



KWAZULU-NATAL PROVINCE

**HEALTH
REPUBLIC OF SOUTH AFRICA**

Quotation Advert

Opening Date: 24/06/2024
Closing Date: 28/06/2024

Closing Time: 11:00

INSTITUTION DETAILS

Institution Name: Eshowe Hospital
Province: KwaZulu-Natal
Department of entity: Department of Health
Division or section: Central Supply Chain Management
Place where goods/ **Eshowe district Hospital**
Date Submitted: 21/06/2024

ITEM CATEGORY AND DETAILS

Quotation number: ESW/67/25
Item Category: Goods
Item Description: Maternity case peri-operative record
Quantity (if supplies): Per quote

COMPULSORY BRIEFING SESSION / SITE VISIT

Select Type: Not applicable
Date: N/a
Time: N/a
Venue: N/a

QUOTES CAN BE COLLECTED FROM: Eshowe hospital and KZN health Website

QUOTES SHOULD BE DELIVERED TO: Eshowe Hospital tender box next to security pedestrian gate

ENQUIRIES REGARDING ADVERT MAY BE DIRECTED TO:

Name: Mr B Zulu/ Qwabe N
Email: Nonhlanhla.qwabe@kznhealth.gov.za
Contact number: 035 473 4597/4664

Finance Manager: Ms Z.N.B/ Khanyile

Finance Manager Signature

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 institution?² **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 cancelling 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.

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.**

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

(i) 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 will not take place.

(ii) Date: _____ / _____ / _____ Time: _____ : _____ Place: _____

Institution Stamp:	Institution Site Inspection / briefing session Official:
	Full Name: _____
	Signature: _____
	Date: _____

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
- (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

12.1. 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. **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.

4. 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:

$$\begin{array}{ccc}
 \text{80/20} & & \text{90/10} \\
 P_s = 80 \left(1 - \frac{P_t - P_{min}}{P_{min}} \right) & \text{OR} & P_s = 90 \left(1 - \frac{P_t - P_{min}}{P_{min}} \right)
 \end{array}$$

Where

- P_s = Points scored for price of tender under consideration
- P_t = Price of tender under consideration
- P_{min} = 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:

$$\begin{array}{ccc}
 \text{80/20} & & \text{90/10} \\
 P_s = 80 \left(1 + \frac{P_t - P_{max}}{P_{max}} \right) & \text{OR} & P_s = 90 \left(1 + \frac{P_t - P_{max}}{P_{max}} \right)
 \end{array}$$

Where

- P_s = Points scored for price of tender under consideration
- P_t = Price of tender under consideration
- P_{max} = 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)
RDP Goal: Full points allocated to promote enterprises manufacturing or producing in the Province of Kwa-Zulu Natal	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: _____



health

Department: Health
REPUBLIC OF SOUTH AFRICA

Maternity Care Peri-operative record

This record must be completed for all person's requiring surgery during pregnancy or the puerperium. Once completed, it must be placed within the Maternity Case Record to be filed at the hospital where the delivery took place. Procedures done at a facility where delivery did not occur must be filed in the patient records. Use a new record for every operation.

Name and ID number of patient or place large patient sticker here

Name of medical practitioner booking the procedure

- Procedure: Caesarean section Tubal ligation Laparotomy Emergency hysterectomy
- Other _____

URGENCY OF PROCEDURE (select only 1)

- RED: Immediate delivery (life threatening to mother and/or fetus)
- YELLOW: Urgent delivery (Maternal/fetal compromise not immediate life threatening)
- GREEN: Scheduled urgent delivery (need early delivery but no maternal/fetal compromise)
- ELECTIVE Scheduled at a time to suit mother/staff

Best describe the reason/indication for the caesarean section/ procedure:

Booking arrangements

Discussed case with senior colleague/consultant (name and time):

Discussed with anaesthetic doctor (name and time):

Discussed with neonatal staff (name and time):

Date and time procedure scheduled:

URGENCY OF CAESAREAN DELIVERY (examples)

