

Government of Nepal
Ministry of Health
Department of Health Services
Logistics Management Division (LMD)
Teku, Kathmandu, Nepal

BIDDING DOCUMENT for Procurement of Medical & Surgical Instruments for PHC

IFB No: DOHS/G/NCB-155/2016-17

Issued on:
Issued to:
Dispatch no:
Date:



Abbreviations

BDS	Bid Data Sheet
BD	Bidding Document
DCS	Delivery and Completion Schedule
DP	.Development Partner
EQC	. Evaluation and Qualification Criteria
GCC	. General Conditions of Contract
GoN	Government of Nepal
ICC	International Chamber of Commerce
IFB	. Invitation for Bids
Incoterms	. International Commercial Terms
ITB	. Instructions to Bidders
LGRS	. List of Goods and Related Services
NCB	. National Competitive Bidding
PAN	Permanent Account Number
PPMO	Public Procurement Monitoring Office
SBD	Standard Bidding Document
SBQ	Schedule of Bidder Qualifications
SCC	Special Conditions of Contract
SR	Schedule of Requirements
TS	Technical Specifications
UNCITRAL	.United Nations Commission on International Trade Law
VAT	Value Added Tay



Table of Contents

Government of Nepal Ministry of Health	3
INVITATION FOR BIDS	3
Section I. Instructions to Bidders	5
Section II. Bid Data Sheet	25
Section III. Evaluation and Qualification Criteria	32
Section IV. Bidding Forms	35
Price Schedules	45
Section V. Schedule of Requirements	50
Section VIII. Contract Forms	127



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Pachali, Teku, Kathmandu

INVITATION FOR BIDS

No: DOHS/G/NCB-155/2016-17

Procurement of Medical & Surgical Instruments for PHC

Date of publication: 19..Falgun 2073 (02 March-2017)

- Government of Nepal has allocated the fund and intends to apply part of the funds to cover eligible
 payments under the Contract for Procurement of Medical & Surgical Instruments For PHC by
 Department of Health Services.
- 2. The Ministry of Health, Department of Health Services, Logistics Management Division invites sealed bids or electronic bids from eligible bidders for the **Procurement of Medical & Surgical Instruments For PHC** as listed below under National Competitive Bidding. Contract shall be done as a single lot
- 3. Interested Eligible Bidders may obtain further information and inspect the bidding documents at the office of the Department of Health Services, Logistics Management Division, Pachali, Teku, Kathmandu, Tel: 01-4261768, Fax: 01-4261413 from 2073-Falgun- 19. (02 March, 2017) to 2073-Chaitra 18- (March 31, 2017) during the office hours or at their own convenience on the LMD website, www.dohslmd.gov.np and PPMO website www.bolpatra.gov.np. Interested bidders must first register on the PPMO website to access the document and choose the electronic bidding procedure.
- 4. Bidding documents may be downloaded from the PPMO website or LMD website or be purchased from the above office of Department of Health Services, Logistics Management Division by eligible Bidders on the submission of a written application, along with the copy of company/firm registration certificate, and upon payment of a non-refundable fee of NPR. 5000.00. The method of payment will be in the form of a cash deposit certificate in Revenue Title no 14227, Office code no. 27-370-11, Account no. Ka-1-1-001 of Rastriya Banijya Bank, Teku Branch. If so requested, the Bidding Documents can also be sent by post/courier services upon payment of additional cost of NRs. 4000. However, the Employer will not be responsible for delay or non-delivery of the documents so sent. Bidding documents can be purchased up to office hours of 2073-Chaitra-18 (31 March 2017).
- 5. For the purpose who choose to submit their bid electronically through e-procurement section of PPMO website: http://www.bolpatra.gov.np, the bidders may either purchase the hard copy of bidding documents or may choose to download the necessary part of bidding documents, prepare their bids and submit their electronic bids as specified in the Instructions to Bidders. In case of bidder who choose to download and submit bid electronically, the bidder shall be required to deposit the cost of bidding document as specified above in the above mentioned Account No. of LMD at Rastriya Banijya Bank, Teku Branch and electronic scanned copy (pdf format) of the Bank deposit voucher/tele transfer receipt shall also be submitted along with the electronic bid files.
- 6. Sealed Bids must be submitted to the above office of Department of Health Services, Logistics Management Division on or before 12:00 hour (Local time) on 2073-Chaitra- 20 (02-April-2017). Alternatively, bidders may submit their bid electronically through e-procurement section in PPMO website: http://www.bolpatra.gov.np, before the above deadline and as specified in the Instructions to Bidders. Documents received after this deadline shall not be accepted.



- 7. Bids shall be opened in the presence of Bidders' representatives who choose to attend at 13:00 hour (Local time) on 2073-Chaitra -20 (02-April-2017). (at the office of Department of Health Services, Logistics Management Division. Bids must be valid for a period of 90 days counting from the day of bid opening that is valid till 2074-Ashadh-16 (30-June-2017). All bids must be accompanied by bid security not less than NPR 16,50,000.00, which shall be valid for minimum 30 days beyond the bid validity period i.e. valid upto 2074- Shrawan-15 (30 Jully-2017).
- 8. If the last date of purchasing, submission and opening falls on a government holiday then the next working day shall be considered the last day. In such a case the bid validity and bid security validity shall be recognized with effect from the original bid submission deadline unless amended.
- 9. The Purchaser reserves the right to accept or reject, wholly or partly any or all the bids without assigning reason, whatsoever.
- 10. Particulars of the requirement are:

SN	Items	Quantity
1	Delivery set	240
2	Episiotomy set	240
3	Peri light	240
4	Suction (electric)	120
5	Nebulizer	120
6	Otoscope	120
7	IUD & Implant insertion and removal set	240
8	Dental Instrument Set	120
9	I/D Set	240
10	Suture Set	240
11	BP set	600
12	Stethoscope	600
13	Suture removal set	240
14	Dressing Set	240
15	Protoscope	120
16	Foetoscope, Plastic	480
17	Digital thermometer	600
18	Tongue depresor	360
19	Weighing machine (digital)	240
20	Laryngoscope	120
21	LSCS set	240



Section I. Instructions to Bidders

Table of Contents

A.	Ge	neral	7
	1.	Scope of Bid	7 7
	2.	Source of Funds	7
	3.	Fraud and Corruption	7
	4.	Eligible Bidders	9
	5.	Eligible Goods and Related Services	11
	6.	SiteVisit	12
B.	Co	ntents of Bidding Document	13
	7.	Sections of the Bidding Document	13
	8.	Clarification of Bidding Document/Pre-bid meeting	13
	9.	Amendment of Bidding Document	14
C.	Pre	eparation of Bids	14
	10.	Cost of Bidding	12
	11.	Language of Bid	12
	12.	Documents Comprising the Bid	12
	13.	Bid Submission Sheet and Price Schedules	13
	14.	Alternative Bids	13
	15.	Bid Prices and Discounts	13
	16.	Currencies of Bid	14
	17.	Documents Establishing the Eligibility of the Bidder	14
	18.	Documents Establishing the Conformity of the Goods and Related Services	
		to the Bidding Document	14
	19.	Documents Establishing the Qualifications of the Bidder	14
	20.	Period of Validity of Bids	15
	21.	Bid Security	15
	22.	Format and Signing of Bid	16
D.	Sul	omission and Opening of Bids	17
	23.	Sealing and Marking of Bids	17
	24.	Deadline for Submission of Bids	17
	25.	Late Bids	17
	26.	Withdrawal, or Modification of Bids	18
	27.	Bid Opening	18



E.Evaluation and Comparison of Bids		
28.	Confidentiality	19
29.	Clarification of Bids	20
30.	Deviations, Reservations, and Omissions	20
31.	Determination of Responsiveness	20
32.	Non-material Non-conformi-ties	20
33.	Correction of Arithmetical Errors	21
34.	Domestic Preference	21
35.	Evaluation and Comparison of Bids	21
36.	Post-qualification of the Bidder	21
37.	Purchaser's Right to Accept Any Bid, and to Reject Any or All Bids	22
F.Award	of Contract	22
38.	Award Criteria	22
39.	Purchaser's Right to Vary Quantities at Time of Award	22
40.	Notification of Intention to Award	22
41.	Performance Security	22
42.	Signing of Contract	23
43.	Complaint and Review	23



Section I. Instructions to Bidders

A. General

1. Scope of Bid

- 1.1 The Purchaser *indicated in the BDS* issues this Bidding Document for the supply of Goods and Related Services incidental thereto as specified in Section V, Schedule of Requirements.
- 1.2 Throughout this Bidding Document :
 - (a) the term "in writing" means communicated in written form with proof of receipt;
 - (b) if the context so requires, singular means plural and vice versa; and
 - (c) "day" means calendar day.

2. Source of Funds

- 2.1 GoN Funded: In accordance with its annual program and budget, approved by the GoN, the Purchaser intends to apply a portion of the allocated budget to eligible payments under the contract(s) *indicated in the BDS* for which this Bidding Document is issued.
 - DP Funded: The GoN has applied for or received financing (hereinafter called "funds") from the Development Partner (hereinafter called "the DP") *indicated in the BDS* toward the cost of the project *named in the BDS*. The GoN intends to apply a portion of the funds to eligible payments under the contract(s) for which this Bidding Document is issued.
- 2.2 DP Funded: Payment by the DP will be made only at the request of the GoN and upon approval by the DP in accordance with the terms and conditions of the financing agreement between the GoN and the DP (hereinafter called the "Loan Agreement"), and will be subject in all respects to the terms and conditions of that Loan Agreement. No party other than the GoN shall derive any rights from the Loan Agreement or have any claim to the funds.
- 2.3 Public Entity's Resources Funded.

3. Fraud and Corruption

- 3.1 The Government of Nepal (GON) requires that the Purchasing Entities as well as bidders, suppliers, and contractors and their subcontractors under GoN/DP-financed contracts, observe the highest standard of ethics during the procurement and execution of such contracts. In pursuance of this policy, this bidding document;
 - (a) defines, for the purposes of this provision, the terms set forth below as follows:
 - (i) "corrupt practice" means the offering, giving, receiving, or soliciting, directly or indirectly, anything of value to influence improperly the actions of another party;
 - (ii) "fraudulent practice" means any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;



- (iii) "coercive practice" means impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party;
- (iv) "Collusive practice" means an arrangement between two or more parties designed to achieve an improper purpose, including influencing improperly the actions of another party.
- (v) "obstructive practice" means:
 - (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a GoN/DP investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or
 - (bb) Acts intended to materially impede the exercise of the GoN's/DP's inspection and audit rights provided for under sub-clause 3.5 below.
- (b) will reject bid(s) if it determines that the bidder has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive, or obstructive practices in competing for the contract in question;
- (c) will sanction a firm or individual, including declaring ineligible, for a stated period of time, to be awarded a GoN/DP-financed contract if it at any time determines that the firm has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive, or obstructive practices in competing for, or in executing, a GoN/DP-financed contract.
- 3.2 The Bidder shall not carry out or cause to carry out the following acts with an intention to influence the implementation of the procurement process or the procurement agreement:
 - (a) give or propose improper inducement directly or indirectly,
 - (b) distortion or misrepresentation of facts,
 - (c) engaging in corrupt or fraudulent practice or involving in such act,
 - (d) interference in participation of other competing bidders,
 - (e) coercion or threatening directly or indirectly to cause harm to the person or the property of any person to be involved in the procurement proceedings,
 - (f) collusive practice among bidders before or after submission of bids for distribution of works among bidders or fixing artificial/uncompetitive bid pricewith an intention to deprive the Employer the benefit of open competitive bid price,
 - (g) Contacting the Employer with an intention to influence the Employer with regards to the bids or interference of any kind in examination and evaluation of the bids during the



period from the time of opening of the bids until the notification of award of contract.

- 3.3 PPMO, on the recommendation of the Procuring Entity may <u>blacklist</u>a Bidder for its conduct for a period of one (1) to three (3) years on the following grounds and seriousness of the act committed by the bidder:
 - (a) if convicted by a court of law in a criminal offence which disqualifies the Bidder from participating in the contract,
 - (b) If it is proved that the bidder has committed an act contrary to ITB 3.2and ITB 19.5.
- 3.4 A bidder declared blacklisted and ineligible by the GoN, Public procurement Monitoring Office (PPMO), and/or the DP in case of DP funded project, shall be ineligible to bid for a contract during the period of time determined by the GoN, PPMO and/or the DP.
- 3.5 The Supplier shall permit the GoN/DP to inspect the Supplier's accounts and records relating to the performance of the Supplier and to have them audited by auditors appointed by the GoN/DP, if so required by the GoN/DP.
- 3.6 DP Funded: In pursuance of the fraud and corruption policy, the DP.
 - (a) will reject a proposal if it determines that the bidder recommended for award has directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive, or obstructive practices in competing for the contract in question;
 - (b) will cancel the portion of the loan/ credit/ grant allocated to a contract if it determines at any time that representative(s) of the GoN or of a beneficiary of the fund engaged in corrupt, fraudulent, collusive, or coercive practices during the procurement or the execution of that contract, without the GoN having taken timely and appropriate action satisfactory to the DP to address such practices when they occur.

4. Eligible Bidders

- 4.1 This Invitation for Bids is open to eligible Bidders from all countries, except for any *specified in the BDS*.
- 4.2 A Bidder may be a natural person, private entity, government-owned entity (subject to ITB 4.4) or any combination of them with a formal intent to enter into an agreement or under an existing agreement in the form of a Joint Venture (JV). In the case of a JV:
 - (a) all parties to the JV shall be jointly and severally liable; and
 - (b) a JV shall nominate a representative who shall have the authority to conduct all businesses for and on behalf of any and all the parties of the JV during the bidding process and, in the event the JV is awarded the Contract, during contract execution.
- 4.3 A Bidder shall not have a conflict of interest. All Bidders found to have a conflict of interest shall be disqualified. A Bidder may be considered to be in a conflict of interest with one or more



parties in this bidding process if, including but not limited to:

- (a) have controlling shareholders in common;
- (b) receive or have received any direct or indirect subsidy from any of them;
- (c) have the same legal representative for purposes of this Bid;
- (d) have a relationship with each other, directly or through common third parties, that puts them in a position to have access to information about or influence on the Bid of another Bidder, or influence the decisions of the Purchaser regarding this bidding process;
- (e) a Bidder participates in more than one bid in this bidding process. Participation by a Bidder in more than one Bid will result in the disqualification of all Bids in which it is involved. However, this does not limit the inclusion of the same subcontractor, not otherwise participating as a Bidder, in more than one bid; or
 - a Bidder or any of its affiliates participated as a consultant in the preparation of the design or technical specifications of the goods and services that are the subject of the bid.
- 4.4 A Bidder that is under a declaration of ineligibility by the GoN/DP in accordance with ITB 3.4, at the date of the deadline for bid submission or thereafter, shall be disqualified.
- 4.5 A GoN-owned enterprise may also participate in the bid if it is legally and financially autonomous, it operates under commercial law, and it is not dependent agency of the Purchaser.
- 4.6 Bidders shall provide such evidence of their continued eligibility satisfactory to the Purchaser, as the Purchaser shall reasonably request.
- 4.7 Firms shall be excluded in any of the cases, if
 - (a) by an act of compliance with a decision of the United Nations Security Council taken under Chapter VII of the Charter of the United Nations prohibits any import of goods or Contracting of works or services from that country or any payments to persons or entities in that country.
 - (b) DP Funded: as a matter of law or official regulation, GoN prohibits commercial relations with that country, provided that the DP is satisfied that such exclusion does not preclude effective competition for the supply of goods or related services required;
 - (c) DP Funded: a firm has been determined to be ineligible by the DP in relation to their guidelines or appropriate provisions on preventing and combating fraud and corruption in projectsfinanced by them.
- 4.8 A bidder and all parties constituting the Bidder shall have the nationality of an eligible country as defined by the concerned DP for DP funded projects.
- 4.9 The domestic Bidder who has obtained Permanent Account Number (PAN) and Value Added Tax (VAT) registration certificate(s) and Tax clearance certificate or proof of submission of tax return from



the Inland Revenue Office shall only be eligible. The foreign bidder submitting the documents *indicated in the BDS* at the time of bid submission and a declaration to submit the document(s) *indicated in the BDS* at the time of contract agreement shall only be eligible

- 5. Eligible Goods and Related Services
- 5.1 All goods and related services to be supplied under the contract are eligible, unless their origin is from a country *specified in the BDS*.
- 5.2 For purposes of this clause, "origin" means the place where the goods are mined, grown, or produced, or the place from which the related services are supplied
- 5.3 The origin of goods and services is distinct from the nationality of the Bidder.
- 6. Site Visit
- 6.1 For goods contracts requiring installation/ commissioning/ networking or similar services at site, the Bidder, at the Bidder's own responsibility and risk, is encouraged to visit and examine the Site and obtain all information that may be necessary for preparing the Bid and entering into a contract for the supply of goods and related services.
- 6.2 The Bidder should ensure that the Purchaser is informed of the visit in adequate time to allow it to make appropriate arrangements.
- 6.3 The costs of visiting the Site shall be at the Bidder's own expense.

B. Contents of Bidding Document

7. Sections of the Bidding Document

7.1 The Bidding Document consist of Parts 1, 2, and 3, which include all the Sections indicated below, and should be read and construed in conjunction with any Addenda issued in accordance with ITB 9.

PART 1 Bidding Procedures

- Section I. Instructions to Bidders (ITB)
- Section II. Bid Data Sheet (BDS)
- Section III. Evaluation and Qualification Criteria
- Section IV. Bidding Forms

PART 2 Supply Requirements

• Section V. Schedule of Requirements

PART 3 Conditions of Contract and Contract Forms

- Section VI. General Conditions of Contract (GCC)
- Section VII. Special Conditions of Contract (SCC)

Section VIII. Contract Forms

- 7.2 The Purchaser will reject any Bid submission if the Bidding Document was not purchased directly from the Purchaser, or through its assigned office *as stated in the BDS*.
- 7.3 The Bidder is expected to examine all instructions, forms, terms, and specifications in the Bidding Document as well as in Amendments, if any. Failure to furnish all information or documentation required by the Bidding Document may result in the rejection of the Bid.
- 7.4 The Invitation for Bids issued by the Purchaser is not part of the Bidding Document



8. Clarification of Bidding Document/Prebid meeting

8.1

- A prospective Bidder requiring any clarification of the Bidding Document shall contact the Purchaser in writing at the Purchaser's address *indicated in the BDS*. The Purchaser will respond in writing to any request for clarification, provided that such request is received within the time limit *specified in the BDS* prior to the deadline for submission of Bids. The Purchaser shall forward copies of its response to all Bidders who have acquired the Bidding Document directly from it, including a description of the inquiry but without identifying its source. Should the Purchaser deem it necessary to amend the Bidding Document as a result of a clarification, it shall do so following the procedure under ITB 9 and 24.2.
- 8.2 The purchaser may organize a pre-bid meeting of Bidders at least ten (10) days before the deadline for submission of Bids at the place, date and time as *specified in the BDS* to provide information relating to Bidding Documents, Technical specifications and the like matters. Should the purchaser deem it necessary to amend the Bidding Document as a result of a clarification, it shall do so following the procedure under ITB 9 and ITB 24.2.

9. Amendment of Bidding Document

- 9.1 At any time prior to the deadline for submission of the Bids, the Purchaser may amend the Bidding Document by issuing addenda.
- 9.2 Any addendum issued shall be part of the Bidding Document and shall be communicated in writing to all who have obtained the Bidding Document directly from the Purchaser.
- 9.3 To give prospective Bidders reasonable time in which to take an addendum into account in preparing their Bids, the Purchaser may, at its discretion, extend the deadline for the submission of the Bids, pursuant to ITB 24.2.

C. Preparation of Bids

10. Cost of Bidding

10.1 The Bidder shall bear all costs associated with the preparation and submission of its Bid, and the Purchaser shall not be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.

11. Language of Bid

11.1 The Bid, as well as all correspondence and documents relating to the Bid exchanged by the Bidder and the Purchaser, shall be written in the language *specified in the BDS*. Supporting documents and printed literature that are part of the Bid may be in another language provided they are accompanied by an accurate translation of the relevant passages in the language *specified in the BDS*, in which case, for purposes of interpretation of the Bid, such translation shall govern.

12. Documents Comprising the Bid

- 12.1 The Bid shall comprise the following:
 - (a) Bid Submission Sheet and the applicable Price Schedules, in accordance with ITB Clauses 13, 15, and 16;
 - (b) Bid Security in accordance with ITB 21;
 - (c) alternative bids, if permissible, in accordance with ITB 14;
 - (d) written confirmation authorizing the signatory of the Bid to commit the Bidder, in accordance with ITB 22;



- (e) documentary evidence in accordance with ITB 17 establishing the Bidder's eligibility to bid;
- (f) documentary evidence in accordance with ITB Clauses 18 and 31, that the Goods and Related Services conform to the Bidding Document;
- (g) documentary evidence in accordance with ITB 19 establishing the Bidder's qualifications to perform the contract if its Bid is accepted; and
- (h) any other document required in the BDS.

