MPUMALANGA PROVINCIAL GOVERNMENT

DEPARTMENT OF HEALTH

BID NUMBER: HEAL/203/21/MP

SUPPLY AND DELIVERY OF SURGICAL SUNDRIES FOR A PERIOD OF TWO (02) YEARS (24 CONSECUTIVE MONTHS)

ISSUED BY:

Department of Health
Private Bag X11285
Mbombela
1200

NAME OF BIDDER: .....

TOTAL BID PRICE (all inclusive) :.....
(Also in words): .....
.....

PART A
INVITATION TO BID

YOU ARE HEREBY INVITED TO BID FOR REQUIREMENTS OF THE DEPARTMENT OF HEALTH					
BID NUMBER:	HEAL/203/21/MP	CLOSING DATE:	06 DECEMBER 2021	CLOSING TIME:	12H00
DESCRIPTION	SUPPLY AND DELIVERY OF SURGICAL SUNDRIES FOR A PERIOD OF TWO (02) YEARS (24 CONSECUTIVE MONTHS)				

THE SUCCESSFUL BIDDER WILL BE REQUIRED TO FILL IN AND SIGN A WRITTEN CONTRACT FORM (SBD7).

BID RESPONSE DOCUMENTS MAY BE DEPOSITED IN THE BID BOX SITUATED AT
(STREET ADDRESS)

MBOMBELA, Riverside Government Complex, Building No 9, Government Boulevard, Mbombela, 1200, **PIET RETIEF,** No. 11 Measroch Street, Piet Retief Office, **KWAMHLANGA,** KwaMhlanga Government Complex, Department of Finance, Building No. 12, Computer Centre **EVANDER,** 10 Cornell Road (previously occupied by Evander Home Affairs Offices), Evander, 2280, **BUSHBUCKRIDGE,** Bushbuckridge Advice Centre, Department of Finance, Protea building (old Telkom building), **MIDDELBURG,** Department of Public Works, Cnr. Lillian Ngoyi and Dr Beyers Naudé Streets – Old TPA Building, Upper ground floor, Office numbers A20, 21 and 25, **MALELANE,** 24 Air Street, Malelane, **ELUKWATINI** ,Elukwatini Sub Regional offices, Office numbers A49 and A50 (opposite Elukwatini Community Hall) Stand number 12 Extension A, Elukwatini.

SUPPLIER INFORMATION

NAME OF BIDDER					
POSTAL ADDRESS					
STREET ADDRESS					
TELEPHONE NUMBER	CODE		NUMBER		
CELLPHONE NUMBER					
FACSIMILE NUMBER	CODE		NUMBER		
E-MAIL ADDRESS					
VAT REGISTRATION NUMBER					
	TCS PIN:		**OR** CSD No:		
B-BBEE STATUS LEVEL VERIFICATION CERTIFICATE [TICK APPLICABLE BOX]	☐ Yes ☐ No		B-BBEE STATUS LEVEL SWORN AFFIDAVIT	☐ Yes ☐ No	
IF YES, WHO WAS THE CERTIFICATE ISSUED BY?					

AN ACCOUNTING OFFICER AS CONTEMPLATED IN THE CLOSE CORPORATION ACT (CCA) AND NAME THE APPLICABLE IN THE TICK BOX		
	☐	AN ACCOUNTING OFFICER AS CONTEMPLATED IN THE CLOSE CORPORATION ACT (CCA)
	☐	A VERIFICATION AGENCY ACCREDITED BY THE SOUTH AFRICAN ACCREDITATION SYSTEM (SANAS)
	☐	A REGISTERED AUDITOR
		NAME:

[A B-BBEE STATUS LEVEL VERIFICATION CERTIFICATE/SWORN AFFIDAVIT(FOR EMEs& QSEs) MUST BE SUBMITTED IN ORDER TO QUALIFY FOR PREFERENCE POINTS FOR B-BBEE]

ARE YOU THE ACCREDITED REPRESENTATIVE **IN SOUTH AFRICA FOR THE GOODS /SERVICES /WORKS OFFERED?**	☐Yes ☐No [IF YES ENCLOSE PROOF]	ARE YOU A FOREIGN BASED SUPPLIER FOR **THE GOODS /SERVICES /WORKS OFFERED?**	☐Yes ☐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.)			

BIDDING PROCEDURE ENQUIRIES MAY BE DIRECTED TO:		**TECHNICAL INFORMATION MAY BE DIRECTED TO:**	
DEPARTMENT/ PUBLIC ENTITY	HEALTH	CONTACT PERSON	Ms BC THELA
CONTACT PERSON	Mr VS MANZINI	TELEPHONE NUMBER	013 283 9000
TELEPHONE NUMBER	013 766 3513	FACSIMILE NUMBER	
CELL. NUMBER			
FACSIMILE NUMBER		E-MAIL ADDRESS	BabalwaT@mpuhealth.gov.za
E-MAIL ADDRESS	Vincentma@mpuhealth.gov.za		

PART B
TERMS AND CONDITIONS FOR BIDDING

1. BID SUBMISSION:

1.1. BIDS MUST BE DELIVERED BY THE STIPULATED TIME TO THE CORRECT ADDRESS. LATE BIDS WILL NOT BE ACCEPTED FOR CONSIDERATION.

1.2. ALL BIDS MUST BE SUBMITTED ON THE OFFICIAL FORMS PROVIDED–(NOT TO BE RE-TYPED) OR ONLINE

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). B-BBEE CERTIFICATE OR SWORN AFFIDAVIT FOR B-BBEE MUST BE SUBMITTED TO BIDDING INSTITUTION.

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. B-BBEE CERTIFICATE OR SWORN AFFIDAVIT FOR B-BBEE MUST BE SUBMITTED TO BIDDING INSTITUTION.

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.

1.6. OFFER TO BE VALID FOR 90 DAYS FROM THE CLOSING DATE OF BID

2. TAX COMPLIANCE REQUIREMENTS

2.1 BIDDERS MUST ENSURE COMPLIANCE WITH THEIR TAX OBLIGATIONS.

2.2 BIDDERS ARE REQUIRED TO SUBMIT THEIR UNIQUE PERSONAL IDENTIFICATION NUMBER (PIN) ISSUED BY SARS TO ENABLE THE ORGAN OF STATE TO VIEW THE TAXPAYER'S PROFILE AND TAX STATUS.

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.

2.4 BIDDERS MAY ALSO SUBMIT A PRINTED TCS TOGETHER WITH THE BID.

2.5 IN BIDS WHERE CONSORTIA / JOINT VENTURES / SUB-CONTRACTORS ARE INVOLVED, EACH PARTY MUST SUBMIT A SEPARATE PROOF OF TCS / PIN / CSD NUMBER.

2.6 WHERE NO TCS IS AVAILABLE BUT THE BIDDER IS REGISTERED ON THE CENTRAL SUPPLIER DATABASE (CSD), A CSD NUMBER MUST BE PROVIDED.

3. QUESTIONNAIRE TO BIDDING FOREIGN SUPPLIERS

3.1.	IS THE BIDDER A RESIDENT OF THE REPUBLIC OF SOUTH AFRICA (RSA)?	☐ YES ☐ NO
3.2.	DOES THE BIDDER HAVE A BRANCH IN THE RSA?	☐ YES ☐ NO
3.3.	DOES THE BIDDER HAVE A PERMANENT ESTABLISHMENT IN THE RSA?	☐ YES ☐ NO
3.4.	DOES THE BIDDER HAVE ANY SOURCE OF INCOME IN THE RSA?	☐ YES ☐ NO

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 REGISTER AS PER 2.3 ABOVE.

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

TENDER SPECIFICATION

MPUMALANGA DEPARTMENT OF HEALTH

SPECIFICATION FOR THE APPOINTMENT OF SERVICE PROVIDER (S) FOR SUPPLY AND DELIVERY OF SUGICAL SUNDRIES FOR THE PERIOD OF TWO (02) YEARS (24 CONSECUTIVE MONTHS)

SUGICAL SUNDRIES

SECTION A: SPECIAL CONDITIONS OF CONTRACT

This bid and all contracts emanating there from will be subject to the General Conditions of Contract issued in accordance with Treasury Regulation 16A published in terms of the Public Finance Management Act, 1999 (Act 1 of 1999). The Special Conditions of Contract are supplementary to that of the General Conditions of Contract. Where, however, the Special Conditions of Contract are in conflict with the General Conditions of Contract, the Special Conditions of Contract prevail.

SPECIAL CONDITIONS:

I. BIDDER CAPACITY AND PERFORMANCE MANAGEMENT

1. The bidder/s shall indemnify the department herewith from any claim from a third party and all cost or legal expenses in regard to such a claim for loss or damage resulting from the death, injuries or ailment of any person, or the damage of property of the bidder(s) or any other person that may result from or be related to the execution of this contract.
2. The performance of bidders will be evaluated throughout the contract period. If performance is not satisfactory and the bidder is unable to remedy any of the breaches within the stipulated period, the Mpumalanga Department of Health will terminate the contract.
3. If it is shown that errors or shortcomings exist with the bidder, the bidder shall be notified in writing and shall be required to perform corrective services to remedy such errors at no cost to the Mpumalanga Department of Health. Such errors or shortcomings shall be corrected/redressed within five (5) working days.
4. The Mpumalanga Department of Health reserves the right to request further information from the bidder/s anytime;
5. The Mpumalanga Department of Health reserves the right to verify information and documentation of the bidder/s.

II. PRICING

1. Tender prices are to INCLUDE DELIVERY COST TO THE RELEVANT INSTITUTIONS AND/OR ANY OTHER NOMINATED DESTINATION. The delivery price of items must include all delivery costs such as packing, off-loading, products handling.
2. Value Added Tax (VAT) is to be INCLUDED in the tender price.
3. Prices must be fixed for each of the two years (duration of the contract). Note that NO requests for price escalations will be considered.
4. In the event that there are vast differences / discrepancies in the prices of the recommended bidders, the Mpumalanga Department of Health will conduct market analysis therefrom a mark-up not exceeding 25% will be added to the averaged prices to cover profit, transportation and all other related costs.
5. The quantities reflected in the bid forms are **estimated quantities** for the contract period and no guarantee is given or implied as to the actual quantity which will be procured during the contract period

III. APPOINTMENT OF BIDDER/S

1. The Mpumalanga Department of Health reserves the right to make sure that the bidder(s) have at their disposal the necessary infrastructure to execute the contract to the satisfaction of the Mpumalanga Department of Health prior to the awarding of the contract;
2. The Mpumalanga Department of Health reserves the right to inspect the operation and verify supporting documentations during the evaluation phase of the Bid;
3. The Mpumalanga Department of Health reserves the right to conduct an inspection of the business premises (offices, vehicles and warehouse) before and after awarding the bid.
4. The Mpumalanga Department of Health reserves the right to appoint or not to appoint service provider(s).
5. The Mpumalanga Department of Health reserves the right to invite short listed bidders to make presentations
6. The Mpumalanga Department of Health reserves the right to appoint more than one service provider
7. The Mpumalanga Department of Health reserves the right to not to award, to award the bid in full or in part including split for product

8. All directors of companies will be subjected to verification if they are Government employees.

IV. PACKAGING

1. All deliveries made against this contract, in all modes of transport, are to be packed in suitable containers, which will be acceptable for further dispatch
2. All products must be labelled with the full product description, Batch/Lot number; expiry date and quantity in a pack appearing clearly on the packing. Packing and cartons must be clearly labelled. The information must be clearly printed on all inner and outer packaging in English
3. Products must have the same batch/lot number on the outer box as on the inner box.
4. For products that require sterility, packaging must state if the product supplied is sterile or mon-sterile.
5. The material and design of a peel-apart (unit) pack shall be such as to ensure easy opening with the fingers. Once opened, the package must be such that cannot be easily re-sealed, and it shall be obvious that the unit package has been opened. A peel-apart pack shall tear cleanly without the formation of loose fluff or loose fibres.

V. ORDERS DELIVERIES AND PAYMENTS

1. Official orders will be placed during the course of the contract period, as and when required.
2. Delivery after receipt of an official order shall be strictly within 14 days
3. The consignment must be accompanied by an original invoice and delivery note with a clear order number being fulfilled.
4. Payment will be effected only after receipt of a detailed original invoice and a signed delivery note to the nominated destination has been received.
5. The shelf life should not be less than eighteen months. If the shelf life is less than eighteen months, the product may only be delivered if there is a written agreement between both parties and the protection letter by the bidder to unconditionally uplift the expired/short dated stock to replace or credit within the period of 30 days after expiry

6. On delivery, consignments must be grouped according to product specification and batches. No mixing of products or batches.
7. All deliveries must be received in suitable packaging.
8. Consignments must be delivered in a closed, water tight vehicle which will be able to maintain temperature during transit below 25°C.
9. Bulk consignments must be delivered on pallets.
10. All Cardio Vascular Catheter **(CVC)** products must comply with ISO: 11070:2014.
11. Compliance certificates must accompany the bid document wherever ISO compliance is required

VI. TERMINATION OF CONTRACT

1. The Mpumalanga Department of Health reserves the right to suspend / terminate the contract if the successful bidder/s does not comply with any stipulations contained in the contract.
2. The Mpumalanga Department of Health reserves the right to reject services / goods / works that does not meet the required standard and to engage a different service provider to complete the work. The bidder shall be served with a seven (07) days written notice for termination of contract in case of unsatisfactory performance.
3. Bidders committing similar errors for three (03) times in a period of twelve consecutive months shall have their contracts terminated.

PRODUCT LIST AND SPECIFICATIONS: Items must comply with item specifications as stated in the bid document. Failure to comply with this condition will invalidate your bid for the items concerned.