	RED Emergency- immediate threat to life of person or her fetus	YELLOW Maternal or fetal compromise which is not immediately life threatening	GREEN Needing early delivery, but no maternal or fetal compromise)
Target time (decision to incision)	Ideally within 30 minutes	Ideally within 60 minutes	Ideally within 3 hours
Fetal condition (examples)	Fetal distress (pathological CTG)	Suspicious CTG	Fetal anomaly or compromise that need daytime delivery for paediatric management (arrange necessary skilled team as needed)
	Cord prolapse	Cord presentation; patient in labour	
	Footling breech- with ruptured membranes	Footling breech, membranes still intact, patient in labour	
Clinical presentation (examples)	Abruptio placentae; baby alive and viable	Poor progress in labour	Eclampsia, failed induction of labour or vaginal delivery not possible
	Placenta praevia- massive bleeding	Unsuccessful attempt at VBAC	Failed induction of labour: urgent indication for delivery
	Uterine rupture/dehiscence	Cephalo-pelvic disproportion	2 or more previous CS/previous classical CS in early labour
	Transverse lie, in labour	Prolonged second stage	One previous CS, patient not for VBAC, in early labour
	Abandoned instrumental delivery	Twin pregnancy; delivery of second twin	Any GREEN indication presenting in active labour
Maternal condition	Severe maternal disease		

IMPORTANT INFORMATION FOR ANAESTHETIC TEAM:

Haemoglobin:	NPO since:	Latest platelet count if pre-eclampsia:
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Maternal medical condition (select all that is applicable)

- Healthy
- Severe pre-operative blood loss (antepartum haemorrhage)
- Abruptio placentae
- Placenta praevia
- Morbidly adherent placenta
- Pre-eclampsia
- Decreased level of consciousness
- Acute severe hypertension
- Maternal diabetes
- BMI 40-50
- BMI >50
- Cardiac disease
- Active respiratory disease
- Currently on MgSO₄
- Currently on anti-coagulative drugs
- Allergies: _____
- Medical history _____
- Surgical history _____
- Other _____

WARD PREPARATION FOR THEATRE AND TRANSFER

Planned procedure

Procedure date/time Pickup date/time

Known allergies

	WARD			Theatre		
	Yes	No	N/A	Yes	No	N/A
Informed consent signed						
Medical alert band/ chain in situ						
Make-up/varnish removed						
Artificial nails removed						
Jewelry removed						
Dentures removed						
Contact lenses removed						
Patient is nil per mouth since ___h__						
Dressed in theatre garment						
Urine catheter in-situ						
List pre-medication drugs:						
Premed administered by						
Signature						
Patient prepared by						
Signature						
Date /time	Left ward			Arrive OT		
Received in theatre by						
Signature						
Vital signs on arrival OT		Documents received OT				
Blood pressure			Maternity case record book			
Pulse			Prescription chart			
Respiration rate			Laboratory results			
Urine disptix			X-Rays			
Catheter						
Fetal heart						

CONSENT TO MEDICAL OR SURGICAL PROCEDURE

I, Dr _____ have explained the nature, risks & possible consequences of the medical /surgical procedure to the undersigned patient or her legal guardian.

Signature _____

Date _____

Circle whichever is applicable

Procedure explained:	Personally	Via Interpreter
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NATURE OF PROCEDURE:

Where applicable indicate side of procedure (Right or Left)

Circle whichever is applicable

Type of anaesthetic:	Local	Spinal	General	Procedural Sedation
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CONSENT TO USE OF BLOOD and/or blood products if necessary during the course of the procedure

Consent granted by Patient/Guardian :

Consent withheld by
Patient/Guardian:

Signature

Signature

I consent to a sample of my blood being taken and tested for Hepatitis B and the Human Immunodeficiency Virus (HIV) should contamination of a health care worker by my bodily fluids occur during the procedure.

Patient's / Guardian's Signature _____

Full Name of Patient		I, the undersigned, hereby consent to the performance of, and understand the nature, risks and possible outcomes of the above procedure. The doctors who perform the above may carry out additional or alternative measures (including general anaesthesia) if considered necessary. In the case of a sterilisation procedure, I understand that pregnancy may occur in exceptional cases, in which case I shall not hold the Department of Health and/or its personnel responsible. I also accept that alternative methods of birth control are still available to me.
Signature/Thumb	Date	
Print of patient		

COMPLETE THIS SECTION IF CONSENT IS GIVEN BY A PERSON ON BEHALF OF THE PATIENT

Print Name _____

Signature _____

Date _____

Relationship to patient _____

Means by which consent was given:

Personally

Telephonically

NAMES AND SIGNATURES OF WITNESSES TO THE PATIENT'S / GUARDIAN'S SIGNATURE ON THIS DOCUMENT

Witness 1

Witness 2

Print Name _____

Print Name _____

Signature _____

Signature _____

CONSENT TO CAESAREAN DELIVERY

NATURE OF PROCEDURE: CAESAREAN SECTION*

Contact details (if patient wishes to discuss options later)

I have introduced myself by name and explained the nature, risks and possible consequences of a caesarean delivery to the undersigned patient or person legally competent to give consent. In particular, I have explained the following:	Print name	NAME OF DOCTOR (To be filled in by a registered health professional with appropriate knowledge of the proposed procedure)
	Signature	

Intended benefit:

Delivery of her baby (or babies) through a cut in the tummy and the uterus (womb) in a situation where the risks of the baby being born through the vagina is more than the risk of the delivery by Caesarean section.

Frequent risks:

Bleeding during or after the operation, infection in the wound or in the womb (sepsis), persistent pain and discomfort over the scar, risk of repeat caesarean delivery in following pregnancies, re-admission to hospital, minor cuts to the baby during delivery.

Serious risks (uncommon):

Emergency requiring removal of the womb (hysterectomy), increased risk of a tear in the womb in future pregnancies, development of a blood clot in the legs or lungs, injury to the bladder or bowel.

I have also discussed what the procedure is likely to involve, the benefits and risks of any available alternative treatments (including no treatment) and any particular concerns of this patient.

THE PROCEDURE WILL INVOLVE (one or more):	General anaesthesia <input type="checkbox"/>	Regional anaesthesia (epidural or spinal) <input type="checkbox"/>	Local anaesthesia <input type="checkbox"/>
---	--	--	--

CONSENT TO USE OF BLOOD and/or BLOOD PRODUCTS	I have counselled the patient on the use and dangers of blood products and the undersigned patient hereby Grants or Withholds consent for the use of blood and/or blood products should it become necessary during the procedure. TICK the appropriate box	
<table style="width: 100%;"> <tr> <td style="width: 50%; border: 1px solid black; padding: 5px;"> I grant consent <input type="checkbox"/> </td> <td style="width: 50%; border: 1px solid black; padding: 5px;"> I withhold consent <input type="checkbox"/> Signature _____ </td> </tr> </table>		I grant consent <input type="checkbox"/>
I grant consent <input type="checkbox"/>	I withhold consent <input type="checkbox"/> Signature _____	

I, the undersigned patient hereby agree that a sample of my blood can be taken and tested and tested for Hepatitis B and Human Immunodeficiency Virus (HIV) should an incident of contamination of a health care worker by bodily fluids occur during the procedure. TICK whichever is applicable.

<input type="checkbox"/> I agree	<input type="checkbox"/> I do not agree
----------------------------------	---

FULL NAME OF PATIENT _____	I, the undersigned, hereby consent to the performance of, and understand the nature, risks and possible consequences of the above procedure. The doctors who perform the procedure may increase the reasonable scope thereof or carry out additional or alternative measures (including general anaesthesia) if considered necessary.	
<table style="width: 100%;"> <tr> <td style="width: 50%; vertical-align: top;"> SIGNATURE or THUMB PRINT OF PATIENT </td> <td style="width: 50%; vertical-align: top;"> Date </td> </tr> </table>		SIGNATURE or THUMB PRINT OF PATIENT
SIGNATURE or THUMB PRINT OF PATIENT	Date	

PERSON LEGALLY COMPETENT TO GIVE CONSENT	Print name			This section to be filled in if a person other than the patient gives consent.
	Signature		Date	
	Capacity or relationship to patient			
	Means by which consent was given	Personally <input type="checkbox"/>	Telephonically <input type="checkbox"/>	

WITNESS 1	Print name _____ Signature _____	Names and signatures of witness to the signing of this document by the patient or a person legally competent to give consent on behalf of the patient.
WITNESS 2	Print name _____ Signature _____	

*A separate consent form should be used for sterilisation procedures.

*A separate consent form should be used if any additional procedures are planned during the time of the Caesarean section (e.g. hysterectomy).