13. Bid Submission Sheet and Price Schedules

- 13.1 The Bidder shall submit the Bid Submission Sheet using the form furnished in Section IV, Bidding Forms. This form must be completed without any alterations to its format, and no substitutes shall be accepted. All blank spaces shall be filled in with the information requested.
- 13.2 The Bidder shall submit the Price Schedules for Goods and Related Services, according to their origin as appropriate, using the forms furnished in Section IV, Bidding Forms

14. Alternative Bids

14.1 Unless otherwise *indicated in the BDS*, alternative bids shall not be considered.

15. Bid Prices and Discounts

- 15.1 The Bidder shall complete the appropriate Price Schedule and the sources of Goods schedules included herein, stating the unit prices, total cost per item, the total Bid amount and the expected countries of origin of the Goods to be supplied under the contract.
- 15.2 Prices quoted in the Price Schedules shall be entered separately in the following manner:
 - i. the price of the goods quoted EXW (ex works, ex factory, ex warehouse, ex showroom, or off-the-shelf, as applicable), including all customs duties and sales and other taxes already paid or payable: (i) on the components and raw materials used in the manufacture or assembly of goods quoted ex works or ex factory; or (ii) on the previously imported goods of foreign origin quoted ex warehouse, ex showroom or off-the-shelf;
 - ii. the price for inland transportation, insurance, and other costs incidental to delivery of the goods to their final destination, if **specified in the BDS**;
 - iii. the price of other (incidental) services, if any, *listed in the BDS*.
- 15.3 Prices quoted by the Bidder shall be fixed during the Bidder's performance of the Contract and not subject to variation on any account, unless otherwise *specified in the BDS*. A Bid submitted with an adjustable price quotation shall be treated as nonresponsive and shall be rejected, pursuant to ITB 31. However, if in *accordance with the BDS*, prices quoted by the Bidder shall be subject to adjustment during the performance of the Contract, a Bid submitted with a fixed price quotation shall not be rejected, but the price adjustment shall be treated as zero.
- 15.4 The terms EXW shall be governed by the rules prescribed in the current edition of INCOTERMS published by the International Chamber of Commerce, Paris.
- 15.5 The Bidder's separation of price components in accordance with



- ITB 15.1 above will be solely for the purpose facilitating the comparison of bids by the Purchaser and will not in any way limit the Purchaser's right to contract on any of the terms offered.
- 15.6 If the Bidder intends to offer any unconditional discount, it shall always be expressed in fixed percentage and that shall not vary as the quantity varies and be applicable to each unit rate. The methodology for its application shall be provided in bid submission sheet.
- 16. Currencies of Bid
- 16.1 All Prices shall be quoted in Nepalese Rupees.
- 17. Documents
 Establishing
 the Eligibility
 of the Bidder
- 17.1 To establish their eligibility in accordance with ITB 4, Bidders shall:
 - (a) complete the eligibility declarations in the Bid Submission Sheet, included in Section IV, Bidding Forms; and
 - (b) if the Bidder is an existing or intended JV in accordance with ITB 4.2, submit a copy of the JV Agreement, or a letter of intent to enter into such an Agreement. The respective document shall be signed by all legally authorized signatories of all the parties to the existing or intended JV, as appropriate.
 - (c) submit the copy of the documents as specified in BDS.
- 18. Documents
 Establishing
 the Conformity
 of the Goods
 and Related
 Services to the
 Bidding
 Document
- 18.1 To establish the conformity of the Goods and Related Services to the Bidding Document, the Bidder shall furnish as part of its Bid the documentary evidence that the Goods and Related Services conform to the requirements specified in Section V, Supply Requirements.
- 18.2 The documentary evidence may be in the form of literature, drawings or data, and shall consist of a detailed item-by-item description of the essential technical and performance characteristics of the Goods and Related Services, demonstrating substantial responsiveness of the Goods and Related Services to those requirements, and if applicable, a statement of deviations and exceptions to the provisions of Section V, Schedule of Requirements.
- 18.3 Standards for workmanship, process, material, and equipment, as well as references to brand names or catalogue numbers specified by the Purchaser in the Section V, Schedule of Requirements, are intended to be descriptive only and not restrictive. The Bidder may offer other standards of quality, brand names, and/or catalogue numbers, provided that it demonstrates, to the Purchaser's satisfaction, that the substitutions ensure substantial equivalence or are superior to those specified in Section V, Schedule of Requirements.
- 19. Documents
 Establishing
 the
 Qualifications
 of the Bidder
- 19.1 The documentary evidence of the Bidder's qualifications to perform the contract, if its bid is accepted, shall establish to the Purchaser's satisfaction that the Bidder meets each of the qualification criterion specified in Section III, Evaluation and Oualification Criteria.
- 19.2 If so *required in the BDS*, a Bidder that does not manufacture or produce the Goods it offers to supply shall submit the Manufacturer's Authorization using the form included in Section



- IV, Bidding Forms to demonstrate that it has been duly authorized by the manufacturer or producer of the Goods to supply these Goods in Nepal and take care of the warranty provided.
- 19.3 If so *required in the BDS*, a Bidder that does not conduct business within Nepal shall submit evidence that it will be represented by an Agent in Nepal equipped and able to carry out the Supplier's maintenance, repair and spare parts-stocking obligations prescribed in the Conditions of Contract and/or Technical Specifications.
- 19.4 A foreign Bidder wishing to have or already having a local agent shall state the following:
 - a. Name and address of the Agent/Representative,
 - b. The Agent/Representative providing type of services,
 - c. Amount of commission if the Agent/Representative is entitled to get such payment and if it participates in the procedure of payment,
 - d. Other agreement with Agent/Representative, if any,
 - e. Bidder shall certify in the Letter of Authorization as follows: "We certify that the statement and disclosure made by us on the above are complete and true to the best of our knowledge and belief",

If the agent has not been appointed:

- f. Source of information about tender invitation,
- g. The remuneration given to the individual or firm/company or organization to work on its behalf for submitting tender, representation in the bid opening and other required action in connection with the tender,
- h. Transfer or handover an evidence of foreign currency exchanged which required to be submitted with the tender,
- i. If the bank account of any Nepali citizen has been used for the exchange of foreign currency specify the name of the individual and his address. If the foreign currency has been exchanged by self then the certificate of currency exchange.
- 19.5 If a foreign Bidder in its Bid, has not provided the information mentioned in ITB 19.4 or has submitted its bid stating that the Bidder does not have a local agent and later it is proved that the bidder has a local agent or it is proved that the commission mentioned in the Bid is less than the commission received by the local agent then the Purchaser shall initiate proceedings to blacklist such bidder in accordance with ITB 3.2.
- 20. Period of Validity of Bids
- 20.1 Bid shall remain valid for a period *specified in the BDS* after the bid submission deadline date prescribed by the purchaser. A bid valid for a shorter period shall be rejected by the purchaser as nonresponsive.
- 20.2 In exceptional circumstances, prior to the expiration of the bid validity period, the Purchaser may request Bidders to extend the period of validity of their Bids. The request and the responses shall be made in writing. If a Bid Security is requested in accordance with ITB 21, it shall also be extended for a corresponding period. A Bidder may refuse the request without forfeiting its Bid Security. A Bidder granting the request shall not be required or permitted to modify its Bid.
- 21. Bid Security
- 21.1 The Bidder shall furnish as part of its bid, in original form a Bid Security as *specified in the BDS*.



- 21.2 If a bid security is specified pursuant to ITB 21.1, the bid security shall be a demand guarantee in any of the following forms at the Bidder's option:
 - (a) original copy of an unconditional bank guarantee from "A" class commercial bank or;
 - (b) original copy of cash deposit voucher in the Employer's Account as *specified in BDS*.

In case of a bank guarantee, the Bid Security shall be submitted using the Bid Security Form included in Section IV, Bidding Forms. The form must include the complete name of the Bidder. The Bid Security shall be valid for minimum thirty (30) days beyond the end of the validity period of the bid. This shall also apply if the period for bid validity is extended.

The bid security issued by any foreign Bank outside Nepal must be counter guaranteed by an "A" class Commercial Bank in Nepal.

- 21.3 If a bid Security is required in accordance with ITB 21.1, any Bid not accompanied by an enforceable and compliant Bid Security in accordance with ITB 21.2, shall be rejected by the Purchaser as nonresponsive. In case of e-submission, if the scanned copy of an acceptable bid security letter is not uploaded with the eletronic bid then bid shall be rejected.
- 21.4 If a Bid Security is specified pursuant to ITB 21.1, the Bid Security of unsuccessful Bidders shall be returned within three (3) days upon the successful Bidder furnishingthe Performance Security and signing the Contract Agreement pursuant to ITB 41 and ITB 42.
- 21.5 If a Bid Security is specified pursuant to ITB 21.1, the Bid Security of the successful Bidder shall be returned as promptly as possible once the successful Bidder has signed the Contract Agreement and furnished the required Performance Security.
- 21.6 The Bid Security may be forfeited:
 - (a) if a Bidder requests for withdrawal or modification of its bid as against of the ITB clause 26.3 during the period of bid validity specified by the bidder on the bid submission for except as provided in ITB 20.2.
 - (b) if the successful Bidder fails to:
 - (i) sign the Contract in accordance with ITB 42; or
 - (ii) furnish a Performance Security in accordance with ITB 41
- 21.7 The Bid Security of a JV must be in the name of the JV that submits the bid. If the JV has not been legally constituted at the time of bidding, the Bid Security shall be in the names of all future partners as named in the letter of intent mentioned in ITB 17.1.

22. Format and Signing of Bid

22.1 The Bidder shall prepare one original of the documents comprising the Bid as described in ITB 12 and clearly mark it "ORIGINAL." In addition, the Bidder shall submit copies of the Bid, in the number *specified in the BDS* and clearly mark them "COPY." In the event of any discrepancy between the original and the copies, the original shall prevail.



- 22.2 The original and all copies of the Bid shall be typed or written in indelible ink and shall be signed by a person duly authorized to sign on behalf of the Bidder. This authorization shall be attached to the Bid.
- 22.3 Any amendments such as interlineations, erasures, or overwriting shall be valid only if they are signed or initialed by the person signing the Bid.

D. Submission and Opening of Bids

23. Sealing and Marking of Bids

- 23.1 Bidders may always submit their bids by mail or by hand or by courier, but in any means bid must be delivered within the deadline of submission as mentioned in ITB 24. When so *specified in the BDS*, Bidders have the option of submitting their bids electronically. Bidders submitting bids electronically shall follow the electronic bid submission procedures *specified in the BDS*.
- 23.2 Bidders submitting bids by mail or by hand or by courier shall enclose the original and each copy of the Bid, including alternative bids, if permitted in accordance with ITB 14, in separate sealed envelopes, duly marking the envelopes as "ORIGINAL", "ALTERNATIVE" and "COPY." These envelopes containing the original and the copies shall then be enclosed in one single envelope. The rest of the procedure shall be in accordance with ITB 23.3 and 23.4.
- 23.3 The inner and outer envelopes shall:
 - (a) bear the name and address of the Bidder;
 - (b) be addressed to the Purchaser in accordance with ITB 23.1; and
 - (c) bear a warning "NOT TO OPEN BEFORE THE TIME AND DATE FOR BID OPENING".
- 23.4 If all envelopes are not sealed and marked as required, the Purchaser will assume no responsibility for the misplacement or premature opening of the bid.

24. Deadline for Submission of Bids

- 24.1 Bids must be received by the Purchaser at the address and no later than the date and time *indicated in the BDS*. In case of esubmission, the standard time for e-submission is Nepal Standard Time as set out in the server. The e-procurement system will accept the e-submission of bid from the date of publishing of notice and will automatically not allow the e-submission of bid after the deadline for submission of bid.
- 24.2 The Purchaser may, at its discretion, extend the deadline for the submission of Bids by amending the Bidding Document in accordance with ITB 9, in which case all rights and obligations of the Purchaser and Bidders previously subject to the deadline shall thereafter be subject to the deadline as extended.

25. Late Bids

25.1 The Purchaser shall not consider any Bid that arrives after the deadline for submission of Bids, in accordance with ITB 24. Any Bid received by the Purchaser after the deadline for submission of Bids shall be declared late, rejected, and returned unopened to the Bidder.



26. Withdrawal, or Modification of Bids

- 26.1) A bidder may withdraw, or modify its bid after it has been submitted either in hard copy or by e-Submission. Procedures for withdrawal or modification of submitted bids are as follows:
 - (i) Bids submitted in hard Copy
 - a) Bidders may withdraw or modify its bids by sending a written notice in a sealed envelope, duly signed by an authorized representative, and shall include a copy of the authorization in accordance with ITB 20.2 before 24 hours prior to the last deadline of submission of bid. The corresponding modification of the bid must accompany the respective written notice. All notices must be:(aa) prepared and submitted in accordance with ITB 20 and ITB 21, and in addition, the envelopes shall be clearly respective "WITHDRAWAL", "MODIFICATION;" and (bb) received by the Employer 24 hours prior to the deadline prescribed for submission of bids, in accordance with ITB 22.
 - ii) E-submitted bids.
 - a) Bidder may submit modification or withdrawal prior to the deadline prescribed for submission of bids through e-GP system by using the forms and instructions provided by the system. Once a Bid is withdrawn, bidder shall not able to submit another bid for the same bid.
- 26.2 Bids requested to be withdrawn in accordance with ITB 26.1 shall be returned unopened to the Bidders.
- 26.3 In case of bids submitted in hard copy no bid shall be withdrawn or modified in the interval between 24 hours prior time of the deadline for submission of bids and the expiration of the period of bid validity specified by the Bidder on the letter of bid or any extension thereof.

In case of e-submitted bids no bids shall be withdrawn or modified in the interval between deadline for submission of bids and the expiration of the period of bid validity specified by the Bidder on the bid submission form or any extension thereof.

27. Bid Opening

27.1 The Purchaser shall conduct the bid opening in public in the presence of bidder or its representative who choose to attend at the address, date and time **specified in the BDS.** The purchaser shall download the e-submitted bid files. The e-procurement system allows the purchaser to download the e-submitted bid files (report) only after bid opening date and time after login simultaneously by at least two members of the bid opening committee.

Electronically submitted bid shall be opened at first in the same time and date as specified above. Electronic Bids shall be opened one by one and read out. The e-submitted bids must be readable through open standards interfaces. Unreadable and or partially submitted bid files shall be considered incomplete.

27.2 Before opening the bids the purchaser shall separate the envelopes of the bids received after the deadline of bid submission, the envelopes containing an application given for WITHDRAWAL, MODIFICATION of bids and the envelopes of bids duly registered. The bids received after the deadline of



submission shall be returned to the concerned bidder unopened. Then envelopes marked "WITHDRAWAL" shall be opened first, read out, and recorded, and the envelope containing the corresponding Bid shall not be opened, but returned to the Bidder. If the withdrawal notice is not accompanied by a copy of the valid authorization pursuant to ITB 22.2, the withdrawal shall not be permitted and the corresponding Bid will be opened. Envelopes marked "MODIFICATION" shall be opened, read out, and recorded with the corresponding Bid. No Bid shall be modified unless the corresponding Modification Notice contains a valid authorization to request the modification and is read out and recorded at bid opening. Only envelopes that are opened, read out, and recorded at bid opening shall be considered further.

- 27.3 All other envelopes shall be opened one at a time, and the following read out and recorded: the name of the Bidder and whether there is a modification; the Bid Prices (per lot if applicable), any discounts and alternative offers; the presence of a Bid Security, if required; if there is discrepancy between figure and words, description of such discrepancy; whether the bid form is signed by the bidder or his agent; and any other details as the Purchaser may consider appropriate. Only discounts and alternative offers read out and recorded at bid opening shall be considered for evaluation. No Bid shall be rejected at bid opening except for late bids, in accordance with ITB 25.1.
- 27.4 The Purchaser shall prepare a record of the bid opening that shall include, as a minimum: the name of the Bidder and whether there is a withdrawal, or modification; the Bid Price, per lot if applicable, any discounts and alternative offers if they were permitted; and the presence or absence of a Bid Security. The Bidders' representatives who are present shall be requested to sign the record. The omission of a Bidder's signature on the record shall not invalidate the contents and effect of the record. A copy of the record shall be distributed to all Bidders who submitted bids in time, and posted on line when electronic bidding is permitted. The Bidders' representatives who are present shall also be requested to sign an attendance sheet.

E. Evaluation and Comparison of Bids

28. Confidentiality

- 28.1 Information relating to the examination, evaluation, comparison, and post-qualification of Bids, and recommendation of contract award, shall not be disclosed to Bidders or any other persons not officially concerned with such process until publication of the Contract award; thereafter, information will be disclosed in accordance with ITB 42.2.
- 28.2 Any attempt by a Bidder to influence the Purchaser in the examination, evaluation, comparison, and post-qualification of the Bids or Contract award decisions may result in the rejection of its Bid.
- 28.3 Notwithstanding ITB 28.2, from the time of bid opening to the time of Contract award, if any Bidder wishes to contact the Purchaser on any matter related to the bidding process, it should do so in writing.



29. Clarification of Bids

29.1 To assist in the examination, evaluation, comparison and post-qualification of the Bids, the Purchaser may, at its discretion, ask any Bidder for a clarification of its Bid. Any clarification submitted by a Bidder with regard to its Bid and that is not in response to a request by the Purchaser shall not be considered. The Purchaser's request for clarification and the response shall be in writing. No change in the prices or substance of the Bid shall be sought, offered, or permitted, except to confirm the correction of arithmetic errors discovered by the Purchaser in the evaluation of the Bids, in accordance with ITB 33.

30. Deviations, Reservations, and Omissions

30.1 During the evaluation of bids, the following definitions apply:

- (a) "Deviation" is a departure from the requirements specified in the Bidding Document;
- (b) "Reservation" is the setting of limiting conditions or withholding from complete acceptance of the requirements specified in the Bidding Document; and
- (c) "Omission" is the failure to submit part or all of the information or documentation required in the Bidding Document.

31. Determination of Responsiveness

- 31.1 The Purchaser's determination of the responsiveness of a Bid is to be based on the contents of the Bid itself, as defined in ITB 12.
- 31.2 A substantially responsive bid is one that meets the requirements of the Bidding Document without material deviation, reservation, or omission. A material deviation, reservation, or omission is one that.
 - (a) if accepted, would:
 - (i) affect in any substantial way the scope, quality, or performance of the Goods and Related Services specified in Section V, Schedule of Requirements; or
 - (ii) limits in any substantial way, inconsistent with the Bidding Document, the Purchaser's rights or the Bidder's obligations under the proposed Contract; or
 - (b) if rectified, would unfairly affect the competitive position of other Bidders presenting substantially responsive bids.
- The Purchaser shall examine the technical aspects of the bid in particular, to confirm that all requirements of Section V, Schedule of Requirements have been met without any material deviation or reservation.

32. Non-material Non-conformities

- 32.1 The Purchaser may regard a Bid as responsive even if it contains minor deviations that do not materially alter or depart from the characteristics, terms, conditions and other requirement set forth in the Bidding Document or if it contains errors or oversights that are capable of being corrected without affecting the substance of the Bid.
- 32.2 Provided that a Bid is substantially responsive, the Purchaser may request that the Bidder submit the necessary information or documentation, within a reasonable period of time, to rectify non-material non-conformities or omissions in the Bid related to documentation requirements. Requesting information or



- documentation on such non-conformities shall not be related to any aspect of the price of the Bid. Failure of the Bidder to comply with the request may result in the rejection of its Bid.
- 32.3 Provided that a Bid is substantially responsive, the Purchaser shall rectify non-material non-conformities or omissions. To this effect, the Bid Price shall be adjusted, for comparison purposes only, to reflect the price of the missing or non-conforming item or component. The adjustment shall be made using the method indicated in Section III, Evaluation and Qualification Criteria.
- 32.4 If small differences are found such as in technical specification, description, feature which does not make the bid to be rejected, then the cost, which is calculated to the extent possible due to such differences, shall be included while evaluating bid.
- 32.5 If the value is found fifteen percent more than the quoted amount of the bidder on account of small differences pursuant to ITB 31.4, such bid shall be considered irresponsive in substance and shall not be considered for evaluation.

33. Correction of Arithmetical Errors

- 33.1 Provided that the Bid is substantially responsive, the Purchaser shall correct arithmetical errors on the following basis:
 - (a) if there is a discrepancy between the unit price and the total price that is obtained by multiplying the unit price and quantity, the unit price shall prevail and the total price shall be corrected, unless in the opinion of the Purchaser there is an obvious misplacement of the decimal point in the unit price, in which case the total price as quoted shall govern and the unit price shall be corrected;
 - (b) if there is an error in a total corresponding to the addition or subtraction of subtotals, the subtotals shall prevail and the total shall be corrected; and
 - (c) if there is a discrepancy between words and figures, the amount in words shall prevail, unless the amount expressed in words is related to an arithmetic error, in which case the amount in figures shall prevail subject to (a) and (b) above.
- 33.2 If the Bidder that submitted the lowest evaluated Bid does not accept the correction of errors, its Bid shall be rejected.

34. Domestic Preference

- 34.1 If the price of goods manufactured in Nepal, are higher up to ten percent than that of foreign goods, a margin of preference up to ten percent to the goods manufactured in Nepal shall be provided in the evaluation of the Bids.

 (This Clause shall be applicable only for GoN funded
 - (This Clause shall be applicable only for GoN funded procurement.)

35. Evaluation and Comparison of Bids

- 35.1 The Purchaser shall evaluate and compare each Bid that has been determined, up to this stage of the evaluation, to be substantially responsive.
- 35.2 To evaluate a Bid, the Purchaser shall only use all the criteria and methodologies defined in this Clause and in Section III, Evaluation and Qualification Criteria. No other criteria or methodology shall be permitted.
- 36. Postqualification of the Bidder
- 36.1 The Purchaser shall determine to its satisfaction whether the Bidder that is selected as having submitted the lowest evaluated and substantially responsive Bid is qualified to perform the



- Contract satisfactorily.
- 36.2 The determination shall be based upon an examination of the documentary evidence of the Bidder's qualifications submitted by the Bidder, pursuant to ITB 19.
- 36.3 An affirmative determination shall be a prerequisite for award of the Contract to the Bidder. A negative determination shall result in disqualification of the Bid, in which event the Purchaser shall proceed to the next lowest evaluated bid to make a similar determination of that Bidder's capabilities to perform satisfactorily.
- 37. Purchaser's
 Right to Accept
 Any Bid, and to
 Reject Any or
 All Bids
- 37.1 The Purchaser reserves the right to accept or reject any Bid, and to annul the bidding process and reject all Bids at any time prior to Contract award, without thereby incurring any liability to the Bidders.

F. Award of Contract

- 38. Award Criteria
- 38.1 The Purchaser shall select to award the Contract to the Bidder whose offer has been determined to be the lowest evaluated Bid and is substantially responsive to the Bidding Document, provided further that the Bidder is determined to be qualified to perform the Contract satisfactorily.
- 39. Purchaser's
 Right to Vary
 Quantities at
 Time of Award
- 39.1 At the time the Contract is awarded, the Purchaser reserves the right to increase or decrease the quantity of Goods and Related Services originally specified in Section V, Schedule of Requirements, provided this does not exceed the percentages *indicated in the BDS*, and without any change in the unit prices or other terms and conditions of the Bid and the Bidding Document.
- 40. Notification of Intention to Award
- 40.1 The Purchaser shall notify the concerned Bidder whose bid has been selected in accordance with ITB 38.1 within seven days of the selection of the bid, in writing that the Purchaser has intention to accept his/her bid and shall Inform via the Letter of Intention included in the Contract Forms and the information of name, address and amount of selected bidder shall be given to all other bidders who submitted the bid.
- 40.2 If no bidder submits an application pursuant to ITB 43.1 within a period of seven days of providing the notice under ITB 40.1 the Purchaser shall accept the bid selected in accordance with ITB 38.1 prior to the expiry of bid validity period, and notification of award shall be communicated to the bidder to furnish the performance security and sign the contract within fifteen days.
- 41. Performance Security
- 41.1 Within fifteen (15) days of the receipt of notification of award from the purchaser, the successful Bidder shall furnish the Performance Secutiry in accordance with the GCC, using for that purpose the Performance Security Form included in Section VIII, contract forms, or another form acceptable to the purchaser.
 - i) If bid price of the bidder selected for acceptance is up to 15 (fifteen) percent less than the approved cost estimate, the performance security amount shall be 5 (five) percent of the bid price.
 - ii) For the bid price less than 15 percent of the cost estimate, the performance security amount shall be determined as follows:



Performance Security Amount = $[(0.85 \times Cost Estimate - Bid Price) \times 0.5] + 5\%$ of Bid Price.

The Bid Price and Cost Estimate shall be inclusive of Value Added Tax.

The performance security issued by any foreign Bank outside Nepal must be counter guaranteed by an "A" class Commercial Bank in Nepal.

- 41.2 Failure of the successful Bidder to submit the above-mentioned Performance Security or sign the Contract Agreement shall constitute sufficient grounds for the annulment of the award and forfeiture of the Bid Security. In that event the Purchaser may award the Contract to the next lowest evaluated Bidder whose offer is substantially responsive and is determined by the Purchaser to be qualified to perform the Contract satisfactorily.
- 42.1 The successful Bidder shall sign the contract in the form included in section VIII after the submission of performance security in accordance with ITB 41.
- 42.2 At the same time, the Purchaser shallaffix a public notice on the result of the award on its notice board and make arrangement for causing such notice to be pasted on the notice board also of the District Development Committee, District Administration Office, and District Treasury and Controller Office. The Purchaser may make arrangements to post the notice into its website, if it has; and if it does not have, into the website of the Public Procurement Monitoring Office, identifying the bid and lot numbers and the following information: (i) name of each Bidder who submitted a Bid; (ii) bid prices as read out at Bid Opening; (iii) name and evaluated prices of each Bid; (iv) name of bidders whose bids were rejected and the reasons for their rejection; and (v) name of the winning Bidder, and the Price it offered, as well as the duration and summary scope of the Contract awarded.
- 42.3 The Purchaser shall promptly respond in writing to any unsuccessful Bidder who, within thirty days from the date of issuance of notification of award in accordance with ITB 40.2, requests in writing the grounds on which its bid was not selected.
- 42.4 If the bidder whose bid is accepted fails to sign the contract as stated ITB 39.1, the Public Procurement Monitoring Office shall blacklist the bidder on recommendation of the Public Entity.

43. Complaint and Review

42. Signing of

Contract

- 43.1 If a Bidder dissatisfies with the Procurement proceedings or the decision made by the Purchaser in the intention to award the Contract, it may file an application to the Chief of the concerning Public Entity of the Purchaser within seven (7) days of having, receipt of such notice or decision making, for review of the proceedings stating the factual and legal grounds.
- 43.2 An application filed after the deadline pursuant ITB 43.1 shall not be processed.
- 43.3 The chief of Public Entity of the Purchaser shall, within five (5) days after receiving the application, give its decision with reasons, in writing pursuant to ITB 43.1:
 - (a) whether to suspend the procurement proceeding and the



procedure for further proceedings to be adopted; or

(b) whether or not to reject a application.

No application can be submitted before the Review Committee for review against the decision made by the chief of the Public Entity for the Bid amount up to the value as stated in BDS.