ITEM NO.	MP DEPOT STOCK CODE	ITEM DESCRIPTION	ITEM SPECIFICATION	PACK SIZE	ESTIMATED QUANTITY FOR THE PERIOD	UNIT PRICE YEAR 1	UNIT PRICE YEAR 2
A-1	2754	Bag Drainage Urine Meter	Bag drainage and urine meter, 2000ml drainage bag, pre-connected to 500ml urine meter, both drainage bag and urine meter must be graduated hang meter with universal double hook to hang on bed frame, urine connecting tube 1200mm, sampling urine from the container, empty container into bag, drain bag, sampling urine from catheter connector to insert needle-free luer slip syringe into the port, caution: conventional needles and blunt cannulas are not recommended and may damage the needle-free port, single use, do not use if package is open or damaged, it must be individually wrapped. Each	Each	1 898		
A-2	1465	Bag Urine Drainage S3 Double End S3	Bag urine drainage Plasticised medical grade non-toxic PVC The finish of which prevents adhesion between front and back of bag Special use - drainable Double ended with bottom drain. Graduated volume of 2000ml with outlet tap and with non-return valve at upper end. Internal diameter of inlet and outlet	Each	229 796		

			tube 5mm-8mm. Sterile, individually packed. Each				
A-3	1464	Bag Urine Drainage; S4 Double End S4; Type 2	Bag urine drainage. Plasticised medical grade non-toxic PVC The finish of which prevents adhesion between front and back of bag. Special use - drainable Type 2 - Double ended with bottom drain. Graduated volume of 2000ml with urine sampling sleeve with outlet tap and with non-return valve at upper end Internal diameter of inlet and outlet tube 5mm-8mm Sterile, individually packed. Each	Each	167 760		
A-4	3218	Bag Urine Leg Bag (Long Tube Non-Sterile) 750ml	Urine Leg Bag (Long Tube Non-Sterile) capacity 750Ml graduated, Rectangular sleeve with Leg bag strap - comfortable texture surface grip straps, side fix -adjustable slips, inlet with non-reflex valve, outlet twist tap/ push-pull outlet, silicon/ siliconised bag to prevent anaphylactic reaction. Labelling must have the following information: item description, size, batch number, expiry date, manufacturing date, quantity, individually wrapped peel packed. Each	Each	4693		
A-5	4543	Catheter - Haemodialysis Chronic 15Fx19Cm	• Long Term haemodialysis catheter 15Fx19cm in sterile pack • Radiopaque polyurethane with Graduated catheter • Kink resistant	Each	1		

			• Side holes to separate and to reduce recirculation • Internal lumen 15G able to withstand blood flow of up to 450ml/min • Spring Guide 97mmx100cm • Introducer needle 18Gx6.35cm • Dilator 12-14FR • Scalpel blade size 11 • Arterial / venous extension must be colour coded and marked with lock volume • It must be supplied in a sealed sterile pack (comply with ISO standard ISO 10555-3:2013) • Label with batch number manufacture date, expiry date • none pyrogenic fluid parts • for single use only				
A-6	4542	Catheter - Haemodialysis chronic 15Fx27Cm	• Long Term haemodialysis catheter 15Fx27cm in sterile pack • Radiopaque polyurethane with Graduated catheter • Kink resistant • Side holes to separate and to reduce recirculation • Internal lumen 15G able to withstand blood flow of up to 450ml/min • Spring Guide wire 97mm x 100cm • Dilator12-14FR	Each	13		

			• Needle 18GAx 6.3mm • Scalpel blade 11 • Arterial / venous extension must be colour coded and marked with lock volume • It must be supplied in a sealed sterile pack (comply with ISO standard ISO 10555-3:2013) • Label with batch number manufacture date, expiry date, • none pyrogenic fluid parts • for single use only				
A-7	4544	Catheter – Haemodialysis Chronic 15Fx31Cm	• Long Term haemodialysis catheter 15Fx31cm in sterile pack • Radiopaque polyurethane with Graduated catheter • Kink resistant • Side holes to separate and to reduce recirculation • Internal lumen 15G able to withstand blood flow of up to 450ml/min • Spring Guide wire 97mm x100mm • Introducer needle 18Ga x 6.35mm • Dilator 12-14fr • Scalpel blade size 11 • Arterial / venous extension must be colour coded and marked with lock volume	Each	1		

			• It must be supplied in a sealed sterile pack (comply with ISO standard ISO 10555-3:2013) • Label with batch number manufacture date, expiry date, • none pyrogenic fluid parts • for single use only				
A-8	2746	Catheter - Suprapubic Drainage kit, 10Fg x 12cm	CATHETER SUPRAPUBIC, Silicone (100%)Proximal end: tapered one end with hollow spigot with luer lock connector to fit syringe and a cap, One balloon port, with non-return valve and fitting for a syringe, Distal end: open tip atraumatic, With lateral drainage eyes, distal to a 5ml integrated balloon Size: 10FG Length not less than 120mmTo comply with SANS 51616Sterile, individually peel packed, double wrapped inner pouch unsealed	Each	390		
A-9	2314	Catheter - Suprapubic Drainage kit, 10Fg x 8cm	CATHETER SUPRAPUBIC, Silicone (100%)Proximal end: tapered one end with hollow spigot with luer lock connector to fit syringe and a cap, One balloon port, with non-return valve and fitting for a syringe, Distal end: open tip atraumatic, With lateral drainage eyes, distal to a 5ml integrated balloon Size: 10FG Length not less than 80mmTo comply with SANS 51616Sterile, individually peel packed, double wrapped inner pouch unsealed	Each	536		

A-10	2727	Catheter - Suprapubic Drainage kit, 15Fg x 12cm	CATHETER SUPRAPUBIC, Silicone (100%)Proximal end: tapered one end with hollow spigot with luer lock connector to fit syringe and a cap, One balloon port, with non-return valve and fitting for a syringe, Distal end: open tip atraumatic, With lateral drainage eyes, distal to a 5ml integrated balloon Size: 15FG Length not less than 120mmTo comply with SANS 51616Sterile, individually peel packed, double wrapped inner pouch unsealed	Each	18		
A-11	4106	Catheter Haemodialysis – Chronic 15Fx23Cm	• Long Term haemodialysis catheter 15Fx23cm in a sterile pack • Radiopaque polyurethane with Graduated catheter • Kink resistant • Side holes to separate and to reduce recirculation • Internal lumen 15G able to withstand blood flow of up to 450ml/min • Spring Guide wire 97mmx100cm • Dilator 12-14FR • Scalpel blade size 11 • Arterial / venous extension must be colour coded and marked with lock volume • It must be supplied in a sealed sterile pack (comply with ISO standard ISO 10555-3:2013)	Each	265		

			• Label with batch number manufacture date, expiry date, • none pyrogenic fluid parts • for single use only				
A-12	3867	Catheter Haemodialysis - Double-Lumen (12Ga) 12fr X 16 Cm	• Radiopaque polyurethane with Graduated catheter • Kink resistant • Rotatable suture wing • Side holes to separate and to reduce recirculation • Internal lumen 12G able to withstand blood flow of up to 350ml/min • Spring Guide wire 97mmx100mm • Dilator 12-14FR • Scalpel blade size 11 • Arterial / venous extension must be colour coded and marked with lock volume • Optional curved extension for jugular placement / patient comfort • It must be supplied in a sealed sterile pack (comply with ISO standard ISO 10555-3:2013) • Label with batch number manufacture date, expiry date, • none pyrogenic fluid parts • for single use only	Each	3200		
B-1	4184	18G Green Safety cannula with Wings and Injection Port 1.3x45mm	Intravenous cannula, safety with port and wings, introducer needle, radio-opaque, Teflon/polyurethane, with	Box of 100	676		

			double-tapered tip and stylet. 3-facetted J-point bevelled needle Length: Not less than 45mm, Size: 18G colour: Green Flow rate: 105ml/min. Sterile, Single use, individually peel packed, To comply with the latest issue of SANS327, Box (100 box per)				
B-2	4185	20G Pink Safety cannula with Wings and Injection Port 1.3x45mm	Intravenous cannula, safety with port and wings, introducer needle, radio-opaque, Teflon/polyurethane, with double-tapered tip and stylet. 3-facetted J-point bevelled needle Length: Not less than 33mm, Size: 20G colour: Pink Flow rate: 61ml/min ml/min, sterile, Single use, individually peel packed, To comply with the latest issue of SANS327, Box (100 box per)	Box of 100	312		
B-3	1311	Intravenous Cannula – 18G - Green Radio-Opaque	Intravenous cannula, introducer needle, radio-opaque, Teflon/polyurethane, with double-tapered tip and stylet. 3-facetted J-point bevelled needle Length: Not less than 40mm, Size: 18G colour: Green Flow rate: 105ml/min. Sterile, Single use, individually peel packed,	Each	7189		
B-4	4799	Needle Spinal 25G Pencil Point 103mm Needle Blade 0.5mm	Needle, Spinal, quincke bevel, luer lock. The needle must be colour coded at the stylet cap and the size must be clearly stated. Must have a clear/ transparent needle hub to allow easy identification of the spinal fluid flashback. The flashback must	Each	788		

			be quick relative to the needle gauge. Must have a key/slot arrangement of the stylet and hub and must be clearly indicate the bevelled orientation. Have a stable stylet that does not shake or quiver to allow for easy re-insertion into the cannula. Must align preciously with the stylet point. Must be bevelled at 220degree, Needle size: 25G with a needle length of 90mm excluding hub, Needle and stylet must be manufactured from good quality stainless steel. Plastic hub must be manufacture from medical grade plastic and all the components must be pyrogenic and latex free, sterile and individually peel packed, Pack size: Box(Box of 25)				
B-5	0985	Syringe – Hypodermic 20ml plastic syringe with an eccentric nozzle, sterile	Syringe, three-part, plastic, latex free, non-pyrogenic and DEPH free. Type 3 (luer slip), with eccentric nozzle. Normal capacity 20ml, must have clear graduation markings that are not easily removed with spirits and allow for permanent marker markings without rubbing off. Sterile, Single use, Individually peel packed, To comply with the latest issue of SANS 1124-2 Box of 50	Each	24 472		
C-1	2513	Admin set - extension line, coiled, for pump sets, OD:1.0-2.0mmx300mm	Admin set - extension line, single lumen, coiled, for pump sets, Male/Female Luer-lock connector points. Diameter: 1.0 x 2.0mm x	Each	3822		

			300mm. Material: Polyethylene, Latex and DEHP free, Non-pyrogenic. Sterile individual peel pack for single use				
C-2	3204	Catheter - Femoral Arterial 18G x 18cm (4Fg)	A sterile set with: • **radiopaque polyurethane** Arterial Catheter 18 Gauge 18cm long 4Fr **indwelling catheter for Femoral and/or brachial arteries** • 19G introducer needle 68mm Long • Straight guide wire 46cm Long with a diameter of 0,71mm • Catheter with a distal luer lock 18cm long • 3-way extension, extension 30cm long • 5ml syringe Pyrogenic free Supplied in a sealed Sterile pack with a sterile White Drape paper covering the contents Labelled with: ✓ Batch No ✓ Manufacturing Date ✓ Expiry Date	Each	410		
C-3	3205	Catheter - Femoral Arterial 20G x 88cm (3Fg)	A sterile set with: • **radiopaque polyurethane** Arterial Catheter 20 Gauge 8cm long 3Fr **indwelling catheter for Femoral and/or brachial arteries**	Each	520		

			• 19G introducer needle 68mm Long • Straight guide wire 46cm Long with a diameter of 0,71mm • Catheter with a distal luer lock 18cm long • 3-way extension, extension 30cm long • 5ml syringe Pyrogenic free Supplied in a sealed Sterile pack with a sterile White Drape paper covering the contents Labelled with: ✓ Batch No ✓ Manufacturing Date ✓ Expiry Date				
C-4	3209	Catheter - Radial Arterial 22G x 8cm (3-4Fr)	A sterile set with: • **1x radiopaque polyurethane** Arterial Catheter 20 Gauge 8cm long 3Fr- 4Fr • 1x 20G introducer needle 38mm Long • 1x Metal Straight guide wire 20cm -30cm Long with a diameter 0.5mm • 1x Catheter with transparent hub that prevents blood spillage and infection control • 3-way extension, extension 30cm long and minimizes kinking • 5ml syringe	Each	400		

			Supplied in a sealed Sterile pack with covered contents to be kept pyrogenic free Labelled with: ✓ Batch No ✓ Manufacturing Date ✓ Expiry Date ✓ List and number of the contents described in full				
C-5	2715	Central Venous Catheter - Double Lumen Set 4 Frx5cm	• 2-Lumen calibrated central venous catheter set 4Fr x 5cm long • Both lumen must be 22G • Radiopaque Polyurethane with blue flexible tip • The set must contain a spring wire guide with J-flexible tip +- 45cm long and diameter of 0.4mm • 20G 4cm Introducer Needle • 25G Introducer Needle • Vessel Dilator 1.8mm x 7.5cm • Scalpel blade size 11 • Heparinized 5ml Syringe • The pack must be sterile • The pack must indicate the number of days the catheter must be in the body • The pack must have a batch number, manufacturing date and an expiry date	Each	481		
C-6	3432	Central Venous Catheter - Double Lumen Set 4 Frx8cm	• 2-Lumen calibrated central venous catheter set 4Fr x 8cm long	Each	1		

			• Both lumen must be 22G • Radiopaque Polyurethane with blue flexible tip • The set must contain a spring wire guide with J-flexible tip +- 45cm long and diameter of 0.4mm • 20G 4cm Introducer Needle • 25G Introducer Needle • Vessel Dilator 1.8mm x 7.5cm • Scalpel blade size 11 • Heparinized 5ml Syringe • The pack must be sterile • The pack must indicate the number of days the catheter must be in the body • The pack must have a batch number, manufacturing date and an expiry date				
C-7	2979	Central Venous catheter - Double lumen set 7fr x 16cm	• 2 lumen calibrated catheter set, 7fr 16cm long • The distal lumen must be 18 and the proximal must be 14G • Radiopaque Polyurethane with blue flexible tip • The set must contain a spring wire guide with J-flexible tip +- 45cm long and diameter of 0.4mm • 18G Introducer Needle • Vessel Dilator- 20G 7.6FR • Scalpel blade size 11 • Heparinized 5ml Syringe	Each	2181		

			• The pack must be sterile • The pack must indicate the number of days the catheter must be in the body • The pack must have a batch number, manufacturing date and an expiry date				
C-8	3107	Central Venous catheter - Triple lumen (15G, 16G and 14G) set – 8,5Fr x 20cm	• 3 lumen calibrated catheter set, 8.5fr 20cm long • The lumens size must be 15G, 16G and 14G • Radiopaque Polyurethane with blue flexible tip • The set must contain a spring wire guide with J-flexible tip +- 45cm long and diameter of 0.4mm • 18G Introducer Needle • Vessel Dilator- 20G 7.6FR • Scalpel blade size 11 • Heparinized 5ml Syringe • The pack must be sterile • The pack must indicate the number of days the catheter must be in the body • The pack must have a batch number, manufacturing date and an expiry date	Each	447		
C-9	2635	Central Venous Catheter - Triple Lumen Paediatric 5.5 Fr 13cm	• Three (3) lumen calibrated paediatric catheter set, 5.5fr 13cm long • The distal lumen must be 20G The other two lumen must be 22G	Each	182		

			• Radiopaque Polyurethane with blue flexible tip • The set must contain a spring wire guide with J-flexible tip +- 45cm long and diameter of 0.4mm • 20G Introducer Needle • Vessel Dilator- 20G 7.6FR • Scalpel blade size 11 • Heparinized 5ml Syringe • The pack must be sterile • The pack must indicate the number of days the catheter must be in the body • The pack must have a batch number, manufacturing date and an expiry date				
C-10	2696	Peripheral IV Catheter 22G, 4Cm	Peripheral IV catheter – radiopaque polyurethane 22G catheter with fixation wings on the hub and integrated extension 4 cm long • Catheter internal diameter 0.5 External diameter 0.7 • Introducer needle 22G/21G and 42mm long • Straight Metal guide wire 0.46mm diameter x 230mm long • Must allow flow rate of up to 17ml per minute • The pack must be sterile • The pack must have a batch number, manufacturing date and an expiry date	Each	2314		

			• Must be able to be used for children (arterial / venous line) and adults (VENOUS / JUGULAR LINE)				
D-1	3960	Ortho-phthalaldehyde 0.55% solution	Ortho-phthalaldehyde 0.55% solution for use as high level disinfectant Container size min of 3.8L	Each	94		
D-2	3961	Ortho-phthalaldehyde 0.55% solution Test Strips	Ortho-phthalaldehyde 0.55% solution. Test Strips to provide a rapid and easy measure of the minimum effective concentration (MEC) of 0.3% to enable optimal reuse of Ortho-phthalaldehyde 0.55% solution for high-level disinfection. Must be in series with the Ortho-phthalaldehyde 0.55% solution above	Each	5		
E	1431	Plastic Medicine Spoon 5ml Measure	Clear plastic medicine measure spoon Graduated Capacity: 2.5ml - 5ml Packaging: Each	Each	65 500		
F-1	1341	Apron Surgeons; Light Green; Autoclave 120cm Long	Apron Surgeons Autoclavable Re-usable Material: - Semi-transparent - Waterproof PVC Gauge: 6 - 8 with seams with soft, adjustable neck bands and tapes for waist Protective length: ± 120 cm Colour: Light Green Packaging: Each	Each	48 077		
F-2	4842	Gloves - purple nitrile extended cuff, powder free, non-sterile - Small	Glove, examination, non-sterile, type 2 (gloves made primarily from nitrile rubber latex, polychloroprene rubber solution, styrene-butadiene rubber emulsion or thermoplastic-elastomer solution), single use, Powder free,	Box of 100 gloves	3 000		