OBSTETRIC ANAESTHETIC RECORD

Proposed Operation: _____		Details of Anaesthetist	
Surgeon: _____	Grade: _____	Name and HPCSA nr and highest qualification	
Date: _____	Consent obtained: _____	Grade:	Intern <input type="checkbox"/>
Nil by mouth since (Time): _____		Comm. Service MO <input type="checkbox"/>	
What was eaten/drank? _____		GP/MO < 2 years <input type="checkbox"/>	
History: _____		GP/MO ≥ 2 years <input type="checkbox"/>	
Previous Anaesthetic History: _____		Registrar <input type="checkbox"/>	
		Specialist <input type="checkbox"/>	

Medication: _____ Allergies: _____

General Examination: Height (m) Mass (kg) BP Pulse

Heart: _____

Chest: _____

Airway Examination: Mallampati Score:

Jaw mobility Loose/awkward teeth: Yes No Pharynx: _____ Neck: _____

ASA rating: 1 2 3 4 5 E

Investigations: Hb Platelets Urea & Electrolytes: _____

Chest X-Ray: Normal Abnormal

Details: _____

Urine: _____

Other: _____

Premedication:	To be given at:	Ordered by	Given at:	By
0.3 Molar sodium citrate 30 mL per os	<input type="checkbox"/>			
Metoclopramide 10 mg iv	<input type="checkbox"/>			
Ranitidine 150 mg per os	<input type="checkbox"/>			
Other:				

Pre-anaesthesia check: Freely running iv Suction Machine check

Technique: Spinal Epidural CSE General Sedation Standby

<p>Regional anaesthesia: Spinal interspace: <input style="width: 40px;" type="text"/></p> <p>Number of attempts: <input style="width: 40px;" type="text"/></p> <p>Position of patient: Lateral <input type="checkbox"/> Sitting <input type="checkbox"/></p> <p>Spinal needle: Type: Atraumatic <input type="checkbox"/></p> <p>Size (gauge): <input style="width: 40px;" type="text"/></p> <p>Epidural needle: Type: Tuohy <input type="checkbox"/> Other <input type="checkbox"/></p> <p>Size (gauge): <input style="width: 40px;" type="text"/></p> <p>Epidural space location: _____</p> <p>Loss of resistance: To air <input type="checkbox"/> To saline <input type="checkbox"/></p> <p>Other (describe): _____</p> <p>Epidural catheter: Size (gauge): <input style="width: 40px;" type="text"/></p> <p>Length within epidural space (cm): <input style="width: 40px;" type="text"/></p> <p>Sensory height (to cold) of block pre-incision: <input style="width: 40px;" type="text"/></p>	<p>General anaesthesia: Induction sequence: Preoxygenation <input type="checkbox"/></p> <p>Cricoid pressure <input type="checkbox"/></p> <p>Laryngoscopy and rapid tracheal intubation with a cuffed tube <input type="checkbox"/></p> <p>Check stomach <input type="checkbox"/> Size of tracheal tube (mm): <input style="width: 40px;" type="text"/></p> <p>Air Entry: L <input type="checkbox"/> R <input type="checkbox"/> Length inserted (cm): <input style="width: 40px;" type="text"/></p> <p>Alternative airway management: Face mask <input type="checkbox"/></p> <p>Laryngeal mask <input type="checkbox"/></p> <p>Awake intubation** <input type="checkbox"/></p> <p>Surgical airway <input type="checkbox"/></p> <p>Combitube Other <input type="checkbox"/> (specify) <input style="width: 40px;" type="text"/></p> <p>Ventilation: Spontaneous <input type="checkbox"/></p> <p>Controlled <input type="checkbox"/></p> <p>Circuit: _____</p> <p>Ventilator: _____</p> <p>FiO₂: _____</p> <p>O₂/Air <input style="width: 40px;" type="text"/></p> <p>O₂/Nitrous Oxide <input style="width: 40px;" type="text"/></p> <p>** Details: _____</p>
---	--

Remarks and Complications

OBSTETRIC ANAESTHETIC RECORD

Time:				Times:
Drugs:				Induction (I)
Prophylactic antibiotics:				Uterine incision (U)
Oxytocin				Cord clamp (D)
Other uterotonics				I-D (min.)
				U-D (sec.)
Agent % (Inspired)				
IV Fluids:				Totals:
Blood Loss:				
Urine output:				