- If the Bidder is not satisfied with the decision of the Public Entity in accordance with ITB 43.3, or the decision by the Public Entity is not given within five (5) days of receipt of application pursuant to ITB 43.1, it can, within seven (7) days of receipt of such decision, file an application to the Review Committee of the GoN, stating the reason of its disagreement on the decision of the chief of Public Entity and furnishing the relevant documents, provided that its Bid amount is above the amount as stated in ITB 43.3. The application may be sent by hand, or by post, or by courier, or by electronic media at the risk of the Bidder itself.
- 43.5 Late application filed after the deadline pursuant to ITB 43.4 shall not be processed.
- Within three (3) days of the receipt of application from the Bidder, pursuant to ITB 43.4, the Review Committee shall notify the concerning Public Entity of the Purchaser to furnish its procurement proceedings and comments on the issue, pursuant to **ITB** 43.3.
- Within three (3) days of receipt of the notification pursuant to ITB 43.6, the Public Entity shall furnish the copy of the related documents along with its comment or reaction of complaint to the Review Committee.
- The Review Committee, after inquiring from the Bidder and the Public Entity, if needed, shall give its decision within one (1) month after receiving the application filed by the Bidder, pursuant to ITB 43.4.
- The Bidder, filing application pursuant to ITB 43.4, shall have to furnish a cash amount or Bank guarantee as stated in BDS with the validity period of at least ninety (90) days from the date of the filing of application pursuant to ITB 43.4. Application filed without furnishing the security deposit shall not be processed.
- 43.10 If the claim made by the Bidder pursuant to ITB 43.4 is justified, the Review Committee shall have to return the security deposit to the applicant, pursuant to ITB 43.9, within seven (7) days of such decision made.
- 43.11 If the claim made by the Bidder pursuant to ITB 43.4 is rejected by the Review Committee, the security deposit submitted by the Bidder pursuant to ITB 43.9 shall be forfeited.

and PPR

44. Provision of PPA If any provision of this document are inconsistent with Public Procurement Act (PPA), 2063 or Public Procurement Regulations (PPR), 2064, the provision of this documents shall be void to the extent of such inconsistency and the provision of PPA and PPR shall prevail.



Section II. Bid Data Sheet

The following specific data for the goods to be procured shall complement, supplement, or amend the provisions in the Instructions to Bidders (ITB). Whenever there is a conflict, the provisions herein shall prevail over those in ITB.

	A. Introduction
ITB 1.1	Name of the Purchaser:
115 1.1	Government of Nepal
	Ministry of Health
	Department of Health Services
	Logistics Management Division
	Teku, Kathmandu.
ITB 2.1	Name of the Contract: Procurement of Medical & Surgical Instruments For PHC.
110 2.11	Contract Identification No.: DOHS/G/NCB-155/2016-17
ITB 2.1	Name of the Project: Procurement of Medical & Surgical Instrumnts For PHC
	Name of the DP: GON/Pool Fund
	Implementing Agency:
	Government of Nepal,
	Ministry of Health
	Department of Health Services, Logistics Management Division
	Teku, Kathmandu.
ITB 4.1	Bidders from the following countries are not eligible: Not Applicable
ITB 4.9	The foreign Bidder at the time of bid submission:
	shall submit:
	a) Up to date Firm or Company Registration Certificate duly permitted to
	deal with the offered goods, b) Manufacturers/Suppliers from eligible source country.
	c) A written declaration made by the Bidder stating that the Bidder is not
	ineligible to participate in the Bid; has no conflict of interest in the
	proposed procurement proceedings, and has not been punished for a
	profession or business related offence. d) Joint Venture Authorization/Agreement (if any)
	shall declare to submit at the time of contract agreement:
	Appointment of a local agent to represent in all aspect on behalf of the bidder.
ITB 5.1	Goods and related services to be supplied from following countries are not
	eligible: NA
	B. Bidding Document
ITB 8.1	For clarification purposes only, the Purchaser's address is:
	Attention: The Director
	Name of the Purchaser: Government of Nepal
	Ministry of Health, Department of Health Services
	Logistics Management Division.
	City/Town: Teku, Kathmandu
	District: Kathmandu



	G .	NT 1		
	Country: Nepal			
	Telephone: 01 4261768			
	Facsimile Number: 01 4261413 Electronic Mail Address: drtinkari@gmail.com			
ITB 8.1	that suc	rchaser will resp th request is recen mission of bid.	boond in writing to any request the bived no later than ten (10) days	for clarification provided prior to the deadline date
ITB 8.2	Pre-Bid	meeting shall n	ot be organized.	
		C. I	Preparation of Bids	
ITB 11.1		guage of the Bid		
ITB 12.1 (h)	•	 The Bidder shall submit the following additional documents with its Bid: Original manufacturer's authorization or Authorization of the authorized dealer in Nepal to bid on the IFB. Audit reports of the last 3 fiscal years (2070/71, 2071/72, 2072/73) Information on past supplies of similar goods and services in last three years with contract amount, supply date, name and address of the purchaser. 		
ITB 14.1	Alterna	tive Bids are no	t permitted	
ITB 15.2 (i)	The price quoted shall be: The prices shall include all duties, taxes, other levies, transportation, and incidental services to the final destination at Logistics Managment Division, Central Store, Teku, Kathmandu.			
ITB 15.2 (ii)			insportation : NA	
ITB 15.2 (iii)		The price of other incidental services: NA		
ITB 15.3	The prices quoted by the Bidder shall be: Fixed during the contract period and			
			rice Adjustment.	P
ITB 15.4		oterms edition is		
ITB 17.1 (c)		dders shall submi	it: gistration Certificate	
	 Copy of Business Registration Certificate 			
	 Copy of VAT and PAN Registration Certificate, 			
	 Copy of Tax Clearance Certificate for the F/Y 2072/73 			
ITB 19.2	A Manufacturer's Authorization letter is required for all the items listed in Section V Schedule of the Requirements.			
	Alternatively, if the authorized dealer is not participating on this bid a bidder must submit an original authorization letter to participate on this bid from the authorized dealer, along with the certificate of the authorised dealer in Nepal.			
ITB 19.3	The Bidder is required to include with its bid, evidence that it will be represented by an Agent in Nepal, if the bidder does not conduct business in Nepal.			
ITB 20.1		l validity period shadh-16(30-June	shall be 90 days from the date e-2017).	of bid opening (i.e. upto
ITB 21.1	The Bio	der shall furnish	a bid security, from "A" class co	ommercial bank of Nepal
	The am	ount of the Bid S	Security shall be:	•
	S,N.	Slice no.	Description-Qty	Required Bid Security (NPR)
		NCB-155	As mentioned in Schedule of requirement	16,50,000.00



	The Bid Security must be valid for thirty (30) days from the bid validity period (i.e. till 2074- Shrawan-15.; 30-July-2017).		
ITB 21.2 (b)	There is no provision of Bid Security in the form of cash.		
ITB 22.1	In addition to the original of the Bid, the number of copies is: One (1) . (That is one original and one copy) In case of e-submission of bid:		
	The Bidder shall submit his bid electronically in PDF files in the manner as specified in ITB Clause 23.1, and submission of hard copy of "original bid" (as Stated above) is not mandatory.		
	In case, if both the electronic bid and original bid in hard copy are submitted to the Purchaser within the bid submission deadline, the Bidder's electronic bid and original bid in hard copy will be accepted for evaluation, provided if the facts and figures in hard copy confirm to the PDF files in electronic bid. If there is any major discrepancy in fact and figures in the electronic bid and original bid in hard copy it will be treated as two separate bids from one Bidder and hence, both the electronic bid and original bid in hard copy shall be disqualified.		
	However, for electronically submitted bid in PDF files, the Bidder shall be required to submit original Bid Security letter and all documentswithin 7 days of bid opening.		
ITB 22.2	The written confirmation of Authorization to sign on behalf of the Bidder shall consist of:		
	(a) Power of Attorney (A separate letter from authorized signatory delecating power to the nominated person, his / her designation in the instutition and an authorization to sign the contract.)		
	(b) In the case of Bids submitted by an existing or intended JV, an undertaking signed by all parties (i) stating that all parties shall be jointly and severally liable, and (ii) nominating a representative who shall have the authority to conduct all business for and on behalf of any and all the parties of the JV during the bidding process and, in the event the JV is awarded the Contract, during contract execution.		
	D. Submission and Opening of Bids		
ITB 23.1	Bidders have the option of submitting their bids electronically.		
ITB 23.1	If bidders submit their bids electronically, the electronic bidding submission procedures shall be:		
	A) Bid submission procedure through electronically (e-submission):		
	i) Interested bidders may either purchase the Bidding documents from the Purchaser's office as specified in the IFB Notice or choose to download the bidding documents from the e-procurement section of PPMO website http://www.bolpatra.gov.np. In case, the bidder choose to download the bidding documents, prepare his/her bids on downloaded documents, and submit his/her bid electronically, the Bidder is required to deposit the cost of bidding document (as specified in the bid notice) in the Purchaser's account as specified in the notice. In addition, electronic scanned copy (PDF format) of the Bank deposit voucher/tele transfer receipt shall also be required to be submitted along with the electronic bid files.		



- ii) The Bidder shall fill the following documents and forms (in hard copy of issued bid documents or downloaded bid documents for specific bid), signed by the authorized representative and with seal of the company:
- iii) The Bidder shall then scan the completed original documents, forms in PDF files with appropriate filename as shown in the table below. PDF (Adobe Acrobat) version must be 4.0 or above.

S. No.	Document	PDF File Name	Requirement	Remarks
1	Form of Bid as of Section IV	Bid form -1	Mandatory	
2	Bid Security (Bank Guarantee) as of Section IV	Bid security-2	Mandatory	
3	Company Registration	Company reg-3	Mandatory	All firms in case of JV
4	VAT Registration	VAT reg-4	Mandatory For Nepali firms	All firms in case of JV
5	Tax Clearances Certificate	Tax-5	Mandatory For Nepali firms	All firms in case of JV
6	Power of Attorney of Bid signatory	Power of att-6	Mandatory	
7	Joint Venture Agreement	JV doc-7	Mandatory	In case of JV
8	Bidder's Qualification Information as per section IV	Qualification-8	Mandatory	
9.	Price Schedule with Rate, Amount and Total Amount	Price-9	Mandatory	
10.	Declaration Form	Declaration-10	Mandatory	
11.	Manufacturer's Authorization or Authorization from Authorised Distributor as of Section IV	Authorization-11	Mandatory	
11.	Bank deposit Voucher/tele transfer receipt for bid document purchasing	Bank Voucher- 12	Mandatory	In case the bid doc. is downloaded electronically

Note: Mandatory means the mentioned files must be included in e-submission and non submission of such file shall be considered as non-responsive bid.

- iv) For e-submission purpose the Bidder shall, at first, register in the e-procurement section of PPMO website http://www.bolpatra.gov.np.
- v) After preparing all the required bidding documents in PDF scan files as specified in (ii) and (iii), the Bidder shall upload the PDF bid files and submit his complete bid online



through e-procurement section of PPMO website http://www.bolpatra.gov.np. Within the specified date and time.

vi) Bidders are advised to download the bid submission report to ensure that all the documents/ files are up to date and complete.

B) Requirements and Conditions for e-submission of bid:

- 1) The e-submitted bids must be readable through Adobe Acrobat Reader. Unreadable and or incomplete bid files (not complying as per ITB Clause 23.1) shall be considered incomplete and rejected for further bid evaluation.
- 2) In addition to electronically submitted PDF files, the Bidder shall be required to submit original Bid security letter and all documents within 7 days of bid opening. Non submission of original Bid security letter, documents within specified time may cause forfeiture of Bid Security.
- In case of major discrepancy found between electronically submitted PDF bid files and documents provided by the Bidder as hard copies shall not be considered for further evaluation.
- 4) Proposed facility for submission of bid electronically through e-submission is to increase transparency, non-discrimination, equality of access, and open competition. The Bidders are fully responsible to use the e-submission facility properly in e-procurement section of http://www.bolpatra.gov.np in specified procedures and in no case the Purchaser shall be held liable for Bidder's inability to use this facility.
- 5) When a Bidder submits electronic bid by downloading the bidding documents from the http://www.bolpatra.gov.np webpage it is assumed that the Bidder prepares his bid by studying and examining all the Bidding documents including specifications and conditions of contract.
- 6) In case, the Bidder chooses to download the bidding documents and deposit the cost of bidding document (as specified in the bid notice) in the account of LMD such deposited amount shall be verified by the office during bid evaluation process. The bid shall be non-responsive and shall not be evaluated if the specified cost for bidding document is not deposited in the specified account of LMD.

ITB 24.1 For bid submission purposes only, the Purchaser's address is:

Government of Nepal Ministry of Health Department of Health Services Logistics Management Division

Teku, Kathmandu.

ITB 24.1 The deadline for bid submission is:

Date: 2073-Chaitra-20. (02-April-2017). Time: 12:00 hours Nepal Standard Time

Add following paragraph at the end of Sub-clause 24.1:

In case of e-submission of bid:

The Purchaser's address for the purpose of electronic bid submission is EProcurement section of **http://www.bolpatra.gov.np**. The bidder should note the following:

i) The e-procurement system will accept the e-submission of bid from the date after publishing of notice and will automatically not allow the e-submission of bid after the



	deadline for submission of bid.
	deadilite for submission of bid.
	ii) The standard time for e-submission is Nepalese Standard Time as set out in the server of PPMO
ITB 24.1	If the last date of purchasing, submission and opening of Bid falls on a government holiday then the next working day shall be considered as the last day without any change in the time and place as fixed.
ITB 25	Add ITB Sub-clause 25.2 as follows:
	In case of e-submission of bids, the e-procurement system will, automatically, not allow the e-submission of bid after the deadline for submission of bid.
ITB 26	Add an ITB Sub-clause 26.4 as follows: Withdrawal or Modification of the bid shall be accompanied by a written Power of Attorney in favor of the person / signatory applying for Withdrawal or Modification, duly signed by Authorized Representative(s) of the firm / all authorized Joint Venture partners.
	When a bidder submits his bid in hard copy, the e-procurement section of PPMO website does not allow the bidder to submit his Withdrawal or Modification through e-submission.
	In case of e-submitted bid: (i) Bidders may submit the Withdrawal or Modification either in hard copy or through e-submission.
	(ii) For Withdrawal or Modification of the bid the Bidder is required to submit scanned copy (in PDF file) of their Withdrawal or Modification letter along with a written Power of Attorney in favor of the person / signatory applying for Withdrawal or Modification, duly signed by Authorized Representative(s) of the firm / all authorized Joint Venture partners.
ITB 27.1	The bid opening shall take place at: Department of Health Services, Logistics Management Division Teku, Kathmandu
	Date: 2073-Chaitra-20 (02-April-2017).
	Time: 13:00 hours
ITB 27.1	If electronic bid submission is permitted in accordance with ITB 23.1, the specific bid opening procedures shall be: In case of e-submitted bid: i) Electronically submitted bid shall be opened at first at the same time and on date as specified above.
	ii) The e-procurement system allows the Purchaser to download the esubmitted bid files from the bidders only after the time for opening the bids.
	ii) The e-submitted bids must be readable through Adobe Acrobat Reader. Unreadable and/or partially submitted bid files (not complying as per ITB Sub-clause 23.1) shall be considered incomplete and rejected for further bid evaluation.
	iii) After opening of e-submitted bids files, all files shall be printed and recorded at the time of bid opening.



E. Evaluation and Comparison of Bids		
	Add the following paragraph at the end of Sub-clause 29.1:	
ITB 29.1	In case of e-submission of bid, for verification of information submitted in its bid, the bidder shall submit the originals of its Bid security, Power of Attorney of the Authorized signatory, parts of the completed bid and other clarifications within 7 days from the date of bid opening.	
	Add an ITB Sub-clause 29.2 as follows:	
	If the Bidder does not provide clarification of its bid by the date and time set in the Purchaser's request for clarification, its bid may be rejected.	
	Add Sub-clause 35.3 as follows:	
ITB 35	35.3 In case of e-submission bids, the Employer evaluates the bid based on the information as per electronically submitted bid files. For verification and acceptance of the bid, the bidder shall submit the original bid security and on the request of the Purchaser the bidder need to submit other documents/clarifications as specified in ITB Sub-clause 29.1.	
	In case, if the Bidder can not substantiate or provide evidence to prove the information provided in e-submitted bid through documents/clarifications as per ITB Sub-clause 29.1, the bid shall not be considered for further evaluation.	
	F. Award of Contract	
ITB 39.1	The maximum percentage by which quantities may be increased is: 15% in number or, in value	
	The maximum percentage by which quantities may be decreased is: 15% in number or, in value	
ITB 43.3	No application can be submitted before the Review Committee for review against the decision made by the chief of the Public Entity for the bid amount below the value of Nepalese Rupees 20,000,000 (Twenty million)	
ITB 43.9	The Bidder, filing application pursuant to ITB 43.4, shall have to furnish a cash amount or Bank guarantee equal to 0.5% of its bid price.	



Section III. Evaluation and Qualification Criteria



Evaluation Criteria

Criteria for Bid evaluation are:

Bids shall be considered non-responsive, if:

- 1. The bid is not submitted in the bid document issued by the DoHS, Logistics Management Division.
- 2. The bid is not submitted in the complete Bid Document issued in the name of the bidder itself.
- 3. The bid is not sealed.
- 4. The bid is not submitted with the Bid Form duly filled and signed in the complete document.
- 5. The bid is not submitted within the specified date / time for submission of bids.
- 6. The bid is not submitted along with the Bid security as specified in ITB 21.
- 7. The bid is submitted without the information as specified in ITB 19.
- 8. The bid does not comply with the instructions as specified in the Invitation for Bids.
- 9. Terms of payment and Destination of Delivery of the materials supplied are different from those specified in the Bid Document.
- 10. All prices quoted are not either firm (not estimated one) or conditional or not valid for the period specified in the Bid Document.
- 11. There is a major deviation in specification of the goods proposed by the bidder from that specified in the Technical Specifications and Schedule of Requirement.

Evaluation of a Bid will take into account, in addition to the Bid price quoted in accordance with Clause 15, one or more of the following factors in the manner and to the extent as specified in the Bidding Data.

a) Delivery schedule: Relevant parameters of delivery:

Within the days/weeks mentioned on the Schedule of Requirements from the date of contract signing

No credit will be given to deliveries before the earliest date, and bids offering delivery after the final date shall be treated as non responsive.

b) Reduction in Bid Price for

Deviation in payment schedule: Not Applicable.

c) Compliance with Specification:

All offered goods thereof must be in compliance with the requirements of Technical Specifications, which will be reviewed by Technical Experts. Offeres not meeting the required Technical Specifications will be rejeted as non responsive.

d) Evaluation will be done for the complete package as lowest responsive total bid Price of all items.



Qualification Criteria

(a) Financial Capability:

Bidder's average annual turnover of the last three years should not be less than **6 Crore**. The bidder shall demonstrate compliance with this requirement through submission of certified copies of the annual audit report and tax clearance certificate for the last three years. Non-submission of annual audit statement will be rejected as non-responsive.

(b) Experience and Technical Capacity:

i The bidder must have been supplying the similar goods for last one year. The bidder shall provide documentary evidence demonstrating that it has at least one (1) year of experience in supply of **similar goods** and related services to any corporate organisations.

To substantiate the experience for the goods, the bidder must submit at least two numbers of end user certificates issued by the user administration. The end user certificates must be on the letterhead of the user and shall include but not be limited to relevant information such as description, year of purchase, quantity, name and address of user, name of contact person, telephone and fax numbers.

Not meeting the above qualification will be considered as non-responsive.



Section IV. Bidding Forms

Table of Forms

1.	Bid Submission Form	36
2.	Bidder's Information Form	
3.	Joint Venture Information Form (if applicable only)	39
4.	Financial Situation Form	40
5.	Average Annual Turnover Form	41
6.	Financial Resources Form (if applicable only)	42
7.	Pending Litigation Form (if applicable only)	43
8.	Specific Experience Form (optional to the bidder)	44
9.	Price Schedule For Goods	46
10.	Bid Security	48
11.	Manufacturer's Authorization Letter (Not Applicable)	49



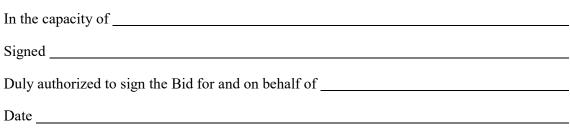
1. Bid Submission Form

(The Bidder shall accomplish the Bid Submission Form in its Letter Head Clearly showing the Bidders Complete name and address)

	Date:
	Contract No.:
	Invitation for Bid No.:
То:	
We	, the undersigned, declare that:
(a)	We have examined and have no reservations to the Bidding Document, including Addenda No.:
(b)	We offer to supply in conformity with the Bidding Document and in accordance with the delivery schedule specified in the Schedule of Requirements, the following Goods and Related Services:
(c)	The total price of our Bid, excluding any discounts offered in item (d) below is:
(d)	The discounts offered and the methodology for their application are:
(e)	Our Bid shall be valid for a period of days from the date fixed for the bid submission deadline in accordance with the Bidding Document, and it shall remain binding upon us and may be accepted at any time before the expiration of that period;
(f)	If our Bid is accepted, we commit to obtain a Performance Security in the amount of percent of the Contract Price for the due performance of the Contract;
(g)	We are not participating, as Bidders, in more than one Bid in this bidding process, other than alternative offers in accordance with the Bidding Document;
(h)	Our firm, its affiliates or subsidiaries, including any subcontractors or suppliers for any part of the Contract, has not been declared includible by the GoN:



(i)	The following commission paid with respect to the bide			or are to be		
	Name of Recipient	Address	Reason	Amount		
	<u> </u>		_			
	(If none has been paid or	r is to be paid, indicate	e "none.")			
(j)	We understand that this Bid your notification of award formal Contract is prepared	, shall constitute a bi				
(k)	k) We understand that you are not bound to accept the lowest evaluated bid or any other bid that you may receive.					
(1)	We declare that, we have interest in the proposed pro an offense relating to the co	ocurement proceedings	and we have not been j			
(m)) We declare that the docum We are fully responsible an			nd authentic.		
(n)	We agree to permit GoN/Dl and other documents relational auditors appointed by the G	ing to the bid submis				
Na	me					





2. Bidder's Information Form

[Insert date (as day, month and year) of Bid Submission]

[The Bidder shall fill in this Form. No alterations to its format shall be permitted and no substitutions shall be accepted. In case of joint venture, each partner shall fill the information in separate form.]

Date:

		Page	of	pages
1.	Bidder's Legal Name			
1.	Bidder & Legar Parite			
2	Bidder's Address:			
3	Bidder's Country of Registration:			
4.	Bidder's Year of Registration:			
5.	Bidder's Legal Address in Country of Registration			
6.	Bidder's Authorized Representative Information:			
	.Name:			
	Address:			
	Telephone/Fax numbers:			
	Email Address			
7	Bidder's Telephone/Fax numbers:			
8	Bidder's Email Address:			
	Attached are copies of the following original documents.			
	1. Firm Registration Certificate			
	2. Authorization to represent the firm			



3. Joint Venture Information Form (if applicable

Lead Name of the Lead Partner in Joint Venture: Partner Share of the Lead Partner: Place of Firm Registration: Place of Business Registration: Percentage of Partnership: Partner Name of the Partner in Joint Venture: Share of the Lead Partner: Place of Firm Registration: Place of Business Registration: Percentage of Partnership: Name of the Partner in Joint Venture: Partner Share of the Lead Partner: Place of Firm Registration: Place of Business Registration: Percentage of Partnership: Name of the partner authorized to sign the Bid:



4. Financial Situation Form

Financial Data for Previous 3 Years (in NRs)					
Year 2070/71	Year 2071/72	Year 2072/73			

Information from Balance Sheet

Total Assets		
Total Liabilities		
Net Worth		
Current Assets		
Current Liabilities		

Information from Income Statement

Total Revenues		
Profits Before Taxes		
Profits After Taxes		

- Attached are copies of financial statements (balance sheets including all related notes, and income statements) for the last three or above years, as indicated above, complying with the following conditions?
 - Historic financial statements must be audited by a certified accountant.
 - Historic financial statements must be complete, including all notes to the financial statements.
 - Historic financial statements must correspond to accounting periods already completed and audited (no statements for partial periods shall be requested or accepted).



5. Average Annual Turnover Form

The information supplied should be the Annual Turnover of the Bidder in terms of the amounts billed to clients for each year for work in progress or completed to NRs at the end of the period reported.

Annual Turnover Data for the Last 3 Years				
Year	Amount (in NRs)			
2070/71				
2071/72				
2072/73				
Average Annual Turnover				



6. Financial Resources Form (if applicable only)

Specify proposed sources of financing, such as liquid assets, unencumbered real assets, lines of credit, and other financial means, available to meet the total cash flow requirements of the subject contract

	Financial Resources					
No.	Source of financing	Amount (in NRs)				
1						
2						
3						

Note:

The letter from the Bank must be unconditional.



7. Pending Litigation Form (if applicable only)

Each Bidder or member of a JV must fill in this form

Year	Matter in Dispute	Value of Pending Claim in NRs	Value of Pending Claim as a Percentage of Net Worth



Bidder's Legal Name:		Date: IFB No.:	
		Page of	pages
Similar Contract		Information	
Contract Identification			
Award date Completion date			
Role in Contract	Contractor	Management Contractor	Subcontract
Total Contract amount			Currency
Description of the works performed by the Bidder			
If partner in a JV or subcontractor, specify participation of total Contract amount	%		Currency
Employer's Name:			
Employer's Address:			
Employer's Telephone/fax number: Employer's E-mail:			

The Bidder shall complete this form for each contract completed/in progress.



Price Schedules



9. Price Schedule For Goods

Name of Bidder	Contract Identification Number	: DOHS/G/NCB-155/2016-17

Item		Country of Ur Origin		Unit Quantity	Unit Price Delivery at LMD, Teku (in NPR)		Total Price
		Origin			In Figure	In Words	(In figure) NPR
1	2	3	4	5	-	6	$7 = 5 \times 6$
1	Delivery set		Unit	240			
2	Episiotomy set		Unit	240			
3	Peri light		Unit	240			
4	Suction (electric)		Unit	120			
5	Nebulizer		Unit	120			
6	Otoscope		Unit	120			
	IUD & Implant insertion and		Unit				
7	removal set			240			
8	Dental Instrument Set		Unit	120			
9	I/D Set		Unit	240			
10	Suture Set		Unit	240			
11	BP set		Unit	600			
12	Stethoscope		Unit	600			
13	Suture removal set		Unit	240			
14	Dressing Set		Unit	240			
15	Protoscope		Unit	120			



16	Foetoscope, Plastic	Unit	480		
17	Digital thermometer	Unit	600		
18	Tongue depressor	Unit	360		
19	Weighting machine (digital)	Unit	240		
20	Laryngoscope	Unit	120		
21	LSCS set	Unit	240		
				Total	
				VAT	
				Grand Total	

Total amount in words:
Name:
In the capacity of:
Signed:
Duly authorized to sign the Bid for and on behalf of:
Date:
Note: 1. Unit price shall include all custom duties and taxes, transportation cost to the final destination and insurance cost except VAT
2. If there is any discrimency between unit price and total price, unit price shall proveil

2. If there is any discripancy between unit price and total price, unit price shall prevail3. If there is any discripancy between amount in figure and amount in words, amount in words shall prevail.



10. Bid Security

[This is the format for the Bid Security to be issued on the letterhead by a "A" class commercial bank specified by Nepal Rastra Bank] [insert Bank's Name, and Address of Issuing Branch or Office]

Date:[insert date]

Beneficiary: [insert Name and Address of Purchaser]

BID GUARANTEE No.: [insert number]

We have been informed that *[insert name of the Bidder]* (hereinafter called "the Bidder") intends to submit its bid to you (hereinafter called "the Bid") for the execution of *[insert name of contract]* under Invitation for Bids No. *[insert IFB number]* ("the IFB").

Furthermore, we understand that, according to your conditions, bids must be supported by a bid guarantee.

At the request of the Bidder, we *[insert name of Bank]* hereby irrevocably undertake to pay you any sum or sums not exceeding in total an amount of *[insert amount in figures][insert amount in words]* upon receipt by us of your first demand in writing accompanied by a written statement stating that the Bidder is in breach of its obligation(s) under the bid conditions, because the Bidder:

- (a) has withdrawn its Bid during the period of bid validity specified by the Bidder in the Form of Bid; or
- (b) having been notified of the acceptance of its Bid by the *Purchaser* during the period of bid validity, (i) fails or refuses to execute the Contract, if required, or (ii) fails or refuses to furnish the performance security, in accordance with the ITB.

This guarantee will expire: (a) if the Bidder is the successful Bidder, upon our receipt of copies of the contract signed by the Bidder and the performance security issued to you upon the instruction of the Bidder; and (b) if the Bidder is not the successful Bidder, upon the earlier of (i) our receipt of a copy of your notification to the Bidder of the name of the successful Bidder; or (ii) thirty (30) days after the expiration of the Bidder's bid which comes to be *[insert the date]*.

Consequently, any demand for payment under this guarantee must be received by us at the office on or before that date.

Name	
In the capacity of	
Signed	
Duly authorized to sign the Bid Security for and on behalf of	
Date	



11. Manufacturer's Authorization Letter

	ald be on the letterhead of the mar roper authority to sign documents manufacturer] Date:	that are b	oinding on the
	IFB No.:		
To:			
WHEREAS	manufacturers		who are
		having	factories at
a Bid in relation to the Invitat	ion for Bids indicated above, the following Goods, mand to subsequently negotiat	he purpos anufacture	se of which is ed by us
•	antee and warranty in accordanc with respect to the Goods offer		
Name			
Signed			
	orization for and on behalf of		
Date			



Section V. Schedule of Requirements

Contents

1.	List of Goods and Related Services	51
2.	Delivery and Completion Schedule	51
3.	Technical Specifications	52



1. List of Goods and Related Services

The Goods and Related Services are shown in Price Schedule.

2. Delivery and Completion Schedule

Delivery shall take place in compliance with the dates, duration, and locations indicated as below:

The delivery period shall start as of the date of signing the contract.

S. No.	Description of Goods	Quantity	Unit for Bid	Acceptable Delivery Date	Final Destination as specified in BDS
1	2	3	4	5	6
NCB 155	Medical & Surgical Instruments for PHC	As mentioned in price Schedule	1 Package	Within 60 days from the date of Contract Signing	Central Store, Department of Health Services, Logistics Management Division, Pathalaiya /Teku, Kathmandu



3. Technical Specifications

The specification and equipment order list contains 2 sections A, B. These sections are an integral part of the specification and equipment order list and complement each other. The two sections are namely:

- A. General points and notes
- B. Slice wise specifications under different Slices (altogether 20 slices)

The right hand blank side must be completed by the bidder with the technical specifications of the equipment offered with supplementary documents enclosed.

A. GENERAL POINTS AND NOTES

1. Dimensions

Dimensions have been included in the specifications and are intended for GUIDANCE ONLY to match the type of size required. Where there are particular parameters to observe, minimum and maximum sizes have been quoted.

2. Mains Electrically Powered Items

All mains electrically powered items should be suitable for operation on the electrical system within Nepal 220/230 volts.

110-volt units, which work through transformers, are NOT acceptable.

3. Instruction/Operating Manuals

Each set of equipment must be supplied with detailed operating and maintenance manuals and technical information in the English language.

4. Sensitive Nature

All the equipment and instruments are of a sensitive nature so they will only be procured from recognized, medical equipment/instrument manufacturers who have an established history or the manufacturers of whose products meet international quality standards.

5. Quality Assurance and Product Conformity

Manufacturers of all medical devices (equipment and instruments) must have a quality assurance system certified under the following standards:

- ISO 13485:2003 (for manufacturing of medical equipment and instruments)
- ISO 9001:2008 (for manufacturing of all other goods).

As may be further specified in the following Technical Specifications per item, medical equipment and instruments proposed and supplied must conform to specific product certification, namely CE mark (certifying compliance with the Medical Devices Directive (MDD)93/42/EEC, with subsequent amendments) or equivalent.

A Certificate of Conformity to the Test Parameters and date of manufacturing shall be available to the Purchaser for all the instruments and equipment.

6. Product Information

All the information provided in the bid should be substantiated by attached product data sheets/technical catalogues and relevant Standards such as International Standards Organisation (ISO), European Norms (EN),



Indian Standard Institute (IS), Nepal Standard (NS), British Standards Institute (BS), American National Standards Institute (ANSI), .

7. Standard Accessories

All equipment should be supplied with their standard accessories as normally provided by the manufacturers in addition to those accessories that are specifically mentioned in the specifications. The cost of these accessories must be included in the bid price.

8. Availability of Spare Parts and Consumables

The bidder shall supply equipment with a start-up supply of consumables allowing for testing and commissioning and approximately six months of normal operation.

9. Technical Trial

Technical trials may be conducted for all the instruments and equipment prior to purchase and the supplier is fully responsible to provide all facilities needed to conduct the Technical trial on the same model of equipment quoted.

10. Installation and commissioning

All equipment which is so specified in the bidding documents must be installed and commissioned by the Supplier at the final destination(s), including, any base plates or connecting devices to the floor/foundation, utility connection to the equipment within the location, calibration and commissioning. The Supplier will also provide and install the latest version of complete programme software required for the installation, commissioning and its functioning for the diagnostic use of the equipment. The Supplier will also provide licensed copies of all such software for future reference and use of the Purchaser. Such software is to be Original Equipment Manufacturer (OEM). The Supplier will also be required to make available the updated version(s) of such OEM program software for use in the same equipment.

The Health Facility at final destination shall be responsible to ensure that a suitable location (room) is made available, including required connections up to the location for electricity, water, air, oxygen, nitrous oxide, drainage, etc. as applicable for the particular equipment.

All other equipment shall be delivered by the supplier in fully assembled operational condition.

The specifications per item also specify whether or not installation and commissioning is required.

12. User training

The Supplier shall conduct user training for equipment to enable operators to use the equipment properly. The training shall include the use of all operational functions of the equipment, as well as routine checks and maintenance expected by users.

For fixed equipment, the training shall be conducted at the site of the equipment, following installation and commissioning. For other equipment the training will be conducted at a central location (Kathmandu or capital of one or more of the regions) in consultation with the Purchaser.

The specifications per item also specify whether or not user training is required.

13. Maintenance service during warranty period

For items as specified in the following individual Technical Specifications and 2 List of Related Services and Completion Schedule, preventive and corrective maintenance must be provided by the supplier during the



period of warranty and included in his bid. The cost of spare parts will be separately payable by the user, except cases covered under warranty.

13. Right to Reject

All the information provided should be accurate and sufficient to convince fully the Purchaser that all the offered goods fully meet the technical specifications and output quality. If such complete information is not provided and which leads to doubts about the technical compliance of the item(s), the Purchaser retains the right to reject the corresponding item.

Bidders are to offer a standard production model most closely matching the specification below and provide details of the offer. The offer must be for brand new equipment.

These specifications are for the **minimum requirement.** Bidders may offer higher specifications but they are to highlight these in the Statement of Compliance column. Units are to be stated using the SI system. For example length in metres (m). Multiples and sub-multiples to be used are: 10^6 mega (M); 10^3 kilo (k); 10^{-3} milli (m) and 10^{-6} micro (μ).

Bidders must enter their offered specifications against each parameter of this Technical Specifications Form (TSF), comment as necessary, and sign and stamp each page. Failure to complete this statement of compliance may result in the offer being rejected.

The bidder shall mention clause by clause comment of the required specification. The bidder shall state:-

- a. "FULLY COMPLIANT" if the item offered fully meet the quotation requirement.
- b. "PARTIALLY COMPLIANT" if the item offered meet the requirement partially. The bidder shall state the reason why the offer is partially compliant. In such cases, the bidder shall clearly mention the extent to which other specifications are offered.
- c. "NON COMPLIANT" if the item cannot meet the requirements. The bidder shall also state reasons for it.

The Statement of Compliance must be substantiated with authenticated catalogue/data sheet/ manual with the page number of original catalogue/datasheet/manual of the relevant parameters indicated.



1. Delivery set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Delivery set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Delivery set for obs/ gyn. purpose	
2	Operational Requirements	
2.1	The Instruments must be made of	
2.1	highest quality materials e.g.	
	stainless steel (S/S) for metal	
	devices. The surgical instruments	
	must be CE marked / USFDA	
	approved. The bidder must	
	submit quality assurance (QA)	
2.2	certificates.	
2.2	Instruments must be made from	
	surgical quality, preferably non-	
	magnetic stainless steel and must	
	be matt surface finish. Quality	
	must comply with EN 46002 and	
	ISO 9002 and / or their latest	
2.2	amendments.	
2.3	Each pack must be packaged in a	
	hospital grade cotton wrapper	
	(autoclave-able) as a complete	
	pack. Bulk loose instrument	
	supply is NOT acceptable. Each	
	of the individual instruments of a	
	set must be packed in a labelled	
	clear plastic wrapper for easy	
	identification. All individually	
	packed instruments of a set shall	
	then be packed together in a	
	larger clear plastic wrapper	
	labelled with the name of the set	
2.4	for easy identification.	
2.4	Instrument surfaces must NOT be	
	stamped, indented or scratched. It	
	is preferred if the suppliers	
	labelled GoN name in anodised	
	form of labelling. It shall carry	
	clear anodised labelling/ marking	



S.N.	Purchaser's Specifications	Bidder's Remarks
5.11.	of manufacturer's name/ brand	
	and the part number/ model	
	number of instruments on the	
	surface of each piece of	
	instruments.	
2.5	Particular attention must be paid to the	
	quality of box joints to ensure	
	that they are smooth and interlock	
	well, and to teeth and grips to	
	ensure that they meet and	
	interlock well. Finger rings must	
	be of proper size and shape for	
	maximum utility and comfort.	
	The inside of finger rings must be	
	well rounded and free of sharp	
	edges, rough areas and grinding	
	marks, cracks, overlaps, burrs.	
2.6	Jaw serration must be well cut and	
	defined and must mesh properly	
	when the jaws are fully closed.	
	The edges of the serration must	
	be well chamfered and must not	
	contain burrs and sharp edges.	
	Teeth must be sharp (unless	
	otherwise specified), of proper	
	size and shape, free of rough	
	edges or burrs, and must mesh	
	with sufficient accuracy to ensure	
	proper performance for the use	
2.5	intended.	
2.7	Ratchet and ratchet catches must be	
	properly aligned and undercut for	
	safe locking. Ratchets must be of	
	such design as to ensure easy and	
	positive engagement and proper	
	disengagement. Ratchets and ratchet catches must be free of	
2.8	burrs and sharp edges.	
2.0	Locks, forceps and similar instruments must be of the box lock type or	
	lap joint type. All type of locks	
	must be accurately fitted, without	
	stiffness and without crevices,	
	burrs or sharp edges anywhere in	
	the construction.	
2.9	Screws of screw lock scissors and other	
2.7	Belews of sciew fock seissofs and other	



S.N.	Purchaser's Specifications	Bidder's Remarks
5.11.	instruments must be the	
	concentrically mustered type,	
	countersunk, flush with, or	
	slightly below the surface or	
	rounded, smooth and flush at the	
	periphery, but not riveted. The	
	screw must retain their position	
	after setting without binding or	
	loosening during use.	
	Scissors	
2.10	The ROCKWELL hardness of the	
	finished instruments must be	
	within the range from 50 HRC to	
	58 HRC. Opposite blades must	
	not vary in hardness by more than	
	4 units on the ROCKWELL C	
	harness scale.	
2.11	Scissors must have joints, which move	
	smoothly and must be neither too	
	loose nor too tight: it must be	
	possible to close and reopen the	
	instrument easily with two	
	fingers.	
2.12	The cutting ability of the instrument	
2.12	must be tested. The instrument	
	must cut clearly without tearing.	
2.13	The finish and all edges and surfaces	
2.13	must be uniform and free of	
	burrs, sharp edges (except where	
	required), pores, crevices, gins	
	1 / 1	
	marks, rough areas, cracks and	
2	overlaps.	
3.1	System Configuration	
	Delivery Set	
4	Technical Specifications	
4.1	The instruments required are listed	
	below , bidder MUST provide full	
	description of all instruments	
	(description includes: full name,	
	type, shape, design, full length,	
	volume and etc.) required below	
	for the evaluation.	
4.2	Instruments Required	
	1 x Sponge Holding Forceps Rampley	
	9.5"	



CN	Durahasan's Specifications	Bidder's Remarks
S.N.	Purchaser's Specifications 1 x Kidney Dish 10"	Diduct 5 Nelliai K5
	1 x Bowl 200mm	
	1 x Umbilical Scissors,	
	American, Straight 4.5"	
	1 x Needle Holder Mayo Hegar	
	Straight 6.5"	
	1 x Episiotomy Scissors 14.5 cm	
	1 x Cusco Vaginal Speculum	
	Medium	
	1 x Hartmann Mosquito Artery	
	Forceps Straight 120mm	
	1 x Metal Catheter	
	1 x Uterine Curette	
	1 x Treves Dissecting Forceps,	
	Plain, 200mm	
	1 x Spencerwells Artery Forceps	
	Curved 6"	
	1 x Kocher Artery Forcep,	
	Toothed Straight 6"	
	1 x Amniotomy Forceps 260mm	
	1 x Scissors Mayo Curved 6.5"	
4.3	1 x Auvard Vaginal Speculum	
4.3	All instruments must be supplied free of residual scale, acid, grease and	
	grinding and polishing materials	
	and workmanship must be first	
	class throughout. Instruments	
	must be free of defects, which	
	detract from their appearance or	
	impair serviceability, proper	
	functioning and intended use.	
4.4	Bidder MUST attach product	
	catalogues with photos for all	
	instruments as mentioned. These	
	catalogues/photos MUST clearly	
	and correctly mark with non-	
	erasable making pen their	
	respective parameter line number	
	(shown on the left column) and	
	instrument name.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables	
	and parts required to operate the	
	equipment, including all standard	
	tools and cleaning and lubrication	
	materials, to be included in the	



S.N.	Purchaser's Specifications	Bidder's Remarks
	offer. Bidders must specify the	
	quantity of every item included in	
	their offer (including items not	
	specified above).	
6	Operating Environment	
6.1	The instrument offered shall be	
	designed to store and to operate	
	normally under the conditions of	
	the purchaser's country. The	
	conditions include Climate,	
	Temperature, Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007	
	for Medical Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA	
	approved product certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to	
	use.	
12	Documentation	
12.1	s/Instructions manual shall be provided	
	in English.	



2. Episiotomy set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Episiotomy set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Episiotomy also known as perineotomy, is a	
	surgical incision of the perineum and the	
	posterior vaginal wall generally done by	
	a midwife or obstetrician during second	
	stage of labour to quickly enlarge the	
	opening for the baby to pass through.	
2	Operational Requirements	
2.1	The Instruments must be made of highest quality	
	materials e.g. stainless steel (S/S) for metal	
	devices. The surgical instruments must be	
	CE marked / USFDA approved. The bidder	
	must submit quality assurance (QA)	
	certificates.	
2.2	Instruments must be made from surgical quality,	
	preferably non-magnetic stainless steel and	
	must be matt surface finish. Quality must	
	comply with EN 46002 and ISO 9002 and /	
	or their latest amendments.	
2.3	Each pack must be packaged in a hospital grade	
	cotton wrapper (autoclave-able) as a	
	complete pack. Bulk loose instrument supply	
	is NOT acceptable. Each of the individual	
	instruments of a set must be packed in a	
	labelled clear plastic wrapper for easy	
	identification. All individually packed	
	instruments of a set shall then be packed	
	together in a larger clear plastic wrapper	
	labelled with the name of the set for easy	
2.4	identification.	
2.4	Instrument surfaces must NOT be stamped,	
	indented or scratched. It is preferred if the	
	suppliers labelled GoN name in anodised	
	form of labelling. It shall carry clear	
	anodised labelling/ marking of	
	manufacturer's name/ brand and the part	
	number/ model number of instruments on	



S.N.	Purchaser's Specifications	Bidder's Remarks
	the surface of each piece of instruments.	
2.5	Particular attention must be paid to the quality of	
	box joints to ensure that they are smooth and	
	interlock well, and to teeth and grips to	
	ensure that they meet and interlock well.	
	Finger rings must be of proper size and	
	shape for maximum utility and comfort. The	
	inside of finger rings must be well rounded	
	and free of sharp edges, rough areas and	
	grinding marks, cracks, overlaps, burrs.	
2.6	Jaw serration must be well cut and defined and	
	must mesh properly when the jaws are fully	
	closed. The edges of the serration must be	
	well chamfered and must not contain burrs	
	and sharp edges. Teeth must be sharp (unless	
	otherwise specified), of proper size and	
	shape, free of rough edges or burrs, and must	
	mesh with sufficient accuracy to ensure	
	proper performance for the use intended.	
2.7	Ratchet and ratchet catches must be properly	
	aligned and undercut for safe locking.	
	Ratchets must be of such design as to ensure	
	easy and positive engagement and proper	
	disengagement. Ratchets and ratchet catches	
	must be free of burrs and sharp edges.	
2.8	Locks, forceps and similar instruments must be of	
	the box lock type or lap joint type. All type	
	of locks must be accurately fitted, without	
	stiffness and without crevices, burrs or sharp	
	edges anywhere in the construction.	
2.9	Screws of screw lock scissors and other	
	instruments must be the concentrically	
	mustered type, countersunk, flush with, or	
	slightly below the surface or rounded,	
	smooth and flush at the periphery, but not	
	riveted. The screw must retain their position	
	after setting without binding or loosening	
	during use.	
	Scissors	
2.10	The ROCKWELL hardness of the finished	
	instruments must be within the range from	
	50 HRC to 58 HRC. Opposite blades must	
	not vary in hardness by more than 4 units on	
	the ROCKWELL C harness scale.	
2.11	Scissors must have joints, which move smoothly	
	and must be neither too loose nor too tight: it	



S.N.		der's Remarks
	must be possible to close and reopen the	
	instrument easily with two fingers.	
2.12	The cutting ability of the instrument must be	
	tested. The instrument must cut clearly	
0.10	without tearing.	
2.13	The finish and all edges and surfaces must be	
	uniform and free of burrs, sharp edges	
	(except where required), pores, crevices, gins marks, rough areas, cracks and	
	gins marks, rough areas, cracks and overlaps.	
3	System Configuration	
3.1	Episiotomy Set	
4	Technical Specifications	
4.1	The instruments required are listed below,	
'''	bidder MUST provide full description of all	
	instruments (description includes: full name,	
	type, shape, design, full length, volume and	
	etc.) required below for the evaluation.	
4.2	Instruments Required	
	INSTRUMENTS BOX STAINLESS STEEL 25x12x6cm / 11x5x2½	4" 1
	FINE OPERATING IRIS SCISSORS 10 Cm / 4", STRAIGHT	1
	BRAUN-STADLER EPIOSOTOMY SCISSORS 14.0cm / 5 1/2"	1
	MAYO-STILLE OPERATING SCISSORS 17.0cm / 7", CURVED	1
	TISSUE FORCEPS 16.0cm / 6 1/4", 1 X 2 TEETH	1
	STANDARD DRESSING FORCEPS 16.0cm / 6 1/4"	1
	MAYO-HEGAR NEEDLE HOLDERS 16.0cm / 6 1/4"	1
	GROSS-MAIER DRESSING AND COTTON SWAB FORCEPS CVD	20.0cm / 8", 1
	DOUBLE BUTTONNED PROBES 16.0cm / 6 1/4", 2mm Ø	1
	ROCHESTER-PEAN ARTERY FORCEPS 18.0cm / 7", STRAIGH	Т 2
	PROBES 16.0cm / 6 1/4"	1
	ADSON RETRACTOR 16.0cm / 6 1/4", 3 X 4 TEETH, BLUNT	1
	BACKHAUS TOWEL FORCEPS 11.0cm / 4 ½"	4
4.3	All instruments must be supplied free of residual	
	scale, acid, grease and grinding and	
	polishing materials and workmanship must	
	be first class throughout. Instruments must	
	be free of defects, which detract from their	
	appearance or impair serviceability, proper	
4.4	functioning and intended use. Bidder MUST attach product catalogues with	
4.4	photos for all instruments as mentioned.	
	These catalogues/photos MUST clearly and	
	correctly mark with non-erasable making	
	pen their respective parameter line number	
	The man of the man of	



S.N.	Purchaser's Specifications	Bidder's Remarks
	(shown on the left column) and instrument	
ı	name.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables and parts	
İ	required to operate the equipment, including	
İ	all standard tools and cleaning and	
İ	lubrication materials, to be included in the	
İ	offer. Bidders must specify the quantity of	
İ	every item included in their offer (including	
	items not specified above).	
6	Operating Environment	
6.1	The instrument offered shall be designed to store	
İ	and to operate normally under the conditions	
İ	of the purchaser's country. The conditions	
ı	include Climate, Temperature, Humidity,	
7	etc.	
7.1	Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for	
/.1	Medical Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA approved	
1.4	product certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During Warranty Period	
10.1	Standard warranty conditions are applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to use.	
12	Documentation	
12.1	s/Instructions manual shall be provided in English.	



3. Peri-Light

S.N.	Purchaser's Specifications	Bidder's Remarks
100210	Peri-Light	
	Manufacturer	
	Brand	
	Type/Model	
	Countryof Origin	
1	Description of Function	
1.1	To be used in labour room in hospital and for	
	Postpartum Perineal care of patient.	
2	Operational Requirements	
2.1	Shall operate on mains AC supply.	
3	System Configuration	
3.1	Peri-light, complete unit.	
4	Technical Specifications	
4.1	Light weight and easy to carry.	
4.2	The lamp can easily be positioned up to 40 degrees	
	backwards.	
4.3	Power: 50 Watt or more with extra focus. Bidder to	
	specify the power of lamp.	
4.4	Shall have on/off switch.	
4.5	Shall have facility to focus the light on specific area.	
4.6	Shall have long lifespan of lamp. Bidder to specify	
	the lifespan of lamp.	
5	Accessories, spares and consumables	
5.1	Accessories:	
	• Spare lamp: 01 no.	
5.2	All standard accessories, consumables and parts	
	required to operate the equipment, including all	
	standard tools and cleaning and lubrication	
	materials, to be included in the offer. Bidders	
	must specify the quantity of every item included	
	in their offer (including items not specified	
	above).	
6	Operating Environment	
6.1	The system offered shall be designed to store and to	
	operate normally under the conditions of the	
	purchaser's country. The conditions include	
	Power Supply, Climate, Temperature,	
	Humidity, etc.	
6.2	Power supply: 220-240V/ 50 Hz AC Single phase	
	fitted with appropriate plug to meet purchaser's	
	country requirements. The power cable must be	
	minimum 3 metres long.	



7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007 for Medical	
	Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA approved	
	product certificate.	
7.3	Electrical safety conforms to standards for electrical safety	
	IEC 60601-1 General requirement for Electrical	
_	safety of Medical Equipment.	
8	User Training	
8.1	Must provide user training (including how to use and	
	maintain the equipment).	
9	Warranty	
9.1	Comprehensive warranty for 1 year.	
10	Maintenance Service During Warranty Period	
10.1	Standard warranty conditions are applicable.	
11	Installation, Inspections and Commissioning	
11.1	Must supply preassembled unit ready to use.	
12	Documentation	
12.1	User (Operating) manual in English.	
12.2	Service (Technical / Maintenance) manual in English.	
12.3	List of important spare parts and accessories with	
	their part numbers and costing.	

4. Electric Suction Pump (Surgical Aspirator)

S.N.	Purchaser's Specifications	
	Electric Suction Pump, Twin type (Surgical Aspirator)	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	To extract fluid from the body during surgery or	
	emergency treatments.	
2	Operational Requirements	
2.1	An electric double jar suction pump for surgical use.	
3	System Configuration	
3.1	Suction machine with two bottles and accessories.	
4	Technical Specifications	
4.1	It shall be mounted on four robust, fully 360 degree	
	swivelling, antistatic, non-marking grey tires	
	castors, minimum size 75 mm with at least 2	
	diagonal brakes.	
4.2	Come with suction controller and vacuum gauge /	



S.N.	Purchaser's Specifications	
	indicator.	
4.3	The pump shall be oil free vacuum pump where the	
1.5	pumped liquid shall be sealed off from the	
	pump.	
4.4	Come with overflow control valves.	
	Bidder shall provide technical design and details of	
	the pump with this TSF	
4.5	Vacuum rate shall be from 0 to not less than 640	
	mmHg (0.85 bars).	
4.6	Air flow rate shall be at least 25 l/min.	
4.7	The pump shall come fitted with twin unbreakable,	
	transparent, autoclaveable polycarbonate	
4.0	suction bottles minimum 2 litre each.	
4.8	The bottles shall be incorporated with an automatic	
	suction cut-off mechanism when they become full.	
4.9	The suction bottles shall come with overflow lid.	
4.10	Noise level: not more than 55 dBA.	
4.11	Air discharge from pump shall be filtered by a 0.3	
	micron bacterial hydrophobic filter.	
5	Accessories, spares and consumables	
5.1	Accessories:	
	• Electrical cable: 1 minimum 3 meter	
	length	
	• Clear suction tubing: 1 set of 5 meter length	
	• Bacterial filter: 0.3 micron, 10 pcs	
	• Spare unbreakable, transparent, autoclave able	
	polycarbonate suction bottle 2L: 1pc	
	• Complete connection tubing set: 1 set	
	• Hand switch & foot switch with cables for	
	operating easily.	
6	Operating Environment	
6.1	The product offered shall be designed to be stored and to operate normally under the conditions of	
	the purchaser's country. The conditions include	
	_ · · · · · · · · · · · · · · · · · · ·	
1	Power Supply. Climate. Temperature	
	Power Supply, Climate, Temperature, Humidity, etc.	
6.2	Power Supply, Climate, Temperature, Humidity, etc. Must operate on 220-240V AC as well as	
6.2	Humidity, etc.	
6.2	Humidity, etc. Must operate on 220-240V AC as well as	
	Humidity, etc. Must operate on 220-240V AC as well as rechargeable batteries. Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for Medical	
7 7.1	Humidity, etc. Must operate on 220-240V AC as well as rechargeable batteries. Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for Medical Devices AND	
7	Humidity, etc. Must operate on 220-240V AC as well as rechargeable batteries. Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for Medical	



S.N.	Purchaser's Specifications	
7.3	Shall meet IEC-60601-1-2 General Requirements of	
	Safety for equipment.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During Warranty Period	
10.1	During warranty period supplier must ensure	
	corrective/breakdown maintenance whenever	
	required.	
11	Installation, Inspections and Commissioning	
11.1	Must supply preassembled unit, ready to use.	
11.2	Inspections to verify the compliance of the offered	
	equipment as per specifications will be	
	conducted by the technical team appointed by	
	the purchaser.	
12	Documentation	
12.1	User (Operating) and Service	
	(Technical/Maintenance) manuals to be	
	supplied in English.	
12.2	Certificate of calibration and inspection.	
12.3	List of important spare parts and accessories with	
	their part numbers and costing	



5. Nebuliser with Diaphragm Type Compressor

S.N	Purchaser's Specifications	
	Nebuliser with Diaphragm Type Compressor	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Nebuliser is a device used to administer medication	
	to people in forms of a liquid mist to the	
	airways. It is commonly used in treating cystic	
	fibrosis, asthma, and other respiratory diseases	
2	Operational Requirements	
2.1	Lightweight and compact, equipped with Diaphragm	
	type compressor.	
3	System Configuration	
3.1	Nebuliser with Diaphragm type Compressor with	
	complete accessories.	
4	Technical Specifications	
4.1	Shall be compact, lightweight and low noise.	
4.2	Durable long life Diaphragm compressor. Suitable	
	for heavy duty/ institutional (hospital) use,	
	must be able to run uninterruptedly for one hour.	
4.3	Must have Maximum air pressure Between 25-40	
7.5	psi.	
4.4	Operating Air Pressure Between 8-14.5 psi	
4.5	Must produce particle of size 0.5-10μm	
4.6	Must have a dust filter.	
4.7	Must be able to deliver a flow rate > 7 l/min.	
4.8	Must have a check valve to protect the device against	
	contamination due to backward inhalation.	
4.9	Must be compatible for continuous use.	
4.10	Shall provide ABS case with handle for easy	
	carrying.	
4.11	Medication capacity:- 5ml	
4.12	Average nebulization rate:- Approximate 0.2 ml/min.	
5	Accessories, spares and consumables	
5.1	Accessories:	
	 Nebuliser bulb reusable, autoclaveable- 01 	
	no.	
	 Adult and child face mask reusable, 	
	autoclaveable- 02 each.	



S.N	Purchaser's Specifications	
	• T piece, Mouthpiece, Nosepiece, reusable, autoclaveable- 01 each.	
	Mouthpiece- 01 no. Noscepiese 01 no.	
	• Nosepiece- 01 no.	
	• 1 x 200 cm. tubing	
5.2	• Spare filters- 10 nos.	
5.2	All standard accessories, consumables and parts	
	required to operate the equipment, including all standard tools and cleaning and lubrication	
	materials, to be included in the offer. Bidders	
	must specify the quantity of every item	
	included in their offer (including items not	
	specified above).	
6	Operating Environment	
6.1	The system offered shall be designed to operate	
	normally under the conditions of the	
	purchaser's country. The conditions include	
	Power Supply, Climate, Temperature,	
	Humidity, etc.	
6.2	Power supply: 220 - 240 VAC, 50Hz fitted with	
	appropriate plug. The power cable must be at	
	least 3 metre in length.	
7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007 for Medical	
	Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA approved	
	product certificate.	
7.3	Electrical safety conforms to standards for electrical	
	safety IEC 60601-1 General requirement for	
8	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment.	
8	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training	
8 8.1	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and	
8.1	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment).	
	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and	
8.1 9	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty	
8.1 9 9.1	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance.	
8.1 9 9.1 10	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period	
8.1 9 9.1 10	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period During the warranty period supplier must ensure	
8.1 9 9.1 10	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period During the warranty period supplier must ensure corrective/breakdown maintenance whenever required. Installation and Commissioning	
8.1 9 9.1 10 10.	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period During the warranty period supplier must ensure corrective/breakdown maintenance whenever required. Installation and Commissioning Supplier must accomplish proper commissioning of	
8.1 9 9.1 10 10.	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period During the warranty period supplier must ensure corrective/breakdown maintenance whenever required. Installation and Commissioning Supplier must accomplish proper commissioning of equipment onsite.	
9.1 10.	safety IEC 60601-1 General requirement for Electrical safety of Medical Equipment. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year after acceptance. Maintenance Service During Warranty Period During the warranty period supplier must ensure corrective/breakdown maintenance whenever required. Installation and Commissioning Supplier must accomplish proper commissioning of	



S.N	Purchaser's Specifications	
12.	Service (Technical / Maintenance) manual in English.	
12.	List of important spare parts and accessories with their part number and costing.	
12.	Certificate of calibration and inspection from factory.	



6. Otoscope

S.N.	Purchaser's Specifications	Bidder's Remarks
	Otoscope	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	An otoscope is a medical device which is used to	
	look into the ears. It is used to screen for	
	illness during regular check-ups and also to	
	investigate ear symptoms.	
	An otoscope potentially gives a view of the	
	ear canal and tympanic membrane or	
	eardrum.	
2	Operational Requirements	
2.1	Otoscope with fibreoptic illumination.	
3	System Configuration	
3.1	Otoscope complete with Halogen bulb, Pneumatic	
	bag, Reusable and autoclavable speculum	
	set, Heavy duty handles with charger.	
4	Technical Specifications	
4.1	It should consist of Otoscope with fibreoptic	
	illumination.	
4.2	3.5 volts Halogen bulb.	
4.4	Magnification, 3 or 4 times.	
4.5	Pneumatic bag for Sieglisation of tympanic	
	membrane	
4.6	Reusable and autoclavable speculum set of 4 or 5.	
4.7	Heavy duty handles with charger and chargeable	
	long life battery. Charging should be	
	possible through direct main line supply.	
5	Accessories, spares and consumables	
5.1	Accessories:	
	One spare battery.	
	• One spare 3.5 V Halogen bulb.	
	Reusable and autoclavable speculum set of	
	4 or 5 (2 sets for each otoscopes).	
5.2	All standard accessories, consumables and parts	
	required to operate the equipment, including	
	all standard tools and cleaning and	
	lubrication materials, to be included in the	
	offer. Bidders must specify the quantity of	
	every item included in their offer (including	



Purchaser's Specifications	Bidder's Remarks
items not specified above).	
Operating Environment	
The system offered shall be designed to be stored	
and to operate normally under the	
1	
•	
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1 2	
and maintain the equipment).	
Warranty	
Comprehensive warranty for 1 year.	
Maintenance Service During Warranty Period	
Must supply preassembled unit, ready to use.	
Documentation	
User (Operating) manual in English.	
Service (Technical / Maintenance) manual in	
English.	
List of important spare parts and accessories with	
their part numbers and costing.	
	Items not specified above). Operating Environment The system offered shall be designed to be stored and to operate normally under the conditions of the purchaser's country. The conditions include, Climate, Temperature, Humidity, etc. Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for Medical Devices AND CE (93/42 EEC Directives) or USFDA approved product certificate. User Training Must provide user training (including how to use and maintain the equipment). Warranty Comprehensive warranty for 1 year. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Must supply preassembled unit, ready to use. Documentation User (Operating) manual in English. Service (Technical / Maintenance) manual in English. List of important spare parts and accessories with

7. IUD & Implant Insertion and Removal Kit

S.N.	Purchaser's Specifications	
	Implant Insertion and Removal Kit	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	It is used in the procedure of implants insertion and removal.	
2	Operational Requirements	
2.1	All instruments must be made of highest quality materials e.g. stainless steel (S/S) for metal devices. The surgical instruments must be CE marked / USFDA approved. The bidder must submit quality assurance (QA) certificates.	



	73	
S.N.	Purchaser's Specifications	
2.2	Instruments must be made from surgical quality,	
	preferably non-magnetic stainless steel and	
	must be matt surface finish. Quality must	
	comply with EN 46002 and ISO 9002	
	and/or their latest amendments.	
	The material shall be fret free. The material must	
	be totally resistant to corrosion and pitting.	
2.3	Each pack must be packaged in a hospital grade	
	cotton wrapper (autoclaveable) as a	
	complete pack and the wrapped packs must	
	be packaged in a labelled clear plastic box.	
	The remaining packs shall be in the cotton	
	wrapper only. Bulk loose instrument	
	supply is NOT acceptable. Each of the	
	individual content of the packs must be in a	
	clear plastic wrapper labelled on the outside	
	for easy identification of the individual	
	instruments.	
3	System Configuration	
3.1	The instruments must comprise of:	
	 Implant insertion and removal set 	
	consisting different kinds of instruments	
	as described below.	
	 IUD implant insertion and removal set 	
	consist of different kinds of instruments as	
	described below	
4	Technical Specifications	
	Detail list of Instruments under different	Qty
	packs NB: All sizes indicating length	Per
	may vary +/- 3mm, other sizes +/- 2mm	Pack
4.1	Implant Insertion and Removal Kit	
4.1.1	Gallis pot; Galli pot – 4 onz, 1.5" height; SS	1
4.1.2	Kidney tray 10" (Medium size), SS	1
4.1.3	Sponge holder, 9.5", straight	2
4.1.4	Volselum, 9.5", curved	1
4.1.5	Dressing forceps (Tooth) 6"	1
4.1.6	Dressing forceps (Non tooth) 6"	1
4.1.7	Speculum (Graves or Sim's) medium size	1
4.1.8	Tubal hook	1
4.1.9	Forceps Baby Babcock, 6.5"	2
4.1.10	Elevator, Uterine, Ramathobodi, SS (Guard 1"	1
	diameter, tip length 2")	
4.1.11	Metallic catheter no. 14	1
4.1.12	Retractor (Richardson-Eastman), single blade	1
4.1.13	Retractor (Richardson-Eastman), double blade	2
4.1.13	rectactor (rectactor Eastman), acaste clade	-



S.N.	Purchaser's Specifications	
4.1.14	Forceps Allis, 6.5", SS (Tooth)	2
4.1.15	Mosquito forceps 5" straight	3
4.1.16	Mosquito forceps 5" curved	3
4.1.17	Mayo operating scissors, 7", curved end	1
4.1.18	Metzenbaum scissors (Straight) 6"	1
4.1.19	Surgical handle no. 3	1
4.1.20	Surgical blade no. 10	1
4.2	IUD Insertion / Removal Kit	Qty
		Per
		Pack
	Insertion	
4.2.1	Gallis pot; Galli pot – 4 onz, 1.5" height; SS	1
4.2.2	Kidney tray 10" (Medium size), SS	1
4.2.3	Forcep arterty, Pean, straight 8 ½ "	1
4.2.4	Volselum, 9.5", curved	1
4.2.5	Sponge holder, 9.5", straight	1
4.2.6	Mayo dissecting scissors SS curved 7"	1
4.2.7	Uterine sound, 12.5" metallic	1
4.2.8	Bi-valve Vaginal speculum medium	1
	Removal	1
4.2.9	Gallis pot; Galli pot – 4 onz, 1.5" height; SS	1
4.2.10	Kidney tray 10" (Medium size), SS	1
4.2.11	Forcep arterty, Pean, straight 8 ½ "	1
4.2.12	Volselum, 9.5", curved	1
4.2.13	Sponge holder, 9.5", straight	1
4.2.14	Bi-valve Vaginal speculum medium	1
4.3	All instruments must be supplied free of residual	
	scale, acid, grease and grinding and	
	polishing materials and workmanship must	
	be first class throughout. Instruments must	
	be free of defects, which detract from their	
	appearance or impair serviceability, proper	
	functioning and intended use.	
4.4	Bidder MUST attach product catalogues with	
	photos for all instruments as mentioned.	
	These catalogues/photos MUST clearly and	
	correctly mark with non-erasable making	
	pen their respective parameter line number	
	(shown on the left column) and instrument	
_	name.	
5	Accessories, spares and consumables	
5.1	Shall provide storage case of good quality for all	
5.0	above surgical instruments.	
5.2	All standard accessories, consumables and parts	
	required to operate the equipment,	



S.N.	Purchaser's Specifications	
50110	including all standard tools and cleaning	
	and lubrication materials, to be included in	
	the offer. Bidders must specify the quantity	
	of every item included in their offer	
	(including items not specified above).	
6	Operating Environment	
6.1	The instrument offered shall be designed to store	
	and to operate normally under the	
	conditions of the purchaser's country. The	
	conditions include Climate, Temperature,	
	Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007 for	
	Medical Devices and must meet ISO 7153-	
	1:1991 for surgical instruments and ASTM	
	F899 - 12b AND	
7.2	CE (93/42 EEC Directives) or USFDA approved	
	product certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year after acceptance.	
10	Maintenance Service During Warranty Period	
10.1	Standard warranty conditions are applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to use.	
12	Documentation	
12.1	's (Operating) and /or Technical/Maintenance	
	manual shall provide in English.	



8. Dental Instruments Set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Dental Instruments Set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Instrument set for dental examination,	
	dental surgery and dental	
	treatment	
2	Operational Requirements	
2.1	The Dental Instruments must be made	
	of highest quality materials e.g.	
	stainless steel (S/S) for metal	
	devices. The surgical instruments	
	must be CE marked / USFDA	
	approved. The bidder must submit	
	quality assurance (QA)	
	certificates.	
2.2	Instruments must be made from surgical	
	quality, preferably non-magnetic	
	stainless steel and must be matt	
	surface finish. Quality must	
	comply with EN 46002 and ISO	
	9002 and / or their latest	
	amendments.	
2.3	Each pack must be packaged in a	
	hospital grade cotton wrapper	
	(autoclave-able) as a complete	
	pack. Bulk loose instrument	
	supply is NOT acceptable. Each of	
	the individual instruments of a set	
	must be packed in a labelled clear	
	plastic wrapper for easy	
	identification. All individually	
	packed instruments of a set shall	
	then be packed together in a larger	
	clear plastic wrapper labelled with	
	the name of the set for easy	
	identification.	
2.4	Instrument surfaces must NOT be	
	stamped, indented or scratched. It	
	is preferred if the suppliers	



S.N.	Purchaser's Specifications	Bidder's Remarks
ו1 (•	labelled GoN name in anodised	DAWN DAWN AND
	form of labelling. It shall carry	
	clear anodised labelling/ marking	
	of manufacturer's name/ brand	
	and the part number/ model	
	number of instruments on the	
	surface of each piece of	
	instruments.	
2.5	Particular attention must be paid to the	
	quality of box joints to ensure that	
	they are smooth and interlock	
	well, and to teeth and grips to	
	ensure that they meet and	
	interlock well. Finger rings must	
	be of proper size and shape for	
	maximum utility and comfort. The	
	inside of finger rings must be well	
	rounded and free of sharp edges,	
	rough areas and grinding marks,	
	cracks, overlaps, burrs.	
2.6	Jaw serration must be well cut and	
	defined and must mesh properly	
	when the jaws are fully closed.	
	The edges of the serration must be	
	well chamfered and must not	
	contain burrs and sharp edges.	
	Teeth must be sharp (unless	
	otherwise specified), of proper	
	size and shape, free of rough	
	edges or burrs, and must mesh	
	with sufficient accuracy to ensure	
	proper performance for the use	
	intended.	
2.7	Ratchet and ratchet catches must be	
	properly aligned and undercut for	
	safe locking. Ratchets must be of	
	such design as to ensure easy and	
	positive engagement and proper	
	disengagement. Ratchets and	
	ratchet catches must be free of	
2.0	burrs and sharp edges.	
2.8	Locks, forceps and similar instruments	
	must be of the box lock type or lap	
	joint type. All type of locks must	
	be accurately fitted, without	
	stiffness and without crevices,	



S.N.	Purchaser's Specifications	Bidder's Remarks
10 12 10	burrs or sharp edges anywhere in	
	the construction.	
2.9	Screws of screw lock scissors and other	
	instruments must be the	
	concentrically mustered type,	
	countersunk, flush with, or	
	slightly below the surface or	
	rounded, smooth and flush at the	
	periphery, but not riveted. The	
	screw must retain their position	
	after setting without binding or	
	loosening during use.	
	Scissors	
2.10	The ROCKWELL hardness of the	
	finished instruments must be	
	within the range from 50 HRC to	
	58 HRC. Opposite blades must not	
	vary in hardness by more than 4	
	units on the ROCKWELL C	
	harness scale.	
2.11	Scissors must have joints, which move	
	smoothly and must be neither too	
	loose nor too tight: it must be	
	possible to close and reopen the	
2.12	instrument easily with two fingers.	
2.12	The cutting ability of the instrument	
	must be tested. The instrument	
2.12	must cut clearly without tearing.	
2.13	The finish and all edges and surfaces	
	must be uniform and free of burrs,	
	sharp edges (except where required), pores, crevices, gins	
	marks, rough areas, cracks and	
	overlaps.	
3	System Configuration	
3.1	Dental Instruments Set	
4	Technical Specifications	
4.1	The instruments required are listed	
	below, bidder MUST provide full	
	description of all instruments	
	(description includes: full name,	
	type, shape, design, full length,	
	volume and etc.) required below	
	for the evaluation.	



S.N.	Purchaser's Specifications	Bidder's Remarks
4.2	Instruments Required	
I	Explorer	
II	Mouth mirror with handle	
III	Tweezers	
IV	Surgery (Extraction of teeth)	
i	Adult forceps:	
	Upper	
	 Upper Incisor forceps 	
	 Upper premolar forceps 	
	 Upper right molar forceps 	
	 Upper left molar forceps 	
	• Upper last molar (Wisdom teeth)	
	forceps (Bennet forceps)	
	Lower	
	 Lower incisor forceps 	
	 Lower premolar forceps 	
	 Lower molar forceps 	
ii	Paediatric forceps:	
	Upper Incisor	
	• Upper molar (right & left)	
	 Lower incisor 	
	 Lower molar 	
V	Elevators:	
	 Periosteal elevator (small & 	
	large size)	
	Straight Elevator (small & large)	
	size)	
	Cryer elevator (right & left)	
	• Cross bar elevator (right & left)	
	• Excavator (small & large)	
	Surgical Blade Handle	
VI	Instruments for filling:	
	Glass slab	
	Cement Spatula	
	Plastic Instrument	
VII	Periodontal Instruments:	
	Periodontal probe	
	Interdental scalar	
	Periscaler (Right & left)	
	• Sickle scalar (right & left)	
	• Curette-small size (right & left)	
	Curette-big size (right & left)	
VIII	Instrument tray, stainless steel of	
	suitable size.	



S.N.	Purchaser's Specifications	Bidder's Remarks
4.3	All instruments must be supplied free of	
	residual scale, acid, grease and	
	grinding and polishing materials	
	and workmanship must be first	
	class throughout. Instruments	
	must be free of defects, which	
	detract from their appearance or	
	impair serviceability, proper	
	functioning and intended use.	
4.4	Bidder MUST attach product catalogues	
	with photos for all instruments as	
	mentioned. These	
	catalogues/photos MUST clearly	
	and correctly mark with non-	
	erasable making pen their	
	respective parameter line number	
	(shown on the left column) and	
	instrument name.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables	
3.1	and parts required to operate the	
	equipment, including all standard	
	tools and cleaning and lubrication	
	materials, to be included in the	
	offer. Bidders must specify the	
	quantity of every item included in	
	their offer (including items not	
	specified above).	
6	Operating Environment	
6.1	The instrument offered shall be	
0.1	designed to store and to operate	
	normally under the conditions of	
	the purchaser's country. The	
	conditions include Climate,	
	Temperature, Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007	
/.1	for Medical Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA	
'	approved product certificate.	
8	User Training	
8.1	S	
	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	



S.N.	Purchaser's Specifications	Bidder's Remarks
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to use.	
12	Documentation	
12.1	s/Instructions manual shall be provided in	
	English.	



9. Incision and Drainage set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Incision and Drainage set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Incision and drainage and clinical	
	lancing are minor surgical	
	procedures to release pus or	
	pressure built up under the skin,	
	such as from an abscess, boil, or	
	infected paranasal sinus. This	
	allows the pus fluid to escape by	
	draining out through the incision.	
2	Operational Requirements	
2.1	It is performed by treating the area with	
	an antiseptic, such as iodine-	
	based solution, and then making a	
	small incision to puncture the	
	skin using a sterile instrument	
	such as a sharp needle, a	
	pointed scalpel or a lancet.	
3	System Configuration	
3.1	I & D Set	
4	Technical Specifications	
4.1	Items Required	
	• Artery Forcep (9x12)	•
	• Blade Holder 4'	
	• Kidney Tray (9x12)	
	 Tooth forcep/ thamb forcep 	
	 Thamb forcep plane 	
	Bowl small plane	
	Scissor plane	
	• Eye towel with rapping cloth	
	• Spoke	
	Surgical blade no 11	
4.4	Bidder MUST attach product	
	catalogues with photos for all	
	instruments as mentioned. These	
	catalogues/photos MUST clearly	
	and correctly mark with non-	



S.N.	Purchaser's Specifications	Bidder's Remarks
	erasable making pen their	20 20 20 20 20 20 20 20 20 20 20 20 20 2
	respective parameter line number	
	(shown on the left column) and	
	instrument name.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables	
	and parts required to operate the	
	equipment, including all standard	
	tools and cleaning and lubrication	
	materials, to be included in the	
	offer. Bidders must specify the	
	quantity of every item included in	
	their offer (including items not	
	specified above).	
6	Operating Environment	
6.1	The instrument offered shall be	
	designed to store and to operate	
	normally under the conditions of	
	the purchaser's country. The conditions include Climate.	
7	Temperature, Humidity, etc.	
7.1	Standards and Safety Requirements Must submit ISO13485:2003/AC:2007	
/.1	for Medical Devices AND	
7.2	CE (93/42 EEC Directives) or USFDA	
1.2	approved product certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to	
	use.	
12	Documentation	
12.1	s/Instructions manual shall be provided	
	in English.	



10. Suture set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Suture set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Complete instrument set for minor surgery:	
	suturing of wounds. Can be used as a	
	complementary set for isolated	
	skeletal traction or split skin grafting	
2	Operational Requirements	
2.1	The Instruments must be made of highest	
	quality materials e.g. stainless steel	
	(S/S) for metal devices. The surgical	
	instruments must be CE marked /	
	USFDA approved. The bidder must	
	submit quality assurance (QA)	
2.2	certificates.	
2.2	Instruments must be made from surgical	
	quality, preferably non-magnetic stainless steel and must be matt	
	surface finish. Quality must comply	
	with EN 46002 and ISO 9002 and / or	
	their latest amendments.	
2.3	Each pack must be packaged in a hospital	
2.3	grade cotton wrapper (autoclave-able)	
	as a complete pack. Bulk loose	
	instrument supply is NOT acceptable.	
	Each of the individual set must be	
	packed in a labelled clear plastic	
	wrapper for easy identification. All	
	individually packed instruments of a	
	set shall then be packed together in a	
	larger clear plastic wrapper labelled	
	with the name of the set for easy	
	identification.	
3	System Configuration	
3.1	Suture Set	
4	Technical Specifications	
4.1	The instruments required are listed	
	below, bidder MUST provide full	



S.N.	Purchaser's Specifications	Bidder's Remarks
	description of all items (description	
	includes: full name, type, shape,	
	design, full length, volume and etc.)	
	required below for the evaluation.	
4.2	Items Required	
	KIDNEY DISH, large, 275x150x45mm, stainless steel =1	
	SCALPEL, HANDLE, No 3 (for blades 10/11/15)=1	
	SCISSORS, MAYO, 17 cm, curved=1 FORCEPS, DRESSING, BLANK, 14.5 cm, atraumatic serration =1	
	FORCEPS, TISSUE, LANE, 14cm, 1x2 teeth=1	
	NEEDLE HOLDER, MAYO-HEGAR, 15 cm, standard=1	
	FORCEPS, HEMOSTATIC, CRILE, 14 cm, curved=4 FORCEPS, HEMOSTATIC, KOCHER, 14 cm/1x2 teeth, straight=2	
	FORCEPS, TOWEL CLAMP, BACKAUS, 13 cm=5 FORCEPS, SPONGE, FOERSTER, 24cm, serrated jaws, straight=2	
	BOWL, ROUND, 100 ml, 80 x 35 mm, stainless =1steel	
4.3	Bidder MUST attach product catalogues	
	with photos for all instruments as	
	mentioned. These catalogues/photos	
	MUST clearly and correctly mark	
	with non-erasable making pen their	
	respective parameter line number	
	(shown on the left column) and	
	instrument name.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables and	
	parts required to operate the	
	equipment, including all standard	
	tools and cleaning and lubrication	
	materials, to be included in the offer.	
	Bidders must specify the quantity of	
	every item included in their offer	
	(including items not specified above).	
6	Operating Environment	
6.1	The instrument offered shall be designed to	
	store and to operate normally under	
	the conditions of the purchaser's	
	country. The conditions include	
	Climate, Temperature, Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit ISO13485:2003/AC:2007 for	
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S.N.	Purchaser's Specifications	Bidder's Remarks
7.2	CE (93/42 EEC Directives) or USFDA	
	approved product certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During Warranty	
	Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to use.	
12	Documentation	
12.1	s/Instructions manual shall be provided in	
	English.	

11. Sphygmomanometer (BP apparatus)

S.N.	Purchaser's Specifications	Bidder's Remarks
	Sphygmomanometer (BP apparatus)	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Sphygmomanometer is a device used to	
	measure blood pressure, composed	
	of an inflatable cuff to restrict blood	
	flow, and a mechanical manometer	
	to measure the pressure.	
2	Operational Requirements	
2.1	Aneroid sphygmomanometer having a	
	dial to show clear numbers and	
	pointer / needle for measurement of	
	pressure.	
3	System Configuration	
3.1	 Aneroid sphygmomanometer 	
	 Cuffs for child size and for adult 	
	size (regular)	
	 Inflation bulb 	
	 Carrying pouch 	
4	Technical Specifications	
4.1	Packed in easy carrying high quality	
	pouch made of waterproof cloth to	
	accommodate cuff, and inflation bulb.	
	0 00101	



S.N.	Purchaser's Specifications	Bidder's Remarks
4.2	Gauge to be calibrated in 2 mm Hg units.	Didder 5 Remarks
4.3	Must provide blood pressure cuffs for	
	child size and for adult size	
	(regular).	
5	Accessories, spares and consumables	
5.1	Allstandard	
	accessories/consumables/parts	
	required for the proper operation of	
	the above item shall be included in	
	the offer. Bidders shall specify, in a	
	separate Excel worksheet, the	
	quantity and details of any items	
	included in this offer which have	
	not been specified in this Technical	
	Specifications Form.	
6	Operating Environment	
6.1	The product offered shall be designed to	
	be stored and to operate normally under the conditions of the	
	purchaser's country. The conditions	
	include Climate, Temperature,	
	Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit ISO 9001 or ISO	
' ' '	13485:2003/AC: 2007.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1year.	
10	Maintenance Service During Warranty	
	Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit, ready to	
	use.	
12	Documentation	
12.1	User's manual in English	



12. Stethoscope

	12. Stetnoscope	
S.N.	Purchaser's Specifications	Bidder's Remarks
	Stethoscope	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	The stethoscope is used for listening to	
	the beating heart of a human, or the	
	lungs. It is also used for listening to	
	the flow of the blood in the	
	surrounding area of the heart.	
2	Operational Requirements	
2.1	Dual type stethoscope - Physician's	
	stethoscope.	
3	System Configuration	
3.1	Stethoscope, dual cup/bell	
3.1	Tubes	
4	Technical Specifications	
4.1	Dual, cup/bell and diaphragm head	
4.1	Head and ear tube assembly to be made	
4.2	of non-ferrous metal,	
4.3	·	
4.3	Tubes to be synthetic material and ear tubes to have shaped plastic	
	cushion ends.	
_		
5 5.1	Accessories, spares and consumables All standard	
3.1		
	accessories/consumables/parts	
	required for the proper operation of the above item shall be included in	
	the offer. Bidders shall specify, in a separate Excel worksheet, the	
	quantity and details of any items included in this offer which have	
	not been specified in this Technical	
	Specifications Form.	
6	Operating Environment The product offered shall be designed to	
0.1		
	be stored and to operate normally under the conditions of the	
	purchaser's country. The conditions	
	include Climate, Temperature,	
	Humidity, etc.	



7	Standards and Safety Requirements	I
7.1	Must submit ISO 9001 or ISO	1
	13485:2003/AC: 2007.	
8	User Training	
8.1	Not applicable.	Ī
9	Warranty	Ī
9.1	Warranty for 1 year.	Ī
10	Maintenance Service During Warranty	Ī
	Period	
10.1	Standard warranty conditions are	Ī
	applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit, ready to	Ī
	use.	
12	Documentation	
12.1	User's manual in English	

13. Suture removal set

S.N.	Purchaser's Specifications	
	Suture set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Complete instrument set for suture	
	removal	
2	Operational Requirements	
2.1	The Instruments must be made of	
	highest quality materials e.g.	
	stainless steel (S/S) for metal	
	devices. The surgical	
	instruments must be CE marked /	
	USFDA approved. The bidder	
	must submit quality assurance	
	(QA) certificates.	
2.2	Instruments must be made from	
	surgical quality, preferably non-	
	magnetic stainless steel and must	
	be matt surface finish. Quality	
	must comply with EN 46002 and	
	ISO 9002 and / or their latest	
	amendments.	
2.3	Each pack must be packaged in a	
	hospital grade cotton wrapper	



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S.N.	Purchaser's Specifications	
	(autoclave-able) as a complete	
	pack. Bulk loose instrument	
	supply is NOT acceptable. Each	
	of the individual set must be	
	packed in a labelled clear plastic	
	wrapper for easy identification.	
	All individually packed	
	instruments of a set shall then be	
	packed together in a larger clear	
	plastic wrapper labelled with the	
	name of the set for easy	
	identification.	
3	System Configuration	
3.1	Suture Set	
4	Technical Specifications	
4.1	The instruments required are listed	
1.1	below, bidder MUST provide	
	full description of all items	
	(description includes: full name,	
	type, shape, design, full length,	
	volume and etc.) required below	
	for the evaluation.	
4.2	Items Required	
1.2	Dressing Forceps Serr 5"	1
	Spencer Stitch Scissors 3 1/2"	1
	Tray suture removal	1
	Staple remover	1
4.4	Bidder MUST attach product	1
4.4	catalogues with photos for all	
	instruments as mentioned. These	
	catalogues/photos MUST clearly	
	and correctly mark with non-	
	and correctly mark with non-	
	erasable making nen their	
	erasable making pen their	
	respective parameter line number	
	respective parameter line number (shown on the left column) and	
5	respective parameter line number (shown on the left column) and instrument name.	
5	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables	
5 5.1	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables	
	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables and parts required to operate the	
	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables and parts required to operate the equipment, including all	
	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and	
	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be	
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	respective parameter line number (shown on the left column) and instrument name. Accessories, spares and consumables All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be	



Purchaser's Specifications	1
(including items not specified	
above).	
Operating Environment	
The instrument offered shall be	
designed to store and to operate	
normally under the conditions of	
the purchaser's country. The	
conditions include Climate,	
Temperature, Humidity, etc.	
Standards and Safety Requirements	
Must submit ISO13485:2003/AC:2007	
for Medical Devices AND	
CE (93/42 EEC Directives) or USFDA	
approved product certificate.	
User Training	
Not applicable.	
Warranty	
Warranty for 1 year.	
Maintenance Service During	
Warranty Period	
Standard warranty conditions are	
applicable.	
Installation and Commissioning	
Must supply complete unit, ready to	
use.	
Documentation	
s/Instructions manual shall be provided	
in English.	
	Above). Operating Environment The instrument offered shall be designed to store and to operate normally under the conditions of the purchaser's country. The conditions include Climate, Temperature, Humidity, etc. Standards and Safety Requirements Must submit ISO13485:2003/AC:2007 for Medical Devices AND CE (93/42 EEC Directives) or USFDA approved product certificate. User Training Not applicable. Warranty Warranty for 1 year. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Must supply complete unit, ready to use. Documentation s/Instructions manual shall be provided



14. Dressing set

S.N.	Purchaser's Specifications	Bidder's Remarks
	Dressing set	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	A dressing is a sterile pad or	
	compress applied to	
	a wound to promote healing	
	and protect the wound from	
	further harm.	
2	Operational Requirements	
2.1	A dressing is designed to be in	
	direct contact with	
	the wound , as distinguished	
	from a bandage , which is	
	most often used to hold a	
	dressing in place	
3	System Configuration	
3.1	Dressing Set	
4	Technical Specifications	
4.1	The instruments required are	
	listed below, bidder MUST	
	provide full description of all	
	items (description includes:	
	full name, type, shape, design,	
	full length, volume and etc.)	
	required below for the	
	evaluation.	
4.2	Items Required	
	Thumb forceps/tooth forceps	
	Artery forceps	
	Bowl (120ml)	
	Instrument tray with cover	
	Wrapper	
	Scissor plane with suture cutting	
	scissor	
	Foreign Body Removal Set	
	Mosquito Forceps	
	Mosquito Artery Forceps	
	Nasal FB Removal Forceps	



S.N.	Purchaser's Specifications	Bidder's Remarks
D-14.	Thumb Tissue Forceps	Didder 5 Reliiai K5
	Blade holder	
	Nasal Speculum	
4.4		
4.4	_	
	catalogues with photos for all	
	instruments as mentioned.	
	These catalogues/photos	
	MUST clearly and correctly	
	mark with non-erasable	
	making pen their respective	
	parameter line number	
	(shown on the left column)	
	and instrument name.	
5	Accessories, spares and	
	consumables	
5.1	All standard accessories,	
	consumables and parts	
	required to operate the	
	equipment, including all	
	standard tools and cleaning	
	and lubrication materials, to	
	be included in the offer.	
	Bidders must specify the	
	quantity of every item	
	included in their offer	
	(including items not specified	
	above).	
6	Operating Environment	
6.1	The instrument offered shall be	
	designed to store and to	
	operate normally under the	
	conditions of the purchaser's	
	country. The conditions	
	include Climate,	
	Temperature, Humidity, etc.	
7	Standards and Safety	
	Requirements	
7.1	Must submit	
	ISO13485:2003/AC:2007 for	
	Medical Devices AND	
7.2	CE (93/42 EEC Directives) or	
	USFDA approved product	
	certificate.	
8	User Training	
8.1	Not applicable.	
U.1	1.00 appiroacio.	



S.N.	Purchaser's Specifications	Bidder's Remarks
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to	
	use.	
12	Documentation	
12.1	s/Instructions manual shall be provided in English.	



16. Proctoscope

S.N.	Purchaser's Specifications	Bidder's Remarks
5.11.	Proctoscope	Diduct 5 Remarks
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	A proctoscope is a hollow, tube-	
1.1	like speculum that is used for	
	visual inspection of the	
	rectum.	
	T Gottam.	
2	Operational Requirements	
2.1	Non-Disposable Kelly's rectal	
	speculum.	
3	System Configuration	
3.1	Kelly's rectal speculum with obturator	
	and proctoscope sheath.	
4	Technical Specifications	
4.1	It should consist of obturator and	
	proctoscope.	
4.2	Sizes of 22x100 mm.	
4.4	Material: High grade fully stainless	
	steel, corrosion resistance.	
4.5	Finish: Mirror finish.	
5	Accessories, spares and	
	consumables	
5.1	Accessories:	
	Not applicable.	
5.2	All standard accessories, consumables	
	and parts required to operate the	
	equipment, including all	
	standard tools and cleaning and	
	lubrication materials, to be	
	included in the offer. Bidders	
	must specify the quantity of	
	every item included in their	
	offer (including items not	
(specified above).	
6.1	Operating Environment	
0.1	The system offered shall be designed	
	to be stored and to operate	
	normally under the conditions of	
	the purchaser's country. The	



S.N.	Purchaser's Specifications	Bidder's Remarks
	conditions include, Climate,	
	Temperature, Humidity, etc.	
7	Standards and Safety Requirements	
7.1	Must submit	
	ISO13485:2003/AC:2007 for	
	Medical Devices AND	
8	User Training	
8.1	Must provide user training (including	
	how to use and maintain the	
	equipment).	
9	Warranty	
9.1	Comprehensive warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit, ready	
	to use.	
12	Documentation	
12.1	User (Operating) manual in English.	
12.2	Service (Technical / Maintenance)	
	manual in English.	
12.3	List of important spare parts and	
	accessories with their part	
	numbers and costing.	



17. Digital Thermometer

Digital Thermometer Manufacturer			1 Hermometer
Manufacturer Brand Type / Model Country of Origin	S.N.	Purchaser's Specifications	
Type / Model Country of Origin Description of Function 1.2 For measuring temperatures and displaying it with LCD/LED Operational Requirements 2.1 Portable, battery operated system is required 3. System Configuration 3.1 Digital Thermometer, portable, battery operated Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy: ± 0.1 °C 5 Accessories, spares and consumables 5.1 Accessories: • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements		Digital Thermometer	
Type / Model Country of Origin Description of Function 1.2 For measuring temperatures and displaying it with LCD/LED 2 Operational Requirements 2.1 Portable, battery operated system is required 3. System Configuration 3.1 Digital Thermometer, portable, battery operated 4 Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy: ± 0.1 °C 5 Accessories, spares and consumables 5.1 Accessories: • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The condition include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements		Manufacturer	
Country of Origin		Brand	
Country of Origin		Type / Model	
1.2 Por measuring temperatures and displaying it with LCD/LED 2 Operational Requirements 2.1 Portable, battery operated system is required 3. System Configuration 3.1 Digital Thermometer, portable, battery operated 4 Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy: ± 0.1 °C 5 Accessories; • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements			
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2.1 Portable, battery operated system is required 3. System Configuration 3.1 Digital Thermometer, portable, battery operated 4 Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy : ± 0.1 °C 5 Accessories, spares and consumables 5.1 Accessories: ■ Temperature probe-(surface and internal probes)-1 each parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements	2	Operational Requirements	
3.1 Digital Thermometer, portable, battery operated 4 Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy: ± 0.1 °C 5 Accessories, spares and consumables 5.1 Accessories: • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements	2.1		
4.1 Technical Specifications 4.1 Temperature measurement range: - 40 °C to 210 °C 4.2 LCD readout 4.3 Temperature measurement accuracy: ± 0.1 °C 5 Accessories, spares and consumables 5.1 Accessories: • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements	3.	System Configuration	
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5.1 Accessories: • Temperature probe-(surface and internal probes)-1 each 5.2 All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements	4.3	Temperature measurement accuracy : ± 0.1 °C	
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offer (including items not specified above). 6 Operating Environment 6.1 The system offered must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Climate, temperature and relative humidity. 6.2 Battery Operated 7 Standards and Safety Requirements			
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6.2 Battery Operated 7 Standards and Safety Requirements		· ·	
7 Standards and Safety Requirements		temperature and relative numbers.	
1	6.2	Battery Operated	
	7	Standards and Safety Requirements	
7.4 Must submit ISO 9001 or	7.4	Must submit ISO 9001 or	
ISO13485:2003/AC:2007 for Medical		ISO13485:2003/AC:2007 for Medical	
Devices AND			
7.5 CE or USFDA approved product certificate.			
7.6 Shall meet IEC-60601-1-2:2001 General	7.6	Shall meet IEC-60601-1-2:2001 General	



	Requirements of Safety for	
	Electromagnetic Compatibility.	
8	User Training	
8.1	Must provide user training (including how to	
	use and maintain the equipment).	
9	Warranty	
9.1	Comprehensive warranty for 1 year after	
	acceptance.	
10	Maintenance Service during Warranty	
	Period	
10.1	Standard warranty conditions are applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit, ready to use.	
12	Documentation	
12.1	User (Operating) manual in English	
12.2	Service (Technical / Maintenance) manual in	
	English	
12.3	List of important spare parts and accessories	
	with their part numbers and costing.	
12.4	Certificate of calibration and inspection from	
	factory.	



18. Tongue Depressor, Reusable

S.N.	Purchaser's Specifications	Bidder's Remarks
	Tongue Depressor, Reusable	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.3	A tongue depressor is used to depress the	
	tongue to allow for examination of	
	the mouth and throat.	
2	Operational Requirements	
2.2	Wieder / Andrew / Lack's, tongue	
	depressor, reusable.	
3	System Configuration	
3.1	Set of Tongue Depressors, stainless steel.	
4	Technical Specifications	
4.1	Sizes:	
	• Adult: 22mmwide.	
	Adolescence: 19mm wide.	
	• Children: 13mm wide.	
4.2	Material: High grade fully stainless steel,	
	corrosion resistance.	
4.3	Workmanship: Both ends slightly curved	
	in opposite direction.	
4.4	Finish: Bright polish, smooth surface	
	without any burr, pits and scratches.	
4.5	Autoclaveable.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables	
	and parts required to operate the	
	equipment, including all standard	
	tools and cleaning and lubrication	
	materials, to be included in the	
	offer. Bidders must specify the	
	quantity of every item included in	
	their offer (including items not	
_	specified above).	
6	Operating Environment	
6.1	The system offered must be designed to	
	store and be operated normally	
	under the condition of the	
	purchaser's Country. The conditions	
	include Climate, temperature and	
	relative humidity.	



S.N.	Purchaser's Specifications	Bidder's Remarks
7	Standards and Safety Requirements	
7.7	Must submit ISO 9001 or	
	ISO13485:2003/AC:2007 for	
	Medical Devices AND	
7.8	CE or USFDA approved product	
	certificate.	
8	User Training	
8.1	Must provide user training (including how	
	to use and maintain the equipment).	
9	Warranty	
9.1	Warranty for 1 year after acceptance.	
10	Maintenance Service during Warranty	
	Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit, ready to	
	use.	
12	Documentation	
12.1	User (Operating) manual in English.	
12.2	Service (Technical / Maintenance) manual	
	in English.	



19. Digital scale, Adult Weighing (Bathroom Type)

S.N.	Purchaser's Specifications	Bidder's Remarks
1000	Adult Weighing Scale, Bathroom	
	Type	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	Weighing the mass of a patient or user	
2	Operational Requirements	
2.1	Digital type adult weighing scale	
3	System Configuration	
3.1	Digital weighing scale-Adult	
	bathroom type with large LCD Display.	
4	Technical Specifications	
4.1	Battery operated.	
4.2	Battery power saving by switching off while no weight on the	
4.0	scale.	
4.3	Equipment must be simple to use, operate and maintain.	
4.4	Scale to weigh 0 to 150 Kg in division of 100g.	
5	Accessories, spares and	
	consumables	
5.1	All standard accessories/consumables/parts required for the proper operation of the above item shall be included in the offer. Bidders shall specify, in a separate Excel worksheet, the quantity and details of any items included in this offer which have not been specified in this Technical Specifications Form.	
6	Operating Environment	
6.1	The product offered shall be designed to be stored and to operate normally under the conditions of the purchaser's	



S.N.	Purchaser's Specifications	Bidder's Remarks
	country. The conditions	
	include Climate, Temperature,	
	Humidity, etc.	
7	Standards and Safety	
	Requirements	
7.1	Must submit ISO 9001 or ISO	
	13485:2003/AC: 2007.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply preassembled unit,	
	ready to use.	
12	Documentation	
12.1	Instructions manual supplied in	
	English.	



20. Laryngoscope, Magnifying Rigid (with Light Source)

S.N.	Purchaser's Specifications	Bidder's Remarks
	Laryngoscope, Magnifying Rigid	
	(with Light	
	Source)	
	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	A rigid laryngoscope is used for	
	direct laryngoscopy	
	(visualizing the larynx). A	
	magnifying glass with light	
	source is used with this to	
	magnify and illuminate the	
	anatomical structure under	
	observation.	
2	Operational Requirements	
2.1	Small and lightweight system is	
	required.	
3	System Configuration	
3.1	Magnifying rigid Laryngoscope with	
	light source, complete unit.	
4	Technical Specifications	
4.1	Rigid, metallic fibre-optic light	
	system with inbuilt magnifying	
	system.	
4.2	Viewing angle 90 degrees.	
4.3	Light source: Halogen bulb	
4.4	Autoclaveable.	
4.5	Shall work on inbuilt rechargeable	
	battery.	
5	Accessories, spares and	
	consumables	
5.1	Accessories:	
	Spare halogen bulb	
5.2	All standard accessories,	
	consumables and parts required	
	to operate the equipment,	
	including all standard tools and	
	cleaning and lubrication	
	materials, to be included in the	
	offer. Bidders must specify the	



S.N.	Purchaser's Specifications	Bidder's Remarks
51111	quantity of every item included	Didder 5 remains
	in their offer (including items	
	not specified above).	
6	Operating Environment	
6.1	The system offered shall be designed	
0.1	to operate normally under the	
	conditions of the purchaser's	
	country. The conditions include	
	Power Supply, Climate,	
	Temperature, Humidity, etc.	
6.2	Rechargeable battery operated	
0.2	system. Charger to be provided	
	if integrated charger is not	
	there.	
7	Standards and Safety	
′	Requirements	
7.1	Must submit ISO13485:2003/ AC:	
	2007 for Medical Devices AND	
7.2	CE (93/42 EEC Directives) or	
	USFDA approved product	
	certificate.	
8	User Training	
8.1	Must provide user training (including	
	how to use and maintain the	
	equipment).	
	equipilient).	
9	Warranty	
9 9.1	1 1	
9.1	Warranty Comprehensive warranty for 1 year from acceptance.	
	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During	
9.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period	
9.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are	
9.1 10 10.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable.	
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9.1 10 10.1 11 11.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Supplier must accomplish proper commissioning of the equipment on site.	
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9.1 10 10.1 11 11.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Supplier must accomplish proper commissioning of the equipment on site. Documentation User (Operating) manual in English Service (Technical / Maintenance)	
9.1 10.1 11.1 12.1 12.2	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Supplier must accomplish proper commissioning of the equipment on site. Documentation User (Operating) manual in English Service (Technical / Maintenance) manual in English	
9.1 10 10.1 11 11.1 12 12.1	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Supplier must accomplish proper commissioning of the equipment on site. Documentation User (Operating) manual in English Service (Technical / Maintenance) manual in English List of important spare parts and	
9.1 10.1 11.1 12.1 12.2	Warranty Comprehensive warranty for 1 year from acceptance. Maintenance Service During Warranty Period Standard warranty conditions are applicable. Installation and Commissioning Supplier must accomplish proper commissioning of the equipment on site. Documentation User (Operating) manual in English Service (Technical / Maintenance) manual in English	

21. Lower (uterine) segment Caesarean section (LSCS) set



S.N.	Purchaser's Specifications	Bidder's Remarks
D.11.	Lower (uterine) segment	Diduct 5 Remarks
	Caesarean section (LSCS)	
	set	
	Manufacturer	
	Brand	
	Type / Model	
1	Country of Origin	
1.1	Description of Function	
1.1	LSCS set for a transverse cut just	
	above the edge of the bladder	
	for resulting in less blood loss	
	and should be easier to repair	
	than other types of Caesarean	
	sections.	
2	Operational Requirements	
2.1	The Instruments must be made of	
	highest quality materials e.g.	
	stainless steel (S/S) for metal	
	devices. The surgical	
	instruments must be CE	
	marked / USFDA approved.	
	The bidder must submit	
	quality assurance (QA)	
	certificates.	
2.2	Instruments must be made from	
	surgical quality, preferably	
	non-magnetic stainless steel	
	and must be matt surface	
	finish. Quality must comply	
	with EN 46002 and ISO 9002	
	and / or their latest	
	amendments.	
2.3	Each pack must be packaged in a	
	hospital grade cotton wrapper	
	(autoclave-able) as a complete	
	pack. Bulk loose instrument	
	supply is NOT acceptable.	
	Each of the individual	
	instruments of a set must be	
	packed in a labelled clear	
	plastic wrapper for easy	
	identification. All	
	individually packed	
	instruments of a set shall then	
	be packed together in a larger	
	clear plastic wrapper labelled	



S.N.	Purchaser's Specifications	Bidder's Remarks
	with the name of the set for	
	easy identification.	
2.4	Instrument surfaces must NOT be	
	stamped, indented or	
	scratched. It is preferred if the	
	suppliers labelled GoN name	
	in anodised form of labelling.	
	It shall carry clear anodised	
	labelling/ marking of	
	manufacturer's name/ brand	
	and the part number/ model	
	number of instruments on the	
	surface of each piece of	
	instruments.	
2.5	Particular attention must be paid to	
	the quality of box joints to	
	ensure that they are smooth	
	and interlock well, and to	
	teeth and grips to ensure that	
	they meet and interlock well.	
	Finger rings must be of	
	proper size and shape for	
	maximum utility and comfort. The inside of finger rings	
	must be well rounded and free	
	of sharp edges, rough areas	
	and grinding marks, cracks,	
	overlaps, burrs.	
2.6	Jaw serration must be well cut and	
2.0	defined and must mesh	
	properly when the jaws are	
	fully closed. The edges of the	
	serration must be well	
	chamfered and must not	
	contain burrs and sharp edges.	
	Teeth must be sharp (unless	
	otherwise specified), of	
	proper size and shape, free of	
	rough edges or burrs, and	
	must mesh with sufficient	
	accuracy to ensure proper	
	performance for the use	
	intended.	
2.7	Ratchet and ratchet catches must be	
	properly aligned and undercut	
	for safe locking. Ratchets	



C N	D 1 C 'C' 1'	P'II. I. D I.
S.N.	Purchaser's Specifications	Bidder's Remarks
	must be of such design as to	
	ensure easy and positive	
	engagement and proper	
	disengagement. Ratchets and	
	ratchet catches must be free of	
	burrs and sharp edges.	
2.8	Locks, forceps and similar	
	instruments must be of the	
	box lock type or lap joint	
	type. All type of locks must	
	7.2	
	be accurately fitted, without	
	stiffness and without crevices,	
	burrs or sharp edges	
	anywhere in the construction.	
2.9	Screws of screw lock scissors and	
	other instruments must be the	
	concentrically mustered type,	
	countersunk, flush with, or	
	slightly below the surface or	
ı	rounded, smooth and flush at	
l	the periphery, but not riveted.	
	The screw must retain their	
	position after setting without	
	binding or loosening during	
	use.	
2.10	Scissors Company of the Process of t	
2.10	The ROCKWELL hardness of the	
	finished instruments must be	
	within the range from 50	
	HRC to 58 HRC. Opposite	
	blades must not vary in	
	hardness by more than 4 units	
	on the ROCKWELL C	
	harness scale.	
2.11	Scissors must have joints, which	
	move smoothly and must be	
	neither too loose nor too tight:	
	it must be possible to close	
	and reopen the instrument	
2.12	easily with two fingers.	
2.12	The cutting ability of the instrument	
	must be tested. The	
	instrument must cut clearly	
	without tearing.	
2.13	The finish and all edges and	
	surfaces must be uniform and	



C NI	Dunahagan's Charifications	Diddon's Domonks
S.N.	Purchaser's Specifications	Bidder's Remarks
	free of burrs, sharp edges	
	(except where required),	
	pores, crevices, gins marks,	
	rough areas, cracks and	
	overlaps.	
3	System Configuration	
3.1	Delivery Set	
4	Technical Specifications	
4.1	The instruments required are listed below, bidder MUST provide full description of all instruments (description includes: full name, type, shape, design, full length, volume and etc.) required below for the evaluation.	
4.2	Instruments Required	
	1 Sponge Holding Forceps 25cm	
	4 Green Armytage Forceps	
	6 Artery Forceps cvd 15cm	
	6 Artery Forceps St 15cm	
	4 Allice Tissue Forceps Medium	
	15cm	
	4 Allice Tissue Forceps Large 15cm	
	2 Bebcock Tissue Forceps 15cm	
	2 Toothed Forceps 15 cm	
	2 Dissecting Forceps Toothed 15cm	
	2 Dissecting Forceps Non Toothed	
	15cm	
	1 Needle Holder Mayo Heger 15	
	cm	
	4 Kell's Clamp 15cm	
	1 Suction Tip	
	1 Tissue Cutting Scissors 17.5cm	
	2 B.P.Handle no4	
	4 Towel Clip Cross Action	
	1 Bowl 20"	
	1 Doyen Retractor 5 cm	
	1 Morr's Retractor 5cm	
	1 Self Retaining Retractor	
	1 Dever's Retractor 5 cm	
	1 Instruments Bag	
4.3	All instruments must be supplied	
	free of residual scale, acid,	



S.N.	Purchaser's Specifications	Bidder's Remarks
D-1 1-	grease and grinding and	Didder 5 Remarks
	polishing materials and	
	workmanship must be first	
	class throughout. Instruments	
	must be free of defects, which	
	detract from their appearance	
	or impair serviceability,	
	proper functioning and	
	intended use.	
4.4	Bidder MUST attach product	
	catalogues with photos for all	
	instruments as mentioned.	
	These catalogues/photos	
	MUST clearly and correctly	
	mark with non-erasable	
	making pen their respective	
	parameter line number	
	(shown on the left column)	
	and instrument name.	
5	Accessories, spares and	
3	consumables	
5.1	All standard accessories,	
3.1	consumables and parts	
	required to operate the	
	equipment, including all	
	standard tools and cleaning	
	and lubrication materials, to	
	be included in the offer.	
	Bidders must specify the	
	quantity of every item	
	included in their offer	
	(including items not specified	
	above).	
6	Operating Environment	
6.1	The instrument offered shall be	
0.1	designed to store and to	
	operate normally under the	
	conditions of the purchaser's	
	country. The conditions	
	include Climate,	
	Temperature, Humidity, etc.	
7	Standards and Safety	
,	Requirements Salety	
7.1	Must submit	
,	ISO13485:2003/AC:2007 for	
	Medical Devices AND	
	Medical Devices AND	



S.N.	Purchaser's Specifications	Bidder's Remarks
7.2	CE (93/42 EEC Directives) or	
	USFDA approved product	
	certificate.	
8	User Training	
8.1	Not applicable.	
9	Warranty	
9.1	Warranty for 1 year.	
10	Maintenance Service During	
	Warranty Period	
10.1	Standard warranty conditions are	
	applicable.	
11	Installation and Commissioning	
11.1	Must supply complete unit, ready to	
	use.	
12	Documentation	
12.1	s/Instructions manual shall be	
	provided in English.	



Section VI. General Conditions of Contract

Table of Clauses

1.	Definitions	112
2.	Contract Documents	112
3.	Fraud and Corruption	113
4.	Interpretation	
5.	Language	
6.	Joint Venture, Consortium or Association	115
7.	Notices	
8.	Governing Law	
9.	Settlement of Disputes	
10.	Scope of Supply	
11.	Delivery	
12.	Supplier's Responsibilities	
13.	Purchaser's Responsibilities	
14.	Contract Price	
15.	Terms of Payment	
16.	Taxes and Duties	116
17.	Performance Security	116
18.	Copyright	116
19.	Copyright	116
20.	Subcontracting	117
21.	Specifications and Standards	117
22.	Packing and Documents	118
23.	Insurance	118
24.	Transportation	118
25.	Inspections and Tests	118
26.	Liquidated Damages	
27.	Warranty	
28.	Patent Indemnity	
29.	Limitation of Liability	120
30.	Change in Laws and Řegulations	121
31.	Force Majeure	121
32.	Change Orders and Contract Amendments	121
33.	Extensions of Time	122
34.	Termination	122
35.	Assignment	



Section VI. General Conditions of Contract

1. Definitions

- 1.1 The following words and expressions shall have the meanings hereby assigned
 - (a) "Contract" means the Agreement entered into between the Purchaser and the Supplier, together with the Contract Documents referred to therein, including all attachments, appendices, and all documents incorporated by reference therein.
 - (b) "Contract Documents" means the documents listed in the Agreement, including any amendments thereto.
 - (c) "Contract Price" means the price payable to the Supplier as specified in the Agreement, subject to such additions and adjustments thereto or deductions there from, as may be made pursuant to the Contract.
 - (d) "Day" means calendar day.
 - (e) "Delivery" means the transfer of the Goods from the Supplier to the Purchaser in accordance with the terms and conditions set forth in the Contract.
 - (f) "Completion" means the fulfillment of the Related Services by the Supplier in accordance with the terms and conditions set forth in the Contract.
 - (g) "GCC" means the General Conditions of Contract.
 - (h) "Goods" means all of the commodities, raw material, machinery and equipment, and/or other materials that the Supplier is required to supply to the Purchaser under the Contract.
 - (i) "Purchaser's Country" is the country specified in the Special Conditions of Contract (SCC).
 - (j) "Purchaser" means the entity purchasing the Goods and Related Services, as specified in the SCC.
 - (k) "Related Services" means the services incidental to the supply of the goods, such as insurance, installation, training and initial maintenance and other similar obligations of the Supplier under the Contract.
 - (l) "SCC" means the Special Conditions of Contract.
 - (m) "Subcontractor" means any natural person, private or government entity, or a combination of the above, including its legal successors or permitted assigns, to whom any part of the Goods to be supplied or execution of any part of the Related Services is subcontracted by the Supplier.
 - (n) "Supplier" means the natural person, private or government entity, or a combination of the above, whose bid to perform the Contract has been accepted by the Purchaser and is named as such in the Agreement, and includes the legal successors or permitted assigns of the Supplier.
 - (o) "GoN" means the Government of Nepal.
 - (p) "The Site," where applicable, means the place named in the SCC.

2. Contract Documents

2.1 Subject to the order of precedence set forth in the Agreement, all documents forming the Contract (and all parts thereof) are intended to be correlative,



complementary, and mutually explanatory.

3 Fraud and Corruption

- 3.1If the Purchaser determines that the Supplier has engaged in corrupt, fraudulent, collusive, coercive or obstructive practices, in competing for or in executing the Contract, then the Purchaser may, after giving 14 days notice to the Supplier, terminate the Supplier's employment under the Contract and the provisions of Clause 34.1 shall apply.
- 3.2 Without prejudice to any other rights of the Purchaser under this Contract, GoN may blacklist the Bidder/Supplier for its conduct up to three (3) years on the following grounds and seriousness of the act committed by the Bidder/Supplier:
 - (a) if it is established that the Supplier has committed substantial defect in implementation of the Contract or has or has not substantially fulfilled its obligations under the Contract

For the purposes of this Sub-Clause:

- (i) "corrupt practice" is the offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party;
- (ii) "fraudulent practice" l is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;
- (iii) "collusive practice" 2 is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party;
- (iv) "coercive practice" is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party;
- (v) "obstructive practice" is
 - (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a GoN/DP investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or
 - (bb) acts intended to materially impede the exercise of the GoN/DP's inspection and audit rights provided for under ITB Clause 3.5 and GCC Clause 25.
- 3.3 Without prejudice to any other rights of the Purchaser under this Contract, GoN may **blacklist** a Bidder/Supplier for its conduct for a period of one (1) to three (3) years on the following grounds and seriousness of the act committed by the bidder:

³a "party" refers to a participant in the procurement process or contract execution.



¹a "party" refers to a public official; the terms "benefit" and "obligation" relate to the procurement process or contract execution; and the "act or omission" is intended to influence the procurement process or contract execution.

² "parties" refers to participants in the procurement process (including public officials) attempting to establish bid prices at artificial, non competitive levels.

if it is established that the Supplier committed acts specified in ITB 3.2,

if it is established later that the Bidder has committed substantial defect in implementation of the contract or has not substantially fulfilled its obligations under the contract or the completed work is not of the specified quality as per the contract.

4. Interpretation

4.1If the context so requires it, singular means plural and vice versa.

4.2 Incoterms

- (a) The meaning of any trade term and the rights and obligations of parties there under shall be as prescribed by Incoterms.
- (b) EXW shall be governed by the rules prescribed in the current edition of Incoterms, published by the International Chamber of Commerce at the date of the Invitation for Bids or as specified in the SCC.

4.3 Entire Agreement

The Contract constitutes the entire agreement between the Purchaser and the Supplier and supersedes all communications, negotiations and agreements (whether written or oral) of parties with respect thereto made prior to the date of Contract.

4.4 Amendment

No amendment or other variation of the Contract shall be valid unless it is in writing, is dated, expressly refers to the Contract, and is signed by a duly authorized representative of each party thereto.

4.5 Nonwaiver

- (a)Subject to GCC Sub-Clause 4.5(b) below, no relaxation, forbearance, delay, or indulgence by either party in enforcing any of the terms and conditions of the Contract or the granting of time by either party to the other shall prejudice, affect, or restrict the rights of that party under the Contract, neither shall any waiver by either party of any breach of Contract operate as waiver of any subsequent or continuing breach of Contract.
- (b) Any waiver of a party's rights, powers, or remedies under the Contract must be in writing, dated, and signed by an authorized representative of the party granting such waiver, and must specify the right and the extent to which it is being waived.

4.6 Severability

If any provision or condition of the Contract is prohibited or rendered invalid or unenforceable, such prohibition, invalidity or unenforceability shall not affect the validity or enforceability of any other provisions and conditions of the Contract.

5. Language

- 5.1 The Contract as well as all correspondence and documents relating to the Contract exchanged by the Supplier and the Purchaser, shall be written in the language specified in the SCC. Supporting documents and printed literature that are part of the Contract may be in another language provided they are accompanied by an accurate translation of the relevant passages in the language specified in the SCC, in which case, for purposes of interpretation of the Contract, this translation shall govern.
- 5.2 The Supplier shall bear all costs of translation to the governing language and



all risks of the accuracy of such translation.

6. Joint Venture, Consortium or Association

6.1 If the Supplier is a joint venture, consortium, or association, all of the parties shall be jointly and severally liable to the Purchaser for the fulfillment of the provisions of the Contract and shall designate one party to act as a leader with authority to bind the joint venture, consortium, or association. A bidder can submit only one bid either as a partner of the joint venture or individually. The composition or the constitution of the joint venture, consortium, or association shall not be altered without the prior consent of the Purchaser.

7. Notices

- 7.1 Any Notice given by one party to the other pursuant to the Contract shall be in writing to the address specified in the SCC. The term "in writing" means communicated in written form with proof of receipt.
- 7.2 A Notice shall be effective when delivered or on the Notice's effective date, whichever is later.

8. Governing Law

8.1 The Contract shall be governed by and interpreted in accordance with the laws of Nepal.

9. Settlement of Disputes

- 9.1 The Purchaser and the Supplier shall make every effort to resolve amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the Contract.
- 9.2 Any dispute between the parties as to matters arising pursuant to this contract which cannot be settled amicably within thirty (30) days after receipt by one party of the other party's request for such amicable settlement may be referred to Arbitration within 30 days after the expiration of amicable settlement period.

10. Scope of Supply

- 10.1 Subject to the SCC, the Goods and Related Services to be supplied shall be as specified in Section V, Schedule of Requirements.
- 10.2 Unless otherwise stipulated in the Contract, the Scope of Supply shall include all such items not specifically mentioned in the Contract but that can be reasonably inferred from the Contract as being required for attaining Delivery and Completion of the Goods and Related Services as if such items were expressly mentioned in the Contract.

11. Delivery

11.1 Subject to GCC Sub-Clause 31.1, the Delivery of the Goods and Completion of the Related Services shall be in accordance with the Delivery and Completion Schedule specified in the Section V, Schedule of Requirements. The details of documents to be furnished by the Supplier are specified in the SCC.

12. Supplier's Responsibilitie

12.1 The Supplier shall supply all the Goods and Related Services included in the Scope of Supply in accordance with GCC Clause 10, and the Delivery and Completion Schedule, as per GCC Clause 11.

13. Purchaser's Responsibilitie

- 13.1 Whenever the supply of Goods and Related Services requires that the Supplier obtain permits, approvals, and import and other licenses from public authorities in Nepal, the Purchaser shall, if so required by the Supplier, make its best effort to assist the Supplier in complying with such requirements in a timely and expeditious manner.
- 13.2 The Purchaser shall pay all costs involved in the performance of its responsibilities, in accordance with GCC Sub-Clause 13.1.



14. Contract Price

- 14.1 The Contract Price shall be as specified in the Agreement subject to any additions and adjustments thereto, or deductions there from, as may be made pursuant to the Contract.
- 14.2 Prices charged by the Supplier for the Goods delivered and the Related Services performed under the Contract shall not vary from the prices quoted by the Supplier in its bid, with the exception of any price adjustments authorized in the SCC.

15. Terms of Payment

- 15.1 The Contract Price shall be paid in Nepalese Currency.
- 15.2 The Supplier's request for payment shall be made to the Purchaser in writing, accompanied by invoices describing, as appropriate, the Goods delivered and Related Services performed, and by the documents submitted pursuant to GCC Clause 11 and upon fulfillment of all the obligations stipulated in the Contract.
- 15.3 Payments shall be made promptly by the Purchaser, no later than thirty (30) days after submission of an invoice or request for payment by the Supplier, and the Purchaser has accepted it.

16. Taxes and Duties

16.1 For goods supplied, the Supplier shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted Goods to the Purchaser

17. Performance Security

- 17.1 The Supplier shall, within fifteen (15) days of the receipt of notification of Contract award, provide a Performance Security for the due performance of the Contract in the amounts and currencies specified in the SCC.
- 17.2 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 its obligations under the Contract.
- 17.3 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 forms stipulated by the Purchaser in the SCC, or in another form acceptable to the Purchaser.
- 17.4 The Performance Security shall 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 specified otherwise in the SCC.

18. Copyright

18.1 The copyright in all drawings, documents, and other materials containing data and information furnished to the Purchaser by the Supplier herein shall remain vested in the Supplier, or, if they are furnished to the Purchaser directly or through the Supplier by any third party, including suppliers of materials, the copyright in such materials shall remain vested in such third party.

19. Confidential Information

19.1 The Purchaser and the Supplier shall keep confidential and shall not, without the written consent of the other party hereto, divulge to any third party any documents, data, or other information furnished directly or indirectly by the other party hereto in connection with the Contract, whether such information has been furnished prior to, during or following completion or termination of the Contract. Notwithstanding the above, the Supplier may furnish to its Subcontractor such documents, data, and other information it receives from the Purchaser to the extent required for the Subcontractor to perform its work



- under the Contract, in which event the Supplier shall obtain from such Subcontractor an undertaking of confidentiality similar to that imposed on the Supplier under GCC Clause 19.
- 19.2 The Purchaser shall not use such documents, data, and other information received from the Supplier for any purposes unrelated to the Contract. Similarly, the Supplier shall not use such documents, data, and other information received from the Purchaser for any purpose other than the design, procurement, or other work and services required for the performance of the Contract.
- 19.3 The obligation of a party under GCC Sub-Clauses 19.1 and 19.2 above, however, shall not apply to information that:
 - (a) the Purchaser or Supplier need to share with the Donor for Donor funded project or other institutions participating in the financing of the Contract;
 - (b) now or hereafter enters the public domain through no fault of that party;
 - (c) can be proven to have been possessed by that party at the time of disclosure and which was not previously obtained, directly or indirectly, from the other party; or
 - (d) otherwise lawfully becomes available to that party from a third party that has no obligation of confidentiality.
- 19.4 The above provisions of GCC Clause 19 shall not in any way modify any undertaking of confidentiality given by either of the parties hereto prior to the date of the Contract in respect of the Supply or any part thereof.
- 19.5 The provisions of GCC Clause 19 shall survive completion or termination, for whatever reason, of the Contract.