			Ambidextrous. SANS 11193-1, Box of 100 gloves, Size Small				
F-3	4843	Gloves - purple nitrile extended cuff, powder free, non-sterile - Medium	Glove, examination, non-sterile, type 2 (gloves made primarily from nitrile rubber latex, polychloroprene rubber solution, styrene-butadiene rubber emulsion or thermoplastic-elastomer solution), single use, Powder free, Ambidextrous. SANS 11193-1, Box of 100 gloves, Size Medium	Box of 100 gloves	8 000		
F-4	4844	Gloves - purple nitrile extended cuff, powder free, non-sterile - Large	Glove, examination, non-sterile, type 2 (gloves made primarily from nitrile rubber latex, polychloroprene rubber solution, styrene-butadiene rubber emulsion or thermoplastic-elastomer solution), single use, Powder free, Ambidextrous. SANS 11193-1, Box of 100 gloves, Size Large	Box of 100 gloves	15 000		
F-5	4845	Gloves - purple nitrile extended cuff, powder free, non-sterile – extra-large	Glove, examination, non-sterile, type 2 (gloves made primarily from nitrile rubber latex, polychloroprene rubber solution, styrene-butadiene rubber emulsion or thermoplastic-elastomer solution), single use, Powder free, Ambidextrous. SANS 11193-1, Box of 100 gloves, Size Extra-large	Box of 100 gloves	6 000		
F-6	2404	Gloves Double System Sterile Size 6.0	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 6.5 Inner	Box of 25 pairs	1170		

			glove size: 6 Size: 6 Packaging: 2 pairs, primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1				
F-7	2740	Gloves Double System Sterile Size 6.5	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 6.5 Inner glove size: 7 Size: 6.5 Packaging: 2 pairs, primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1	Box of 25 pairs	2 790		

F-8	2737	Gloves Double System Sterile Size 7.0	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 7 Inner glove size: 7.5 Size: 7 Packaging: 2 pairs primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1	Box of 25 pairs	5 333		
F-9	2594	Gloves Double System Sterile Size 7.5	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 7.5 Inner glove size: 8 Size: 7.5 Packaging: 2 pairs, primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves	Box of 25 pairs	4 633		

			reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1				
F-10	2595	Gloves Double System Sterile Size 8	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 8.5 Inner glove size: 8 Size: 8 Packaging: 2 pairs, primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1	Box of 25 pairs	780		
F-11	2405	Gloves Double System Sterile Size 8.5	Glove surgical natural rubber latex hypo-allergenic Powder and pyrogenic free Sterile Must be a double gloving system Consisting of a coloured inner glove and a standard	Box of 25 pairs	546		

			outer glove Residual antigen protein shall be less than 5µg/g Length: 275mm Outer glove size: 9 Inner glove size: 8.5 Size: 8.5 Packaging: 2 pairs, primary packaging of the inner glove must be individually wrapped in a purse style paper with left and right hand clearly marked with sleeves reversed up to the thumb, Secondary packaging must be a sterilised peel pack, supplied in a box of 25 pairs, labelling on the peel pack and box: Item description, size, quantity, batch/ lot number, manufacturing date, expiry date and sterile for single use, latex warning, ISO: 11193-1				
F-12	1898	Mask surgical, flat pleated WITH anti-fog coated and anti-reflective face shield. Tie on.	Shield, eye, protection, a full face mask must be attached to eye shield, must provide protection against blood and body fluids. Features of shield: -Superior anti-fog coating -High quality optical grade 0.07 mm film -Zero optical distortion -Lightweight -One size fits all -Disposable -No eye wear frame to obstruct vision -A strip of foam rubber at the top to prevent eye contamination -Two self-adhesive foam strips to attach to face mask -Have four tie backs for fastening to head	Each	22 581		

			-Made of high efficiency, fluid resistant surgical type. -Bacterial filtration efficiency to be greater than 99%. - Compliance certificate must be submitted. Box of 20.				
G-1	1470	Catheter - Suction Respiratory 8Fg (Control Airport) 350Mm	CATHETER, SUCTION, RESPIRATORY, medical grade non-toxic PVC smooth, flexible but sufficiently rigid to pass through endotracheal tube. Proximal end: finger control airport, tapered connector, Distal end: open atraumatic tip with Two lateral staggered eyes, Length not less than 350mm, Size: 8FG. Blue, Sterile, Individually peel Packed, To comply with the latest issue of ISO 8836 EACH	Each	91 588		
G-2	1471	Catheter Suction Respiratory 10Fg (Control Airport)350mm	CATHETER, SUCTION, RESPIRATORY, medical grade non-toxic PVC smooth, flexible but sufficiently rigid to pass through endotracheal tube. Proximal end: finger control airport, tapered connector, Distal end: open atraumatic tip with Two lateral staggered eyes, Length not less than 350mm, Size: 10FG. Black, Sterile, Individually peel Packed, To comply with the latest issue of ISO 8836 EACH	Each	112 177		
G-3	1472	Catheter Suction Respiratory 12Fg (Control Airport)500mm	CATHETER, SUCTION, RESPIRATORY, medical grade non-toxic PVC smooth, flexible but sufficiently rigid to pass through endotracheal tube. Proximal end: finger control airport, tapered	Each	29 471		

			connector, Distal end: open atraumatic tip with Two lateral staggered eyes, Length not less than 500mm, Size: 12FG. White, Sterile, Individually peel Packed, To comply with the latest issue of ISO 8836 EACH				
G-4	1473	Catheter Suction Respiratory 14Fg (Control Airport) 500mm	CATHETER, SUCTION, RESPIRATORY, medical grade non-toxic PVC smooth, flexible but sufficiently rigid to pass through endotracheal tube. Proximal end: finger control airport, tapered connector, Distal end: open atraumatic tip with Two lateral staggered eyes, Length not less than 500mm, Size: 14FG. Green, Sterile, Individually peel Packed, To comply with the latest issue of ISO 8836 EACH	Each	31 697		
G-5	1476	Catheter Thoracic With Trocar 12Fg; Paediatric	CATHETER, THORACIC DRAINAGE, WITH TROCAR, Trocar: anodized aluminium with handle Catheter: Medical grade non-toxic PVC, non-collapsible , kink- resistant Radio-opaque line, Graduated every 50mm Proximal end: tapered with connector Distal end: rigid open tip with Two lateral staggered eyes, Size 12FG – Paediatric, Length not less than 225mm Sterile, Individually peel packed EACH	Each	1214		
G-6	1534	Drainage Bottle– Chest Underwater Drainage Bottle	DRAINAGE SYSTEM, CHEST, UNDERWATER, CONTAINER ONLY Adult - Calibrated above water seal line in 50ml increments	Each	500		

			- Capacity 1500ml – 2000ml (will be regarded as series with container item 21.2) EACH				
G-7	1532	Drainage System – Chest Underwater drainage system	Drainage system, chest, underwater, set adult, tube: 1 milkable tube with: micro frosted surface for minimal friction, one end attached to the water seal rigid pipe, one end attached to a stepped connector for attachment to various sizes of intercostal drainage catheters, container: calibrated above water seal line in 500ml increments, capacity 1500ml - 2000ml, clear plastic, rigid enough to hold safe vacuum for patient, threaded neck to accommodate screw cap, screw cap: with gasket to prevent leakage, integral suction port / air vent, water seal rigid pipe connected at one end to screw cap and tubing and having its other bevelled end reaching below the water seal line indicated on the container, clamp to attach to patient clothing or bedding/ straps to carry, system to be supplied complete with adaptors and protectors, set, adult, container: sterile, individually double peel packed. Each	Each	8 020		
G-8	2741	Tube Aspirating Specimen Container 8ml	TUBE, ASPIRATING, SPECIMEN CONTAINER, With two variable level ports attached to latex suction	Each	1		

			tubing: Internal diameter: not less than 4.76mm, Wall 1.59m. Removable stopper, Capacity not less than 8ml. Sterile, individually peel packed, EACH				
H-1	1132	Bandage Elastic Adhesive Porous 25mmx1m	Bandage, Porous, Conforming, Surgical. To consist with a woven cotton cloth backed with a porous, zinc oxide filled hypoallergenic adhesive, BP, Bandage to have strong support with moderate compression Size: 25mm x not less than 2.7m unstretched 4.5m stretched individually wrapped Roll	Each	549		
H-2	1151	Dressing Absorbent surgical pad Non- Adherent Pad, 200x200mm	Pad, Surgical, Dissecting, type 1, X-ray detectable. 100 swabs, Size 10mm materials: Outer layer – The outer layer shall be gauze, the cotton wool shall be cotton wool grade 2, Construction: The cotton wool shall be packed in single layer of gauze x-ray detectable material of size 5mm x 5mm to be placed between the cotton wool and the outer layer of gauze so as to be visible from the outside. The gauze must be secured with a cotton threat of acceptable quality, The swabs shall be loosely stitched together in bundles of five by means of an uneasy to break cotton threat pack of 100 swabs	Each	1		
H-3	3236	Dressing charcoal impregnated 19X10.5 cm Absorbent Non-Adhere	Dressing wound absorbent non-adherent sterile 100% pure activated charcoal impregnated with silver	Box of 10	866		

			Must absorb odours and have broad-spectrum antimicrobial activity Size: 105mm X 190mm Packaging: box of 10 sachets (Dressing must be sterile, individually peel packed)				
H-4	2231	Dressing Hydro Capillary Absorbent Non -Adherent Sheet ±75mm x ±75mm	Dressing Wound, hydro-capillary, woven Absorbent, layered dressing to assist in the removal of exudate and dead tissue, be non-adherent to wound surface, sterile. Should have description on outer pack Sterile, individually peel packed Size Dressing: ±75mm x 75mm Each	Each	1		
H-5	4430	Dressing Hydro fibre Sheet with Silver 150mm x150mm	Primary wound dressing made from sodium Carboxymethyl cellulose and silver. Flat non-woven pad. The dressings are presented individually packed in peel pouches, Size: 150mm x150mm	Each	5785		
H-6	2318	Dressing Hydrocolloid-Sodium Carboxymethyl Cellulose, Thin, Transparent, 15 x15cm	Dressing, hydrocolloid, Transparent, Sodium Carboxymethyl cellulose, Pectin, gelatin, thin, hydrocolloid layer bonded to a polyurethane film, transparent for visual inspection, flexible dressing. MCC registration, Sterile, Individually peel packed, Size of Dressing: ±150mm x 150mm, Each	Each	75		
H-7	2563	Dressing Hydrocolloid-Sodium Carboxymethyl Cellulose, Thin, Transparent, Approx.: 5 x 20cm	Dressing, hydrocolloid, Transparent, Sodium Carboxymethyl cellulose, Pectin, gelatin, thin, hydrocolloid layer bonded to a polyurethane film,	Each	104		

			transparent for visual inspection, flexible dressing. MCC registration, Sterile, Individually peel packed, Size of Dressing: ±50mm x 200mm, Each				
H-8	2857	Dressing Hydrogel Sterile 15g Squeeze	Amorphous Hydrogel Dressing gel, desloughing agent, colourless transparent gel with adhesion properties that moisturises wounds. Sterile 15g Squeeze tube with twist cap. Each	Each	1 820		
H-9	3250	Dressing hydro-polymer foam 15 X 20 cm	Dressing wound absorbent non-adherent with adhesive border Layered dressing to be: waterproof and bacterial proof Borders to prevent leakage Highly absorbent centre to accommodate large volume of exudate and odours Size: 15cm X 20cm Packaging: box of 10 dressings	Box of 10	2 582		
H-10	2237	Dressing Sterile, Hydro Capillary, Non-Adherent, Absorbent 20Cmx1m	Dressing Wound, hydro capillary, woven. Absorbent, layered dressing to assist in the removal of exudate and dead tissue, be non-adherent to wound surface, sterile. Should have description on outer pack. Sterile, individually peel packed Size Dressing: ±200mm x 1m Each"	Each	1986		
H-11	2235	Dressing Sterile, Hydro Capillary, Non-Adherent, Absorbent 15Cmx20cm	Dressing Wound, hydro capillary, woven. Absorbent, layered dressing to assist in the removal of exudate and dead tissue, be non-adherent to wound surface, sterile. Should have description on outer pack. Sterile, individually peel packed. Size Dressing: ±150mm x 200mm. Each"	Each	1304		

H-12	4030	Dressing Wound Absorbent Non-Adhere 18X18 Adhesive	Dressing wound absorbent non-adherent with adhesive border Layered dressing to be: waterproof and bacteria proof non-adherent super-absorbent layer Moisture vapour permeable with a waterproof polyurethane backing Size: 18cm X 18cm Packaging: box of 10 dressings (individually peel packed, sterile)	Box of 10	117		
H-13	4031	Dressing Wound Absorbent Non-Adherent 12X12	Dressing wound absorbent non-adherent with adhesive border Layered dressing to be: waterproof and bacteria proof non-adherent super-absorbent layer Moisture vapour permeable with a waterproof polyurethane backing Size: 12cm X 12cm Packaging: box of 10 dressings (sterile, individually peel packed,)	Box of 10	39		
H-14	1167	Dressing Wound Adhesive. Size 280X450mm 10	Dressing wound adhesive transparent to adhere to incision/wound edge Size 280mm X 450mm 10 dressings, each dressing must be sterile, individually wrapped peel packed, it must be water vapour permeable, water resistance, both primary and secondary labelling must have the following information: item description, size, batch number, expiry date, manufacturing date, quantity	Pack of 10	2 363		
H-15	1903	Eye Shield Elastic; Flesh-Coloured Plastic;	Eye shield, elastic for left or right eye, with an elastic strap size: Approximate 5x7cm.	Each	4285		

H-16	1175	Gauze Ribbon Absorbent Cotton; Size 12mmx10m	Gauze, high degree of absorbency cotton, type 1, ribbon, lint free with no loose edges, Size: 12mm x 10m, Individually packed, non-sterile, To comply with the latest issue of SANS 446 compliance certificate to be submitted with bid, Each	Each	156		
H-17	1176	Gauze Ribbon Absorbent Cotton; Size 25mmx10m	Gauze, high degree of absorbency cotton, type 1, ribbon, lint free with no loose edges, Size: 25mm x 10m, Individually packed, non-sterile, To comply with the latest issue of SANS 446 compliance certificate to be submitted with bid, Each	Each	1534		
H-18	1177	Gauze Ribbon Absorbent Cotton; Size 50mmx10m	Gauze, high degree of absorbency cotton, type 1, ribbon, lint free with no loose edges, Size: 50mm x 10m, Individually packed, non-sterile, To comply with the latest issue of SANS 446 compliance certificate to be submitted with bid, Each	Each	2088		
H-19	1178	Gauze Ribbon Absorbent Cotton; Size 75mmx10m	Gauze, high degree of absorbency cotton, type 1, ribbon, lint free with no loose edges, Size: 75mm x 10m, Individually packed, non-sterile, To comply with the latest issue of SANS 446 compliance certificate to be submitted with bid, Each	Each	1		
H-20	1187	Haemostatic sponge, gelatine, absorbent, 70x50x1mm	Sponge gelatin absorbable haemostatic B.P./U.S.P., STERILE, water-insoluble, malleable, porcine gelatin absorbable sponge, 1 sponge 70mm x 50mm x1mm sterile peel packed 1 sponge	Each	83		