Monitoring every 5 mins:																																									
ECG	<input type="checkbox"/>	210																																							
Oximetry	<input type="checkbox"/>	200																																							
Capnograph	<input type="checkbox"/>	190																																							
NIBP	<input type="checkbox"/>	180																																							
CVP	<input type="checkbox"/>	170																																							
Arterial line	<input type="checkbox"/>	160																																							
N-M block	<input type="checkbox"/>	150																																							
Urine	<input type="checkbox"/>	140																																							
Temp	<input type="checkbox"/>	130																																							
PIP _(AW)	<input type="checkbox"/>	120																																							
O ₂ Analyser	<input type="checkbox"/>	110																																							
	<input type="checkbox"/>	100																																							
Position:		90																																							
Supine tilt	<input type="checkbox"/>	80																																							
Wedge	<input type="checkbox"/>	70																																							
Other	<input type="checkbox"/>	60																																							
Lithotomy	<input type="checkbox"/>	50																																							
Trendel.	<input type="checkbox"/>	40																																							
SBP	∇	Sat %																																							
DBP	∧	ET _{CO2}																																							
HR	•	CVP																																							
CVP	x	T °C																																							
		FiO ₂																																							

Recovery Room Record

Time	BP	Pulse	Respiratory pattern and rate	Output			Drugs & iv therapy	State of consciousness	Signature
				Urine	Vomitus	Wound			

- 1 = Complete block (unable to move feet or knees)
- 2 = Almost complete block (able to move feet only)
- 3 = Partial block (just able to move knees)
- 4 = Detectable weakness of hip flexion (between scores 3 and 5)
- 5 = No detectable weakness of hip flexion while supine (full flexion of knees)
- 6 = Able to perform partial knee bend

Complications in recovery room	
Transfer from recovery room authorised by _____	Time _____
Transferred to ward _____	Time _____
Received by _____	Time _____

OPERATION		INTRA-OPERATIVE RECORD						
		NB: Complete or mark in space given				THEATRE NR:		
Operation Time:		From:		To:		Duration:		
Type of Anaesthesia:				Anaesthetist:				
Surgeon:		Assistant:						
SECTION: B - SURGEON COMPLETES THIS SECTION								
Nature of Operation:								
Surgeon:		Name in Print:		Signature:		Qualification:		
Procedure code:								
INTRAOPERATIVE PATIENT CARE		SECTION C: PROFESSIONAL NURSE COMPLETES THIS SECTION						
		PATIENT POSITION:		(MARK x)	Supine		Prone	
Left Lateral			Right Lateral		Trendellen- burg		Other	
BONY PROMINENCES		Checked:	YES:	NO:	Padded:	YES:	NO:	
WARMING BLANKET		YES:	NO:					
ANY ABNORMALITIES OBSERVED (Describe shortly)								
DIATHERMY:		Diathermy used		YES:	NO:	Checked	YES:	NO:
Plate site:		ARM:	LEG:	OTHER:	LEFT:		RIGHT:	
WOUND CLASSIFICATION:					CLEAN:			
INFECTED:		CONTAMINATED:			CLEAN CONTAMINATED:			
SKIN PREPARATION					Chlorhexidine in Water			
Chlorhexidine in Alcohol		Povidone-iodine			Other:			
INFILTRATION		YES:	NO	Type:				
X-RAYS USED:		YES:	NO	C-Arm used	YES:	NO		
					Contrast used	YES:	NO	
SWAB/INSTRUMENT/SHARP CONTROL								
We, the undersigned, hereby declare that the instruments, needles and swabs in respect of the above-mentioned operation were counted before, during and after the operation and that the totals were found correct.								
		COMPLETE		TOTAL:	N.A.	PLUGS:	YES:	NO:
		YES	NO					
Abdominal						Type:		
Raytec						Size:		
Dissecting						Tapes/Other	YES:	NO:
Other						Type:		
						Clips	YES:	NO:
						SKIN SUTURE		
CATHETERS/ DRAINS				YES:	NO:	SIZE:		
Urine								
Nasal tube								
Thoracic drain								
Pencil drain								
Other								