20. Subcontractin

- 20.1 The Supplier shall notify the Purchaser in writing of all subcontracts awarded under the Contract if not already specified in the Bid. Subcontracting shall in no event relieve the Supplier from any of its obligations, duties, responsibilities, or liability under the Contract.
- 20.2 Subcontracts shall comply with the provisions of GCC Clauses 3.

21. Specifications and Standards

- 21.1 Technical Specifications and Drawings
 - (a) The Supplier shall ensure that the Goods and Related Services comply with the technical specifications and other provisions of the Contract.
 - (b) The Supplier shall be entitled to disclaim responsibility for any design, data, drawing, specification or other document, or any modification thereof provided or designed by or on behalf of the Purchaser, by giving a notice of such disclaimer to the Purchaser.
 - (c) The Goods and Related Services supplied under this Contract shall conform to the standards mentioned in Section V, Schedule of Requirements and, when no applicable standard is mentioned, the standard shall be equivalent or superior to the official standards whose application is appropriate to the country of origin of the Goods.
- 21.2 Wherever references are made in the Contract to codes and standards in accordance with which it shall be executed, the edition or the revised



version of such codes and standards shall be those specified in the Section V, Schedule of Requirements Requirements. During Contract execution, any changes in any such codes and standards shall be applied only after approval by the Purchaser and shall be treated in accordance with GCC Clause 32.

22. Packing and Documents

- 22.1 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. During transit, the packing shall be sufficient to withstand, without limitation, rough handling and exposure to extreme temperatures, salt and precipitation, and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the final destination of the Goods and the absence of heavy handling facilities at all points in transit.
- 22.2 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 the SCC, and in any other instructions ordered by the Purchaser.

23. Insurance

- 23.1 Unless otherwise specified in the SCC, 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.
- 24. Transportatio
- 24.1 Unless otherwise specified in the SCC, obligations for transportation of the Goods shall be in accordance with the Incoterms specified in Sections V, Schedule of Requirements.

25. Inspections and Tests

- 25.1 The Supplier shall at its own expense and at no cost to the Purchaser carry out all such tests and/or inspections of the Goods and Related Services as are specified in Sections V, Schedule of Requirements.
- 25.2 The inspections and tests may be conducted on the premises of the Supplier or its Subcontractor, at point of delivery, and/or at the final destination of the Goods, or in another place in Nepal as specified in the SCC. Subject to GCC Sub-Clause 25.3, if conducted on the premises of the Supplier or its Subcontractor, all reasonable facilities and assistance, including access to drawings and production data, shall be furnished to the inspectors at no charge to the Purchaser.
- 25.3 The Purchaser or its designated representative shall be entitled to attend the tests and/or inspections referred to in GCC Sub-Clause 25.2, provided that the Purchaser bear all of its own costs and expenses incurred in connection with such attendance including, but not limited to, all traveling and board and lodging expenses.
- 25.4 Whenever the Supplier is ready to carry out any such test and inspection, it shall give a reasonable advance notice, including the place and time, to the Purchaser. The Supplier shall obtain from any relevant third party or manufacturer any necessary permission or consent to enable the Purchaser or its designated representative to attend the test and/or inspection.
- 25.5 The Purchaser may require the Supplier to carry out any test and/or inspection not required by the Contract but deemed necessary to verify that the characteristics and performance of the Goods comply with the technical specifications, codes and standards under the Contract, provided that the



Supplier's reasonable costs and expenses incurred in the carrying out of such test and/or inspection shall be added to the Contract Price. Further, if such test and/or inspection impede the progress of manufacturing and/or the Supplier's performance of its other obligations under the Contract, due allowance will be made in respect of the Delivery Dates and Completion Dates and the other obligations so affected.

- 25.6 The Supplier shall provide the Purchaser with a report of the results of any such test and/or inspection.
- 25.7 The Purchaser may reject any Goods or any part thereof that fail to pass any test and/or inspection or do not conform to the specifications. The Supplier shall either rectify or replace such rejected Goods or parts thereof or make alterations necessary to meet the specifications at no cost to the Purchaser, and shall repeat the test and/or inspection, at no cost to the Purchaser, upon giving a notice pursuant to GCC Sub-Clause 25.4.
- 25.8 The Supplier agrees that neither the execution of a test and/or inspection of the Goods or any part thereof, nor the attendance by the Purchaser or its representative, nor the issue of any report pursuant to GCC Sub-Clause 25.6, shall release the Supplier from any warranties or other obligations under the Contract.

26. Liquidated Damages

26.1 Except as provided under GCC Clause 31, if the Supplier fails to deliver any or all of the Goods or perform the Related Services within the period specified in the Contract, the Purchaser may without prejudice to all its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to the percentage specified in the SCC of the Contract Price for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the percentage specified in the SCC. Once the maximum is reached, the Purchaser may terminate the Contract pursuant to GCC Clause 34.

27. Warranty

- 27.1 The Supplier warrants that all the Goods are new, unused, and of the most recent or current models, and that they incorporate all recent improvements in design and materials, unless provided otherwise in the Contract.
- 27.2 Subject to GCC Sub-Clause 21.1, the Supplier further warrants that the Goods shall be free from defects arising from any act or omission of the Supplier or arising from design, materials, and workmanship, under normal use in the conditions prevailing in Nepal.
- 27.3 Unless otherwise specified in the SCC, the 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 SCC.
- 27.4 The Purchaser shall give Notice to the Supplier stating the nature of any such defects together with all available evidence thereof, promptly following the discovery thereof. The Purchaser shall afford all reasonable opportunity for the Supplier to inspect such defects.
- 27.5 Upon receipt of such Notice, the Supplier shall, within the period specified in the SCC, expeditiously repair or replace the defective Goods or parts thereof, at no cost to the Purchaser.
- 27.6 If having been notified, the Supplier fails to remedy the defect within the period specified in the SCC, the Purchaser may proceed to take within a reasonable period 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.

28. Patent Indemnity

- 28.1 The Supplier shall, subject to the Purchaser's compliance with GCC Sub-Clause 28.2, indemnify and hold harmless the Purchaser and its employees and officers from and against any and all suits, actions or administrative proceedings, claims, demands, losses, damages, costs, and expenses of any nature, including attorney's fees and expenses, which the Purchaser may suffer as a result of any infringement or alleged infringement of any patent, utility model, registered design, trademark, copyright, or other intellectual property right registered or otherwise existing at the date of the Contract by reason of:
 - (a) the installation of the Goods by the Supplier or the use of the Goods in the country where the Site is located; and
 - (b) the sale in any country of the products produced by the Goods.

Such indemnity shall not cover any use of the Goods or any part thereof other than for the purpose indicated by or to be reasonably inferred from the Contract, neither any infringement resulting from the use of the Goods or any part thereof, or any products produced thereby in association or combination with any other equipment, plant, or materials not supplied by the Supplier, pursuant to the Contract.

- 28.2 If any proceedings are brought or any claim is made against the Purchaser arising out of the matters referred to in GCC Sub-Clause 28.1, the Purchaser shall promptly give the Supplier a notice thereof, and the Supplier may at its own expense and in the Purchaser's name conduct such proceedings or claim and any negotiations for the settlement of any such proceedings or claim.
- 28.3 If the Supplier fails to notify the Purchaser within thirty (30) days after receipt of such notice that it intends to conduct any such proceedings or claim, then the Purchaser shall be free to conduct the same on its own behalf.
- 28.4 The Purchaser shall, at the Supplier's request, afford all available assistance to the Supplier in conducting such proceedings or claim, and shall be reimbursed by the Supplier for all reasonable expenses incurred in so doing.
- 28.5 The Purchaser shall indemnify and hold harmless the Supplier and its employees, officers, and Subcontractors from and against any and all suits, actions or administrative proceedings, claims, demands, losses, damages, costs, and expenses of any nature, including attorney's fees and expenses, which the Supplier may suffer as a result of any infringement or alleged infringement of any patent, utility model, registered design, trademark, copyright, or other intellectual property right registered or otherwise existing at the date of the Contract arising out of or in connection with any design, data, drawing, specification, or other documents or materials provided or designed by or on behalf of the Purchaser.

29. Limitation of Liability

29.1 Except in cases of gross negligence or willful misconduct :

(a)neither party shall be liable to the other party 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 liquidated 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, or to any obligation of the Supplier to indemnify the Purchaser with respect to patent infringement.
- 30. Change in Laws and Regulations
- 30.1 Unless otherwise specified in the Contract, if after the date of the Invitation for Bids, any law, regulation, ordinance, order or bylaw having the force of law is enacted, promulgated, abrogated, or changed in Nepal where the Site is located (which shall be deemed to include any change in interpretation or application by the competent authorities) that subsequently affects the Delivery Date and/or the Contract Price, then such Delivery Date and/or Contract Price shall be correspondingly increased or decreased, to the extent that the Supplier has thereby been affected in the performance of any of its obligations under the Contract. Notwithstanding the foregoing, such additional or reduced cost shall not be separately paid or credited if the same has already been accounted for in the price adjustment provisions where applicable, in accordance with GCC Clause 14.

31. Force Majeure

- 31.1 The Supplier shall not be liable for forfeiture of its Performance Security, liquidated damages, or termination for default if and to the extent that its delay in performance or other failure to perform its obligations under the Contract is the result of an event of Force Majeure.
- 31.2 For purposes of this Clause, "Force Majeure" means an event or situation beyond the control of the Supplier that is not foreseeable, is unavoidable, and its origin is not due to negligence or lack of care on the part of the Supplier. Such events may include, but not be limited to, acts of the Purchaser in its sovereign capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions, and freight embargoes.
- 31.3 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.

32. Change Orders and Contract Amendments

- 32.1 The Purchaser may at any time order the Supplier through Notice in accordance GCC Clause 7, to make changes within the general scope of the Contract in any one or more of the following:
 - (a) drawings, designs, or specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the Purchaser;
 - (b) the method of shipment or packing;
 - (c) the place of delivery; and
 - (d) the Related Services to be provided by the Supplier.
- 32.2 If any such change causes an increase or decrease in the cost of, or the time required for, the Supplier's performance of any provisions under the Contract, an equitable adjustment shall be made in the Contract Price or in



the Delivery and Completion Schedule, or both, and the Contract shall accordingly be amended. Any claims by the Supplier for adjustment under this Clause must be asserted within thirty (30) days from the date of the Supplier's receipt of the Purchaser's change order.

32.3 Prices to be charged by the Supplier for any Related Services that might be needed but which were not included in the Contract 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.

33. Extensions of Time

- 33.1 If at any time during performance of the Contract, the Supplier or its Subcontractors should encounter conditions impeding timely delivery of the Goods or completion of Related Services pursuant to GCC Clause 11, the Supplier shall promptly, and at least seven (7) days before the expiry of procurement contract, notify the Purchaser in writing of the delay, its likely duration, and its cause. As soon as practicable after receipt of the Supplier's notice, the Purchaser shall evaluate the situation and may at its discretion extend the Supplier's time for performance, in which case the extension shall be ratified by the parties by amendment of the Contract.
- 33.2 Except in case of Force Majeure, as provided under GCC Clause 31, a delay by the Supplier in the performance of its Delivery and Completion obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC Clause 26, unless an extension of time is agreed upon, pursuant to GCC Sub-Clause 34.1.

34. Termination

34.1 Termination for Default

- (a) The Purchaser, without prejudice to any other remedy for breach of Contract, by Notice of default sent to the Supplier, may terminate the Contract in whole or in part:
 - (i) if the Supplier fails to deliver any or all of the Goods within the period specified in the Contract, or within any extension thereof granted by the Purchaser pursuant to GCC Clause 33; or
 - (ii) if the Supplier fails to perform any other obligation under the Contract.
- (b) In the event the Purchaser terminates the Contract in whole or in part, pursuant to GCC Clause 34.1(a), the Purchaser may procure, upon such terms and in such manner as it deems appropriate, Goods or Related Services similar to those undelivered or not performed, and the Supplier shall be liable to the Purchaser for any additional costs for such similar Goods or Related Services. However, the Supplier shall continue performance of the Contract to the extent not terminated.
- (c) if the Supplier, in the judgment of the Purchaser has engaged in corrupt, fraudulent, collusive, coercive or obstructive practices, as defined in GCC Clause 3, in competing for or in executing the Contract.

34.2 Termination for Insolvency

The Purchaser may at any time terminate the Contract by giving Notice to the Supplier if the Supplier becomes bankrupt or otherwise insolvent. In such event, termination will be without compensation to the Supplier,



provided that such termination will not prejudice or affect any right of action or remedy that has accrued or will accrue thereafter to the Purchaser.

34.3 Termination for Convenience

- (a) The Purchaser, by written Notice sent to the Supplier, may terminate the Contract, in whole or in part, at any time for its convenience. The Notice of termination shall specify that termination is for the Purchaser's convenience, the extent to which performance of the Supplier under the Contract is terminated, and the date upon which such termination becomes effective.
- (b) The Goods that are complete and ready for shipment within seven (7) days after the Supplier's receipt of the Notice of termination shall be accepted by the Purchaser at the Contract terms and prices. For the remaining Goods, the Purchaser may elect:
 - (i) To have any portion completed and delivered at the Contract terms and prices; and/or
 - (ii) To cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Related Services and for materials and parts previously procured by the Supplier.

35. Assignment

35.1 Neither the Purchaser nor the Supplier shall assign, in whole or in part, their obligations under this Contract, except with prior written consent of the other party.



Section VII. Special Conditions of Contract

The following Special Conditions of Contract (SCC) shall supplement the General Conditions of Contract (GCC). Whenever there is a conflict, the provisions herein shall prevail over those in the GCC.

GCC 1.1(i)	The Purchaser's country is: Nepal
GCC 1.1(j)	The Purchaser is:
	Ministry of Health Department of Health Services
	Logistics Management Division
	Teku, Kathmandu
GCC 1.1 (p)	The Site is:
	Logistics Management Division
	Central Store, Teku, Kathmandu
GCC 4.2 (b)	The version of Incoterms shall be: NA
GCC 5.1	The language shall be: English
GCC 7.1	For <u>notices</u> , the Purchaser's address shall be:
	Name and Address of the Purchaser:
	Ministry of Health
	Department of Health Services
	Logistics Management Division
	Teku, Kathmandu
	Telephone number: 014261768
	Facsimile number: 014261413
	e-mail Address: drtinkari@gmail.com
	For <u>notices</u> , the Suppliers's address shall be:
	Name and Address of the Supplier:
	Telephone number:
	Facsimile number:
	e-mail Address:
GCC 10.1	The Scope of Supply shall be defined in: "Section V, Schedule of Requirements"
GCC 11.1	Upon delivery of the Goods to the transporter, the Supplier shall notify the Purchaser and send the following documents to the Purchaser:
	a) Copies of the Supplier's invoice showing the description of the Goods, quantity, unit price, and total amount;
	b) Copy of packing list indentifying the contents of each package;



	125
	c) Delivery note, railway receipt, or truck receipt;(as appropriate)
	d) Manufacturer's or Supplier's warranty certificate;
	e) Other documents (if any).
	The Purchaser shall receive the above documents before the arrival of the Goods and, if not received, the Supplier will be responsible for any consequent expenses.
GCC 14.2	Price adjustment shall not be applicable.
GCC 15.1	The terms of payment to be made to the Supplier under the contract shall be as follows:
	1. The payment shall be made:
	(a) through Finance Section of the Purchaser
GCC 15.1	a) Payments shall be made in Nepalese Rupees in the following manner:
	b) Advance Payment: Not Applicable .
	c) On Delivery and acceptance: Hundred (100) percent. of the Contract Price of the Goods delivered shall be paid within thirty (30) days of receipt of the Goods and upon submission of a claim supported by the documents specified in GCC 11.1
	d) 1.5% shall be deducted from the total billed amount against advance income tax.
GCC 17.1	The Supplier shall provide a Performance Security as per ITB 41. The amount of the Performance Security shall be in Nepalese Rupees,
	The Supplier shall provide a Performance Security as follows:
	i) If the contract price of bidder selected for acceptance is within 15 (fifteen) percent less than the approved cost estimate, the performance security amount shall be 5 (five) percent of the bid price.
	ii) If the contract price is below 15 percent of the cost estimate, the performance security amount shall be determined using the following formula:
	Performance Security = [(0.85 x Cost Estimate - Contract Price) x 0.5 + Contract Price x 0.05]
	The Performance Security shall be denominated in the currencies of the Contract, or in a freely convertible currency acceptable to the Purchaser.
	and shall be valid for the period of one year from the date of delivery of goods
GCC 17.3	The types of acceptable Performance Securities are: A Bank guarantee issued by "A" class commercial Bank located in Nepal or reputable Bank located abroad, acceptable to the Purchaser, in the format included in Section VIII, Contract Forms, Performance Security issued by foreign Bank must be counter – guaranteed by "A" class commercial Bank in Nepal.
GCC 17.4	Discharge of the Performance Security shall take place: After completion of the Supplier's warranty obligations in accordance with GCC Clause 27.3.



GCC 22.2	A complete packing list indicating the content of each package shall
	be enclosed in a water proof envelope and shall be secured to the
	outside of the packing case. In addition, each package shall be
	marked with indelible ink/paint in bold letters, as follows:
	a. Contract number:
	b. Name and address of the Purchaser:
	c. Country of origin,
	d. Gross weight
	e. Net weight
	f. Package number of total number of packages
	g. Brief description of content
	Upright markings, where appropriate, shall be placed on all four
	vertical sides of the package.
	All materials used for packing shall be environmentally neutral.
GCC 23.1	The insurance coverage shall be in an amount equal to 110 percent of the EXW price of the Goods from "warehouse" to "warehouse" on "All Risks" basis, including War Risks, riots and/or Strikes.
GCC 24.1	Obligations for transportation of the Goods shall be in accordance with:
	The supplier is required under the contract to transport the Goods to a specified place of final destination, defined as the project site. Transport to such place of destination including insurance and storage, as specified in the contract, shall be arranged by the supplier, and related costs shall be included in the contract price.
GCC 25.2	All offered goods thereof must be in compliance with the requirements of Technical Specifications, which will be reviewed by Technical Experts. Supplies not meeting the required Technical Specifications will be rejeted, which has to be replaced with a new quality product aceptable to the purchaser within 30 days.
GCC 26.1	The applicable rate of liquidated damages shall be: 0.05 percent of the Contract Price per day.
GCC 26.1	The maximum amount of liquidated damages shall be: Ten (10) percent of the Contract Price.
GCC 27.3	The period of validity of the Warranty shall be: 12 month after delivery and acceptance of goods at the final destination.
	For the purposes of the Warranty, the place of final destination shall be: Department of Health Services, Logistics Management Division, Teku.
GCC 27.5	The Supplier shall correct any defects covered by the Warranty within: 15 days of being notified by the Purchaser of the occurrence of such defects



Section VIII. Contract Forms

Table of Forms

Letter of Intent	125
Letter of Acceptance	126
Agreement Form	127
Performance Security	128
Advance Payment Security	129



Letter of Intent

	date
То:	name and address of the Contractor
Subject:	. Issuance of letter of intent to award the contract
datedfor identification number, amount in fig modified in accorda	u that, it is our intention to award the contract execution of the
	Authorized Signature:
	Name:
	Title

CC:



Letter of Acceptance

date
To: name and address of the Contractor
Subject: Notification of Award
This is to notify that your Bid dated date for execution of the
You are hereby instructed to contract this office to sign the formal contract agreement within 15 days. As per the Conditions of Contract, you are also required to submit Performance Security, as specified in SCC, consisting of a Bank Guarantee in the format included in Section VIII (Contract Forms) of the Bidding Document.
The Employer shall forfeit the bid security, in case you fail to furnish the Performance Security and to sign the contract within specified period.
Authorized Signature:
Name and Title of Signatory:



Agreement Form

THIS AGREEMENT made on the [insert number] day of [insert month], [insert year], between [insert complete name of Purchaser] of [insert complete address of Purchaser] (hereinafter "the Purchaser"), of the one part, and [insert complete name of Supplier] of [insert complete address of Supplier] (hereinafter "the Supplier"), of the other part:

WHEREAS the Purchaser invited Bids for certain Goods and Related Services, viz., [insert brief description of the Goods and Related Services] and has accepted a Bid by the Supplier for the supply of those Goods and Related Services in the sum of NRs[insert amount of contract price in words and figures including taxes] (hereinafter "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

- 1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
- 2. The following documents shall be deemed to form and be read and construed as part of this Agreement, viz.:
 - (a) the Purchaser's Notification to the Supplier of Award of Contract;
 - (b) the Bid Submission Form and the Price Schedules submitted by the Supplier;
 - (c) the Special Conditions of Contract;
 - (d) the General Conditions of Contract;
 - (e) the Schedule of Requirements; and
 - (f) Technical Specification of Contract

This Contract shall prevail over all other Contract documents. In the event of any discrepancy or inconsistency within the Contract documents, then the documents shall prevail in the order listed above.

- 3. In consideration of the payments to be made by the Purchaser to the Supplier as indicated in this Agreement, the Supplier hereby covenants with the Purchaser to provide the Goods and Related Services and to remedy defects therein in conformity in all respects with the provisions of the Contract.
- 4. The Purchaser hereby covenants to pay the Supplier in consideration of the provision of the Goods and Related Services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the Contract at the times and in the manner prescribed by the Contract.

IN WITNESS whereof the parties hereto have caused this Agreement to be executed in accordance with the laws of "Nepal" on the day, month, and year indicated above.

Signed by [insert authorized signature for the Purchaser] (for the Purchaser)

Signed by [insert authorized signature for the Supplier] (for the Supplier)



Performance Security

[insert complete name and number of Contract]

To: [insert complete name of Purchaser]

WHEREAS [insert complete name of Supplier] (hereinafter "the Supplier") has received the notification of award for the execution of [insert identification number and name of contract](hereinafter "the Contract").

AND WHEREAS it has been stipulated by you in the aforementioned Contract that the Supplier shall furnish you with a security [insert type of security] issued by a reputable guarantor for the sum specified therein as security for compliance with the Supplier's performance obligations in accordance with the Contract.

AND WHEREAS the undersigned [insert complete name of Guarantor], legally domiciled in [insert complete address of Guarantor], (hereinafter the "Guarantor"), have agreed to give the Supplier a security:

THEREFORE WE hereby affirm that we are Guarantors and responsible to you, on behalf of the Supplier, up to a total of [insert currency and amount of guarantee in words and figures] and we undertake to pay you, upon your first written demand declaring the Supplier to be in default under the Contract, without cavil or argument, any sum or sums within the limits of [insert currency and amount of guarantee in words and figures] as aforesaid, without your needing to prove or to show grounds or reasons for your demand or the sum specified therein.

This security is valid until the [insert day, month, year].

Name: [insert complete name of person signing the Security]

In the capacity of: [insert legal capacity of person signing the Security]

Signed: [insert signature of person whose name and capacity are shown above]

Duly authorized to sign the security for and on behalf of: [insert seal and complete name of Guarantor]

Date: [insert date of signing]



Advance Payment Security - Not Applicable

[insert complete name and number of Contract]

To: [insert complete name of Purchaser]

In accordance with the payment provision included in the Contract, in relation to advance payments, [insert complete name of Supplier] (hereinafter called "the Supplier") shall deposit with the Purchaser a security consisting of [indicate type of security], to guarantee its proper and faithful performance of the obligations imposed by said Clause of the Contract, in the amount of [insert currency and amount of guarantee in words and figures].

We, the undersigned [insert complete name of Guarantor], legally domiciled in [insert full address of Guarantor] (hereinafter "the Guarantor"), as instructed by the Supplier, agree unconditionally and irrevocably to guarantee as primary obligator and not as surety merely, the payment to the Purchaser on its first demand without whatsoever right of objection on our part and without its first claim to the Supplier, in the amount not exceeding [insert currency and amount of guarantee in words and figures].

This security shall remain valid and in full effect from the date of the advance payment being received by the Supplier under the Contract until [(insert day, month, year) Contract completion date may be a basis for this date].

Name: [insert complete name of person signing the Security]

In the capacity of: [insert legal capacity of person signing the Security]

Signed: [insert signature of person whose name and capacity are shown above]

Duly authorized to sign the security for and on behalf of: [insert seal and complete name of Guarantor]

Date: [insert date of signing]

