An instruction leaflet to describe the use of the Toxicology Sampling Kit, detail to be provided by the client and printed by the supplier package/ bucket, and having a unique identifier per container, eg a, b, c etc.
9. Each specimen container (jars and McCartney bottles) should contain a unique serial number (barcode) mirrored by that of the package/ bucket.
10. Each specimen jar should have a self-sealing lid that is leak-free with a watertight fit
11. 3 x Post Mortem Toxicology Referral Form, detail to be provided by the client and printed by the supplier of the kit (see example in Annexure B).
12. A zip seal plastic bag affixed to the lid of the bucket, big enough to accommodate Annexure B when folded in half (to hold Annexure B after completion by Forensic Medical Practitioner).
13. All the content must be sealed into a strong clear plastic bag.
14. The bag mentioned in 28 must be labelled with:
 - TOXICOLOGY SAMPLING KIT;
 - The unique serial number of the kit package;
 - A brief list of the contents;
 - The relevant manufacturing and sterilization date;
 - The batch number;
 - The expiry date;

Sampling Kit

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- Confirmation of quality control performed.	
15. A molded base or molded base insert to secure the filled jars and McCartney bottles during transport.	
16. 2 x Strips of tamper evident tape or better alternative, bearing a unique serial number with the prefix T e.g. T000001 (bar-coded and numerical) that must be recorded with full serial tracking as per ISO standard. It must be easy to see if a seal or the packaging has undergone tampering.	
17. 1 x tamper evident exhibit sealing bag with unique sequential reference number.	
ADDITIONAL REQUIREMENTS	
1. All containers in the Toxicology Sampling Kit must be clean and sterilized to avoid the possibility of contamination.	
2. A clinically clean environment and stringent quality control procedure are critical to ensure clean containers and kits.	
3. The production of the POTOX should happen in a Bio Hazard safety zone, and should be gamma-irradiated after filling, to ensure sterility.	
4. The bidder must include a detailed procedure describing the preparation of the kit for the evaluation by the client.	
5. An example of the complete Toxicology Sampling Kit must be delivered for evaluation with the bid documents. Offers without examples to test, will not be considered.	
6. The expiry date of both the kit and its contents (where applicable) should not be less than 24 months.	
NB: THE SERVICE PROVIDER SHOULD PROVIDE THE SAMPLE AS PART OF THE EVALUATION CRITERIA.	
Bidder's Comment:	
.....	
.....	

1. The bidder is requested to put a company stamp on this specification document.
2. The bidder is requested to complete "Mandatory requirements to be completed by bidder" segment

NB: FAILURE TO COMPLY WITH THE ABOVE WILL RESULT FOR A BIDDER TO BE DISQUALIFIED.

Bidder's Official Stamp



Bidder Signature: _____

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