H-21	1932	Nasal Tampon 80mm with Cannula	Sponge, Surgical, Polyvinyl alcohol type X-ray detectable, Nasal tampon 80mm – double compressed, sterile peel packed, Each	Each	975		
H-22	1186	S Haemostatic sponge, gelatine, absorbent, 70x50x10mm	Sponge gelatin absorbable haemostatic B.P./U.S.P., STERILE, water-insoluble, malleable, porcine gelatin absorbable sponge, 1 sponge 70mm x 50mm x10mm sterile peel packed 1 sponge	Each	1131		
H-23	1186	Sponge Gelatin Absorbable. Haemostatic 70X50x10mm 2'S	Absorbable haemostatic gelatin sponge, sterile haemostat for surgical use, labelling must have the following information: item description, size, batch/ lot number, expiry date, manufacturing date, single use. Pack of 2	Pack of 2	11 131		
H-24	1192	Swabs Cotton Abdominal X-Ray detectable, Autoclavable 4 ply Tied in 5 , 350 x350mm	Swabs, abdominal, woven, cotton, X-Ray detectable, autoclavable, non-sterile, 4ply 5 x 5 swabs, banded in 5 and clearly marked with the size and quantity, Size: 350mm x 350mm Scope All layers to consist of white open wove bandage cloth as per the LATEST issue of the SANS 443 Compliance certificate to be submitted with bid Construction Edges: All cut edges of the material to be folded in and stitched leaving no frayed edges Cotton tape: Length: 400mm Width of the tape: 12mm The cotton tape must be embedded in a	Each	4150		

			50mm pocket with an X (i.e. 100mm in length of tape folded double to 50mm stitched into the corner of swab leaving a 300mm length of the cotton tape clear.) Minimum breaking strength of tape 7,95 kg. X-ray detectable thread shall be enclosed by the outer layer of material which can withstand autoclaving. Packing: The swabs shall be bound in bundles of five by means of either a paper band consisting of one piece secured by autoclavable glue (and not by a separable adhesive tape) or by means of an acceptable tying material which can withstand autoclaving. packed in a sterile peel packed pouch. Pack of 100				
H-25	1200	Swabs Cotton Dissecting balls X- Ray detectable - 10mm diameter Tied in 5	Swabs, surgical, dissecting, type 1, X-ray detectable, 100 swabs Size 10mm Materials: Outer Layer - The outer layer shall be gauze The cotton wool shall be cotton wool Grade 2 Construction: The cotton wool shall be packed in single layer of gauze X-ray detectable material of size 5mm X 5mm to be placed between the cotton wool and the outer layer of gauze so as to be visible from the outside The gauze must be secured with cotton threat of acceptable	Pack of 100	57		

			quality The swabs shall be loosely stitched together in bundles of five by means of an uneasy to break cotton thread. Pack of 100 swabs				
H-26	1195	Swabs Gauze Absorbable X-Ray detectable 8Ply x100x100mm	Swabs, abdominal, cotton, type 2, x-ray detectable, Autoclavable, sterile, size 8 ply x 10cm x 10cm, scope all layers to consist of white open bandage cloth as per the latest issue of the SANS 443, compliance to be submitted with the bid construction, edges: all cut edges of the material must be folded in and stitches leaving no frayed edges, cotton tape length: 400mm, width of the tape: 12mm, the cotton tape must be embedded in 50mm pocket with an x (i.e. 100mm in length of tape folded double to 50mm stitched into the corner of the swab leaving a 300mm length of the cotton tape clear.), minimum breaking strength of the tape 7.95kg, x-ray detectable thread shall be enclosed by the outer layer of the material which can withstand autoclaving., packing the swab must be bounded in bundle of five by means of either a paper band consisting of one piece secured by Autoclavable glue (and not by a separable adhesive tape) or by means of an acceptable tying material which can withstand autoclaving, packed in	Pack of 100	44 881		

			a sterile peel packed pouch. Pack of 100				
H-27	4001	Tray, Surgical Pre-Packed for minor Surgical Procedures	Tray, surgical, pre-packed, for minor surgical procedures. The tray must be sterile. Must have a +/- 10mm width seal. Must open with relative ease. Must have rounded corners with no serrated edges Tray must contain the minimum requirements and be packed in the following in this manner: 1X Plastic disposable bag +/- 400X250mm; 2X Medical towel serviette +/-330X300mm; 2X Latex gloves medium; 1X Plastic non-toothed forceps +-130mm; 2pair X Protection sheet +/-400X400mm; 5X Gauze swabs plain +/- 100X100mm 8ply; 5X Cotton balls 1g; 1X compartment +/- 110X40mm depth +/-40mm; 1X compartment +/- 110X100mm depth +/-40mm; 1 Tray +/-175X150mm with two compartments Large Tray. Each	Each	123 846		
I-1	1127	Bandage P.O.P. Roll; 100mmx3.5m	Bandage, Cast, Plaster of Paris, Leno weave base cloth, Type 1, Size: 100mm x 3.5m, Individually wrapped and Packed in a sealed containers, SANS 1308, Table 1 and 2	Each	1836		
I-2	1126	Bandage P.O.P.Roll;50Mmx3m 12'S	Bandage, Cast, Plaster of Paris, Leno weave base cloth, Type 1, Size: 50mm x 3.5m, Individually wrapped and Packed in a sealed containers, SANS 1308, Table 1 and 2	Each	653		

I-3	3049	Blanket Rescue Paediatric	Blanket Rescue Paediatric, material: plastic, metallised with aluminium, measure: 160 x 210 cm, colour gold/ silver, weight: approx. 60g, CE certificate medical device ac. To 93/42/EEC, highly reflective, extremely thin and tear-resistant, prevent heat loss, protects against hypothermia, and stops largely the thermal radiation, good visibility at accident site. Each. Individually wrapped.	Each	213		
I-4	3360	Collar, Stiff neck Adult	Adjustable stiff neck single piece collar/brace, with adjustable Velcro straps, PVC covered with foam, chin support, for single use, prior diagnostic, must have an open tracheal window. Each. Individually wrapped.	Each	723		
I-5	3360	Collar, Stiff neck Adult	Adjustable stiff neck single piece collar/ brace with adjustable Velcro straps, PVC covered with foam, Chin support for single use prior diagnostic, the must be an open tracheal window, Size Adult. Individually wrapped.	Each	723		
I-6	3358	Collar, Stiff neck Paediatric	Adjustable stiff neck single piece collar/brace, with adjustable Velcro straps, PVC covered with foam, chin support, for single use, prior diagnostic, there must have an open tracheal window. Individually wrapped. Each	Each	20		

I-7	3358	Collar, Stiff neck Paediatric	Adjustable stiff neck single piece collar/ brace with adjustable Velcro straps, PVC covered with foam, Chin support for single use prior diagnostic, the must be an open Individually wrapped tracheal window, Size Paediatric.	Each	803		
I-8	1139	Stockinette tubular orthopaedic bandage 100Mmx25m	Bandage Tubular, orthopaedic, knitted, Cotton Type, stockinette, unbleached. Size Width 100mm, Length 25m, Roll Individually boxed	Each	1014		
I-9	1119	Stockinette tubular orthopaedic bandage 150mm x25m	Bandage Tubular, orthopaedic, knitted, Cotton Type, stockinette, unbleached. Size Width 150mm, Length 25m, Roll Individually boxed	Each	1		
I-10	1121	Stockinette tubular orthopaedic bandage 250Mmx25m	Bandage Tubular, orthopaedic, knitted, Cotton Type, stockinette, unbleached. Size Width 250mm, Length 25m, Roll Individually boxed	Each	68		
I-11	1133	Stockinette tubular orthopaedic bandage 50Mmx25m	Bandage Tubular, orthopaedic, knitted, Cotton Type, stockinette, unbleached. Size Width 50mm, Length 25m, Roll Individually boxed	Each	403		
I-12	1134	Stockinette tubular orthopaedic bandage 80Mmx25m	Bandage Tubular, orthopaedic, knitted, Cotton Type, stockinette, unbleached. Size Width 80mm, Length 25m, Roll Individually boxed	Each	325		
J-1	1588	Bag, Rebreathing with reservoir, 22mm, Size: 0.5L	Bag, rebreathing, Size 0.5 litre latex free, tapered neck with collar, 15mm ID female connector, sing use, it must not twist or kink on itself as demonstrated in a clinical setting. Size 0.5 litre with 15mm F connector, Compliance: 30-60cmH20 filled to 4 x	Each	1		

			nominal volume, must hold at least 10x nominal volume before it will burst, must not twist or kink on itself – designed to prevent this and demonstrate in a clinical setting. Must retain original volume and shape (withi10%) at deflation. Must be antistatic, non-conductive and must not adhere to itself				
J-2	1589	Bag, Rebreathing with reservoir, 22mm, Size: 1	Bag, rebreathing, Size 1 litre latex free, tapered neck with collar, 15mm ID female connector, sing use, it must not twist or kink on itself as demonstrated in a clinical setting. Size 1 litre with 22mm F connector, Compliance: 30-60cmH20 filled to 4 x nominal volume, must hold at least 10x nominal volume before it will burst, must not twist or kink on itself – designed to prevent this and demonstrate in a clinical setting. Must retain original volume and shape (withi10%) at deflation. Must be antistatic, non-conductive and must not adhere to itself	Each	**1**		
J-3	1590	Bag, Rebreathing with reservoir, 22mm, Size: 2L	Bag, rebreathing, Size 2 litre latex free, tapered neck with collar, 15mm ID female connector, sing use, it must not twist or kink on itself as demonstrated in a clinical setting. Size 2 litre with 22mm F connector, Compliance: 30-60cmH20 filled to 4 x nominal volume, must hold at least 10x nominal volume before it will	Each	512		

			burst, must not twist or kink on itself – designed to prevent this and demonstrate in a clinical setting. Must retain original volume and shape (withi10%) at deflation. Must be antistatic, non-conductive and must not adhere to itself				
J-4	3001	Catheter Suction 6Fr Neonatal	Close suction system, siliconised medically safe PVC, Atraumatic, bevelled tip. A single lumen endotracheal suction. System to seal airway from atmospheric contamination and maintain uninterrupted ventilation. Must have standard 15mmtaper fittings. Must be fitted with durable flexible plastic sheath for storage, to avoid contamination. Must be provided with lateral irrigation port with a one-way valve. Radio – opaque line on catheter tip. Must be able to maintain positive pressure ventilation. Single use 24 hours, Sterile, Size 6FR	Each	907		
J-5	4473	Close suction intubation catheter - Endotracheostomy - 10Fg x 50cm (min)	Close suction intubation catheter - Endotracheostomy - 10Fg x 50cm (min). Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included.	Each	1		

J-6	4474	Close suction intubation catheter - Endotracheostomy - 12Fg x 50cm (min)	Close suction intubation catheter - Endotracheostomy - 12Fg x 50cm (min). Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included.	Each	52		
J-7	3125	Close suction intubation catheter - Endotracheostomy - 14Fg x 50cm (min)	Close suction intubation catheter - Endotracheostomy - 14Fg x 50cm (min). Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included.	Each	4212		
J-8	4471	Close suction intubation catheter - Endotracheostomy - 8Fg x 35cm	Close suction intubation catheter - Endotracheostomy - 8Fg x 35cm. Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included. Use up to 72 hours. ET tube adaptors for 3mm, 3.5m and 4mm	Each	1		
J-9	4472	Close suction intubation catheter - Tracheostomy - 10Fg x 30cm	Close suction intubation catheter - tracheostomy - 10Fg x 30cm (min). Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included.	Each	44		

J-10	3126	Close suction intubation catheter - Tracheostomy - 14Fg x 30cm (min)	Close suction intubation catheter - Tracheostomy - 14Fg x 30cm (min). Latex and DEHP free, one way or dedicated port for saline lavage, Straight with Suction Control Valve Double swivel elbow. 15mmx22mm flex connector. Day stickers should be included.	Each	1690		
J-11	3047	E/T Intubation Stylet Small	Intubation stylet, Neonatal: Purpose to pre-form an endotracheal tube to a desired shape to aid intubation. Size Neonatal to fit endotracheal tube size 2.0 – 3.5 mmm. Length 225mm. Manufactured from low friction atraumatic outer coating around malleable aluminium core. Soft atraumatic tip, U-bent or other mechanism at proximal end to limit depth. Latex free. Clinically clean packaging. Individually packed in peel pouch.	Each	1		
J-12	1581	Guedel, Airway size 0 - Grey	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic implant tested EVA or medically safe PVC surgical clean. Size:0, length 50mm. Must be individually wrapped	Each	4238		
J-13	1582	Guedel, Airway size 00 - Blue	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic	Each	2155		

			implant tested EVA or medically safe PVC surgical clean. Size:00, length 40mm. Must be individually wrapped				
J-14	1584	Guedel, Airway size 1 - White	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic implant tested EVA or medically safe PVC surgical clean.Size:1, length 70mm. Must be individually wrapped	Each	2265		
J-15	1585	Guedel, Airway size 2 - Green	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic implant tested EVA or medically safe PVC surgical clean.Size:2, length 80mm. Must be individually wrapped	Each	10 299		
J-16	1586	Guedel, Airway size 3 - Orange/Yellow	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic implant tested EVA or medically safe PVC surgical clean.Size:3, length 90mm. Must be individually wrapped	Each	17 950		
J-17	1587	Guedel, Airway size 4 - Red	Airway, oropharyngeal, with rigid bite block, smooth rounded distal end (Guedel). Anatomically appropriate contour. Flanged buccal end. manufactured from non-toxic implant tested EVA or medically safe PVC surgical clean.Size:4, length	Each	11 661		