INTRA-OPERATIVE RECORD CONTINUED						
NB: Mark applicable given spaces						
Unplanned events	UNUSUAL INCIDENT REPORT WRITTEN?		YES:		NO:	
	Intraoperative bleeding					
	Source of bleeding		Blood Loss			
INTRA-OPERATIVE	ROUTE CHART COMPLETED:		YES		NO	
	SPECIMEN OBTAINED	YES:	NO:		NUMBER:	
	TYPE:					
	OPERATING TEAM MEMBERS:		NAME IN PRINT		SIGNATURE	
	REGISTERED SCRUB NURSE:					
	SUPERVISOR: (If theatre student/ new PN)					
	CO-CHECKER/ CIRCULATING NURSE:					
ANAESTHETIC NURSE:						
POST-OPERATIVE	POST OPERATIVE CHECKLIST					
	Post-operative skin/pressure areas check:		Intact		Skin Lesion:	
	Short description of skin lesion:					
	PATIENT TRANSFERRED TO: (Date/Time)					
	RECOVERY ROOM					
	Professional authorising release of patient from theatre					
	Date/Time	Name	Signature			
	Professional receiving patient from Theatre					
		Date/Time	Name	Signature		
WARD:						
CRITICAL CARE:						
HIGH CARE:						

COUNSELLING CHECKLIST PRIOR TO POST PARTUM TUBAL LIGATION

For persons capable of signing their own consent

I have discussed the following with this person:

- Her reason for choosing sterilization.
- Alternative long acting effective contraceptive methods.
- Sterilisation is a permanent and irreversible method of contraception.
- Stability of relationship and possibility of regret due to change in circumstances, such as possible loss of child/children/partner or remarriage.
- Consider option of male or female sterilization. (Male procedure is smaller, safer and more effective).
- The sterilization procedure. Local or general anaesthetic, surgical approach, type of tubal closure.
- Risk of anaesthesia/surgery and possibility of additional surgery if complications occur.
- The risk of failure: 1 in 200 lifetime risk of pregnancy in a female
- If pregnancy occurs after sterilisation, there is a slight risk of ectopic pregnancy and the symptoms to report are lower abdominal pain, missed period and irregular bleeding.
- The menstrual cycle will revert to what it was before pregnancy.
- No effect on long term health.
- Sterilisation does not protect against STI/HIV transmission.
- I have answered the person's questions and given a pamphlet

Date _____ Counselling by _____

I, (patient name)

with ID/Passport/other number.....

Hereby states that I have requested a sterilisation (permanent family planning).
This was my own choice and I was not forced to make this decision.
I understand that I will not be able to have any pregnancies in the future and that the operation is permanent.

Signed (patient).....

Witness 1.....

