



**KWAZULU-NATAL PROVINCE**  
HEALTH  
REPUBLIC OF SOUTH AFRICA

BID DOCUMENT NUMBER: **ZNB 6685/2/2021-H:**

DESCRIPTION: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.

Name of Bidder.....

Central Supplier's Database Registration Number.....

Income Tax Reference Number.....

**BIDDER TO NOTE THE FOLLOWING**

**CLOSING DATE AND TIME:**

Date: 04 MAY 2021

Time: 11: 00AM

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# SECTION A: INVITATION TO BID

## PART A

<b>YOU ARE HEREBY INVITED TO BID FOR REQUIREMENTS OF THE KWAZULU-NATAL DEPARTMENT OF HEALTH</b>					
BID NUMBER:	ZNB 6685/2/2021-H	CLOSING DATE:	04 MAY 2021	CLOSING TIME:	11: H 00 AM
DESCRIPTION	THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.				
<b>THE SUCCESSFUL BIDDER WILL BE REQUIRED TO FILL IN AND SIGN A WRITTEN CONTRACT FORM (SBD7).</b>					
BID RESPONSE DOCUMENTS MAY BE DEPOSITED IN THE BID BOX SITUATED AT (STREET ADDRESS)					
CENTRAL SUPPLY CHAIN MANAGEMENT DIRECTORATE					
OLD BOYS SCHOOL, 310 JABU NDLOVU STREET					
PIETERMARITZBURG					
3201					
<b>SUPPLIER INFORMATION</b>					
NAME OF BIDDER					
POSTAL ADDRESS					
STREET ADDRESS					
TELEPHONE NUMBER	CODE		NUMBER		
CELLPHONE NUMBER					
FACSIMILE NUMBER	CODE		NUMBER		
E-MAIL ADDRESS					
VATREGISTRATION NUMBER					
	TCS PIN:		OR	CSD No:	
STATUS LEVEL VERIFICATION CERTIFICATE [TICK APPLICABLE BOX]	<input type="checkbox"/> Yes <input type="checkbox"/> No		STATUS LEVEL SWORN AFFIDAVIT		<input type="checkbox"/> Yes <input type="checkbox"/> No
IF YES, WHO WAS THE CERTIFICATE ISSUED BY?					
AN ACCOUNTING OFFICER AS CONTEMPLATED IN THE CLOSE CORPORATION ACT (CCA)	<input type="checkbox"/>	AN ACCOUNTING OFFICER AS CONTEMPLATED IN THE CLOSE CORPORATION ACT (CCA)			
CCA) AND NAME THE APPLICABLE IN THE TICK BOX	<input type="checkbox"/>	A VERIFICATION AGENCY ACCREDITED BY THE SOUTH AFRICAN ACCREDITATION SYSTEM (SANAS)			
	<input type="checkbox"/>	A REGISTERED AUDITOR			
		NAME:			
<b>[A STATUS LEVEL VERIFICATION CERTIFICATE/SWORN AFFIDAVIT (FOR EMEs&amp; QSEs) MUST BE SUBMITTED IN ORDER TO QUALIFY FOR PREFERENCE POINTS FOR ]</b>					
ARE YOU THE ACCREDITED REPRESENTATIVE IN SOUTH AFRICA FOR THE GOODS /SERVICES /WORKS OFFERED?	<input type="checkbox"/> Yes <input type="checkbox"/> No  [IF YES ENCLOSE PROOF]		ARE YOU A FOREIGN BASED SUPPLIER FOR THE GOODS / SERVICES / WORKS OFFERED?		<input type="checkbox"/> Yes <input type="checkbox"/> No  [IF YES ANSWER PART B:3 BELOW]
SIGNATURE OF BIDDER	.....		DATE		
CAPACITY UNDER WHICH THIS BID IS SIGNED (Attach proof of authority to sign this bid; e.g. resolution of directors, etc.)					
TOTAL NUMBER OF ITEMS OFFERED			TOTAL BID PRICE (ALL INCLUSIVE)		

BIDDING PROCEDURE ENQUIRIES MAY BE DIRECTED TO:		TECHNICAL INFORMATION MAY BE DIRECTED TO:	
DEPARTMENT	KZN Department of Health	DEPARTMENT	KZN Department of Health
CONTACT PERSON	Tenders@kznhealth.gov.za	CONTACT PERSON	Dr R Groenewald
TELEPHONE NUMBER	033 815 8361	TELEPHONE NUMBER	033 395 4200
FACSIMILE NUMBER		FACSIMILE NUMBER	
E-MAIL ADDRESS	Tenders@kznhealth.gov.za	E-MAIL ADDRESS	Edendale.Anaesthetics@kznhealth.gov.za

## PART B: TERMS AND CONDITIONS FOR BIDDING

<b>1. BID SUBMISSION:</b>	
<p>1.1. BIDS MUST BE DELIVERED BY THE STIPULATED TIME TO THE CORRECT ADDRESS. LATE BIDS WILL NOT BE ACCEPTED FOR CONSIDERATION.</p> <p>1.2. ALL BIDS MUST BE SUBMITTED ON THE OFFICIAL FORMS PROVIDED – (NOT TO BE RE-TYPED) OR ONLINE</p> <p>1.3. BIDDERS MUST REGISTER ON THE CENTRAL SUPPLIER DATABASE (CSD) TO UPLOAD MANDATORY INFORMATION NAMELY: (BUSINESS REGISTRATION/ DIRECTORSHIP/ MEMBERSHIP/IDENTITY NUMBERS; TAX COMPLIANCE STATUS; AND BANKING INFORMATION FOR VERIFICATION PURPOSES). CERTIFICATE OR SWORN AFFIDAVIT FOR MUST BE SUBMITTED TO BIDDING INSTITUTION.</p> <p>1.4. WHERE A BIDDER IS NOT REGISTERED ON THE CSD, MANDATORY INFORMATION NAMELY: (BUSINESS REGISTRATION/ DIRECTORSHIP/ MEMBERSHIP/IDENTITY NUMBERS; TAX COMPLIANCE STATUS MAY NOT BE SUBMITTED WITH THE BID DOCUMENTATION. CERTIFICATE OR SWORN AFFIDAVIT FOR MUST BE SUBMITTED TO BIDDING INSTITUTION.</p> <p>1.5. THIS BID IS SUBJECT TO THE PREFERENTIAL PROCUREMENT POLICY FRAMEWORK ACT 2000 AND THE PREFERENTIAL PROCUREMENT REGULATIONS, 2017, THE GENERAL CONDITIONS OF CONTRACT (GCC) AND, IF APPLICABLE, ANY OTHER LEGISLATION OR SPECIAL CONDITIONS OF CONTRACT AND ANY AMENDMENTS THERETO.</p>	
<b>2. TAX COMPLIANCE REQUIREMENTS</b>	
<p>2.1 BIDDERS MUST ENSURE COMPLIANCE WITH THEIR TAX OBLIGATIONS.</p> <p>2.2 BIDDERS ARE REQUIRED TO SUBMIT THEIR UNIQUE PERSONAL IDENTIFICATION NUMBER (PIN) ISSUED BY SARS TO ENABLE THE DEPARTMENT TO VIEW THE TAXPAYER'S PROFILE AND TAX STATUS.</p> <p>2.3 APPLICATION FOR TAX COMPLIANCE STATUS (TCS) OR PIN MAY ALSO BE MADE VIA E-FILING. IN ORDER TO USE THIS PROVISION, TAXPAYERS WILL NEED TO REGISTER WITH SARS AS E-FILERS THROUGH THE WEBSITE WWW.SARS.GOV.ZA.</p> <p>2.4 BIDDERS MAY ALSO SUBMIT A PRINTED TCS TOGETHER WITH THE BID.</p> <p>2.5 IN BIDS WHERE CONSORTIA/ JOINT VENTURES/ SUB-CONTRACTORS ARE INVOLVED, EACH PARTY MUST SUBMIT A SEPARATE PROOF OF TCS / PIN / CSD NUMBER.</p> <p>2.6 WHERE NO TCS IS AVAILABLE BUT THE BIDDER IS REGISTERED ON THE CENTRAL SUPPLIER DATABASE (CSD), A CSD NUMBER MUST BE PROVIDED.</p>	
<b>3. QUESTIONNAIRE TO BIDDING FOREIGN SUPPLIERS</b>	
<p>3.1. IS THE BIDDER A RESIDENT OF THE REPUBLIC OF SOUTH AFRICA (RSA)? <input type="checkbox"/> YES <input type="checkbox"/> NO</p> <p>3.2. DOES THE BIDDER HAVE A BRANCH IN THE RSA? <input type="checkbox"/> YES <input type="checkbox"/> NO</p> <p>3.3. DOES THE BIDDER HAVE A PERMANENT ESTABLISHMENT IN THE RSA? <input type="checkbox"/> YES <input type="checkbox"/> NO</p> <p>3.4. DOES THE BIDDER HAVE ANY SOURCE OF INCOME IN THE RSA? <input type="checkbox"/> YES <input type="checkbox"/> NO</p> <p><b>IF THE ANSWER IS "NO" TO ALL OF THE ABOVE, THEN, IT IS NOT A REQUIREMENT TO OBTAIN A TAX COMPLIANCE STATUS/ TAX COMPLIANCE SYSTEM PIN CODE FROM THE SOUTH AFRICAN REVENUE SERVICE (SARS) AND IF NOT REGISTERED AS PER 2.3 ABOVE.</b></p>	

**NB: FAILURE TO PROVIDE ANY OF THE ABOVE PARTICULARS MAY RENDER THE BID INVALID.**

## **SECTION B: SPECIAL INSTRUCTIONS AND NOTICES TO BIDDERS REGARDING THE COMPLETION OF BIDDING FORMS**

PLEASE NOTE THAT THIS BID IS SUBJECT TO TREASURY REGULATIONS 16A ISSUED IN TERMS OF THE PUBLIC FINANCE MANAGEMENT ACT, 1999, THE KWAZULU-NATAL SUPPLY CHAIN MANAGEMENT POLICY FRAMEWORK AND THE GENERAL CONDITIONS OF CONTRACT. REFER TO THE GENERAL CONDITIONS OF CONTRACT AT THE FOLLOWING WEB ADDRESS:

<http://www.treasury.gov.za/divisions/ocpo/ostb/contracts/default.aspx>

1. Unless inconsistent with or expressly indicated otherwise by the context, the singular shall include the plural and visa versa and with words importing the masculine gender shall include the feminine and the neuter.
2. Under no circumstances whatsoever may the bid forms be retyped or redrafted. Photocopies of the original bid documentation may be used, but an original signature must appear on such photocopies.
3. The bidder is advised to check the number of pages and to satisfy himself that none are missing or duplicated.
4. Bids submitted must be complete in all respects.
5. Bids 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 bid documents.
6. Each bid shall be addressed in accordance with the directives in the bid documents and shall be lodged in a separate sealed envelope, with the name and address of the bidder, the bid number and closing date indicated on the envelope. The envelope shall not contain documents relating to any bid other than that shown on the envelope. If this provision is not complied with, such bids may be rejected as being invalid.
7. All bids received in sealed envelopes with the relevant bid numbers on the envelopes are kept unopened in safe custody until the closing time of the bids. Where, however, a bid is received open, it shall be sealed. If it is received without a bid number on the envelope, it shall be opened, the bid number ascertained, the envelope sealed, and the bid number written on the envelope.
8. A specific box is provided for the receipt of bids, and no bid found in any other box or elsewhere subsequent to the closing date and time of bid will be considered.
9. No bid sent through the post will be considered if it is received after the closing date and time stipulated in the bid documentation, and proof of posting will not be accepted as proof of delivery.
10. No bid submitted by telefax, telegraphic or other electronic means will be considered.
11. Bidding documents must not be included in packages containing samples. Such bids may be rejected as being invalid.
12. Any alteration made by the bidder must be initialled.
13. Use of correcting fluid is prohibited.
14. Bids will be opened in public as soon as practicable after the closing time of bid.
15. Where practical, prices are made public at the time of opening bids.
16. 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.
17. The bidder must initial each and every page of the bid document.