			100mm. Must be individually wrapped				
J-18	2377	Mask Laryngeal Airway Size 4 (12Mmx20mm)	Laryngeal mask airway device, orolaryngeal type. Reusable, autoclavable up to 40 times. Silicone tube and cuff, epiglottis protection bars. Size: 4 (50 to 70Kg) Adult. ID: 12mm, OD: 20 mm.	Each	1037		
J-19	1602	Mask, Anaesthetic with inflatable silicone rim, 22mm connector, Size 2	Mask, Anaesthetic, clear silicon round shade, soft Inflatable cushion rim with a 22mm connector. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 2. Re-usable and Autoclavable. Individually packed and surgical clean,	Each	1		
J-20	2521	Mask, Anaesthetic soft malleable silicone rim, 22mm connector, Size 3	Mask, Anaesthetic, clear silicon round shade, soft rim with a 22mm connector. Malleable but not collapsible. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 4. Re-usable and autoclavable. Individually packed and surgical clean	Each	81		
J-21	1605	Mask, Anaesthetic with inflatable silicone rim, 22mm connector, Size 0	Mask, Anaesthetic, clear silicon round shade, soft Inflatable cushion rim with a 22mm connector. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 0. Re-usable	Each	101		

			and autoclavable. Individually packed and surgical clean				
J-22	1606	Mask, Anaesthetic with inflatable silicone rim, 22mm connector, Size 1	Mask, Anaesthetic, clear silicon round shade, soft Inflatable cushion rim with a 22mm connector. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 1. Re-usable and autoclavable. Individually packed and surgical clean	Each	73		
J-23	2657	Mask, Anaesthetic with inflatable silicone rim, 22mm connector, Size 3	Mask, Anaesthetic, clear silicon round shade, soft Inflatable cushion rim with a 22mm connector. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 3. Re-usable and autoclavable. Individually packed and surgical clean	Each	23		
J-24	2658	Mask, Anaesthetic with inflatable silicone rim, 22mm connector, Size 3	Mask, Anaesthetic, clear silicon round shade, soft rim with a 22mm connector. Malleable but not collapsible. Able to create a good seal between the mask and patients face, Good fit between mask connector and breathing circuit. Latex free. Size 5. Re-usable and autoclavable. Individually packed and surgical clean	Each	304		
J-25	3983	Tracheostomy tube - Cuffed, Size 5 - Low pressure	Tracheostomy tube - Cuffed, Size 5 - Low Pressure PVC/ polyurethane. Outer cannula with a cuff, inner cannula with connector Perforated	Each	1		

			obturator and wide neck strap. With sizes close to following ID-IC: 5mm; OD-OC 8.6mm; OD-OC bf: 10.1mm Length TL: 66mm Length OB: 73mm BA: 100° CD: 18mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer cannula behind the flange; Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)				
J-26	4443	Tracheostomy tube - Fenestrated, Size 6 - Low pressure with 2 inner cannulas	Tracheostomy tube - Fenestrated, Size 6 - Low Pressure, PVC/ polyurethane. Outer cannula, fenestrated with a cuff, inner cannula, fenestrated with connector Perforated obturator and wide neck strap. With sizes close to following ID-IC: 6mm; OD-OC 9.2mm; OD-OC bf: 10.8mm Length TL: 72mm Length OB: 80mm BA: 95° CD: 23mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer	Each	18		

			cannula behind the flange; Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)				
J-27	4445	Tracheostomy tube - Fenestrated, Size 8 - Low pressure with 2 inner cannulas	Tracheostomy tube - Fenestrated, Size 8 - Low Pressure, PVC/ polyurethane. Outer cannula, fenestrated with a cuff, inner cannula, fenestrated with connector Perforated obturator and wide neck strap. With sizes close to following ID-IC: 8mm; OD-OC 11.4mm; OD-OC bf: 12.7mm Length TL: 76mm Length OB: 83mm BA: 90° CD: 28mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer cannula behind the flange; Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)	Each	1		
J-28	3987	Tracheostomy tube - non-cuffed, Size 6 - Low pressure	Tracheostomy tube - Non-Cuffed, Size 6 - Low Pressure PVC/ polyurethane. Outer cannula with a cuff, inner cannula with connector	Each	1		

			Perforated obturator and wide neck strap. With sizes close to following ID-IC: 6mm; OD-OC 9.2mm; OD-OC bf: 10.8mm Length TL: 72mm Length OB: 80mm BA: 95° CD: 23mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer cannula behind the flange; Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)				
J-29	3988	Tracheostomy tube - non-cuffed, Size 7 - Low pressure	Tracheostomy tube - Non-Cuffed, Size 7 - Low Pressure PVC/ polyurethane. Outer cannula with a cuff, inner cannula with connector Perforated obturator and wide neck strap. With sizes close to following ID-IC: 7mm; OD-OC 10.4mm; OD-OC bf: 12.0mm Length TL: 74mm Length OB: 80mm BA: 95° CD: 26mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer cannula behind the flange;	Each	52		

			Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)				
J-30	3993	Tracheostomy tube - non-cuffed, Size 9 - Low pressure	Tracheostomy tube - Non-Cuffed, Size 8 - Low Pressure PVC/ polyurethane. Outer cannula with a cuff, inner cannula with connector Perforated obturator and wide neck strap. With sizes close to following ID-IC: 9mm; OD-OC 12.5mm; OD-OC bf: 14.2mm Length TL: 78mm Length OB: 88mm BA: 90° CD: 30mm (ID-IC: Inside diameter at bottom of inner cannula; OD-OC bc: Outside diameter at bottom of outer cannula; OD-OC bf: Outside diameter of outer cannula behind the flange; Length TL: Length along the centre line from start of neck flange to bottom of tube; Length OB: Length along outer bend BA: Bending angle CD: Cuff diameter)	Each	21		
J-31	1838	Tube, E/T nasal, preformed cuff ID: 7mm	Tube, endotracheal, cuffed medically safe PVC, thermo sensitive. Preformed for nasal use. Murphy tip and eye. Radio opaque line. Tube to have clear depth markings, all components to be latex free. 15mm	Each	65		

			OD connector. Individual sterile peel pack 7.0 mm ID				
J-32	1840	Tube, E/T nasal, preformed cuff ID: 8mm	Tube, endotracheal, cuffed medically safe PVC, thermo sensitive. Preformed for nasal use. Murphy tip and eye. Radio opaque line. Tube to have clear depth markings, all components to be latex free. 15mm OD connector. Individual sterile peel pack 8.0 mm ID	Each	215		
J-33	2734	Tube, E/T oral non-cuff ID: 4.0mm x 230mm	Tube, Endotracheal, Non-Cuffed, Oral/Nasal/with a specially designed tip to reduce airways trauma. High volume and low-pressure. Barrel shaped cuff to facilitate optimal pressure dispersion against the tracheal wall. Shall incorporate a soft Flexible point in the midline so as to pass easy between the vocal chords without impaction thereof. The tube shall incorporate two murphy eyes for additional safety, the tube must be able to demonstrate a reduction in intubation, induced airways trauma. Tube must be marked with numbered depth marking of 1cm intervals, starting from the tip of the tube. The marking on the tube (tube size, length, ID, OD) must be clear and not rub-off. Size 4.0 mm x 230 mm. Sterile individually peel packed.	Each	130		
J-34	2732	Tube, E/T oral non-cuff ID: 4.5mm x 230mm	Tube, Endotracheal, Non-Cuffed, Oral/Nasal/with a specially designed tip to reduce airways trauma. High	Each	130		

			volume and low-pressure. Barrel shaped cuff to facilitate optimal pressure dispersion against the tracheal wall. Shall incorporate a soft Flexible point in the midline so as to pass easy between the vocal chords without impaction thereof. The tube shall incorporate two murphy eyes for additional safety, the tube must be able to demonstrate a reduction in intubation, induced airways trauma. Tube must be marked with numbered depth marking of 1cm intervals, starting from the tip of the tube. The marking on the tube (tube size, length, ID, OD) must be clear and not rub-off. Size 4.5mm x 230mm. Sterile individually peel packed.				
K-1	4435	Foetal Delivery Device, Vacuum Assisted. With a Force / Traction Indicator, Disposable.	Complete vacuum assisted foetal delivery device, should have a flexible stem and low profile cup to enable placement over the flexion point, Integral vacuum pump with vacuum strength indicator and release button, Integrated traction handle with foam filter with a 60mm diameter and 20mm cup depth. Must be Latex free, must be sterile and individually packed.	Each	2751		
L-1	4533	CSF drainage bag - 600ml	Cerebrospinal fluid drainage bag with integral drip chamber in a 600ml translucent drainage bag without a vent. The bag should contain an access port with stopcock and	Each	1		

			removable cap at the bottom end. The top should have a drip chamber, 30cm tubing length with a clamp and a male Luer lock connector. The bag should be graduated in both ml and 360mmH_2O Adjustable cord at the top of the bag to allow for regulation of drainage from a single point of suspension				
L-2	4532	CSF drainage system - external 600ml drainage bag	Cerebrospinal fluid drainage system for aseptic shunting from the ventricular cavities or lumbar subarachnoid spaces Integral drip chamber in a 600ml translucent drainage bag without a vent. A striped tubing patient line of approximately 165cm for easy identification with a female luer lock connector, clamp, anti-reflux valve, a y-shaped injection port and 3-way stopcock with 10cm proximal tubing and male luer lock connector. The bag should contain an access port with stopcock and removable cap at the bottom end. The top should have a drip chamber, 30cm tubing length with a clamp and a male Luer lock connector. The bag should be graduated in both ml and 360mmH_2O Adjustable cord at the top of the bag to allow for regulation of drainage from a single point of suspension	Each	104		
L-3	1148	Drape Incise: Adhesive area 600 x 450mm, Drape size 900 x 450mm	Drape, Surgical, Adhesive patch with circle aperture centrally situated, Size	Each	169		

			400mm x 400mm, Adhesive area: 600mm x 450mm, Circle Aperture: 900mm x 450mm Diameter Material, Adhesive and handles as per drip surgical incise adhesive above, Each				
L-4	4487	Electrosurgical Diathermy Dual plate for Adult use with a REM plug	Electrosurgical Diathermy plate for adults. Dual plate with water based hydrogel elongated fills/zones in a vertical split. Should have an electrode insulating safety ring. Cable length of 3m with a REM plug connector	Each	11 929		
L-5	4537	Electrosurgical Diathermy Dual plate for Paediatric use with a REM plug	Electrosurgical Diathermy plate for Paediatric use. Dual plate with water based hydrogel elongated fills/zones in a vertical split. Should have an electrode insulating safety ring. Cable length of 3m with a REM plug connector	Each	520		
L-6	2630	Electrosurgical Diathermy pencil with a 70mm blade electrodes attachment. Hand controls with large buttons and a 3m flexible cable with 3 point connector.	Electrosurgical Diathermy pencil with 70mm blade. Slim, sleek ergonomically grip with large button controls for both cutting (yellow) and coagulation (blue). Pencil length 170mm, Shape of tip: Blade, Length of blade: 70mm Diameter of blade: 2.3 mm. 3m long flexible cable with standard 3-point connector. Latex free. Must be compatible and use with interchangeable cutting tips/electrodes that locks safely into place but easily exchanged Individually sterile wrapped peel packets	Each	17 332		

L-7	4535	Electrosurgical electrode - loop 15x12mm	Electrosurgical Diathermy tip/electrode replacement attachment for standard hand control Diathermy Pencil. Tip Shape: Loop. Pen length 130mm. Loop size: 15x12mm Individually sterile wrapped peel packets	Each	910		
L-8	4536	Electrosurgical electrode - loop 15x15mm	Electrosurgical Diathermy tip/electrode replacement attachment for standard hand control Diathermy Pencil. Tip Shape: Loop. Pen length 130mm. Loop size: 15x15mm Individually sterile wrapped peel packets	Each	172		
L-9	4534	Electrosurgical tip/electrode - blade 150mm	Electrosurgical Diathermy tip/electrode replacement attachment for standard hand control Diathermy Pencil. Tip Shape: Blade. Blade length 150mm Individually sterile wrapped peel packets	Each	234		
L-10	2819	Knife A-Ok Full Handle 15º	Micro scalpels - ophthalmic stab knife with full handle, blade 15° (degrees) double bevelled. Stainless steel. Sterile and individually wrapped.	Each	109		
L-11	1657	Plain Catgut taper point, 2/0, 90cm ½ circle 45mm	Type: Plain Catgut - Absorbable Suture size: 2/0 Suture Length: 90cm Needle size: 45mm Needle shape: ½ circle Cut Type: Taper point Box: 12 per box	Pack of 12	26		
L-12	2941	Suture - Braided black silk, Reverse cut x2, 6/0, 45cm, ⅜ circle 13mm	Type: Braided black silk Suture size: 6/0	Pack of 12	1		

			Suture Length: 45cm Needle size: 13mm Needle shape: ⅜ circle Cut Type: Reverse cut x2 Box: 12 per box				
L-13	2362	Suture - Chromic Catgut, reverse cut, 2/0, 75-90cm, ⅜ circle 36-37mm	Type: Chromic Catgut Suture size: 2/0 Suture Length: 75-90cm Needle size: 36-37mm Needle shape: ⅜ circle Cut Type: Reverse cut Box: 12 per box	Pack of 12	1		
L-14	1680	Suture - Chromic Catgut, reverse cut, 2/0, 75cm, ⅜ circle 30mm	Type: Chromic Catgut Suture size: 2/0 Suture Length: 75cm Needle size: 30mm Needle shape: ⅜ circle Cut Type: Reverse cut Box: 12 per box	Pack of 12	1245		
L-15	3605	Suture - Chromic Catgut, taper cut, 4/0, 75cm, ½ circle 22mm	Type: Chromic Catgut Suture size: 4/0 Suture Length: 75cm Needle size: 22mm Needle shape: ½ circle Cut Type: Taper cut Box: 12 per box	Pack of 12	1		
L-16	2367	Suture - Chromic Catgut, taper heavy, 2, 75cm, ½ circle 50mm	Type: Chromic Catgut Suture size: 2 Suture Length: 75cm Needle size: 50mm Needle shape: ½ circle Cut Type: Taper Heavy Box: 12 per box	Pack of 12	75		

L-17	1725	Suture - Nylon/Polyamide, Conventional cut, 0, 100cm, ⅜ circle, 90mm	Type: Nylon/Polyamide Suture size: 0 Suture Length: 100cm Needle size: 90mm Needle shape: ⅜ circle Cut Type: Conventional cut Box: 12 per box	Pack of 12	1303		
L-18	1742	Suture - Nylon/Polyamide, Reverse cut, 2/0, 45cm, ⅜ circle, 26mm	Type: Nylon/Polyamide Suture size: 2/0 Suture Length: 45cm Needle size: 26mm Needle shape: ⅜ circle Cut Type: Reverse cut Box: 12 per box	Pack of 12	3762		
L-19	2398	Suture - Nylon/Polyamide, Spatula,10/0, 30cm, ⅜ circle, 6mm	Type: Nylon/Polyamide Suture size:10/ 0 Suture Length: 30cm Needle size: 6mm Needle shape: ⅜ circle Cut Type: Spatula Box: 12 per box	Pack of 12	78		
L-20	1726	Suture - Nylon/Polyamide, Straight cut, 2/0, 100cm, ½ circle, 55mm	Type: Nylon/Polyamide Suture size: 2/0 Suture Length: 100cm Needle size: 55mm Needle shape: ½ circle Cut Type: Straight cut Box: 12 per box	Pack of 12	125		
L-21	1745	Suture - Nylon/Polyamide, Taper cut, 0, 150cm loop, ½ circle, 45mm	Type: Nylon/Polyamide Suture size: 0 Suture Length: 150cm loop Needle size: 45mm Needle shape: ½ circle	Pack of 12	140		

			Cut Type: Taper cut Box: 12 per box				
L-22	1756	Suture - Nylon/Polyamide, Taper cut, 1, 100cm, ½ circle, 30mm	Type: Nylon/Polyamide Suture size: 1 Suture Length: 100cm Needle size: 30mm Needle shape: ½ circle Cut Type: Taper cut Box: 12 per box	Pack of 12	21		
L-23	1736	Suture - Plain Catgut taper point, 2/0, 75cm ½ circle 36mm	Type: Plain Catgut - Absorbable Suture size: 2/0 Suture Length: 75cm Needle size: 36mm Needle shape: ½ circle Cut Type: Taper point Box: 12 per box	Pack of 12	140		
L-24	1714	Suture - Polybutester blue, Colt, 3/0, 100cm, ⅜ circle 80-90mm	Type: Polybutester blue Suture size: 3/0 Suture Length: 100cm Needle size: 80-90mm Needle shape: ⅜ circle Cut Type: Colt Box: 12 per box	Pack of 12	1		
L-25	3371	Suture - Polydioxanone monofilament, Taper point, 4/0, 70cm, ½ circle, 22mm	Type: Polydioxanone monofilament Suture size: 4/0 Suture Length: 70cm Needle size: 22mm Needle shape: ½ circle Cut Type: Taper point Box: 12 per box	Pack of 12	68		
L-26	3369	Suture - Polydioxanone monofilament, Taper point/heavy, 1, 90cm, ½ circle, 40mm	Type: Polydioxanone monofilament Suture size: 1 Suture Length: 90cm Needle size: 40mm	Pack of 12	73		

			Needle shape: ½ circle Cut Type: Taper point/heavy Box: 12 per box				
L-27	2675	Suture - Polyester braided, Taper cut, 2/0, 75cm, ½ circle, 17mm	Type: Polyester braided Suture size: 2/0 Suture Length: 75cm Needle size: 17mm Needle shape: ½ circle Cut Type: Taper cut Box: 12 per box	Pack of 12	1		
L-28	3029	Suture - Polyester braided, Taper point, 3/0, 90cm, ½ circle, 17.4mm	Type: Polyester braided Suture size: 3/0 Suture Length: 90cm Needle size: 17.4mm Needle shape: ½ circle Cut Type: Taper point Box: 12 per box	Pack of 12	23		
L-29	1702	Suture - Polyglycolic acid, Taper cut, 4/0, 70-75cm, ½ circle 20-22mm	Type: Polyglycolic acid Suture size: 4/0 Suture Length: 70-75cm Needle size: 20-22mm Needle shape: ½ circle Cut Type: Taper cut Box: 12 per box	Pack of 12	117		
L-30	1705	Suture - Polyglycolic acid, Taper fine point, 2/0, 70-75cm, ½ circle 26mm	Type: Polyglycolic acid Suture size: 2/0 Suture Length: 70-75cm Needle size: 26mm Needle shape: ½ circle Cut Type: Taper fine point Box: 12 per box	Pack of 12	2233		
L-31	2406	Suture - Polypropylene monofilament, reverse cut, 5/0, 45cm, ⅜ circle, 17,4mm	Type: Polyproylene monofilament Suture size: 5/ 0 Suture Length: 45cm	Pack of 12	52		

			Needle size: 17,4mm Needle shape: ⅜ circle Cut Type: Reverse cut Box: 12 per box				
L-32	1701	Suture - Vicryl/Polyglycolic acid, Reverse cut, 5/0, 45cm, ⅜ circle 17.6mm	Type: Vicryl/Polyglycolic acid Suture size: 5/0 Suture Length: 45cm Needle size: 17.6mm Needle shape: ⅜ circle Cut Type: Reverse cut Box: 12 per box	Pack of 12	57		
L—33	2721	Suture - Vicryl/Polyglycolic acid, Taper cut, 0, 75cm, ½ circle 30mm	Type: Vicryl/ Polyglycolic acid Suture size: 0 Suture Length: 75cm Needle size: 30mm Needle shape: ½ circle Cut Type: Taper cut Box: 12 per box	Pack of 12	5		
M-1	1565	Tube – Feeding tube – nasal 5FG	TUBE, NASAL, FEEDING Medical grade non-toxic PVC, Proximal end: Luer connector and plug, Distal end: closed round tip with TWO lateral eyes, Length: not less than 400mm. Size: 5 FG (outer diameter1,7mm) Sterile, Individually peel packed, EACH	Each	53 316		
N-1	4839	Administration Set Autofusion 20Drps Y-Clave Hydrophobic Cap Auto	Admin Autofusion set 20 drops with Y-clave, Auto shut off and hydrophobic prime cap 200cm. Sterile individual peel pack for single use.	Each	300		
N-2	4889	Admin Infusion Set 20 drop with 0.2micron filter 250cm.	Admin Infusion Set 20 drop with 0.2micron filter 250cm. Sterile individual peel pack for single use	Each	1		

N-3	4830	Admin Infusion Set Opaque Y-Site 180Cm	Admin Infusion Set Opaque Y-site 180cm. Sterile individual peel pack for single use	Each	260		
N-4	4838	Administration Set 20 Drops 200Cm 2Mic Filter Hydrophobic Cap	Admin set 20 drop, 200cm, with 2 micron filter and hydrophobic prime cap for Paxitaxol 011CH5050TX Sterile individual peel pack for single use	Each	390		
N-5	4840	Administration Set Air Vent 20 Drps,72mm Chamber, Y-Clave No Needle	Admin set with air vent, Light sensitive,20 drops, long chamber(72mm) 200cm clear DEHP free tube, Y Clave, Imported retractable rotating luer lock with cap. Sterile individual peel pack for single use	Each	260		

EVALUATION CRITERIA

1. First stage: Pre - Evaluation on compulsory returnable
2. Second stage: Preferential Procurement Policy Framework Act (PPPFA) points scoring.
3. Third stage: Product Technical Compliance. This will only be required from bidders who advanced through the first and second stage

Returnable Documents

Each bid shall comprise of a clearly indexed proposal with the tender documents as follows:

Section	Compulsory Returnable Documents	Attached YES / NO
a.	SBD 1 - Invitation to bid	
b.	SBD 4 - Declaration of interest	
c.	SBD 6.1 - Preference points claim form in terms of the Preferential Procurement Regulations 2017	
d.	SBD 8 - Declaration of Bidder's Past Supply Chain Management Practices.	
e.	SBD 9 - Certificate of Independent Bid Determination	
f.	Fully completed pricing schedule	
g.	Proof of registration on Central Supplier Database (detailed report not summary report).	
h.	A Letter of Good Standing, issued by the Compensation Fund in terms of the Compensation for Occupational Injuries and Diseases Act, 1993 must be attached. The certificate must be valid by closing date of the bid. The letter of intention to issue a letter of good standing by the Compensation Commissioner is not acceptable.	

i.	Valid UIF compliance certificate issued by the Department of Labour. The certificate must me valid as at the closing date of the bid.	
j.	Letter of approval by Executing Authority to do business if the entity has member / members who is / are a Government employees.	
k.	List of products offered sourced from third party together with signed agreement between the bidder and the third party (Manufacturer)	
l.	Letter of undertaking from the third party for supply assurance/guarantee	
m.	Certified copy of the SAHPRA license as a Manufacturer and/or Distributor of medical device	
n.	Certified copy of ISO standard: 11070:2014 for CVC products. Certificate of compliance with ISO Standards as stated in the item specification for items that require such compliance	

N.B. BIDDERS WHO FAIL TO ATTACH ONE OF THE COMPULSORY REQUIREMENTS ABOVE WILL BE AUTOMATICALLY DISQUALIFIED

Section	Supporting Returnable Documents	Attached YES / NO
a.	Original or certified copy of B-BBEE Verification Certificate from a Verification Agency accredited by the South African National Accreditation System (SANAS), or sworn affidavit indicating the level of preferential points to be claimed as contemplated on the amended Code of good practice of the B-BBEE certification sworn affidavit only applies to Exempted Micro Enterprise (EME). Bidder in a joint venture, partnership consortium must attach consolidated original or certified copy of the B-BBEE certificate or sworn affidavit if EME. The date on the certificate copies must not be older than one (01) months as at closing date of the bid.	
b.	Sworn affidavit reflecting the annual turnover of the bidder in order to determine whether the bidder is an EME.	

SECTION B - BIDDING PROCESS IN TERMS OF PPPFA

1. Preferential points in terms of PPPFA

The contract shall be awarded in terms of the Preferential Procurement Policy Framework Act, 2000 (Act 5 of 2000) and Regulation of 2017, responsive bids shall be evaluated and adjudicated by the Mpumalanga Department of Health on the 80/20 preference point system in terms of which points are awarded to bidder (s) on the basis of:

Point allocation for price and B-BBEE status level of contribution:

Price	80
B-BBEE Status level of Contributor	20

A maximum of (20) points shall be awarded to a bidder/s in respect of Broad-based Black Economic Empowerment (BBBEE) contribution contemplated in sub-regulation (2) must be added to the points scored for price as calculated in accordance with sub-regulation (1).

Subject to regulation 7, the contract must be awarded to the tenderer who scores the highest total number of points.
Subject to sub-regulation (3) points must be awarded to a tenderer for attaining their BBBEE status level of contributor in accordance with the table below:

B-BBEE Status level of Contributor	Number of points
1	20
2	18
3	14
4	12
5	8
6	6
7	4
8	2
Non-compliant contributor	0

Failure on the part of a bidder to fill in and / or to sign the SBD 6.1 form and submit a BBBEE Verification Certificate from a Verification Agency accredited by the South African National Accreditation System (SANAS) together with the bid, will be interpreted to mean that preference points for BBBEE status level of contribution are not claimed.

TAX CLEARANCE

TCC 001

Application for a Tax Clearance Certificate

Purpose

Select the applicable option ..... Tenders | Good standing

If "Good standing", please state the purpose of this application

Particulars of applicant

Name/Legal name (Initials & Surname or registered name)

Trading name (if applicable)

ID/Passport no

Company/Close Corp. registered no

Income Tax ref no

PAYE ref no 7

VAT registration no 4

SDL ref no L

Customs code

UIF ref no U

Telephone no

Fax no

E-mail address

Physical address

Postal address

Particulars of representative (Public Officer/Trustee/Partner)

Surname

First names

ID/Passport no

Income Tax ref no

Telephone no

Fax no

E-mail address

Physical address

Particulars of tender (If applicable)

Tender number

Estimated Tender amount R ,

Expected duration of the tender year(s)

Particulars of the 3 largest contracts previously awarded

Date started	Date finalised	Principal	Contact person	Telephone number	Amount

Audit

Are you currently aware of any Audit investigation against you/the company? ..... YES NO

If "YES" provide details

Appointment of representative/agent (Power of Attorney)

I the undersigned confirm that I require a Tax Clearance Certificate in respect of Tenders or Goodstanding.

I hereby authorise and instruct to apply to and receive from SARS the applicable Tax Clearance Certificate on my/our behalf.

Signature of representative/agent Date

Name of representative/ agent

Declaration

I declare that the information furnished in this application as well as any supporting documents is true and correct in every respect.

Signature of applicant/Public Officer Date

Name of applicant/ Public Officer

Notes:

1. It is a serious offence to make a false declaration.
2. Section 75 of the Income Tax Act, 1962, states: Any person who
   (a) fails or neglects to furnish, file or submit any return or document as and when required by or under this Act; or
   (b) without just cause shown by him, refuses or neglects to-
      (i) furnish, produce or make available any information, documents or things;
      (ii) reply to or answer truly and fully, any questions put to him ...

   As and when required in terms of this Act ... shall be guilty of an offence ...
3. **SARS will, under no circumstances, issue a Tax Clearance Certificate unless this form is completed in full.**
4. Your Tax Clearance Certificate will only be issued on presentation of your South African Identity Document or Passport (Foreigners only) as applicable.

SBD 4

DECLARATION OF INTEREST

1. Any legal person, including persons employed by the state[1], 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 a price quotation, advertised competitive bid, limited bid or proposal). 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 (director, trustee, shareholder[2]): .....

2.4 Company Registration Number: .....

2.5 Tax Reference Number: .....

2.6 VAT Registration Number: .....

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

[1]"State" means –

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

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

2.7	Are you or any person connected with the bidder presently employed by the state?	**YES / NO**
2.7.1	If so, furnish the following particulars:	
	Name of person / director / trustee / shareholder/ member: Name of state institution at which you or the person connected to the bidder is employed : Position occupied in the state institution:	....
	Any other particulars:	
2.7.2	If you are presently employed by the state, did you obtain the appropriate authority to undertake remunerative work outside employment in the public sector?	**YES / NO**
2.7.2.1	If yes, did you attached proof of such authority to the bid document? (Note: Failure to submit proof of such authority, where applicable, may result in the disqualification of the bid.	**YES / NO**
2.7.2.2	If no, furnish reasons for non-submission of such proof:	
2.8	Did you or your spouse, or any of the company's directors / trustees / shareholders / members or their spouses conduct business with the state in the previous twelve months?	**YES / NO**
2.8.1	If so, furnish particulars:	
2.9	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**

2.9.1 If so, furnish particulars.

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

2.10 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**

2.10.1 If so, furnish particulars.

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

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

2.11.1 If so, furnish particulars:

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

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

Full Name	Identity Number	Personal Tax Reference Number	State Employee Number / Persal Number

4 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 IN TERMS OF PARAGRAPH 23 OF THE GENERAL CONDITIONS OF CONTRACT SHOULD THIS DECLARATION PROVE TO BE FALSE.

.....	.....
Signature	Date
.....	.....
Position	Name of bidder

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 (B-BBEE) Status Level of Contribution

NB: BEFORE COMPLETING THIS FORM, BIDDERS MUST STUDY THE GENERAL CONDITIONS, DEFINITIONS AND DIRECTIVES APPLICABLE IN RESPECT OF B-BBEE, 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

a) The value of this bid is estimated to not exceed R50 000 000 (all applicable taxes included) and therefore the **80/20** preference point system shall be applicable; or

b) Either the 80/20 or 90/10 preference point system will be applicable to this tender (*delete whichever is not applicable for this tender*).

1.3 Points for this bid shall be awarded for:

(a) Price; and

(b) B-BBEE Status Level of Contributor.

1.4 The maximum points for this bid are allocated as follows:

	POINTS
PRICE	80
B-BBEE STATUS LEVEL OF CONTRIBUTOR	20
Total points for Price and B-BBEE must not exceed	**100**

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

1.6 The purchaser 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 purchaser.

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) **"Broad-Based Black Economic Empowerment Act"** means the Broad-Based Black Economic Empowerment Act, 2003 (Act No. 53 of 2003);

(e) "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;

(f) **"functionality"** means the ability of a tenderer to provide goods or services in accordance with specifications as set out in the tender documents.

(g) **"prices"** includes all applicable taxes less all unconditional discounts;

(h) **"proof of B-BBEE status level of contributor"** means:

1) B-BBEE Status level certificate issued by an authorized body or person;

2) A sworn affidavit as prescribed by the B-BBEE Codes of Good Practice;

3) Any other requirement prescribed in terms of the B-BBEE Act;

(i) **"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;

(j) **"rand value"** means the total estimated value of a contract in Rand, calculated at the time of bid invitation, and includes all applicable taxes;

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**

$$Ps = 80\left(1 - \frac{Pt - P\min}{P\min}\right) \quad \text{or} \quad Ps = 90\left(1 - \frac{Pt - P\min}{P\min}\right)$$

Where

Ps = Points scored for price of bid under consideration

Pt = Price of bid under consideration

Pmin = Price of lowest acceptable bid

4. POINTS AWARDED FOR B-BBEE 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 B-BBEE status level of contribution in accordance with the table below:

B-BBEE 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 B-BBEE Status Level of Contribution must complete the following:

6. B-BBEE STATUS LEVEL OF CONTRIBUTOR CLAIMED IN TERMS OF PARAGRAPHS 1.4 AND 4.1

6.1 B-BBEE Status Level of Contributor: . = ...(maximum of 10 or 20 points)

(Points claimed in respect of paragraph 7.1 must be in accordance with the table reflected in paragraph 4.1 and must be substantiated by relevant proof of B-BBEE status level of contributor.

7. SUB-CONTRACTING

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

(***Tick applicable box***)

YES		NO	

7.1.1 If yes, indicate:

i) What percentage of the contract will be subcontracted.....%
ii) The name of the sub-contractor.....
iii) The B-BBEE status level of the sub-contractor.....
iv) Whether the sub-contractor is an EME or QSE
(Tick applicable box)

YES		NO	

v) 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		
OR		
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 B-BBEE status level of contributor has been claimed or obtained on a

fraudulent basis or any of the conditions of contract have not been fulfilled, the purchaser may, in addition to any other remedy it may have –

(a) disqualify the person from the bidding process;

(b) recover costs, losses or damages it has incurred or suffered as a result of that person's conduct;

(c) cancel the contract and claim any damages which it has suffered as a result of having to make less favourable arrangements due to such cancellation;

(d) recommend that the bidder or contractor, its shareholders and directors, or only the shareholders and directors who acted on a fraudulent basis, be restricted by the National Treasury from obtaining business from any organ of state for a period not exceeding 10 years, after the *audi alteram partem* (hear the other side) rule has been applied; and

(e) forward the matter for criminal prosecution.

WITNESSES

1. .....

2. .....

.....
SIGNATURE(S) OF BIDDERS(S)

DATE: .....

ADDRESS .....

.....

.....

DECLARATION CERTIFICATE FOR LOCAL PRODUCTION AND CONTENT FOR DESIGNATED SECTORS

This Standard Bidding Document (SBD) must form part of all bids invited. It contains general information and serves as a declaration form for local content (local production and local content are used interchangeably).

Before completing this declaration, bidders must study the General Conditions, Definitions, Directives applicable in respect of Local Content as prescribed in the Preferential Procurement Regulations, 2017, the South African Bureau of Standards (SABS) approved technical specification number SATS 1286:2011 (Edition 1) and the Guidance on the Calculation of Local Content together with the Local Content Declaration Templates [Annex C (Local Content Declaration: Summary Schedule), D (Imported Content Declaration: Supporting Schedule to Annex C) and E (Local Content Declaration: Supporting Schedule to Annex C)].

1. General Conditions

1.1. Preferential Procurement Regulations, 2017 (Regulation 8) make provision for the promotion of local production and content.

1.2. Regulation 8.(2) prescribes that in the case of designated sectors, organs of state must advertise such tenders with the specific bidding condition that only locally produced or manufactured goods, with a stipulated minimum threshold for local production and content will be considered.

1.3. Where necessary, for tenders referred to in paragraph 1.2 above, a two stage bidding process may be followed, where the first stage involves a minimum threshold for local production and content and the second stage price and B-BBEE.

1.4. A person awarded a contract in relation to a designated sector, may not sub-contract in such a manner that the local production and content of the overall value of the contract is reduced to below the stipulated minimum threshold.

1.5. The local content (LC) expressed as a percentage of the bid price must be calculated in accordance with the SABS approved technical specification number SATS 1286: 2011 as follows:

$$LC = [1 - x / y] * 100$$

Where
x is the imported content in Rand
y is the bid price in Rand excluding value added tax (VAT)

Prices referred to in the determination of x must be converted to Rand (ZAR) by using the exchange rate published by South African Reserve Bank (SARB) at 12:00 on the date of advertisement of the bid as indicated in paragraph 4.1 below.

The SABS approved technical specification number SATS 1286:2011 is accessible on http:/www.thedti.gov.za/industrial development/ip.jsp at no cost.

1.6. A bid may be disqualified if this Declaration Certificate and the Annex C (Local Content Declaration: Summary Schedule) are not submitted as part of the bid documentation;

2. **The stipulated minimum threshold(s) for local production and content (refer to Annex A of SATS 1286:2011) for this bid is/are as follows:**

Description of services, works or goods	Stipulated minimum threshold
_____	_____%
_____	_____%
_____	_____%

3. Does any portion of the goods or services offered have any imported content?
(*Tick applicable box*)

YES		NO	

3..1 If yes, the rate(s) of exchange to be used in this bid to calculate the local content as prescribed in paragraph 1.5 of the general conditions must be the rate(s) published by SARB for the specific currency at 12:00 on the date of advertisement of the bid.

The relevant rates of exchange information is accessible on www.reservebank.co.za

Indicate the rate(s) of exchange against the appropriate currency in the table below (refer to Annex A of SATS 1286:2011):

Currency	Rates of exchange
US Dollar	
Pound Sterling	
Euro	
Yen	
Other	

NB: Bidders must submit proof of the SARB rate (s) of exchange used.

4. Where, after the award of a bid, challenges are experienced in meeting the stipulated minimum threshold for local content the dti must be informed accordingly in order for the dti to verify and in consultation with the AO/AA provide directives in this regard.

LOCAL CONTENT DECLARATION (REFER TO ANNEX B OF SATS 1286:2011)

LOCAL CONTENT DECLARATION BY CHIEF FINANCIAL OFFICER OR OTHER LEGALLY RESPONSIBLE PERSON NOMINATED IN WRITING BY THE CHIEF EXECUTIVE OR SENIOR MEMBER/PERSON WITH MANAGEMENT RESPONSIBILITY (CLOSE CORPORATION, PARTNERSHIP OR INDIVIDUAL)

IN RESPECT OF BID NO. .....

ISSUED BY: (Procurement Authority / Name of Institution):
.....

NB

1 The obligation to complete, duly sign and submit this declaration cannot be transferred to an external authorized representative, auditor or any other third party acting on behalf of the bidder.

2 Guidance on the Calculation of Local Content together with Local Content Declaration Templates (Annex C, D and E) is accessible on http://www.thdti.gov.za/industrial development/ip.jsp. Bidders should first complete Declaration D. After completing Declaration D, bidders should complete Declaration E and then consolidate the information on Declaration C. **Declaration C should be submitted with the bid documentation at the closing date and time of the bid in order to substantiate the declaration made in paragraph (c) below.** Declarations D and E should be kept by the bidders for verification purposes for a period of at least 5 years. The successful bidder is required to continuously update Declarations C, D and E with the actual values for the duration of the contract.

I, the undersigned, ... (full names), do hereby declare, in my capacity as ..... of .....(name of bidder entity), the following:

(a) The facts contained herein are within my own personal knowledge.

(b) I have satisfied myself that:

(i) the goods/services/works to be delivered in terms of the above-specified bid comply with the minimum local content requirements as specified in the bid, and as measured in terms of SATS 1286:2011; and

(c) The local content percentage (%) indicated below has been calculated using the formula given in clause 3 of SATS 1286:2011, the rates of exchange indicated in paragraph 4.1 above and the information contained in Declaration D and E which has been consolidated in Declaration C:

Bid price, excluding VAT (y)	R
Imported content (x), as calculated in terms of SATS 1286:2011	R
Stipulated minimum threshold for local content (paragraph 3 above)	
Local content %, as calculated in terms of SATS 1286:2011	

If the bid is for more than one product, the local content percentages for each product contained in Declaration C shall be used instead of the table above.
The local content percentages for each product has been calculated using the formula given in clause 3 of SATS 1286:2011, the rates of exchange indicated in paragraph 4.1 above and the information contained in Declaration D and E.

(d) I accept that the Procurement Authority / Institution has the right to request that the local content be verified in terms of the requirements of SATS 1286:2011.

(e) I understand that the awarding of the bid is dependent on the accuracy of the information furnished in this application. I also understand that the submission of incorrect data, or data that are not verifiable as described in SATS 1286:2011, may result in the Procurement Authority / Institution imposing any or all of the remedies as provided for in Regulation 14 of the Preferential Procurement Regulations, 2017

promulgated under the Preferential Policy Framework Act (PPPFA), 2000 (Act No. 5 of 2000).

SIGNATURE: _____ **DATE:** _____

WITNESS No. 1 _____ **DATE:** _____

WITNESS No. 2 _____ **DATE:** _____

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 as companies or persons prohibited from doing business with the public sector? **(Companies or persons who are listed on this database were informed in writing of this restriction by the National Treasury after the *audi alteram partem* rule was applied).**	Yes ☐	No ☐
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)? **To access this Register enter the National Treasury's website, www.treasury.gov.za, click on the icon "Register for Tender Defaulters" or submit your written request for a hard copy of the Register to facsimile number (012) 3265445.**	Yes ☐	No ☐
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 ☐	No ☐
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 ☐	No ☐
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**

SBD 9

CERTIFICATE OF INDEPENDENT BID DETERMINATION

1 This Standard Bidding Document (SBD) must form part of all bids[1] 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).[2] 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 This SBD serves as a certificate of declaration that would be used by institutions to ensure that, when bids are considered, reasonable steps are taken to prevent any form of bid-rigging.

5 In order to give effect to the above, the attached Certificate of Bid Determination (SBD 9) must be completed and submitted with the bid:

[1] Includes price quotations, advertised competitive bids, limited bids and proposals.

[2] 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.

SBD 9

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[3] 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.

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

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

Js914w 2

THE NATIONAL TREASURY

Republic of South Africa

GOVERNMENT PROCUREMENT:

GENERAL CONDITIONS OF CONTRACT

July 2010

GOVERNMENT PROCUREMENT

GENERAL CONDITIONS OF CONTRACT
July 2010

NOTES

The purpose of this document is to:

(i) Draw special attention to certain general conditions applicable to government bids, contracts and orders; and

(ii) To ensure that clients be familiar with regard to the rights and obligations of all parties involved in doing business with government.

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

- The General Conditions of Contract will form part of all bid documents and may not be amended.

- Special Conditions of Contract (SCC) relevant to a specific bid, should be compiled separately for every bid (if (applicable) and will supplement the General Conditions of Contract. Whenever there is a conflict, the provisions in the SCC shall prevail.

TABLE OF CLAUSES

General Conditions of Contract

1. Definitions

1. The following terms shall be interpreted as indicated:

1.1 "Closing time" means the date and hour specified in the bidding documents for the receipt of bids.

1.2 "Contract" means the written agreement entered into between the purchaser and the supplier, as recorded in the contract form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.

1.3 "Contract price" means the price payable to the supplier under the contract for the full and proper performance of his contractual obligations.

1.4 "Corrupt practice" means the offering, giving, receiving, or soliciting of any thing of value to influence the action of a public official in the procurement process or in contract execution.

1.5 "Countervailing duties" are imposed in cases where an enterprise abroad is subsidized by its government and encouraged to market its products internationally.

1.6 "Country of origin" means the place where the goods were mined, grown or produced or from which the services are supplied. Goods are produced when, through manufacturing, processing or substantial and major assembly of components, a commercially recognized new product results that is substantially different in basic characteristics or in purpose or utility from its components.

1.7 "Day" means calendar day.

1.8 "Delivery" means delivery in compliance of the conditions of the contract or order.

1.9 "Delivery ex stock" means immediate delivery directly from stock actually on hand.

1.10 "Delivery into consignees store or to his site" means delivered and unloaded in the specified store or depot or on the specified site in compliance with the conditions of the contract or order, the supplier bearing all risks and charges involved until the supplies are so delivered and a valid receipt is obtained.

1.11 "Dumping" occurs when a private enterprise abroad market its goods on own initiative in the RSA at lower prices than that of the country of origin and which have the potential to harm the local industries in the

RSA.

1.12 "Force majeure" means an event beyond the control of the supplier and not involving the supplier's fault or negligence and not foreseeable. Such events may include, but is not restricted to, acts of the purchaser in its sovereign capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions and freight embargoes.

1.13 "Fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of any bidder, and includes collusive practice among bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to deprive the bidder of the benefits of free and open competition.

1.14 "GCC" means the General Conditions of Contract.

1.15 "Goods" means all of the equipment, machinery, and/or other materials that the supplier is required to supply to the purchaser under the contract.

1.16 "Imported content" means that portion of the bidding price represented by the cost of components, parts or materials which have been or are still to be imported (whether by the supplier or his subcontractors) and which costs are inclusive of the costs abroad, plus freight and other direct importation costs such as landing costs, dock dues, import duty, sales duty or other similar tax or duty at the South African place of entry as well as transportation and handling charges to the factory in the Republic where the supplies covered by the bid will be manufactured.

1.17 "Local content" means that portion of the bidding price which is not included in the imported content provided that local manufacture does take place.

1.18 "Manufacture" means the production of products in a factory using labour, materials, components and machinery and includes other related value-adding activities.

1.19 "Order" means an official written order issued for the supply of goods or works or the rendering of a service.

1.20 "Project site," where applicable, means the place indicated in bidding documents.

1.21 "Purchaser" means the organization purchasing the goods.

1.22 "Republic" means the Republic of South Africa.

1.23 "SCC" means the Special Conditions of Contract.

1.24 "Services" means those functional services ancillary to the supply of the goods, such as transportation and any other incidental services, such as installation, commissioning, provision of technical assistance, training, catering, gardening, security, maintenance and other such

obligations of the supplier covered under the contract.

1.25 "Written" or "in writing" means handwritten in ink or any form of electronic or mechanical writing.

2. Application

2.1 These general conditions are applicable to all bids, contracts and orders including bids for functional and professional services, sales, hiring, letting and the granting or acquiring of rights, but excluding immovable property, unless otherwise indicated in the bidding documents.

2.2 Where applicable, special conditions of contract are also laid down to cover specific supplies, services or works.

2.3 Where such special conditions of contract are in conflict with these general conditions, the special conditions shall apply.

3. General

3.1 Unless otherwise indicated in the bidding documents, the purchaser shall not be liable for any expense incurred in the preparation and submission of a bid. Where applicable a non-refundable fee for documents may be charged.

3.2 With certain exceptions, invitations to bid are only published in the Government Tender Bulletin. The Government Tender Bulletin may be obtained directly from the Government Printer, Private Bag X85, Pretoria 0001, or accessed electronically from www.treasury.gov.za

4. Standards

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

5. Use of contract documents and information; inspection.

5.1 The supplier shall not, without the purchaser's prior written consent, disclose the contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the purchaser in connection therewith, to any person other than a person employed by the supplier in the performance of the contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.

5.2 The supplier shall not, without the purchaser's prior written consent, make use of any document or information mentioned in GCC clause 5.1 except for purposes of performing the contract.

5.3 Any document, other than the contract itself mentioned in GCC clause 5.1 shall remain the property of the purchaser and shall be returned (all copies) to the purchaser on completion of the supplier's performance under the contract if so required by the purchaser.

5.4 The supplier shall permit the purchaser to inspect the supplier's records relating to the performance of the supplier and to have them audited by auditors appointed by the purchaser, if so required by the purchaser.

6. Patent rights

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

7. Performance security

7.1 Within thirty (30) days of receipt of the notification of contract award, the successful bidder shall furnish to the purchaser the performance security of the amount specified in SCC.

7.2 The proceeds of the performance security shall be payable to the purchaser as compensation for any loss resulting from the supplier's failure to complete his obligations under the contract.

7.3 The performance security shall be denominated in the currency of the contract, or in a freely convertible currency acceptable to the purchaser and shall be in one of the following forms:

(a) a bank guarantee or an irrevocable letter of credit issued by a reputable bank located in the purchaser's country or abroad, acceptable to the purchaser, in the form provided in the bidding documents or another form acceptable to the purchaser; or

(b) a cashier's or certified cheque

7.4 The performance security will be discharged by the purchaser and returned to the supplier not later than thirty (30) days following the date of completion of the supplier's performance obligations under the contract, including any warranty obligations, unless otherwise specified in SCC.

8. Inspections, tests and analyses

8.1 All pre-bidding testing will be for the account of the bidder.

8.2 If it is a bid condition that supplies to be produced or services to be rendered should at any stage during production or execution or on completion be subject to inspection, the premises of the bidder or contractor shall be open, at all reasonable hours, for inspection by a representative of the Department or an organization acting on behalf of the Department.

8.3 If there are no inspection requirements indicated in the bidding documents and no mention is made in the contract, but during the contract period it is decided that inspections shall be carried out, the purchaser shall itself make the necessary arrangements, including payment arrangements with the testing authority concerned.

8.4 If the inspections, tests and analyses referred to in clauses 8.2 and 8.3 show the supplies to be in accordance with the contract requirements, the cost of the inspections, tests and analyses shall be defrayed by the purchaser.

8.5 Where the supplies or services referred to in clauses 8.2 and 8.3 do not comply with the contract requirements, irrespective of whether such supplies or services are accepted or not, the cost in connection with these inspections, tests or analyses shall be defrayed by the supplier.

8.6 Supplies and services which are referred to in clauses 8.2 and 8.3 and which do not comply with the contract requirements may be rejected.

8.7 Any contract supplies may on or after delivery be inspected, tested or

analyzed and may be rejected if found not to comply with the requirements of the contract. Such rejected supplies shall be held at the cost and risk of the supplier who shall, when called upon, remove them immediately at his own cost and forthwith substitute them with supplies which do comply with the requirements of the contract. Failing such removal the rejected supplies shall be returned at the suppliers cost and risk. Should the supplier fail to provide the substitute supplies forthwith, the purchaser may, without giving the supplier further opportunity to substitute the rejected supplies, purchase such supplies as may be necessary at the expense of the supplier.

8.8 The provisions of clauses 8.4 to 8.7 shall not prejudice the right of the purchaser to cancel the contract on account of a breach of the conditions thereof, or to act in terms of Clause 23 of GCC.

9. Packing

9.1 The supplier shall provide such packing of the goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packing, case size and weights shall take into consideration, where appropriate, the remoteness of the goods' final destination and the absence of heavy handling facilities at all points in transit.

9.2 The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the contract, including additional requirements, if any, specified in SCC, and in any subsequent instructions ordered by the purchaser.

10. Delivery and documents

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

10.2 Documents to be submitted by the supplier are specified in SCC.

11. Insurance

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

12. Transportation

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

13. Incidental services

13.1 The supplier may be required to provide any or all of the following services, including additional services, if any, specified in SCC:

(a) performance or supervision of on-site assembly and/or commissioning of the supplied goods;
(b) furnishing of tools required for assembly and/or maintenance of the supplied goods;
(c) furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied goods;

(d) performance or supervision or maintenance and/or repair of the supplied goods, for a period of time agreed by the parties, provided that this service shall not relieve the supplier of any warranty obligations under this contract; and

(e) training of the purchaser's personnel, at the supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied goods.

13.2 Prices charged by the supplier for incidental services, if not included in the contract price for the goods, shall be agreed upon in advance by the parties and shall not exceed the prevailing rates charged to other parties by the supplier for similar services.

14. Spare parts

14.1 As specified in SCC, the supplier may be required to provide any or all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the supplier:

(a) such spare parts as the purchaser may elect to purchase from the supplier, provided that this election shall not relieve the supplier of any warranty obligations under the contract; and

(b) in the event of termination of production of the spare parts:

(i) Advance notification to the purchaser of the pending termination, in sufficient time to permit the purchaser to procure needed requirements; and

(ii) following such termination, furnishing at no cost to the purchaser, the blueprints, drawings, and specifications of the spare parts, if requested.

15. Warranty

15.1 The supplier warrants that the goods supplied under the contract are new, unused, of the most recent or current models, and that they incorporate all recent improvements in design and materials unless provided otherwise in the contract. The supplier further warrants that all goods supplied under this contract shall have no defect, arising from design, materials, or workmanship (except when the design and/or material is required by the purchaser's specifications) or from any act or omission of the supplier, that may develop under normal use of the supplied goods in the conditions prevailing in the country of final destination.

15.2 This warranty shall remain valid for twelve (12) months after the goods, or any portion thereof as the case may be, have been delivered to and accepted at the final destination indicated in the contract, or for eighteen (18) months after the date of shipment from the port or place of loading in the source country, whichever period concludes earlier, unless specified otherwise in SCC.

15.3 The purchaser shall promptly notify the supplier in writing of any claims arising under this warranty.

15.4 Upon receipt of such notice, the supplier shall, within the period specified in SCC and with all reasonable speed, repair or replace the defective goods or parts thereof, without costs to the purchaser.

15.5 If the supplier, having been notified, fails to remedy the defect(s) within the period specified in SCC, the purchaser may proceed to take

such remedial action as may be necessary, at the supplier's risk and expense and without prejudice to any other rights which the purchaser may have against the supplier under the contract.

16. Payment

16.1 The method and conditions of payment to be made to the supplier under this contract shall be specified in SCC.

16.2 The supplier shall furnish the purchaser with an invoice accompanied by a copy of the delivery note and upon fulfillment of other obligations stipulated in the contract.

16.3 Payments shall be made promptly by the purchaser, but in no case later than thirty (30) days after submission of an invoice or claim by the supplier.

16.4 Payment will be made in Rand unless otherwise stipulated in SCC.

17. Prices

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

18. Contract amendments

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

19. Assignment

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

20. Subcontracts

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

21. Delays in the supplier's performance

21.1 Delivery of the goods and performance of services shall be made by the supplier in accordance with the time schedule prescribed by the purchaser in the contract.

21.2 If at any time during performance of the contract, the supplier or its subcontractor(s) should encounter conditions impeding timely delivery of the goods and performance of services, the supplier shall promptly notify the purchaser in writing of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the supplier's notice, the purchaser shall evaluate the situation and may at his discretion extend the supplier's time for performance, with or without the imposition of penalties, in which case the extension shall be ratified by the parties by amendment of contract.

21.3 No provision in a contract shall be deemed to prohibit the obtaining of supplies or services from a national department, provincial department, or a local authority.

21.4 The right is reserved to procure outside of the contract small quantities or to have minor essential services executed if an emergency arises, the

supplier's point of supply is not situated at or near the place where the supplies are required, or the supplier's services are not readily available.

21.5 Except as provided under GCC Clause 25, a delay by the supplier in the performance of its delivery obligations shall render the supplier liable to the imposition of penalties, pursuant to GCC Clause 22, unless an extension of time is agreed upon pursuant to GCC Clause 21.2 without the application of penalties.

21.6 Upon any delay beyond the delivery period in the case of a supplies contract, the purchaser shall, without canceling the contract, be entitled to purchase supplies of a similar quality and up to the same quantity in substitution of the goods not supplied in conformity with the contract and to return any goods delivered later at the supplier's expense and risk, or to cancel the contract and buy such goods as may be required to complete the contract and without prejudice to his other rights, be entitled to claim damages from the supplier.

22. Penalties

22.1 Subject to GCC Clause 25, if the supplier fails to deliver any or all of the goods or to perform the services within the period(s) specified in the contract, the purchaser shall, without prejudice to its other remedies under the contract, deduct from the contract price, as a penalty, a sum calculated on the delivered price of the delayed goods or unperformed services using the current prime interest rate calculated for each day of the delay until actual delivery or performance. The purchaser may also consider termination of the contract pursuant to GCC Clause 23.

23. Termination for default

23.1 The purchaser, without prejudice to any other remedy for breach of contract, by written notice of default sent to the supplier, may terminate this contract in whole or in part:

(a) if the supplier fails to deliver any or all of the goods within the period(s) specified in the contract, or within any extension thereof granted by the purchaser pursuant to GCC Clause 21.2;

(b) if the Supplier fails to perform any other obligation(s) under the contract; or

(c) if the supplier, in the judgment of the purchaser, has engaged in corrupt or fraudulent practices in competing for or in executing the contract.

23.2 In the event the purchaser terminates the contract in whole or in part, the purchaser may procure, upon such terms and in such manner as it deems appropriate, goods, works or services similar to those undelivered, and the supplier shall be liable to the purchaser for any excess costs for such similar goods, works or services. However, the supplier shall continue performance of the contract to the extent not terminated.

23.3 Where the purchaser terminates the contract in whole or in part, the purchaser may decide to impose a restriction penalty on the supplier by prohibiting such supplier from doing business with the public sector for a period not exceeding 10 years.

23.4 If a purchaser intends imposing a restriction on a supplier or any

person associated with the supplier, the supplier will be allowed a time period of not more than fourteen (14) days to provide reasons why the envisaged restriction should not be imposed. Should the supplier fail to respond within the stipulated fourteen (14) days the purchaser may regard the intended penalty as not objected against and may impose it on the supplier.

23.5 Any restriction imposed on any person by the Accounting Officer / Authority will, at the discretion of the Accounting Officer / Authority, also be applicable to any other enterprise or any partner, manager, director or other person who wholly or partly exercises or exercised or may exercise control over the enterprise of the first-mentioned person, and with which enterprise or person the first-mentioned person, is or was in the opinion of the Accounting Officer / Authority actively associated.

23.6 If a restriction is imposed, the purchaser must, within five (5) working days of such imposition, furnish the National Treasury, with the following information:

(i) the name and address of the supplier and / or person restricted by the purchaser;
(ii) the date of commencement of the restriction
(iii) the period of restriction; and
(iv) the reasons for the restriction.

These details will be loaded in the National Treasury's central database of suppliers or persons prohibited from doing business with the public sector.

23.7 If a court of law convicts a person of an offence as contemplated in sections 12 or 13 of the Prevention and Combating of Corrupt Activities Act, No. 12 of 2004, the court may also rule that such person's name be endorsed on the Register for Tender Defaulters. When a person's name has been endorsed on the Register, the person will be prohibited from doing business with the public sector for a period not less than five years and not more than 10 years. The National Treasury is empowered to determine the period of restriction and each case will be dealt with on its own merits. According to section 32 of the Act the Register must be open to the public. The Register can be perused on the National Treasury website.

24. Anti-dumping and countervailing duties and rights

24.1 When, after the date of bid, provisional payments are required, or anti-dumping or countervailing duties are imposed, or the amount of a provisional payment or anti-dumping or countervailing right is increased in respect of any dumped or subsidized import, the State is not liable for any amount so required or imposed, or for the amount of any such increase. When, after the said date, such a provisional payment is no longer required or any such anti-dumping or countervailing right is abolished, or where the amount of such provisional payment or any such right is reduced, any such favourable difference shall on demand be paid forthwith by the contractor to the State or the State may deduct such amounts from moneys (if any) which may otherwise be due to the contractor in regard to supplies or services which he delivered or rendered, or is to deliver or render in terms of the contract or any other contract or any other amount which

may be due to him

25. Force Majeure

25.1 Notwithstanding the provisions of GCC Clauses 22 and 23, the supplier shall not be liable for forfeiture of its performance security, damages, or termination for default if and to the extent that his delay in performance or other failure to perform his obligations under the contract is the result of an event of force majeure.

25.2 If a force majeure situation arises, the supplier shall promptly notify the purchaser in writing of such condition and the cause thereof. Unless otherwise directed by the purchaser in writing, the supplier shall continue to perform its obligations under the contract as far as is reasonably practical, and shall seek all reasonable alternative means for performance not prevented by the force majeure event.

26. Termination for insolvency

26.1 The purchaser may at any time terminate the contract by giving written notice to the supplier if the supplier becomes bankrupt or otherwise insolvent. In this event, termination will be without compensation to the supplier, provided that such termination will not prejudice or affect any right of action or remedy which has accrued or will accrue thereafter to the purchaser.

27. Settlement of Disputes

27.1 If any dispute or difference of any kind whatsoever arises between the purchaser and the supplier in connection with or arising out of the contract, the parties shall make every effort to resolve amicably such dispute or difference by mutual consultation.

27.2 If, after thirty (30) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the purchaser or the supplier may give notice to the other party of his intention to commence with mediation. No mediation in respect of this matter may be commenced unless such notice is given to the other party.

27.3 Should it not be possible to settle a dispute by means of mediation, it may be settled in a South African court of law.

27.4 Mediation proceedings shall be conducted in accordance with the rules of procedure specified in the SCC.

27.5 Notwithstanding any reference to mediation and/or court proceedings herein,

(a) the parties shall continue to perform their respective obligations under the contract unless they otherwise agree; and
(b) the purchaser shall pay the supplier any monies due the supplier.

28. Limitation of liability

28.1 Except in cases of criminal negligence or willful misconduct, and in the case of infringement pursuant to Clause 6;

(a) the supplier shall not be liable to the purchaser, whether in contract, tort, or otherwise, for any indirect or consequential loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the supplier to pay penalties and/or damages to the purchaser; and

(b) the aggregate liability of the supplier to the purchaser, whether under the contract, in tort or otherwise, shall not exceed the total contract price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment.

29. Governing language

29.1 The contract shall be written in English. All correspondence and other documents pertaining to the contract that is exchanged by the parties shall also be written in English.

30. Applicable law

30.1 The contract shall be interpreted in accordance with South African laws, unless otherwise specified in SCC.

31. Notices

31.1 Every written acceptance of a bid shall be posted to the supplier concerned by registered or certified mail and any other notice to him shall be posted by ordinary mail to the address furnished in his bid or to the address notified later by him in writing and such posting shall be deemed to be proper service of such notice

31.2 The time mentioned in the contract documents for performing any act after such aforesaid notice has been given, shall be reckoned from the date of posting of such notice.

32. Taxes and duties

32.1 A foreign supplier shall be entirely responsible for all taxes, stamp duties, license fees, and other such levies imposed outside the purchaser's country.

32.2 A local supplier shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted goods to the purchaser.

32.3 No contract shall be concluded with any bidder whose tax matters are not in order. Prior to the award of a bid the Department must be in possession of a tax clearance certificate, submitted by the bidder. This certificate must be an original issued by the South African Revenue Services.

33. National Industrial Participation (NIP) Programme

33.1 The NIP Programme administered by the Department of Trade and Industry shall be applicable to all contracts that are subject to the NIP obligation.

34 Prohibition of Restrictive practices

34.1 In terms of section 4 (1) (b) (iii) of the Competition Act No. 89 of 1998, as amended, an agreement between, or concerted practice by, firms, or a decision by an association of firms, is prohibited if it is between parties in a horizontal relationship and if a bidder (s) is / are or a contractor(s) was / were involved in collusive bidding (or bid rigging).

34.2 If a bidder(s) or contractor(s), based on reasonable grounds or evidence obtained by the purchaser, has / have engaged in the restrictive practice referred to above, the purchaser may refer the matter to the Competition Commission for investigation and possible imposition of administrative penalties as contemplated in the Competition Act No. 89 of 1998.

34.3 If a bidder(s) or contractor(s), has / have been found guilty by the Competition Commission of the restrictive practice referred to above, the purchaser may, in addition and without prejudice to any other remedy provided for, invalidate the bid(s) for such item(s) offered, and / or terminate the contract in whole or part, and / or restrict the bidder(s) or contractor(s) from conducting business with the public sector for a period not exceeding ten (10) years and / or claim damages from the bidder(s) or contractor(s) concerned.