Witness 2

CAESAREAN DELIVERY SAFETY CHECKLIST

SIGN IN (To be said out loud before induction of anaesthesia)	TIME OUT (To be said out loud before skin incision)	SIGN OUT (To be said out loud before patient leaves the operation room)
<p>Patient has confirmed</p> <p><input type="checkbox"/> Identity</p> <p><input type="checkbox"/> Procedure</p> <p><input type="checkbox"/> Consent</p> <p><input type="checkbox"/> Anaesthesia safety check completed (Equipment and medication)</p> <p><input type="checkbox"/> Neonatal safety check completed (Equipment and medication)</p> <p><input type="checkbox"/> Pulse oximeter on patient and functioning</p> <p>Is a difficult airway anticipated?</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes and equipment and assistance is available</p> <p>Does patient have a known allergy</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes</p> <p>Assess bleeding risk (Pre op Hbg/dl)</p> <p>Risk factors for PPH. <input type="checkbox"/> No <input type="checkbox"/> Yes (i.e. prolonged labour, multiple pregnancy, big baby, polyhydramnios, grand multiparity, clotting dysfunction, PPH in the past). If yes,</p> <p><input type="checkbox"/> There is adequate IV access?</p> <p>Is emergency blood available? <input type="checkbox"/> No <input type="checkbox"/> Yes</p> <p>Are there any concerns about the placental site</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes</p>	<p><input type="checkbox"/> Confirm all team members have introduced themselves by name and role</p> <p>To Surgeon</p> <p>Are there any potential problems the team should be aware of?</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes</p> <p><input type="checkbox"/> Mothers rhesus status known</p> <p>Does cord blood need to be taken?</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes</p> <p>To Anaesthetist:</p> <p><input type="checkbox"/> Wedge placed?</p> <p><input type="checkbox"/> Any patient specific concerns?</p> <p>To Scrub Sister</p> <p><input type="checkbox"/> Sterility of instruments confirmed</p> <p><input type="checkbox"/> Any equipment issues / concerns</p> <p><input type="checkbox"/> Diathermy and suction functional</p>	<p>Practitioner verbally confirms with the team:</p> <p><input type="checkbox"/> Name of the procedure and any additional procedure has been recorded?</p> <p><input type="checkbox"/> Instruments, swabs and sharp counts are correct?</p> <p><input type="checkbox"/> Specimens have been labelled?</p> <p><input type="checkbox"/> Blood loss has been recorded?</p> <p>Obstetrician, Anaesthetist and Scrub Nurse have discussed:</p> <p><input type="checkbox"/> Concerns for recovery and further management?</p> <p><input type="checkbox"/> Need for post-operative VTE prophylaxis?</p> <p><input type="checkbox"/> Need for postoperative antibiotics?</p> <p><input type="checkbox"/> Equipment problems that have been identified?</p> <p><input type="checkbox"/> Oxytocin 20 IU in 1000mls IV ready to be administered?</p>
<p><input type="checkbox"/> Antibiotic prophylaxis given in the last hour?</p> <p><input type="checkbox"/> Appropriate / recent antacid prophylaxis given?</p> <p><input type="checkbox"/> Urinary catheter is draining</p> <p>Are any additional procedures planned?</p> <p><input type="checkbox"/> IUCD</p> <p><input type="checkbox"/> BTL</p> <p><input type="checkbox"/> N/A</p> <p>Is the foetal heart present?</p> <p><input type="checkbox"/> No <input type="checkbox"/> Yes</p>	<p>Patient Name: _____</p> <p>Patient Surname: _____</p> <p>Date of Birth: _____</p> <p>Hospital number: _____</p> <p>Date of Surgery: _____</p>	<p>Midwife has confirmed that</p> <p><input type="checkbox"/> Baby/ies been correctly labelled?</p> <p><input type="checkbox"/> Relevant cord bloods have been taken?</p>
NAME AND SIGNATURE OF HEALTHCARE WORKER	NAME AND SIGNATURE OF HEALTHCARE WORKER	NAME AND SIGNATURE OF HEALTHCARE WORKER



ESHOWE HOSPITAL
SPEC NO.: ESHW 97- 2024/25

ESW NO.:.....

Quote Number: _____

Item Description: **MATERNITY CASE PERI-OPERATIVE RECORD BOOK**

Department/Section: **SCM - LOGISTICS**

Purpose of Item: **INVENTORY**

key note:

- The Bidder is required to make comments on section A and B about service(s) and item(s) the Bidder is prepared to offer:
- All pages must be fully signed
- Failure to sign and comment may disqualify the Bidder
- Fully completed form must be returned back with the Quotation Documents and other supporting documents

SECTION A: SPECIFICATION

1. ITEM(S) ON THIS BID

- **MATERNITY CASE PERI-OPERATIVE RECORD BOOK**
 - MCR 2018 Surgery
 - Front cover: labelled/printed to show contents and item number.
 - All Printing on file must be in the English language only.
 - NB: Hard copy sample on request.

NB: The Bidder is required to also specify the expiry date of the item(s) willing to offer with comments below
BIDDER'S COMMENTS:

NB: The Bidder is required to also specify the expiry date of the item(s) willing to offer with comments below
BIDDER'S COMMENTS:

2. All items offered should be according to Health Standards/ WHO Standards/ SABS Standards / ISO/CIDB Standards and approved.