## SECTION C: AUTHORITY TO SIGN A BID

### A. COMPANIES

If a Bidder is a company, a certified copy of the resolution by the Board of Directors, personally signed by the Chairperson of the Board, authorising the person who signs this bid to do so, as well as to sign any contract resulting from this bid and any other documents and correspondence in connection with this bid and/or contract on behalf of the company must be submitted with this bid, that is before the closing time and date of the bid

#### AUTHORITY BY BOARD OF DIRECTORS

By resolution passed by the Board of Directors on.....20.....,  
..... (Full name)  
(whose signature appears below) has been duly authorised to sign all documents in connection with this bid on behalf of  
.....(Name of Company).

IN HIS/ HER CAPACITY AS: .....

SIGNED ON BEHALF OF COMPANY: ..... (PRINT NAME)

SIGNATURE OF SIGNATORY: ..... DATE: .....

WITNESSES: 1 ..... DATE: .....

2 ..... DATE: .....

---

### B. SOLE PROPRIETOR (ONE - PERSON BUSINESS)

I, the undersigned..... (Full name)  
hereby confirm that I am the sole owner of the business trading as:  
.....(Name of Business)

SIGNATURE..... DATE.....

---

### C. PARTNERSHIP

The following particulars in respect of every partner must be furnished and signed by every partner:

FULL NAME OF PARTNER	RESIDENTIAL ADDRESS	SIGNATURE

We, the undersigned Partners in the business trading as .....  
.....(name of partnership)

hereby authorise ..... (full name) to sign this bid as well as any contract resulting from the bid and any other documents and correspondence in connection with this bid and/ or contract on behalf of

.....  
**SIGNATURE**

.....  
**SIGNATURE**

.....  
**SIGNATURE**

.....  
**DATE**

.....  
**DATE**

.....  
**DATE**

---

#### **D. CLOSE CORPORATION**

In the case of a Close Corporation submitting a bid, a certified copy of the Founding/ Amended Founding Statement of such corporation shall be included with the bid, together with the resolution by its members authorising a member or other official of the corporation to sign the documents on their behalf.

By resolution of members at a meeting on ..... 20.....

....., (Full name)

whose signature appears below, has been authorised to sign all documents in connection with this bid on behalf of

.....(Name of Close Corporation)

Trading as .....(Trading name).

**IN HIS/ HER CAPACITY AS:** .....

**SIGNED ON BEHALF OF THE CLOSE CORPORATION:** .....  
(PRINT NAME)

**SIGNATURE OF SIGNATORY:** ..... **DATE:** .....

**WITNESSES:** 1 ..... **DATE:** .....

2 ..... **DATE:** .....

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#### **E. CO-OPERATIVE**

A certified copy of the Constitution of the co-operative must be included with the bid, together with the resolution by its members authoring a member or other official of the co-operative to sign the bid documents on their behalf.

By resolution of members at a meeting on ..... 20.....

..... (full name) whose signature

appears below, has been authorised to sign all documents in connection with this bid on behalf of .....

.....(Name of cooperative)

**SIGNATURE OF AUTHORISED REPRESENTATIVE/SIGNATORY:**

.....

**IN HIS/ HER CAPACITY AS:** .....

**DATE:** .....

**SIGNED ON BEHALF OF CO-OPERATIVE:** .....

**FULL NAME IN BLOCK LETTERS:** .....

**WITNESSES:** 1 .....

**DATE:** .....

2 .....

**DATE:** .....

---

## **F. JOINT VENTURE**

If a bidder is a Joint Venture, a certified copy of the resolution/ agreement passed/ reached, signed by the duly authorised representatives of the entities, authorising the representatives who sign this bid to do so, as well as to sign any contract resulting from this bid and any other documents and correspondence in connection with this bid and /or contract on behalf of the Joint Venture must be submitted with this bid, before the closing time and date of the bid.

### **AUTHORITY TO SIGN ON BEHALF OF THE JOINT VENTURE**

By resolution/agreement passed/reached by the Joint Venture partners  
on.....20.....

..... (Full name)

..... (Full name)

..... (Full name)

..... (Full name)

whose signatures appear below have been duly authorised to sign all documents in connection with this bid on behalf of:  
..... (Name of Joint Venture)

**IN HIS/ HER CAPACITY AS:** .....

**SIGNED ON BEHALF OF (ENTITY NAME):** .....

**SIGNATURE:** ..... **DATE:** .....

**IN HIS/ HER CAPACITY AS:** .....

**SIGNED ON BEHALF OF (ENTITY NAME):** .....

**SIGNATURE:** ..... **DATE:** .....

**IN HIS/ HER CAPACITY AS:** .....



**SIGNED ON BEHALF OF (ENTITY NAME):** .....

**SIGNATURE:** ..... **DATE:** .....

**IN HIS/ HER CAPACITY AS:** .....

**SIGNED ON BEHALF OF (ENTITY NAME):** .....

**SIGNATURE:** ..... **DATE:** .....

**IN HIS/ HER CAPACITY AS:** .....

---

## **G. CONSORTIUM**

If a bidder is a Consortium, a certified copy of the resolution/ agreement passed/ reached, signed by the duly authorised representatives of concerned entities, authorising the representatives who sign this bid to do so, as well as to sign any contract resulting from this bid and any other documents and correspondence in connection with this bid and/ or contract on behalf of the Consortium must be submitted with this bid, before the closing time and date of the bid.

### **AUTHORITY TO SIGN ON BEHALF OF THE CONSORTIUM**

By resolution/agreement passed/reached by the Consortium on.....20.....  
..... (full name)

whose signature appears below have been duly authorised to sign all documents in connection  
with this bid on behalf of:

..... (Name of Consortium)

**IN HIS/ HER CAPACITY AS:** .....

**SIGNATURE:** ..... **DATE:** .....

## SECTION D: DECLARATION OF INTEREST

1. Any legal person, including persons employed by the state, 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 bid (includes an advertised competitive bid, a limited bid, a proposal or written price quotation). In view of possible allegations of favouritism, should the resulting bid, 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 bid(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 bid.

2. In order to give effect to the above, the following questionnaire must be completed and submitted with the bid.

2.1 Full Name of bidder or his or her representative:

.....

2.2 Identity Number: .....

2.3 Position occupied in the Company (Shareholder, Director, Sole Proprietor, Member, Partner, Trustee):

.....

2.4 Registration number of Company, Sole Proprietor, Close Corporation, Partnership, Joint Venture, Consortium or Trust:

.....

2.5 Tax Reference Number: .....

2.6 VAT Registration Number: .....

2.7 The names of all Shareholders/ Directors/ Sole Proprietors, Members, Partners, Trustees, their individual identity numbers, tax reference numbers and, if applicable, employee/ PERSAL numbers must be indicated in paragraph 3 below.

**“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);
- (b) Any municipality or municipal entity;
- (c) Provincial Legislature;
- (d) National Assembly or the National Council of Provinces; or
- (e) Parliament.

**“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.

- 2.8 Are you or any person connected with the bidder presently employed by the State? **YES/NO**
- 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:  
.....  
.....  
.....
- 2.9 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**
- If yes, did you attach proof of such authority to the bid document? **YES/NO**
- (Note: Failure to submit proof of such authority, where applicable, may result in the disqualification of the bid.
- If no, furnish reasons for non-submission of such proof:  
.....  
.....  
.....
- 2.10 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**
- If so, furnish particulars:  
.....  
.....  
.....
- 2.11 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 bid? **YES/NO**
- If so, furnish particulars.  
.....  
.....  
.....
- 2.12 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 bid? **YES/NO**

If so, furnish particulars.

.....

.....

.....

2.13 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

If so, furnish particulars:

.....

.....

.....

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

FULL NAME	IDENTITY NUMBER	PERSONAL INCOME TAX REFERENCE NUMBER	STATE EMPLOYEE NUMBER/ PERSAL NUMBER

**DECLARATION**

I, THE UNDERSIGNED (NAME) .....

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

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

.....  
Signature

.....  
Date

.....  
Position

.....  
Name of Bidder

## SECTION E: DECLARATION OF BIDDER'S PAST SUPPLY CHAIN MANAGEMENT PRACTICES

1. This Standard Bidding Document must form part of all bids invited.
2. It serves as a declaration to be used by institutions in ensuring that when goods and services are being procured, all reasonable steps are taken to combat the abuse of the supply chain management system.
3. The bid of any bidder may be disregarded if that bidder, or any of its directors have-
  - a. abused the institution's supply chain management system;
  - b. committed fraud or any other improper conduct in relation to such system; or
  - c. failed to perform on any previous contract.
4. In order to give effect to the above, the following questionnaire must be completed and submitted with the bid.

ITEM	QUESTION	YES	NO
4.1	Is the bidder or any of its directors listed on the National Treasury's Database of Restricted Suppliers as companies or persons prohibited from doing business with the public sector? <b>(Companies or persons who are listed on this Database were informed in writing of this restriction by the Accounting Officer/Authority of the institution that imposed the restriction after the <i>audi alteram partem</i> rule was applied).</b>  The Database of Restricted Suppliers now resides on the National Treasury's website ( <a href="http://www.treasury.gov.za">www.treasury.gov.za</a> ) and can be accessed by clicking on its link at the bottom of the home page.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
4.1.1	If so, furnish particulars:		
4.2	Is the bidder or any of its directors listed on the Register for Tender Defaulters in terms of section 29 of the Prevention and Combating of Corrupt Activities Act (No 12 of 2004)?  <b>The Register for Tender Defaulters can be accessed on the National Treasury's website (<a href="http://www.treasury.gov.za">www.treasury.gov.za</a>) by clicking on its link at the bottom of the home page.</b>	Yes <input type="checkbox"/>	No <input type="checkbox"/>
4.2.1	If so, furnish particulars:		
4.3	Was the bidder or any of its directors convicted by a court of law (including a court outside of the Republic of South Africa) for fraud or corruption during the past five years?	Yes <input type="checkbox"/>	No <input type="checkbox"/>
4.3.1	If so, furnish particulars:		
4.4	Was any contract between the bidder and any organ of state terminated during the past five years on account of failure to perform on or comply with the contract?	Yes <input type="checkbox"/>	No <input type="checkbox"/>
4.4.1	If so, furnish particulars:		

## CERTIFICATION

I, THE UNDERSIGNED (FULL NAME) .....  
CERTIFY THAT THE INFORMATION FURNISHED ON THIS DECLARATION FORM IS TRUE AND CORRECT.

I ACCEPT THAT, IN ADDITION TO CANCELLATION OF A CONTRACT, ACTION MAY BE TAKEN AGAINST ME  
SHOULD THIS DECLARATION PROVE TO BE FALSE.

.....  
**Signature**

.....  
**Date**

.....  
**Position**

.....  
**Name of Bidder**

**SECTION F: DECLARATION THAT INFORMATION ON CENTRAL SUPPLIER DATABASE IS CORRECT AND UP TO DATE (To be completed by bidder)**

This is to certify that I

.....  
(Name of bidder/authorized representative)

Who represents

.....  
(State name of bidder)

am aware of the contents of the Central Supplier Database with respect to the bidder's details and registration information, and that the said information is correct and up to date as on the date of submitting this bid, and I am aware that incorrect or outdated information may be a cause for disqualification of this bid from the bidding process, and/ or possible cancellation of the contract that may be awarded on the basis of this bid.

.....  
**SIGNATURE OF BIDDER OR AUTHORISED REPRESENTATIVE**

**DATE:** .....

## SECTION G: PREFERENCE POINTS CLAIM FORM IN TERMS OF THE PREFERENTIAL PROCUREMENT REGULATIONS 2017

This preference form must form part of all bids invited. It contains general information and serves as a claim form for preference points for Broad-Based Black Economic Empowerment ( ) Status Level of Contribution

**NB: BEFORE COMPLETING THIS FORM, BIDDERS MUST STUDY THE GENERAL CONDITIONS, DEFINITIONS AND DIRECTIVES APPLICABLE AS PRESCRIBED IN THE PREFERENTIAL PROCUREMENT REGULATIONS, 2017.**

### 1. GENERAL CONDITIONS

- 1.1. The following preference point systems are applicable to all bids:
- 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 value of this bid is estimated not to exceed R50 000 000 (all applicable taxes included) and therefore the 80/20 preference point system shall be applicable.
- 1.3. Points for this bid shall be awarded for:
- (a) Price; and
  - (b) Status Level of Contributor.
- 1.4. The maximum points for this bid are allocated as follows:
- | CATEGORY  | POINTS     |
|---|------------|
| PRICE   | 90 or 80   |
| STATUS LEVEL OF CONTRIBUTOR                       | 10 or 20   |
| <b>Total points for Price and must not exceed</b> | <b>100</b> |
- 1.5. Failure on the part of a bidder to submit proof of Status level of contributor together with the bid will be interpreted to mean that preference points for Status level of contribution are not claimed.
- 1.6. The department reserves the right to require of a bidder, either before a bid is adjudicated or at any time subsequently, to substantiate any claim in regard to preferences, in any manner required by the department.

### 2. DEFINITIONS

- a) **"B-BBEE"** means broad-based black economic empowerment as defined in section 1 of the Broad-Based Black Economic Empowerment Act;
- b) **"B-BBEE status level of contributor"** means the B-BBEE status of an entity in terms of a code of good practice on black economic empowerment, issued in terms of section 9(1) of the Broad-Based Black Economic Empowerment Act;
- c) **"Bid"** means a written offer in a prescribed or stipulated form in response to an invitation by an organ of state for the provision of goods or services, through price quotations, advertised competitive bidding processes or proposals;
- d) **"Black Designated Groups"** has the meaning assigned to it in the codes of good practice issued in terms of section 9(1) of the Broad-Based Black Economic Empowerment Act;
- e) **"Black People"** has the meaning assigned to it in section 1 of the Broad-Based Black Economic Empowerment Act;



- f) **“Broad-Based Black Economic Empowerment Act”** means the Broad-Based Black Economic Empowerment Act, 2003 (Act No. 53 of 2003);
- g) **“Co-operative”** means a co-operative **registered** in terms of section 7 of the Cooperatives Act, 2005 (Act No. 14 of 2005);
- h) **“EME”** means an Exempted Micro **Enterprise** in terms of a code of good practice on black economic empowerment issued in terms of section 9 (1) of the Broad-Based Black Economic Empowerment Act;
- i) **“Functionality”** means the ability of a tenderer to provide goods or services in accordance with specifications as set out in the tender documents.
- j) **“Military Veteran”** has the meaning assigned to it in section 1 of the Military Veterans Act, 2011 (Act No. 18 of 2011);
- k) **“prices” includes** all applicable taxes less all unconditional discounts;
- l) **“proof of status level of contributor” means:**
  - 1) Status level certificate issued by an authorized body or person;
  - 2) A sworn affidavit as prescribed by the Codes of Good Practice;
  - 3) Any other requirement prescribed in terms of the Act;
- m) **“QSE”** means a qualifying small business enterprise in terms of a code of good practice on black economic empowerment issued in terms of section 9 (1) of the Broad-Based Black Economic Empowerment Act;
- n) **“rand value”** means the total estimated value of a contract in Rand, calculated at the time of bid invitation, and includes all applicable taxes; and
- o) **“stipulated minimum threshold”** means the minimum threshold stipulated in terms of regulation 8(1)(b).

### 3. POINTS AWARDED FOR PRICE

#### 3.1 THE 80/20 OR 90/10 PREFERENCE POINT SYSTEMS

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

80/20	or	90/10
$P_s = 80 \left( 1 - \frac{P_t - P_{\min}}{P_{\min}} \right)$	or	$P_s = 90 \left( 1 - \frac{P_t - P_{\min}}{P_{\min}} \right)$

Where

$P_s$  = Points scored for price of bid under consideration  
 $P_t$  = Price of bid under consideration  
 $P_{\min}$  = Price of lowest acceptable bid

#### 4. POINTS AWARDED FOR STATUS LEVEL OF CONTRIBUTOR

- 4.1 In terms of Regulation 6 (2) and 7 (2) of the Preferential Procurement Regulations, preference points must be awarded to a bidder for attaining the status level of contribution in accordance with the table below:

STATUS LEVEL OF CONTRIBUTOR	NUMBER OF POINTS (90/10 SYSTEM)	NUMBER OF POINTS (80/20 SYSTEM)
1	10	20
2	9	18
3	6	14
4	5	12
5	4	8
6	3	6
7	2	4
8	1	2
Non-compliant contributor	0	0

#### 5. BID DECLARATION

- 5.1 Bidders who claim points in respect of Status Level of Contribution must complete the following:

#### 6. STATUS LEVEL OF CONTRIBUTOR CLAIMED IN TERMS OF PARAGRAPHS 1.4 AND 4.1

- 6.1 Status Level of Contributor: = ..... (maximum of 10 or 20 points) (Points claimed in respect of paragraph 6.1 must be in accordance with the table reflected in paragraph 4 and must be substantiated by relevant proof of status level of contributor.

#### 7. SUB-CONTRACTING

- 7.1 Will any portion of the contract be sub-contracted?

**(Tick applicable box)**

YES	<input type="checkbox"/>	NO	<input type="checkbox"/>
-----	--------------------------	----	--------------------------

- 7.1.1 If yes, indicate:

- What percentage of the contract will be subcontracted.....%
- The name of the sub-contractor.....
- The status level of the sub-contractor.....
- Whether the sub-contractor is an EME or QSE

**(Tick applicable box)**

YES	<input type="checkbox"/>	NO	<input type="checkbox"/>
-----	--------------------------	----	--------------------------

- Specify, by ticking the appropriate box, if subcontracting with an enterprise in terms of Preferential Procurement Regulations, 2017:

DESIGNATED GROUP: AN EME OR QSE WHICH IS AT LAST 51% OWNED BY:	EME √	QSE √
Black people		
Black people who are youth		
Black people who are women		
Black people with disabilities		
Black people living in rural or underdeveloped areas or townships		
Cooperative owned by black people		
Black people who are military veterans		
<b>OR</b>		
Any EME		
Any QSE		

## 8. DECLARATION WITH REGARD TO COMPANY/FIRM

8.1 Name of company/firm: .....

8.2 VAT registration number: .....

8.3 Company registration number: .....

8.4 TYPE OF COMPANY/ FIRM

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

[TICK APPLICABLE BOX]

8.5 DESCRIBE PRINCIPAL BUSINESS ACTIVITIES

.....  
.....  
.....

8.6 COMPANY CLASSIFICATION

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

[TICK APPLICABLE BOX]

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

8.8 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 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 .....

.....

.....

## SECTION H: CERTIFICATE OF INDEPENDENT BID DETERMINATION

- 1 This Standard Bidding Document (SBD) must form part of all bids invited.
2. Section 4 (1) (b) (iii) of the Competition Act No. 89 of 1998, as amended, prohibits an agreement between, or concerted practice by, firms, or a decision by an association of firms, if it is between parties in a horizontal relationship and if it involves collusive bidding (or bid rigging). Collusive bidding is a *pe se* prohibition meaning that it cannot be justified under any grounds.
- 3 Treasury Regulation 16A9 prescribes that accounting officers and accounting authorities must take all reasonable steps to prevent abuse of the supply chain management system and authorizes accounting officers and accounting authorities to:
  - a. disregard the bid of any bidder if that bidder, or any of its directors have abused the institution's supply chain management system and or committed fraud or any other improper conduct in relation to such system.
  - b. cancel a contract awarded to a supplier of goods and services if the supplier committed any corrupt or fraudulent act during the bidding process or the execution of that contract.
- 4 In order to give effect to the above, the attached Certificate of Bid Determination (SBD 9) must be completed and submitted with the bid:

**Bid rigging (or collusive bidding)** occurs when businesses, that would otherwise be expected to compete, secretly conspire to raise prices or lower the quality of goods and / or services for purchasers who wish to acquire goods and / or services through a bidding process. Bid rigging is, therefore, an agreement between competitors not to compete.

## CERTIFICATE OF INDEPENDENT BID DETERMINATION

I, the undersigned, in submitting the accompanying bid:

---

(Bid Number and Description)

in response to the invitation for the bid made by:

---

(Name of Institution)

do hereby make the following statements that I certify to be true and complete in every respect:

I certify, on behalf of: \_\_\_\_\_ that:  
(Name of Bidder)

1. I have read, and I understand the contents of this Certificate;
2. I understand that the accompanying bid will be disqualified if this Certificate is found not to be true and complete in every respect;
3. I am authorized by the bidder to sign this Certificate, and to submit the accompanying bid, on behalf of the bidder;
4. Each person whose signature appears on the accompanying bid has been authorized by the bidder to determine the terms of, and to sign the bid, on behalf of the bidder;

5. For the purposes of this Certificate and the accompanying bid, I understand that the word “competitor” shall include any individual or organization, other than the bidder, whether or not affiliated with the bidder, who:
  - a) has been requested to submit a bid in response to this bid invitation;
  - b) could potentially submit a bid in response to this bid invitation, based on their qualifications, abilities or experience; and
  - c) provides the same goods and services as the bidder and/or is in the same line of business as the bidder
6. 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 consortium will not be construed as collusive bidding.
7. In particular, without limiting the generality of paragraphs 6 above, there has been no consultation, communication, agreement or arrangement with any competitor regarding:
  - a) prices;
  - b) geographical area where product or service will be rendered (market allocation)
  - c) methods, factors or formulas used to calculate prices;
  - d) the intention or decision to submit or not to submit, a bid;
  - e) the submission of a bid which does not meet the specifications and conditions of the bid; or
  - f) bidding with the intention not to win the bid.
8. In addition, there have been no consultations, communications, agreements or arrangements with any competitor regarding the quality, quantity, specifications and conditions or delivery particulars of the products or services to which this bid invitation relates.
9. 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.
10. 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.

.....  
Signature

.....  
Date

.....  
Position

.....  
Name of Bidder

**SECTION I: RECORD OF AMENDMENTS TO BID DOCUMENTS**

I / We confirm that the following communications amending the bid documents that I / we received from KwaZulu-Natal Department of Health or their representative before the closing date for submission of bids have been taken into account in this bid.

ADDENDUM NO.	DATE	TITLE OR DETAILS

SIGNATURE: ..... DATE: .....  
(of person authorized to sign on behalf of the Bidder)

**SECTION J: GENERAL CONDITIONS OF CONTRACT**

<http://www.treasury.gov.za/divisions/ocpo/sc/GeneralConditions/General%20Conditions%20of%20Contract.pdf>

❖ I have read, understand and accept the General conditions of the contract which are binding upon me.

.....  
**Signature**

.....  
**Date**

.....  
**Name of Bidder**



## SECTION K: SPECIAL TERMS AND CONDITIONS

The bid is issued in accordance with the following subject to the provisions of the General Conditions of Contract:

- i. Section 217 of the Constitution,
- ii. The PFMA and its Regulations in general,
- iii. The Preferential Procurement Policy Framework Act ( PPPFA) of 2000
- iv. National Treasury guidelines, and
- v. Revised PPPFA Regulations of 2017

The special terms and conditions are supplementary to that of the General Conditions of Contract. Where, however, the special terms and conditions are in conflict with the General Conditions of Contract, the Special Terms and Conditions prevail.

- (a) **Bidder/s must ensure that they are fully aware of all the conditions contained in this bid document.**
- (b) **Only bidders that fully meet the specifications and all conditions will be considered.**

### 1. CONDITIONS OF BID

The bid is issued in accordance with the following conditions:

#### 1.1 ACCEPTANCE OF A BID

- 1.1.1 The Department of Health Bid Adjudication Committee is under no obligation to accept any bid.
- 1.1.2 The financial standing of a bidder and its ability to supply goods or render services may be examined before the bid is considered for acceptance.

#### 1.2 CERTIFICATE OF COMPLIANCE

- 1.2.1 If the bidder submits offers for items that make reference to South African National Standards (SANS) or South African Bureau of Standards (SABS) specifications, a Certificate of Compliance must be submitted with the bid document at the time of closing of the bid. SABS/SANS can be contacted for testing and conformity services at Tel: 031 203 2900/ Fax: 031 203 2907. SANS, SABS AND CKS specifications will be for the account of the prospective bidder.
- 1.2.2 Failure to submit the certificate, where applicable, will result in the bid being disqualified. The Department reserves its rights to contact SABS/SANS/CKS for testing and conformity services.
- 1.2.3 The South African National Accreditation System (SANAS) is recognized by the South African Government as the single National Accreditation Body that gives formal recognition that Laboratory, Certification Bodies, Inspection Bodies, Proficiency Testing Scheme Providers and Good Laboratory Practice (GLP) test facilities are competent to carry out specific tasks. This organization can be contacted as follows: Tel: 012 3943760: Fax: 012 3940526.
- 1.2.4 Prior to an award of the bid being made and/or during the evaluation process, the Department of Health reserves the right to conduct inspections of the premises of the most acceptable bidder. Therefore, premises of the bidder shall be open, at reasonable hours, for inspection by a representative of the Department of Health or organization acting on its behalf.
- 1.2.5 Any specification/s and conformity testing will be for the account of the prospective bidder.

- 1.2.6 In the event of the bidder not being the actual manufacturer and will be sourcing the product(s) from the manufacturer, a letter from the manufacturer confirming firm supply arrangement(s) including lead times in this regard, must accompany the bid at closing date and time. If the bidder is the manufacturer, a letter confirming that the bidder is the manufacturer should accompany the bid at the closing date and time.

### **1.3 COMPLIANCE WITH SPECIFICATION**

- 1.3.1 Offers must comply strictly with the specification.
- 1.3.2 Offers exceeding specification requirements will be deemed to comply with the specification.
- 1.3.3 The quality of services/ supply must not be less than what is specified.

### **1.4 LATE BIDS**

- 1.4.1 Bids are late if they are received at the address indicated in the bid documents after the closing date and time.

### **1.5 MORE THAN ONE OFFER/ COUNTER OFFERS**

- 1.5.1 Should the bidder make more than one offer, where applicable, against any individual item, such offer/s must be detailed in the Schedule of Additional Offer/s. The Department reserves its rights in and to the consideration of any additional offer/s subject to compliance with specification and the bidding conditions.
- 1.5.2 Bidders' attention is drawn to the fact that counter offers with regard to any of the abovementioned Special Terms and Conditions will invalidate such bids.
- 1.5.3 Bidders are at liberty to bid for one, a number of items, or bid for all items. If a bidder is not bidding for all the items, the appropriate price page must reflect: 'nil quote'.

### **1.6 ONLY ONE OFFER RECEIVED**

- 1.6.1 Where only 1 offer is received, the Department of Health will determine whether the price is fair and reasonable. Proof of reasonableness will be determined as follows:
- (i) Comparison with prices, after discounts, to the bidder's other normal clients and the relative discount that the State enjoys;
  - (ii) Where this is not possible, profit before tax based on a full statement of relevant costs; and
  - (iii) In all cases, comparison with previous bid prices where these are available.

### **1.7. AWARD OF BID (S)**

- 1.7.1. The Department of Health Bid Adjudication Committee reserves the right to award the bid to more than one bidder, provided that the respective bidders' offers comply with the specification and meets all the conditions attached to the bid. Bidders must ensure that they quote as per the price page failing which they will be disqualified.
- 1.7.2. Notification of the intention to award the bid shall be in the same media that the bid was advertised.
- 1.7.3. In terms of Practice Note Number: SCM-07 of 2006, Section 5: Appeal Procedure, 5.1 "A bidder aggrieved by a decision of the Departmental Bid Adjudication Committee or a delegate of an accounting officer may appeal to the Bid Appeals Tribunal in the prescribed manner"

- 1.7.4. After all appeals, should they be lodged, have been dealt with by the Bid Appeals Tribunal, the successful bidder (s) shall be notified in writing by a duly authorised official of the Department of Health, Central Supply Chain Management Unit. A formal contract will then be entered into by both parties.

## **1.8. REGISTRATION ON THE CENTRAL SUPPLIER DATABASE (CSD)**

- 1.8.1. A bidder submitting an offer must be registered on the Central Supplier Database. A bidder who has submitted an offer and is not registered on the Central Supplier Database will not be considered.
- 1.8.2. Each party to a joint venture/ consortium must be registered on the Central Suppliers Database at the time of submitting the bid.

**NB.: IF A BIDDER IS FOUND TO BE EMPLOYED BY THE STATE AND IS ON THE CENTRAL SUPPLIER DATABASE, THE BIDDER WILL BE DISQUALIFIED.**

## **1.9. TAX COMPLIANCE REQUIREMENTS**

- 1.9.1. Bidders must ensure compliance with their tax obligations.
- 1.9.2. No award may be made to any bidder who is not tax compliant either on the Central Supplier Database or SARS eFiling system at the time of finalisation of the award of the bid. The Onus is on the bidder to ensure that their tax affairs are in order and is valid on the CSD.

## **1.10. TRUST, CONSORTIUM OR JOINT VENTURE**

- 1.10.1. In terms of the Preferential Procurement Policy Framework Act and Regulations, as amended, a Trust, Consortium or Joint Venture must submit a consolidated Status Level Verification Certificate for every separate bid.
- 1.10.2. A separate B-BBEE Certificate must be submitted by each company participating in the Trust, Consortium or Joint Venture.
- 1.10.3. The non-submission of a B-BBEE Certificate by a Trust, Consortium or Joint Venture will result in zero (0) preference points being allocated for evaluation purposes.
- 1.10.4. Should this bid be submitted by a Joint Venture, the Joint Venture agreement must accompany the bid document.
- 1.10.5. The Joint Venture agreement must clearly specify the percentage of the contract to be undertaken by each company participating therein.
- 1.10.6. The Joint Venture/Consortium must submit a formal agreement that outlines the roles and responsibilities of each member of the Joint Venture/ Consortium, nomination of an authorised person to represent the Joint Venture or Consortium in all matters relating to this bid and the details of the bank account for payments to be affected.
- 1.10.7. No award will be made to a Trust/ Joint Venture/ Consortium that is not tax compliant at the finalisation of the award.
- 1.10.8. For verification purposes, each party must submit separate proof of TCS/ PIN / CSD number.

## **1.11. VALIDITY PERIOD OF BID AND EXTENSION THEREOF**

- 1.11.1. The validity (binding) period for the bid will be **120 days** from close of bid.
- 1.11.2. However, circumstances may arise whereby the department may request bidders to extend the validity (binding) period. Should this occur, the department will request bidders to extend the validity (binding) period under the

same terms and conditions as originally offered for by bidders? This request will be done before the expiry of the original validity (binding) period.

## **2. SPECIAL CONDITIONS OF CONTRACT**

### **2.1 CHANGE OF ADDRESS**

- 2.2.1 Bidders must advise the Department of Health's Central Supply Chain Management Unit, Contract Administration Section, should their ownership and/or address (domicilium citandi et executandi) details change from the time of bidding to the expiry of the contract.

### **2.2 DELIVERY AND PACKAGING**

- 2.2.1 Basis of delivery: Delivery of respiratory aids must be made in accordance with the instructions appearing on the official order form (various institutions).
- 2.2.2 All deliveries must take place from Monday to Friday between 08h00 and 14h00.
- 2.2.3 In emergency cases, the Department of Health reserves the right to request the successful bidder/s to effect deliveries at any given time including Saturdays, Sundays and public holidays.
- 2.2.4 Random inspection and sampling of items will be conducted upon delivery to verify quantity and compare the item against the contract sample and any other quality accreditation that is prescribed.
- 2.2.5 It is the contractor's responsibility to off load the delivery vehicle.
- 2.2.6 Order details must be presented upon delivery on delivery notes.
- 2.2.7 The following information must appear on the outer packaging of the carton/box:
- (a) Name of the manufacturer/supplier
  - (b) Description of item
  - (c) Date of manufacture

### **2.3 DELIVERY CONDITIONS**

- 2.3.1 Delivery of products must be made in accordance with the instructions appearing on the official order form.
- 2.3.2 All deliveries or dispatches must be accompanied by a delivery note stating the official order number against the delivery that has been affected.
- 2.3.3 In respect of items awarded to them, contractors must adhere strictly to the delivery periods stipulated by them in their bid document.
- 2.3.4 The instructions appearing on the official order form regarding the supply, dispatch and submission of invoices must be strictly adhered to.
- 2.3.5 All invoices must be submitted in the original.
- 2.3.6 Deliveries not complying with the order form will be returned to the contractor at the contractor's expense.
- 2.3.7 No locally manufactured product may be substituted during the contract period with an imported product, and vice versa, without prior approval of Contract Management at Central Supply Chain Management, Department of Health.

## **2.4 ENTERING OF HOSPITAL/CLINIC STORES**

- 2.4.1 No representative from a company shall be permitted to enter the hospital/clinic premises, buildings or containers where stores are kept unless he/she is accompanied by the responsible official in charge of stores. Before entering the hospital/clinic premises, buildings or containers where stores are kept, the company representative must in writing, motivate why entry is necessary and written authority must be obtained to enter from the Head of the Institution.

## **2.5 EQUAL BIDS**

- 2.5.1 If two or more tenderers score an equal total number of points, the contract must be awarded to the tenderer that scored the highest points for B-BBEE.
- 2.5.2 If functionality is part of the evaluation process and two or more tenderers score equal total points and equal preference points, the contract must be awarded to the tenderer that scored the highest points for functionality.
- 2.5.3 If two or more tenderers score equal total points in all respects, the award must be decided by the drawing of lots.

## **2.6 FIRM PRICES AND ESCALATIONS**

- 2.6.1 This bid requires that all bid prices offered are firm for the three years of the contract. If a non-firm price is offered, the bidder shall be disqualified for not complying with the conditions of the bid.
- 2.6.2 In respect of rates of exchange, it is mandatory that bidders take forward cover upon award of the contract, for the contract period, with a recognized Financial Institution. Proof of this forward cover must be submitted to the contract management unit upon signing of the contract. Therefore, a price adjustment in respect of a rate of exchange claim will not be considered.

## **2.7 STATEMENT OF SUPPLIES AND SERVICES**

- 2.7.1 The contractor shall, monthly, furnish particulars of supplies delivered or services executed. Such information must be submitted to the Department of Health Supply Chain Management, Contract Management as follows:
- (i) Name of institution.
  - (ii) Orders received – order number & catalogue number & quantity delivered.
  - (iii) Price.
- 2.7.2 Historical value and volume reports may be requested by the Department of Health, Supply Chain Management, during the term of the contract for the following:

### **a) SUPPLIER MEASURES**

- Delivery period adherence
- Quality adherence

- 2.7.3 This information will be submitted at the expense of the contractor.

## **2.8 INSPECTION FOR QUALITY**

- 2.8.1 All deliveries to authorised participants will be subjected to a visual examination and scrutiny by the relevant participants, and/or inspection for quality by Provincial Quality Control Laboratories in the Republic of South

Africa, and/or inspection for quality by an accredited South African National Accreditation Section (SANAS) testing agency.

2.8.2 In the event of products tested, the contractor will bear the cost of any item failing to meet the relevant standard.

## **2.9 INVOICES AND PAYMENTS**

2.9.1 All invoices submitted by the Contractor must be Tax Invoices indicating item description, catalogue number, quantity ordered and quantity delivered, unit price, total price, the amount of tax charged and the total invoice amount.

2.9.2 A tax invoice shall be in the currency of the republic of South Africa and shall contain the following particulars:

- (a) The name, address and registration number of the supplier;
- (b) The name and address of the recipient;
- (c) An individual serialized number and the date upon which the tax invoice is issued;
- (d) A description of the goods or services supplied;
- (e) The quantity or volume of the goods or services supplied
- (f) The value of the supply, the amount of tax charged and the consideration for the supply; or
- (g) Where the amount of tax charged is calculated by applying the tax fraction to the consideration, the consideration for the supply and either the amount of the tax charged, or a statement that it includes a charge in respect of the tax and the rate at which the tax was charged.

2.9.3 A contractor shall be paid by the institution concerned, in accordance with supplies delivered and services rendered.

2.9.4 Should a contractor indicate a special discount on his/her account provided payment is made within a certain time, every effort shall be made to take advantage of such discount.

2.9.5 Any query concerning the non-payment of accounts must be directed to the institution concerned. The following protocol will apply if accounts are queried:

- (i) Contact must be made with the officer-in-charge of stores;
- (ii) If there is no response from stores, the finance manager of the institution must be contacted.

## **2.10 IRREGULARITIES**

2.10.1 Companies are encouraged to advise the Department of Health timeously of any possible irregularities which might come to their notice in connection with this or other contracts.

## **2.11 PERIOD OF CONTRACT**

2.11.1 Three-year contract.

## **2.12 QUALITY CONTROL TESTING OF PRODUCTS**

2.12.1 If it is discovered that the product supplied is not in accordance with the specification the following will occur:

- (i) Testing charges will be for the account of the principal contractor;
- (ii) Possible cancellation of the contract with the principal contractor;
- (iii) Reporting such negligence by the principal contractor to the provincial and national treasury for listing on the Restricted Suppliers' Database.

## **2.13 RATE OF EXCHANGE**

- 2.13.1 All bids involving imported products must use the rate of exchange that was applicable 14 days prior to the closing date indicated in the bid document. If this day falls on a weekend or public holiday, the next working day must be used.
- 2.13.2 Bidders must submit documentary proof (in the form of a certified copy) from their bank or any recognized legal financial Institution, clearly indicating what the rate of exchange was 14 days prior to the closing date, as mentioned above. Information can be sourced from the internet from a financial Institution website.
- 2.13.3 The Department of Health reserves the right to renegotiate the price should there be a reduction of the price in the market.
- 2.13.4 This clause must be read in conjunction with paragraphs 2.6.1 and 2.6.2

## **2.14 SAMPLES**

- 2.14.1 Samples will not be accepted with the closing of the bid document.
- 2.14.2 A sample meeting will be arranged with selected companies whereby the companies will be invited to forward their samples on a specified date and time.
- 2.14.3 Samples must be made available for the sample meeting, failure to provide a sample will reject their bid offer.
- 2.14.4 Samples shall be supplied by the bidder at his/her own cost/risk. Samples must be packaged as per the specification. Failure to do so will render the bid invalid.
- 2.14.5 Representative samples will not be accepted.
- 2.14.6 The Department reserves the right not to return such samples and to dispose of them at its discretion.
- 2.14.7 Samples must be clearly marked: Item number:
  - Brand Name
  - Name of the Company
  - Bid number
  - Name of the manufacturer/supplier
  - Description of item
  - Date of manufacture
- 2.14.8 The award of this bid will be based on the sample submitted from a manufacturer based on a letter of undertaking, which is compliant to specification. If, during the contract, the awarded supplier wishes to change the item being supplied, the service provider shall apply to the Department in writing, giving reasons why they want to change the product being supplied, which the Department shall consider. This process will be subject to the sample being submitted to the technical committee for evaluation and if in order, to the adjudication committee for approval. This will be done via the contract management unit of the Department. If there is a change in the product being supplied, and no prior approval has been granted, the Department reserves its right to cancel the contract.



**N.B Failure to clearly mark the samples submitted shall result in the samples not being evaluated and eliminated from further consideration.**

## **2.15 UNSATISFACTORY PERFORMANCE**

2.15.1 Unsatisfactory performance occurs when performance is not in accordance with the contract conditions.

(i) The institution shall warn the contractor by registered/certified mail that action will be taken in accordance with the contract conditions unless the contractor complies with the contract conditions and delivers satisfactory supplies or services within a specified reasonable time (7 days minimum). If the contractor does not perform satisfactorily despite the warning the institution will:

(a) Take necessary action in terms of its delegated powers.

(ii) When correspondence is addressed to the contractor, reference will be made to the contract number/item number/s and an explanation of the complaint.

## **2.16 PREFERENCES**

2.16.1 Should the Contractor apply for preferences in the submission of his bid, and it is found at a later stage that these applications were incorrect or made under false pretences, the Department may, at its own right:

i. Recover from the Contractor all costs, losses or damages incurred or sustained by the Department as a result of the award of the Contract; and/or

ii. Cancel the contract and claim any damages which the Department may suffer by having to make less favourable arrangements after such cancellation.

iii. The Department may impose penalties, however, only if provision therefore is made in the Special Conditions of Bid.

## **2.17 RESTRICTION OF BIDDING**

The Accounting Officer or his/her delegate must:

a) Notify the supplier and any other person of the intention to restrict it doing business with KZN-DoH by registered mail. The letter of restriction must provide for:

✓ The grounds for restriction;

✓ The period of restriction which must not exceed 10 years;

✓ A period of 14 calendar days for the supplier to provide reasons why the restriction should not be imposed.

b) The Accounting Officer his/her delegate:

✓ May regard the intended penalty as not objected to and may impose such penalty on the supplier, should the supplier fail to respond within the 14 days; and

✓ Must assess the reasons provided by the supplier and take the final decision.

c) If the penalty is imposed, the Accounting Officer must inform National Treasury of the restriction within 7 calendar days and must furnish the following information:

✓ The name and address of the entity/ person to be restricted;

✓ The identity number of individuals and the registration number of the entity; and

- ✓ The period of restriction.
- d) National Treasury will load the details on the Database of Prohibited Vendors.
- e) The restriction period applicable will be based on the value of award/s made to the supplier over a financial year.  
The table below illustrates the restriction period that will be applicable per the award threshold:

## **2.18 CONTRACTOR'S LIABILITY**

- 2.18.1 In the event of the contract being cancelled by the Department in the exercise of its rights in terms of these conditions, the Contractor shall be liable to pay to the Department any losses sustained and/or additional costs or expenditure incurred as a result of such cancellation, and the Department shall have the right to recover such losses, damages or additional costs by means of set-off from moneys due or which may become due in terms of the contract or any other contract or from guarantee provided for the due fulfilment of the contract and, until such time as the amount of such losses, damages or additional costs have been determined, to retain such moneys or guarantee or any deposit as security for any loss which the Department may suffer or may have suffered.
- 2.18.2 The Contractor may be held responsible for any consequential damages and loss sustained which may be caused by any defect, latent or otherwise, in supply or service rendered or if the goods or service as a result of such defect, latent or otherwise, does not conform to any condition or requirement of the contract.

## **2.19 DEPARTMENTAL PROPERTY IN POSSESSION OF A CONTRACTOR**

- 2.19.1 The Department's property supplied to a Contractor for the execution of a contract remains the property of the Department and shall at all times be available for inspection by the Department or its representatives. Any such property in the possession of the Contractor on the completion of the contract shall, at the Contractor's expense, be returned to the Department forthwith.
- 2.19.2 The Contractor shall be responsible at all times for any loss or damages to the Department's property in his possession and, if required, he shall furnish such security for the payment of any such loss or damages as the Department may require.

## **2.20 RIGHTS TO PROCURE OUTSIDE THE CONTRACT**

- 2.20.1 The Department reserves the right to procure goods outside the contract in cases of urgency or emergency or if the quantities are too small to justify delivery costs, or if the goods are obtainable from another organ of State or if the Contractor's point of supply is not situated at or near the place where the goods are required or if the Contractor's goods are not readily available.
- 2.20.2 No provision in a contract shall be deemed to prohibit the obtaining of goods or services from a Department or local authority.

## **2.21 USE OF CONTRACT DOCUMENTS AND INFORMATION INSPECTION**

- 2.21.1 The Contractor shall not, without the Department'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 Department in connection therewith, to any person other than a person employed by the Contractor 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.
- 2.21.2 The Contractor shall not, without the Department's prior written consent, make use of any document or information mentioned in GCC clause 2.21.1 except for purposes of performing the contract.

- 2.21.3 Any document, other than the contract itself mentioned in GCC clause (2.21.1) shall remain the property of the Department and shall be returned (all copies) to the Department on completion of the Contractor's performance under the contract of so required by the Department.
- 2.21.4 The Contractor shall permit the Department to inspect the Contractor's records relating to the performance of the Contractor and to have them audited by auditors appointed by the Department, if so required by the Department.

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

1.	ZNB 6685/2/2021-H	LMA DEVICE – Reusable: Per Unit				
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 01	Laryngeal mask airway device - reusable <b>Size: 1</b>				
	30 301 02	Laryngeal mask airway device - reusable <b>Size: 1.5</b>				
	30 301 05	Laryngeal mask airway device -reusable <b>Size: 3</b>				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

**VARIOUS INSTITUTIONS**

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

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2.	ZNB 6685/2/2021-H		LMA DEVICE WITH GASTRIC DRAIN TUBE - Non-inflatable: Per box of 10			
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 19	Laryngeal mask airway device with gastric drain tube. Non inflatable - single use <b>Size: 2.5</b>				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

VARIOUS INSTITUTIONS

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

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(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

3.	ZNB 6685/2/2021-H	DOUBLE LUMEN BLIND INSERTION AIRWAY DEVICES: Per Unit				
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	DLBI 1	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Small</b>				
	DLBI 2	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Medium</b>				
	DLBI 3	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Large</b>				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by: KZN DEPARTMENT OF HEALTH

-At: **VARIOUS INSTITUTIONS**

Country of origin .....

Brand .....

Delivery period (on order) .....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

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(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

4.	ZNB 6685/2/2021-H		EMERGENCY CRICOTHYROTOMY: Per Unit			
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 45	Emergency Cricothyrotomy Kit – with <b>Varess</b> needle				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3**

**The annual unit price will be the applicable (contractual) price per year per item.**

**Bidders must bid as per the price page failing which they will be disqualified.**

Required by: KZN DEPARTMENT OF HEALTH

-At: **VARIOUS INSTITUTIONS**

Country of origin .....

Brand .....

Delivery period (on order) .....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....	.....	.....	.....
(Signature of Bidder)	Date	(Signature of Witness)	Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
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OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

5.	ZNB 6685/2/2021-H	AIRWAY EXCHANGE CATHETER: Per Unit				
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 47	Airway Exchange Catheter. <b>Infant 8 Fr</b>				
	30 301 50	Airway Exchange Catheter. <b>Adult 11 Fr</b> Extra length and extra firm				
	30 301 51	Airway Exchange Catheter. <b>Adult 14 Fr</b> Extra length and extra firm				
	AEC_SE	Airway Exchange Catheter with guidewire – staged extubation kit.				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3**

**The annual unit price will be the applicable (contractual) price per year per item.**

**Bidders must bid as per the price page failing which they will be disqualified.**

Required by: KZN DEPARTMENT OF HEALTH

-At: **VARIOUS INSTITUTIONS**

Country of origin .....

Brand .....

Delivery period (on order) .....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

..... (Signature of Bidder)	..... Date	..... (Signature of Witness)	..... Date
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**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

6.	ZNB 6685/2/2021-H		INTUBATING INTRODUCER: Per Unit			
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 52	Intubation Catheter <b>19 Fr</b> for use with a flexible fibre-optic scope				
	30 301 53	Intubating introducer – single use <b>14 Fr</b>				
	30 301 54	Intubating introducer – single use <b>8 Fr</b>				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

VARIOUS INSTITUTIONS

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

7.	ZNB 6685/2/2021-H		INTUBATION STYLET: PER BOX OF 100			
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 62	Intubation Stylet, <b>Neonatal</b>				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3**

**The annual unit price will be the applicable (contractual) price per year per item.**

**Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

**VARIOUS INSTITUTIONS**

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder)

.....  
Date

.....  
(Signature of Witness)

.....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

8.	ZNB 6685/2/2021-H		RETROGRADE:PER UNIT			
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 66	Retrograde Intubation Kit.6 Fr				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

**VARIOUS INSTITUTIONS**

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

Name of bidder.....	Bid number: <b>ZNB 6685/2/2021-H</b>
Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

9.	ZNB 6685/2/2021-H	INTUBATION SET:PER UNIT				
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 67	Retrograde Intubation Kit.11 Fr				
	30 301 68	Retrograde Intubation Kit.14Fr				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

VARIOUS INSTITUTIONS

Country of origin

.....

Brand

.....

Delivery period (on order)

.....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder).....  
Date.....  
(Signature of Witness).....  
Date

**SECTION M: PRICING SCHEDULE: refer to specification schedule for item description**

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Closing Time 11:00	Closing Date: <b>04 MAY 2021</b>

OFFER TO BE VALID FOR 120 DAYS FROM THE CLOSING DATE OF BID.

**DESCRIPTION: ZNB 6685/2/2021: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS.**

10.	ZNB 6685/2/2021-H	ORAL FIBRE OPTIC GUIDE: PER BOX OF 10				
	ITEM No.	DESCRIPTION	Unit Price Year 1 (incl. VAT)	Unit Price Year 2 (incl. VAT)	Unit Price Year 3 (incl. VAT)	Sub-Total Price (incl. VAT)
	30 301 74	Oral fibre-optic guide – Berman type 40 mm				
	30 301 75	Oral fibre-optic guide – Berman type 50 mm				
	30 301 76	Oral fibre-optic guide – Berman type 60 mm				
	30 301 77	Oral fibre-optic guide – Berman type 70 mm				
	30 301 78	Oral fibre-optic guide – Berman type 80 mm				
	30 301 79	Oral fibre-optic guide – Berman type 90 mm				
	30 301 80	Oral fibre-optic guide – Berman type 100 mm				
			Total price (incl. of taxes) To be used for evaluation			

AMOUNT IN WORDS.....

**NB: The Sub Total price is the unit price (as per specification packaging) for year 1 + year 2 + Year 3****The annual unit price will be the applicable (contractual) price per year per item.****Bidders must bid as per the price page failing which they will be disqualified.**

Required by:

KZN DEPARTMENT OF HEALTH

-At:

VARIOUS INSTITUTIONS

Country of origin

.....

Brand .....

Delivery period (on order) .....

**Failure to comply with the above shall invalidate the offer received.**

Note: All delivery costs must be included in the bid price, for delivery at prescribed destination.

.....  
(Signature of Bidder)

.....  
Date

.....  
(Signature of Witness)

.....  
Date

**SECTION N: SPECIFICATION FOR ZNB6685/2/2021-H: THE SUPPLY AND DELIVERY OF RESPIRATORY AIDS: SUPRAGLOTTIC AIRWAYS and AIRWAY RESCUE DEVICES FOR VARIOUS INSTITUTIONS FOR A PERIOD OF 3 YEARS**

**LIST OF ITEMS**

NO.	CATEGORY	CATALOGUE NUMBER	PACKAGING	DESCRIPTION
1.	<b>LMA DEVICE - Reusable</b>	30 301 01	Per Unit	Laryngeal mask airway device - reusable <b>Size: 1</b>
2.		30 301 02	Per Unit	Laryngeal mask airway device - reusable <b>Size: 1.5</b>
3.		30 301 05	Per Unit	Laryngeal mask airway device - reusable <b>Size: 3</b>
4.	<b>LMA DEVICE WITH GASTRIC DRAIN TUBE - Non-inflatable</b>	30 301 19	Per Box of 10	Laryngeal mask airway device with gastric drain tube. Non inflatable - single use <b>Size: 2.5</b>
5.	<b>DOUBLE LUMEN BLIND INSERTION AIRWAY DEVICES</b>	DLBI 1	Per Unit	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Small</b>
6.		DLBI 2	Per Unit	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Medium</b>
7.		DLBI 3	Per Unit	Blind insertion airway device – Oesophageal-tracheal double lumen airway - <b>Large</b>
8.	<b>EMERGENCY CRICOTHYROTOMY</b>	30 301 45	Per Unit	Emergency Cricothyrotomy Kit – with <b>Varess</b> needle
9.	<b>AIRWAY EXCHANGE CATHETER</b>	30 301 47	Per Unit	Airway Exchange Catheter. <b>Infant 8 Fr</b>
10.		30 301 50	Per Unit	Airway Exchange Catheter. <b>Adult 11 Fr</b> Extra length and extra firm
11.		30 301 51	Per Unit	Airway Exchange Catheter. <b>Adult 14 Fr</b> Extra length and extra firm
12.		AEC_SE	Per Unit	Airway Exchange Catheter with guidewire – staged extubation kit.
13.	<b>INTUBATING INTRODUCER</b>	30 301 52	Per Unit	Intubation Catheter <b>19 Fr</b> for use with a flexible fibre-optic scope
14.		30 301 53	Per Unit	Intubating introducer –single use <b>14 Fr</b>
15.		30 301 54	Per Unit	Intubating introducer –single use <b>8 Fr</b>
16.	<b>INTUBATION STYLET</b>	30 301 62	Per Box of 10	Intubation Stylet, <b>Neonatal</b>
17.	<b>RETROGRADE</b>	30 301 66	Per Unit	Retrograde Intubation Kit. <b>6 Fr</b>

18.	INTUBATION SET	30 301 67	Per Unit	Retrograde Intubation Kit.11 Fr
19.		30 301 68	Per Unit	Retrograde Intubation Kit.14Fr
20.	ORAL FIBRE OPTIC GUIDE	30 301 74	Per Box of 10	Oral fibre-optic guide – Berman type 40 mm
21.		30 301 75	Per Box of 10	Oral fibre-optic guide – Berman type 50 mm
22.		30 301 76	Per Box of 10	Oral fibre-optic guide – Berman type 60 mm
23.		30 301 77	Per Box of 10	Oral fibre-optic guide – Berman type 70 mm
24.		30 301 78	Per Box of 10	Oral fibre-optic guide – Berman type 80 mm
25.		30 301 79	Per Box of 10	Oral fibre-optic guide – Berman type 90 mm
26.		30 301 80	Per Box of 10	Oral fibre-optic guide – Berman type 100 mm



Specifications for **LMA DEVICE - Reusable**

**LARYNGEAL MASK AIRWAY DEVICE Reusable – COLLECTIVE REQUIREMENTS**

For use proximal to the vocal chords

Must have:

A soft **silicone cuff** with a one-way valve and bulb for inflation and deflation of pharyngeal cuff

- The cuff must be suitable for positive pressure ventilation up to 20 cm/H2O

**Aperture bars** to prevent blockage of the airway by the epiglottis

A kink resistant **airway tube** with a smooth transition to the posterior aspect of the boot and a 15 mm connector

Clear **markings** that do not rub off to indicate

- Trade name
- Size
- Recommended patient weight
- Guideline on posterior aspect to assist with orientation

Must be manufactured from medical grade silicone that is latex free

Must be autoclavable – at least 40 times

Must comply with **ISO 11712** or **equivalent and/or** SAPHRA Certification

The following must be noted on the packaging:

- Trade name of product
- Size and specification
- Manufacturing site
- CE number
- Lot number
- Expiry date

Per unit

ITEM:	DESCRIPTION:
30 301 01	<p>Laryngeal mask airway device - <b>Reusable Size: 1</b></p> <p><b>Purpose:</b> Supra-glottic airway control in a spontaneously breathing or ventilated patient. Can be used as a rescue aid after the failure of either mask ventilation, tracheal intubation or both</p> <p>Size: <b>1</b> Recommended weight: <b>up to 5 kg</b> To allow for ETT: <b>3.5</b> uncuffed</p> <p>See <b>LARYNGEAL MASK AIRWAY DEVICE Reusable – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 02	<p>Laryngeal mask airway device - <b>Reusable Size: 1.5</b></p> <p><b>Purpose:</b> Supra-glottic airway control in a spontaneously breathing or ventilated patient. Can be used as a rescue aid after the failure of either mask ventilation, tracheal intubation or both</p> <p>Size: <b>1.5</b> Recommended weight: <b>5 – 10 kg</b> To allow for ETT: <b>4</b> uncuffed</p> <p>See <b>LARYNGEAL MASK AIRWAY DEVICE Reusable – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 05	<p>Laryngeal mask airway device airway device - <b>Reusable Size: 3</b></p> <p><b>Purpose:</b> Supra-glottic airway control in a spontaneously breathing or ventilated patient. Can be used as a rescue aid after the failure of either mask ventilation, tracheal intubation or both</p> <p>Size: <b>3</b> Recommended weight: <b>30 – 50 kg</b> To allow for ETT: <b>6</b> cuffed</p> <p>See <b>LARYNGEAL MASK AIRWAY DEVICE Reusable</b> – COLLECTIVE REQUIREMENTS</p>

#### Specifications for LMA DEVICE WITH GASTRIC DRAIN TUBE - Non inflatable

LARYNGEAL MASK AIRWAY DEVICE Non inflatable – COLLECTIVE REQUIREMENTS	
<p>For use proximal to the vocal chords</p> <p>Must have:</p> <p>A preformed soft gel-like <b>non- inflatable cuff</b> allowing easy and atraumatic insertion</p> <p>An <b>integral bite block</b> incorporated into the airway tube</p> <p>Must have a buccal cavity stabiliser to aid with insertion and prevent rotation.</p> <p><b>Epiglottic rest</b> to prevent epiglottic obstruction</p> <p>Must be suitable for positive pressure ventilation up to 20 cm/H2O</p> <p>Clear <b>markings</b> that do not rub off to indicate</p> <ul style="list-style-type: none"> <li>- Trade name</li> <li>- Size</li> <li>- Recommended patient weight</li> </ul> <p>Manufactured from medical grade thermoplastic elastomer that is latex free</p> <p>Must be sterile and individually packed in peel pouch that is easy to open</p> <p>For single use only</p> <p>Must comply with <b>ISO 11712</b> or <b>equivalent</b> and/or SAPHRA Certification</p> <p>The following must be noted on the packaging:</p> <p>Trade name of product</p> <p>Size and specification</p> <p>Method of sterilization</p> <p>Manufacturing site</p> <p>CE number</p> <p>Lot number</p> <p>Expiry date</p> <p>Per Box of <b>10</b></p>	

ITEM:	DESCRIPTION:
30 301 19	<p>Laryngeal mask airway device airway device with gastric drain tube. <b>Non-inflatable - single use Size: 2.5</b></p> <p><b>Purpose:</b> Supra-glottic airway control in a spontaneously breathing or ventilated patient. Can be used as a rescue aid after the failure of either mask ventilation, tracheal intubation or both</p> <p>Size: <b>2.5</b> Recommended weight: <b>25 - 35 kg</b></p> <p>See <b>LARYNGEAL MASK AIRWAY DEVICE Non-inflatable</b> – COLLECTIVE REQUIREMENTS</p>

## Specifications for DOUBLE LUMEN BLIND INSERTION AIRWAY DEVICES

DOUBLE LUMEN BLIND INSERTION DEVICE – COLLECTIVE REQUIREMENTS	
<p>Consists of</p> <p>A <b>cuffed, double-lumen tube</b> with a soft non-traumatic tip.</p> <p>A <b>high volume proximal cuff</b> for oro and nasopharyngeal occlusion. The tube must be held firmly in place by the proximal cuff and it must prevent the escape of gases through the nose or mouth</p> <p>A <b>low volume distal cuff</b> for oesophageal occlusion</p> <p><b>Colour coded pilot balloons</b> and <b>2 syringes</b> prefilled with the correct amount of air for the inflation and deflation of the proximal and distal cuffs</p> <p><b>Bilateral ventilation ports</b> between the two cuffs</p> <p>The tube must allow for rapid intubation of difficult airways and should be a viable airway whether it is placed in the trachea or in the oesophagus</p> <p>It should also allow for the use of a fibre-optic device or tube exchanger into the trachea, and for a suction catheter into the oesophagus</p> <p>Manufactured from medical grade plastic</p> <p>All components must be latex free, sterile and individually packed in peel pouch that is easy to open</p> <p>For single use only</p> <p>Must comply with <b>ISO 11712</b> or <b>equivalent</b> or <b>SAPHRA Certification</b></p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>• Trade name</li> <li>• Size and specification</li> <li>• Method of sterilization</li> <li>• Manufacturing site</li> <li>• CE number</li> <li>• Lot number</li> <li>• Expiry date</li> </ul> <p>Per unit</p>	

ITEM:	DESCRIPTION:
DLBI 1	<p>Blind insertion airway device – <b>Oesophageal-tracheal double lumen airway - Small</b></p> <p><b>Purpose:</b> Dual lumen tube designed for difficult or emergency airway intubation and ventilation without the use of a laryngoscope</p> <p>Size: <b>28 Fr</b> <b>Small Patients: 90 - 130 cm tall</b></p> <p>See <b>DOUBLE LUMEN BLIND INSERTION DEVICE – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
DLBI 2	<p>Blind insertion airway device – <b>Oesophageal-tracheal double lumen airway - Medium</b></p> <p><b>Purpose:</b> Dual lumen tube designed for difficult or emergency airway intubation and ventilation without the use of a laryngoscope</p> <p>Size: <b>37 Fr Medium</b> Patients: <b>120 - 180 cm tall</b></p> <p>See <b>DOUBLE LUMEN BLIND INSERTION DEVICE – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
DLBI 3	<p>Blind insertion airway device – <b>Oesophageal-tracheal double lumen airway - Large</b></p> <p><b>Purpose:</b> Dual lumen tube designed for difficult or emergency airway intubation and ventilation without the use of a laryngoscope</p> <p>Size: <b>41 Fr Large</b> Patients: <b>&gt; 130 cm tall</b></p> <p>See <b>DOUBLE LUMEN BLIND INSERTION DEVICE – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 45	<p>Emergency Cricothyrotomy Kit – with <b>Varess</b> needle</p> <p><b>Purpose:</b> Used for emergency airway access when conventional endotracheal intubation or ventilation can't be performed</p> <p>Pack consists of:</p> <p>Pre-assembled tube on a varess needle.</p> <p>Varess needle to confirm entry into the trachea and indicate any subsequent contact with the posterior tracheal wall.</p> <p>A <b>6 mm</b> cricothyroidotomy tube - cuffed, radio-opaque with a 15 mm connector</p> <p><b>10 ml syringe, Scalpel Size 15</b>, Lubricating gel, Nylon sutures, HME filter</p> <p>Manufactured from medical grade metal and plastic</p> <p>Latex free</p> <p>Sterilised with ethylene oxide, single use</p> <p>Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Method of sterilization</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul> <p>Per unit</p>

## Specifications for **AIRWAY EXCHANGE CATHETER**

### **WAY EXCHANGE CATHETER – COLLECTIVE REQUIREMENTS**

Consists of a catheter manufactured from medical grade plastic

- Must be radio-opaque
- Must have a lumen for constant oxygenation of the patient, with distal side ports to ensure adequate airflow
- Must be supplied with either a 15 mm connector or a luer lock connector for attachment of a ventilatory device.
- Must have a blunt tip for atraumatic intubation, centimetre markings for accurate tube placement.

All components must be latex free, sterile and individually packed in peel pouch that is easy to open

For single use only

The following must be noted on the packaging:

- Trade name
- Size and specification
- Method of sterilization
- Manufacturing site
- CE number
- Lot number
- Expiry date

Per unit

ITEM:	DESCRIPTION:
30 301 47	<p>Airway Exchange Catheter. <b>Infant 8 Fr</b></p> <p><b>Purpose:</b> For simple, atraumatic endotracheal tube exchange whilst providing oxygenation</p> <p>Size: <b>8 Fr</b> ID: <b>1.6 mm</b> OD: <b>2.6 mm</b> To fit endotracheal tubes <math>\geq 3.0</math> mm</p> <p>Length: <b>450 mm</b></p> <p>See <b>AIRWAY EXCHANGE CATHETER – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 50	<p>Airway Exchange Catheter. <b>Adult – Extra length and extra firm 11 Fr</b></p> <p><b>Purpose:</b> For simple, atraumatic endotracheal tube exchange for double lumen tubes, whilst providing oxygenation.</p> <p>Size: <b>11 Fr</b> ID: <b>2.3 mm</b> OD: <b>&lt; 4 mm</b></p> <p>Length: <b>1000 mm</b></p> <p>See <b>AIRWAY EXCHANGE CATHETER – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 51	<p>Airway Exchange Catheter. <b>Adult – Extra length and extra firm 14 Fr</b></p> <p><b>Purpose:</b> For simple, atraumatic endotracheal tube exchange for double lumen tubes, whilst providing oxygenation.</p> <p>Size: <b>14 Fr</b> Size: <b>11 Fr</b> ID: <b>3.0 mm</b> OD: <b>&lt; 4 mm</b> Length: <b>1000 mm</b></p> <p>See <b>AIRWAY EXCHANGE CATHETER – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
AEC_SE	<p>Airway Exchange Catheter with guidewire – staged extubation kit.</p> <p><b>Purpose:</b> For extubation of ICU patients with cardiorespiratory dysfunction and potentially difficult reintubation that may be required up to 12 hours post-extubation.</p> <p>Catheter: Size: <b>14 Fr</b> ID: <b>2.3 mm</b> OD: <b>&lt; 4 mm</b> Length: 8-900mm Guidewire: Diameter: 0.8-1mm Length: 145cm</p> <p>See <b>AIRWAY EXCHANGE CATHETER – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 52	<p>Intubation Catheter – for use with a flexible <b>fibre-optic scope</b></p> <p><b>Purpose:</b> Used for fibre-optic guided placement of endotracheal or nasotracheal tubes. Also used for the exchange of laryngeal mask airway and endotracheal tubes using a fibre-optic bronchoscope.</p> <p>Size: <b>19 Fr</b> ID: <b>4.7 mm</b> OD: <b>&lt; 5 mm</b> To fit endotracheal tubes <b>7.0 mm</b> Length: <b>560 mm</b></p> <p>Consists of a catheter manufactured from medical grade plastic</p> <ul style="list-style-type: none"> <li>- Must have a lumen for constant oxygenation of the patient, with distal side ports to ensure adequate airflow</li> <li>- Must be supplied with either a 15 mm connector or a luer lock connector for attachment of a ventilatory device.</li> <li>- Must have a blunt tip for atraumatic intubation, centimetre markings for accurate tube placement.</li> </ul> <p>All components must be latex free, clinically clean and individually packed in peel pouch that is easy to open For single use only</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Method of sterilization</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul>

	Per unit
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#### INTUBATION INTRODUCER Single Use – COLLECTIVE REQUIREMENTS

Consists of a **hollow inner tube** manufactured from medical grade plastic that is flexible but kink resistant

- Must have a removable malleable **stylet** which acts as a stiffening cannula.
- Must have a **ventilatory adaptor** - to allow oxygenation via either 15 mm port or jet ventilator
- Must have a curved, blunt atraumatic tip to facilitate glottic introduction and give feedback of tracheal placement
- Tip must have **side ports**
- Must be graduated in cm marking,

All components must be latex free, clinically clean and individually packed in peel pouch that is easy to open  
For single use only

The following must be noted on the packaging:

- Trade name
- Size and specification
- Manufacturing site
- CE number
- Lot number
- Expiry date

Per unit

ITEM:	DESCRIPTION:
30 301 53	<p>Intubating introducer – single use <b>14 Fr</b></p> <p><b>Purpose:</b> To facilitate endotracheal intubation in patients where visualization of the glottis is inadequate whilst allowing oxygenation</p> <p>Size: <b>14 Fr</b> To fit endotracheal tubes <math>\geq 6.0</math> mm Length: <b>700 mm</b></p> <p>See <b>INTUBATION INTRODUCER Single Use – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 54	<p>Intubating introducer – single use <b>8 Fr</b></p> <p><b>Purpose:</b> To facilitate endotracheal intubation in patients where visualization of the glottis is inadequate whilst allowing oxygenation</p> <p>Size: <b>8 Fr</b> ID: <b>1.6 mm</b> To fit endotracheal tubes <math>&gt; 3</math> mm <math>&lt; 6</math> mm Length: <b>350 mm</b></p> <p>See <b>INTUBATION INTRODUCER Single Use – COLLECTIVE REQUIREMENTS</b></p>

**INTUBATION STYLET – COLLECTIVE REQUIREMENTS**

Consists of a malleable aluminium core coated with high density polyurethane/ Ivory PVC / medical grade plastic

- Must have a soft atraumatic tip
- Must have a U-bend proximally or other mechanism at proximal end to limit depth

All components must be latex free, clinically clean and individually packed in peel pouch that is easy to open  
For single use only

The following must be noted on the packaging:

- Trade name
- Size and specification
- Manufacturing site
- CE number
- Lot number
- Expiry date

Per Box of 10

ITEM:	DESCRIPTION:
30 301 62	<p>Intubation Stylet, <b>Neonatal</b></p> <p><b>Purpose:</b> To pre-form an endotracheal tube to a desired shape to aid intubation</p> <p>Size: <b>Neonatal</b> OD: <b>2.0 – 2.2 mm</b> To fit endotracheal tubes <b>2.5 – 4.0 mm</b> Total Length: <b>225 mm</b></p> <p>See <b>INTUBATION STYLET – COLLECTIVE REQUIREMENTS</b></p>

**RETROGRADE INTUBATION KIT – COLLECTIVE REQUIREMENTS**

Consists of a

Introducer needle or Catheter introducer needle: **18 G**

**Wire guide:** Size: **0.038 "** ; Length **110 cm**

Made from stainless steel with two positioning marks 20 cm from each end; atraumatic tips; one end straight and the other a flexible J tip. Plastic tapered introducing guide.

**Radio-opaque catheter:**

With distal side ports and a 15 mm connector for ventilatory devices

**Syringe** with luer slip

Seldinger technique access to the airway via the cricothyroid membrane. Retrograde (cephalad) advancement of wire towards the oral/ nasal cavities. Antegrade advancement of a hollow catheter with central & peripheral side ports for optimal ventilation.

Manufactured from medical grade plastic and stainless steel

Latex free

Single use, Sterile

Individually packed in peel pouch that is easy to open



The following must be noted on the packaging:

- Trade name
- Size and specification
- Method of sterilization
- Manufacturing site
- CE number
- Lot number
- Expiry date

Per unit

ITEM:	DESCRIPTION:
30 301 66	<p>Retrograde Intubation Kit. <b>6 Fr</b></p> <p><b>Purpose:</b> Used to assist in the placement of an endotracheal tube using retrograde techniques during a difficult airway access procedure.</p> <p>Catheter size: <b>6 Fr</b> for ETT with ID <b>2.5 mm</b></p> <p>Length: <b>50 cm</b></p> <p>See <b>RETROGRADE INTUBATION KIT – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 67	<p>Retrograde Intubation Kit. <b>11 Fr</b></p> <p><b>Purpose:</b> Used to assist in the placement of an endotracheal tube using retrograde techniques during a difficult airway access procedure.</p> <p>Size: <b>11 Fr</b> for ETT with ID <b>4 mm</b> or larger</p> <p>Length: <b>70 cm</b></p> <p>See <b>RETROGRADE INTUBATION KIT – COLLECTIVE REQUIREMENTS</b></p>

ITEM:	DESCRIPTION:
30 301 68	<p>Retrograde Intubation Kit. <b>14Fr</b></p> <p><b>Purpose:</b> Used to assist in the placement of an endotracheal tube using retrograde techniques during a difficult airway access procedure.</p> <p>Size: <b>14 Fr</b> for ETT with ID <b>5 mm</b> or larger</p> <p>Length: <b>70 cm</b></p> <p>See <b>RETROGRADE INTUBATION KIT – COLLECTIVE REQUIREMENTS</b></p>

### Specifications for ORAL FIBRE OPTIC GUIDE

ITEM:	DESCRIPTION:
30 301 74	<p>Oral fibre-optic guide – <b>Berman type 40 mm</b></p> <p><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p>Size: <b>40 mm</b></p>

	<p>Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p>Made from medical grade plastic Latex free</p> <p>Single use, clinically clean Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>• Trade name</li> <li>• Size and specification</li> <li>• Manufacturing site</li> <li>• CE number</li> <li>• Lot number</li> <li>• Expiry date</li> </ul> <p>Per Box of <b>10</b></p>
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ITEM:	DESCRIPTION:
30 301 75	<p>Oral fibre-optic guide – <b>Berman type 50 mm</b> <b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p>Size: <b>50 mm</b></p> <p>Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p>Made from medical grade plastic Latex free</p> <p>Single use, clinically clean Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>• Trade name</li> <li>• Size and specification</li> <li>• Manufacturing site</li> <li>• CE number</li> <li>• Lot number</li> <li>• Expiry date</li> </ul> <p>Per Box of <b>10</b></p>

ITEM:	DESCRIPTION:
30 301 76	<p>Oral fibre-optic guide – <b>Berman type 60 mm</b></p> <p><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p>Size: <b>60 mm</b></p> <p>Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p>Made from medical grade plastic Latex free</p> <p>Single use, clinically clean Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul> <p>Per Box of <b>10</b></p>

ITEM:	DESCRIPTION:
30 301 77	<p>Oral fibre-optic guide – <b>Berman type 70 mm</b></p> <p><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p>Size: <b>70 mm</b></p> <p>Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p>Made from medical grade plastic Latex free</p> <p>Single use, clinically clean Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul> <p>Per Box of <b>10</b></p>

ITEM:	DESCRIPTION:
30 301 78	<p data-bbox="280 197 766 230"><b>Oral fibre-optic guide – Berman type 80 mm</b></p> <p data-bbox="280 230 1347 297"><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p data-bbox="280 331 416 365">Size: 80 mm</p> <p data-bbox="280 398 1023 499">Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p data-bbox="280 533 632 600">Made from medical grade plastic Latex free</p> <p data-bbox="280 633 638 701">Single use, clinically clean Individually packed in peel pouch</p> <p data-bbox="280 734 782 768">The following must be noted on the packaging:</p> <ul data-bbox="331 768 612 1126" style="list-style-type: none"> <li data-bbox="331 768 507 801">● Trade name</li> <li data-bbox="331 835 612 869">● Size and specification</li> <li data-bbox="331 902 576 936">● Manufacturing site</li> <li data-bbox="331 969 501 1003">● CE number</li> <li data-bbox="331 1037 501 1070">● Lot number</li> <li data-bbox="331 1104 501 1137">● Expiry date</li> </ul> <p data-bbox="280 1193 435 1227">Per Box of <b>10</b></p>

ITEM:	DESCRIPTION:
30 301 79	<p>Oral fibre-optic guide – <b>Berman type 90 mm</b></p> <p><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p>Size: <b>90 mm</b></p> <p>Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p>Made from medical grade plastic Latex free</p> <p>Single use, clinically clean Individually packed in peel pouch</p> <p>The following must be noted on the packaging:</p> <ul style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul> <p>Per Box of <b>10</b></p>

ITEM:	DESCRIPTION:
30 301 80	<p data-bbox="277 197 778 230">Oral fibre-optic guide – <b>Berman type 100 mm</b></p> <p data-bbox="277 230 1347 297"><b>Purpose:</b> Modified oral airway specifically designed for oral fibre-optic intubation. Used to guide the tracheal tube and scope around the tongue and protect the scope should the patient bite down</p> <p data-bbox="277 331 432 365">Size: <b>100 mm</b></p> <p data-bbox="277 398 1023 499">Groove to allow removal of scope upon intubation. Long &amp; tubular (modified Guedel) to overcome upper airway oedema. Integral bite block</p> <p data-bbox="277 533 632 600">Made from medical grade plastic Latex free</p> <p data-bbox="277 633 636 701">Single use, clinically clean Individually packed in peel pouch</p> <p data-bbox="277 734 783 768">The following must be noted on the packaging:</p> <ul data-bbox="331 768 612 1126" style="list-style-type: none"> <li>● Trade name</li> <li>● Size and specification</li> <li>● Manufacturing site</li> <li>● CE number</li> <li>● Lot number</li> <li>● Expiry date</li> </ul> <p data-bbox="277 1193 427 1227">Per Box of <b>10</b></p>

## SECTION O: EVALUATION CRITERIA

Evaluation will be based on the following:

- Phase 1: Minimum Compulsory Requirements
- Phase 2: Technical Evaluation
- Phase 3: Price and Preference Points

### Phase 1: Minimum Compulsory Requirements

The Bidder shall complete and submit the following returnable schedules and documents:

NO.	SECTION/ SCHEDULE	COMPULSORY (YES / NO) NON- SUBMISSION WILL RENDER BIDDERS NON- RESPONSIVE	COMPULSORY (YES / NO) FOR BID EVALUATION PURPOSES	FOR OFFICIAL USE ONLY		
				YES	NO	N/A
Prospective Bidders MUST ensure that the following Sections of the bid document MUST be completed in ALL respects to qualify for the next stage of evaluation:						
1	Section A: Invitation to Bid	Yes	Yes			
2	Section B: Special Instructions	Yes	Yes			
3	Section C: Authority to Sign the Bid	Yes	Yes			
4	Section D: Declaration of Interest	Yes	Yes			
5	Section E: Declaration of Bidder's Past SCM Practices	Yes	Yes			
6	Section F: Declaration that CSD is Updated with Latest Bidder's Details	Yes	Yes			
7	Section G: Preference Points Claimed	Yes	Yes			
8	Section H: Certificate of Independent Bid Determination	Yes	Yes			
9	Section I: Record of Amendments to Bid Documents	Yes	Yes			
10	Section J: General Conditions of Contract	Yes	Yes			
11	Section K: Special Terms and Conditions	Yes	Yes			
12	Section L: Compulsory Briefing Session	No	No			
13	Section M: Pricing Schedule	Yes	Yes			
Prospective Bidders MUST provide the following as per the Mandatory Requirements:						
1	Copy of the Consortium/ Joint Venture/ Partnership agreement, if applicable	Yes If Applicable				
2	A Status Level Verification Certificate/Sworn Affidavit (For EMEs& QSEs) must be Submitted in order to qualify for Preference Points	Yes	Yes			



NO.	SECTION/ SCHEDULE	COMPULSORY (YES / NO) NON- SUBMISSION WILL RENDER BIDDERS NON- RESPONSIVE	COMPULSORY (YES / NO) FOR BID EVALUATION PURPOSES	FOR OFFICIAL USE ONLY		
				YES	NO	N/A
3	Relevant compliance certificates, applicable to each item	Yes; If applicable	Yes, If applicable			
4	Letter of undertaking if not the manufacturer of the Equipment	Yes	Yes			
5.	SAPHRA certification	Yes, If applicable	Yes, If applicable			

### Phase 2: Technical Evaluation of Bid

The prospective bidder will be required to provide a sample for evaluation purposes as required in terms of clause 2.14 of the special terms and conditions of the bid. Samples per an item must be accompanied with it relevant Certification (SANS Certificate or equivalent). Offers must comply strictly with the specification. Offers exceeding specification requirements will be deemed to comply with the specification. The quality of services/ supply must not be less than what is specified.

### Phase 3: Price and Preference Points

The evaluation of the price and preference will be done per category as per the pricing pages item and it is intended that this bid will be awarded as a multiple award bid. The total price for year 1-3 (VAT inclusive) per category shall be used for price and preference evaluation.

The following preference point systems are applicable to all bids:

- 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).

Points for this bid shall be awarded for:

- (a) Price; and
- (b) Status Level of Contributor.

The maximum points for this bid are allocated as follows:

CATEGORY	POINTS
PRICE	80 or 90
STATUS LEVEL OF CONTRIBUTOR	20 or 10
<b>Total points for Price and must not exceed</b>	<b>100</b>

Failure on the part of a bidder to submit proof of Status level of contributor together with the bid will be interpreted to mean that preference points for Status level of contribution are not claimed.

The department reserves the right to require of a bidder, either before a bid is adjudicated or at any time subsequently, to substantiate any claim in regard to preferences, in any manner required by the department.