NB: Proof must be attached
BIDDER'S COMMENTS:

3. The successful Supplier will be required to deliver the item on this bid within three weeks from order date
BIDDER'S COMMENTS:

4. The Sample is required and must be made available when requested in writing for all items on this Bid for Evaluation

BIDDER'S COMMENTS:

5. The guarantee period should be a minimum of 12 months from the delivery date
BIDDER'S COMMENTS:

BIDDER'S AUTHORIZED REPRESENTATIVE SIGNATURE:

SECTION B: CONDITIONS OF THIS BID

THE BIDDER IS REQUIRED TO COMMENT BY TICKING ONLY ONE BLOCK PER ROW (CLAUSE) BELOW

Clause No.	Description	Bidder's Comment: Offer will Comply?	
		TICK ONE(v)	
		YES	NO
CA1	The Supplier must supply product that is new, no defect, product of the most recent or current models, and that incorporate all recent improvements in design and materials.		
CA2	Bidders must submit their offers in line with the bid specifications. Offers exceeding specification are also deem to be in line with the specification. NB: Failure to comply shall invalidate the bid		
CA3	The Supplier must honour the proposed delivery date and any delays must be approved by Eshowe District Hospital. The Eshowe District Hospital may then extend the delivery date, if and as it deems fit. Should the Contractor fail to supply the offer within the time stated on the bid, or within the extended time allowed to him/her, the Eshowe District Hospital reserves the right to cancel the order		
CA4	Product(s) shall be delivered on Weekdays between 07:30 and 16:00. No goods or service will be received on Saturdays, Sundays and public holidays.		
CA5	The Supplier must comply with the General Conditions of Contract by Treasury Department, Environmental Management Act no.107 of 1998, Occupational Health and Safety Act no.83 of 1995, Food, Cosmetics and Disinfectants Act no.54 of 1972, Hazardous Substances Act no.15 of 1973 and other relevant regulations		
CA6	The Bidder must specify the product code or catalogue number and Brand name of the item willing to offer		
CA7	The Bidder must specify the expiry date of the item(s) willing to offer		
CA8	-All applicable Items on this bid (quote) must be SABS and ISO Approved. -Proof Must Be attached NB: Failure to comply shall invalidate the bid		
CA9	The Bidder must specify the country of Manufacture on Official Price Page		
CA10	The successful Bidder should at no extra cost provide additional ongoing advise, updates or training to End-user on item(s) offered.		
Clause No.	Description	Bidder's Comment: Offer will Comply?	
		TICK ONE(v)	
		YES	NO
CA11	Where the sample is required, it must be submitted at the Bidders' expense and risk for the purpose of visual screening of products during the evaluation phase. Hence, all samples submitted for visual screening must be a true representation of the product which will be supplied. NB: Sample should be marked with : Company name and contacts, ZNQ number, Brand (Manufacture's) Name, Product Code		

BIDDER'S AUTHORIZED REPRESENTATIVE SIGNATURE:

CA12	<p>The Sample is required for all items on this Bid for Evaluation</p> <p>NB: Failure to comply shall invalidate the bid</p>		
CA13	<p>Packaging: The following information must be clearly and indelibly printed on all inner and outer packaging:-</p> <ul style="list-style-type: none"> ➤ The product name, product code as relevant, quantity of contents (e.g box of 100units), expiry date (if applicable). ➤ Size of the product, date of manufacture and trademark or trade name of the manufacture ➤ All products must be packed in acceptable containers/packets/box/bale, where applicable, specifically developed for the product ➤ The word "sterile" (where applicable) ➤ The warning information, instructions and conditions under which the product must be stored ➤ Labels for consumable items must be clearly marked for occupational health and safety purposes 		

6. Penalties to be noted by the suppliers:

- a. 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.

7. The evaluation criteria / special terms and conditions

List evaluation criteria / special terms and conditions to be advertised (if applicable)		
1.	Pre-qualification criteria	Does the offer meet the pre-qualification criteria?
2.	Administrative	Does the offer comply with stipulated administrative requirements?
3.	Conformance:	Was the product made or service performed to specifications?
4.	Performance:	Will/does the product/service fulfil its performance obligation, in a manner that releases the supplier from all liabilities under the contract?
5.	Features:	What characteristics does the product or service have?
6.	Reliability:	How long can a product go between failures and the need for maintenance? (guarantee)
7.	Durability:	What is the useful life for the product? How will the product hold up under extended use?
8.	Serviceability:	How easy is it to repair, maintain or support the product or service? (customer support)
9.	Ability & Capacity	The ability and capacity of the vendor to execute the contract
10.	Preference points	Preferential Procurement System (20 points for specific goals)

BIDDER'S AUTHORIZED REPRESENTATIVE SIGNATURE: